

Juvenile Idiopathic Arthritis

This hospital or doctor's surgery is taking part in a piece of work to help to see where care can be made better for future children and young people who have **Juvenile Idiopathic Arthritis (JIA)**.

The organisation collecting the data is called **NCEPOD**.

NCEPOD is a national organisation that uses data to help **improve the quality of care for future patients** in the UK. We do this by collecting data on patients from hospitals and other places where they may be cared for. For this work we are collecting data on **children and young adults aged 0-24 years old**.

For over 35 years **NCEPOD** has run studies across the UK and these have led to lots of changes in how healthcare is provided.

How you can get involved



If you are a **young person** or **parent carer** of a young person, who has JIA, and would like to **take part in a group discussion**, or fill in an **online survey** to **share your experiences** to help future patients, please contact us by post, phone or email:

arthritis@ncepod.org.uk

*Please ask permission from your parent/carer if you are under 16**

If you are aged **0-24 years old**, your healthcare data might be included in our work. We will keep your data very safe until the end of the work, when it will all be safely deleted.

If you (if aged 16 or over) or your parent/carer (if you are aged under 16)* do not want your data used please tell:

Marisa Mason, Chief Executive, by post, phone or email
mmason@ncepod.org.uk

How your data might be used, and how you can say NO!

**A parent or carer can also contact us on behalf of a young person aged 16 or over if they are their court appointed Care and Welfare Deputy.*

NCEPOD
www.ncepod.org.uk

  @ncepod

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