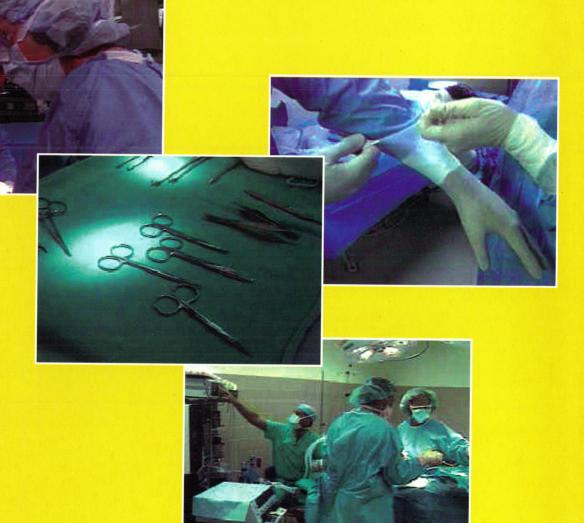
The Report of the National Confidential Enquiry into Perioperative Deaths

1996 / 1997

Compiled by

• A J G GRAY • R W HOILE • G S INGRAM • K M SHERRY





THE REPORT OF THE NATIONAL CONFIDENTIAL ENQUIRY

INTO

PERIOPERATIVE DEATHS

1996/1997

(1 April 1996 to 31 March 1997)

Compiled by

A J G Gray R W Hoile G S Ingram K M Sherry

MB BChir FRCA MS FRCS MBBS FRCA MBBS FRCA

Contents

Forewor	d	7
Acknow	ledgements	8
Introduc	etion	9
Clinical	Contributors	10
Recomm	nendations	13
Data col	lection and review	14
Gynaeco	ology	18
Head an	d neck surgery	25
	The obstructed airway in head and neck surgery	27
	Pharyngeal pouch surgery	32
	Anaesthesia for carotid endarterectomy	33
Minimal	ly invasive surgery (excluding gynaecology and urology)	38
	Minimally invasive abdominal surgery may not be minimally stressful to the patient	40
	Percutaneous Endoscopic Gastrostomy (PEG)	43
	Laparoscopic general surgery	46
Oesopha	geal surgery	50
	Management of patients undergoing oesophagectomy	57
	Preoperative assessment of oesophageal malignancy and preparation for oesophageal surgery	62
	Proposed facilities for the specialist practice of oesophageal surgery	65
	Oesophageal stenting	68
Spinal sı	urgery	70
	The management of metastatic spinal tumours	73
Urology		75

(continued overleaf)

Contents 5

Miscellaneous

	Central venous pressure monitoring during anaesthesia and surgery			
	Admission on the day of surgery			
	Complications associated with postoperative analgesia			
	Non-steroidal anti-inflammatory drugs			
	Tra	tinee anaesthetists supervising sedation for sick patients	90	
Patholog	\mathbf{y}		92	
Referenc	es		105	
Appendi	ces			
	A	Glossary	109	
	В	Abbreviations	111	
	C	NCEPOD corporate structure	113	
	D	Protocol	115	
	E	General data 1996/97	118	
	F	Anaesthetic questionnaire	124	
	G	Surgical questionnaire	137	
	Н	Pathology proforma	149	
	J	Participants (anaesthetists)	151	
	K	Participants (surgeons and gynaecologists)	154	
	Τ.	Local reporters	158	

The analysis of data from completed anaesthetic and surgical questionnaires is not included in full in this report. A supplement containing detailed tables is available free of charge from the NCEPOD office:

NCEPOD, 35-43 Lincoln's Inn Fields, London, WC2A 3PN

Telephone 0171 831 6430 Fax 0171 430 2958 E-mail info@ncepod.org.uk

Foreword

The information for this report, which covers the year from 1 April 1996 up to and including 31 March 1997, was analysed in detail by the clinical coordinators and advisors during the following year. The presentation of all the information differs from previous reports in that it concentrates on an overview and discussions of frequent or recurring clinical issues. It was approved by the Steering Group in July 1998.

The total number of deaths reported to NCEPOD for this study was 19496. Just over 47% of deaths occurred on the day of operation or within the next five days after the operation. Out of the 19496 deaths reported, 7097 (36%) of the patients who died were aged 80 or over. The number of cases for the detailed review was 2541 (13% of the total). These were subdivided into six groups of which four (head and neck surgery, minimally invasive surgery, oesophageal surgery and spinal surgery) involved consultants from different specialties, and two single subject disciplines (gynaecology and urology). The overall return rate of return of questionnaires on a purely voluntary basis by surgeons was 71% and by anaesthetists 76%. However, the rate of return by gynaecologists was 80% and by their associated anaesthetists 86%, by urologists 78% and their associated anaesthetists 80%. In each of the six groups of operations reviewed it will be seen that there was usually a high degree (over 90%) of consultant involvement in the decision-making and they were also often involved in the operations. Only 728 (43%) of these deaths were considered by surgeons at an audit meeting, and only 345 (27%) by the anaesthetists. These figures unfortunately do not take into account the fact that many of the reasons for, or the factors in, these deaths are similar and repetitious as will, I hope, immediately become clear on reading this report.

During the two-and-a-half year cycle of collection and review of the questionnaires and the publication of this report there have been many changes, or proposed changes, to medical practice. There is more emphasis on audit, openness and communication between patient, hospital and doctor, greater expectation of a successful outcome, longer survival with an ageing population. It is generally acknowledged that NCEPOD over the past decade has proved its worth and improved the standards of care by highlighting some of the deficiencies in the facilities, practice or provision of anaesthetic and surgical care. This is despite the fact that everyone involved in the preparation of the NCEPOD reports is acutely aware that the basic denominator data (numbers of patients and operations) are not available nationally.

Finally I would like to thank all those who have contributed to this report, from the local reporters to those members of the permanent staff at NCEPOD who work tirelessly behind the scenes to provide the data for analysis and are so ably led by our Chief Executive, Anne Campling.

V R Tindall CBE FRCS FRCOG Chairman, National Confidential Enquiry into Perioperative Deaths

Foreword 7

Acknowledgements

This is the eighth report of National CEPOD and we continue to be indebted to the local reporters and administrative and clinical staff who provide the initial data on perioperative deaths.

Many consultant and trainee anaesthetists and surgeons have, once again, voluntarily contributed to the Enquiry by completing detailed questionnaires.

The Steering Group members wish to record here their appreciation of the efforts of the NCEPOD staff: Peter Allison, Anne Campling, Fatima Chowdhury, Paul Coote, Jennifer Drummond, Sean Gallimore and Dolores Jarman. All data analysis was carried out by Sean Gallimore.

Introduction

The National Confidential Enquiry into Perioperative Deaths (NCEPOD) is based on case collection and peer review. The protocol under which the Enquiry functions charges it with identifying remediable factors in the practice of anaesthesia and surgery. This has resulted in previous years in recommendations on organisational issues: the provision of theatre time for urgent and emergency cases, the availability of HDU and ICU beds and the experience and seniority of staff providing out-of-hours cover; these have all been recurring themes. In this report we seek to turn attention more closely to the clinical aspects of the cases reviewed.

By using death as the starting point on which to base an enquiry, it is inevitable that the population studied will primarily be old, sick and more often than not will suffer from conditions that offer little prospect of recovery. But for those operating and providing anaesthesia and perioperative care, they represent a formidable challenge. Although the individual patient's outlook may not be good, in skilled and experienced hands, for each patient who dies many more will survive and some will do well. In looking only at those patients who died we are directing our attention to one end of a spectrum. The NCEPOD coordinators and the advisors have tried to keep this in mind during the preparation of this report.

Surgical and anaesthetic practice is constantly evolving. Improvement comes through the application of scientific method in research based studies, through the introduction of new drugs and techniques and through advances in technology. But there is delay as these advances permeate into everyday clinical practice. This may to an extent be no bad thing, a certain scepticism is probably sensible but too much conservatism can result in the perpetuation of outmoded and unsatisfactory practice. The questionnaires and associated information returned to NCEPOD contain a great deal of clinical information. During discussion, aspects of clinical care become the subject of recurring debate amongst the advisors. The sample on which this report is based related to particular types of surgery and, not surprisingly, specific issues arose in relation to these. Examples are airway problems in the head and neck surgery cases and the surgical management of metastatic secondaries in spinal surgery. As discussion developed around these problems it was decided to focus attention on them and to use the expertise available amongst the advisors and coordinators to write commentaries about them. In addition there were other issues which arose more broadly across anaesthesia and surgery but which it was felt would benefit from a similar approach; for instance the use of NSAIDs for postoperative pain relief.

The commentaries in this report have therefore been prepared following this broad debate by the anaesthetic and surgical advisors. Whereas previously the disciplines have tended to work largely independently in the preparation of NCEPOD Reports, on this occasion every opportunity has been taken to bring together the anaesthetic and surgical advisors with an interest in a particular area to develop a common view. The resulting commentary may then either have been written by a subgroup of coordinators and advisors, sometimes combining anaesthetists and surgeons, or sometimes from a single discipline and in some cases just a single individual, the criterion in each case being a specific interest in that particular field. The purpose has been to present basic details of the key clinical cases involved in order to give an overall view of aspects of their management. These are opinions, commentaries based on facts as recorded, written by practising clinicians with which readers may take issue and indeed we would be disappointed if they did not. The reality of day-to-day medicine, whatever the hindsight of medico-legal opinion may believe, is not black and white but complex with far more subtle shades.

In the past NCEPOD has sometimes been seen, particularly in the use of vignettes describing specific cases, as seeking to criticise individual unidentified clinicians. In this report details of cases are given because they relate to the clinical issue being discussed, not because of the quality of care given. Critical examination of current clinical practice can teach us much. It is hoped that this form of qualitative review can help to identify where we need to re-evaluate our clinical practice, replacing long established custom with a fresh approach. If in reading this report clinicians ask themselves, "what would I do in a similar situation?" or "what is my view on this issue?" and if it stimulates debate at local meetings; then NCEPOD will have succeeded in its intended purpose.

Ron Hoile and Stuart Ingram, Principal Clinical Coordinators

Clinical Contributors

NCEPOD Coordinators

A J G Gray Clinical Coordinator, NCEPOD and Consultant Anaesthetist, Norfolk and Norwich

Health Care

R W Hoile Principal Clinical Coordinator, NCEPOD and Consultant General Surgeon, Medway

G S Ingram Principal Clinical Coordinator, NCEPOD and Consultant Anaesthetist, University

College London Hospitals

K M Sherry Clinical Coordinator, NCEPOD and Consultant Anaesthetist, Northern General Hospital

V R Tindall Chairman, NCEPOD and Emeritus Professor of Gynaecology and Obstetrics, University

of Manchester

Specialty advisors

Anaesthesia

I Baguley Consultant Anaesthetist, Royal Shrewsbury Hospitals*

I Conacher Newcastle upon Tyne Hospitals A E Cooper Rotherham General Hospitals

M Davies Royal Liverpool and Broadgreen University Hospitals

G L Greenslade Frenchay Healthcare

N D Groves Llandough Hospital and Community

R M Haden Alexandra Healthcare

R Kishen Consultant, Anaesthesia and Intensive Care, Salford Royal Hospitals*

S J Power Greenwich Healthcare

N Schofield The Oxford Radcliffe Hospital

V Sidhu St Mary's Hospital
I E Tweedie Aintree Hospitals
J R Wace Portsmouth Hospitals

^{*} author of personal commentary

Gynaecology

D D Boyle Consultant Obstetrician and Gynaecologist, The Royal Group of Hospitals and Dental

Hospital Health and Social Services

L Cardozo Professor of Urogynaecology, King's Healthcare

S G Crocker Consultant Obstetrician and Gynaecologist, Norfolk and Norwich Health Care

R Garry Consultant Gynaecologist, South Tees Acute Hospitals

J Lane Consultant Gynaecologist and Obstetrician, Royal Shrewsbury Hospitals
J H Shepherd Consultant Gynaecological Surgeon and Oncologist, The Royal Hospitals

J Tidy Senior Lecturer in Gynaecology, Northern General Hospital

Head and neck surgery

M Beck Consultant Ophthalmologist, University Hospital of Wales Healthcare A M S Brown Consultant Maxillofacial Surgeon, University Hospital Birmingham

M J K M Brown Consultant ENT Surgeon, Glan Hafren

K B Hughes Consultant ENT Surgeon, The Doncaster Royal & Montagu Hospital

D Jonathan Consultant ENT Surgeon, Frimley Park Hospital

N R McLean Consultant Head and Neck/Plastic and Reconstructive Surgeon, Newcastle upon Tyne

Hospitals

A Resouly Consultant ENT Surgeon, Portsmouth Hospitals*

A P Walby Consultant ENT Surgeon, The Royal Group of Hospitals and Dental Health and Social

Services

M Wickstead Consultant ENT Surgeon, Norfolk and Norwich Health Care

Minimally invasive surgery

W A Brough Consultant General Surgeon, Stockport Acute Services

W Fountain Consultant Thoracic Surgeon, Royal Brompton & Harefield Hospitals

B I Rees Consultant General Surgeon, University of Wales Healthcare*

R D Rosin Consultant Surgeon in General and Oncological Medicine, St Mary's Hospital*

R E Sayer Consultant Thoracic Surgeon, St George's Healthcare

Oesophageal surgery

D Alderson Professor of Gastrointestinal Surgery, United Bristol Healthcare

A K Deiraniya Consultant Cardiothoracic Surgeon, South Manchester University Hospitals*

S M Griffin Consultant General Surgeon, Newcastle upon Tyne Hospitals*

S Raimes Consultant Surgeon and Gastroenterologist, Carlisle Hospitals*

R Vaughan Consultant Thoracic Surgeon, Norfolk and Norwich Health Care

Spinal surgery

A-T Cross Consultant Orthopaedic Surgeon, City Hospitals Sunderland

G F G Findlay Consultant Neurosurgeon, The Walton Centre for Neurology and Neurosurgery

R W Gullan Consultant Neurosurgeon, King's Healthcare

T R Morley Consultant Orthopaedic Surgeon, Royal National Orthopaedic Hospital*

(continued overleaf)

* author of personal commentary

Clinical Contributors

Specialty advisors (continued)

Urology

J G W Feggetter Consultant Urologist, Northumbria Healthcare

M Fordham Consultant Urologist, The Royal Liverpool and Broadgreen University Hospitals

A D Joyce Consultant Urologist, United Leeds Teaching Hospitals P O'Boyle Consultant Urologist, Taunton & Somerset

P O'Boyle Consultant Urologist, Taunton & Somerset K K Sethia Consultant Urologist, Norfolk and Norwich Health Care

Pathology

P N Cooper Senior Lecturer in Forensic Pathology, Newcastle upon Tyne Hospitals

C M Corbishley Consultant/Senior Lecturer in Histopathology, St George's Healthcare

N Francis Consultant Histopathologist, The Hammersmith Hospitals
J D Hemming Consultant Histopathologist, Newcastle upon Tyne Hospitals

S Rogers Consultant Histopathologist, The Doncaster Royal & Montagu Hospital

R D Start Consultant Histopathologist, Chesterfield and North Derbyshire Royal Hospital

Recommendations

General comment:

It is a surgical skill to recognise when surgery will be too adventurous, ill advised or futile, given the condition of the patient. It is difficult to resist pressure to operate, whether this comes from the patient, relatives or medical colleagues but it must be recognised that surgery cannot solve every problem (page 38).

- A fibreoptic intubating laryngoscope should be readily available for use in all surgical hospitals. Several anaesthetists working in a department should be trained for, and competent at, awake fibreoptic intubations (page 27).
- The maintenance of an adequate blood pressure through the operative and postoperative period is an essential part of *anaesthesia* for patients undergoing carotid endarterectomy surgery. This requires invasive blood pressure monitoring and particular care in patients with poor cardiovascular reserve (page 33).
- Laparoscopic abdominal surgery may take place through a small incision but it still requires *anaesthesia* and the physiological onslaught of a pneumoperitoneum. High risk patients may not be able to tolerate this stress (page 40).
- Morbidity/mortality meetings should take place in all anaesthetic departments. Regular review of
 mortality following operations is an essential part of anaesthetic practice (page 123).
- There are many aspects around the care of patients undergoing *anaesthesia and surgery* for oesophageal disease, which are of major concern. A fundamental re-examination of the arrangements for the care of these patients is urgently required (pages 50, 57 and 65).
- The technique of tracheostomy should be taught to trainee *surgeons*. The indications for performing this procedure under local or general anaesthesia should also be taught (page 30).
- Pharyngeal pouch is a benign condition but appears to have a significant mortality. Surgical subspecialisation for this condition within otolaryngology departments is required (page 32).
- More detailed preoperative investigation and assessment may prevent radical spinal *surgery*, which is unhelpful for individual patients with advanced malignant disease (page 71).
- Surgeons need to be clear about the aims of treatment and benefits for the patient when planning surgery for advanced malignancy (pages 23, 73 and 76).
- Patients and their relatives need to recognise the limits of *surgery* in advanced malignant disease. A decision to operate may not be in the best interests of the patient (page 38).
- The hospital postmortem rate of 8% was unacceptably low. The reasons for this low rate need to be examined (page 95).

These recommendations have been selected from the report as it is felt that they should receive early implementation. Although the page numbers may relate to a personal commentary, in all cases these recommendations originated from strong views expressed by the groups of advisors. They are therefore based on the opinions of clinicians practising in particular surgical and anaesthetic specialties. As with all NCEPOD recommendations, in the absence of denominator data, they have not been statistically validated.

Data collection and review

The National Confidential Enquiry into Perioperative Deaths (NCEPOD) reviews clinical practice and aims to identify remediable factors in the practice of anaesthesia, all types of surgery and, from 1 April 1998, other invasive procedures. We consider the *quality* of the delivery of care and not specifically causation of death. Data are supplied on a voluntary basis; consultant clinicians in the relevant specialties are invited to participate. We also review requests for and reporting of postmortem examinations. The commentary in the reports is based on peer review of the data, questionnaires and notes submitted to us: it is not a research study based on differences against a control population, and does not attempt to produce any kind of comparison between clinicians or hospitals.

Scope

All National Health Service and Defence Medical Services hospitals in England, Wales and Northern Ireland, and public hospitals in Guernsey, Jersey and the Isle of Man are included in the Enquiry, as well as hospitals managed by BUPA Hospitals Limited, General Healthcare Group PLC (until April 1998), Nuffield Hospitals, St Martins Hospitals Limited, Benenden Hospital and The London Clinic.

Reporting of deaths

NCEPOD collects basic details on all deaths in hospital within 30 days of a surgical procedure, through a system of local reporting (see Appendix L). The local reporters in each hospital are predominantly consultant clinicians, but this role is increasingly being taken on by information and clinical audit departments who are often able to provide the data from hospital information systems. In the independent sector, hospital or nursing managers provide the data. When incomplete information is received, the NCEPOD administrative staff contact the appropriate medical records or information officer, secretarial or clinical audit staff.

Deaths of patients *in hospital* within 30 days of a surgical procedure (excluding maternal deaths) are included. If local reporters are aware of postoperative deaths at home, they report them to us. A surgical procedure is defined by NCEPOD as;

"any procedure carried out by a surgeon or gynaecologist, with or without an anaesthetist, involving local, regional or general anaesthesia or sedation".

Reporters provide the following information:

Name of authority/trust

Sex/hospital number/NHS number of patient

Name of hospital in which the death occurred (and hospital where surgery took place, if different)

Dates of birth, final operation and death

Surgical procedure performed

Name of consultant surgeon

Name of anaesthetist

Since April 1998, NCEPOD has also begun to review data relating to interventional radiological procedures.

Sample for more detailed review

The data collection year runs from 1 April to 31 March. Each year, a sample of the reported deaths is reviewed in more detail. The sample selection varies for each data collection year, and is determined by the NCEPOD Steering Group (see Appendix C).

NCEPOD may collect data about patients who have survived more than 30 days after a procedure. These data are used for comparison with the data about deaths, or to review a specific aspect of clinical practice. Data from other sources may also be used.

The perioperative deaths which fell within the sample group for 1996/97 were those where the patient's final operation before death fell within the group of surgical procedures listed below:

Gynaecological surgery
Head and neck surgery
Minimally invasive surgery
Oesophageal surgery
Spinal surgery
Urological surgery

(For more detailed definitions of these groups see Appendix A).

For each sample case, questionnaires (see Appendices F and G) were sent to the consultant surgeon or gynaecologist and consultant anaesthetist. These questionnaires were identified only by a number, allocated in the NCEPOD office. Copies of operation notes, anaesthetic records and fluid balance charts and postmortem reports were also requested. Surgical questionnaires were sent directly to the consultant surgeon or gynaecologist under whose care the patient was at the time of the final operation before death. When the local reporter had been able to identify the relevant consultant anaesthetist, the anaesthetic questionnaire was sent directly to him or her. However, in many cases this was not possible, and the local tutor of the Royal College of Anaesthetists was asked to name a consultant to whom the questionnaire should be sent.

Consultants

We hold a database, regularly updated, of all consultant anaesthetists, gynaecologists and surgeons in England, Wales and Northern Ireland.

Analysis and review of data

The NCEPOD administrative staff manage the collection, recording and analysis of data. The data are aggregated to produce the tables and information in the reports; further unpublished aggregated data may be available from the NCEPOD offices on request. All data are aggregated to regional or national level only so that individual trusts and hospitals cannot be identified.

Advisory groups

The NCEPOD clinical coordinators (see Appendix C), together with the advisory groups for anaesthesia and surgery, review the completed questionnaires and the aggregated data. The members of the advisory groups are drawn from hospitals in England, Wales and Northern Ireland. The advisory group in pathology reviews postmortem data from the surgical questionnaires as well as copies of postmortem reports.

Production of the report

The advisory groups comment on the overall quality of care within their specialty and on any particular issues or individual cases which merit attention. These comments form the basis for the published report, which is prepared by the coordinators, with contributions from the advisors. The report is reviewed and agreed by the NCEPOD Steering Group prior to publication.

Confidentiality

NCEPOD is registered with the Data Protection Registrar and abides by the Data Protection Principles. All reporting forms, questionnaires and other paper records relating to the sample are shredded once an individual report is ready for publication. Similarly, all patient-related data are removed from the computer database.

Before review of questionnaires by the clinical coordinators or any of the advisors, all identification is removed from the questionnaires and accompanying papers. The source of the information is not revealed to any of the coordinators or advisors.

Contents (pages 18 to 105)

Gynaeco	logy	18
Head an	d neck surgery	25
	The obstructed airway in head and neck surgery	27
	Pharyngeal pouch surgery	32
	Anaesthesia for carotid endarterectomy	33
Minimal	ly invasive surgery (excluding gynaecology and urology)	38
	Minimally invasive abdominal surgery may not be minimally stressful to the patient	40
	Percutaneous Endoscopic Gastrostomy (PEG)	43
	Laparoscopic general surgery	46
Oesopha	geal surgery	50
	Management of patients undergoing oesophagectomy	57
	Preoperative assessment of oesophageal malignancy and preparation for oesophageal surgery	62
	Proposed facilities for the specialist practice of oesophageal surgery	65
	Oesophageal stenting	68
Spinal su	ırgery	70
	The management of metastatic spinal tumours	73
Urology		75
Miscellar	1eous	
	Central venous pressure monitoring during anaesthesia and surgery	79
	Admission on the day of surgery	82
	Complications associated with postoperative analgesia	85
	Non-steroidal anti-inflammatory drugs	87
	Trainee anaesthetists supervising sedation for sick patients	90
Patholog	y	92
Referenc	es	105

Gynaecology

Key messages

- There was a high involvement of consultants in the decision-making and operating.
- Surgery for benign disease can have a poor outcome if the age, physical status of the patient and the type of hysterectomy are not taken into account (see page 20).
- Infection was a significant factor in more deaths than expected.
- There were examples of inadequate investigation, which could have avoided unnecessary or unhelpful surgery particularly in cases of pelvic or abdominal masses.
- Guidelines from the Royal College of Obstetricians & Gynaecologists on thromboembolic prophylaxis are not always followed.
- Some questionnaires were poorly completed. The confidential nature of NCEPOD means that when key information is omitted from the forms further information cannot be requested, thus weakening any conclusions.

Written by Mr Ron Hoile MS, FRCS and Emeritus Professor Vic Tindall FRCOG in consultation with the gynaecology advisors (see page 11)

List of procedures

Procedure (may be multiple for individual patients)		
Benign		
Total abdominal hysterectomy and bilateral salpingo-oophorectomy	13	
Vaginal hysterectomy (+/- vaginal repair)	13	
Unilateral salpingo-oophorectomy	5	
Laparotomy for postoperative haemorrhage	4	
Diagnostic laparotomy	4	
Colposuspension	2	
Vaginal repair	2	
Laparotomy and drainage of abscess	2	
Bilateral salpingo-oophorectomy	2	
EUA	2	
Miscellaneous (one each of: change of shelf pessary, diagnostic laparoscopy, D & C,	17	
hysteroscopy, Manchester repair, partial vaginectomy, excision urethral mucosa, cystoscopy, simple vulvectomy, anterior resection of rectum, division of adhesions, small bowel resection, appendicectomy, sigmoid colectomy, abdominal rectopexy, cholecystectomy, manual evacuation of faeces)		

Gynaecology

Procedure (may be multiple for individual patients)		
Malignant		
Total abdominal hysterectomy, bilateral salpingo-oophorectomy and omentectomy	45	
'Open and shut' laparotomy (+/- biopsies and drainage of ascites)	43	
Salpingo-oophorectomy	26	
EUA	16	
Cystoscopy	12	
Debulking pelvic mass	9	
D & C	8	
Hysteroscopy	7	
Defunctioning colostomy	7	
Laparoscopy	5	
Subtotal hysterectomy	4	
Ileo-colic bypass	3	
Sigmoid colectomy	3	
Local excision vulval lesion	3	
Radical vulvectomy and bilateral inguinal node dissection	2	
Hartmann's procedure	2	
Anterior resection of rectum	2	
Bilateral inguinal node dissection	2	
Sigmoidoscopy	2	
Repair small bowel tears	2	
Wertheim's hysterectomy	2	
Small bowel resection	2 2	
Total colectomy and ileostomy	2	
Resuture abdominal wound dehiscence	2	
Miscellaneous (one each of: right hemicolectomy, defunctioning ileostomy, reimplantation of ureter, removal of packs, removal of shelf pessary, excision of clitoral tumour, total	14	
vaginectomy, simple vulvectomy, ovarian cystectomy, appendicectomy, laparotomy for		
postoperative haemorrhage, insertion of ureteric stent, splenectomy, suturing of leg ulcer)		

Total number of patients = 205 (164 with malignant disease and 41 with benign disease)

Consultant involvement

A consultant gynaecologist was involved, in some way, in the care of 95% (194/205) of the women having a gynaecological procedure. A consultant was the most senior person in the operating room in 92% (188/205) and was the operating surgeon in 87% (179/205) of cases.

Age

Twelve of the women who died were aged less than 50 years. Seven of these women had a malignancy (five cases of ovarian cancer, one of endometrial cancer and the youngest woman aged 26 had secondary ovarian cancer from a primary tumour arising in the stomach). The five benign conditions being treated were respectively related to endometriosis, dysfunctional haemorrhage, prolapse, menorrhagia in a Jehovah's Witness and a case of suspected ectopic pregnancy in a woman who arrived at hospital in a moribund state, but at laparotomy was found to have overwhelming sepsis.

A total of 131 women who died were aged 70 years and over, of whom 78% (102/131) had malignant disease. Two of the women who died following operations for benign gynaecological disease had a previous history of successful treatment of breast cancer. Sixty (29%) women were aged 80 years and over.

Hysterectomy and benign disease

There were seven deaths from haemorrhage and/or infection in women who had a hysterectomy for benign disease. Five of these deaths occurred after vaginal hysterectomy and two after abdominal hysterectomy. Currently more than 70% of hysterectomies in the UK are performed abdominally. This equates to an approximate death rate of four per 100,000 following abdominal hysterectomy and 25 per 100,000 following vaginal hysterectomy. Although discrepancies in the mean ages of the two groups may contribute to the observed differences, these relative risks should be considered when selecting the type of hysterectomy operation to be performed in patients with benign disease.

Malignant disease

All cases of ovarian malignancy, whether primary or secondary, were classified as having gynaecological malignancy. Only 68% (112/164) of the patients with gynaecological malignancy were admitted initially to a gynaecological ward compared with 76% (31/41) of the women with benign disease. Just over 50% of the women with benign and malignant disease had two or more co-existing problems present.

Inadequate information

It was observed that the quality of information provided by gynaecologists to the NCEPOD study was poor in comparison to that which they supply as obstetricians to the Confidential Enquiries into Maternal Deaths. This often applied to lack of histological confirmation of the diagnosis and the failure to obtain a postmortem examination when the diagnosis was uncertain. The advisors also commented on the failure nationally to obtain the number of individual operations done.

ASA classification

This system of classification for physical status is not well known by gynaecologists and many cases classified as ASA 1 should have been graded as 2 or 3 i.e. more at risk.

Death was apparently not anticipated by the gynaecological surgeons in 31% (64/205) of all cases. The figures for benign disease were 51% (21/41) compared with 26% (43/164) of the women with malignant disease. Whilst the disease process in benign gynaecological conditions may not be life-threatening, the outcome will be poor if the age, comorbidities present and overall physical status of the patient is not taken into account. Do gynaecologists need to assess their patients, particularly those with benign disease, more thoroughly?

Thromboembolism

Despite the existence of Royal College of Obstetricians and Gynaecologists Guidelines,² only 33% (54/164) of women with malignant disease undergoing major surgery received anticoagulant therapy compared with 56% (23/41) of women with benign disease. Overall, 48% (98/205) received anticoagulant therapy despite 94% (193/205) of the women being aged 50 years and over, 84% (172/205) aged 60 years and over, and 64% (131/205) aged 70 years and over with a high incidence of malignant disease in each age group. There were four cases of confirmed pulmonary embolism, two of whom died despite all the appropriate prophylactic measures, although in both these cases the postmortem examination revealed evidence of antemortem thrombosis. In one case, anticoagulant therapy was inadvertently omitted and in the fourth case there was inadequate information in the questionnaire to allow evaluation.

Gynaecology 20

On the information available, anticoagulation in gynaecological surgery, whether using conventional unfractionated heparin or low molecular weight heparin, as prophylaxis against thromboembolism in malignant disease associated with major surgery (i.e. where there are additional risk factors) does not appear to be as clinically effective as in other specialties.³ It is nevertheless important that mechanical antithromboembolic measures (physiotherapy, leg pumps, TED stockings etc.) should be used in all cases and in combination with anticoagulation in some patients.

There were three sudden deaths in which, due to lack of a postmortem examination, it was impossible to determine whether the death was the result of a pulmonary embolism or a myocardial infarction.

Haemorrhage

There were seven deaths primarily due to haemorrhage. Three of the cases were associated with malignant disease, and four deaths were associated with benign disease (one woman a Jehovah's witness). The operations for the benign conditions were two vaginal hysterectomies and repair operations, a subtotal hysterectomy for menorrhagia on a Jehovah's Witness and the final one a laparotomy for secondary bleeding in the retropubic space shortly after a laparoscopic colposuspension operation. The advisors regarded the deaths of Jehovah's Witness patients as medically unavoidable given the wishes of the patients and their relatives.

Infection

There were seven cases in which infection was a major factor in the eventual outcome.

Age	ASA	Procedure	Comment
39	5	Laparotomy	Three-day illness referred in collapsed state. Urgent laparotomy – possible ectopic pregnancy. Rapid collapse and death. Actual diagnosis – pelvic cellulitis and streptococcal A septicaemia.
42	3	Vaginal hysterectomy and repair	Operation followed by septicaemic shock on fourth day.
47	2	TAH and BSO	Routine operation. Collapsed third day with wound infection. Plan to reoperate but arrested during intubation. Streptococcal septicaemia. No antibiotic prophylaxis.
51	5	Diagnostic laparotomy	Laparotomy for ovarian cyst, admitted to ICU due to bronchial aspiration. Transferred to ward because of shortage of beds. Readmitted to ICU, further laparotomy (inconclusive) followed by septicaemia and paralytic ileus.
55	1	Total vaginectomy and pelvic lymphadenectomy	Previous endometrial cancer with recurrence. Operation complicated by a subphrenic abscess. Death from sepsis.
74	4	Laparotomy for suspected postoperative bleeding	Initially vaginal hysterectomy and repair. Laparotomy because of suspected bleeding. None found. Subsequent septicaemia and death. Was there failure to recognise infection earlier?
76	1	Vaginal hysterectomy, anterior and posterior repair	Routine surgery. Died four days later in ICU with septicaemia. No prophylactic antibiotics given.

There appeared to be a failure in many of these cases to appreciate the potential dangers of infection in postoperative cases. The pelvis is potentially contaminated with urogenital and colorectal organisms, let alone commensals on the skin and perineum. There is therefore every justification for preoperative and intraoperative antibiotic usage in gynaecological surgery. Individual units should draw up protocols based on audited experience and after taking local microbiological advice.

Inadequate preoperative assessment

There were three cases where there were doubts expressed by the advisors about the preoperative assessment or management of these individual cases.

Age	ASA	Procedure	Comment
68	1	BSO and omentectomy	A case of prolapse and a possible fibroid. At operation there was found to be an ovarian cyst and a radical procedure was done. Patient had COPD and a previous MI. An ultrasound examination preoperatively might have prevented the radical surgery for a benign ovarian cyst in a high-risk patient.
82	1	Vaginal hysterectomy and repair of prolapse	A case of apparently longstanding procidentia. A difficult vaginal hysterectomy and repair was done when it was commented that the bladder was very adherent to the uterus. At postmortem examination it was revealed that a diagnosis of carcinoma of the cervix had been missed and carcinomatosis was present.
89	3	TAH and BSO	This patient had carcinoma of the breast and was on tamoxifen. She had persistent and heavy bleeding for which a TAH and BSO were done. As far as could be established there was no consideration of stopping the tamoxifen for a period of observation before opting for surgery.

Was the operation justified?

There were three cases in which the justification for part or all of the operation was doubtful.

Age	ASA	Procedure	Comment
66	3	Laparotomy, relief of obstruction and debulking	A case of advanced ovarian cancer presenting with intestinal obstruction. It is debatable whether relief of obstruction combined with debulking is justified.
69	1	Diagnostic laparoscopy. Large ovarian tumour. Laparotomy & bilateral oophorectomy. Very haemorrhagic procedure.	Admission as day case for laparoscopy for pelvic mass by general surgeon. Trainee surgeon throughout. Was it appropriate to proceed to laparotomy on patient unprepared (and possibly not consented) for major surgery?
89	4	Hysteroscopy. Laparotomy and debulking of tumour	Diagnosis of endometrial cancer had been established. Was anaesthesia and laparotomy for stage 4 cancer justified for a woman in cardiac failure?

Pelvic or abdominal masses

There were 20 cases in which a woman presented with or had a pelvic or abdominal mass. The actual diagnoses were: 11 ovarian cancers, one peritoneal cancer, three bowel cancers, one of whom also had an unrecognised endometrial cancer, two cases of high-grade lymphoma, one grossly over-distended bladder and one case with negative findings at laparotomy.

The advisors felt that some general points should be made as a consequence of analysing the details of these cases.

The management of a pelvic or abdominal mass is a complex challenge and warrants considered preoperative investigations and consultation. This report has identified that over 50% of the women with malignant disease (who subsequently died) had two or more coexisting medical problems at the time of presentation. Consultation between relevant medical specialists, including anaesthetists and cardiopulmonary physicians, is essential. These patients will often require extensive preoperative investigations, so that appropriate risk assessment can be made before any planned surgery. In patients with a suspected pelvic or abdominal mass a full examination of the patient, including vaginal and rectal examination, is invaluable. Investigations that are helpful include a full haematological profile, CA 125 estimation and an ultrasound scan of the pelvis, particularly if ovarian cancer is suspected. If there is doubt over the origin of the mass or the pathology then a preoperative CT or MRI scan of the pelvis with guided biopsies may be helpful in obtaining a diagnosis. There is little justification in performing surgery on women with disseminated abdominal tumours. Prior to surgery there are a number of determinants (symptoms/pain/haemorrhage /quality of life/risk of the surgery/long term outcome) which should influence the options available. Consultation with the woman and relatives is helpful in deciding the appropriate management policy. This report has identified that at the time of surgery for ovarian cancer there were examples of inappropriate attempts to debulk the disease, which would still result in suboptimal treatment. Having diagnosed, assessed and staged the disease at laparotomy it is only then appropriate to decide what would constitute suitable debulking surgery. If the debulking can only be considered to be palliative it is not justifiable to continue with procedures associated with a high morbidity and mortality. In these cases, appropriate chemotherapy, rather than inadequate and futile surgery, will influence the outcome for the patient.

23

Individual departments can construct their own guidelines for the management of ovarian cancer based on national advice.⁴ An agreed and unified approach would improve patient care, teaching, audit and promote better communication both within and without the department. Careful consideration should be given to who carries out any operative procedure and where the patient is managed. It is prudent to either refer to or take advice from a multidisciplinary gynaecological oncology team in a cancer centre. Guidelines for referral and clinical practice will be determined from these multidisciplinary teams. General guidelines for the management of gynaecological malignancies are in preparation by the NHS Executive along the lines of the comments already released for breast, lung and colorectal cancers.

Head and Neck Surgery

Key messages

• In this mixed group of specialist procedures, consultant surgeons were involved in the management of 90% of the cases.

Total

- The management of the obstructed airway gave cause for concern.
- The advisors were surprised by the deaths from pharyngeal pouch surgery.

Written by Mr Ron Hoile MS, FRCS in consultation with the head and neck surgery advisors (see page 11)

These may be multiple for individual patients and many of the miscellaneous procedures,

Head and neck procedures

which may appear inappropriate, were done together with a procedure such as a tracheostomy	number
Tracheostomy	74
Carotid endarterectomy	25
Panendoscopy	23
Unilateral block dissection of neck	15
Excision biopsy cervical node	15
Laryngoscopy and biopsy	12
Flap reconstruction	11
Surgery for pharyngeal pouch	8
Excision skin tumour	7
Tongue biopsy	7
Fixation of mandibular fractures	6
Debridement of necrotic flaps	6
Exploration of cervical mass	5
Incision and drainage of abscess (all sites)	5
Nasal packing for epistaxis	5
Biopsy of nasopharynx	5
Tonsillectomy/biopsy for malignancy	4
Dilatation of oesophageal stricture	4
Biopsy mouth	4
Surgery for carotid blow-out	4
Debulking malignant thyroid mass	3
Percutaneous tracheostomy	3
Exploration of neck for haemorrhage (traumatic or postoperative)	3
Pharyngo-laryngo-oesophagectomy	3
Teflon injection of vocal cord	3
Total glossectomy	3
Parotidectomy	3
Exploration of ear	2
Percutaneous endoscopic gastrostomy (PEG)	2
Parathyroidectomy	2
Rigid oesophagoscopy	2
Laryngectomy	2
Partial thyroidectomy	2
Total thyroidectomy	2
Elevation of depressed skull fracture	2
Dental extraction/clearance	2
Excision of ethmoid tumour	2
(continued overleaf)	

Local excision oropharyngeal tumour	2
Revision of tracheostomy	2
Miscellaneous (one each of: closure of pharyngeal fistula, ligation of maxillary artery for epistaxis,	26
replacement of tracheal stent, craniectomy, mandibulectomy, biopsy of clavicle, bronchoscopy,	
cricothyroidotomy, resuture of anastomosis, FNAC, mandibular biopsy, myringotomy, MUA,	
hemiglossectomy, arytenoidectomy, open gastrostomy, tympanoplasty & mastoidectomy, excision	
submandibular gland, evacuation of haematoma, repair of severe neck lacerations, resuture	
abdominal dehiscence, repair AAA, excision sebaceous cyst, split skin graft, biopsy thyroid, repair	
recurrent inguinal hernia)	

Ophthalmic procedures

Cataract extraction and intra-ocular lens implantation	20
Cataract extraction	6
Repair of ruptured globe	3
Trabeculectomy	2
Miscellaneous (one each of: bilateral cataract extraction & intraocular lens	9
implantation, ptosis surgery, removal of suture, tracheostomy, canthoplasty,	
capsulotomy, tarsorrhaphy, enucleation, vitrectomy)	

Total number of patients 315 (276 head and neck, 39 ophthalmic)

General comments

This group of patients and procedures is heterogeneous and it is difficult to draw general conclusions. There are some broad observations to be made and areas of concern.

The specialties involved in head and neck surgery were otorhinolaryngology, ophthalmic surgery, neurosurgery, general, vascular, maxillofacial, plastic, cardiothoracic and orthopaedic surgery. The ages of the patients varied between specialties but were noticeably lower in neurosurgery (all seven patients were under 60 years of age) and higher in ophthalmic surgery (64%, 25/39 being over 80 years of age). For the group as a whole 62% (195/315) of the patients were over 70 years old.

Review of the overall ASA status showed that 77% (244/315) were ASA class 3 or higher. Interestingly, despite the higher age groups, patients undergoing ophthalmic surgery were in lower ASA classes (as judged by surgeons). There were coexisting medical problems in 87% (273/315) of the patients.

Consultants were directly involved in the care of these patients in 90% (282/315) of cases and were the operating surgeon in 68% (215/315). There were 54 instances where surgery was performed by a non-consultant without the immediate availability of a supervising consultant; these cases were reviewed and only one case was considered to be possibly inappropriate. This was an exploration of the neck, debulking of a thyroid tumour and tracheostomy done by a fourth year specialist registrar.

In ophthalmic surgery, 36% (14/39) procedures were done solely under local anaesthesia (with or without sedation) administered by the surgeon. There were no problems which were specific to this group but it should be emphasised that a surgeon cannot monitor the patient's condition whilst performing the operation. It is recommended that an anaesthetist should be available in order to monitor the patient's general condition throughout the operation and to give resuscitation should it be required. This policy is contained in a document published jointly by the Royal Colleges of Anaesthetists and Ophthalmologists in 1993.⁵

Areas of concern included deaths after pharyngeal pouch surgery, the management of the obstructed airway (see commentaries on pages 27 to 32) and the poor case selection and preoperative preparation seen in some cases, especially as 68% (215/315) of procedures were elective or scheduled with plenty of time to properly assess and prepare patients for surgery.

The obstructed airway in head and neck surgery

Key messages

• Management of the partly obstructed airway should be a carefully planned procedure between *senior* surgical and anaesthetic staff. This is a clinical situation for which a local protocol could be devised.

Clinical management

- Wherever possible the level and the full extent of the airway difficulty should be defined by preoperative investigations including the use of fibreoptic nasendoscopy.
- Early consultation between surgeon and anaesthetist is essential, with consideration of all options and the formation of a management plan.
- Awake fibreoptic intubation and tracheostomy using local anaesthesia should be considered amongst the
 options.

Personnel and training

- Appropriately trained assistance should be available for anaesthetists and surgeons preparing for a difficult intubation.
- Fibreoptic intubation is an established part of anaesthetic practice. Several individuals within a department need to ensure that they maintain these skills.

Equipment

• A fibreoptic intubating laryngoscope should be readily available for use in all surgical hospitals.

Written by Dr Kathy Sherry FRCA and Mr Ron Hoile MS, FRCS in consultation with surgical and anaesthetic advisors (see pages 10 and 11)

From the deaths reported to NCEPOD, anaesthetic and surgical advisors with an interest in head and neck surgery reviewed the questionnaires of 30 patients who had been referred for management of lesions in the upper airway or who presented with stridor. This report is based on their discussions and opinions with reference to the NCEPOD questionnaires.

The cases highlighted both good and poor practice in airway management. In most cases the airway management did not contribute to the patient's death which was related to progress of the underlying disease, for example, malignancy.

The following cases with known airway obstruction illustrate a variety of airway managements.

	Age	Diagnosis	Operation	Airway control	Outcome
1	85	Retrosternal goitre	Resection	Awake fibreoptic intubation	Inoperable, extubated and returned to the ward
2	58	Subglottic and thyroid secondaries from primary renal tumour	Subglottic resection	Propofol/vecuronium, grade four laryngoscopy	Extubated, had an undiagnosed pharyngeal mass, airway obstruction followed by cardiac arrest and death in recovery area
3	69	Cricoarytenoid joint fixation	Tracheostomy	Gaseous induction, no relaxant, tracheal intubation	Uneventful
4	76	Retrosternal goitre in a patient with rheumatoid arthritis and fixed cervical spine flexion	Resection	Thiopentone/ suxamethonium two doses, failed intubation, severe hypoxia	45 minutes later, on mask ventilation and with hypoxia, difficult tracheostomy, cardiac arrest and died in theatre
5	83	Subglottic tumour	EUA	Awake fibreoptic intubation	Inoperable, extubated and returned to the ward
6	41	Supraglottic stenosis	Tracheostomy	Gaseous induction, assisted spontaneous ventilation adequate so no oral intubation	Uneventful
7	79	Subglottic tumour	Tracheostomy	Propofol/atracurium	Uneventful
8	85	Ca larynx	EUA	Gaseous induction, no relaxant, tracheal intubation	Extubated under deep anaesthesia, uneventful
9	83	Thyroid tumour	Debulking and tracheostomy	Failed awake fibreoptic intubation, intubated by awake direct laryngoscopy	Haemodynamic instability peroperatively, ventilated in recovery area overnight (ICU was full)
10	64	Extensive tumour in supraglottis and hypopharynx	EUA	Gaseous induction, "difficult mask ventilation", then mivacurium given, two anaesthetists and surgeon failed to intubate. Airway oedema and hypoxia supervened	Cricothyroid puncture and oxygenation using Sanders injector, then tracheostomy in the anaesthetic room. Pneumothorax diagnosed in the recovery area. CVA immediately postoperatively, died day two
11	81	Lymphomatous swelling R side of neck and stridor	Tracheostomy	Propofol/suxamethonium	Undiagnosed mid-tracheal obstruction, initially ventilated through orotracheal tube, then unable to ventilate via tracheostomy and subsequently by endotracheal tube, died in theatre
12	73	Thyroid goitre and obesity (112kg.)	Thyroidectomy	Etomidate/fentanyl/ suxamethonium two doses "nightmare intubation"	Uneventful

The preoperative management

When admission of a patient with airway pathology is *planned*, assessment starts during the outpatient visit. The history and examination should produce a high index of suspicion that the airway might be obstructed. When this is suspected an adequate view of the larynx should form part of the preoperative assessment, either by indirect laryngoscopy or by fibreoptic nasendoscopy. If an abnormality exists, a lateral soft tissue X-ray film of the neck and a PA chest X-ray film should follow the examination. Many surgeons and anaesthetists now consider that a CT scan of the upper airway is indicated. In five of the cases reviewed the patients had disseminated malignant disease with a possibility of more than one airway lesion (for examples see cases 2 and 11). If, from the history and chest X-ray, this possibility exists then a CT scan to demonstrate the *full* length of the airway from the teeth to the carina should be obtained.

In planned cases, on or before admission of the patient, there should be direct communication between the surgeon and anaesthetist to determine management strategies. The risks and planned airway management should be discussed with the patient as part of the informed consent process; this should include discussion of the alternative strategies that might be used.

It is recognised that, in *emergency* cases of critical airway obstruction, for example oedema of the glottis and stridor at rest, radiological investigations may be dangerous and, in these circumstances, may not be appropriate.

Experience

A team should decide the management of the obstructed airway. A consultant surgeon and a consultant anaesthetist should lead the team.

In this group, 21/30 (70%) of patients were anaesthetised by consultants and, in total, 26/30 (86%) by experienced anaesthetists (SpR 4 and above). In contrast an SHO anaesthetist in the first year of training sedated and supervised a patient with severe cardiovascular disease and a fractured mandible for plating of his jaw under local anaesthesia. Was an SHO appropriate for this sort of case?

The difficult airway is a training opportunity for both anaesthetists and surgeons. Solo anaesthetists managed eight patients. Experienced anaesthetists and surgeons working with trained assistants should be able to manage a difficult airway. However, it is important for *trainee* anaesthetists and surgeons to learn airway management under supervision and, whenever possible, a trainee attachment should be sought.

Some anaesthetists encounter airway obstruction infrequently and both the patient and anaesthetist may benefit by referral to a colleague with specialised experience. For one patient the consultant anaesthetist had sought advice from a colleague, for another patient the consultant anaesthetist enlisted help from a second consultant with special experience in management of the difficult airway.

The location

The trachea was intubated in the anaesthetic room in 19 cases and in the operating theatre in nine. In two cases there was no attempt to intubate the trachea. The management of a difficult airway requires a room of adequate size and appropriately equipped for a difficult intubation. A trained surgeon and surgical facilities should be immediately available. The surgical instrument pack for a tracheostomy should be open, checked and *ready for use*; there is no time for delay if difficulties arise.

The site of airway obstruction may be sub divided into: 1) supraglottic and glottic; 2) subglottic and 3) lower third of the trachea.

- 1 For supraglottic or glottic problems it is prudent to avoid relaxants unless mask ventilation is easy or direct laryngoscopy indicates that tracheal intubation will be straightforward. The caution with relaxants does not necessarily apply to lesions lower down the airway.
- 2 For subglottic obstruction a determined attempt should be made to ascertain the diameter of the airway preoperatively.
- Lesions of the lower third of the trachea may be missed, or their extent under-estimated, if CT scanning is not done or it is limited to the head and neck. Low tracheal lesions require carefully planned surgery. There were two patients where the standard tracheostomy tubes did not get past lesions in the lower third of the trachea (see case 11).

Anaesthetic management

Not all patients need oro- or naso-tracheal intubation.

One patient had surgery to the mandible under local anaesthesia. When the airway is obstructed the use of local anaesthesia is appropriate if the lesion is accessible and the patient is cooperative; however, there must be an alternative strategy in case of failure.

One patient (case 6) had inhalation induction and assisted spontaneous mask ventilation until tracheostomy. The experienced anaesthetist sensibly commented "he had a good airway and I thought it was wise <u>not</u> to attempt endotracheal intubation".

Awake fibreoptic laryngoscopic intubation was used in only 6/30 (20%) of patients. Awake fibreoptic tracheal intubation is widely recognised as a safe method; however, it does require the anaesthetist to be trained and practised in its use. If the anaesthetist managing the patient lacks this skill then consideration should be given to drawing on the experience of other anaesthetic or surgical colleagues. A fibreoptic intubating laryngoscope should be readily available for use in all surgical hospitals.

Inhalation induction alone was used in 11 patients and combined with sedation in three. Sevoflurane for inhalation induction was used in eight cases, halothane in five and enflurane in one.

The laryngeal mask airway, now part of difficult intubation management⁶ may not be an effective airway in the presence of supraglottic obstruction. The use of a laryngeal mask airway was reported in four cases. On three occasions, all with supraglottic lesions and airway distortion, it could not be well positioned. In the fourth case with normal supraglottic anatomy and subglottic stenosis the laryngeal mask airway effected a conduit for fibreoptic laryngoscopy to assess the airway.

Surgical management

An elective tracheostomy under local anaesthesia should always be considered as an option in cases where significant upper airway obstruction is diagnosed. This is not an easy operation and trainees in anaesthesia and surgery should be encouraged to learn the appropriate local anaesthetic and surgical techniques, and to see this rare procedure.

Needle cricothyroidotomy and jet ventilation may allow a brief period of oxygenation prior to tracheostomy, providing that appropriate driving gas pressures are used and the upper airway is sufficiently patent to allow expiration. It was used in one patient (case 10).

The decision to perform an emergency tracheostomy should be taken expeditiously.

Two patients (cases 4 and 10) had an emergency surgical tracheostomy. In both patients it was accompanied by complications. In one patient (case 4) the medical staff could have anticipated that it would be technically difficult to do a tracheostomy in an emergency situation.

A conventional tracheostomy tube will not always pass through mid and lower tracheal lesions. Two patients had lesions extending into the lower third of the trachea which were not crossed by the tracheostomy tube (including case 11). The extent of these lesions was not anticipated, the management of the complication was unplanned and, because of this, the surgical options were limited.

A surgeon experienced in rigid bronchoscopy may be able to visualise the larynx using a more lateral approach through the mouth when conventional laryngoscopy gives no view and provide either a temporary airway or position a bougie. This solution was not used in any of these cases.

When intubation of the trachea was particularly complicated it appeared that, often, it was not a single cause but a chain of factors resulting in difficulties. For example: preoperative investigations were incomplete, there was no alternative strategy in case the initial management failed, there was repeated use of short-acting relaxants, long-acting relaxants had been used, the intubation attempt continued until the vulnerable airway was critically oedematous or the team were locked into repeated attempts at orotracheal intubation and failed to stop and think of the alternatives. The difficult questions are always "how long to persevere before surgical intervention?" and "how long before hazardous airway oedema will develop?" There is great variation between patients. In one case the cords were seen only once before critical oedema supervened.

The postoperative management

Extubation of the airway also requires planning. For glottic and especially supraglottic lesions patients will often need good airway control and in these cases awake tracheal extubation is indicated. For subglottic and lower lesions, coughing on the tracheal tube may exacerbate oedema and consideration should be given to extubation of the trachea under deep anaesthesia. In all patients with airway obstruction care should be taken to ensure that they are returning to an area staffed by appropriately skilled nurses. When the airway has been secured by a tracheostomy then patients should be in an area with nursing staff skilled in tracheostomy management at all times.

The advisors considered the following, along with other cases, to be an example of good clinical practice.

An 85-year-old man who presented with stridor from disseminated thyroid carcinoma and invasion of the airway was referred for a tracheostomy. He also had dysphagia, hypercalcaemia, renal impairment and was wheelchair-bound due to rib and spinal secondaries. Preoperative assessment included CT scan and barium swallow. A bilateral vocal cord palsy was identified. His initial airway management was by two consultant anaesthetists, one with special experience of the difficult airway. In theatre he was sedated with three increments of midazolam of 0.5mg and fentanyl 10,10 and 5 microgrammes. Topical anaesthesia of the nose and pharynx enabled fibreoptic intubation of the trachea, despite almost no identifiable structures. A 5.5mm nasal microlaryngoscopy tube was passed and its position confirmed by capnography before inhalation induction of anaesthesia. A tracheostomy was then done by an experienced consultant otolaryngologist. After the tracheostomy the patient was monitored on the HDU overnight.

Pharyngeal pouch surgery

Key messages

- Pharyngeal pouch surgery is an uncommon procedure associated with a significant mortality (1-2%).
- In the year of study eight questionnaires were received. Seven deaths occurred after open surgery and one followed endoscopic stapling.
- Endoscopic stapling is a quicker, lesser invasive procedure associated with earlier postoperative discharge from hospital.
- Subspecialisation within otolaryngology departments for this type of surgery would seem appropriate.

Personal commentary by Mr Adel Resouly FRCS in consultation with the head and neck surgery advisors (see page 11)

The total number of operations for pharyngeal pouch performed in England range between 437-504 per annum over the five years up to 1995/96. Surgical treatment is associated with a significant mortality rate, which in the last five years ranged between 0.6-2%. Hospital Episode Statistics (HES) data⁷ and information on deaths collected by NCEPOD (albeit without denominator data) suggest that the mortality rate for pharyngeal pouch surgery may be higher than previously recorded.

In the year under study, there were eight questionnaires returned following pharyngeal pouch surgery, seven of which were by open techniques and one by endoscopic stapling. The cause of death in most cases was reported as either respiratory or cardiac failure.

The majority of patients undergoing pharyngeal pouch surgery are elderly and represent high risk as elderly medicine physicians often consider them to be too frail for surgery. The mortality rates observed are a reflection of this frailty.

There are several open surgical procedures in current practice. The relatively new procedure of endoscopic stapled diverticulotomy is quick, less invasive, bloodless and safe. 9,10 A wider use of this procedure could transform the outcome for these patients, as there is a reduced anaesthetic time, return to normal swallowing in 24 hours and a reduced hospital stay. Specialist head and neck surgeons should perform this procedure where appropriate and there should be one or two surgeons within a department of otolaryngology who are trained and competent in the surgery of pharyngeal pouch. Should more general surgeons, especially endoscopists, wish to do this procedure then it would be essential to obtain proper training. Long term results are not yet published and individual departments are urged to audit their results.

Anaesthesia for carotid endarterectomy

Key messages

- Intra-arterial pressure monitoring should be routine for carotid endarterectomy.
- Peroperative blood pressure control needs to be excellent.
- Approximately half the deaths associated with carotid endarterectomy are due to causes other than stroke.
- Record keeping was often surprisingly poor considering the risks associated with this operation.

Written by Dr Anthony Gray FRCA and Mr Ron Hoile MS, FRCS in consultation with the anaesthetic advisors (see page 10)

The frequency of carotid endarterectomy is increasing. There were 1,748 carotid endarterectomies performed in 1993/94 and 2,781 in 1995/96 in England (Hospital Episode Statistics). Two large trials have compared carotid endarterectomy coupled with medical treatment against medical treatment alone. These trials showed that patients with severe symptomatic carotid artery stenosis benefited from surgery. This conclusion may not be applicable in units where the peroperative complication rate is higher than in these trials. In the North American Symptomatic Carotid Endarterectomy Trial, 5.8% of patients allocated to surgery had a major stroke or died between randomisation and 30 days after surgery. In the MRC European Carotid Surgery Trial, 7.0% of patients had a major stroke or died within 30 days of surgery.

Surgical techniques are constantly being modified particularly in relation to the maintenance of cerebral circulation during surgery. Most operations will involve the use of some form of monitoring to assess cerebral circulation such as the patient's conscious level (if under local anaesthesia), transcranial Doppler (TCD), electroencephalography (EEG) or near infra-red spectroscopy (NIRS). Alternatively surgeons will use an arterial shunt. Anaesthetic techniques are chosen so as to maintain cerebral blood flow whilst minimising cardiac morbidity.

NCEPOD received details of 30 patients who died after carotid endarterectomy. In one case no surgical questionnaire was returned; in 11 cases no anaesthetic questionnaire was returned. Hospital Episode Statistics report 35 deaths in England in 1993/94 and 34 in 1995/96,⁷ so it is likely that most deaths following carotid endarterectomy were reported to NCEPOD.

Comment on selected cases

Age	ASA grade	Previous medical history	Anaesthetic/surgical management	Outcome
53	3	CVA, hypertension, diabetes	Pre-induction BP 210/95. Prolonged intra- operative hypotension with BP of 100/60. Hypertensive crisis postoperatively	Postoperative CVA
70	3	COPD, left ventricular hypertrophy	Gap in monitoring of 12 minutes when BP 75/40, possibly when moving into theatre	Intra-operative CVA, pneumonia
76	3	Resolving pneumonia , angina, pacemaker	Pre-operative ward BP 140/80. Systolic BP of 30 after induction	Postoperative acute LVF, cardiac arrest, successful resuscitation, pneumonia
72	2	Asthma, previous vascular surgery	Very poor records	Postoperative CVA
66	4	Asthma, previous MI, crescendo TIAs	Chest pain in morning, not thought to be cardiac. Asystole for 30 seconds post-induction	Postoperative CVA
76	3	Angina, hypertension	One episode of bradycardia of 30 bpm	Postoperative CVA
74	2	Previously fit	Pre-operative ward BP 130/80, 95/40 for most of the operation	Postoperative CVA
59	2	Previous CVA, MI, diabetes	Pre-induction BP 170/80. Intra-operatively BP varied between 120/65 and 100/55	Postoperative MI, cardiac failure
81	3	Previous CVA, MI, angina	Pre-operative ward BP 170/80. Down to 100/60 after induction. BP of 240/110 in recovery	Cardiac arrest
54	3	Previous CVA, asthma, hypertension	Uneventful	Postoperatively failed to breathe, developed signs of CVA and pneumonia
63	3	COPD, previous CABG	Uneventful	Postoperative CVA
63	3	Previous CVA and CABG, chronic renal failure	Uneventful	Haematoma in neck. Returned to theatre for tracheostomy, went into ventricular fibrillation
66	4	TIAs from right and left carotids	Uneventful	Postoperative CVA
58	3	Previous CVA, diabetes	Uneventful	Cardiogenic shock
65	3	Previously fit	Uneventful	Postoperative CVA

75	3	Ischaemic heart disease	Poor records	Returned to theatre with haematoma in the neck. Good recovery. Found dead after return to the ward, postmortem report not available to NCEPOD
82	3	MI, paroxysmal AF	Uneventful	CVA after return to theatre for bleeding from hernia repair done concurrently with carotid endarterectomy
74	2	Previous CVA	Uneventful	Ruptured intra-cranial aneurysm
63	2	TIAs. Occluded right common carotid artery and tight stenosis left internal carotid	Left carotid endarterectomy. Hypertensive during procedure.	Postoperative CVA

Grade of the anaesthetist

All the patients were anaesthetised by an experienced anaesthetist.

Consultant	17
Associate specialist	1
Specialist registrar year 4	1

Thirteen of the consultants were accompanied by another anaesthetist:

Specialist registrar year 4	1
Specialist registrar year 3	5
Specialist registrar year 2	5
Clinical assistant	1
Senior house officer year 1	2

Pre-operative assessment

Nine out of the 19 patients for whom anaesthetic questionnaires were received had the ward blood pressure recorded on the anaesthetic assessment form. The blood pressure was recorded before induction in 12 patients. In four patients the first blood pressure recording was made only after the induction of anaesthesia. It is difficult to manage a patient's blood pressure without knowledge of its usual level. All patients had the expected pre-operative investigations including ECG. Echocardiography was performed in three patients.

Monitoring

Monitoring of ECG, pulse oximetry and capnography was reported in all cases. Direct intra-arterial pressure monitoring was used in 16 cases out of 19, and in 11 of these, recording was started before induction. In one of the three cases where the blood pressure was measured non-invasively an arterial catheter was inserted but then became dislodged and the anaesthetist was unable to site a replacement.

Capnography was used in 18 cases, although in one case no readings were recorded. In five cases average end-tidal CO_2 recordings were below 4.5 kPa, in eight cases between 4.5 and 5.4 kPa, and in five cases were

above 5.5 kPa. The patients had a mean age of 67 and a third had respiratory disease, so one could anticipate an unpredictable disparity between the end-tidal CO₂ and arterial PaCO₂. Nonetheless, no capnograph recordings were checked against blood gas analysis. The one patient managed without a capnograph had the blood gases measured on two occasions during their operation.

Cerebral function monitoring with transcranial Doppler (TCD) was used in two cases, and "EEG/Cerebral Function Monitoring" in another. Two patients were monitored with near infra-red spectroscopy (NIRS) and TCD.

Anaesthesia

One patient was managed with local anaesthesia alone. All other patients received general anaesthesia. In two patients this was supplemented with a cervical plexus block.

Many cases were marked by episodes of hypotension. Some patients were hypotensive throughout the operation. Four patients had marked falls in blood pressure after induction (one patient had a period of asystole). The management of intra-operative hypotension sometimes appeared illogical; small doses of vasopressors were given without effect and hypotension was then tolerated without any further intervention.

Grade of the surgeon (supervision in theatre)

Twenty-eight procedures were done by an experienced surgeon. Consultants were involved in the decision-making for all cases but were not physically present for two procedures* done by trainees.

Consultant	27
Accredited specialist registrar	1
Specialist registrar year 4*	1
Specialist registrar year 3*	1

Postoperative care

These patients require a high level of supervision continuing into the postoperative period, to monitor blood pressure and to detect impending carotid occlusion. Some patients were discharged from the recovery area to a general ward before their condition had stabilised. Is it not advisable that after operation all such patients should be managed in an area with at least the facilities of a High Dependency Unit, where intra-arterial monitoring can be continued for some hours after the end of surgery?

Record keeping

Record keeping was poor in some cases; it is difficult to understand such a casual approach to a potentially hazardous anaesthetic and surgical procedure. Is it wise to make only six recordings of the blood pressure in 2½ hours of anaesthesia? Automated record-keeping would be expected to help with the common clinical scenario of the anaesthetist trying to keep good records whilst carrying out several anaesthetic interventions. Unfortunately, the automated charts received at NCEPOD often had periods without any readings lasting several minutes, at times of cardiovascular instability. These gaps may have occurred when patients were being moved from anaesthetic room to theatre. Operative records were disappointingly poor and did not comply with guidelines for record keeping.¹³

Cause of death in all 30 cases reported to NCEPOD

Fifteen patients died from a stroke. Ten died from cardiac causes, four from respiratory causes, and one from rupture of an intra-cranial aneurysm. The postmortem examination of one patient who developed atrial fibrillation after operation followed by cardiac arrest showed that the patient had had a large myocardial infarction in the days preceding his operation. This patient also suffered from diabetes mellitus. Another patient had already suffered an established stroke and had aspiration pneumonia. His pneumonia worsened after surgery leading to septicaemia and death.

Conclusions

It would appear that in England, Wales and Northern Ireland, at the time of data collection, carotid endarterectomy was usually performed under general anaesthesia. Publications suggest that in North America a much greater number are performed under local anaesthesia. A review by the Cochrane Collaboration¹⁴ found that whilst non-randomised trials suggested potential benefits from using local anaesthetic techniques, there are no good controlled trials to assist in deciding whether local or general anaesthesia is preferable.

Carotid endarterectomy poses important problems for the anaesthetist and the surgeon with patients almost always having significant concurrent medical conditions. Attention to the minimisation of risk factors for cardiovascular complications is as important as preserving cerebral oxygenation. The data reported to NCEPOD reflect published work that shows that approximately half the mortality from this operation is due to causes other than strokes. The care of many patients was exemplary. In others, it is impossible to say whether deaths would have been prevented by changes in the anaesthetic and surgical management but improvements were possible in

- pre-operative assessment
- monitoring
- peroperative management of blood pressure
- peroperative management of CO₂ levels
- record keeping

Minimally invasive surgery (excluding gynaecology and urology)

Key messages

- Minimal access surgery is not synonymous with a minimal procedure.
- Few deaths were related to diagnostic endoscopy but rather to subsequent major surgery.
- There was a high incidence of consultant involvement in decision making and operating.

Written by Mr Ron Hoile MS, FRCS in consultation with the minimally invasive surgery advisors (see page 11)

The term "minimally invasive surgery" covers a wide range of procedures (see glossary, page 109) and the sample included many procedures, such as gastroscopy and diagnostic thoracoscopy, which were simply a part of the patient's management rather than the final procedure (the specialties involved were mainly general and cardiothoracic surgery). In addition, many of these patients were elderly, frail with advanced malignancy or exsanguinating haemorrhage. A minimally invasive procedure was often used to obtain a diagnosis in the simplest way or in an attempt to arrest haemorrhage without major surgery. There were few deaths directly related to these procedures. However, these endoscopies often led to more major procedures from which the patient died. These subsequent operations and resections sometimes seemed inappropriate and aggressive. It is a surgical skill to recognise when surgery will be too adventurous, ill advised or futile, given the condition of the patient. It is difficult to resist pressure to operate, whether this comes from the patient, relatives or medical colleagues but it must be recognised that surgery cannot solve every problem.

Review of this sample produced several areas worthy of comment such as misconceptions about fitness for laparoscopy, the need to understand the indications for (and role of) a PEG, and the management of biliary stones in the elderly. These areas are addressed in short articles later in this section.

List of procedures

Procedure (may be multiple for individual patients)	Number
Percutaneous endoscopic gastrostomy (PEG)	17
Laparoscopic procedures	
Diagnostic laparoscopy (+/- biopsy) Cholecystectomy Inguinal hernia repair (two with adhesiolysis) Gastrojejunostomy Suture of perforated peptic ulcer Adhesiolysis Miscellaneous (one each of: loop colostomy, hiatus hernia repair, right hemicolectomy and drainage of retroperitoneal abscess)	35 14 4 2 2 2 2 4
Procedures following or as a consequence of laparoscopy "Open and shut" laparotomy Open cholecystectomy Miscellaneous (one each of: small bowel resection, pancreatoduodenectomy, re-exploration for bleeding, subtotal gastrectomy, division of adhesions, gastroscopy, Hartmann's procedure, oversewing of duodenal ulcer, Roux-en-Y repair of hepatic duct injury, right hemicolectomy and pyloroplasty)	2 2 11

Endoscopy

Oesophagogastroduodenoscopy (+/- biopsy or injection)	95
Endoscopic retrograde cholangiopancreatography (+/- sphincterotomy and stenting)	29
Rigid sigmoidoscopy (+/- biopsy)	27
Colonoscopy (+/- polypectomy)	20
Flexible sigmoidoscopy	13
Arthroscopy (knee)	5
Transanal endoscopic resection of tumour	3
Thoracoscopy	2
Bronchoscopy	2
Miscellaneous (one each of: arthroscopic menisectomy and percutaneous cholangiography and	2
stent)	
Procedures following or as a consequence of endoscopy	
Laparotomy and oversewing of bleeding ulcer	6
Sigmoid colectomy (perforation)	2
Laparotomy for bleeding oesophageal varices	2
Transverse colostomy	2
"Open and shut" laparotomy	2
Miscellaneous (one each of: partial gastrectomy, right hemicolectomy, haemorrhoidectomy,	7
laparotomy for gastrointestinal bleeding [cause unknown], sigmoid colectomy [volvulus],	
laparotomy for ileo-ileal fistula and drainage of abscess)	

General comment

The majority of patients (80%, 250/310) had an ASA status of 3 or higher, i.e. were of high risk, and 62% (192/310) were over the age of 69 years. Deaths were mainly due to advancing malignancy and severe haemorrhage in the presence of extensive comorbidity and senescence. Most surgeons involved in these cases declared a special interest, which was appropriate for the case concerned; there were only 35/310 cases in which the surgeon perceived him/herself as a 'general surgeon'. The involvement of seniors was commendably high with 93% of cases having some input from consultants. There were 21 cases in which no consultant involvement was recorded. A consultant was personally operating in 66% (207/310) of cases and overall a consultant was the most senior person present in the operating theatre in 73% (226/310) of cases Direct comparison with other NCEPOD reports cannot be made but this appears to be a good level of consultant input. The advisors identified only one case where the grade of surgeon was possibly inappropriate; a case of iatrogenic colonic perforation following colonoscopy in which a laparotomy and colonic resection was done by a specialist registrar (year 3) after discussion with a consultant. Where the procedure was considered inappropriate it was the initial case selection which appeared to be suspect, coupled with a failure to adopt an holistic approach.

Another positive and encouraging area was the small number of cases where any form of non-clinical delay occurred (3%, 10/310); one could speculate that this may be due to the wider introduction of emergency theatres, often referred to as 'CEPOD' theatres. The use of audit and postmortem examinations in this group was also adequate with 74% (230/310) of cases undergoing investigation by a combination of local audit and/or postmortem examination as appropriate.

Specific consideration of the deaths following cardiothoracic surgery raised several general points. There were no deaths directly related to the minimally invasive procedures but the subsequent resections were sometimes inappropriate and aggressive. Two procedures in particular were considered to require direct consultant involvement. These were pneumonectomy in the elderly (over 75-year-old) patient with a high ASA class, and mediastinoscopy.

Minimally invasive abdominal surgery may not be minimally stressful to the patient

Key message

Surgeons, as well as anaesthetists, need to be aware of the effects of peritoneal distension and the
position of the patient. Despite careful anaesthetic management, unexpected physiological changes often
occur.

Personal commentary written by Dr Ian Baguley FRCA and Dr Anthony Gray FRCA

Increasing numbers of surgeons and anaesthetists believe that laparoscopic general surgery may be advantageous in high risk patients and the elderly. It is thought that surgical trauma and postoperative pain are reduced, fewer respiratory and cardiac sequelae are seen, and the patients return to normal activities earlier. However, laparoscopic surgery can be associated with physiological changes which compromise both the respiratory and cardiovascular systems. These include high positive airways pressures, raised intra-abdominal pressure, and effects on venous return due to Trendelenburg and reversed Trendelenburg position, increased intrathoracic pressure, and compression of the inferior vena cava and iliac veins. Measuring, interpreting and correcting changes to these systems during surgery is essential but can be difficult.

Anaesthetic questionnaires were returned for 21 deaths in general surgical patients undergoing laparoscopic procedures in whom the pathology was benign.

Age	Operation	Days to death	Preoperative condition	Peroperative course
42	Laparoscopic cholecystectomy converted to open	1	Cardiac murmur, mild mitral regurgitation on echocardiogram	Uneventful anaesthetic. Cardiac arrest in ventricular fibrillation next day. Postmortem examination found no cardiac abnormality.
63	Laparoscopic cholecystectomy	3	COPD, untreated hypertension and ischaemic heart disease (graded ASA 1 by the anaesthetist)	Hypertensive in recovery ward. Three days later, severe hypotension followed after one hour by cardiac arrest. Postmortem showed pulmonary oedema cerebral infarction and severe coronary atheroma but no cardiac infarction.
63	Laparoscopic cholecystectomy	17	Insulin controlled diabetes. Previous CVA. AF, CCF, hypertension	Developed hypotension in recovery ward, admitted to ICU and treated with inotropes. Responded well and returned to the general ward, but continued to have episodes of cardiac failure and died.
66	Laparoscopic cholecystectomy	0	NIDDM. No cardiac symptoms. ECG showed some ischaemic changes	Became hypotensive on induction of anaesthesia, responded to ephedrine. Patient then stable until placed in head-up position. Became hypotensive with fall in expired carbon dioxide, progressing to cardiac arrest. Postmortem examination showed severe coronary atheroma.
68	Laparoscopy and drainage of abscess	9	NIDDM, poor general condition	Tachycardic throughout operation. CVA fourth postoperative day with progressive deterioration.
74	Laparoscopic cholecystectomy	1	Hypertrophic cardiomyopathy, AF, LVF	Previously turned down because of cardiac condition, but patient wished to proceed with surgery. Elective postoperative admission to ICU, but developed intractable LVF.

75	Laparoscopic cholecystectomy	9	Hypertension	Arrested on ward. Postmortem showed pulmonary emboli and severe coronary atheroma.
75	Laparoscopic cholecystectomy	30	COPD	Bronchospasm in recovery ward. Admitted to ICU 26 days after operation in acute respiratory failure. No information provided on events before ICU admission.
76	Laparoscopic cholecystectomy	18	Asthma. Angina, previous coronary angioplasty	Initially did well. Re-admitted seven days after operation with MI, and progressively deteriorated.
77	Laparoscopic cholecystectomy	10	Angina, hypertension, COPD	Hypotension intra-operatively when patient positioned head-up. Postoperatively required two unit blood transfusion. Found dead in bed four days after discharge. Postmortem confirmed presence of coronary artery disease.
77	Laparoscopic cholecystectomy	14	"? chest infection"	Intra-operative hypotension. Required doxapram infusion postoperatively. Developed LVF and bronchopneumonia.
78	Inguinal hernia repair and adhesiolysis	5	Angina	Left ventricular failure two days after operation. Developed intractable cardiogenic shock. Postmortem examination showed patient had had an MI one week before surgery which had extended.
80	Bilateral inguinal hernia repair	2	Fit	Hypertensive on return to ward. Hypotensive first postoperative day, treated with fluid. Cardiac arrest second day. No postmortem examination.
80	Bilateral inguinal hernia repair	11	Angina	MI second postoperative day. Developed LVF, and despite transfer to ICU on seventh day, died.
80	Laparoscopic cholecystectomy	8	Hypertension. Mild renal failure	Pulmonary embolus.
84	Laparoscopic cholecystectomy plus inguinal	26	Previously fit	Initially did well, discharged on day seven. Admitted 12 days later with urinary retention, renal failure, DVT and pulmonary embolism from which he died.
84	hernia repair Laparoscopic cholecystectomy	2	Mild CCF	Delayed recovery from anaesthesia, required admission to ICU, followed by an episode of sepsis requiring inotropic support. After discharge from ICU developed pneumonia and CCF.
84	Laparoscopic cholecystectomy	21	Fit	Labile blood pressure intra-operatively. Myocardial infarction first postoperative day. Subsequently developed left ventricular failure, biliary collection, renal failure, and gastro-intestinal bleeding. Postmortem showed moderate coronary atheroma.
85	Suture of perforated peptic ulcer	1	Angina, hypertension, with peritonitis	Two hour operation. Hypotensive five hours after surgery, transferred to ICU in LVF. Died 20 hours after operation. Postmortem showed severe coronary atheroma.
86	Laparoscopy and peritoneal washout	6	No information	No intra-operative problems. Died from peritonitis from perforated diverticulum.
88	Laparoscopic adhesiolysis	14	COPD, angina, renal failure, CCF	Operation for abdominal pain. Progressive deterioration postoperatively with development of pulmonary oedema.

The respiratory effects of pneumoperitoneum can be inferred from changes in ventilator pressure, expired carbon dioxide content and peripheral oxygen saturation. As muscle relaxation is generally used, changes to the pattern of ventilation can overcome the respiratory embarrassment. After surgery the reduced compliance of the chest and the increased airways resistance seen during pneumoperitoneum revert to pre-operative values within approximately one hour depending on the site and extent of surgery. Tests of respiratory function are abnormal only at the extremes of respiratory effort. Laparoscopic surgery appears to be suitable for patients with both normal and compromised respiratory systems, with only one patient in the table above dying from respiratory disease exclusively. Nevertheless, patients with severe respiratory disease may still be at risk of postoperative respiratory complications.

Respiratory events contributed to death in two patients; one died from bronchopneumonia, and a second from chronic obstructive pulmonary disease.

The cardiovascular effects of pneumoperitoneum cannot easily be detected by routine non-invasive monitoring. Changes in cardiac output and vascular resistance are not usually reflected in the capnograph, oximeter, ECG or blood pressure recordings.

General anaesthesia invariably produces a transient fall in cardiac output and blood pressure which most patients tolerate well. Positioning the patient head-up also reduces cardiac output. The introduction of pneumoperitoneum with direct compression of femoral veins and the vena cava further reduces flow, venous return and cardiac output. This may promote iliac vein thrombosis. The pneumoperitoneum splints the diaphragm which increases mean intrathoracic pressure, compresses the pulmonary vessels and increases the work of the right ventricle. Stimulation of the renin-angiotensin system contributes to an increase in systemic vascular resistance which increases the work of the left ventricle. The blood pressure is maintained or may even rise, giving the appearance that all is well even though cardiac output may be markedly reduced. Volatile anaesthetic agents which cause vasodilatation will ameliorate the increase in systemic vascular resistance. Vasopressors, when used to treat a decrease in blood pressure which has been caused by a fall in cardiac output, will further increase vascular resistance. These changes have been confirmed in both healthy and high risk patients undergoing laparoscopic cholecystectomy although the subject numbers are small and more research is needed.

These adverse physiological events may have been responsible for the episodes of hypotension and hypertension seen on reviewing the questionnaires. Laparoscopic surgery did not protect patients from developing severe cardiac complications during or after surgery. Many of these patients did not have overt cardiac disease when assessed pre-operatively.

In summary, laparoscopic surgery and its associated anaesthesia should be treated with respect. Recovery after laparoscopic surgery is probably improved compared to open procedures, but during the operation itself, laparoscopic procedures may induce as much physiological stress. Respiratory problems seem to be uncommon after laparoscopic procedures. However, when there is any suggestion of cardiac failure or widespread ischaemic heart disease there may be significant risk to the patient.

Percutaneous Endoscopic Gastrostomy (PEG)

Key messages

- There should be careful patient selection prior to the insertion of a PEG.
- PEG tubes are inappropriate in patients with a very short life expectancy.
- A PEG does not protect a patient from aspiration of gastric contents.

Written by Mr R David Rosin MS, FRCS and Mr Ron Hoile MS, FRCS in consultation with the minimally invasive surgery advisors (see page 11)

There were 17 questionnaires received concerning deaths after the endoscopic insertion of a percutaneous gastrostomy (PEG), 12 of which occurred within three days of the procedure. Details of the cases are given below followed by a commentary. NCEPOD has no information as to the number of PEGs that are appropriately used with advantages for the patient. The cases described below suggest that some clinicians are under the misconception that a PEG can be used to prevent aspiration of gastric contents, which it does not. In other cases a PEG was inserted in situations where symptom control might have been more relevant. Bronchopneumonia was often referred to as 'the old man's friend'. Is that statement still true in modern hospital practice or do we strive too hard to keep patients alive when the prognosis is hopeless and death is imminent?

Age	Indication	ASA	Comorbidity	Anaesthesia	Grade of surgeon	Day and cause of death
21	Brain stem astrocytoma.	?	Asthma & respiratory failure	LA + sedation	Consultant. Does 3/year	Died same day. Cardiac arrest
37	Carcinomatosis. Nasogastric tube not tolerated. PEG inserted for gastric drainage.	4		GA	Accredited specialist registrar. Does 20/year	Day 1. Carcinomatosis
43	Advanced carcinoma of prostate.	4		LA + sedation	Consultant. Does 6/year	Day 2. Carcinomatosis
50	Neurological problems (not specified).	4		LA + sedation	Specialist registrar year 4. No experience	Day 6. Pulmonary embolus and aspiration pneumonia.
61	CVA. No gag reflex	4		LA + sedation	Consultant. Does 6/year	Day 4. Cardiac failure and anaphylaxis (drugs)
61	Multiple sclerosis. Nasogastric feeding not tolerated.	4		LA + sedation	Consultant. Does 6/year	Day 7. Chest infection and septicaemia
63	Motor neurone disease. Unable to swallow. Nursed in GP unit.	4		LA + sedation	Basic surgical trainee. Does less than 6/year	Day 3. Aspiration pneumonia.

71	CVA. Unable to swallow. Two previous PEGs.	3		LA + sedation	Specialist registrar year 4 (supervised). Does 40/year.	Day 9. Aspiration pneumonia
75	CVA	3	Respiratory. Extension of CVA at end of procedure.	LA + sedation	Consultant. Does 12/year	Day 2. Progressive CVA.
76	CVA.	4	Carcinoma of colon, bleeding duodenal ulcer and peripheral vascular disease.	GA	Specialist registrar year 4	Day 16. Respiratory failure.
77	CVA	4		LA + sedation	Consultant. Does 5/year	Day 5. General deterioration and further CVA
79	RTA. Comatose with cerebral haematoma.	4	Demented	LA + sedation	Associate specialist. Does 50/year	Day 6. General deterioration
83	CVA. Aspiration pneumonia and dysarthria	4	·	LA + sedation	Consultant surgeon and anaesthetist present. Surgeon does 3/year	Day 2. Aspiration pneumonia.
85	Senile dementia. Facial fractures	4	Respiratory, cardiac, renal and neurological disease.	LA + sedation	Consultant. Does 50/year	Day 2. Aspiration pneumonia
86	Fractured femoral neck	4	Chest infection, ischaemic heart disease, cardiac failure and Parkinson's disease.	LA + sedation	Consultant. Does 30/year	Day 3. Bronchopneumonia
90	Multiple sclerosis	4	Chest infection	LA + sedation	Basic surgical trainee. Does 6/year	Day 5. Bronchopneumonia
92	CVA	4	Chest infection	LA + sedation	Consultant. Does 20/year	Day 7. Respiratory failure

Percutaneous endoscopic gastrostomy (PEG) has gained wide acceptance for patients who require prolonged tube feeding support. It is a well-established procedure for creating a feeding port in patients who are unable to take oral nutrition. The patients in this sample were often elderly, with multiple ailments and therefore mortality and morbidity was, in the main, due to comorbidities rather than the placement of the gastrostomy. Bronchial aspiration, which was a feature in many of the cases, is known to be reduced when feeding via a PEG but is certainly not abolished.

Percutaneous endoscopic gastrostomy is used to provide nutrition for patients who are unable to eat but have a functionally intact gut. Clinical guidelines for a PEG are uncertain and have been derived mainly from referral practices. PEGs are mainly used for patients with neurological disease, head and neck cancers, acquired immuno-deficiency syndromes, critically injured patients and in-patients with dementia. We were surprised at the number of deaths following the insertion of a PEG, especially the twelve occurring within three days of the procedure. Many of the deaths were due to aspiration pneumonia and there were other respiratory complications. It would seem that most early deaths were due to pulmonary complications whereas later deaths were usually due to the reason for using the PEG (especially in patients with malignant disease).

Review of the literature confirms that younger patients have a lower mortality and morbidity following insertion of a PEG. One paper shows that all patients under 40 years of age survived whereas the mortality in patients over 40 was 59%. A paper looking at predictive factors for early mortality after a PEG identified a subgroup of patients with a high mortality rate after the procedure patients with urinary tract infection and previous aspiration were much more likely to die within a week after insertion of a PEG. Patients who had both risk factors had a 48.4% probability of dying within seven days whereas those who had no risk factors had a 4.3% probability of death. Urinary tract infection, previous aspiration and an age greater than 75 years were also predictive factors for death at one month after a PEG. Patients who had all three risk factors had a 67.1% probability of death at one month while those who had no risk factors had a 10% probability of death at this point. The authors of this paper concluded that there was a need to evaluate less invasive ways of providing nutritional support for these patients.

In an audit of PEG in long term enteral feeding in a nursing home¹⁷ it was shown that only 20% of patients with a PEG were free of cutaneous problems around the insertion site and eight abscesses occurred during the first week. Pulmonary complications occurred in 39% of patients with a PEG.

In a large audit of long term outcomes, 7369 patients with a PEG inserted between 1990 and 1992 were reviewed. ¹⁸ The mean age was 68.1 years and 98.6% were men. The indications were cerebrovascular disease, other organic neurological disease and head and neck cancer. The authors pointed out that the complication rate of the procedure itself was low (4%) but because of the severity of the underlying diseases, 23.5% of the patients died during the hospitalisation during which the PEG tube was inserted. The study documented widespread placement of PEG tubes in severely ill patients, half of whom were in the terminal phase of their illness. Further studies are needed to determine whether these patients benefit from PEG tube placement in terms of their quality of life and survival. Complications related to a PEG occur in over a third of patients and realistic expectations of what a PEG can accomplish should be a factor in the decision to place a PEG tube in nursing home patients.

A study of sixty-four patients receiving a PEG recorded that seven died within thirty days, four of complications of the PEG and three from the primary illness. ¹⁹ There were nineteen complications (32%), four wound complications occurred and nine patients developed aspiration pneumonia within three days of the procedure (four of whom died in hospital). Of the twenty-four patients with a history of aspiration, nine experienced aspiration during or after insertion of the PEG. Finally a retrospective comparison of long term enteral feeding by PEG compared to naso-enteric tubes has been done. ²⁰ This study concluded that a PEG offers no substantial advantages over a naso-enteric tube with respect to patient nutrition, performance or survival. Aspiration pneumonia occurred within fourteen days of tube placement in 6% of those with a PEG and 24% of patients with a naso-enteric tube.

In summary, it is our opinion that patients must be much more carefully selected for the insertion of a PEG to establish enteral nutrition. It would appear that some of the patients in whom a PEG was inserted had a very short life expectancy and maybe these tubes were used inappropriately. We should also repeat that the placement of a PEG does not protect that patient from aspiration of gastric contents.

Laparoscopic general surgery

Key messages

- Senior surgeons or surgeons under supervision performed all the procedures.
- There is at present difficulty in knowing whether there is higher mortality following a laparoscopic procedure when compared to an open operation.

Written by Mr Brian I Rees FRCS and Mr Ron Hoile MS, FRCS in consultation with the advisors in minimally invasive surgery (see page 11)

Deaths following laparoscopic cholecystectomy and hernia repair were of special interest. Details of some cases are given below together with the estimated experience (as stated by the surgeon). Laparoscopic abdominal surgery is physiologically challenging for the patient and, whilst it may be preferable to open surgery for patients with lung disease, it may not be advisable in patients with cardiac disease. However, on occasions surgery is necessary despite all efforts to avoid it. The fifth patient described below (with a mitral valve replacement) was a known operative risk, and was managed conservatively for five years, but developed an empyema of the gall bladder and an intra-abdominal abscess. Surgery was required, complications developed and, despite a second laparotomy, the patient died. The intention to attempt laparoscopic surgery would seem irrelevant in this case.

Age	Indication	ASA	Comorbidity	Procedure	Grade of surgeon and experience	Length of operation	Death
43	Gallstones. Biliary colic	2	Downs syndrome	Laparoscopic cholecystectomy, converted to open procedure	Consultant. Does 60/year	90 mins	Day 1. Cardiac arrest
61	Gallstones. Recurrent cholecystitis	1	None	Laparoscopic cholecystectomy	Visiting SpR (supervised). Does 12/year	80 mins	20 hours postop. CO ₂ embolus – confirmed at postmortem
63	Gallstones	3	Hepatic cirrhosis, cardio- respiratory disease	Laparoscopic cholecystectomy	Consultant. Does 60-70/year	55 mins	Day 4. CVA and ischaemic heart disease
63	Gallstones and jaundice. ERCP preoperatively	3	Diabetes mellitus. Ischaemic heart disease and atrial fibrillation. On Warfarin.	Laparoscopic cholecystectomy	Visiting specialist registrar. Does 50/year	45 mins	Day 15. Cardiac failure

	Gallstones. Empyema of gallbladder	3	Mitral valve replacement and Warfarin therapy. Rheumatoid arthritis. Previous ERCP	Laparoscopic cholecystectomy (converted). Open cholecystectomy, Roux-en-Y repair of common hepatic duct. Further laparotomy for bleeding.	Consultant. Does 60/year	10 hours	Day 7. Multiorgan failure
70	Gallstones. Cholecystitis	3	Ischaemic heart disease. Previous abdominal aortic aneurysm repair	Laparoscopic cholecystectomy	Consultant. Does 50/year	60 mins	Day 21. Cardiac failure
70	Gallstones. Biliary colic	3	Cardiac and musculoskeletal problems	Laparoscopic cholecystectomy	Consultant. Does 80/year	40 mins	Day 8. Pulmonary embolus (at home)
75	Gallstones. Cholecystitis	2	Hypertension. Temporal arteritis	Laparoscopic cholecystectomy	Specialist registrar year 4. Experience not reported	40 mins	Day 9. Pulmonary embolus
75	Gallstones. Biliary colic	3	Heart disease. On Warfarin and Digoxin. Initially refused surgery	Laparoscopic cholecystectomy	Consultant. Does 40/year	95 mins	Day 2. Myocardial infarct
76	Gallstones. Biliary colic	4	Ischaemic heart disease. Previous coronary angioplasty. Advanced breast cancer.	Laparoscopic cholecystectomy and operative cholangiogram	Consultant. Does 120/year	80 mins	Day 18. Myocardial infarct
77	Gallstones. Biliary colic	2	None	Laparoscopic cholecystectomy complicated by bleeding (not converted). Transfused two units	Specialist registrar. Does 10/year	90 mins	Day 1. Myocardial infarct
80	Bilateral inguinal hernias	2	Scrotal hydrocoeles	Laparoscopic hernia repair (bilateral) and scrotal surgery to repair hydrocoeles	Consultant. Does 100/year	75 mins	Day 2. Cardiac arrest

Age	Indication	ASA	Comorbidity	Procedure	Grade of surgeon and experience	Length of operation	Death
80	Bilateral recurrent inguinal hernias	3	Ischaemic heart disease. Previous TURP	Laparoscopic hernia repair (bilateral)	Consultant. (Experience not recorded)	75 mins	Day 11. Myocardial infarct
82	Gallstones. Cholecystitis	2	None	Laparoscopic cholecystectomy and open repair of inguinal hernia	Consultant. Does 25/year	90 mins	Day 26. Pulmonary embolus.
84	Gallstones. Three previous ERCPs – no stones seen at latest examination	1	None	Laparoscopic cholecystectomy	Specialist registrar year 3. Does 25/year	75 mins	Day 21. Myocardial infarct. Bleeding DU (inoperable in view of MI)
85	Inguinal hernia	3	Cardiac failure. On Warfarin	Laparoscopic hernia repair and division of adhesions	Consultant. Does more than 100/year	110 mins	Day 1. Myocardial infarct

Laparoscopic cholecystectomy is now an accepted procedure for the treatment of gallstones despite a paucity of prospective randomised controlled trials (due to the ethical problems in the constitution of such a trial). If death is a reasonable end-point for evaluation, any audit of laparoscopic data needs to be against an historical audit of open cholecystectomies prior to 1990. Unfortunately mortality figures for open cholecystectomy are few. The relevant study on open cholecystectomy²¹ compares the complications (including mortality) between two periods, 1990-91 and 1994. The overall mortality during 1990-91 (in England and Wales) was 0.7 % (55 deaths out of 8035 procedures) as compared to 0.5% (5 out of 1019 procedures) in 1994. Pneumoperitoneum with carbon dioxide gas produces a physiological challenge to the cardiovascular system (see commentary on anaesthesia for laparoscopic surgery, pages 40 to 42). This challenge and the subsequent physiological changes are immediate and subsequent prolonged surgery is not contributory to death or increased morbidity.²² There should be judicious invasive monitoring and careful interpretation of the findings in patients in ASA grades 3 to 4.

There is a higher mortality following laparoscopic cholecystectomy in patients who have a compromised cardiorespiratory system. ²³ One might justifiably argue that careful case selection, restricting laparoscopy to patients in ASA grades 1 to 2 would reduce mortality. This policy would also raise questions on the use of a cut-off age limit. There were no deaths in a series of 411 laparoscopic cholecystectomies in a group with a reported median age of 52 years (range 40-64 years). ²⁴ Acute cholecystitis was not a contraindication and the duration of surgery had no relation to the postoperative morbidity in this group. Absence of mortality in that study was in all probability related to the lower age but the study represents the results in a group of selected patients.

We have listed above the details of sixteen patients who died, thirteen following laparoscopic gall bladder operations and three following laparoscopic hernia repairs. The median age in both groups was similar at 82 years. Indications for surgery in the cholecystectomy group were biliary colic or inflammatory episodes. Of the thirteen deaths in the cholecystectomy group, three were due to pulmonary embolus, nine due to cardiac causes and a carbon dioxide embolus caused one death. It is unclear whether routine thromboembolic prophylaxis should be used during laparoscopic surgery but the number of pulmonary emboli suggest that we should consider such a policy. The case for establishing the use of routine thromboembolic prophylaxis during laparoscopic surgery will require further research and/or analysis.

In this sample of deaths after gall bladder surgery, the advisors were reassured to find that:

- indications for surgery were appropriate
- senior surgeons or surgeons under supervision performed all the procedures.
- grades of operator did not influence the outcome
- deaths were entirely due to cardiopulmonary problems

There is at present difficulty in knowing whether there is higher mortality following a laparoscopic procedure when compared to an open operation. There is some indication (on comparison of historical data) of a higher mortality if laparoscopy is performed for all age groups and all ASA grades. One may argue in favour of restricting the procedure to those who have no pre-existing cardiopulmonary problems (ASA 1 to 2) and to those aged less than 70 years. Such a policy would require further evaluation on a prospective basis. Applying such criteria would automatically have excluded the patients in this sample who presented with hernia.

Oesophageal surgery

Key messages

- Oesophageal surgery requires teamwork supported by the availability of a full range of treatment modalities and critical care services.
- Risk stratification using a system such as POSSUM might identify poor risk cases.
- Centres offering superspecialised services are required for the management of some oesophageal diseases.
- Timing of surgery after chemotherapy is crucial, as some patients were clearly immunosuppressed at the time of surgery.
- Bleeding oesophageal varices were poorly managed.
- Attention to postoperative respiratory care was essential to achieve a good outcome.
- There were few postmortem examinations and when done surgical attendance was poor.
- In some cases surgeons showed a lack of insight and a poor knowledge of thoracic surgical anatomy.

Written by Mr Ron Hoile MS, FRCS in consultation with the oesophageal surgery advisors (see page 11)

There were 273 questionnaires received concerning patients who died following oesophageal procedures. The definition of this group was such that endoscopic procedures were included alongside major resections. The full list of procedures is given below and includes multiple procedures for individual patients.

List of procedures

Malignant disease

Procedure (may be multiple for individual patients)	Number
Oesophagogastroduodenoscopy (including biopsy and dilatation)	57
Thoracoabdominal oesophagogastrectomy	56
Oesophagogastroduodenoscopy and intubation/stenting of tumour	32
Transabdominal oesophagogastrectomy	9
Transhiatal oesophagectomy	7
Laparotomy and open intubation of oesophageal tumour	7
Feeding jejunostomy	7
Bronchoscopy	6
Splenectomy	6
Three-stage oesophagectomy (cervical anastomosis)	5
Insertion of percutaneous endoscopic gastrostomy	5
Oesophagoscopy and laser destruction of tumour	4
Cervical oesophagostomy (anastomotic breakdown)	4
Tracheostomy	3
Rigid oesophagoscopy	3
Minitracheostomy	2
Palliative bypass for oesophageal tumour	2
Resection of necrotic jejunal loop	2
"Open and shut" laparotomy	2
Miscellaneous (one each of: staging laparoscopy, ligation of thoracic duct, drainage of mediastinum, open removal of oesophageal tube, transthoracic oesophagectomy, feeding gastrostomy, laparotomy	7
for lymphatic leak)	

Benign disease

Procedure (may be multiple for individual patients)	Number
Oesophagogastroduodenoscopy	52
Oesophageal transection for bleeding varices	9
Oesophagogastroduodenoscopy and injection of oesophageal varices	8
Oesophagogastrectomy (rupture or bleeding)	6
Feeding jejunostomy	4
Repair of oesophagus (spontaneous rupture)	2
Repair of iatrogenic perforation of oesophagus	2
Miscellaneous (one each of: diagnostic laparotomy, partial gastrectomy, right hemicolectomy, sigmoid colectomy, laparotomy for postoperative bleeding, drainage of cervical abscess, vagotomy and pyloroplasty, drainage of empyema, revision of gastroenterostomy, laparotomy and repair of Mallory-Weiss tear, splenectomy)	11

These patients were under the care of a variety of surgeons, mostly general surgeons, who professed many differing interests. Of the 217 deaths under the care of general surgeons, there were only five cases where the surgeon specifically identified an interest in oesophageal surgery. Thoracic surgeons cared for 52 patients, a specialty where there is usually extensive experience of oesophageal surgery. In most cases the operating surgeon was a consultant (81%, 222/273) or a consultant was the most senior person in the operating theatre (87%, 239/273). In a few instances the grade of surgeon was not appropriate and comment on these cases appears below. Overall, consultants were involved in the care of 264/273 patients.

The management of patients with oesophageal disease (the majority of deaths occurred in patients with malignant disease) requires special knowledge, understanding and skills. This applies not only to the surgeon and anaesthetist but also to the large number of professions who support them and form part of the team (such as physicians, intensivists, nurses, physiotherapists, nutritional experts). A successful outcome for the patient depends on the provision of this team and the ability to offer a full range of treatments, supportive therapies and backup. Comment on the facilities for the specialist practice of oesophageal surgery appears below on page 65.

Questions arose when the details of the patients who died were reviewed. These included:

- What is required for the basic assessment of patients and their disease prior to major oesophageal surgery and how can this information be linked to a more holistic approach and an attempt at risk stratification?
- Can we improve case selection for surgery based on this assessment?
- How should anaesthesia for oesophageal surgery be conducted?
- How can the number of inappropriate, poorly conceived and often unorthodox operations be reduced?
- What is the current role of oesophageal transection for bleeding oesophageal varices?
- How should a department offering a surgical service for the treatment of oesophageal disease be staffed and organised?
- Is there any place for the open insertion of oesophageal tubes or stents?

These questions were prompted during the review of specific cases. Examples of the defective delivery of care are given below followed by broader discussion of some specific issues addressing the questions above.

Inappropriate operations

There were examples of inappropriate operations, often linked to poor decision-making in general. Some cases involved incorrect choice of procedure due to lack of experience or the inability to provide a wide range of procedures. In other instances there were illogical and unorthodox approaches to a problem or failure to acknowledge the frailty of the patient. Other cases demonstrated an incorrect choice of general anaesthesia for endoscopic procedures and, overall, there was a general absence of an holistic approach.

Age	Operation	Problems	Comments by the advisors
36	Oesophagectomy (achalasia)	Complicated case following multiple procedures. Died from haemorrhage on day of operation	Surgical plan questionable. General surgeon operating when superspecialist needed. Knowledge of thoracic anatomy lacking.
52	Oesophagogastrectomy.	Severe cachexia ASA 3.	Technical error using feeding gastrostomy for inadequate time preop. No staging or assessment. Not a suitable case for surgery.
58	Oesophagogastroduodenoscopy ?gastric volvulus. Later right hemicolectomy.	Died from multiorgan failure 19 days after laparotomy.	Inappropriate procedure, delayed diagnosis.
64	Initially three-stage oesophagectomy and splenectomy. Final procedure was feeding jejunostomy.	Recurrent tumour in upper third after irradiation. COPD. 17 days in ICU. Anastomotic leak. Died two days after jejunostomy – carcinomatosis.	Initial resection through tumour (operation seems illogical in view of previous assessment), technique described unorthodox. Decision to feed too late.
66	Oesophageal transection for bleeding varices.	COPD, IHD, hepatic and renal failure.	Patient was unlikely to survive.
67	Thoracoabdominal oesophagogastrectomy.	Incomplete resection. MRSA sepsis.	Gross disease remained. Better to abandon procedure.
70	Laparotomy to reposition oesophageal tube.		Endoscopic expertise to change tube not available.
71	Oesophagogastrectomy and division of pharyngeal web	Inadequate preoperative assessment. Incomplete resection.	Gross disease remained. Better to abandon procedure.
72	Resection of necrotic jejunal graft.	Oesophagogastrectomy. Free jejunal graft for reconstruction became necrotic. Resection margins positive.	First operation inappropriate and inadequate. 48-hour delay after leak recognised. ASA 5 i.e. moribund.
76	Attempt to intubate recurrent tumour at anastomosis two years after transhiatal oesophagectomy.	Cachectic (wt 30kg) and breathless. Cardiac arrest on table.	Poor case selection and unorthodox technique.
85	Laparotomy for open insertion of Mousseau-Barbin tube. Abandoned.	Elderly patient with extensive tumour. Died day five –carcinomatosis.	Outmoded inappropriate management. Palliation should be possible without open surgery.
86	Palliative bypass lower oesophagus via thoracotomy.	Previous total cystectomy. Died first postoperative day.	No attempt at endoscopic intubation. Careful thought needed in patients aged over 80.
88	Repair of iatrogenic oesophageal perforation.	Tear caused by endoscopic attempts to remove large gallstone impacted in pylorus. Death two weeks later.	Ill-advised endoscopic procedure.

Poor decision-making regarding intubation

Age	Procedure	Problems	Comments by the advisors
68	Oesophagogastroduodenoscopy and insertion of stent	Known liver metastases. Died six days after procedure.	Was symptom control more appropriate?
71	Laparotomy and insertion of Mousseau-Barbin tube.	Iatrogenic perforation of distal squamous carcinoma. Died of asphyxiation six days later when tube dislodged.	Fourth year specialist registrar operating without supervision. Postmortem examination showed operable lesion. Inappropriate decision to intubate a fit patient without proper assessment of stage of disease. No senior help available.
76	Oesophagogastroduodenoscopy and insertion of nasogastric feeding tube.	Repeated endoscopic laser treatment of tumour. Tracheo-oesophageal fistula.	Insertion of feeding tube delayed for 26 days. A stent might have been more appropriate.
80	Repositioning of oesophageal tube under general anaesthesia.	Very frail, ASA 4. Died 11 days later – carcinomatosis.	Why give a general anaesthetic? Was procedure appropriate for terminally ill patient?
80	Open insertion of oesophageal tube for stricture.	Large preoperative haematemesis, anaemia, thrombocytopenia and hypoalbuminaemia.	No mention that stricture was impassable. Was endoscopic treatment considered?
91	Oesophageal dilatation and insertion of tube.	Diagnosis made eight months previously. Difficult procedure. Died 15 days later.	Why give a general anaesthetic? Survival was expected to be short. Was this procedure necessary?

Case selection

Some circumspection might help surgeons identify those patients who were clearly not going to benefit from the procedure at the time it took place.

Age	Procedure	Problems	Comments by the advisors
37	Oesophagogastroduodenoscopy.	Perforated oesophagus. Died 14 days postoperatively	Young patient with vomiting and weight loss. Benign disease likely but inadequate work-up.
64	Laparotomy for lymphatic leak following oesophagogastrectomy.	Known cirrhotic. Died four days after surgery. Multiorgan failure.	Ascites not leak. Underestimated comorbidity.
66	Oesophagogastrectomy.	Bronchopneumonia and hypokalaemia preoperatively. Died 13 days postoperatively.	Cause of death was bronchopneumonia. This patient was not a suitable candidate for this operation.
71	Bronchoscopy and minitracheostomy following oesophagogastrectomy.	No tissue diagnosis before original operation. Extensive comorbidity with cardiorespiratory disease. Left lung collapse.	Death due to respiratory failure. Extent of comorbidity appeared to be contraindication to surgery.

Age	Procedure	Problems	Comments by the advisors
79	Ivor-Lewis oesophagogastrectomy.	Tumour inadequately staged and found to invade left bronchus and trachea. Resection palliative.	Poor preoperative assessment and subsequent judgement. Thoracotomy should have been "open and close".
83	Ivor-Lewis oesophagogastrectomy and splenectomy.	Thrombocytopenia, hypertension, ischaemic heart disease and aortic aneurysm. Anaesthetic assessment one hour before surgery.	Died from haemorrhage 24 hours after surgery. No cause identified clinically and no postmortem examination.
85	Oesophageal dilatation & insertion of percutaneous endoscopic gastrostomy.	Third attempt at endoscopy and dilatation in demented frail patient. Radiotherapy apparently considered.	Died with bronchopneumonia three days after surgery.

Inadequate preoperative assessment and preparation

Age	Procedure	Problems	Comments by the advisors
69	Oesophagogastrectomy.	Poor preoperative assessment in patient with ischaemic heart disease, thoracic aneurysm, hypertension, TIAs and peripheral vascular disease.	Disease understaged and resection margins involved with tumour. Subsequent anastomotic breakdown. Inadequate operation by occasional oesophageal surgeon.
70	Oesophagogastrectomy.	No antibiotic prophylaxis. Fast atrial fibrillation postoperatively due to anastomotic breakdown.	No preoperative assessment apparent from questionnaire. Inadequate management of surgical complication. No postmortem examinations. Implications ignored for six days.
78	Oesophagogastroduodenoscopy and injection bleeding duodenal ulcer.	Patient with dense hemiplegia, Hb 6.5 g/dl and no transfusion.	Lack of initial resuscitation and inappropriate endoscopy by inexperienced vascular surgeon. Died after rebleed at three days.

Lack of ICU beds or early discharge from ICU

Age	Procedure	Problems	Comments by the advisors
59	Transhiatal oesophagectomy.	High risk case with COPD, rheumatoid arthritis and peripheral vascular disease.	Surgery commenced without ICU bed available. Should procedure have been started? Died with respiratory failure.
67	Oesophagogastrectomy.	Preoperative chemotherapy.	ICU full so nursed on general ward. Chylothorax and anastomotic breakdown. Failed to reoperate when complications recognised.

Age	Procedure	Problems	Comments by the advisors
72	Oesophagogastrectomy.	Initially fine but discharged from ICU early because of pressure on beds. Chest infection with pseudomonas and ARDS. Readmitted to ICU on day seven after bronchoscopy.	Died on day 13 from respiratory failure.
74	Oesophagogastrectomy.	Fit patient not admitted to ICU until late stages of respiratory failure.	No preparation, no ICU, inappropriate anaesthetist, and surgeon had never done this procedure before. Died on day nine from cardiorespiratory failure.
75	Oesophagogastrectomy.	Known ischaemic heart disease. ECG changes noted during surgery.	No ICU bed therefore nursed in theatre postoperatively. Invasive monitoring inserted postoperatively. Died in theatre – myocardial infarction.

Problems with the management of oesophageal varices

Age	Procedure	Problems	Comments by the advisors
40	Laparotomy and insertion of Minnesota tube	Massive bleeding from varices with known metastases in liver (previous breast carcinoma).	Heroic and inappropriate surgery in terminally ill patient.
55	Oesophagogastroduodenoscopy	Massive bleeding from varices.	Research registrar operating. No attempt at therapeutic procedures.
57	Oesophageal transection.	Coagulopathy and hepatic failure. No attempt at endoscopic therapy.	General surgeon with an interest in breast surgery had only performed this procedure once before.
66	Laparotomy, stapling of oesophagus and gastrostomy.	Attempt to underrun bleeding vessels failed. Oesophagus divided and stapled off above bleeding varices. Died on day two from continued bleeding.	Wrong operation. No postmortem examination. Why not refer to specialist centre?
66	Oesophageal transection for bleeding varices.	COPD, ischaemic heart disease, renal and hepatic failure. Died on day four with multi-organ failure.	This patient was unlikely to survive.
69	Oesophageal transection.	Bleeding varices in patient with lobar pneumonia, cardiac and renal failure.	Basic trainee anaesthetist. Died on operating table. Inappropriate operation.
72	Oesophageal transection & splenectomy.	Bleeding varices and coagulopathy. No blood products given to correct deficiencies. No attempt at endoscopic treatment prior to surgery.	Inadequate treatment. Surgeon had never done procedure before and anaesthetist was very junior basic trainee.

General problems

Age	Procedure	Problems	Comments by the advisors
55	Oesophagogastrectomy.	Ankylosing spondylitis.	Poor fluid management with overload.
72	Total gastrectomy.	Preoperative chemotherapy. Anastomotic and duodenal stump leak.	Inexperienced registrar operating; not assisted by consultant.
74	Oesophagogastrectomy.	Adenocarcinoma in Barrett's oesophagus. Preoperative chemotherapy.	Surgical mishap. Splenectomy. Blood loss six litres. Poor surgery and poor management of blood replacement. Sepsis and multi-organ failure.
75	Oesophagogastrectomy.	COPD & CCF. Poor respiratory function measured preoperatively. Never came off ventilator postoperatively.	Inappropriate grade of surgeon and counselling/consent questionable.

Reoperation for early complications

Sometimes complications occurred, were recognised but then corrective surgery was either delayed or did not occur. Examples are included in other sections above where, often, multiple failures of management occurred.

Management of patients undergoing oesophagectomy

Key messages

- Preoperative resuscitation was often inadequate after oesophageal rupture.
- Monitoring of direct arterial pressure, central venous pressure, urine output and temperature is routine
 practice; in selected cases a pulmonary artery catheter may be useful.
- Anaesthetists using one lung ventilation (OLV) should ensure that they have appropriate training and experience in this technique
- Chest infection, from whatever cause, is a major precursor of sepsis and death. This is an area of potential improvement.

Written by Dr Kathy Sherry FRCA in consultation with surgical and anaesthetic advisors (see pages 10 and 11)

Hospital Episode Statistics⁷ report that about 2000 oesophagectomies are performed annually in England and in the last five years the death rate has been between 9.5-10.5%. From the 1996/7 sample NCEPOD received 78 completed anaesthetic questionnaires for review. Most oesophagectomies were done "in-hours" and by senior anaesthetists.

Grade of the most senior anaesthetist (working alone)

Consultant	67	(20)
Associate specialist	1	(1)
Staff grade	3	(2)
Specialist registrar year 4	4	(2)
Specialist registrar year 3	2	(1)
Specialist registrar year 2	1	(1)

Age of patient (years)

<50	1
50 - 59	11
60 - 69	29
70 – 79	35
>79	2

Forty-three (55%) patients had concurrent disease. Twenty-nine (37%) patients had cardiac disease, five had respiratory disease and a further nine had both cardiac and respiratory disease.

Oesophageal rupture

Five patients underwent surgery for oesophageal rupture; all were in poor condition and "likely to die". These included a 65-year-old patient with an oesophageal rupture between 10 and 22 days previously who was septic, on positive pressure ventilation and receiving inotropic support, and an 85-year-old patient whose oesophagus had ruptured 15 days previously, and who was initially treated conservatively and was receiving intravenous fluids. The other three, despite their poor condition, had not received any form of resuscitation preoperatively. Two were anaemic (Hb 9.3g.dl⁻¹) and one had a Hb 17.2g.dl⁻¹, bicarbonate 17m.mol.l⁻¹, pH 7.18 and glucose 12.3m.mol.l⁻¹. Spontaneous oesophageal rupture is infrequent and its

diagnosis is sometimes delayed. These patients require preoperative resuscitation in the same way as a similarly dehydrated patient with intra-abdominal perforation.

Operative monitoring

Standard operative monitoring included direct arterial pressure, central venous pressure, urine output and temperature. In 45 (58%) patients, all four variables were monitored. Direct arterial pressure was monitored in 73 (94%), CVP in 69 (88%) and urine output in 73 (94%) patients, temperature was monitored in only 51 (65%) of cases. One patient, a 74-year-old man with severe cardiac disease and anaesthetised by a specialist registrar year 4, had none of these. His trachea was extubated following surgery. His temperature was 34°C in the recovery area where he was given neostigmine, doxapram and naloxone before he was sent to the general ward. Two patients had only the direct arterial monitoring, one anaesthetised by a consultant and one by a specialist registrar year 4. Of the patients reviewed none had a pulmonary artery catheter inserted. In this group of elderly patients with a high incidence of cardiac disease and undergoing major surgery including large fluid fluxes a pulmonary artery catheter can be useful. It requires practice to insert and interpret. However, it allows a more accurate assessment of fluid loading and the facility to optimise the cardiac output.

Double lumen tube

One-lung ventilation (OLV) is associated with several well-recognised complications. Anaesthetists using this technique should ensure that they have appropriate training and experience to use this technique safely.

In 66 (86%) patients a double lumen tube was used or its placement attempted. In 30% of these cases problems were encountered. These are presented in the accompanying table.

Thirteen (20%) of the 66 patients suffered arterial desaturation at some time during one-lung ventilation; in some, high airway pressures accompanied this. The two main causes of arterial desaturation during one-lung ventilation are an incorrectly positioned double lumen tube and intrapulmonary shunting of blood.

Correct placement of a double lumen tube requires practice and is more likely to be achieved by the anaesthetists who use them regularly. For some anaesthetists, when working with a general surgeon who has a small oesophageal practice, this experience is acquired slowly. No anaesthetist reported using a fibreoptic bronchoscope to confirm the position of the tube, either before or during surgery, even when the tube was evidently incorrectly placed. The fibreoptic bronchoscope is now widely available and with practice it can aid correct placement and verify the tube position. In four out of the five cases (numbers 1-5 in the table below) when the experienced anaesthetist recognised poor positioning *before* surgery an alternative, more satisfactory, tube was used. A trainee who did not try an alternative tube managed the fifth case. *During* surgery, when it was suspected that arterial desaturation was related to the tube position, there was a reluctance to change to an alternative tube. In some cases the anaesthetist was prepared to tolerate the unsatisfactory situation.

The management of arterial desaturation due to intrapulmonary shunting forms part of the basic training in one-lung anaesthesia. The response should be to increase inspired oxygen, start non-dependent lung CPAP, differential dependent lung PEEP with non-dependent lung CPAP, intermittent lung inflation and, when these fail, resuming two-lung ventilation, finally returning to the supine position. Dependent lung PEEP alone has been shown to be of little use. Of these manoeuvres only the increase of inspired oxygen and two-lung ventilation were reported as having been used.

Whatever the cause, if arterial desaturation or inadequate ventilation occurs during anaesthesia it cannot be ignored. The immediate action to correct may need to take priority over surgical considerations and the surgery interupted.

In two patients haemodynamic instability was reported. In one (case 20) it was related to surgical manipulation around the mediastinum; the hypotension and arrhythmias were allowed to continue uncorrected. During oesophageal surgery a surgical cause of haemodynamic instability is well recognised and it was commented on in several questionnaires. The severity can be limited by the co-operation of the informed surgeon, and appropriate treatment to optimise the circulation.

Case	Problem	Action	Resolution
1	Larynx too small to admit double- lumen tube.	Changed to orotracheal tube	No further problems
2	Failed to place a Robertshaw tube	Changed for a Carlens tube	Good position.
3	Failure to ventilate either lung through the double lumen tube during transfer from the anaesthetic room to theatres. Then, failure to collapse L lung during surgery	Tube replaced twice for these problems. Still could not oxygenate on one-lung ventilation	Finally ventilated on two lungs (TLV) by the third double-lumen tube
4	Failed to place a L side double- lumen tube	R side double-lumen tube placed	Good position
5	Failed to place a L side Robertshaw tube	R side Robertshaw tube placed	Good position
6	Arterial desaturation during OLV	None	Resolved when TLV was resumed
7	Arterial desaturation during OLV	None	Arterial saturation 88-90% for one hour
8	Double-lumen tube via tracheostome, high airway pressures on OLV.	None	PaCO ₂ 11.2kPa on blood gases which were taken two hours later
9	Arterial desaturation during OLV	None	Tube changed for orotracheal tube between the abdominal and thoracic stages
10	Arterial desaturation during OLV	Tube repositioned	Continued hypoxia (6.5kPa) and hypercarbia (10kPa), pH 7.16, finally developed fulminating pulmonary oedema
11	Arterial desaturation during OLV	Tube repositioned	Resolved
12	Arterial desaturation and high airway pressures some time after starting OLV, bronchospasm diagnosed	Nebulised salbutamol	Resolved
13	Arterial desaturation and hypotension during OLV	Reverted to TLV	Resolved
14	Arterial desaturation during OLV	Increased FiO ₂	Arterial saturation increased to 92% during OLV
15	Arterial desaturation and high airway pressures during OLV	Reverted to TLV	Resolved.
16	Arterial desaturation during OLV	PEEP applied to the dependent lung without accompanying CPAP on the non-dependent lung	Poor documentation but arterial saturation 90% for four to six hours. No blood gases recorded

(continued overleaf)

Case	Problem	Action	Resolution
17	Arterial desaturation during OLV to 78%	FiO ₂ increased to 1.0	On FiO ₂ 0.8, saturation 90-95% for 1½hrs.
18	Arterial desaturation (71%) 25 minutes after starting OLV	After 15 minutes of severe arterial desaturation resumed TLV.	Arterial saturation stayed at 88-92% despite trying to reposition the tube. Postoperative X-ray showed whiteout of a lung
19	Hypotension and bradycardia during OLV	Atropine	Resolved
20	Obstruction to ventilation during surgical manipulation	None	11/4 hr of hypotension and arrhythmias

Postoperative anaesthetic management

Thirty-two (42%) of the 78 patients had their trachea extubated immediately postoperatively. Two of these 32 were suffering respiratory inadequacy and had their trachea reintubated straight away. Forty-four (58%) of the 78 patients were ventilated, either electively or because of hypoxaemia or unstable haemodynamics. Two patients died in theatre.

Fifty-eight (74%) of the 78 patients had epidural analgesia postoperatively.

Postoperative sepsis

Fifty-three (68%) patients died of sepsis and multiple organ failure. In 36 (46%) patients the source of sepsis was a chest infection, fifteen had an anastomotic leak and two had abdominal infections. Only seven patients died primarily from a cardiac event.

It is of concern that oesophagectomy is associated with a high mortality and that the main cause of death is systemic sepsis, secondary to respiratory complications or anastomotic breakdown. Amongst these patients NCEPOD could not detect obvious common precipitating causes. However, we can suggest factors that may improve outcome.

Preoperative precipitating factors

Compromised preoperative immunity. Some patients in this sample had evidence of preoperative poor nutritional state, often as a result of their swallowing difficulties but two had documented alcoholism. Others had undergone a course of chemotherapy shortly before surgery. It is advisable to leave an interval after chemotherapy so that the immune system can recover and the white cell and platelet count return to normal.

Preoperative respiratory function. There is a high incidence of smoking amongst patients with oesophageal carcinoma, thus a high incidence of chronic obstructive pulmonary disease. In these patients a transhiatal approach for carcinomas of the lower oesophagus which are surgically accessible by this incision has been advised. There is also an incidence of preoperative respiratory infection consequent upon over-spill and, in these patients, a few days of preoperative chest physiotherapy combined with appropriate antibiotics may well influence outcome.

Operative and postoperative factors

Respiratory system. One-lung ventilation, intrathoracic surgery and a misplaced double lumen tracheal tube can all lead to areas of lung collapse and so increase the potential for postoperative chest infection. To reduce this problem:

- correct double lumen tube placement is essential and if doubt exists the position should be re-confirmed with a fibrescope.
- surgical manipulation and retraction of the non-dependent lung should be minimal and gentle in order to limit lung trauma.
- continuous positive airway pressure on the non-dependent lung may reduce atelectasis and it does reduce intrapulmonary shunting and hypoxaemia.
- the lung should be fully re-expanded on chest closure.

Postoperative respiratory care hinges on chest physiotherapy. Physiotherapists should be aware of the high incidence of postoperative life-threatening respiratory infection following oesophagectomy. They have a pivotal role in the patient's treatment, both pre- and postoperatively. Would physiotherapists consider the management of patients following oesophagectomy suitable for locally devised guidelines? It is recognised that good postoperative analgesia facilitates chest physiotherapy and good chest expansion. Epidural analgesia was the most popular form of postoperative pain control.

When chest infection or sepsis is diagnosed the early involvement of the microbiology department is likely to produce antibiotic regimens closely targeted at the causative organism(s).

Cardiovascular system. Oesophagectomy is a major operation involving large fluid fluxes that continue into the postoperative period. During mobilisation of their oesophagus patients often have arrhythmias provoked by surgical manipulation and the haemodynamic instability may be compounded by poor management of epidurals peroperatively. For these reasons patients are at risk of inappropriate fluid loading (under- or over-filling) and impaired cardiac output; those with pre-existing cardiovascular disease are especially vulnerable. Low cardiac output can potentially reduce oxygen delivery to and adversely affect the oesophago-gastric anastomosis, the oxygenation of which has already been compromised by vascular dissection and manipulation. Arterial pressure and central venous pressure monitoring can give no information on cardiac output or ventricular filling. It is therefore surprising that none of these patients had a pulmonary artery catheter inserted peroperatively to facilitate monitoring during surgery or the early postoperative period. Pulmonary artery catheters were only inserted in those patients who had developed established sepsis, in general several days postoperatively.

Conclusion

There is a clear need to reduce the incidence of postoperative sepsis, chest infection and anastomotic breakdown. To achieve this, more research is required in order to predict the vulnerable patients, and to determine the most appropriate management by all those involved in the patient's care.

No patient with an active chest infection should be submitted to surgery until antibiotic therapy and physiotherapy have cleared the infection. Patients with an FEV1 of less than one litre are unlikely to survive or benefit from a radical oesophageal resection as the interstitial oedema resulting from disruption of the pulmonary lymphatics, especially in those with a low serum albumin, further compromises a precarious respiratory status. Rarely should a patient aged 80 or over be subjected to radical resection in view of the steep rise in mortality with extremes of age.

No patient who is unfit to withstand a major surgical assault should be subjected to radical surgery and no surgeon with inadequate training (in all aspects of oesophageal surgery including preoperative assessment, intraoperative management and postoperative care) should undertake the care of such patients independently. A well conducted, patient-tailored operation supported by skilled anaesthesia, including invasive monitoring of haemodynamic and respiratory parameters, will produce a better outcome for patients undergoing major oesophageal resections.

Further reading

Ellis HJF et al. Oesophagogastrectomy for carcinoma of the oesophagus and cardia. Journal of Thoracic and Cardiovascular Surgery. 1997; 113:836-848

Proposed facilities for the specialist practice of oesophageal surgery

Key messages

A multidisciplinary team within a specialist centre should manage patients with oesophageal disease.

Personal commentary written by Mr S Michael Griffin MD, FRCS

Introduction

The majority of the workload and use of resources in a department of oesophageal surgery will be in the management of oesophago-gastric cancer. A multidisciplinary team within a specialist centre providing all investigative and treatment modalities must manage patients with oesophageal cancer. The concentration of cases within such a unit would allow accurate audit of outcome and ensure appropriate recruitment to clinical trials. Such a unit would be sited within a cancer centre.

Multi-disciplinary team

The core members of the multidisciplinary team dealing with oesophageal conditions include:

- · oesophago-gastric surgeons including thoracic surgeons with expertise in oesophageal surgery
- specialist oesophageal anaesthetists
- diagnostic and interventional radiologists
- · clinical and medical oncologists
- specialised upper GI histopathologist
- upper GI nurse specialist and database manager
- palliative care specialist

Other members who are key to providing this service but not core staff include:

- specialist intensivist with experience in management of patients following oesophageal surgery
- medical gastroenterologist
- General Practitioner
- nutritionist
- · dedicated physiotherapist

The specialist centre would comprise expertise and specialisation as outlined above.

All investigative modalities must be available. These include:

- video endoscopy
- pulmonary physiology laboratory
- · oesophageal physiology laboratory
- computerised tomography (ideally spiral)
- endoluminal ultrasound

The specialist centre would require adequate inpatient facilities in terms of beds on a ward with specialist nurses, theatre time and experienced anaesthesia with ready access to intensive care and high dependency units.

Full back up with an identified interventional radiologist with suitable imaging including access to spiral CT on a 24-hour basis.

Surgeons practising in the centre must have specialist expertise in the field of oesophageal cancer. Surgeons should work in teams of at least two and preferably four to ensure continuity of care and to provide a consultant-based service throughout the year. These surgeons would devote all their fixed sessions to gastro-oesophageal surgery, which would include a service for benign diseases.

Upper GI nurse specialists would act as a focus for the patients referred to the centre and as a contact point in times of difficulty or anxiety. These nurse specialists would help to coordinate therapy, to liaise with all members of the team and to direct the collation of data for regular audit. Dietary problems occur in almost all patients postoperatively, especially whilst undergoing neo-adjuvant therapy. The upper GI nurse specialist and dietician are essential to diagnose these feeding problems and treat them early. The upper GI nurse specialist would form a vital link in primary care and would visit patients in the postoperative and post-therapeutic period.

The multi-disciplinary team would meet on a weekly basis to discuss all cases and to formulate management plans.

Most patients would present to smaller units where the diagnosis will be made. Protocols could be disseminated to ensure similar standards of diagnosis and urgent referral. All potentially curable oesophageal cancer patients should be referred to specialist centres. Upper GI surgeons working single-handed should not continue. They should be encouraged to be part of the central team with sessions in a specialist unit performing surgery. For maximum efficiency over 250 new cases of oesophago-gastric cancer should be assessed in such a centre. This would serve a population of at least one million.

Better investigative facilities and centralisation of surgical expertise would result in better preoperative assessment to enable patients to avoid unnecessary surgery and receive all appropriate treatment for their general health and the stage of their disease. Many more patients with T2 and T3 tumours will be considered for neo-adjuvant chemo-radiotherapy. These patients would be included in clinical trials. There is much evidence to support the role of the multidisciplinary team and the specialist centre in the management of oesophageal disease. Oesophago-gastric surgery carries the highest postoperative mortality of any elective surgery. A recent survey of the world literature³² showed an average mortality of 15%. Audited results from specialist centres in the United Kingdom have brought mortality figures down to below 5% despite the treatment of high risk patients.³³ In specialist centres the five-year survival figures for patients undergoing curative surgery for cancer is in the region of 25-30%, similar to the best figures in the world. Overall survival in the United Kingdom is well below this figure.

The complications of oesophageal surgery are significant. Over 30% of patients suffer major complications following thoraco-abdominal surgery. Consultant input and access on site to interventional radiology on a 24-hour basis is essential to treat these complications. A high level of manpower and expertise is essential to manage these patients effectively. This can only be achieved in a specialist centre with at least two and preferably four surgeons and two interventional radiologists.

The stage of the disease and its assessment preoperatively are crucially important in the management of oesophageal cancer. Spiral CT and endoluminal ultrasound are pivotal to this accurate assessment. These modalities are relatively expensive and endoluminal ultrasound is only available in a few large centres. Resources need to be concentrated in specialist centres so that there is no duplication resulting in financial wastage.

Multi-modality therapy, particularly including neo-adjuvant chemo-radiotherapy may well improve outcome in operative patients. The cost of radiotherapy equipment restricts this service to cancer centres where the specialist centre should be based. Furthermore the use of expensive palliative measures in those patients with oesophageal cancer not suitable for curative surgery should be concentrated in these centres and not distributed in smaller hospitals resulting in duplication of expensive equipment.

Setting up such centres would prevent the occasional surgeon performing less than five cases of oesophageal surgery or resections per year. Although specialist centres always raise the spectre of increased cost there would be potential savings by concentrating expensive equipment (spiral CT and endoscopic ultrasound) and resources (intensive care and high dependency units) on fewer sites. There would be a reduction in operative mortality. In addition the early recognition of complications and their treatment would reduce ICU stay and prevent a later transfer to a specialist unit with associated prolonged stay.

Oesophageal stenting

Key messages

- Surgeons, endoscopists and interventional radiologists may be involved in procedures to insert
 oesophageal stents. They should all be aware of the indications, risks and overall management of
 patients requiring this procedure.
- Patients should receive detailed information about life with a stent.

Personal commentary written by Mr Simon Raimes MD FRCS

There are now a variety of oesophageal prostheses that require different techniques of insertion. Surgeons, endoscopists and interventional radiologists may be involved individually or as teams in procedures to insert stents and should equally be aware of the indications, risks and overall management of the patient if mortality is to be minimised.

The primary indication for oesophageal stent insertion is relief of malignant dysphagia in patients who are unsuitable for, or have failed, other techniques to improve their swallowing. It is presently not regarded as first-line treatment except in very elderly or unfit patients. None of the presently available stents consistently allow completely normal swallowing and should not be used unless there is significant unrelieved dysphagia except in two circumstances. The first is in the treatment of an aero-digestive fistula involving the oesophagus and the other is the rare situation where there is uncontrolled blood loss from an oesophageal cancer.

Oesophageal stent insertion is a palliative procedure in patients expected to have a limited life span. It does not affect the natural history of the underlying disease although the resulting better nutrition and morale of the patient may improve survival. It is not a treatment for a patient in the last phase of a terminal illness where other palliative measures are more appropriate.

There are presently two types of stent in common use. Stents inserted at open operation are now only very rarely indicated. Rigid reinforced tube stents are inserted endoscopically using a pulsion technique. The newer self-expandable covered stents are inserted either endoscopically or under radiological control alone. Randomised trials are being conducted to determine the possible advantages of the self-expandable stents since they are five to six times more expensive to purchase. Whichever stent is used the procedure should be regarded as an operation and the patient and their relatives fully informed of the risks and likely benefits of the procedure. It is still not certain whether the significant mortality of rigid stent insertion is avoided by use of the self-expandable stents. High standards of care before, during and after the procedure are required to minimise the mortality.

Preoperative preparation should include correction of dehydration and electrolyte disturbances. Patients with complete dysphagia are usually significantly malnourished and often have chronic or acute aspiration into the respiratory tract. Antibiotics and chest physiotherapy are essential in preparing the patient with any evidence of infection.

Stenting should be performed under heavy intravenous sedation with a short-acting benzodiazepine combined with a short-acting opiate. There is no indication for the use of general anaesthesia but it is sensible for an anaesthetist to administer the sedation to these very sick patients. The heavily sedated patient must receive oxygen, have monitoring of oxygen saturation and ECG and have their airway cared for and protected until fully awake. Insertion of the stent and pre-dilatation, if required, must be performed under radiological control.

Many patients experience chest pain after stent insertion and may require strong analgesia for up to 72 hours. Oesophageal perforation during insertion is a recognised risk of these procedures and if suspected should be confirmed by clinical examination, an erect chest X-ray film and possibly a water-soluble contrast swallow. In most cases the tube will occlude the leak, but a short course of intravenous broad-spectrum antibiotics is advisable. If the clinical signs deteriorate or the leak is uncontained then a decision on further management has to be made in light of the overall condition of the patient. Stents that cross the gastro-oesophageal junction can produce significant problems. Uncontrolled reflux is a frequent problem and the recovering patient must be nursed in the sitting position to prevent aspiration. In the longer term prescription of an antisecretory drug and a prokinetic agent are important to decrease oesophageal damage and chronic aspiration. Stents in this position may impact on the greater curve of the stomach resulting in an ongoing inability to eat that can be very difficult to solve.

Patients should receive detailed information about life with a stent. They will also need expert dietetic advice. The patient and their family should be offered contact with appropriate cancer support services in view of the poor prognosis.

Further reading

Blazely JM, Alderson D. *Palliative treatments of carcinoma of the oesophagus and stomach*. In:Upper Gastrointestinal Surgery, eds. Griffin SM, Raimes SA. W B Saunders, London 1997 pp 213-244.

Sturgess RP, Morris AI. Metal stents in the oesophagus. Gut 1995; 37: 593-594.

Spinal surgery

Key messages

- There was a high consultant involvement in this group of patients and surgery was done by appropriate specialists in all cases.
- In benign disease, difficult revisional surgery for conditions such as scoliosis are best referred to centres with experience of this type of procedure and its difficulties.
- For surgery in metastatic disease there need to be some clear, well understood criteria for selection and for justifying the extent of the surgical approach.

Written by Mr Ron Hoile MS, FRCS in consultation with the spinal surgery advisors (see page 11)

List of procedures

Benign disease

Procedure (may be multiple for individual patients)	Number
Decompressive cervical surgery with or without fusion/stabilisation	14
Decompressive lumbar surgery with or without fusion/stabilisation	9
Cervical fusion for fracture/dislocation and/or instability (all levels)	8
Application of halo traction vest	4
Transoral odontoidectomy	2
Debridement of thoracolumbar vertebrae for osteomyelitis (pyogenic 1, tuberculous 1)	2
Decompression of thoracic epidural abscess	2
Revision of cervical spinal fusion	2
Fixation of thoracolumbar spinal fractures with or without cord injury	2
Miscellaneous: (one each of – revision of scoliosis surgery/removal of rods, cervical	5
microdiscectomy, decompressive thoracic surgery, debridement of wound, tracheostomy)	

Metastatic malignant disease

Procedure (may be multiple for individual patients)	Number
Decompressive thoracic surgery with or without fusion/stabilisation	13
Decompressive cervical surgery with or without fusion/stabilisation	7
Decompressive lumbar surgery with or without fusion/stabilisation	6
Biopsy of lumbar vertebral body	1

Total number of patients = 76

General comments

There was a high consultant involvement in this group of patients and there were only two cases in which there was no involvement whatsoever. Surgery was done by appropriate specialists in all cases (orthopaedic surgeons 38, neurosurgeons 31, specialised spinal surgeons 6 and general surgeon 1). Eighty-nine percent (68/76) of these patients were aged over 50 years, 72% (55/76) were classed as ASA 3 or greater and in 80% (61/76) there were coexisting problems.

Patient assessment and selection is crucial in this group of patients and it appeared that the patient as a whole was often not considered, especially in the presence of known metastatic malignant disease. Where selection was appropriate, surgery for metastatic disease was justifiable because, although it is not curative, it produces neurologically good results and improves quality of life (see personal commentary by Mr Timothy Morley, page 73). However, many procedures appeared to be a fruitless waste of resources. There may be pressure on surgeons from oncologists but proper discussion about patient management must take place.

In benign disease, difficult revisional surgery for conditions such as scoliosis are best referred to centres with experience of this type of procedure and its difficulties. Patient selection was often poor. Results depend on selection, which in turn requires experience. In the case of trauma there is time to transfer patients to specialist units and there is little need for operations out of hours.

Specific cases

Preoperative assessment

Age	Procedure	Comments by the advisors
51	Cervical and thoracic spinal decompression and stabilisation (anterior approach).	Colonic carcinoma with cerebral and hepatic metastases. A thoracic and an orthopaedic surgeon operated jointly but was the patient properly assessed? No cerebral or hepatic imaging despite clinical signs.
54	Anterior removal of L4 body and graft (tumour).	Tumour involved iliolumbar veins. Uncontrollable haemorrhage. Poor assessment with no MR imaging. Was surgeon experienced in this field?
64	Posterior stabilisation D8-L3 with laminectomy D12-L1.	Surgery for uncontrolled pain from suspected but unproven myeloma. Known cardiac and respiratory disease. No proper work-up. Final diagnosis carcinoma prostate. Aggressive surgery could have been avoided if preoperative diagnosis made.
73	Laminectomy C3-4.	Cervical myelopathy. Known ischaemic heart disease and hypertension. Hypotensive during surgery. Postoperative CVA. Poor selection.

Missed abdominal pathology

Age	Procedure	Comments
39	L5-S1 fenestration and excision of disc.	Surgery appropriate. Died of general condition in specialist ward. Was avoidable if condition had been recognised. Suspected adynamic ileus but taking opiates and was constipated leading to stercoral ulceration and perforation.
Not known	C1-C2 fusion (displaced odontoid fracture).	Transfer to regional unit said to be too risky. Original operation rather precipitate but two surgeons operating together. High steroid dosage. No NSAIDs. Missed abdominal condition – perforated duodenal ulcer – in ICU on fourth day.

Inappropriate procedures

Age	Procedure	Comments
49	Removal of Hartshill rectangle, revision of spinal decompression and posterior fusion.	Five litre blood loss. Patient died same day. Pulmonary embolus found at postmortem examination. Difficult case worthy of referral to specialised centre.
71	Thoracotomy and vertebrectomy.	Metastasis in D12. No primary identified. Failed irradiation. Died on third day - respiratory failure. Indication for surgery in terminally ill patient unclear.
73	Open biopsy L3.	Metastatic tumour. ASA 4. Why not do needle biopsy rather than open procedure?
76	L4 laminectomy (metastatic prostatic carcinoma).	Three to four hour operation by locum consultant who blames trainees for death. Anaesthetist felt pressurised. Surgeons missed gross renal failure.
85	C2-C3 laminectomy.	Rapidly progressive myelopathy. Hypertensive and rheumatoid arthritis. Registrar operation. No ICU bed. Respiratory failure. Too heroic.
90	Anterior vertebrectomy L4.	Metastatic disease, anaemia, ischaemic heart disease and established chest infection.

Spinal surgery 72

The management of metastatic spinal tumours

Key messages

- The frequency with which spinal metastases present to spinal surgeons is increasing.
- A logical approach needs to be taken, both in justifying surgery for the individual and also in deciding the extent of surgery.

Personal commentary written by Mr Timothy Morley FRCS

Metastatic spread to the spinal column is most common from primaries occurring in the breast, lung, kidney, thyroid and prostate.³⁴ Involvement of the spine by metastatic disease occurs in 60% of patients dying of breast cancer ³⁵ and as many as 5% of patients with systemic cancer will develop spinal cord compression.³⁶

With recent advances in oncological management, patients are surviving longer and the frequency with which spinal metastases present to the spinal surgeons is increasing. The goal of the spinal surgeon should be to relieve pain and preserve neurological function. This has been achieved by an aggressive approach of spinal decompression followed by stabilisation using implants. ³⁷⁻⁴² By adopting this approach there is no doubt that the outcome has been significantly improved when compared with the more classical approach of irradiation or laminectomy and irradiation.

It is important however, to temper enthusiasm by the fact that published results have recently shown that outcomes have fallen short of expectation. ⁴³⁻⁴⁵ This is because in all series it is clear that there are difficulties in deciding the right strategy in individual cases. ^{46,47}

When planning surgical intervention, consideration must be given to the patient's potential survival, and this is dependent on the primary site, the extent of spread to other distant sites as well as the spread within the spinal column. The general condition of the patient and the degree of neural involvement are also relevant. It is important also to consider the likely benefit to be expected in terms of relief of pain and recovery of neurological function and also to improve the quality of life. It is well recognised that patients who develop rapid onset of paraparesis (less than three days) do badly whatever strategy is chosen. In order to avoid undertaking major reconstructive surgery in patients with a poor prognosis, it is important to be aware of the relative significance of these factors. A number of authors have suggested criteria for decision making, but all of these fail to take into account all modalities which require consideration. Tokuhashi et al⁴⁵ tried to obtain a more accurate prognostic score based on six factors;

- General condition
- Number of extra-spinal bone metastases
- Number of metastases in the vertebral body
- Metastases to the major internal organs
- Primary site of the cancer
- Spinal cord involvement

This score is calculated preoperatively with each parameter scoring from 0-2 giving a maximum score of 12. Excisional surgery would be indicated for patients with a total score of 9 or more, whereas palliation is indicated if the score is less than 5. This approach has recently been validated by Enkaova et al ⁴³ and they applied the criteria retrospectively to 71 patients presenting with vertebral metastases. Their study confirmed the prognostic value, except in cases where the primary tumour site was unknown, in which case the outcome was significantly worse than predicted. A further prospective trial of this scoring system is at present being undertaken.

With improvement in surgical techniques it is also important to consider less radical surgical decompression. For example, it is quite possible to debulk an anterior tumour mass via the pedicle before stabilisation with posterior implants.

Early postoperative mortality following spinal surgery is clearly to be expected in this difficult group of patients. However, in the sample reviewed by the NCEPOD advisors, the histories and surgical procedures undertaken in patients dying in the early postoperative period strongly suggest that a more logical approach needs to be taken, both in justifying surgery and also in the extent of the surgical approach.

Further reading

British Association of Surgical Oncology. Guidelines for the Management of Metastatic Bone Disease in Breast Cancer in the United Kingdom. European Journal of Surgical Oncology 1998 (in press).

Urology

Key messages

- Team assessment is required for all elderly patients, particularly when contemplating radical surgery for malignant disease, because of the high incidence of other medical problems.
- A balance has to be maintained between too aggressive surgery and the quality of life when carrying out surgery in the presence of metastases or following previous radiotherapy.
- In urology there was a high involvement of Consultants.

Written by Mr Ron Hoile MS, FRCS and Emeritus Professor Vic Tindall FRCOG in consultation with the urology advisors (see page 12)

List of procedures

Procedure (may be multiple for individual patients)	Mumber
Transurethral resection of prostate (benign and malignant)	106
Cystoscopy (+/- biopsy)	96
Transurethral resection of bladder tumour	80
Nephrectomy (malignant and benign)	45
Testicular & scrotal procedures	25
Ureteric stenting	18
Cystectomy	16
Ileal conduit	14
Other procedures on bladder (e.g. repair of rupture/perforation)	13
Bladder neck incision	12
Open prostatic procedures	12
Miscellaneous ureteric procedures (e.g. reimplantation, ureterolysis etc.)	8
Procedures for ureteric/bladder calculus	8
Miscellaneous renal procedures (e.g. re-exploration for bleeding, removal of packs, nephrostomy)	7
Urethra dilatation	7
Miscellaneous urethra procedures e.g. drainage of periurethral abscess)	7
Evacuation of bladder clots	6
Cystodiathermy	6
Combined transurethral resection of prostate and bladder tumour	5
Penile procedures	4
Laparotomy	4
Percutaneous nephrolithotomy	4
Resuture of wound	2
Repair inguinal hernia	2 2
Right hemicolectomy	2
Total abdominal hysterectomy	2
Transverse colostomy	2
Miscellaneous (one each of: cholecystectomy, splenectomy, hemiarthroplasty, manual evacuation	6
of faeces, excision retroperitoneal nodes, pelvic exenteration)	

Total number of patients = 518

Urology 75

Review of cases

In many of the cases reviewed the patients had died of progressive disease and the urological intervention seemed to be part of the overall care of the patient, with death occurring perhaps a week or two after surgery. Most deaths occurred in patients undergoing surgery for cancer; it is gratifying to see how few deaths resulted from the management of patients with septic stone disease or conditions such as female urinary incontinence or benign prostatic enlargement in men. There was also a high involvement of senior surgeons. In 92% of the 518 cases reviewed the consultant was involved in the preoperative decision making. Consultants were present in theatre in 79% of all cases and in 71% of cases were the principal operators.

Elderly patients

The majority of the deaths were in elderly patients; 77% (399/518) were aged 70 and over, of whom 46% (184/399) were aged 80 and over.

Comorbidity

Co-existing problems were present in 87% (449/518) of the patients and there were two or more problems in 60% (271/449) of these patients. Nevertheless death was not anticipated or expected (by the surgeons) in 42% (219/518) of the cases.

Postmortem examinations

There were several cases in which it was considered that a postmortem examination should have been performed and there was no real explanation as to why this was not recommended.

Recognised hazards

Haemorrhage is a recognised risk of nephrectomy and/or difficult major surgery and was the major factor in five deaths.

Pulmonary embolism was the cause of death in only 1% (7/518) of the cases reviewed.

Inappropriate surgery or management

There were nine deaths in which the advisors considered that the surgery for the particular patient was inappropriate. Often it was inadequate preoperative assessment which led to inappropriate surgery, for example proper assessment and CT scanning might have prevented surgery in four of the cases detailed below. Major urological cancer surgery in patients with an ASA classification of 3 or more may be technically feasible from a surgical point of view but, as in the cases presented below, is sometimes not tolerated by the patient. In such cases less aggressive surgery and an early referral for palliative care assessment would seem to represent a better management choice. Criticism of these nine cases, where it was agreed that surgery was generally inappropriate, should not be seen out of context. They represent a small percentage (2%, 9/518) of the cases studied and (although NCEPOD does not have the denominator data) an even smaller percentage of urological surgery in general.

Age and sex	ASA grade	Procedure	Comments by the advisors
62F	3	Total cystectomy	Invasive TCC. Advised cystectomy despite being hypertensive and was on warfarin because of previous aortic valve replacement. Previous CVA. Postoperative persistent hypertension and overwhelming sepsis. Considered to be an inappropriate operation.
70M	3	Nephrectomy, cystectomy and ileal conduit	Anaplastic tumour of kidney infiltrating perirenal tissues. Palliative care was considered to be more appropriate.
71M	4	Initial operation was radical left nephrectomy. Final operation was laparotomy for bleeding from body of pancreas	This man had a stable bronchogenic cancer. A left adrenal mass involving the kidney, aorta and coeliac axis was detected. Delayed haemorrhage and CVA followed an inappropriate radical operation in this high-risk patient.
71F	3	Cystectomy and ileal neo- bladder	A poor choice of operation in view of the extensive disease present and ASA grade.
72M	4	Total cystectomy	Salvage cystectomy post irradiation. Postoperative haemorrhage and subsequent sepsis. Were the risks of this type of surgery justified?
74F	3	Total cystectomy and ileal loop diversion	Reason for the procedure in the presence of lung metastases not clear. An operative blood loss of three litres was related to her gross obesity.
74M	4	Cystoscopy	Poor risk with COPD. Inappropriate cystoscopy in the presence of known metastatic disease and the absence of symptoms.
76F	3	Right hemicolectomy, total cystectomy, hysterectomy and ileal conduit	Inappropriate radical urological surgery was carried out by a general surgeon with an interest in gastroenterology when a partial cystectomy was all the urological surgery necessary in the management of a caecal carcinoma involving the dome of the bladder.
84M	3	Radical nephrectomy	High medical risk – previous MI and DVT; inpatient for three weeks prior to operation. No anticoagulant prophylaxis. Locum consultant surgeon. Postmortem examination revealed lung metastases and massive pulmonary embolism. Considered poor case assessment.

Minor problems

The preoperative assessment or work-up was thought to be less than ideal in two cases and in one of these cases the postmortem examination revealed an undetected carcinoma of the bladder and secondary lung metastases. The other patient was considered to be unsuitable for the day case surgery which was done in view of known morbid obesity and previous major anaesthetic problems. Did the unit in question have a protocol for the selection of patients suitable for day case surgery?

Postoperative problems

The postoperative care was identified to be of poor quality in eight cases (fluid overload in two, sepsis in two cases and poor continuity of care by general medical team in four cases).

The problem of patients being on general surgical or medical wards without the speciality trained nurses was considered a factor which influenced the outcome in several of these cases. One patient, for example, who was unable to be put on ICU after a pelvic exenteration, was admitted to a medical CCU and subsequently died from intravenous fluid overload. NCEPOD has commented on similar points in a previous report.⁴⁸

Central venous pressure monitoring during anaesthesia and surgery

Key messages

- CVP monitoring may be indicated for patients with acute or chronic medical conditions.
- Equipment for CVP monitoring should be available in operating theatres in which major surgery is performed.
- CVP insertion is a core anaesthetic skill that needs regular practice.

Personal commentary written by Dr Roop Kishen FRCA

Anaesthesia and surgery have undergone great changes in the last two decades. Major and increasingly more complex surgery is being undertaken in physiologically compromised patients who present for surgery with co-morbidity, especially respiratory and cardiac disease (reference 49, page 41). In some of these patients surgery would not have been contemplated but for several important developments which enhance patient safety. These include the use of appropriate and timely monitoring and the support services of intensive care and high dependency units. For routine and emergency surgery minimal monitoring standards have been established. However, in many situations the safe conduct of anaesthesia and surgery requires additional, more invasive monitoring such as central venous pressure (CVP), intra-arterial blood pressure and

Equipment

The equipment for insertion and monitoring of CVP and suitably trained assistants should be available in all operating theatres.

Training

CVP monitoring is easy to learn and teach. Anaesthetists learn CVP line insertion during their basic training and all should be familiar with the technique. However, it is an acquired skill and, to be proficient, it has to be practised. With practice CVP line insertion is not time consuming, it can be placed easily and has few complications providing well known basic rules are followed. In anaesthetised ventilated patients the internal jugular is preferable to the subclavian route, it is associated with fewer complications e.g. pneumothorax and concealed haemorrhage. However, if the internal jugular vein cannot be accessed, the subclavian route can be used providing the benefits of monitoring outweigh the risks of complications.

Patient selection

Some types of surgery are associated with the use of CVP routinely. CVP monitoring is well established for cardiac surgery. Most anaesthetists will use invasive monitoring during major vascular surgery, e.g. aortic aneurysm repair (reference 49, page 68). NCEPOD 1993/1994⁴⁹ reported that 79 out of 85 patients undergoing emergency aortic aneurysm repair had a CVP line inserted for monitoring purposes. The report goes on to say that "the role of invasive monitoring is undisputed; central venous and arterial pressures are recognised to be valuable aids to safe management of patients during and after these operations".

In other patients who are undergoing anaesthesia and surgery, however major, routine use 6. CVP monitoring is not yet established (reference 49, page 79). The reasons for this are not clear. It is possible that the small risks of CVP catheter insertion, when balanced against its benefits are not fully appreciated. All anaesthetists recognise the importance of minimal monitoring of patients during anaesthesia and surgery. CVP monitoring, in selected cases and for patients at risk, should be regarded as an extension of this monitoring.

Which patients would benefit from CVP monitoring?

Surgical indications include major thoracic surgery (e.g. oesophagectomy), major abdominal surgery (massive resections, major liver surgery, pelvic clearance etc.), surgery for complex fistulae, resection of phaeochromocytoma, major orthopaedic reconstruction, major plastic and burns surgery, sepsis, trauma etc. - all cases in which large fluid shifts, major blood losses or circulatory instability are expected during their surgery and postoperative course.

A group of patients which appear to the advisors less readily recognised are those with acute or chronic medical conditions that may compromise their tolerance of blood loss and fluid infusions. Examples of these include patients with a history of cardiac failure, those at risk of developing postoperative acute renal failure, patients with dialysis dependent chronic renal failure etc. This list is not exhaustive; any patient at risk is a candidate for CVP monitoring.

Age itself is a relative indication. As longevity increases, more and more elderly patients will present for surgery. This population is physiologically less competent and has many co-morbidities including ischaemic heart disease, hypertension (increasingly treated with potent ACE inhibitors), diabetes and poor renal function. For safe conduct of anaesthesia and surgery, for appropriate fluid management and timely cardiovascular support in these patients, CVP monitoring should be considered as routine as other standard monitoring. It is accepted that insertion of a CVP line does not guarantee outcome. However, its use in this population should make their management easier and more logical. This in itself may avoid mortality and morbidity.

In many patients with severe preoperative dehydration or sepsis requiring resuscitation, CVP monitoring is a useful guide to the effectiveness of therapy.

Interpretation of the CVP

In its interpretation, CVP monitoring requires understanding and experience. It must be appreciated that CVP monitoring is not a substitute for the measurement of cardiac output. CVP measurement does not equate with blood volume or left ventricular function. It does not tell us about the right ventricular function either! CVP monitoring is an indirect measure of right heart preload and it indicates the ability of a particular patient to deal with the infused fluid load. In certain circumstances it may also indicate the need for inotropic support. Thus it can aid fluid management in many patients, especially those at risk. NCEPOD has repeatedly pointed out the dangers of inappropriate fluid infusions in 'at risk' patients. In such patients it is possible to avoid large, inappropriate fluid infusions and either provide inotropic support or institute further investigative monitoring such as pulmonary artery catheter or oesophageal Doppler.

Patients undergoing surgery and general anaesthesia without CVP monitoring.

In these patients, amongst many others, the advisors questioned whether CVP monitoring might have been useful.

	Age	ASA	Operation	Co-morbidity	Outcome
Yes	73	3	Radical hysterectomy and node dissection	Previous MI, angina, COPD, heavy smoker, three previous CVAs)	Per- and post-operative hypotension. Returned to the ward. Developed pulmonary oedema
The second	91	3	TAH, bilateral salpingo-oophorectomy	Frail	Returned to the ward. Very positive fluid balance. Developed cardiac and renal failure
	90	4	De-bulk ovarian tumour adherent to bowel	Two previous MIs, CCF, COPD, asthma, ascites, SOB at rest	Per- and post-operative hypotension. Returned to ICU/HDU
	57	4	Bilateral oophorectomy and omentectomy	44kg, cachectic, ascites, SpO ₂ 84%	CVP inserted postoperatively for feeding. Desaturation in recovery. Returned to the ward
i	83	5	Ovarian mass excision, colostomy, omentectomy	Bowel obstruction, 'moribund'	Returned to the ward
}	34	3	TAH, bilateral salpingo- oophorectomy, omentectomy and right hemicolectomy	Mild CCF and AF	Peroperative hypotension, SAP 80mmHg for two hours. Fast AF in recovery. Returned to the ward
7	8	3	Laparoscopic mobilisation of the colon failed, progressed to colon excision via Pfannenstiel incision, nephrectomy via subcostal incision.	IHD, CCF, AF, orthopnoea with five pillows	Five hours of surgery. Epidural top-up at the end of the operation. In recovery hypotension, cardiac arrest and died
71	0	3	Total gastrectomy and splenectomy	Hypertension, AF, IHD	Epidural top-ups at the beginning and end of surgery. Prolonged episodes of hypotension peroperatively. Returned to the ward
76	6		Acute abdomen. Attempted bowel resection	Unwell, on steroids and NSAIDs for vascular and musculoskeletal problems. K ⁺ 5.5mmol.L ⁻¹ , Cr 185micro mol.L ⁻¹ , U 20.9mmol.L ⁻¹	No preoperative fluids. Per-operatively; fluids 1500ml, BP 80/35, pulse 120. Returned to ICU, died 6h later
59) 2		Laparotomy, bilateral ureterolysis	Obese, pneumoconiosis, cor pulmonale, unstable angina, CCF, renal failure	Returned to ICU, then the ward. Developed ischaemic colon, probably embolic

Admission on the day of surgery

Key messages

- The selection of patients for admission on the day of surgery should conform to locally agreed guidelines.
- Patients should be fully assessed and investigated at an outpatient visit preoperatively.
- Some patients need joint decisions between specialists, in particular between consultant surgeons and consultant anaesthetists.
- The timing of admission on the day of surgery should allow a final clinical assessment by both surgeon
 and anaesthetist before the patient enters the theatre complex.

Written by Dr Kathy Sherry FRCA in consultation with the anaesthetic advisors (see page 10)

In this sample of deaths, 77 anaesthetic questionnaires were received, relating to patients who had undergone scheduled or elective surgery and who were also admitted on the day of operation. They were not scheduled for day case surgery.

Classification of final operation*/ASA grade

	ASA 1	ASA 2	ASA 3	ASA 4	ASA 5	Total
Scheduled	-	14	16	6	-	36
Elective	4	18	17	2	-	41
Total	4	32	33	8	-	77

^{*} see glossary for classification and ASA grades

The two elective ASA 4 patients were a 64-year-old man, for a transurethral resection of prostate, who had angina on minimal exertion and at rest, and an 85-year-old man for palliative oesophagoscopy and dilatation.

Age of patient (years)

30 - 39	1
40 - 49	2
50 - 59	3
60 - 69	17
70 -79	31
80 -89	20
90 - 99	3
Total	77

The following seven patients had no preoperative investigations.

Age (years)	Operation	General/local anaesthesia	Medical disorder
91	Cataract removal	Local	None
92	Ectropion repair	General	'Frail'
74	Urethral dilatation	General	None
70	Check cystoscopy	General	Asthma
68	Check cystoscopy	General	COPD and peripheral vascular disease
78	Laparoscopic inguinal hernia repair	General	Angina
65	Palliative oesophagoscopy and dilatation	General	Recurrent ascites

Twenty-one cases are presented in outline to illustrate the range of cases that were reviewed. NCEPOD is not commenting on the suitability of individual cases for admission on the day of surgery but hopes that their presentation here may stimulate useful local debate.

Age 68	ASA 3	Classification Scheduled	Medical problems COPD and peripheral vascular disease*	Operation Check cystoscopy
64	4	Elective	Angina on minimal exertion and at rest	TURP
77	3	Scheduled	Ischaemic heart disease	TAH and bilateral salpingo- oophorectomy
76	4	Scheduled	Ischaemic heart disease, CVA, alcoholism	EUA larynx, pharynx and
78	3	Elective	Angina, obesity (90Kg)*	oesophagus Laparoscopic inguinal hernia repair
67	2	Scheduled	IHD	Thoraco-abdominal oesophagectomy
72	2	Elective	IHD and hypertension, no preoperative CXR, died of thoracic aortic dissection	Vaginal hysterectomy and anterior vaginal repair
88	3	Elective	IHD	TURP
88	2	Elective	'Frail', preoperative ECG only	TURBT
56	3	Elective	Diabetes, angina on minimal exertion and at rest	Colposuspension
76	3	Elective	Angina	TURP
85	3	Scheduled	No preoperative ECG	TURBT
70	3	Elective	Severe angina	Percutaneous nephrolithotomy
81	2	Scheduled	No preoperative ECG, MI 36hr postoperatively	Retropubic prostatectomy
83	4	Scheduled	Severe ischaemic heart disease	TURBT
60	3	Elective	Unstable NIDDM, IHD	Cystoscopy
70	3	Elective	IHD	Total gastrectomy and splenectomy
68	3	Elective	Renal impairment and NIDDM	TURBT
80	2	Elective	IDDM, angina, dementia, admitted one hour preoperatively	Vitrectomy
59	2	Elective	History of difficult previous anaesthetic, obese (115kg), smoked 30/day, hiatus hernia	Check cystoscopy
40	2	Elective	Down's syndrome, mitral regurgitation, no preoperative ECG	Laparoscopic cholecystectomy

^{*} Patient had no preoperative investigations, and is also featured in the previous table.

In a drive to reduce length of hospital stay and increase patient throughput many patients are being admitted on the day of surgery. We do need to recognise that for some patients admission on the day of surgery is not in their best interest.

A large number of the deaths following scheduled or elective surgery were unanticipated. The impact of an unexpected death in the family cannot be measured but it is unlikely to be eased by precipitate admission for a planned operation.

Patients admitted on the day of surgery require meticulous pre-admission preoperative assessment and preparation.

Guidelines on patient suitability, locally agreed between the surgical and anaesthetic staff, should be encouraged.

Preoperative assessment

Assessment should be at a dedicated outpatient clinic allowing time for review of investigations before admission as well as co-ordination of the admission, referral for specialist opinions or special investigations, as required. At this time preoperative preparation including the management of routine medication should be explained and supported by written instruction.

Junior surgical staff often run assessment clinics. However, some individuals will need referral to and assessment by an anaesthetist before admission.

There are no absolute criteria for suitability; assessment should consider the following:

Medical status

Age alone should be considered but is not a contraindication; poor physiological status is. Patients who are healthy or have stable medical conditions are more likely to be suitable. Patients with unstable medical conditions need more careful consideration.

Examination of the table above shows a striking incidence of ischaemic heart disease that was known of or detectable preoperatively. Patients are often anxious about hospital admission and their impending anaesthesia and surgery. It is recognised that myocardial ischaemia is worsened by stress and that preoperative myocardial ischaemia is associated with increased postoperative death. It should be questioned, therefore, whether patients with ischaemic heart disease are suitable for admission on the day of surgery.

Operation

The type of operation alone is not a contraindication. For example, a young, healthy individual having a major operation may be considered suitable for admission on the day of surgery; a patient with an unstable medical condition having a minor operation is likely to be unsuitable. Patients undergoing palliative surgery should be assessed individually.

Social factors

Social circumstances need to be considered. It may be difficult for the patient to get to hospital on time and in a suitably prepared emotional state. Admission on the day of surgery may cause unacceptable anxiety for the patient or their carer.

The anaesthesia and surgery may be by medical staff unfamiliar with the patient or there may have been a change in the patients' condition between preoperative assessment and admission. It is important therefore, that admission on the day of surgery should be timed to allow the surgical and anaesthetic staff to make a final clinical assessment *before* the patient enters the theatre complex.

Complications associated with postoperative analgesia

Written by Dr Kathy Sherry FRCA

Twelve questionnaires reported complications with postoperative analgesia.

Age 50	Operation L4/5 spinal fusion.	Analgesia Intravenous bolus opiate in recovery.	Location and problem In recovery area, acute laryngeal oedema in response to morphine	Intervention Died from the complication
81	Tracheostomy and EUA larynx	Intravenous bolus opiate in recovery, subcutaneous opiate on the ward	In recovery area, transient hypoventilation	Doxapram given, resolved
83	Left nephrectomy	Continuous intravenous opioid	In recovery area, drowsy	Responded to naloxone. Already scheduled for ICU
76	Laparotomy for bleeding gastric ulcer.	Continuous infusion of opiate	In HDU, opiate narcosis	Changed to Tramadol
83	Total abdominal hysterectomy and high anterior resection.	Opiate PCA with background infusion	In HDU, "relative opiate overdose"	No treatment
81	Bilateral oophorectomy and hysterectomy.	Opiate PCA	On the ward, patient unable to use it	Changed to a continuous infusion of opiate
70	Laparotomy and over- sew bleeding duodenal ulcer	Opiate PCA	On the ward, over-sedation resulting from the patient's relatives' inappropriate use.	None
68	Open pyelonephrolithotomy.	Opiate PCA	On the ward, at 21.00 hrs respiratory rate 6/min. and pinpoint pupils.	Responded to naloxone
83	Left nephrectomy	Opiate PCA	On the ward, drowsiness.	Responded to naloxone
70	Posterior thoracic spinal decompression and fusion following a previous anterior decompression and chest drain.	Epidural opiate	Peroperatively the epidural catheter was inserted under direct vision. The patient had infection in the chest drain site. Postoperatively he developed a paraspinal abscess.	None, developed paraplegia
72	Total gastrectomy	Epidural opiate and bupivacaine	Postoperatively on the ward he had breakthrough pain; necrosis of the distal oesophagus was not recognised.	None, patient developed sepsis and respiratory failure
54	Oesophagogastrectomy	Epidural opiate and bupivacaine	Postoperatively developed infarct of the posterior cord and paraplegia.	None

Lessons from the above patients are clear.

It is the responsibility of the anaesthetist to ensure that patients are nursed in an area where both the nursing and medical staff with primary responsibility for the patient have been appropriately trained in the analgesic techniques and are aware of the associated complications and their management.

With epidural analgesia a surgical complication causing breakthrough pain should not be missed.

Respiratory depression can occur with PCA or intravenous opioid analgesia and patients need documented monitoring according to the local protocol.

Two of the complications that were reported -laryngeal oedema following intravenous morphine and cord infarct associated with epidural analgesia - are thankfully very rare.

Non-steroidal anti-inflammatory drugs

Key messages

Non-steroidal anti-inflammatory drugs (NSAIDs) can be very useful analgesics in the perioperative period. However they should be used with caution in patients with:

- renal impairment
- hypertension and cardiac failure
- GI ulceration and bleeding
- asthma

Written by Dr Anthony Gray FRCA in consultation with the anaesthetic advisors (see page 10)

Current thinking is that if a therapy is effective it should be applied in every situation where the patient should benefit; conversely, the therapy should not be used when there are contra-indications or the risks are too great. Wherever possible, decisions on effectiveness, risks and contra-indications should be based on published evidence and guidelines. The Royal College of Anaesthetists⁵⁰ has issued "Guidelines for the Use of Non-steroidal Anti-inflammatory Drugs in the Perioperative Period". These show that NSAIDs are excellent analgesics in the right circumstances.

The questionnaires returned to NCEPOD give an insight into the way that NSAIDs are used particularly by anaesthetists.

NSAIDs were prescribed for postoperative pain for 178 patients out of the total of 1269 (14.0%). NSAIDs were prescribed in combination with IM analgesics (64 patients), PCA (28 patients) and epidural infusions (12 patients). NCEPOD did not ask for copies of prescription charts so it is not possible to say how often these drugs were actually dispensed. These figures indicate that anaesthetists are cautious about prescribing NSAIDs in the cohort of surgical patients examined in this survey, despite their undoubted efficacy in the treatment of postoperative pain.

On the other hand, many patients were prescribed NSAIDs in situations where there was the potential to precipitate adverse effects. The anaesthetic advisors agreed that such a decision was justified when NSAIDs were prescribed as part of palliative care; obviously in such situations, a further deterioration in a patient's general condition may be less important compared to achieving good control of pain. In other cases the advisors questioned their use.

The following patients, whose condition was not regarded as terminal, were prescribed NSAIDs

Age	Operation	Clinical details
78	TURP	Hypertensive, chronic renal failure with a creatinine of 500 μ mol/L. On hydralazine, atenolol and bendrofluazide.
81	EUA, hysteroscopy	BP 205/110 mm Hg, no treatment. No preoperative urea or electrolyte measurement.
84	TURP	Noted preoperatively to have "severe chronic active gastritis and ulceration". Given diclofenac during his operation.
	(continued overleaf)	

Age	Operation	Clinical details
74	Laparotomy for postoperative bleeding	Preoperative renal impairment, creatinine 296 μ mol/L.
74	Laparoscopic liver biopsy	Hepato-renal syndrome, creatinine 142 μmol/L, bilirubin 87.
60	Abdominal hysterectomy	Asthmatic, on beclomethasone, salbutamol and aminophylline, no known allergies. Died from acute gastric dilatation.
87	Nephrectomy	CCF, pleural effusions, NIDDM, on frusemide and amiloride. Creatinine 97μ mol/L, urea 12.9 mmol/L.
76	Incision and drainage knee abscess	Septic arthritis, chronic renal failure, IHD, liver dysfunction with INR of 2.2.
70	Total gastrectomy	Hypertensive, IHD on oxprenolol. Developed oliguria day after surgery, died the following day in acute LVF.
79	Nephrectomy	Diet-controlled diabetes, anaemia.
88	ORIF cervical spine	Frail. AF. Na 123 mmol/L preoperatively. Hypotensive intra- operatively, given 5000 ml of fluid in first 24 hours. No CVP. No record of blood loss.
39	L5/S1 discectomy	Diclofenac and PCA postoperatively. Fluctuating nausea and vomiting, then collapsed eight days after surgery and could not be resuscitated. Postmortem examination showed a small colonic perforation.
83	TURP	Old MI, angina. Hypotensive during GA, given 1500 ml fluid intra- operatively over 30 minutes.
81	Incision and drainage infected hydrocoele	Poor renal function which deteriorated postoperatively
86	Division of adhesions, formation of colostomy	Two previous myocardial infarctions, AF, CRF, anaesthetised by a 2^{nd} year SHO in the middle of the night and sent back to the ward
86	TURP	Three previous MIs, hypertension on atenolol. SHO anaesthetist of more than two years' experience
39	Ileal conduit	Renal impairment, creatinine 189 μmol/L
76	TURBT	Hepatic metastases and renal impairment. Developed haematemesis and melaena postoperatively
75	Balloon dilatation of ureter through ileal conduit	IHD, CCF, and CRF (creatinine 308 μ mol/L). GA. Hypotensive intraoperatively, given ephedrine and 2,500 ml of fluid over 90 minutes

Age

Age is not a specific contraindication to the use of NSAIDs. Many of the risks associated with NSAIDs do increase with age, for example GI bleeding and the risk of concurrent renal impairment, hypertension or cardiac failure. In this group of patients, the youngest patient prescribed NSAIDs was 36 and the oldest was 92 years. The proportion prescribed NSAIDs did not change with the patient's age until they were over 90 years.

Age	Total number	NSAIDs	Percentage
<60	183	24	13.1
60-69	241	35	14.5
70-79	474	66	13.9
80-89	338	51	<i>15.1</i>
>90	33	2	6.1

Renal impairment and known renal pathology

To quote from the Royal College of Anaesthetists (RCA) guidelines⁵⁰ (p. 10) "...it is clear that in the great majority of cases, use of NSAIDs for postoperative pain control will not jeopardise renal function. However, practising nephrologists recognise that a high proportion of cases of postoperative renal failure are associated with the use of NSAIDs. Rather than being the primary cause they are usually a contributory factor to the development of ARF (acute renal failure)." The guidelines list situations in which NSAIDs should be avoided, including renal impairment, hypovolaemia and circulatory failure. They also recommend caution in the use of NSAIDs in patients receiving ACE inhibitors, potassium-sparing diuretics and β-adrenergic blocking drugs. Renal function should be checked whenever renal impairment might be suspected.

Hypertension and cardiac failure

The RCA guidelines recommend that patients receiving treatment for hypertension or cardiac failure should be monitored closely if they are prescribed NSAIDs because NSAIDs cause fluid and water retention. In addition, concomitant diuretic administration increases the risk of renal toxicity.

GI ulceration and bleeding

GI problems are usually associated with long term use. However, GI bleeding or perforation should be a prominent differential diagnosis in all patients receiving NSAIDs. They should not be administered to patients with a history of GI ulceration or bleeding.

Haematological effects

NSAIDs increase bleeding time. The precise clinical significance of this is not clear.

Asthma

NSAIDs should not be used in patients with aspirin-sensitive asthma. They should be used with caution in other asthmatics.

It is impossible to assert that a patient would have recovered if NSAIDs had not been prescribed but clinicians must respect the contra-indications to the use of NSAIDs. The risks of renal impairment especially seem to be ignored. This needs to be appreciated by anaesthetists and surgeons. Some patients with raised creatinine levels were receiving diclofenac preoperatively on the ward. Other patients were prudently not prescribed NSAIDs by the anaesthetist, only for them to be prescribed by surgical ward staff postoperatively. Sick patients, especially if they are elderly, need immaculate care if they are to survive surgery.

Trainee anaesthetists supervising sedation for sick patients

Key message

• When deciding on the appropriate grade of anaesthetist the patient's condition should always be considered, together with the type of procedure and anaesthetic proposed.

Written by Dr Anthony Gray FRCA

Some patients requiring operation were correctly assessed as being very sick. It was assumed that general anaesthesia would impose great risks so their operation was carried out under local anaesthesia. In some of these, a junior anaesthetist was asked to provide sedation.

A 92-year-old man required decompression of a sigmoid volvulus. He had had a previous CVA. His current medical problems included an acute confusional state, atrial fibrillation, hypovolaemia, prerenal renal failure, and a white blood count of $31.8 \times 10^9 \, \text{L}^{-1}$. He was classed as ASA 4. A first year specialist registrar did not discuss the case with anyone. He gave "analgesia and oxygen only". The patient died after 12 days from continuing pseudo-obstruction.

An 81-year-old man required open reduction and fixation of a fractured mandible. He had had a previous CVA, and suffered from ischaemic heart disease, congestive cardiac failure, diabetes, and chronic renal failure. Investigations showed a white count of $20.9 \times 10^9 L^{-1}$, a serum sodium of 126 mmol.L⁻¹, serum glucose of 21.5 mmol.L⁻¹ and a creatinine of 209 micromol.L⁻¹. He was reported to be hypoxic on air, and was classed as "ASA 3-4". A first year SHO did not discuss the case with anyone, although he wrote on the chart that it was "an anaesthetic nightmare". He gave midazolam and ketamine whilst the operation was done under local anaesthesia. The patient died two days later.

A 76-year-old patient was scheduled for brachial embolectomy. He had a history of chronic obstructive pulmonary disease, congestive cardiac failure, and non-insulin dependent diabetes. His serum glucose was 23 mmol.L⁻¹, and his oxygen saturation was 85% despite oxygen being administered by a face mask. After discussion with a consultant, a second year SHO gave the patient 1 mg of midazolam during the operation, which finished at 13.00 hrs. In the recovery ward, the patient was unresponsive and noted to be "dusky". He was returned to the ward. He died three hours later.

A 75-year-old patient with rheumatoid arthritis, chronic obstructive pulmonary disease and congestive cardiac failure developed rectal bleeding and required an oesophago-gastroscopy. She was attended by an SHO who had started training in anaesthesia five years before. He did not discuss the case with anybody. The patient was given 1 mg of midazolam and 40 mg of propofol. She died the next day.

A 54-year-old man had an inoperable rectal cancer with obstructive renal failure and a creatinine of 899 micromol.L⁻¹. His INR was 2.2. A first-year SHO attended whilst a ureteric stent was inserted. He discussed the case with the second on call. He gave the patient alfentanil, midazolam and pethidine. The man died 19 days after the operation.

An 84-year-old woman with chronic obstructive pulmonary disease, ischaemic heart disease and atrial fibrillation, assessed as ASA 4, required gastroscopy. No results of investigations were available. A second year SHO did not discuss the case with anyone. He sedated the woman with alfentanil 300 mcg and midazolam 1 mg and administered oxygen. She died the next day.

All of these patients were very ill and the anaesthetists were of limited experience. In some cases it is not clear what benefit the patient was expected to derive from the operation. In only two out of the six cases above did the trainee discuss the patient with a more senior colleague. Presumably an anaesthetist was asked to attend during the procedures because of the patients' poor general condition, the high probability that the patient would become over-sedated and unconscious, and the risk that a local anaesthetic might not be effective. However, in all of these cases it would have been quite inappropriate for the patients to have received a general anaesthetic from anaesthetists of such limited experience, so the anaesthetist present would be precisely the wrong person to deal with this complication. In addition, the anaesthetists were unaccompanied and so unable to obtain any educational benefit from the encounter.

As in all other areas of anaesthetic practice, if the surgical team request assistance, the decision as to the experience of anaesthetist required should be based on the patient's condition and not on the type of anaesthetic suggested by the surgeon.

Pathology

Advisors

Dr P N Cooper

(Newcastle upon Tyne)

Dr C M Corbishley

(London)

Dr N Francis

(London)

Dr J D Hemming

(Hexham)

Dr S Rogers

(Doncaster)

Dr R D Start

(Chesterfield)

Key messages

- The overall postmortem rate of 28% was very low.
- The hospital postmortem rate of 8% was unacceptably low.
- A small proportion of postmortem reports still did not include a clinicopathological correlation which should be present in all cases.
- Eighty-seven percent of postmortem reports were graded by the advisors as satisfactory or better.
- Histology is frequently not necessary particularly in Coroner's postmortem examinations.
- There is a need for a set of guidelines and some standardisation in formulating an OPCS cause of death in postoperative death cases.
- Variation in Coronial practice makes it impossible to build a single logical framework for deciding whether a case should be referred to HM Coroner.

Review process and sample

The members of the advisory group of consultant pathologists met on five occasions and reviewed a total of 301 postmortems of which 237 were performed for Her Majesty's Coroners and 64 for hospital doctors with the consent of the relatives of the deceased. A total of 474 cases out of the overall total of 1697 appeared from the surgical questionnaire to have had a postmortem performed so only 63% (301/474) of postmortem reports had been submitted for review. The overall postmortem rate was 28%.

The diverse nature of the six groups of cases reviewed this year makes any examination of the combined figures somewhat artificial. The value of comparison with previous years where different case types were selected is questionable. It would be useful to look at the same groups again in say 10 years time to see if there have been any changes or improvements. The groups will be referred to individually when appropriate.

Pathology 93

Deaths reportable to the coroner and postmortem examination requests

The question numbers refer to those in the surgical questionnaire (see Appendix G).

Table PM1, question 50 Was the death reported to the Coroner?

Yes	742	44%
No	824	49%
Not answered	131	
Total	1697	

Deaths reported to the Coroner/specialty group

	Yes	No	Not	Total
			answered	
Gynaecology	91	96	18	205
Head and neck surgery	126	143	46	315
Minimally invasive surgery	114	175	21	310
Oesophageal surgery	132	134	7	273
Spinal surgery	49	19	8	76
Urology	230	257	31	518
Total	742	824	131	1697

Table PM2, question 50a

If yes, was a Coroner's postmortem ordered and performed?

Yes	371	50%
No	343	46%
Not answered	28	
Total	742	

Forty-four percent of deaths were reported to the Coroner, a lower figure than in previous years. ^{48,51-53} The minimally invasive surgery group in particular had a very low referral rate, whereas the spinal surgery group had a high referral rate. Higher referral rates in previous years probably reflect different sample criteria. For example, all the cases in 1994/95, where the referral rate was 82%, were of deaths within three days of surgery. ⁵³ Doctors are poor at recognising cases which should be referred to H.M. Coroner. ⁵⁴ This is partly as a result of a lack of medicolegal education in the undergraduate curriculum. ⁵⁵ A potentially even larger problem is the enormous variation in individual Coronial practice which means that it is not possible for a doctor moving from one Coroner's area to another over his career to build up a single logical decision-making framework when it comes to referral of a postoperative death to HM Coroner.

Referral would be greatly aided by a nationally agreed set of guidelines, probably by the centralisation of death-related matters within hospitals and possibly a routine review of individual deaths by a consultant pathologist.⁵⁶

The percentage of cases referred which were accepted by the Coroner and a postmortem examination ordered (50%) is similar to that in previous years but did vary a little according to surgery type.

Table PM3, question 51

Other

Not stated

Not known/not recorded

If a Coroner's postmortem examination was not performed, was a hospital postmortem examination undertaken?

102

20%

22%

3%

209

32

Yes	103		0/0
No	961		72%
Not answered	262		
Total	1326		
If no, why not? (answers may be multiple)			
Cause of death/diagnosis known/no indication		434	45%
Relative-related		198	21%
"Not requested"/patient died under care of another	97	10%	
Coroner felt not necessary		17	2%
Other		15	2%

Of those cases where a Coroner's postmortem was not performed only 8% had a hospital consent examination. This is similar to previous years remaining very low. Certainty as to the clinical diagnosis was again given as the commonest justification for not obtaining a postmortem despite a known high rate of discrepancies between such diagnoses and the results of postmortem examination. One would have hoped that, particularly in a relatively new area such as minimally invasive surgery, the clinicians would be more than usually interested in asking the relatives for permission for a postmortem examination. Factors likely to improve the hospital postmortem rate such as centralisation of death certification and regular clinicopathological meetings and a positive attitude to postmortem examinations by clinicians have previously been identified.⁵⁷ One reason for the low postmortem rate may be the fear of litigation based upon the findings. However, the availability of postmortem findings has on the contrary been shown to reduce both the number of claims and the cost of settlement where liability existed. 58,59

Review of postmortem examination reports

The pathology advisory group again used the "Guidelines for Postmortem Reports" published by the Royal College of Pathologists in 199360 as the gold standard. The one exception to this was that some reports were still graded as excellent or very good in the absence of any histology. The guidelines recommend that histology should be performed on every postmortem but in some of the cases reviewed histology appeared unnecessary to the advisory group.

Clinical history

Table PM4

Is a clinical history	provided	?				
·	Corone		Hospit	al	Total	
Yes	202	85%	62	97%	264	88%
No	35	15%	2	3%	37	12%
Total	237		64		301	

When present, the clinical history is:

Unacceptably brief, obscure, uninformative	10 24	4% 9%
Poor Satisfactory	68	26%
Good	60	23%
Fully detailed, clear, informative	102	38%
Total	264	

95 Pathology

Once again, all cases were typewritten. A clinical history was provided in all but two of the hospital postmortem examinations but was absent in 15% of the Coroner's postmortem examinations. This must in part reflect the fact that some Coroners forbid or discourage their pathologists from including a clinical history and there is perhaps increasing concern amongst pathologists that they might be said to be overstepping their remit by interpreting clinical evidence. When present, the clinical history was satisfactory or better in 87% of cases.

External examination

Table PM5

The description of external appearance is:

Unacceptably brief, inadequately detailed	15	5%
Poor	41	14%
Satisfactory	95	31%
Good	78	26%
Fully detailed, clear, informative	72	24%
Total	301	
Table PM6		
Were scars and incisions measured?		
Yes	172	57%
No	69	23%
Not applicable	60	20%

The external examination was described as poor or inadequate in almost one fifth of cases (similar figures for both Coroner's and hospital postmortem examinations). Scars and incisions were measured in over two thirds of the cases where relevant. There was disagreement amongst the advisory group as to whether it was necessary to measure scars.

301

Internal examination

Table PM7

Total

The gross description of internal organs is:

Unacceptably brief, inadequately detailed	8	2%
Poor	35	12%
Satisfactory	87	29%
Good	96	32%
Fully detailed, clear, informative	. 75	25%
Total	301	

The description of internal organs was satisfactory or better in 85% of cases, similar to previous years.

Table PM8
Number of organs weighed (paired organs count as 1)

None	21	7%
1	11	4%
2	14	5%
3	14	5%
4	19	6%
5	43	14%
6	141	47%
7	17	5%
8	14	5%
9	6	2%
more than 9	1	<1%
Total	301	

Table PM9

Were the skull and brain examined?

	Corone	r's	Hospit	al	Total	
Yes	218	92%	54	84%	272	90%
No	17	7%	10	16%	27	9%
Not known	2				2	
Total	237		64		301	

Sixteen percent of hospital postmortem examinations did not include examination of the skull and brain, at least some of which were cases where permission specifically excluded this examination. The 7% of Coroner's postmortem examinations where the skull and brain were not examined is identical to that from 1994/95.⁵³

Table PM10

Is the operation site described?

Yes	237	79%
No	44	15%
Full notes not available	20	
Total	301	

The operation site was not described in 15% of cases. A postmortem examination report should always include this information but in some of the cases this year it may have been missed out simply because of the relatively minor nature of the surgery (endoscopy, tracheostomy, TURP).

Table PM11

Is the gross examination appropriate to the clinical problem?

Yes	289	96%
No	10	3%
Full notes not available	2	
Total	301	

Additional investigations

Table PM12 (answers may be multiple)

Samples were taken for:

	Coroner's		Hospi	ital	Total	
Histology	72	30%	31	48%	103	34%
Microbiology	4		3		7	2%
Toxicology	4		-		4	1%
None of these investigations	162		33		195	65%
Total	237		64		301	

Table PM13 (answers may be multiple)

A thorough postmortem examination in this case would have called for:

	Coroner's				Hospital	Total			
	Called for	Sample taken		Called for	Sample taken		Called for	Sample taken	
Histology	234	71	30%	62	29	47%	296	100	34°
Microbiology	7	3		9	3		16	6	
Toxicology	2	2		-	-		2	2	
Total	237			64			301		

Table PM14
Is a histological report included with the postmortem examination report?

Yes	Coroner's		Hospit	al	Total		
	49	21%	24	31%	73	24%	
No	188	79%	40	69%	228	76%	
Total	237		64		301		

When present, the histological report is:

Unacceptably brief, inadequately detailed	-	
Poor	3	4%
Satisfactory	24	33%
Good	13	18%
Fully detailed, clear, informative	33	45%
Total	73	

When absent, does the lack of histology detract significantly from the value of this report?

	Coroner's		Hospital		Total	
Yes	28	15%	16	40%	44	19%
No	160	85%	24	60%	184	81%
Total	188		40		228	

Samples were taken for histology in 48% of hospital cases and 30% of Coroner's cases, higher figures than in preceding years but a histology report was included with the postmortem report in a much smaller proportion of cases. There was often evidence that histology had been taken and in many cases additional histology reports were probably issued but not sent to NCEPOD. The figures in Table PM13 assume that histology should have been done in all cases as suggested by the College guidelines but the figures in Table PM14 show that in Coroners' postmortem examinations, in the opinion of the advisors, the absence of histology was probably not important in 85% of cases. In 40% of hospital cases where it had not been performed it was thought that it would have been useful. There were many cases where surgical specimens had been removed in life and therefore some histological evidence had been gathered. There is considerable discrepancy here between the value of histology as seen by the pathology advisory group and that recommended by the Royal College of Pathologists. Whatever the postmortem examination type, histology has to be cost effective.⁶²

Pathology 98

The postmortem report

1. SUMMARY

Table PM15

Is a summary of the lesions present?

Yes	100	33%
No	201	67%
Total	301	

When present, does this correspond accurately to the text report?

Yes	95	95%
No	2	2%
Full notes not available	3	
Total	100	

A separate summary of lesions was only present in 33% of cases but in some cases a form of summary was included within the text of the clinicopathological correlation. The percentage where a summary was present was considerably higher than the preceding year. The College guidelines recommend that such a summary is always present. Perhaps the guidelines are having an effect.

2. THE CAUSE OF DEATH

Table PM16 Is an OPCS cause of death present?

Yes	283	94%
No	18	6%
Total	301	

When present, does it correspond accurately to the text report?

_	Coroner's		Hospit	al	Total	
Yes	218	92%	45	96%	263	93%
No	14	6%	2	4%	16	6%
Not clear	4		-		4	
Total	236		47		283	

Does it follow OPCS formatting rules?

	Coroner's		Hospit	al	Total	
Yes	227	96%	46	98%	273	96%
No	9	4%	1	2%	10	4%
Total	236		47		283	

An OPCS format cause of death was present in 94% of cases. Seventeen of the 18 cases where it was absent were hospital postmortem examinations, some of which were only partial examinations. The guidelines for postmortem examination reports state that it is good practice to include an OPCS cause of death. In many hospital postmortem examinations the death certificate is completed prior to the examination and sometimes the pathologist may feel it inappropriate to formulate a cause of death particularly if the examination is limited.

The cause of death when present was regarded as well formulated and accurate. However, it was apparent that there was widespread variation in practice between pathologists, including the advisors, particularly with respect to where and whether the operative procedure was mentioned in the cause of death. There was frequent use of the formula whereby the disease for which the operation was performed was given as the cause of death with "operated on" ... in brackets after the disease. This variation presumably reflected local practice, particularly the preference of the Coroner.

Formulating the cause of death can be difficult. There is the additional pressure on the pathologist of knowing that often the manner in which the cause of death is formulated decides whether there will be an inquest. Postoperative deaths are frequently complicated but the pathologist's position is particularly difficult if he has inadequate clinical information, has to interpret complex clinical events or has not had the opportunity of discussing the case with the surgeon. It has been argued that where the cause of death requires interpretation of the clinical history, a pathologist should refrain from giving a cause of death; such a task is for the court.⁶¹ It is certainly not the responsibility of the pathologist to decide whether an inquest is necessary.

Depending on local Coronial practice when formulating a cause of death the pathologist may need to take into account whether the surgery was emergency or elective, whether an error occurred, whether the fatal complication is regarded as "recognised" and whether there are allegations of negligence. These matters provoked lively discussion amongst the members of the group with little agreement. One basic problem is the lack of a consistent, clear purpose within the Coroners' system⁶³ perhaps particularly in relation to the investigation of postoperative deaths.

197

65%

3. CLINICOPATHOLOGICAL CORRELATION

Table PM17 Is a clinicopathological correlation present?

Yes

1 03		
No	104	35%
Total	301	
If yes, the clinicopathological correlation is:		
Unacceptably brief, obscure, uninformative	3	1%
Poor	9	5%
Satisfactory	63	32%
Good	57	29%
Fully detailed, clear, informative	65	33%
Total	197	

A clinicopathological correlation was present in a higher percentage of cases than previously (65%) but absent in 34% of hospital cases surprisingly considering that in many cases it is the only part of the report read by the clinician. In a postoperative death every postmortem examination report should contain a clinicopathological correlation, even if only a sentence or two, indicating whether the procedure played any part in the death and whether anything untoward or unexpected had happened. When present the correlation was felt to be satisfactory or better in virtually all cases and in one third was described as fully detailed, clear and informative, a much higher percentage than previous years.

Pathology 100

Table PM18
Overall score for the postmortem examinations

	Total		
Unacceptable, laying the pathologist open to serious	7	2%	
professional criticism			
Poor	34	11%	
Satisfactory	108	36%	
Good	112	37%	
Excellent, meeting all standards set by RCPath guidelines	40	13%	
Total	301		

Table PM19
Overall score for the postmortem examinations by grade of pathologist (from SQ)

					Overall	Score					
	Unsatisfa	ctory	Poor		Satisfacto	ory	Good		Excell	ent	Total
Consultant	7	3%	30	11%	98	36%	102	37%	36	13%	273
Junior	-	0%	-	0%	7	50%	5	36%	2	14%	14
Not answered	-		3		-		3		2		
Not known /	-		1		3		2		-		
not recorded											
Total	7		34		108		112		40		301

Eighty-seven percent of postmortem examinations were graded as satisfactory or better with the only difference to previous years being that in our figures 13% were graded as excellent, a much higher proportion than before probably reflecting our slightly "looser" interpretation of the need for histology. Seven postmortem examinations, all Coroners', were graded as unacceptable and laid the pathologist open to serious professional criticism. Several cases could not be assessed properly because of the restraints placed upon the pathologist's report by the Coroner, i.e. he only allowed a cause of death to be issued. All the small number of postmortem examinations performed by trainee pathologists were satisfactory or better.

Table PM20 (answers may be multiple)
When the history, antemortem clinical diagnosis and cause of death are compared with the postmortem findings, this postmortem demonstrates:

	Coroner's		Hospital		Total	
A discrepancy in the cause of death or in a major diagnosis, which if known, might have affected treatment, outcome or prognosis	14	6%	3	5%	17	6%
A discrepancy in the cause of death or in a major diagnosis, which if known, would probably not have affected treatment, outcome or prognosis	33	14%	8	13%	41	14%
A minor discrepancy	4	2%	-	0%	4	1%
Confirmation of essential clinical findings	175	74%	52	81%	227	75%
An interesting incidental finding	11	5%	7	11%	18	6%
A failure to explain some important aspect of the clinical problem, as a result of a satisfactory postmortem	4	2%	3	5%	7	2%
A failure to explain some important aspect of the clinical problem, as a result of an unsatisfactory postmortem	10	4%	-	0%	10	3%
Other	1		-		-	

Pathology 101

A small percentage of postmortem examinations showed a discrepancy which if known might have affected treatment, outcome or prognosis with a very large percentage confirming the principal clinical findings. These figures are very similar to previous years. The totally unexpected postmortem examination findings included leakage of TPN fluid into the pericardial sac; an unsuspected acute myocardial infarct; a jejunostomy leaking into the peritoneal cavity; an unexpected dilated cardiomyopathy; peritonitis due to traumatic perforation of the ileum; septicaemia due to an unsuspected perforated duodenal ulcer; unsuspected peritonitis from a gastrostomy tube; acute gastric dilatation; unsuspected pneumonia and peritonitis due to small bowel perforation. Many of these were in relatively minor procedures such as gastrostomy, illustrating the importance of leaving tubes and lines in situ prior to the postmortem examination.

Transmission of the postmortem examinations findings

Table PM21, question 52 Was the surgical team informed of the date and time of the postmortem?

	Coroners' postmortems	Hospital postmortems			All postmortems		
Yes	123	33%	50	49%	173	36%	
No	221		44		265		
Not answered	27		9		36		
Total	371		103		474		

Table PM22, question 52a (answers may be multiple)

If yes, which member of the surgical team attended the postmortem?

None	75	43%
House officer	12	7%
Senior house officer	22	13%
Specialist registrar	25	14%
Staff grade	1	1%
Clinical assistant	2	1%
Associate specialist	4	2%
Consultant	33	19%
Other	6	3%
Not answered	7	4%
Total	173	

Table PM23, question 53 (answers may be multiple)

If a surgeon did not attend the postmortem, why not?

(Only for those cases where the surgical team was informed of the date and time of the postmortem but none of the surgical team attended)

Unavailable/other commitments	45	60%
Nothing to be gained/diagnosis known	10	13%
Other	13	17%
Not answered	4	
Not known/not recorded	3	
Total	75	

Table PM24, question 54 Did the surgical team receive a copy of the postmortem report?

Yes	360	76%
Informal report/verbal message	33	7%
No	73	
Not answered	8	
Total	474	

In only a third of cases was the surgical team informed of the date and time of the postmortem. This was a figure similar to preceding years with Coroners' postmortems less well communicated. This low percentage did raise the issue of cases where the postmortem is done other than at the hospital where the operation was performed. This is, in terms of clinicopathological continuity, an undesirable occurrence. In several cases in the surgical questionnaire the surgeon stated that he had either been forbidden from attending the postmortem or felt that his presence was not allowed as he might influence the pathologist's judgement. In fact the Coroners' Rules (1984)⁶⁴ list the "deceased's' regular medical attendant" and "the hospital" (if the death occurred in hospital) amongst those to be notified by the Coroner when a postmortem is to be performed. These individuals/institutions have the right to be represented at the postmortem by a medical practitioner or in the case of a doctor to attend themselves.⁶⁴

In a quarter of cases the surgical team did not receive a copy of the postmortem report. This figure was very similar for hospital and Coroners' postmortems. There is no obvious explanation for this in the case of hospital postmortems but the attitude of Coroners to the dissemination of postmortem findings does vary. Some will not allow a report to be released. Others only after an inquest has been held. In some cases the surgeon filling in the surgical questionnaire may not have known that a report had been received by another member of the clinical team.

Table PM25, question 54a/question 44 How long after the patient's death was the first information received, giving the results of the postmortem?

	Corono postmo		Hospital All postmor		tmortems	
7 days or less	90	24%	28	27%	118	25%
8 days to 30 days	43	12%	23	22%	66	14%
31 days to 60 days	15	4%	3	3%	18	4%
More than 60 days	28	8%	6	6%	34	7%
Not answered*	195	53%	43	42%	238	50%
Total	371		103		474	

^{*} may indicate that no written information was received

The College guidelines recommend that initial information is transmitted to the clinician within a maximum of two days.

Pathology 103

Table PM26, Question 56

Did the pathological information confirm the clinical impression?

	Coron postm	er's ortems	Hospital All post postmortems		tmortems	
Yes	271	73%	75	73%	346	73%
No	60	16%	14	14%	74	16%
Not known/not recorded	26		8		34	
Not answered*	14		6		20	
Total	371		103		474	

^{*} may indicate that no written information was received

In three-quarters of cases the postmortem examinations confirmed the clinical findings but in 16% of cases there were significant unexpected findings.

References

- Hall V, Overton C, Hargreaves J, Maresh MJA Hysterectomy in the treatment of dysfunctional uterine bleeding. British Journal of Obstetrics & Gynaecology 1998 (vol 105) supp 17; page 16
- 2 Report of the RCOG Working Party on prophylaxis against thromboembolism in gynaecology and obstetrics. Ed: RW Shaw. London, 1995.
- 3 Low molecular weight heparins for venous thromboembolism. Drugs Therapeutics Bulletin. 1998;36:25-29.
- Standing Medical Advisory Committee: Report on the management of ovarian cancer, current clinical practices. 1991.
- *Report of the joint working party on anaesthesia in ophthalmic surgery.* Royal College of Anaesthetists & College of Ophthalmologists. London 1993.
- Benumof JL. *Laryngeal mask airway and the ASA difficult airway algorithm*. Anesthesiology 1996; **84**:686-699
- 7 Data provided to NCEPOD by Hospital Episode Statistics (Department of Health).
- Freeland AP, Bates GJ. *The surgical treatment of a pharyngeal pouch: inversion or excision?* Annals RCS Eng 1987; 69:57-58.
- 6 Koay CB, Bates GJ. Endoscopic stapling diverticulotomy for pharyngeal pouch. Clin Otolaryngol. 1996;21:371-6.
- Baldwin DD, Toma AG. Endoscopic stapled diverticulotomy: a real advance in the treatment of hypopharyngeal diverticulum. Clin Otolaryngol 1998;23:244-47.
- North American Symptomatic Carotid Endarterectomy Trial Collaborators. *Beneficial effect of carotid endarterectomy in symptomatic patients with high grade stenosis.* N Engl J Med 1991;**325**:445-53.
- European Carotid Surgery Trialists' Collaborative Group. Randomised trial of endarterectomy for recently symptomatic carotid stenosis: final results of the MRC European Carotid Surgery Trial (ECST). Lancet 1998;351:1379-87
- Royal College of Surgeons of England. *Guidelines for clinicians on medical records and notes*. London: RCS Eng, 1994.
- Tangkanakul C, Counsell C, Warlow C. Carotid endarterectomy performed under local anaesthetic compared to general anaesthetic (Cochrane Review). In: The Cochrane Library, Issue 3, 1998. Oxford: Update Software.
- Kaw M, Sekas G. Long-term follow-up of consequences of percutaneous endoscopic gastrostomy (PEG) tubes in nursing home patients. Dig Dis Sci 1994;39:738-43.
- 16 Light VL, Slezak FA, Porter JA, Gerson LW. Predictive factors for early mortality after percutaneous endoscopic gastrostomy. Gastrointest Endosc 1995;42:330-5.
- Bourdel-Marchasson I, Dumas F, Pinganaud G, Emeriau JP, Decamps A. *Audit of percutaneous endoscopic gastrostomy in long-term enteral feeding in a nursing home*. Int J Qual Health Care 1997;9:297-302.

References 105

- Rabeneck L, Wray NP, Petersen NJ. Long-term outcomes of patients receiving percutaneous endoscopic gastrostomy tubes. J Gen Intern Med 1996;11:287-93.
- Jarnagin WR, Duh QY, Mulvihill SJ, Ridge JA, Schrock TR, Way LW. The efficacy and limitations of percutaneous endoscopic gastrostomy. Arch Surg 1992;127:261-4.
- Fay DE, Poplausky M, Gruber M, Lance P. Long-term enteral feeding: a retrospective comparison of delivery via percutaneous endoscopic gastrostomy and nasoenteric tubes. Am J Gastroenterol 1991;86:1604-9.
- Nair RG, Dunn DC, Fowler S, McCloy RF. Progress with cholecystectomy: improving results in England and Wales. Br J Surg 1997;84:1396-8
- Wahba RW, Beique F, Kleiman SJ. Cardiopulmonary function and laparoscopic cholecystectomy. Can J Anaes 1995;**42**:51-63.
- Begos DG, Franco KL, Baldwin JC, Lee FA, Revkin JH, Modlin IM. *Optimal timing and indications for cholecystectomy in cardiac transplant patients*. World J Surg 1995;**19**:661-7.
- Dexter SPL, Martin IG, Marton J, McMahon MJ. Long operation and the risks from laparoscopic cholecystectomy. Br J Surg 1997;84:464-6.
- Williamson WA et al. Barrett's oesophagus. Prevalence and incidence of adenocarcinoma. Arch Int Med 1991;151:2212-6
- Streitz JM, Andrews CW, Ellis FHJ. *Endoscopic surveillance of Barrett's oesophagus. Does it help?* Journal of Thoracic and Cardiovascular Surgery 1993;**105**:383.
- Hoff SJ et al. *Prognosis of adenocarcinoma arising in Barrett's oesophagus*. Annals of Thoracic Surgery 1998;**65**:176-9.
- Merger RS, Merger D. Carcinoma of the oesophagus and tylosis. A lethal genetic combination. Cancer 1993; **72**:17.
- 29 Streitz JM, Ellis FH, Gibb PS, Heatley GM. Achalasia and squamous cell carcinoma of the oesophagus. Analysis of 241 patients. Annals of Thoracic Surgery, 1995;59:1604-9.
- 30 Choi TK, Siu KF, Lam KH. Bronchoscopy and carcinoma of the oesophagus. American Journal of Surgery 1984;**147**:757.
- Dittler HJ, Stewart JR. *Role of endoscopic ultrasonography in oesophageal cancer*. Endoscopy 1993;**25**:156.
- Muller JM, Erasmitt, Steizner M, Zieren U, Pichlmaier H. Surgical therapy of oesophageal cancer. Br J Surg 1990; 77: 845-857.
- O'Hanlon DM, Karat D, Shaw I, Scott D, Griffin SM. *Oesophagectomy: results from a single UK centre.* Br J Surg 1997; 84 (supplement 1): 14.
- Harrington KD. (1981). The use of methylmethacrylate for vertebral body replacement and anterior stabilisation of pathologic fracture dislocation of the spine due to malignant disease. J Bone Joint Surgery 63A: 36-47.
- Kostwick JP. (1983). Anterior spinal cord compression for lesions of the thoracic and lumbar spine. Spine 8: 512-531.

- Barron KD, Hirano A, Avaki S, Terry RD, (1989). Experience with metastatic neoplasms involving the spinal cord. Neurology 9: 91-106.
- Constans JP, DeDivitiis E, Dozelli R, Spaziante R, Mider JF, Hayer C. (1983). Spinal metastasis with neurological manifestations. Review of 600 cases. J Neurosurg 59: 11-118.
- DeWald RL, Bridwell KM, Prodromas C, Rodts MF (1985). Reconstruction spinal surgery as palliation for metastatic malignancies of the spine. Spine 10: 21-26.
- Fidler MW, (1986). Anterior decompression and stabilisation of metastatic spinal fractures. J Bone Joint Surgery 68B: 83-90.
- Siegal T, Tigra P, Siegal T. (1985). Vertebral body resection for epidural compression by malignant tumours. J bone Joint Surg 67A: 375-382.
- Tomita K, Toribatake Y, Kawanava N, Ohnari H, Kobe H. (1994). *Total en block spondylectomy and circum spinal decompression for solitary spinal metastasis*. Paraplegia 31: 36-46.
- Turner PL, Prince HG, Webb JK, Sokal MPJW. (1988). Surgery for malignant extradural tumours of the spine. J Bone Joint Surg 70B: 451-456.
- Enkaova E A, Doursounian L, Chatellier G, Mabesoone F, Aimard T, Saillent G. (1997) A critical appreciation of the pre-operative prognostic Takuhashi Score in a series of 71 cases. Spine 22: 2293-2298.
- Hammerberg KW. (1992). Surgical treatment of metastatic spine disease. Spine 17: 1148-1153.
- Takuhashi Y, Matsuzaki H, Toriyama S, Kawano H, Ohsaka S. (1990). Scoring system for the preoperative evaluation of metastatic spine tumour prognosis. Spine 15: 1110-1113.
- Kostwick JP, Errico JN, Gleason TF, Errico CC. (1988) Spinal stabilisation of vertebral body tumours. Spine 13: 250-256.
- 47 McLain R F, Wernstein JN. (1990). Tumours of the spine. Semin. Spine Surg 2: 157-180.
- Campling EA, Devlin HB, Hoile RW, Lunn JN. *The Report of the National Confidential Enquiry into Perioperative Deaths 1991/1992*. London 1993.
- 49 The Report of the National Confidential Enquiry into Perioperative Deaths 1993/1994. NCEPOD. London, 1996.
- The Royal College of Anaesthetists. *Guidelines for the Use of Non-steroidal Anti-inflammatory*Drugs in the Perioperative Period. The Royal College of Anaesthetists, London 1998
- Campling EA, Devlin HB, Hoile RW, Lunn JN. *The Report of the National Confidential Enquiry into Perioperative Deaths 1992/1993*. London, 1995.
- 52 The Report of the National Confidential Enquiry into Perioperative Deaths 1993/1994. NCEPOD. London, 1996.
- Gallimore SC, Hoile RW, Ingram GS, Sherry KM. *The Report of the National Confidential Enquiry into Perioperative Deaths 1994/1995*. London, 1997.
- Start RD, Dalargy-Aziz Y, Dorries CP, et al. Clinicians and the coronial system: ability of clinicians to recognise reportable deaths. British Medical Journal 1993; **306**: 1038-1041.

References 107

- Knight B, Thompson IM. *The teaching of legal medicine in British medical schools*. Medical Education 1986; **20:** 246-258.
- Leadbeatter S, Knight B. Reporting deaths to Coroners: all the legal aspects of dying need reexamining. British Medical Journal 1993; **306:** 1018.
- 57 Champ C, Tyler X, Andrews PS, Coghill SB. *Improve your hospital postmortem rate to 40-50 per cent, a tale of two towns.* Journal of Pathology 1992; **166:** 405-407.
- Lord JT. *Risk management in pathology and laboratory medicine*. Arch Pathol Lab Med 1990; **114:** 1164-1167.
- Valaske MJ. Loss control/risk management: a survey of the contribution of postmortem examination. Arch Pathol Lab Med 1984; 108: 462-468.
- The Royal College of Pathologists. *Guidelines for Postmortem Reports*. London, The Royal College of Pathologists, 1993.
- James DS, Leadbeatter S. *The use of personal health information in the Coroner's inquiry*. Journal of the Royal College of Physicians of London 1997; **31:** 509-511.
- Reid WA. Cost effectiveness of routine postmortem histology. *Journal of Clinical Pathology* 1987; 40: 459-461.
- 63 Cordner SM, Loff B. 800 years of coroners: have they a future? *The Lancet* 1994; **344**: 799-801.
- 64 The Coroners Rules 1984.

Appendix A - Glossary

Definition of the 1996/97 sample groups

Gynaecology – any gynaecological procedure.

Head and neck surgery – any procedure in the head and neck region including the base of skull and pharyngeal pouch surgery but excluding intracranial operations.

Minimally invasive surgery – the sample for this group was difficult to define as much debate remains about the most accurate title for this form of surgery, which is variously called minimally invasive surgery, minimal access surgery, endoscopic surgery, laparoscopic surgery and keyhole surgery. We have included minimal access general abdominal surgery, endoscopic thoracic and orthopaedic surgery and diagnostic or therapeutic endoscopies (excluding gynaecological laparoscopy and urological endoscopy).

Oesophageal surgery – any oesophageal procedure including specific oesophageal endoscopy.

Spinal surgery – any spinal procedure.

Urology - any urological procedure including urological endoscopy.

Admission

Elective - at a time agreed between the patient and the surgical service.

Urgent - within 48 hours of referral/consultation.

Emergency - immediately following referral/consultation, when admission is unpredictable and at short notice because of clinical need.

American Society of Anesthesiologists (ASA) Classification of Physical Status

- **ASA 1** a normal healthy patient.
- ASA 2 a patient with mild systemic disease.
- ASA 3 a patient with severe systemic disease that limits activity but is not incapacitating.
- ASA 4 a patient with incapacitating systemic disease that is a constant threat to life.
- ASA 5 a moribund patient who is not expected to survive for 24 hours with or without an operation.

(NCEPOD) Classification of operations

Emergency

Immediate life-saving operation, resuscitation simultaneous with surgical treatment (e.g. trauma, ruptured aortic aneurysm). Operation usually within one hour.

Urgent

Operation as soon as possible after resuscitation (e.g. irreducible hernia, intussuception, oesophageal atresia, intestinal obstruction, major fractures). Operation within 24 hours.

Scheduled

An early operation but not immediately life-saving (e.g. malignancy). Operation usually within three weeks.

Elective

Operation at a time to suit both patient and surgeon (e.g. cholecystectomy, joint replacement).

Out of hours

NCEPOD's definition of out-of-hours operating includes all operations started between 18.01 and 07.59 on a weekday, as well as operations performed at any time on a Saturday or Sunday.

Recovery and special care areas

(Definitions used by the Association of Anaesthetists of Great Britain and Ireland)

High dependency unit

A high dependency unit (HDU) is an area for patients who require more intensive observation, treatment and nursing care than can be provided on a general ward. It would not normally accept patients requiring mechanical ventilation, but could manage those receiving invasive monitoring.

Intensive care unit

An intensive care unit (ICU) is an area to which patients are admitted for treatment of actual or impending organ failure, especially when mechanical ventilation is necessary.

Recovery area

A recovery area is an area to which patients are admitted from an operating theatre, and where they remain until consciousness has been regained, respiration and circulation are stable and postoperative analgesia is established.

Appendix B - Abbreviations

AAA Abdominal aortic aneurysm

ACE (inhibitors) Angiotensin-converting enzyme

AF Atrial fibrillation

ASA American Society of Anesthesiologists

BP Blood pressure

BSO Bilateral salpingo-oophorectomy

CABG Coronary artery bypass graft

CCF Congestive cardiac failure

CCU Coronary care unit

COAD Chronic obstructive airway disease

COPD Chronic obstructive pulmonary disease

CPAP Continuous positive airways pressure

CRF Chronic renal failure

CT Computerised tomography

CVA Cerebro vascular accident

CVP Central venous pressure

D & C Dilatation and curettage

DU Duodenal ulcer

DVT Deep vein thrombosis

ECG Electrocardiogram

ERCP Endoscopic retrograde cholangio-pancreatography

EUA Examination under anaesthesia

FEV1 Forced expiratory volume, one second

FNAC Fine needle aspiration cytology

GA General anaesthetic

GI Gastrointestinal

GU Genito-urinary

HDU High dependency unit

ICU Intensive care unit

IDDM Insulin-dependent diabetes mellitus

IHD Ischaemic heart disease

IM Intramuscular

INR International normalised ratio

LA Local anaesthetic

LVF Left ventricular failure

MI Myocardial infarction

MRI Magnetic resonance imaging

MRSA Methicillin resistant staphylococcus aureus

MUA Manipulation under anaesthesia

NIDDM Non-insulin-dependent diabetes mellitus

NIRS Near infra-red spectroscopy

NMR Nuclear magnetic resonance imaging
NSAID Non-steroidal anti-inflammatory drug

OCTT Oral cuffed tracheal tube

OGD Oesophago-gastro-duodenoscopy

OLV One lung ventilation

ORIF Open reduction and internal fixation

PA Posterior anterior

PCA Patient controlled analgesia

PE Pulmonary embolism

PEEP Positive end expiratory pressure

PEG Percutaneous endoscopic gastrostomy

PM Postmortem

POSSUM Physiological and operative severity score for enumeration of mortality and morbidity

RTA Road traffic accident

SAP Systolic arterial pressure

SHO Senior house officer

SOB Short of breath

SpR 1,2,3,4 Specialist Registrar, years 1, 2, 3 and 4

TAH Total abdominal hysterectomy

TCC Transitional cell carcinoma

TCD Trans-cranial Doppler

TIA Transient ischaemic attack

TLV Two lung ventilation

TPN Total parenteral nutrition

TURBT Transurethral resection of bladder tumour

TURP Transurethral resection of prostate

Appendix C - NCEPOD Corporate structure

The National Confidential Enquiry into Perioperative Deaths (NCEPOD) is an independent body to which a corporate commitment has been made by the Associations, Colleges and Faculties related to its areas of activity. Each of these bodies nominates members of the Steering Group.

Steering Group (as at 1 October 1998)

Chairman

Emeritus Professor V R Tindall CBE

Other members

Mrs M Beck (Royal College of Ophthalmologists)

Dr J F Dyet (Royal College of Radiologists)

Dr M J Goldacre (Faculty of Public Health Medicine)

Dr H H Gray (Royal College of Physicians of London)

Mr G T Layer (Association of Surgeons of Great Britain and Ireland)

Professor V J Lund (Royal College of Surgeons of England)

Dr J M Millar (Royal College of Anaesthetists)

Dr A J Mortimer (Royal College of Anaesthetists)

Mr J H Shepherd (Royal College of Obstetricians and Gynaecologists)

Dr P J Simpson (Royal College of Anaesthetists)

Mr M F Sullivan (Royal College of Surgeons of England)

Professor P G Toner (Royal College of Pathologists)

Professor T Treasure (Royal College of Surgeons of England)

Dr D J Wilkinson (Association of Anaesthetists of Great Britain and Ireland)

Mr J Ll Williams (Faculty of Dental Surgery, Royal College of Surgeons of England)

Observers

Dr P A Knapman (Coroners' Society of England and Wales)

Dr V Chishty (Department of Health - England)

NCEPOD is a company limited by guarantee, managed by the Trustees.

Trustees

Chairman Professor V R Tindall

Treasurer Dr J N Lunn

Dr J Lumley

Mr J Ll Williams

Clinical Coordinators

The Steering Group appoint the Principal Clinical Coordinators for a defined tenure. The Principal Clinical Coordinators lead the review of the data relating to the annual sample and advise the Steering Group and write the reports. They may also from time to time appoint Clinical Coordinators, who must be engaged in active academic/clinical practice (in the NHS) during the full term of office.

Principal Clinical Coordinators

Anaesthesia

Dr G S Ingram

Surgery

Mr R W Hoile

Clinical Coordinators

Anaesthesia

Dr A J G Gray Dr K M Sherry

Surgery

Mr K G Callum Mr I C Martin

Funding

The total annual cost of NCEPOD is approximately £450,000 (1997/98). We are pleased to acknowledge the continued support of;

Department of Health (England)

Welsh Office

Health and Social Services Executive (Northern Ireland)

States of Guernsey Board of Health

Jersey Group of Hospitals

Department of Health and Social Security, Isle of Man Government

BUPA Hospitals Limited

Benenden Hospital

Nuffield Hospitals

St Martins Hospitals Limited

The London Clinic

This funding covers the *total* cost of the Enquiry, including administrative salaries and reimbursements for clinical coordinators, office accommodation charges, computer and other equipment as well as travelling and other expenses for the coordinators, Steering Group and advisory groups.

Appendix D - Protocol

The National Confidential Enquiry into Perioperative Deaths

(Protocol revised July 1996)

The National Confidential Enquiry into Perioperative Deaths (NCEPOD) is an independent body, launched in 1988, to which a corporate commitment has been made by the Associations, Colleges and Faculties related to its areas of activity. NCEPOD was incorporated in February 1995 as a company limited by guarantee.

NCEPOD is independent of the Departments of Health and of the other governmental and non-governmental organizations which support it financially. The Enquiry reviews clinical practice across England, Wales, Northern Ireland, Jersey, Guernsey and the Isle of Man. All NHS and Defence Medical Services hospitals are included as well as those managed by BUPA Hospitals Ltd, General Healthcare Group PLC, Nuffield Hospitals, St Martins Ltd, Benenden Hospital and Wellington Hospital.

Management of the Enquiry

NCEPOD is overseen by a Steering Group, the members of which are nominated by the following Associations, Colleges and Faculties:

Association of Anaesthetists of Great Britain and Ireland

Association of Surgeons of Great Britain and Ireland

Royal College of Anaesthetists

Royal College of Obstetricians and Gynaecologists

Royal College of Ophthalmologists

Royal College of Physicians of London

Royal College of Pathologists

Royal College of Radiologists

Royal College of Surgeons of England

Faculty of Dental Surgery of the Royal College of Surgeons of England

Faculty of Public Health Medicine of the Royal Colleges of Physicians of the UK

The Steering Group's responsibilities include agreement of the definition and method of the annual sample for detailed study and approval of the reports published by NCEPOD.

The Steering Group members appoint Trustees who are responsible for the management of the Enquiry and its business as specified in the Memorandum and Articles of Association.

Method of the Enquiry

The protocol, agreed in December 1988, was derived from that of the Confidential Enquiry into Perioperative Deaths (CEPOD), the report of which was published in December 1987.

The Enquiry reviews clinical practice and aims to identify remediable factors in the practice of anaesthesia, all types of surgery and now invasive procedures. Data are supplied on a voluntary basis; consultant clinicians in the relevant specialties are invited to participate. NCEPOD also reviews requests for and reporting of postmortem examinations.

The Enquiry collects information on deaths in hospital within 30 days of an invasive procedure involving local, regional or general anaesthetic or sedation. Maternal deaths are not included in the Enquiry (they are covered by the Confidential Enquiry into Maternal Deaths).

Reporting of deaths

NCEPOD depends on local reporters to provide data on deaths in their hospital(s). Most of the reporters in the public sector are consultant clinicians who have devised their own methods of obtaining the information; many have delegated the data collection to administrative staff. In the independent sector, hospital or nursing managers provide the data. When incomplete information is received, the NCEPOD staff contact the appropriate medical records or information officer or secretarial or clinical audit staff.

Reporters provide the following information, which is entered onto the computer database:

Name of authority/trust

Name/sex/hospital number of patient

Name of hospital in which the death occurred (and hospital where surgery took place, if different)

Date of birth, final operation and death

Surgical procedure performed

Name of consultant surgeon

Name of anaesthetist

The data collection year runs from 1 April to 31 March.

Sample for more detailed review

The nature of the sample changes annually, agreed before the start of the data collection. Information about a selected number of deaths is obtained by means of questionnaires. These are sent to consultant clinicians who are also asked to provide photocopies of relevant parts of the patient's notes (see "Confidentiality" below).

NCEPOD may collect data about patients who have survived more than 30 days after a procedure. These data are used for comparison with the data about deaths, or to review a specific aspect of clinical practice. Data from other sources may also be used.

Clinical Coordinators

The Trustees, on the recommendation of the Steering Group, appoint for a defined tenure Clinical Coordinators who lead the review of the data relating to the annual sample and advise the Steering Group and the Trustees.

They may also from time to time appoint Assistant Clinical Coordinators, who must be engaged in active academic/clinical practice (in the NHS) during the full term of office.

Advice

The Clinical Coordinators obtain advice from consultant clinicians in current practice. The Steering Group recommends the method of selection of these advisors.

Analysis and review of data

The collection, recording and analysis of data are managed by the Chief Executive of NCEPOD.

Reports

NCEPOD publishes a report which includes aggregated data and commentary on the information received. The reports are prepared by the Clinical Coordinators, Chief Executive and administrative staff and are reviewed in detail and approved by the Steering Group before publication.

General recommendations are made as well as specific points for individual specialties. Reports may include, as acknowledgements, the names of consultant clinicians who have returned questionnaires and whose names are known to NCEPOD.

Confidentiality

NCEPOD is registered with the Data Protection Registrar and abides by the Data Protection Principles. NCEPOD does not provide individual data to any person or organization outside the NCEPOD staff, coordinators, Trustees and Steering Group other than in the published Report.

All reporting forms, questionnaires and other paper records relating to the sample are shredded once an individual report has been published. Similarly, all patient-related data are removed from the computer database.

Data in the reports are aggregated to regional or national level so that individual Trusts, hospitals, authorities and patients and clinicians (who treat them) cannot be identified.

Before review of questionnaires by the Clinical Coordinators or any of the advisors, all identification is removed from the questionnaires and accompanying papers. The source of the information is not revealed to any of the Coordinators or advisors.

---End----

Appendix E - General data 1996/97

Reporting of deaths

Local reporters (see Appendix L) provided data to NCEPOD about patients who died in hospital between 1 April 1996 and 31 March 1997 within 30 days of a surgical procedure. We were also informed of a few deaths which occurred at home within the 30-day period.

Tables G1 to G4 refer to the total number of 19496 reported deaths. This total does not include nine reports where, despite all efforts, the data were incomplete, 168 reports received too late for inclusion in the analysis, and 969 inappropriate reports (see table G5). The regional breakdown of total deaths reported to NCEPOD, together with the totals of reports for previous years, are shown in table G1. The year-on-year fluctuations are in part caused by local (hospital) problems with data collection, and changes in the regional structure of the NHS. This means that the figures for years before 1996/97 are not directly comparable with those for earlier years.

Table G1

Deaths reported to NCEPOD

				Previous	years			
	1996/97	1995/96	1994/95	1993/94	1992/93	1991/92	1990	1989
Anglia & Oxford	1578	1672	1361	1577	1862	1556	1367	1371
North Thames	2292	2081	1944	2703	2515	2127	2554	2609
North West	2634	2736	2618	2636	2378	2509	2736	2864
Northern & Yorkshire	2870	3110	2549	2637	2671	2267	2464	2685
South & West	2201	2508	2469	2561	2493	1847	1997	2306
South Thames	2330	2166	2246	2531	2445	2465	2457	2840
Trent	2218	2397	2386	2342	2036	2014	1722	28 4 0 1849
West Midlands	1527	1595	1531	1578	1565	1578	1826	1902
Wales	1102	840	933	1078	1072	1079	1102	1162
Northern Ireland	480	469	497	529	474	375	316	380
Guernsey	27	33	12	33	26	18	39	
Jersey	18	26	17	27	32	25	22	32 26
Isle of Man	26	0	0	25	41	25	25	20 7
Defence Medical Services	8	7	17	36	40	75	60	94
Independent sector	185	201	148	149	166	172	130	94 120
Total	19496	19841	18728	20442	19816	18132	18817	20247

Table G2

Calendar days from operation to death (i.e. not 24 hour periods)

		%
0 (i.e. day of operation)	2092	10.7
1	2242	11.5
	1611	8.3
2 3 4 5	1243	6.4 36.9
4	1090	5.6
5	989	5.1
6	893	4.6
7	814	4.2
8	753	3.9 19.4
9	686	3.5
10	633	3.2
		<u></u>
11	598	3.1
12	491	2.5
13	496	2.5 12.7
14	436	2.2
15	451	2.3
16 to 20	1771	9.1
21 to 25	1324	6.8 20.4
26 to 30	883	4.5
		
Total	19496	

Figure G1 (see table G2)

Calendar days from operation to death (i.e. not 24 hour periods)

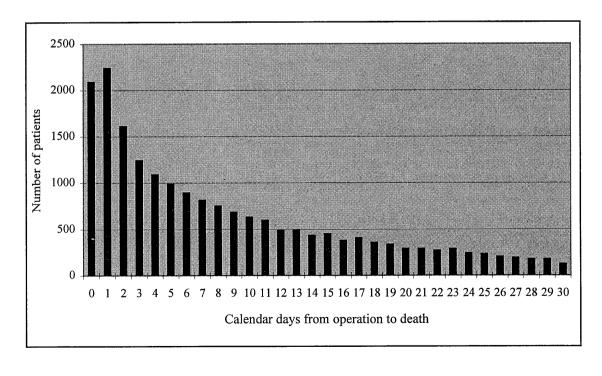


Table G3 **Age/sex distribution of reported deaths**

Ag	ge in :	years	Male	Female	Total
0	to	*4	129	94	223
5	to	9	16	7	23
10	to	14	16	13	29
15	to	19	49	13	62
20	to	24	55	20	75
25	to	29	62	43	105
30	to	34	68	47	115
35	to	39	77	58	135
40	to	44	152	103	255
45	to	49	221	141	362
50	to	54	267	205	472
55	to	59	453	336	789
60	to	64	783	476	1259
65	to	69	1253	817	2070
70	to	74	1840	1334	3174
75	to	79	1775	1476	3251
80	to	84	1441	1792	3233
85	to	89	909	1539	2448
90	to	94	312	785	1097
95	to	99	50	241	291
100	+		6	22	28
Total			9934	9562	19496

^{*} i.e. day of birth to the day preceding the fifth birthday

Table C4

Calendar days between death and receipt of report by NCEPOD

(i.e. not 24 hour periods)

: 1	to	29	4922
30	to	59	4144
60	to	89	2665
90	to	119	2240
120	to	149	1621
150	to	179	1142
180	or	more	2762
Total			19496

Table G5

Inappropriate reports received and not included

- 270 More than 30 days (day of operation to day of death)
- 247 Procedure not performed by a surgeon
- 239 Duplicate report
- 186 No surgical procedure performed or inappropriate procedure (according to NCEPOD criteria)
- 18 Procedure performed in non-participating hospital
- 5 Patient still alive (death wrongly reported)
- 4 Maternal death
- 969 Total

These figures do not include inappropriate reports included in computer printout format. Some hospital systems cannot easily filter out inappropriate reports such as deaths following procedures by physicians, or deaths following procedures excluded by NCEPOD.

Sample for detailed review

From a total of 19496 deaths reported to NCEPOD, the sample for this year comprised 2541 (13.0%) cases (see page 15 and Appendix A for details of the sample criteria)

Surgical questionnaires

Questionnaires were sent to consultant surgeons for further information on all 2541 cases, and 1808 completed questionnaires were returned to NCEPOD. The overall return rate (see table G6) was therefore 71.2% (1808/2541).

After excluding 111 questionnaires which were returned incomplete, related to the wrong operation or patient, or were returned too late to be included, 1697 questionnaires were analysed.

Anaesthetic questionnaires

Questionnaires were sent to consultant anaesthetists for further information on 1721 of the 2541 sample cases. No questionnaire was sent for the remaining 820 cases for the following reasons:

- Name of appropriate consultant anaesthetist unobtainable or notified too late to send questionnaire
- 660 No anaesthetist involved (local anaesthesia or sedation administered solely by the surgeon)

Of the questionnaires distributed, 76.5% (1316/1721) were returned. After excluding 44 questionnaires which were returned incomplete, related to the wrong operation or patient, or were returned too late to be included, 1272 questionnaires were analysed.

Table G6

Distribution and return of questionnaires by NHS region

SQ = Surgical Questionnaire AQ = Anaesthetic Questionnaire		. of Qs tributed		. of Qs turned	% return	-	No. o analy	•	No. ho	
	SQ	AQ	SQ	AQ	SQ	AQ	SQ	AQ	SQ	AQ
Anglia & Oxford	188	132	152	109	80.9	82.6	141	106	23	23
North Thames	266	156	161	108	60.5	69.2	144	103	33	30
North West	391	274	274	186	70.1	67.9	255	179	36	33
Northern & Yorkshire	392	250	279	205	71.2	82.0	267	199	37	37
South & West	275	196	229	163	83.2	83.2	215	155	23	20
South Thames	266	165	177	122	66.5	73.8	168	120	25	26
Trent	329	235	228	177	69.3	<i>75.3</i>	215	170	18	18
West Midlands	176	135	122	103	69.3	76.3	119	101	21	19
Wales	137	97	95	74	69.3	76.2	85	72	16	16
Northern Ireland	85	47	62	38	72.9	80.9	59	37	14	10
Guernsey	5	5	5	4	100.0	80.0	5	4	1	1
Jersey	2	2	2	2	100.0	100.0	2	1	1	1
Isle of Man	4	3	3	2	75.0	66.7	3	2	1	1
Defence Medical Services	2	2	2	2	100.0	100.0	2	2	1	1
Independent sector	23	22	17	21	73.9	95.5	17	21	11	12
Total	2541	1721	1808	1316	71.2	76.5	1697	1272	261	248

It is pleasing to note that in the NHS, several regions/authorities achieved return rates of 80% or higher; there remains considerable room for improvement in the other regions and particularly in North and South Thames.

Table G7
Distribution and return of surgical questionnaires by sample group

	Numb cases in		Numb question return	naires	90 question retur	nnaires	Numb question analy	nnaires
	SQ	AQ	SQ	AQ	SQ	AQ	SQ	AQ
Gynaecology	265	244	213	209	80.4	85.7	205	201
Head and neck surgery	466	335	346	238	74.2	71.0	315	229
Minimally invasive surgery	625	236	354	160	56.6	67.8	310	155
Oesophageal surgery	406	270	288	206	70.9	76.3	273	198
Spinal surgery	99	81	79	59	79.8	72.8	76	57
Urology	680	555	528	444	77.6	80.0	518	432
Total	2541	1721	1808	1316	71.2	76.5	1697	1272

Table G8
Reasons for the non-return of questionnaires

Surgical Questionnaires	Anaesthetic Questionnaires	
459	263	No reason given
102	85	Medical notes lost or unavailable
114	21	Surgeon/anaesthetist judged return of questionnaire 'inappropriate'
27	24	Surgeon/anaesthetist no longer working at the hospital or on sick-leave
7	2	Consultant did not wish to participate
24	10	Other
733	405	Total

Selected data from the completed anaesthetic and surgical questionnaires

Table G9 (q59 – surgical questionnaire)

Has this death been considered (or will it be considered) at a local audit/quality control meeting and/or was a postmortem examination performed?

	PM + audit	PM only	Audit only	Neither	Total
Gynaecological surgery	21	32	47	105	205
Head and neck surgery	61	23	106	125	315
Minimally invasive surgery	64	14	152	80	310
Oesophageal surgery	63	11	156	43	273
Spinal surgery	24	9	24	19	76
Urological surgery	124	28	243	123	518
Total	357	117	728	495	1697

Table G10 (q67a – anaesthetic questionnaire)

Do you have morbidity/mortality review meetings in your department?

Yes	1171
No	92
Not answered	9
Total	1272

If yes, will this case be or has it been discussed at your departmental meeting?

Yes	345
No	802
Not answered	11
Not known/not recorded	13
Total	1171

Although 64% of these deaths were considered at a surgical audit meeting, only 29% were discussed at an anaesthetic departmental meeting. In 7% of returned questionnaires it was stated that there was no anaesthetic morbidity/mortality review meeting at which such discussion could have taken place.

NATIONAL CONFIDENTIAL ENQUIRY INTO PERIOPERATIVE DEATHS

35-43 Lincoln's Inn Fields, London, WC2A 3PN

<u></u>	
9	
ဖွ	
9	
<u></u>	
٠.	
ш	
Œ	
=	
\simeq	
\leq	
z	
0	
Ě	
7	
S	
쁘	
ڀ	
O	
()	
Ē	
ш	
I	
F	
Ś	
Ш	
⋖	
Z	
7	
_	

QUESTIONNAIRE No.	A
ESTIONNAIRE NO.	A

QUESTIONNAIRE COMPLETION

This questionnaire should be completed with reference to the final operation before the death of the patient specified by NCEPOD. If you feel that this was not the main operation in the period before the patient's death, please give additional information.

Please **enclose** a copy of the ANAESTHETIC RECORD and of the FLUID BALANCE CHARTS. These are especially important. Any identification will be removed in the NCEPOD office.

All original copies of correspondence with NCEPOD are confidential (but do not retain copies of your correspondence).

The whole questionnaire will be shredded when data collection is complete.

For further information or for assistance, please contact the NCEPOD office on:

HAVE YOU ENCLOSED COPIES OF THE ANAESTHETIC RECORD AND FLUID BALANCE CHARTS?

Please use this section to provide extra information to help the advisory group of anaesthetists to understand aspects of this case which may not be apparent from your answers to the questions. The advisors who read this questionnaire are not apportioning blame. NCEPOD enquires into clinical practice in order to identify remediable factors in the practice of anaesthesia. It is intended to help clinicians to improve the care of patients.

Please write clearly, and ensure the accuracy of the information given.

							entry ye, di dili didi wa minay waga.	e and a special property.				
								And the state of t				
								-				Court of the Court

PROXY ANAESTHETISTS

If you	If you were not involved in any way with this anaesthetic and have completed the questionnaire on behalf of someone else, please indicate your position.	proce
<	Chairman of Division	Ques
ш ООШ	College tutor Duty consultant Other consultant Other (please specify)	ო
		4
	THE ANAESTHETISTS	5
Wha Plea	What was the grade of the most senior anaesthetist present at the start of the anaesthetic? Please enter the appropriate letter in the box.	
Trair	Trainee grade and years of training*	
C B A	Senior house officer - Year 1 Senior house officer - Year 2 Senior house officer - more than two years	
ОШГОТ	Specialist registrar - Year 1 Specialist registrar - Year 2 Specialist registrar - Year 3 Specialist registrar - Year 4 Specialist registrar - Accredited/CCST	
-	Other trainee - please specify	
Care	Career grades	ဖ
- スト	Consultant Staff grade Associate specialist	
Othe	Other grades	
∑zo	General practitioner Hospital practitioner Other (please specify)	^

We w	We want to procedure.	We want to know about the experience of this most senior anaesthetist in the operating room at the start of this procedure.	g room at the start of this
Ques	stions	Questions 3 to 7 inclusive refer to this anaesthetist.	
က	Year	Year of primary medical qualification	1 9 3a
	Plea	Please state country in which medical qualification awarded	35
4	Year	Year of first full-time anaesthetic training post	1 9 4
Ŋ	Whic	Which higher diploma(s) in anaesthesia were held at the time of the operation?	
			Year of award
	⋖	None	
	ω	Fellowship (Royal College, College or Faculty)	1 9 5b
		(please state country	
	ပ	DA (or Part 1 FRCA)	1 9 5c
	۵	Part 2 FRCA (physiology/pharmacology)	1 9 5d
	ш	Other (please specify)	1 9 5e
ဖ	# # the for #	If the most senior anaesthetist present at the start of the anaesthetic was <u>not</u> in a training grade, please enter the appropriate letters for weekly (i.e. more than 50 operations per year) NHS commitments in anaesthesia for the following:	training grade, please enter nmitments in anaesthesia
	∢	Cardiac surgery	
	ω	Children under 3 years old	9
	ပ	Neurosurgery	
	۵	None of the above	
7	If the	If the most senior anaesthetist at the start of the anaesthetic was <u>not</u> a consultant, where was consultant help available?	, where was consultant
	4 B O	A consultant came to the theatre before the end of the anaesthetic A consultant was available in the operating suite but not directly involved A consultant was available in the hospital, but was not present in the operating suite	ting suite
	ΩШ	A consultant was available by telephone Other (please specify)	

^{*} this is designed as if the Calman proposals were in place. If there is a delay in implementation, please complete the question as if the changes <u>had</u> taken place.

Grades of other anaesthetists present during the procedure.

œ

Please enter the appropriate letter(s) for the anaesthetist(s) present at the start of or during the procedure, excluding the anaesthetist mentioned in question 2. If there was more than one anaesthetist of the same grade, please enter the same letter for <u>each</u> of these individuals.

Tra	Trainee grade and years of training*	Start of anaesthetic	Later in the procedure
4	Senior house officer - Year 1		
∞	Senior house officer - Year 2		
ပ	Senior house officer - more than two years		
Q	Specialist registrar - Year 1		
ш	Specialist registrar - Year 2		
щ	Specialist registrar - Year 3		
g	Specialist registrar - Year 4		T
I	Specialist registrar - Accredited/CCST		
_	Other trainee - please specify		
Care	Career grades		
7	Consultant		
¥	Staff grade	Ţ	
_	Associate specialist		
Othe	Other grades		
Σ	General practitioner		
z	Hospital practitioner		
ο .	Other (please specify)	8a	- 88

Was advice sought, at any time, from another anaesthetist who was not present in the theatre at any time during the anaesthetic?

6

6	
	Iht?
Yes No	s, from whom was advice sought?
∢ ₪	If yes, f

THE PATIENT

9	Date of patient's birth							9
	_			Σ	≥	>	>	
Ξ	Date of admission to the hospital in which the final operation took place	ρ						
								Ξ
		۵	۵	Σ	Σ	>	>	
12	Time of admission to that hospital							12
:] -	esn)	24 hc	(use 24 hour clock)	<u>ફ</u>	
<u>ნ</u>	Was the patient transferred as an inpatient from another hospital?							
	A Yes B No							13a
	If yes, please explain the reason(s)							
					į		1	
	If yes, had the patient's condition apparently deteriorated during transfer?	er?						
	A Yes B No							13b
	If yes, please explain					J		
	THE OPERATION							
4	Please list any related surgical procedures carried out prior to this final operation.	Loper	ation	_				
	Procedure	Date						
							i	
	P	ĺ						
	Please enclose a copy of the anaesthetic records for these operations.							
15	What was the primary pre-operative diagnosis and planned final operation?	tion?						
							I	
							1	

^{*} this is designed as if the Calman proposals were in place. If there is a delay in implementation, please complete the question as if the changes. had taken place.

Co-e and s	Co-existing medical diagnoses at the time of the final operation. Please enter the appropriate letter in a box, and specify the disorder in the space next to the category.	tter	in a box,
⋖	none		∢
æ	respiratory		8
ပ	cardiac		ပ
۵	neurological		Q
ш	endocrine		ш
щ	alimentary		ட
g	renal		g
I	hepatic		I
-	musculoskeletal		_
7	vascular		7
¥	haematological		¥
_	genetic abnormality		_
Σ	obesity		≥
z	sepsis (specify site)		z
0	other (please specify)		0

What procedure(s) was performed (final operation)?

17

Please enclose a copy of the anaesthetic record and fluid balance chart(s).

operation.
the final
ification of
OD class
NCEPC
8

9

We ask you to take particular care with the answers to this question which are crucial to the analysis made by NCEPOD. Please note the specific definition of "Emergency".

•		
		LCCLC
i		3
•		•

8

A NCEPOD Emergency
B NCEPOD Urgent
C NCEPOD Scheduled
D NCEPOD Elective

NCEPOD Definitions - Classification of operation

Emergency

Immediate life-saving operation, resuscitation simultaneous with surgical treatment (e.g. trauma, ruptured aortic aneurysm). Operation usually within one hour.

gent

Operation as soon as possible after resuscitation (e.g. irreducible hernia, intussusception, oesophageal atresia, intestinal obstruction, major fractures). Operation usually within 24 hours.

Scheduled

An early operation, but not immediately life-saving (e.g. malignancy). Operation usually within 3 weeks.

Elective

Operation at a time to suit both patient and surgeon (e.g. cholecystectomy, joint replacement).

Was a record of the patient's weight available?	Which of the following investigations were done before the anaesthetic? This should include tests carried
A Yes B No	out in a referral hospital and for which the results were available before the final operation. Please enter the letter for each test.
If yes, what was this weight	Please write the results in the space next to the test name.
Vas	A None
ight available?	B Haemoglobingm.litre ⁻ '
A Yes	C Packed cell volume (haematocrit)
No yes, what was the height? cm	D White cell count
If no, the estimated height wascm	
Was an anaesthetist consulted by the surgeon (as distinct from informed) before this operation?	
A Yes B No	H Plasma electrolytes Na mol.litre ' m mol.litre '
Did an anaesthetist visit the patient before the final operation?	K m mol.litre¹
A Yes B No	Cl
If no, why not?	m mol.litre
	M Creatinine M
If yes, where?	N Serum albumin a. itre
Ward Outbatient department	O Bilirubin (total)
C Accident and Emergency department D ICU/HDU	P Glucose m mol.litre¹
	Q Amylase
	R Urinalysis (ward or lab)R
If Ves. was this anaesthetist present at the start of the final operation?	S Blood gas analysisS
A Yes	T Chest x-ray
B No	U Electrocardiography
	V Respiratory function tests
	W Special cardiac investigation (eg cardiac catheterization)
	X CAT scan/ultrasound/MRI/NIMR X
	Z Others relevant to anaesthesia (please specify)

		59	Time of start of anaesthetic (enter "X" in boxes if times not recorded)	(use 24 hour clock)
		93	Time of start of surgery	30
ASA	ASA GRADES			(use 24 hour clock)
25	Please enter the patient's ASA status prior to the final operation. (NB we do not use the E subclassification).	31	Time of transfer out of operating room (e.g. to recovery room, ICU)	319
Defir	Definitions of the ASA Grades			(use 24 flour clock)
ASA 1	1 a normal healthy patient		If you are not able to provide the times, please indicate the total duration of the operation (i.e. time of start of anaesthetic to time of transfer):	
ASA 2	2 a patient with mild systemic disease			\exists
ASA 3	3 a patient with severe systemic disease that limits activity but is not incapacitating	8		hours minutes
ASA 4	4 a patient with incapacitating systemic disease that is a constant threat to life	ş	natv	
ASA 5	5 a moribund patient who is not expected to survive for 24 hours with or without an operation			32
	PREPARATION OF THE PATIENT BEFORE THE OPERATION		Associate specialist Clinical assistant	
56	Did the patient receive intravenous fluid therapy in the 12 hours before induction?		G Start grade H Consultation Other release snecify	
	A Yes B No	33	Was there a trained anaesthetist's assistant (i.e. ODA, SODA, anaesthetic nurse) present for this case?	e) present for this case?
	If yes, please send copies of the fluid balance charts.		A Yes	33
27	Was it necessary to delay the anaesthetic to improve the patient's state before operation?		2]
	A Yes			
	If yes, please indicate which system(s) needed attention:			
	A Cardiac			
	B Respiratory 27b			
	C Metabolic			
	D Haematological			
	Please explain the reasons for the delay:			
78	Were premedication drugs prescribed?			
	A Yes B No			

31a

THE ANAESTHETIC

What drug(s) or other therapy was the patient receiving regularly at the time of operation (excluding premedication or drugs for anaesthesia)?

24

31b

Is there an anaesthetic record for this operation in the patient's notes?

FLUIDS DURING OPERATION

∀ 8	Yes No		37a	99	What type of anaesthesia was used?	
ıf yeş	If yes, were monitoring instruments already attached to the patient (i.e. from ICU or A&E)?	or A&E)?]			68
∀ B	Yes No		37b		C regional alone (47 to 48) D general and regional (40 to 48) E general and local infiltration (40 to 46)]
Plea	Please enter appropriate letter(s) in the boxes for monitoring devices used:				G sedation atone (49 to 50) G sedation and local infiltration (49 to 50) H sedation and regional (47 to 50)	
	In place Us already i	Used during induction c	Used during op		Please now answer the questions indicated in brackets, and then continue from question 51.	n 51.
⋖	ECG				GENERAL ANAESTHESIA	
в 0	pulse oximeter indirect BP			40	Was a nasogastric tube in place before induction?	
۵	oesophageal or precordial (chest wall) stethoscope				A Yes	70,
ш	O ₂ analyser		I		oz m	t g
щ	inspired anaesthetic vapour analyser				If yes, was it	
ŋ	expired CO ₂ analyser				a) aspirated before induction?	
I	airway pressure gauge					
7	ventilation volume				DA Yes	40b
¥	ventilation disconnect device	1	37c		(4)	
_	peripheral nerve stimulator				b) removed before induction?	
Σ	temperature (state site)	1			A Yes	400
z	urine output				ON ON]
۵	CVP	1_		41	Was cricoid pressure used at induction of anaesthesia?	
σ	direct arterial BP (invasive)				A Yes	
œ	pulmonary arterial pressure					F
S	intracranial pressure			45	Was the patient formally given oxygen to breathe spontaneously before induction?	
-	EEG/OFAM/evoked responses	Ι			30/A	
>	other (please specify)				S- 02 No	45
				43	Were the lungs inflated with an oxygen-rich mixture before tracheal intubation?	
Did	Did anything hinder full monitoring?				A Yes	43
⋖ Œ	Yes No		38	*	Who contracts and the second s]
1				‡	was suxamemonium given to tacintate tracheal intubation?	
If ye	If yes, please specify (e.g. bilateral arm surgery, radiotherapy, skin pigmentation, inaccessibility non-availability of monitors):	inaccessibilit	×		A Yes B No	44

38

TYPE OF ANAESTHESIA

37 Were monitoring devices used during the management of this anaesthetic?

A face mask (with or without oral airway)	49	Which sedative drugs were given for this procedure (excluding premedication)?	
B larvngeal mask		A inhalant	
D nasotracheal intubation		C benzodiazepine	49
E endobronchial	45	D sub-anaesthetic doses of IV anaesthetic drugs	
F tracheostomy	1	E other (please specify)	
G patient already intubated prior to arrival in theatre suite	20	Was oxygen given during the operation?	
H other (please specify)		A Yes	50a
Were there any problems with airway maintenance or ventilation?		yes	
No No	46	A routine	£
If yes, please explain		B otherwise indicated (please specify indications)	
	- 5	Where did this patient go on leaving the operating room?	
	1	A recovery area or room equipped and staffed for this purpose [∗]	
REGIONAL ANAESTHESIA		B high dependency unit*	<u>.</u>
If the anaesthetic included a regional technique, which method was used?		D specialised ICU	
A epidural - caudal		E ward	
B - lumbar		F another hospital	
C - thoracic		G other (please specify)	
D interpleural	ļ	H died in theatre	
E intravenous regional	52	Were you unable at any time to transfer the patient into an ICU, HDU etc?	
F cranial or peripheral nerve blocks (please specify)		30/X	[
G plexus block (e.g. brachial, 3-in-1 block)		No No	25
H subarachnoid (spinal)		ونماصين مصماما	
J surface (e.g. for bronchoscopy)		ıı yes, please explain	
Which agent was used? Please specify drug(s) and dosage(s):			1.
A local			
B narcotic	* DE	* DEFINITIONS	
C other (please specify)	A rec	A recovery area is an area to which patients are admitted from an operating room, where they remain until	n until

SEDATION (as opposed to General Anaesthesia)

How was the airway established during anaesthesia?

45

46

An intensive care unit (ICU) is an area to which patients are admitted for treatment of actual or impending organ failure, especially when mechanical ventilation is necessary.

A high dependency unit (HDU) is an area for patients who require more intensive observation, treatment and nursing care than can be provided on a general ward. It would not normally accept patients requiring mechanical ventilation, but could manage those requiring intensive monitoring.

RECOVERY ROOM

56 Was controlled ventilation used postoperatively?

Ф

_____56a

11 the patient did not enter a recovery room (option A on question 51), please move to question 56. 33 Were monitoring devices used, or investigations carried out during the management of this patient in the recovery room? A Yess B No If yes, please indicate which monitors were used. Enter the letter(s) in each appropriate box: C Indirect BP D Gesophageal or precordial (chest wall) stethoscope E O ₂ analyser C Indirect BP D Gesophageal or precordial (chest wall) stethoscope B H vanifiation volume J verifiator disconnect device K peripheral newe stimulator C DP D Gesophagea newe stimulator C Indirect BP D Gesophagea or precordial (chest wall) stethoscope B G airway pressure gauge H vanifiator volume J verifiator disconnect device K peripheral newe stimulator C DP D Grect arterial BP (invasive) M Unite output N Orther (please specify) S4 Time of transfer from recovery area: T cardiac output V other (please specify) S54 Time of transfer from recovery area: C intensive area of this patient go next (i.e. after the recovery room)? C intensive area of this patient go next (i.e. after the recovery room)? C intensive area of this patient of possible of CU E another hoosital	A Yes	yes	A routine management	B respiratory inadequacy	C cardiac inadequacy	D control of intracranial pressure or other neurosurgical indications	E part of the management of pain	F poor general condition of patient	G to allow recovery of body temperature																
rem did not enter a recovery room (option A on question 51), please re monitoring devices used, or investigations carried out during the mana overy room? Yes No es, please indicate which monitors were used. Enter the letter(s) in each ECG pulse oximeter indirect BP oesophageal or precordial (chest wall) stethoscope O ₂ analyser expired CO ₂ analyser airway pressure gauge ventilation volume ventilation volume ventilation state site) urine output CVP direct arterial BP (invasive) blood gas analysis pulmonary arterial pressure intracranial pressure cardiac output other (please specify) me of transfer from recovery area: "in boxes if not recorded) here did this patient go next (i.e. after the recovery room)? ward high dependency unit intensive care unit specialised (CU home another hospital	move to question 56.	gement of this patient in the	53		appropriate box:											530							54	(use 24 hour clock)	99
is so an y andone allocations of it is sendent.	ient did not enter a recovery room (option A on question 51), please ı	Were monitoring devices used, or investigations carried out during the manac recovery room?	Yes	No	If yes, please indicate which monitors were used. Enter the letter(s) in each	CCI	n les aximatar	pulse oximater		oesophageal or precordial (chest wall) stethoscope O, analyser	expired CO, analyser	airway pressure gauge	ventilation volume	ventilator disconnect device	peripheral nerve stimulator	temperature (state site)	CVP	direct arterial BP (invasive)	pulmonary arterial pressure	intracranial pressure	cardiac output	other (please specify)	Time of transfer from recovery area:	(enter "X" in boxes if not recorded)	Where did this patient go next (i.e. after the recovery room)? A ward B high dependency unit C intensive care unit D specialised ICU E home F another hospital G died in recovery area G died in recovery area

CRITICAL EVENTS DURING ANAESTHESIA OR RECOVERY

22

58 Was there any mechanical failure of equipment during anaesthesia or recovery?

recov	on any or the following events, which required specific freathent, occur during anaesthesia or immediate recovery (ie the first few hours after the end of the operation)?	Ē	ədiate	
∀ 80	Yes No		57a	
lf yes	If yes, please specify nature by insertion of the appropriate letter(s) in a box.			
⋖	air embolus		4	
6	airway obstruction		6 0	
ပ	anaphylaxis		O	
۵	arrhythmia	<u>.</u>		
ш	bradycardia (to or less than 50% of resting)		5/b E	
щ	bronchospasm		L	
_U	cardiac arrest (unintended)		ڻ ت	
I	convulsions	Ļ	I	
	disconnection of breathing system			
	hyperpyrexia (greater than 40°C or very rapid increase in temperature)		7	
¥	hypertension (increase of more than 50% resting systolic)		¥	
_	hypotension (decrease of more than 50% resting systolic)	L	1	
Σ	hypoxaemia (please state oxygen saturation)		Σ	
z	misplaced tracheal tube		z	
0	pneumothorax		0	
۵	pulmonary aspiration		۵	
σ	pulmonary oedema		σ	
œ	respiratory arrest (unintended)		Œ	
S	tachycardia (increase of 50% or more)		S	
-	unintentional delayed recovery of consciousness		_	
-	ventilatory inadequacy		n	
>	excessive spread of regional anaesthesia (eg total spinal, overextensive epidural)		>	
≥	wrong dose or overdose of drug		*	
×	other (please specify)		×	
Pleas	Please specify location of patient, treatment and outcome.			
		İ		

	⋖ മ	Yes No	58a
	if ye	If yes, please specify:	
	∢	equipment for IPPV	4
	m	suction equipment	a I
	ပ	syringe drivers	ပ
	۵	infusion pump	280 D
	ш	instrumental monitor (please specify)	ш
	ш	other (please specify)	<u>u</u>
	If the	If the patient died in the theatre please move to question 63	
29	What	What were the complications or events after this operation?	
	Pleas	Please enter a letter for each, and specify in the space below each category:	
	∢	ventilatory problems (e.g. pneumonia, pulmonary oedema)	4
	©	cardiac problems (e.g. acute LVF, intractable arrhythmias, post-cardiac arrest)	В
	ပ	hepatic failure	0
	Q	septicaemia	
	ш	renal failure	ш
	ш	central nervous system failure (e.g. failure to recover consciousness, CVA etc.)	<u>ш</u>
	g	progress of surgical condition	<u>o</u>
	I	electrolyte imbalance	I
	_	haematological disorder/coagulopathy	_
	7	other (please specify)	
	Pleas	Please give an account of any adverse events during this period.	

62

REMINDER

Have you enclosed copies of the anaesthetic record and fluid balance charts?

THANK YOU FOR TAKING THE TIME TO COMPLETE THIS QUESTIONNAIRE

You are advised for legal reasons not to keep a copy of this questionnaire, because a copy would form a part of the patient's medical record.

All material sent to NCEPOD is ultimately destroyed.

Please return the questionnaire and accompanying papers in the reply-paid envelope provided.

If you wish to inform the NCEPOD office of any other details of this case, please do so on a separate sheet, but include the number of the questionnaire.

CONSULTANT ANAESTHETISTS ONLY

We would like to publish the names of all consultants who have returned completed questionnaires.

Please help us by providing your initials and surname.

This page will be removed from the questionnaire on receipt.

Surname

Initials

NATIONAL CONFIDENTIAL ENQUIRY INTO PERIOPERATIVE DEATHS

35-43 Lincoln's Inn Fields, London, WC2A 3PN

1996/97
IRE 1
NN
ESTIC
IL QUI
GICAI
SUR

QUESTIONNAIRE COMPLETION

This questionnaire should be completed with reference to the last operation before the death of the patient specified by the NCEPOD office. If you feel that this was not the <u>main</u> operation in the period before the patient's death, you may give additional information.

The whole questionnaire will be shredded when data collection is complete. The information will be filed anonymously.

Neither the questions nor the choices for answers are intended to suggest standards of practice.

Please enclose a copy of all the relevant surgical operation notes, the postmortem examination report and the histology report if available. Any identification will be removed in the NCEPOD office.

Many of the questions can be answered by "yes" or "no". Please insert a tick (</) in the appropriate box.

Where multiple choices are given, please insert the tick(s) in the appropriate box(es).

Where more details are requested for an answer, please write in BLOCK CAPITALS.

In case of difficulty, please contact the NCEPOD office on:

HAVE YOU ENCLOSED COPIES OF THE OPERATION NOTES,
POSTMORTEM EXAMINATION NOTES AND THE
HISTOLOGY REPORT?

PLEASE USE THIS SECTION TO PROVIDE A BRIEF SUMMARY OF THIS CASE, ADDING ANY COMMENTS OR INFORMATION YOU FEEL ARE RELEVANT. (PLEASE WRITE CLEARLY FOR THE BENEFIT OF THE SPECIALIST GROUP WHO WILL BE REVIEWING THE QUESTIONNAIRE).

IF YOU WISH TO INFORM THE NCEPOD OFFICE OF ANY FURTHER DETAILS OF THIS CASE, PLEASE DO SO ON A SEPARATE SHEET, QUOTING THE QUESTIONNAIRE NUMBER.

S
Z
Ξ
三
Ā
Ω

			ADMISSION TO THE HOSPITAL IN WHICH THE OPERATION TOOK PLACE	LACE
Date of birth		4	Date of admission: Date of admission:	4
Date of <u>final</u> operation	α	Ŋ	Admission category:	
			a Elective - at a time agreed between patient and surgical service	a a
			b Urgent - within 48 hours of referral / consultation	
Female b 3			c Emergency - immediately following referral / consultation	0
			If elective, date placed on waiting list or entered into admission diary:	
What diagnosis was recorded in the notes at the time of admission?	ذ.			Sa
		9	What was the pathway for this admission?	
		ю	Transfer as an inpatient from another <u>acute</u> surgical hospital	rd .
		q	Transfer from another non-surgical hospital, nursing home etc.	
		O	Referral from a General Medical or General Dental Practitioner	0
		Ð	Admission following a previous outpatient consultation (please state date of outpatient attendance)	TO
	-		> > = = = = = = = = = = = = = = = = = =	
		Ф	Admission via A&E department	Φ.
		+-	Other (please state)	

A.B.	N.B. If the patient was transferred as an inpatient from another acute surgical hospital, I.e. option "a" in Q6, please answer Q7; otherwise go directly to Q8.	თ	Was the patient originally admitted to "your" hospital (i.e. in which the operation took place) under the care of the surgeon whose team undertook the operation?	ion took place) under the care of
Typ	Type of referring hospital:			Yes
ø	(District) General			
٩	University/Teaching b		If no, what was the source of referral to the Consultant Surgeon?	
ပ	Surgical Specialty c			
p	Other Acute/Partly Acute		a A medical specialty (please specify)	co.
o)	Community			
-	Defence Medical Services			
ō	Independent		b Another surgical specialty (please specify)	ه
모	Other (please specify)			9a
W	Why was the patient transferred?		c Same surgical specialty (please state reason for referral)	0
			d Other (please specify)	
			·	
Did	Did the patient's condition deteriorate during transfer?			
	Yes 7a		Date and time of transfer to surgical team undertaking operation.	
:	ON.			
H ye	If yes , please specify		> > M M Q Q	(use 24 hour clock)
			Date of first consultation following referral:	06
			J 0	W W X
	PREOPERATIVE CARE			
ρ	To what type of area was the patient first admitted in "your" hospital (i.e. in which the final operation took place)?	10	Was there any delay in either the referral or the admission of this patient?	
æ	Surgical ward (including surgical specialties)			Yes 10
q	Gynaecological/Obstetric ward b			S S
ပ	Medical ward			
ъ	Mixed medical/surgical ward d		lf yes, please specify.	
Φ	Geriatric ward			
—	Admission ward			
Б	A&E holding area (or other emergency admission ward)			
-	Day unit h			
-	Direct to theatre			
<u> </u>	lcu i			
×	Coronary care unit (CCU)			
Ε	HDN ndH			
_	Other (please specify)			

Spe	Specialty of Consultant Surgeon in charge <u>at the time of the final operation</u> (pleas tick only <u>one</u> box).	12	What was the grade of the most senior surgeon consulted before this operation?	
ಹ	General		a House Officer	Γ
q	General with special interest in Paediatric Surgery			w T
ပ	General with special interest in Urology			٥
ъ	General with special interest in Vascular Surgery			0
9				0
4				0
ō				4 -
_	Accident and Emergency			0
: . <u>_</u>	Ocracic			ء
			i Visiting Specialist Registrar	 T
. <u>*</u>	Neurosurgery 11		J Locum Appointment - Training (state grade)	. <u>.</u>
_	>		k Locum Appointment - Service (state grade)	. <u>×</u>
Ε	6		m Staff Grade	ε Ε
_			n Clinical Assistant	: c
0	ABOOOD		o Associate Specialist	: (
۵			p Consultant	,
2 م	Plactic			<u>.</u>
,	a citation			-
_				
S	Urology	13	Please state the working diagnosis by the most senior member of the surgical torm:	
	Vascular		(PLEASE USE BLOCK CAPITALS)	
_	Other(please specify)			l
				!
				ŀ
				l
				1
				l
		7		
		<u>:</u>	what operation was proposed by the most senior member of the surgical team? (PLEASE USE BLOCK CAPITALS)	
				ı
				1
				ı
				ſ
				ı
		1 5	What was the immediate indication for the proposed operation?	
			(TERME USE BLOCK CAPITALS)	
				ı
				1

Ξ

Please enter the patient's ASA status prior to the final operation. (NB we do not use the E subclassification).

	16	2		
-	N	က	4	2

American Society Of Anesthesiology (A.S.A.) Classifications Of Physical Status

Definitions

ASA 1 a normal healthy patient

ASA 2 a patient with mild systemic disease.

ASA 3 a patient with severe systemic disease that limits activity but is not incapaciting.

ASA 4 a patient with incapacitating systemic disease that is a constant threat to life.

ASA 5 a moribund patient who is not expected to survive for 24 hours with or without an operation.

Were there any coexisting problems (other than the main diagnosis) at the time of this operation?

1	<u> </u>	
Yes	₈	

ovided.
pace pro
n the s
disorder i
y the
specif
x and s
e box
ropriate bo
n each appr
eac
ţi r
put a tick ir
, please
lf yes,

Respiratory Malignancy Cardiac

ρ ပ σ

- Renal _
 - Gastrointestinal Haematological
 - Vascular Sepsis

17a

Ε

D _

- Diabetes mellitus Other endocrine Musculoskeletal Neurological
- Alcohol-related problems Other (please specify) Genetic abnormality Drug addiction Psychiatric

PREOPERATIVE PREPARATION

recautions or therapeutic manoeuvres were undertaken preoperatively (excluding anaesthetic room ement) to improve the patient's preoperative condition?
What p manage

8

Enter a tick in each appropriate box.

Cardiac support drugs or anti-arrhythmic agents

æ Ф ပ σ

- Intravenous fluids Gastric aspiration
- Correction of hypovolaemia
- Urinary catheterisation Blood transfusion
 - Diuretics

р 1 8

- Anticoagulants
- Vitamin K
- Antibiotics (pre- or intraoperative)

Bowel preparation (specify method used)

- Chest physiotherapy Oxygen therapy
- Airway protection (e.g. in unconscious patients)

Tracheal intubation

- Mechanical ventilation
- Others (please specify) Nutritional support
- Were any measures taken (before, during or after operation) to prevent venous thromboembolism? 9

	19	
>	ß	<u>8</u>

Yes	å	
>	_	

Please list any previous operations. Please send all relevant operation notes.		OPERATION
	53	Were there any delays (between admission and surgery) due to factors other than clinical?
Operation Date Specialty and grade of operating surgeon		Yes No No 23
as a second seco		If yes, please specify:
Q		
	24 (Classify this operation (see definitions below and choose the category most appropriate to the case).
		a Emergency
9	_	
	J	
	ס	Elective
	_	Definitions
Date of decision to operate:	Ø	Emergency Immediate life-saving operation, resuscitation simultaneous with surgical treatment (e.g. trauma, ruptured aortic aneurysm). Operation usually within one hour.
X X W W Q Q	Ω	Urgent Operation as soon as possible after resuscitation (e.g. irreducible hernia, intussusception, oesophageal atresia, intestinal obstruction, major fractures). Operation usually within 24 hours.
What was the anticipated risk of death related to the proposed operation?	O	Scheduled An early operation but not immediately life-saving (e.g. malignancy). Operation usually within 3 weeks.
a Not expected a Not expected b Small but significant rick	σ	
Definite risk		Operation at a time to suit both patient and surgeon (e.g. cholecystectomy, joint replacement).
d Expected		
if death was expected, please specify the anticipated benefit of the operation.		

Time of start of operation: (not including anaesthetic time)	25	We want to know about the experience and qualifications of the most senior operating surgeon (Questions 28 to 31)	verating surgeon.
Duration of operation (not including anaesthetic time):	(2000)		
hrsmins	28	Year of primary medical qualification	9
Which grades of surgeon were present in the operating room during the procedure?			
a House Officer	29 a	Year of first full-time surgical training post	1 9
b Senior House Officer - Year 1	q	Which higher diploma(s) in surgery were held at the time of the operation?	
s Senior House Officer - Year 2	0		
d Specialist Registrar - Year 1	٥		Year of award
e Specialist Registrar - Year 2	Φ	A None	
Specialist Registrar - Year 3	4 -	B Part 1 of Fellowship	6
g Specialist Registrar - Year 4	5		┪┝
n Specialist Registrar - Accredited/CCST	h 20	C Fellowship (Hoyal College, College or Faculty)	1
Visiting Specialist Registrar	. <u> </u>	(please state country if not UK	
Locum Appointment - Training (state grade)	<u>-</u>	D MS/MD	1
Locum Appointment - Service (state grade)	¥	F intercollegiate specially divloma	6
n Staff Grade	Ε		-
n Clinical Assistant	c .	(please state which diploma	
b Associate Specialist	0	F Other (please specify)	1 9
o Consultant	٩		
q Other (please specify)	ь		
	3	How long had this surgeon spent in this grade in this specialty?	
What was the grade of the most senior operating surgeon (as distinct from surgeons present or supervisory capacity)?	s present in an assisting		mths
A House Officer	a E	How many similar procedures had THIS surgeon performed in the previous 12 months?	months?
Senior House Officer - Year 1	٩	(If not known, please enter an estimate)	
Senior House Officer - Year 2	o	sanpaooid	
5 Specialist Registrar - Year 1	70		
Specialist Registrar - Year 2	Ф 32	If the most senior operator was not a consultant, was a more senior surgeon immediately available,	mediately available,
Specialist Registrar - Year 3	<u>_</u>	i.e. in the operating room/suite?	
3 Specialist Registrar - Year 4	Ō		se A
n Specialist Registrar - Accredited/CCST	٤		o Z
Visiting Specialist Registrar	 	if yes, please specify grade and location.	
Locum Appointment - Training (state grade)	, z/	Grade	
C Locum Appointment - Service (state grade)	*		
n Staff Grade	٤	Location	
n Clinical Assistant	C		
Associate Specialist	0		
) Consultant	۵		
q Other (please specify)	ъ		

59

88

56

22

32

Operation undertaken: (PLEASE USE BLOCK CAPITALS)		POSTOPERATIVE CARE	
Ö .	38	Was the patient admitted to an ICU or HDU* immediately after leaving the theatre suite?	
		a Intensive Care Unit	а 38
N.B. PLEASE ENCLOSE A COPY OF THE OPERATION NOTES.		Neither of the above	2 0
If the operation undertaken was different to that <u>proposed,</u> please explain.		If neither, was the patient admitted to an ICU or HDU after an initial period on a routine postoperative ward?	ward?
			a 383
		High Dependency Unit	
		c Neither of the above	O
	38a	If either a or b , after how many days postoperatively?	
Please state the dragnosts established at operation: (PLEASE USE BLOCK CAPITALS)		days days	
		If the patient <u>was</u> admitted to an ICU or HDU please answer questions 39 to 41.	
		If the patient was \underline{not} admitted to an ICU or HDU, please continue from question 42.	
Were there any unanticipated intra-operative problems?			
No No		* DEFINITIONS (as used by the Association of Anaesthetists of Great Britain and Ireland)	
If yes, please specify.		Intensive Care Unit (ICU): an area to which patients are admitted for treatment of actual or impending organ failure, especially when mechanical ventilation is necessary.	ig organ
		High Dependency Unit (HDU): an area for patients who require more intensive observation, treatment and nursing care than can be provided on a general ward. It would not normally accept patients requiring mechanical ventilation, but could manage those receiving invasive monitoring.	int and
Was the procedure performed SOLELY under local anaesthetic or sedation administered by the SURGEON?			
Yes No			
If yes, which of the following were recorded during or immediately after the procedure?			
a Blood pressure			
b Pulse b			
c ECG c ECG c 27a			
Other (please specify)			
None			

37

а 38а b

POSTOPERATIVE CARE

33

34

35

Discriatige from ICU/HDU was due to:		Please specify the postoperative complications:	
Death	c	Languarhann leastern attent blanding requirities transfer attent to the	٦
b Elective transfer to ward b	ס ע	riadinon radiophosoperative oreduing requiring translation	<u></u>
Pressure on beds	2 (•
d Other (please specify)	ד כ		. 42
	5 0	Veries alised sepais Wound infantion/labitopana	B (
		Anastomotic failure	D •
Was the patient subsequently readmitted to an ICU/HDU?	. 0	Cardiac arrest	. •
40 NO	۔	Low cardiac output/other cardiac problems	, <u> </u>
	-	Hepatic failure	. <u>-</u>
If yes , please give details.		Renal failure	<u> </u>
	¥	Endocrine system failure	. <u>~</u>
	_	Stroke or other neurological problems	<u>-</u>
	Ε	Persistent coma	E
	_	Other organ failure (please specify)	-
	0	Problems with analgesia	•
If the patient's condition warranted an admission to an ICU/HDU, were vou at any time unable to transfer the	đ.	DVT	۵
patient into an ICU/HDU within the hospital in which the surgery took place?	σ	Pulmonary embolus	<u> </u>
Se S	_	Fat embolus	_
	S	Orthopaedic prosthetic complication	ς
	+	Pressure sores	-
	Þ	Peripheral ischaemia	
ii yes, wny ?	>	Urinary tract infection	>
	*	Urinary retention/catheter blockage	*
	×	Ureteric injury/fistula	×
	>	Nutritional problems	<u>></u>
	7	Other(please specify)	z
			1
43		Was there a shortage of personnel in this case?	
		Yes	
		2	1
	=	If yes, which?	
	ro	Consultant surgeons	La.
	q	Trainee surgeons	٥
	O	Consultant anaesthetists	o
	р	Trainee anaesthetists	g G
	Φ	Skilled assistants	e 43a
	-	Nurses	-
	D	ODAs	50
	≖ .	Porters	ے ۔
	-	Other (please specify)	<u>-</u>

Discharge from ICU/HDU was due to:

DEATH		CAUSE OF DEATH (this is a facsimile of the death certificate: please complete it accordingly, using BLOCK CAPITALS).
Date of death:	D D M M Y Y	I (a) Disease or condition directly leading to death
Time of death:	(use 24 hour clock)	(b) Other disease or condition, if any, leading to I(a)
Place of death:		
	e	
b Recovery room c Ward	ی م	
d ICU/HDU	, p	(c) Uther disease or condition, if any, leading to (b)
e CCU or other specialised ICU	40 •	
f Home	-	
	D	
h Other (please specify)	۴	Uther significant conditions CONTRIBUTING TO THE DEATH but not related to the
Was cardiopulmonary resuscitation attempted?	Yes	
	S. ON	
	90	Was the death reported to the Coroner?
If no , was this a decision made preoperatively?	Yes A7a No	Yes Son No
		If yes, was a Coroner's postmortem examination ordered and performed?
What was the immediate clinical cause of death (this need not be a duplication of the death certificate)? (PLEASE USE BLOCK CAPITALS)	a duplication of the death certificate)?	Yes S0e No
		If a Coroner's postmortem examination was performed, please answer questions 52 to 58.
	51	If a Coroner's postmortem examination was not performed, was a hospital examination undertaken?
		Yes No
		If no, why not?

N.B. If a postmortem examination was \underline{not} performed, please move to question 59.

25	Was the surgical team informed of the date and time of the postmortem examination?	56	Did the nathological information confirm the clinical impraction?	
	Yes 752	:		Š
	ON			Ž
52a	If yes, which member of the surgical team attended the postmortem examination?		None receive	ě
	a None			
	b House Officer		If no, what was different?	
	c Senior House Officer			
	d Specialist Registrar			
	e Staff Grade 652a			
	f Clinical Assistant	22	Who performed the postmortem examination?	
	g Associate Specialist g			
	Consultant			
	i Other (please specify)		b Junior pathologist c Not known	
53	If a surgeon did not attend the postmortem examination, why not?	28	Please specify the specialty of the pathologist who performed the postmortem examination.	÷
			a General histopathologist	
			b Home Office histopathologist	
54	Did the surgical team receive a copy of the postmortem examination report?		c Neuropathologist	
	A Yes		d Paediatric patriologist e Non-histopathologist (ed. chemical pathologist)	
	Informal report or verbal message c			
54a	If yes, when was this received?			
	>			
22	Please list the relevant findings of the postmortem examination including histology. (PLEASE USE BLOCK CAPITALS)			
	PLEASE SEND A COPY OF ALL POSTMORTEM EXAMINATION REPORTS, POSTMORTEM REQUEST FORM AND HISTOLOGY REPORTS IF AVAILABLE			

c b 57

а с 58 e d d

a b c

Yes None received

Has this death been considered, (or will it be considered) at a local audit/quality control meeting?

29

9

9

Ves 59		Yes 60	I	Yes No 61		В	٩	0	d og	Ф	-	5	پ		Yes 63	,
	Did you have any problems in obtaining the patient's notes (i.e. more than 1 week)?		If yes, how long did they take to reach you?	Were all the notes available?	If no, which part was inadequate/unavailable?	a Preoperative notes	b Operative notes	c Postoperative notes	d Death certificate book	e Nursing notes	f Anaesthetic notes	g Postmortem report	h Other notes (please specify)	Please state the grade of the surgeon who completed this questionnaire	Has the consultant surgeon seen and agreed this questionnaire?	Date questionnaire completed

82

အ

8

THANK YOU FOR TAKING THE TIME TO COMPLETE THIS QUESTIONNAIRE

YOU MUST NOT KEEP A COPY OF THIS QUESTIONNAIRE

NCEPOD 35-43 Lincoln's Inn Fields LONDON WC2A 3PN

Please return it in the reply-paid envelope provided to:

THIS QUESTIONNAIRE IS THE PROPERTY OF NCEPOD

Same Vice Same

Appendix H - Pathology proforma 1996/97

NCEPOD No.	Type of autopsy	☐ Hospital ☐ Coroner ☐ Other
GENERAL FEATURES OF THE A	UTOPSY REPORT	
The report is typewritten		□ Yes □ No
A clinical history is provided When present the clinical history is		☐ Yes ☐ No ☐ 1 Unacceptably brief, obscure, uninformative ☐ 2 Poor ☐ 3 Satisfactory ☐ 4 Good ☐ 5 Fully detailed, clear, informative
A summary of lesions is present When present this corresponds accurate	ely to the text report	☐ Yes☐ No☐ Yes☐ No
An OPCS cause of death is present When present this corresponds accurate When present this follows OPCS forms		 ☐ Yes ☐ No ☐ Yes ☐ No
A clinico-pathological correlation is pr When present the clinico-pathological		☐ Yes ☐ No ☐ 1 Unacceptably brief, obscure, uninformative ☐ 2 Poor ☐ 3 Satisfactory ☐ 4 Good ☐ 5 Fully detailed, clear, informative
SPECIFIC FEATURES OF THE AU	TOPSY REPORT	
The description of external appearance	s is	 □ 1 Unacceptably brief, inadequately detailed □ 2 Poor □ 3 Satisfactory □ 4 Good □ 5 Fully detailed, clear, informative
Scars and incisions are measured		□ Yes □ No □ N/A
The gross description of internal organ	s is	 □ 1 Unacceptably brief, inadequately detailed □ 2 Poor □ 3 Satisfactory □ 4 Good □ 5 Fully detailed, clear, informative
Organs weighed (paired organs score 1) The skull and brain have been examined. The operation site is described. The gross examination is appropriate to Samples have been taken for:	ed	□ 2 □ 3 □ 4 □ 5 □ 6 □ 7 □ 8 □ 9 □ Yes □ No □ N/A □ Yes □ No □ 1 Histology □ 2 Microbiology □ 3 Toxicology □ 4 Other □ 5 None of these

In my	judgement samples should have been taken for:	☐ 1 Histo ☐ 2 Micro ☐ 3 Toxio ☐ 4 Other ☐ 5 None	obiology cology r
	ology report is included with the PM report present the histological report is:	☐ 2 Poor ☐ 3 Satist ☐ 4 Good	· ·
	absent does the lack of histology detract significantly from ue of this report?	□ Yes	□ No
My ove	erall score for this autopsy is:	to se □ 2 Poor □ 3 Satisf □ 4 Good □ 5 Excel	•
When	CAL RELEVANCE the history, antemortem clinical diagnosis and cause of dea demonstrates (more than one answer will often apply):	th are comp	pared with the postmortem findings, this
□ 1	A discrepancy in the cause of death or in a major diagnosis, which if known, might have affected treatment, outcome or prognosis		
□2	A discrepancy in the cause of death or in a major diagnosis, which if known, would probably not have affected treatment, outcome or prognosis		
□ 3	A minor discrepancy		
□ 4	Confirmation of essential clinical findings		
□ 5	An interesting incidental finding		
□ 6	A failure to explain some important aspect of the clinical pro	blem, as a re	esult of a satisfactory autopsy
□ 7	A failure to explain some important aspect of the clinical pro	blem, as a re	esult of an unsatisfactory autopsy

ANY FEATURES WHICH MIGHT BE QUOTED IN THE NCEPOD REPORT:

Appendix J - Participants

Consultant anaesthetists

These consultant anaesthetists returned at least one questionnaire relating to the period 1 April 1996 to March 1997. We are not able to name <u>all</u> of the consultants who have done so as their names are not known to us.

Adams H.G.
Adley R.
Ahmed M.
Aitkenhead A.R.
Akhtar M.
Akinpelu O.E.
Al Quisi N.K.S.
Al-Kassim N.
Al-Shaikh B.Z.
Alagesan K.
Albin M.Z.
Alexander C.A.
Allan D.J.
Amin S.
Ammar T.A.A.
Anandanesan J.
Anderson J.
Anderson J.D.
Andrew D.S.
Andrews C.J.H.
Andrews C.J.H. Andrews J.I.
Ankutse M.
Antrobus J.H.L.
Appleby J.N.
Aquilina R.
Archer P.L.
Armstrong J.
Arnold R.W.
Arrowsmith A.E.
Ashby M.W.
Ashurst N.H.
Aspbury J.N.
Astley B.A.
Ather T.
Atkinson R.E.
Auterson T.
Avery A.F.
Baichoo S.
Bailey C.R.
Bailey D.M.
Bailey P.W.
Baker G.M.
Balakrishnan P.H.
Baldwin L.N.
Ballance P.G.
Ballard P.K.
Balmer H.G.R.
Bansal O.P.
Bardgett D.M.M.
Bardosi L.
Basu S.
Baxter R.C.H.
Bayoumi M.
Beck G.N.
Beechey A.P.G.

n ump
Beers H.T.B.
Bembridge J.L. Benedict M.
Bennett A.
Bennett J.A.
Bennett S.R.
Berry C.
Bexton M.
Bhaskar H.K.
Biggart M.
Bird T.M.
Birks R.J.S.
Bishton I.
Blackburn A. Blake D.
Blogg C.E.
Bloor G.K.
Blundell M.D.
Boaden R.W.
Boobyer M.D.
Bousfield J.D.
Bowley C.J.
Bowman R.A.
Bowring D.
Boyd J.D.
Boyd T.
Boyd V. Boyle A.S.
Brampton W.
Bramwell R.G.B.
Bray B.M.
Breen D.P.
Brice D.D.
Brighouse D.
Brighten P.W.
Brim V.B.
Bristow A.S.E. Broadley J.
Brodrick P.M.
Brooker J.
Brooks A.M.
Brooks R.J.
Brown C.
Brown J.G.
Brown J.J.
Brown R.
Brownlie G.
Brunner M.D.
Bryan G.
Bryant J.D. Buckland R.W.
Duckiana IV. W.

Caddy J.M.
Campbell N.
Campbell W.I.
Canton J.L.
Carlisle R.J.T. Carnie J.C.
Carrie J.C. Carson D.
Carter M.I.
Casey W.F.
Cash T.I.
Cashman J.N.
Catling J.S.
Caunt J.A.
Chadwick I.S.
Chaffe A.G.
Challands J.
Chalmers E.P.D.
Chamberlain M.E.
Chambers J.J.
Chapman J.M.
Charters P.
Charway C.L.
Cheema S.
Chetty S.
Choudhry A.
Chow A.E. Christian A.S.
Chung R.A.
Chung S.K.N.
Church J.J. Clark G.
Clarke K.
Clayton K.C.
Clunie R.W.D.
Coad N.
Cobner P.G.
Cobner P.G. Cockroft S.
Cody M.
Coe A.J.
Coghill J.C.
Cohen D.G.
Cole J.R.
Collis R.
Colville L.J.
Conway M. Conyers A.B.
Cook L.B.
Cook M.H.
Cook P.R.
Cooper A.E.
Cooper A.R.
Cooper A.R. Cooper P.D.
Coote M.J.A.

Cotton B R
Cotton B.R. Counsell D.
Counsell D. Cowan P.
Craddock S.C.
Craddock S.C.
Craig J.F.
Cranston D. Creagh-Barry P.
Creagh-Barry P.
Cross G.D.
Culbert B.
Curran J.P.
Cyna A.
Dann W.L.
Darling R.
Dasey N.
Daum R.E.O.
Davidson D.G.D.
Davies G.
Davies J.R.
Davies K.J.
Davies M.
Davies P.R.F.
Davies P.R.F. Davies R.
Davies K. Davis I.
Day S.
de Courcy J.
De Silva T. Dearden N.M.
Dearden N.M.
Desborough R.C.
Deulkar U.V.
Devine A.
Dewar J.A.
Dhariwal N.K.
Dickson D.E.
Dixon J.
Dobson M.B.
Dodds C.
Doran B.R.H.
Dormon F.M.
Dowdall J.W.
Dowling R.M.
Doyle L.
Drummond R.S.
Du Boulay P.M.
Dua R.
Duggan J.E.
Duncan N H
Duncan N.H. Dunn S.R.
Dunnet J.
Dunnett I.A.R.
Duthie D.J.R.
Dwyer N.
Dyar O.
Earnshaw G.
Lintools (' D

Eatock C.R.

Eddleston J.M.

Disaw C
Edge W.G.
Edmondson L.
Edmondson R.S.
Edmondson W.C.
Edwards A.E.
Edwards G.
Edwards H.
El-Kholy M.
El-Rakshy M.
Elliot J.M.
Elliott D.J.
Elliott R.H.
Elsworth C.
Eltoft M.E.
Eltringham R.J.
Emam M.S.
Emery F.M.
Emmanuel E.R.
Enraght-Moony R.N.
Eppel B.
Erskine R.
Erwin D.C.
Evans C.S.
Evans G.
Evans I.M.
Evans K.
Evans R.J.C.
Ewart M.C.
Fahy L.T.
Fairbrass M.J.
Farling P.A.
Farquharson S.
Ferguson M.R.
Fergusson N.V.
Field L.
Firn S.
Fleming B.G.
Fletcher S.J.
Flynn M.J.
Fogarty D.
Ford P.
Forster D.M.
Forward R.
Francis G.A.
Francis R.I.
Frater R.A.S.
Freeman J.
Freeman R.
Frew R.M.
Frimpong S.
Frost A.R.
Fuzzey G.J.J.
Cabbatt D A

Gabbott D.A.

Gabrielczyk M.R.

Gademsetty M.K.

Coren A.

Cotter J.

Buist R.J.

Bull P.T.

Burns A.

Burchett K.R.

Gadgil P.S.	Harrison C.A.	Jingree M.	Linsley A.	Metias V.F.
Galizia E.J.	Harrison G.R.	Johnson C.J.H.	Linter S.P.K.	Mettam I.
Gallagher L.B.S.	Harrison K.M.	Johnson R.C.	Lintin D.J.	Michael W.
Ganado A.	Hartley M.	Johnston J.R.	Littler C.	Middleton C.M.
Ganepola S.R.	Harvey D.C.	Johnstone R.D.	Loh L.	Mikl B.
Gardner L.G.	Harvey P.B.	Jones D.I.	Longan M.A.	Millican D.L.
Garrett C.P.O.	Hasbury C.R.	Jones K.E.	Loveday R.	Milne B.R.
Gaston J.H.	Haslett W.H.K.	Jordan M.J.	Lowry K.G.	Milne L.A.
Gavin N.J.	Hawkins D.J.	Judkins K.	Ludgrove T.	Mishra K.P.
Gell I.R.	Hawkins T.J.	Kalia P.	Lung C.P.C.	Mitchell R.G.
Gerrish S.P.	Haycock J.C.	Kay P.M.	Luntley J.B.	Mitchell R.W.D.
Ghandour F.M.	Hebden M.W.	Keane T.K.	Lynch L.	Monks P.S.
Ghattas B.F.A.	Hegarty J.E.	Keeler J.	Macartney I.D.	Moon C.J.
Ghazala J. Ghosh N.	Hegde R.T.	Keep P.J.	McBride R.J.	Moore J.K.
Ghosh S.	Heidelmeyer C.F.	Kelly D.R.	McCallum I.J.	Moore K.C.
Gilbertson A.A.	Heining M.P.D.	Kelly E.P.	McCaughey W.	Morgan C.J.
Gill N.	Henderson P.A.L.	Kendall A.P.	McCleane G.	Morgan M.
Gill S.S.	Heneghan C.P.H. Hett D.	Kenny N.T.	McCluskey A.	Morgan R.
Gillbe C.		Keogh B.F.	McCrirrick A.	Morris S.
Glew R.	Higgins D.J. Hill S.	Kerr J.H.	McCulloch D.A.	Morriss G.W.
Goldberg P.		Kesseler G.	McDonnell J.	Mostert M.J.
Goldhill D.	Hitchings G.M. Hoad D.J.	Kestin I.G.	McEwen A.W.	Mowbray M.J.
Goodall J.R.	Hobbs G.	Khalil H.R.I.	McGeachie J.F.	Mudie L.L.
Goodrum D.T.	Hodgson R.M.H.	Khan M.Y.	McGuinness C.	Muirhead R.
Goodwin A.P.E.	Holdcroft A.	Kilpatrick S.M.	McHugh P.	Mulholland D.
Goodwin A.P.L.	Hollis J.N.	King M.	McHutchon A.	Mulrooney P.
Goold J.E.	Hopkinson J.M.	King T.A.	MacIntosh K.C.	Mulvein J.T.
Gothard J.W.W.	Hopkinson R.B.	Kipling R.M. Kneeshaw J.	Mackenzie S.I.P.	Murphy J.
Gough M.B.	Howard R.P.	Knibb A.A.	McKinnon P.	Murphy P.
Goulden P.	Howell R.S.C.	Knights D.	McLaren I.M.	Murray J.M.
Goulding S.	Howes D.	Knights D. Knowles M.G.	Maclaurin R.E.	Myatt J.K.
Govenden V.	Howie J.	Knowles M.G. Kocan M.	McLeod G.F.	Myint Y.
Graham S.	Huddy N.C.	Koehli N.	McPherson J.J.	Nalliah R.S.C.
Grant I.C.	Hughes J.	Kokri M.S.	McSwiney M.	Nandakumar C.G.
Gray A.J.G.	Hughes J.A.	Kong K.L.	McVey F.	Nandi K.
Gray B.M.	Hughes T.J.	Kotting S.C.	Madej T.H. Magee P.T.	Nash J.
Gray D.	Hull J.	Kouzel A.S.	Maher O.A.	Nash P.J.
Gray S.	Hunsley J.E.	Kraayenbrink M.A.	Mahoney A.	Nathanson M.H.
Green B.	Hunter S.J.	Kumar V.	Mahroo A.R.	Navaratnarajah M.N. Naylor A.F.
Green C.P.	Hurwitz D.S.	Lahoud G.	Male C.G.	Neasham J.
Green J.D.	Huss B.K.D.	Lake A.P.J.	Marjot R.	Newby D.M.
Greenbaum R.	Hutchings P.J.G.	Lamb A.S.T.	Marsh R.H.K.	Newman V.J.
Greenwell S.K.	Hutchinson A.	Landon K.	Martin A.J.	Newnam P.T.F.
Greenwood B.K.	Hutchinson H.T.	Langham B.T.	Martin J.W.	Newson C.
Greiff J.	Hutchinson S.E.	Lanham P.R.W.	Martin P.	Newton D.E.F.
Greig A.J.	Hutter C.	Lassey P.D.	Martin V.	Nicholl A.D.J.
Grieff J.	Hutton D.S.	Lavies N.G.	Mason J.S.	Nickalls R.
Griffiths R.	Imrie M.M.	Laxton A.G.P.	Mason R.A.	Noble H.
Groves J.	Inglis M.S.	Leach A.	Masri Z.	Noble J.
Grummitt R.	Iyer D.	Leach A.B.	Masters A.P.	Nolan J.
Gupta A.	Jackson D.G.	Leadbeater M.J.	Matheson H.A.	Norman J.
Gwinnutt C.L.	James P.	Leigh J.	Mathew P.	Normandale J.P.
Hackett G.H. Hamilton-Farrell M.R.	James R.H.	Leigh J.M.	Mathias I.M.J.	Norton A.C.
Hampson J.M.	Jamieson J.R.	Leith S.	Mathur N.K.	Nunn G.F.
Handy J.L.	Jappie A.G.	Lenoir R.	Matson A.M.	O'Connor M.
Hardwick P.B.	Jardine A.D.	Lesser P.J.A.	Matthews A.J.	O'Donoghue B.
Hardy I.	Jarvis A.P. Jasani A.	Levitt M.J.	Matthews R.F.J.	O'Donovan N.P.
Hargreaves C.G.	Jasan A. Jash K.	Levy D.	Mawson P.J.	O'Dwyer J.
Hargreaves R.		Lewis G.A.	Mayall R.M.	O'Keefe V.
Harmer M.	Javed E.B. Jeffs S.	Lewis M.A.H.	Mayor A.H.	O'Riordan J.A.
Harper J.	Jellicoe J.A.	Lewis R.P.	Maz S.	O'Shea P.J.
Harris C.	Jenkins B.J.	Liddle M. Lilburn J.K.	Mazumder J.K.	Oduro-Dominah A.
Harris D.N.F.	Jephcott G.	Lindop M.J.	Meadows D.P.	Okonkwo N.
Harris G.	Jeyapalan I.	Lindop M.J. Lindsay R.G.	Melville C.	Orton J.K.
Harris T.J.B.	Jeyaratnam P.	Lindsay W.A.	Mendel L. Mercer N.P.	Osborne M.A.
	•		MICHOLI IN.F.	Padfield A.

Pais W.A.
Palin P.H.
Panch G.
Papas S. Pappin J.C.
Park W.G.
Parker C.J.R.
Parmar N. Parnell C.J.
Parry H.M.
Parsloe M.
Patel R. Pathiratne L.S.I.
Paul S.
Peacock J.E.
Pearson R.M.G. Peden C.
Peebles D.J.
Pennefather S.H.
Pepperman M.L.
Pereira N.H. Perfettini J.
Perks D.
Perks J.S.
Peterson A.C. Peutrell J.M.
Phillips A.
Phillips A.J.
Phillips D.C. Phillips G.
Phillips K.A.
Phillips P.D.
Pick M.J. Piggott S.E.
Plumley M.H.
Plummer R.B.
Pole Y.L. Power K.J.
Pratt C.I.
Prince G.D.
Protheroe D.T. Pryle B.
Puddy B.R.
Purcell G.
Purcell-Jones G. Putnam E.A.
Quaife J.
Radford P.
Raftery S.M. Raghavjee I.V.
Raithatha H.H.
Raitt D.G.
Ralph S. Ralston S.
Ramachandran A.
Ramsden W.N.
Rana M.S. Ranasinghe D.
Randall N.P.C.
Randall P.J.
Rao D.S. Rao I.N.
Raper J.M.
Ravenscroft P.J.
Rawal S.B. Ray S.
Razis P.A.

Rebstein S.E. Reddy B.K. Redman D.R.O. Redman L.R. Reed P.A. Reeder M. Reid M.F. Reilly C.S. Remington S.A.M. Renshaw A.J.N. Renshaw V. Restall J. Rich P. Richards D.C. Richards E. Richardson J. Richmond M.N. Riddell G.S. Ridley S. Riedel B. Rimell P.J. Ritchie P.A. Rittoo D.B. Roberts C. Roberts D.R.D. Roberts F.L. Roberts P. Robertson S.M. Robinson D.A. Robinson K.N. Robson J.E. Rodgers R.C. Rogers C. Rollin A.M. Rose N. Routh G.S. Roylance J. Royston D. Ruddock J.M. Ruff S.J. Ruiz K. Russell M.H. Russell W. Ryall D.M. Ryan D.A. Ryan T.D.R. Saddler J.M. Sagar D.A. Sahal B.B. Saleh A. Salmon N.P. Salt P.J. Samaan A.A. Samaan A.K. Sanchez A. Sanikop S. Sansome A. Saxena S.N. Scallan M.J.H. Schmulian C.M. Schofield N.M.

Segath M. Seingry D. Sellwood W.G. Seymour A.H. Shah H. Shah M.M. Shah R.K. Shah Z.P. Shakir S.A. Shambrook A.S. Shammas R. Sharawi R. Sharpe T.D.E. Sharples A. Shaw D. Shaw T.C. Sherry K.M. Short S. Sides C.A. Silk J.M. Simpson D.A. Simpson J.C. Simpson M.E. Sinclair M. Singh P. Singh R.K. Skinner J.B. Smalberger J.M. Smith H.S. Smith I.D. Smith M. Smith M.B. Smith P.A. Smyth P.R.F. Sneyd J.R. Snow D. Somanathan S. Somerville I.D. Sowden G.R. Spargo P. Spelina K.R. Spencer E.M. Spencer I. Spiers S.P.W. Spilsbury R.A. Sritharan S. Stacey M.R. Stacey R.G.W. Stacey R.K. Stannard C. Stanton J.M. Steer B. Stevens J. Stock J.G.L. Stoneham J.R. Strang T. Stray C.M. Stride P.C. Strong J.E. Strube P. Stubbing J.F. Sugden J.C.

Tannett P.G. Tarr T.J. Tatham P.F. Taylor C. Taylor M. Teasdale A. Tehan B. Telford R. Thomas A.N. Thomas D.G. Thomas S. Thompson A.R. Thompson E.M. Thompson J.F.W. Thomson J. Thomson S. Thomson W. Thorn J.L. Thorniley A. Thornley B. Till C. Tinloi S.F. Tipping T. Titoria M. Tofte B.C. Tolley M.E. Tomlinson J.H. Tomlinson M.J. Toomey P. Trask M.D. Tring I.C. Trotter T. Turner D.A.B. Turner R.J.N. Turtle M.J. Tweedie D.G. Ulyett I. Uncles D.R. Underhill S. Utting H.J. Valentine J. Valijan A. van Miert M. Van Ryssen M.E.P. Vanner R.G. Vater M. Venkat N. Ventham P.A. Verghese C. Verma R. Vine P.R. Vohra A. Voice A. Wadon C.A. Wait C.M. Walker G. Walker M.B. Walker R.S. Walker T. Wallbank W.A. Walsh E.M.

Walters F.

Walton D.P.

Walton M.

Ward M.E.

Ward R.M.

Ward S.

Wark K.J. Waters J.H. Watkins T.G. Watson D.M. Watson N.A. Watson-Jones E. Watt J.W.H. Watt T. Weller R. West K.I. West S. Westbrook J. Weston G.A. Wheatley R.G. Whelan E. White D. White D.J.K. White M.J. White W.D. Whitehead E.M. Whittle M.J. Wilkey A.D. Wilkinson P.A. Will R. Williams A.C. Williams C. Williams D.J.M. Williams E.G.N. Williams L.J. Williams P.J. Williams S.H. Williams T.I.R. Wilson A.J. Wilson I. Wilson P.T.J. Wiltshire S.J. Windsor J.P.W. Woods S.D. Woodsford P.V. Woollam C.H.M. Wright M.M. Wyatt R. Yanny W.A. Yates D.W. Yeoman P. Young J.D. Young P.N. Youssef H. Zideman D.A.

Schwarz P.A.

Scott-Knight V.

Seagger R.A.

Searle A.E.

Scott A.D.

Scott R.

Summerfield R.J.

Sutton B.A.

Sweeney J.E.

Swinhoe C.F.

Swayne P.

Appendix K - Participants

Consultant surgeons/gynaecologists

These consultant surgeons/gynaecologists returned at least one questionnaire relating to the period 1 April 1996 to March 1997.

41 1DD	n rn	D 115		
Abel P.D.	Beacon J.P.	Browell D.	Cohen G.L.	Dhingra D.
Abrams P.	Bearn P.	Brown A.A.	Cole R.S.	Diamond T.
Ackroyd J.S.	Beeby D.I.	Brown G.J.A.	Colin J.F.	Dickerson D.
Adair H.M.	Beggs F.D.	Brown M.G.	Collins F.J.	Dixon A.R.
Adamson A.	Bell J.C.	Brown R.J.	Colmer M.R.	Docherty P.T.C.
Affifi R.	Bell M.S.	Brown S	Cook A.	Doig R.L.
Afify S.E.	Bell P.R.F.	Brown T.H.	Cook A.I.M.	Donaldson D.R.
Agarwal P.K.	Bellringer J.	Brunskill P.J.	Cooke R.S.	Donaldson R.A.
Agarwal P.N.	Benjamin J.C.	Buchanan G.	Cooksey G.	Donnelly R.J.
Agarwal S.	Benson E.A.	Budd D.W.G.	Cooper J.C.	Donovan R.
Aggarwal V.P.	Berry A.R.	Budd J.S.	Cooper M.J.	Dooley J.F.
Aitken R.H.	Best B.G.	Bull P.D.	Cooper Wilson M.	Dormandy J.A.
Alderman B.	Bett N.J.	Bulman C.H.	Corfield R.	Dornan J.C.
Aldoori M.I.	Betts C.D.	Bunker T.D.	Corrigan A.M.	Dossa M.
Alexander D.J.	Bevis C.R.A.	Burgess N.A.	Costello C.B.	Douglas D.L.
Ali D.	Bewick M.	Busfield H.M.B.	Cowen M.E.	Douglas P.S.
Ali M.L.	Billings P.J.	Butler C.	Cowie A.G.A.	Duke A.B.
Allan S.M.	Birch B.R.	Byrne P.O.	Cowie R.A.	Dunn D.C.
Allen M.	Birchall J.P.	Cade D.	Cox G.J.	Dunn M.
Allen N.	Birzgalis A.	Callaghan P.S.	Cox H.J.	Dunning J.
Alun-Jones T.	Bishop C.C.R.	Callum K.G.	Crabbe D.C.G.	Dunster G.D.
Amanat L.A.	Bishop M.C.	Calvert C.H.	Crane P.W.	Durham L.
Anderson G.E.	Bisson D.	Campbell J.B.	Cranley B.	Durning P.
Anderson I.D.	Black J.	Campbell W.B.	Cranston D.W.	Durrans D.
Anderson J.B.	Black J.E.	Carey D.	Crawshaw C.C.	Duthie J.
Anderson R.J.L.	Black P.D.	Carr R.T.W.	Crisp J.C.	Duthie J.S.
Andrews N.	Blackburn C.W.	Carroll R.N.P.	Cross A.T.	
	Blacklock A.R.E.	Carswell W.		Dyke G.W.
Appleyard I.			Crumplin M.K.H.	Dzumhur S.
Archbold J.A.	Blair S.D.	Carter P.G.	Cullen R.J.	Eardley I.
Archer T.J.	Blake J.R.S.	Case W.G.	Cumming J.A.	Earl P.D.
Armitage T.G.	Bloomfield M.D.	Cawthorn S.J.	Curley P.J.	Eaton A.C.
Armour R.H.	Bolger C.	Cetti N.E.	Curran F.T.	Ebbs S.R.
Ashworth M.F.	Bolger J.	Chan D.	Curry A.R.	Edge C.J.
Aukland P.	Bond E.B.	Chan K.K.	Curwen C.	Edmondson R.
Aylward G.W.	Booth C.M.	Chapman D.F.	Dahar N.A.	Edmondson S.
Badenoch D.F.	Borgaonkar S.S.	Chapman J.	Damale N.	Edwards J.L.
Bailey M.J.	Bose P.P.K.	Chappatte O.A.	Dan A.N.	Edwards M.H.
Bainbridge E.T.	Boughdady I.	Chapple C.R.	Das S.K.	Edwards P.
Baldwin D.L.	Boyd D.A.	Charnock F.M.L.	David V.C.	Eisenstein S.M.
Balfour T.W.	Boyd I.E.	Cheong-Leen P.	Davies M.	El-Shunnar K.S.
Ball A.	Boyd N.R.H.	Chia P.	Davies N.P.	Eldred J.M.
Ball A.J.	Boyd P.J.R.	Chilton C.P.	Dawson P.M.	Eldridge P.R.
Ball C.	Bradbrook R.A.	Chinegwundoh F.I.	Day J.B.	Elias J.
Bancewicz J.	Bradford R.	Chowdhury S.D.	De Bolla A.R.	Ellenbogen S.
Bannister J.J.	Bradley P.J.	Chumas P.D.	Deane A.M.	Ellis B.W.
Bardsley A.F.	Bradpiece H.A.	Clark J.	Deans G.	Emtage L.A.
Barr G.S.	Bramble F.J.	Clarke D.	de Boer F.	England P.C.
Barr H.	Brame K.G.	Clarke D.G.	Debrah S.	English P.J.
Barton D.	Bransom C.J.	Clarke S.	de Cossart L.M.	Enoch B.E.
Bashir E.	Brearley S.	Clarkson P.K.	Deiraniya A.	Evans A.S.
Bashir T.	Britton B.J.	Clayton J.K.	Denton G.W.L.	Evans B.T.
Bateman S.	Britton D.C.	Clearkin L.G.	Desai K.M.	Evans D.A.
Bates R.G.	Brooks J.	Clegg J.F.	Desmond A.D.	Evans G.H.
Batey N.R.	Brooks S.G.	Cobb R.A.	Devereux M.H.	Evans J.
Battersby R.D.E.	Brookstein R.	Cochrane J.P.S.	Dewar E.P.	Evans J.W.H.
Baxter J.N.	Brooman P.J.	Coen L.D.	Dhillon R.	Evans K.L.

Evans-Jones J.C.	Greenall M.	Holmes F.J.	Jones D.R.B.	Lemberger R.J.
Ewah P.O.	Greene D.	Holt S.D.H.	Jones J.A.H.	Lennox J.M.
Eyong E.	Greenway B.	Hopkin N.B.	Jones R.B.	Lennox M.S.
Faber R.G.	Griffiths D.A.	Hopkinson B.R.	Jones R.O.	
Fairbank J.C.T.	Griffiths E.		Jones R.O. Jones S.E.	Leopard P.J.
		Hopper M.S.C.		Leopold P.W.
Farrington W.T.	Grimaldi P.M.G.	Horgan K.	Joseph T.	Leveckis J.
Fawcett I.	Grotte G.J.	Horner J.	Joyce M.	Lewis B.V.
Feggetter J.G.W.	Gudgeon M.	Horrocks M.	Jurewicz A.	Lewis J.L.
Fellows G.J.	Guest J.	Houghton D.J.	Kadow C.	Lewis M.H.
Feneley R.C.L.	Gupta M.C.	Houghton P.W.J.	Kane J.F.	Liston T.G.
Fergus J.N.	Guy P.J.	Howard D.J.	Kapadia C.R.	Little G.
Ferrie B.G.	Gwynn B.R.	Howat J.M.T.	Kapila L.	Liu S.
Ferro M.	Hackman B.W.	Howell G.	Karanjia N.	Livingstone J.
Field E.S.	Hadley J.M.	Hudd C.	Karim O.	Lloyd D.A.
Fielder C.	Haggie S.J.	Huddy S.P.J.	Kay N.J.	Lloyd D.M.
Fiennes A.	Hale J.E.	Hughes-Nurse J.	Keane P.	Longstaff S.
Fife D.G.	Hall J.H.	Hulton N.R.	Kelleher J.P.	Longstan 5. Loosemore T.
Finan P.J.	Hall R.I.	Hunter G.	Kelly J.F.	Lopes A.
Flook D.	Halligan A.	Hurley P.	Kelly S.B.	Lotz J.C.
Fogarty P.	Hallissey M.T.	Hussain F.H.	Kemeny A.A.	Loughridge W.G.G.
Forbes P.B.	Hamer D.B.	Hussein I.Y.	Kennedy C.L.	Love A.
Fordham M.	Hamid S.	Hutchinson C.	Kennedy R.H.	Lowry J.C.
Fossard D.P.	Hammad Z.	Hutchinson D.	Kernohan R.	Lucas M.G.
Foster M.E.	Hammonds J.C.	Hutchinson I.F.	Kerr-Wilson R.	Lund V.
Fountain S.	Haque M.	Iftikhar S.Y.	Khalil-Marzouk J.F.	Lyndon P.J.
Fountain S.W.	Hardcastle J.D.	Ilankovan V.	Khan I.A.	Lyttle J.A.
Fowler C.G.	Hardingham M.	Inglis J.A.	Khan M.A.R.	McClelland R.
Fox G.C.	Harland R.N.L.	Ingoldby C.J.H.	Khan O.	McCloy R.F.
Fox G.C. Fozard J.B.J.	Harris D.L.	Ingram G.	Khazim R.M.	McCollum C.N.
Fraser I.A.	Harris V.G.	Ingram N.P.	Kidd L.C.	MacDermott S.
French M.E.	Harrison D.J.	Ireland D.	Kinder R.B.	MacDonagh R.
Frietes O.N.	Harrison G.S.M.	Ismal M.	Kingston R.E.	MacDonald R.C.
Gajraj H.	Harrison I.D.	Isworth R.A.	Kirby R.S.	McDonnell P.J.
Gallagher P.	Harrison J.D.	Jackson W.D.	Kirkpatrick P.J.	MacFie J.
Gallegos C.R.	Harrison J.M.	Jacob G.	Kitson J.	McGregor A.D.
Gammall M.M.	Harrison S.C.W.	Jacobs I.	Kiwanuka A.I.	McHugh D.
Gardner N.H.N.	Harrison T.A.	Jaganathan R.S.	Klimach O.	McIndoe A.
Garlick D.J.	Hart A.J.L.	Jago R.H.	Knight J.R.	McIntosh I.H.
Garry R.	Hart R.O.	Jain B.K.	Knight M.J.	McKelvey S.T.D.
Gartell P.C.	Hartley M.	Jakubowski J.	Knox A.J.S.	McKelvie P.
Gatehouse D.	Harvey M.H.	James M.J.	Knox R.	McKenna D.M.
	•			
George P.P.	Harvey-Hills N.	James S.E.	Kobbe A.E.R.	MacKenzie I.
Ghanekar D.G.	Hasan S.T.	Jardine-Brown C.P.	Kolar K.M.	McKibbin A.
Gibbens G.L.D.	Hastie K.J.	Jasim S.	Kotta S.	McLatchie G.
Gibbons C.P.	Hawkyard S.J.	Jeffery P.J.	Kourah M.A.	MacLennan I.
Gillatt D.	Hay D.J.	Jenkins B.	Kulkarni R.P.	McLoughlin J.
Gilliland E.L.	Haynes S.	Jenkins D.H.R.	Laing R.	McMahon M.J.
Gillison E.W.	Hendry W.F.	Jennison K.M.	Lake S.P.	McMillan D.L.
Gilroy D.	Hendy-Ibbs P.M.	Jepson K.	Lamb R.J.	McNeal A.D.
Gingell J.C.	Hensher R.	Jeyasingham K.	Lamerton A.J.	McNicholas T.A.
Glazer G.	Hershman M.	Johal B.	Lamki H.	McRae A.R.
Glossop L.P.	Heslip M.R.	John D.G.	Lane I.F.	MacSweeney S.
Gogoi N.K.	Hibbert J.	Johnson A.G.	Langdon J.D.	McVicar I.
Goldstraw P.		Johnson A.O.B.		Mace M.C.
	Higgins A.F.		Langton J.	
Goodall R.J.R.	Higgins J.R.A.	Johnson C.D.	Larvin M.	Machin D.G.
Gooding M.R.	Hill J.T.	Johnson D.A.N.	Lavelle M.A.	Mackle E.J.
Goodman A.J.	Hindmarsh J.R.	Johnson I.R.	Lawrence W.T.	Maclean A.D.W.
Gordon A.	Hobbiss J.H.	Johnson J.R.	Laws I.M.	Macnab G.
Gordon D.J.	Hocken D.B.	Johnson M.G.	Lawson R.A.M.	Mahapatra T.K.
Gordon E.M.	Hodgkins J.	Johnson S.R.	Lea R.E.	Maheson M.V.S.
Gore R.	Hoile R.W.	Johnston F.	Leabeater B.F.	Mair W.S.J.
Gottlieb A.	Holbrook M.C.	Jones A.S.	Lee J.O.	Maiwand O.
Goulbourne I.A.	Holcombe C.	Jones C.	Lee M.D.	Makin C.A.
Gowland-Hopkins N.F.	Holden D.	Jones C.A.	Lee P.W.R.	Malak T.M.
Graham J.M.	Holdsworth P.J.	Jones D.J.	Lees T.A.	Mansfield A.O.
Gray R.F.	Holliday H.W.	Jones D.L.	Leese T.	Marcuson R.W.
Greatrex G.H.	Holme T.C.	Jones D.R.	Le Fur R.	Markham N.I.
GICALICA U.II.	110mmc 1.C.	JUNES D.K.	Lorui K.	iviai Kiiaiii IV.I.

Marsh H.T.	Mufti G.	Pearson R.C.	Rhys-Williams S.	Sells R.A.
Marshall R.W.	Mughal M.M.	Peck J.E.	Richards P.	Sellu D.
Martin I.C.	Mukoyogo J.M.	Peel K.R.	Richardson J.A.	Sengupta B.S.
Martin I.G.	Mulholland R.C.	Peet T.N.D.	Richmond W.D.	Sergeant R.J.
Martin R.H.	Mullan F.	Pentlow B.D.	Ridley D.	Shafi M.I.
Mason J.R.	Muller P.W.S.	Pereira J.H.	Rigby C.C.	Shafiq M.
Massey J.A.	Munson K.W.	Perkins C.	Rigg K.M.	Shaikh N.A.
Masson G.M.	Murphy D.J.	Perks A.G.B.	Ritchie A.	Shan S.
Mathew B.G.	Murphy F.	Perry S.	Ritchie A.W.S.	Shanahan D.J.
Maurice-Williams R.S.	Murrant N.J.	Peters J.	Roberts A.	Sharma N.
			Roberts J.G.	Sharma P.
Mavor A.I.D.	Murray A.	Phelan P.S.		
Maxwell R.J.	Murray A.	Phillips J.G.	Robertshaw J.K.	Sharma S.D.
May A.R.L.	Murty G.	Phillips J.J.	Robertson A.A.	Sharma S.K.
May R.E.	Musgrove B.	Philp N.H.	Robertson C.	Sharp J.
Maybury N.K.	Narula A.	Philp T.	Robinson A.C.	Shaw G.
Mearns A.J.	Nash A.G.	Pickles J.M.	Robinson M.H.	Shepherd R.J.
Mee A.D.	Nashef S.A.M.	Pietroni M.C.	Robinson M.R.G.	Sherlock D.J.
Meehan S.E.	Nasmyth D.G.	Pipe N.G.J.	Roe A.M.	Sherman I.W.
Meer J.A.	Naylor A.R.	Plail R.	Rogawski K.M.	Shieff C.L.
Mehdian S.M.	Needham P.G.	Plumb A.P.	Rogers J.	Shields M.D.
Meleagros L.M.	Neil-Dwyer G.	Pobereskin L.H.	Rooney P.S.	Shinkfield M.
	•	Pocock R.D.	Rose M.B.	Shortridge R.T.J.
Mellon J.K.	Nelson I.W.			•
Menzies D.	Nelson R.J.	Pollard R.	Rosen E.	Sibson D.E.
Mercurius-Taylor L.A.	Nicolaides A.	Pope A.	Rosenberg D.A.	Sill P.R.
Meyrick Thomas J.	Noble A.D.	Poskitt K.R.	Rosin M.D.	Silverman S.H.
Michel M.	Noble J.G.	Poston G.J.	Rosin R.D.	Silvester K.C.
Milewski P.J.	North A.D.	Power R.	Ross A.C.	Simms J.M.
Miller J.F.	Norton R.	Prasad S.V.L.	Ross A.H.M.	Singh A.
Miller S.E.P.	Nuseibeh I.	Premachandra D.J.	Ross S.A.	Singh A.B.
Miller W.A.	Nwabineli N.J.	Price J.H.	Rothera M.P.	Singh G.
Millner J.	O'Bonyo H.B.	Primrose W.J.	Rothwell N.	Sinha K.N.
Mills C.L.	O'Donoghue E.P.N.	Prince H.	Rowlands D.J.	Smallpeice C.J.
Mills M.S.	O'Donoghue G.M.	Pritchett C.J.	Rowse A.D.	Smith E.M.
	_	Pushpanathan R.J.	Rowson N.J.	Smith M.R.
Milner J.C.	O'Flynn K.	-		
Milton C.M.	O'Hare M.F.	Putnam G.D.	Royston C.M.S.	Smith R.B.
Minchin A.J.	O'Reilly B.J.	Pye J.K.	Rundle J.S.H.	Smyth A.G.
Mitchell D.C.	O'Reilly P.H.	Pyper P.C.	Sabanathan S.	Snow D.G.
Modgill V.K.	O'Riordan B.	Queen K.B.	Sabharwal S.	Sole G.M.
Moghissi K.	O'Sullivan D.G.	Rahamim J.	Sabin H.I.	Soorae A.S.
Mohan J.	Oram D.H.	Rainsbury R.M.	Sadler A.P.	Sosnowski A.W.
Moir A.A.	Orr M.M.	Rajasekaran J.	Sagar P.M.	South L.M.
Moisey C.U.	Osborne D.R.	Rajesh P.B.	Sagar S.	Sowinski A.
Monaghan J.M.	Osman I.S.	Ralphs D.N.L.	Sainsbury J.R.C.	Speakman C.T.
Monson J.R.T.	Ostick D.G.	Ramsden C.	Salter M.C.P.	Spearing G.J.
Montgomery A.C.V.	Pace-Balzan A.	Ramus N.I.	Sampath S.A.C.	Spencer S.R.
Montgomery B.S.I.	Padwick M.	Randall C.J.	Samuel P.R.	Spencer T.S.
Mooney P.	Page R.D.	Rangecroft R.G.	Sandeman D.R.	Spychal R.T.
Moore A.J.	Pain J.A.	Ranjit R.	Sanderson C.J.	Spyt T.
		-	Sandhu D.P.S.	Srinivasan V.
Moore K.T.H.	Palmer J.H.	Rao P.N.		Staff W.G.
Moore P.J.	Pantelides M.	Rashid A.	Sansom J.R.	
Moorehead R.J.	Parker C.J.	Ratliff D.A.	Sant Cassia L.J.	Stafford F.W.
Moores W.K.	Parker D.	Ravi M.	Sarkar P.K.	Stafford N.D.
Moran B.	Parr D.C.	Ravichandran G.	Sarmah B.D.	Stansby G.
Morcos M.	Parr N.J.	Reddy P.	Saunders N.R.	Stassen L.F.A.
Morgan M.W.E.	Parrott N.R.	Reed M.W.	Sauven P.	Stebbing M.A.
Morgan S.J.	Parry J.R.W.	Rees A.	Savage P.E.	Steel M.R.
Morgan W.E.	Parsons K.F.	Rees A.E.	Sayer R.E.	Steele R.J.C.
Morgan W.P.	Parvin S.	Rees M.	Sayer T.	Steele S.C.
Morrell M.T.	Parys B.T.	Regan P.	Scott A.D.N.	Stewart D.J.
Morris G.E.	Pastorino U.	Reid B.A.	Scott G.I.	Stewart J.
Morris I.R.	Paterson M.	Reid D.	Scott I.H.K.	Stewart P.A.H.
Morsman J.M.	Paterson M.E.L.	Reilly D.	Scott I.V.	Stewart W.E.
Mortensen N.	Patterson M.	Reilly P.G.	Scott N.	Stoddard C.J.
		Resouly A.	Scott R.A.P.	Stoker T.A.M.
Morton K.E.	Payne E.S.	•	Scott S.D.	Stoodley B.J.
Morton M.E.	Payne S.R.	Reynolds R.		-
Mosley J.G.	Pearson H.J.	Rhind R.	Scriven M.	Stower M.J.
Mosquera D.A.	Pearson M.	Rhodes M.	Sell P.J.	Strachan J.R.

Strachan R.D. Wake M. Stringer R. Walker M.G. Stuart A.E. Walker R.T. Walker S.J. Studley J. Sturzaker H.G. Wallace D.M.A. Sudlow R.A. Wallace R.G.H. Sullivan S.C. Wallwork J. Suresh G. Walsh S. Surtees P. Walsh T.H. Ward D. Sutton R. Swingler G. Ward D.C. Symes J.M. Ward J.K. Ward S.E. Taggart D. Tait W.F. Watkin D.F.L. Talbot R.W. Watkins R.M. Tate J.J.T. Watson C. Watson D.C.T. Taylor A.B.W. Taylor M.C. Watson G.M. Taylor P.H. Watson J.A.S. Taylor S.A. Watson M.G. Taylor W.G. Way B.G. Teddy P.J. Waymont B. Terry T.R. Webb J.K. Thomas D.M. Webb P.J. Thomas D.R. Webber P.A. Thomas M.H. Wedgwood K.R. Thomas W.E.G. Weeden D. Thompson H.H. Wells A.D. Thompson J.F. Wells F.C. Thompson M.H. Wellwood J.M. Thomson A.A.G. Wengraf C.L. Thomson H.J. Westmore G.A. Thorpe A.C. Weston P.M.T. Thorpe J.A.C. Weston Underwood J. Thurston A.V. Westwood C.A. Timoney A. White B.D. Tindall S.F. White C.M. Tolley N. Whitehead E. Toor K.S. Whitehead S.M. Tophill P.R. Whiteley G.S.W. Whiteway J.E. Towler J.M. Whitfield B.C.S. Townsend E.R. Toynton C.J. Whyman M.R. Treacy P.J. Wickham M.H. Treble N.J. Wilkins D.C. Trotter G. Wilkinson M.J.S. Tulloch D.N. Willatt D.J. Turner A.G. Williams G T Turner D.T.L. Williams G. Turner S.M. Williams G.T. Tweedie J.H. Williams H. Tyrrell M.R. Williams H.T. Upadhyay S. Williams J.H. Williams J.L.

Williamson E.P.M. Vandal M.T. Vanwaeyenbergh J. Wills M.I. Vasey D.P. Wilson L. Vaughan E.D. Wilson N.M. Vaughan R. Wilson R.Y.

Upsdell S.M. Urwin G.H.

Vadanan B.S.

Van Vuuren L.

Veitch P.S. Wilson-McDonald J.

Windle R. Vesey S.G. Vetrivel S.M. Winslet M.C. Vipond M. Witherow R.O.N. Virdi J. Womack N.

Woods W. Woodward A. Woodwards R.T. Woolfenden K.A. Worth P.H.L. Worth R.W. Wyatt M.G. Wynne K.S. Yacoub M.H. Yardley M.P.J. Yeo R. Young H.L.

Williams M.R.

Williams T.G.

Williamson B.

Appendix L - Local Reporters

This list shows the local reporters as of August 1998.

We recognise that there are many clinical audit and information departments involved in providing data, although we have in many cases named only the consultant clinician nominated as local reporter.

Anglia & Oxford

Nuffield Orthopaedic Centre

Addenbrooke's Dr D. Wight

Mrs S. Blackley **Bedford Hospital**

Heatherwood & Wexham Park Hospitals Dr M.H. Ali

Hinchingbrooke Health Care Dr M.D. Harris

Mr I.E. Cowles Ipswich Hospital James Paget Hospital Mrs C.L. Eagle

Dr B.E. Gostelow Kettering General Hospital

Miss J.M. Rippon Kings Lynn & Wisbech Hospitals

Dr S.S. Jalloh Milton Keynes General

Norfolk & Norwich Health Care Dr A.J.G. Gray

Northampton General Hospital Dr A.J. Molyneux

Dr P. Millard (John Radcliffe Hospital) Oxford Radcliffe Hospital

Dr M.J. Mahy (Horton General Hospital)

Dr P. Millard

Dr R. Menai-Williams

Dr N. Cary Papworth Hospital

Dr P.M. Dennis Peterborough Hospitals

Dr P. Millard Radcliffe Infirmary

Royal Berkshire & Battle Hospital South Buckinghamshire Dr M.J. Turner

Dr A.F. Padel Stoke Mandeville Hospital

Dr D.A.S. Lawrence The Luton & Dunstable Hospital

Mrs V. Hamilton West Suffolk Hospitals

North Thames

Dr S.G. Subbuswamy Basildon & Thurrock General Hospitals

Dr C.A. Amerasinghe Central Middlesex Hospital

Dr W.H.S. Mohamid Chase Farm Hospitals Ms G. Nuttal Chelsea & Westminster Healthcare

Dr C. Schmulian Ealing Hospital

East Hertfordshire Dr A. Fattah

Essex Rivers Healthcare Mrs A. Bridge

Dr K.M. Thomas Forest Healthcare

Hammersmith Hospitals

Professor G.W.H. Stamp

(No local reporter for Charing Cross Hospital)

Havering Hospitals

Ms C. Colley

Hillingdon Hospital

Dr F.G. Barker

Mid-Essex Hospital Services

Mr A.H.M. Ross (Broomfield Hospital)
Dr S.G. Subbuswamy (St Andrew's Hospital)

Moorfields Eye Hospital

Professor P. Luthert

Mount Vernon & Watford Hospitals

Mrs M. Hill (Mount Vernon Hospital)

Dr W.K. Blenkinsopp (Watford General Hospital)

Newham Healthcare

Dr G. Russell

North Hertfordshire

Dr D.J. Madders

North Middlesex Hospital

Dr K.J. Jarvis

Northwick Park & St Mark's

Dr S. Boyle

Redbridge Healthcare

Dr P. Tanner

Royal Brompton & Harefield Hospitals

Ms J. Byrne (Harefield Hospital)

Professor R. Denison (Royal Brompton Hospital)

Royal Free Hampstead

Dr J.E. McLaughlin

Royal Marsden Hospital

Mr R.J. Shearer

Royal National Orthopaedic Hospital

None

Southend Health Care

Ms L. Bell

St Albans & Hemel Hempstead

Dr A.P. O'Reilly

St Mary's Hospital

Ms R.A. Hittinger

Dr R.G.M. Letcher

The Great Ormond Street Hospital for Children

Dr A. Mackersie

The Homerton Hospital

Ms M. Joseet

The Princess Alexandra Hospital

_ _ _ _

The Royal Hospitals

Dr P.J. Flynn

University College London Hospitals

Ms A.E. Glover (Middlesex Hospital

Mrs J.A. Sullivan (The National Hospital for Neurology and

Neurosurgery)

Wellhouse

Dr J. El-Jabbour

West Middlesex University Hospital

Dr R.G. Hughes

Whittington Hospital

Dr S. Ramachandra

North West

Aintree Hospitals

Dr W. Taylor

Blackburn, Hyndburn & Ribble Valley Healthcare

Mr R.W. Nicholson

Blackpool Victoria Hospital

Dr K.S. Vasudev

Bolton Hospitals

Dr S. Wells

Burnley Health Care

Mr D.G.D. Sandilands

Bury Health Care

Dr E. Herd

Central Manchester Healthcare

Dr E.W. Benbow

Chorley & South Ribble

Dr C. Loyden

North West continued

Christie Hospital None

Countess of Chester Hospital Dr P.R.M. Steele
East Cheshire Dr A.R. Williams
Halton General Hospital Dr K. Strahan

Liverpool Women's Hospital Ms C. Fox

Manchester Children's Hospitals Dr M. Newbould
Mid Cheshire Hospitals Miss H. Moulton

Morecambe Bay Hospitals Dr R.W. Blewitt (Royal Lancaster Infirmary)

Dr V.M. Joglekar (Furness General Hospital)

North Manchester Health Care None

Oldham Dr I. Seddon

Preston Acute Hospitals Dr C.M. Nicholson

Rochdale Healthcare Mr S. Murray
Royal Liverpool & Broadgreen University Hospital Miss K. Scott

Royal Liverpool Children's Hospital Mrs P.A. McCormack Salford Royal Hospitals Mr M. McKenna

South Manchester University Hospitals Dr J. Coyne (Withington Hospital)

Dr P.S. Hasleton (Wythenshawe Hospital)

Southport & Formby Dr S.A.C. Dundas
St Helens & Knowsley Hospitals Mr K. Benyon
Stockport Acute Services Dr P. Meadows

Tameside Acute Care Dr A.J. Yates
The Cardiothoracic Centre Liverpool Mr M. Jackson

Trafford Healthcare Dr B.N.A. Hamid
Walton Centre for Neurology & Neurosurgery Dr J. Broome

Walton Centre for Neurology & Neurosurgery Dr J. Broome
Warrington Hospital Dr M.S. Al-Jafari

West Lancashire Mr A.D. Johnson

Wigan & Leigh Health Services Ms S. Tarbuck
Wirral Hospital Dr M.B. Gillett

Wrightington Hospital Mr A.D. Johnson

Northern & Yorkshire

Airedale Dr J.J. O'Dowd

Bradford Hospitals Dr B. Naylor

Calderdale Healthcare Mr R.J.R. Goodall

Carlisle Hospitals Dr E.D. Long

City Hospitals Sunderland Miss K. Ramsey

Dewsbury Health Care Dr P. Gudgeon

East Yorkshire Community Healthcare Mr G. Britchford

East Yorkshire Hospitals Mr G. Britchford

Gateshead Hospitals Dr I.M.J. Mathias

Harrogate Health Care

Hartlepool & East Durham

Huddersfield

Newcastle upon Tyne Hospitals

North Durham Healthcare

North Tees Health

Northallerton Health Services

Northumbria Healthcare

Pinderfields & Pontefract Hospitals

Royal Hull Hospitals

Scarborough & N E Yorkshire Healthcare

South Durham Healthcare

South Tees Acute Hospitals

South Tyneside Healthcare

United Leeds Teaching Hospitals

West Cumbria Health Care

York Health Services

Miss A.H. Lawson

Mrs A. Lister

Dr H.H. Ali

Dr M.K. Bennett

Miss D. Robson

Dr D. Wood

Dr J. Hoffman

Dr D.C. Henderson

Dr J.A. Henry (Cheviot & Wansbeck)

Dr F. Johri (North Tyneside)

Dr S. Gill (Pinderfields General Hospital)

Dr I.W.C. MacDonald (Pontefract General Infirmary)

Dr M.R.F. Reynolds

Dr A.M. Jackson

Dr D.C.A. Senadhira (Bishop Auckland Hospital)

Ms C. Evans (Darlington Memorial)

Mrs L. Black

Dr K.P. Pollard

Dr C. Abbott (The General Infirmary at Leeds)

Mr S. Knight (St James' University Hospital)

Dr R.G. Ghazala

Dr C. Bates

South Thames

Ashford & St Peter's Hospitals

Brighton Healthcare

Bromley Hospitals

Dartford & Gravesham

Eastbourne Hospitals

Epsom Health Care

Frimley Park Hospital

Greenwich Healthcare

Guy's & St Thomas'

Hastings & Rother

Kent & Canterbury Hospitals

Kent & Sussex Weald

King's Healthcare

Kingston Hospital

Mayday Health Care

Medway

Dr J.C. Dawson (Ashford Hospital)

Mr R.H. Moore (St Peter's Hospital)

Mr M. Renshaw

Dr M.H. Elmahallawy

Dr A.T.M. Rashid

Mr M.D. Bastable

Dr T.J. Matthews

Dr G.F. Goddard

Mr O. Smith

Dr. B. Hartley (Guy's Hospital)

Professor S.B. Lucas (St Thomas' Hospital)

Mr S. Ball

Mr M. Guarino

Dr G.A. Russell

Mr M. Fleming

Mr P. Willson

Mr C. Fernandez

Mrs J.L. Smith

South Thames continued

Mid Kent Healthcare Mr J. Vickers

Mid-Sussex Dr P.A. Berresford (The Princess Royal Hospital)

Dr E.H. Rang

Mr E. Robbin

Mr P.H. Walter (Hurstwood Park Neurology Centre)

Queen Mary's Sidcup Dr E.J.A. Aps

Richmond, Twickenham & Roehampton Mr M. McSweeney

Royal Surrey County & St Luke's Hospitals Mrs M. Davies

South Kent Hospitals Dr C.W. Lawson Dr S. Dilly St George's Healthcare

Surrey & Sussex Healthcare Ms C. Parkinson (East Surrey Hospital)

Dr C. Moon (Crawley Hospital and Horsham Hospital)

Thanet Healthcare Mrs B.M. Smith

The Lewisham Hospital Dr C. Keen

The Oueen Victoria Hospital Mrs D.M. Helme The Royal West Sussex Mr J.N.L. Simson

The St Helier

Worthing & Southlands Hospitals Mrs J. Tofield

South & West

Dorset Community Dr A. Anscombe

East Gloucestershire Dr W.J. Brampton

East Somerset Dr G. Purcell

Frenchay Healthcare Dr N.B.N. Ibrahim

Gloucestershire Royal Dr B.W. Codling

Isle of Wight Healthcare Mr P. Donaldson North Hampshire Hospitals Dr J.M. Finch

Northern Devon Healthcare Dr J. Davies

Plymouth Hospitals Dr C.B.A. Lyons

Poole Hospital Dr D.S. Nicholas

Portsmouth Hospitals Dr N.J.E. Marley

Royal Bournemouth & Christchurch Hospitals

Royal Cornwall Hospitals Dr R. Pitcher

Royal Devon & Exeter Healthcare Dr R.H.W. Simpson

Royal United Hospital Bath None

Salisbury Health Care Dr C.E. Fuller

South Devon Healthcare Dr D.W. Day

Southampton University Hospitals Dr A. Bateman

Southmead Health Services Ms G. Davies

Swindon & Marlborough Hospital Mr M.H. Galea

Taunton & Somerset Mr I. Eyre-Brook

United Bristol Healthcare Dr E.A. Sheffield

West Dorset General Hospitals Dr A. Anscombe Weston Area Health

Winchester & Eastleigh Healthcare

Dr M.F. Lott

Dr R.K. Al-Talib

Trent

Barnsley District General Hospital

Bassetlaw Hospital & Community Services

Central Nottinghamshire Health Care

Central Sheffield University Hospitals

Chesterfield & North Derbyshire Royal Hospital

Southern Derbyshire Hospitals

Doncaster Royal Infirmary & Montagu Hospital

Glenfield Hospital

Grantham & District Hospital

Leicester General Hospital

Leicester Royal Infirmary

Lincoln & Louth

North East Lincolnshire

Northern General Hospital

Nottingham City Hospital

Pilgrim Health

Queen's Medical Centre University Hospital

Rotherham General Hospitals

Scunthorpe & Goole Hospitals

Sheffield Children's Hospital

The King's Mill Centre for Health Care Services

West Lindsey

Dr M.A. Longan

Dr P.A. Parsons

Dr. I. Ross

Dr C Angel

Dr P.B. Gray

Mrs L. Smith (Deby City General Hospital)

Mr J.R. Nash (Derbyshire Royal Infirmary)

Dr J.A.H. Finbow

Mrs S. Clarke

Dr D. Clark

Dr E.H. Mackay

Mr R. Mowbrav

Dr J.A. Harvey (Lincoln County Hospital)

Mr E.O. Amaku (County Hospital Louth)

Dr W.M. Peters

Dr S.K. Suvarna

Dr. J.R. Jones

Ms S. Cosgriff

Dr. J.R. Jones

Mr R.B. Jones

Dr C.M. Hunt Dr I. Barker

Ms J. Jenkins Dr J.A. Harvey

West Midlands

Alexandra Healthcare

Dr L. Brown

Dr F. Raafat

Birmingham Heartlands Hospital

Birmingham Children's Hospital

Dr M. Taylor

Birmingham Women's Healthcare

Dr T. Rollason Dr N. Kasthuri

Burton Hospitals

Dr J. Mercer

George Eliot Hospital Good Hope Hospital

Dr J. Hull

Hereford Hospitals

Dr F. McGinty

Kidderminster Health Care

Dr G.H. Eeles

Mid Staffordshire General Hospitals

Dr V. Suarez

West Midlands continued

North Staffordshire Hospital Centre

Mrs C. McPherson Robert Jones & Agnes Hunt

Royal Shrewsbury Hospitals Dr R.A. Fraser

Sandwell Healthcare Dr J. Simon

South Warwickshire General Hospitals Mr M. Gilbert

None The City Hospital

Dr S. Ghosh The Dudley Group of Hospitals Dr R.A. Fraser The Princess Royal Hospital Mr A. Thomas The Royal Orthopaedic Hospital

The Royal Wolverhampton Hospitals Professor E.L. Jones (Queen Elizabeth Hospital) University Hospital Birmingham

Dr T.A. French

Dr J. Tomlinson

(No local reporter for Selly Oak Hospital)

Dr Y.L. Hock Walsall Hospitals Walsgrave Hospitals Dr J. Macartney

Worcester Royal Infirmary Mr A. Singfield

Northern Ireland

Altnagelvin Hospitals Dr J.N. Hamilton

Armagh & Dungannon Mr B. Cranley

Belfast City Hospital Mr R. J. Hannon

Dr C. Watters Causeway Craigavon Area Hospital Group Mr B. Cranley

Down Lisburn Dr B. Huss (Lagan Valley Hospital)

Mrs M. Gilgunn (Downe Hospital)

Green Park Healthcare Dr J. D. Connolly

Dr P. Gormley Mater Hospital

Mr B. Cranley Newry & Mourne

Dr W. Holmes (Erne Hospital) Sperrin Lakeland

Dr F. Robinson (Tyrone County Hospital)

The Royal Group of Hospitals & Dental Hospitals

Ulster Community & Hospital Dr T. Boyd

Mr I. Garstin (Antrim Area Hospital) United Hospitals

Mr P.C. Pyper (Mid -Ulster Hospital)

Mr D. Gilroy (Whiteabbey Hospital)

Ms M. Toner

Wales

Bridgend & District Dr A.M. Rees

Dr R.B. Denholm Carmarthen & District

Mrs C. Smith Ceredigion & Mid Wales

Dr D. Stock East Glamorgan

Glan Clwyd District General Hospital Dr B. Rogers

Glan Hafren None Glan-y-Mor

Gwynedd Hospitals

Llandough Hospital & Community

Llanelli Dinefwr

Morriston Hospital/Ysbyty Treforys

Nevill Hall & District

North Glamorgan

Pembrokeshire & Derwen

Rhondda Health Care

Swansea

University Hospital of Wales

Wrexham Maelor Hospital

Defence Medical Services

The Duchess of Kent Military Hospital

The Royal Naval Hospital, Haslar

Guernsey / Isle of Man / Jersey

Guernsey

Isle of Man

Jersey

BUPA

BUPA Wellesley Hospital

BUPA Hospital Norwich

BUPA Hospital Leeds

BUPA Hospital Little Aston

BUPA North Cheshire Hospital

BUPA Parkway Hospital

BUPA Hospital Portsmouth

BUPA Roding Hospital

BUPA Hospital Hull & East Riding

BUPA South Bank Hospital

BUPA Hospital Leicester BUPA Hospital Elland

BUPA St. Saviour's Hospital

BUPA Belvedere Hospital

BUPA Hospital Manchester BUPA Hartswood Hospital

The Glen Hospital

Dr A. Dawson

Dr M. Hughes

Dr J. Gough

Dr L.A. Murray

Dr A. Dawson

Dr R.J. Kellett

None

Dr G.R. Melville Jones

Dr D. Stock

Dr S. Williams

Dr A.G. Douglas-Jones (University Hospital of Wales)

Mrs P. Perrott (Cardiff Royal Infirmary)

Dr R.B. Williams

The Commanding Officer

Dr N. Carr

Dr B.P. Gunton-Bunn

Ms E. Clark

Dr H. Goulding

Mrs L. Horner

Ms J. Middows

Mr D. Farrell

Mr K. Smith

Miss A.L. Alexander

Mrs M.T. Hall

Ms J. Ward

Ms D. Brett

Ms A. Meyer

Mrs F. Cox

Mrs C.A. Jones

Ms M.E. Schofield

Mrs E. Biddle

Mr S.J. Greatorex

Ms A. McArdle

Ms N. Howes

Miss M. O'Toole

BUPA continued

BUPA Alexandra Hospital Mr N.R. Permain Ms L. Adair **BUPA Hospital Bushey** BUPA Cambridge Lea Hospital Ms M. Vognsen Mrs D. Wright **BUPA Gatwick Park Hospital** BUPA Murrayfield Hospital Mr S. Milner BUPA Hospital Harpenden Ms B. Hayter Dr A. Gibbs **BUPA Hospital Cardiff BUPA** Fylde Coast Hospital Ms D. Revell **BUPA** Dunedin Hospital Mrs C. Bude Ms M. Wood **BUPA Hospital Clare Park BUPA** Chalybeate Hospital Ms M. Falconer

General Healthcare Group

Dr A.G. Jacobs The Priory Hospital Mrs J. Bevington AMI Princess Margaret Hospital Ms S. Prince Harbour Hospital Mrs P. Hill The Runnymede Hospital The Somerfield Hospital Ms N. Poulson Ms C. Poll Chelsfield Park Hospital Mr R.J. Hoff The Wellington Hospital Mrs C. Pagram Fawkham Manor Hospital Mrs U. Palmer The Sloane Hospital Dr I. Chanarin Clementine Churchill Hospital Mrs J. Cooper The Thornbury Hospital Ms D. Cunliffe The Princess Grace Hospital Ms S. Hill The Chiltern Hospital Ms S. Thomas The Harley Street Clinic Mr C.J. Durkin The Highfield Hospital Ms S. Quickmire Park Hospital Ms V. Power The Blackheath Hospital Miss A.D. Sayburn The Portland Hospital for Women and Mrs P. Enstone Alexandra Hospital

Nuffield

The Chaucer Hospital

The North London Nuffield Hospital Miss J. Ward
The Huddersfield Nuffield Hospital Miss S. Panther
The Grosvenor Nuffield Hospital Mrs J.L. Whitmore
The Lancaster & Lakeland Nuffield Hospital Miss A. Durbin
The Nuffield Hospital Leicester Mrs S. Harriman

Mrs G. Mann

Mid Yorkshire Nuffield Hospital Mrs M. Dunderdale
The Newcastle Nuffield Hospital Miss K.C. Macfarlane

The Hull Nuffield Hospital Ms V. Ward
The North Staffordshire Nuffield Hospital Mr D. Allison
The Nuffield Hospital Plymouth Mrs T. Starling
The Purey Cust Nuffield Hospital Mr J. Gdaniec
The Shropshire Nuffield Hospital Mrs S. Crossland
The Somerset Nuffield Hospital Mrs J.A. Dyer
The Sussex Nuffield Hospital Mrs F. Booty

The Woking Nuffield Hospital Miss B.E. Harrison
The Thames Valley Nuffield Hospital Ms D. Williams
The Acland Hospital Miss C. Gilbert
The Exeter Nuffield Hospital Mrs A. Turnbull
The Tunbridge Wells Nuffield Hospital Mr R. Muddiman
The Wolverhampton Nuffield Hospital Mrs I. Jones

The Wolverhampton Nuffield Hospital Mrs I. Jones
The Wessex Nuffield Hospital Mrs V. Heckford
The Wye Valley Nuffield Hospital Mrs W.P. Mawdesley

HRH Princess Christian's Hospital Ms S. Fisher Mrs P. Shields The Birmingham Nuffield Hospital Mrs S. Jackson The Bournemouth Nuffield Hospital The Chesterfield Nuffield Hospital Miss P.J. Bunker Mrs C. Williams The East Midlands Nuffield Hospital Mrs J.T. Penn The Cotswold Nuffield Hospital Miss S.J. Gardner The Duchy Nuffield Hospital The Cleveland Nuffield Hospital Mrs S. Jelley The Essex Nuffield Hospital Mrs B.M. Parker Mrs J. Worth The Warwickshire Nuffield Hospital Ms Y. Terry London Bridge Hospital

St Martin's Hospitals

The Lister Hospital Mrs J. Johnson

Devonshire Hospital Miss C. Lewington

Other independent hospitals

Benenden Hospital Mr D. Hibler
The London Clinic Mrs K. Perkins