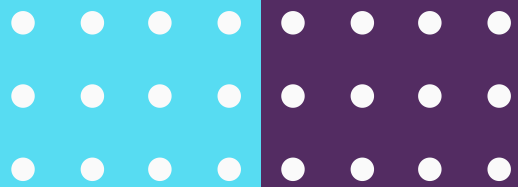


HOW DOES THIS AFFECT ME?

We will need to collect some information about children with a new diagnosis of CRMO/CNO. This includes their medical journey so far, gender, ethnic group and date of birth. The doctors will use information from medical records to complete 2 surveys. Names and addressess will not be included.

WHAT IF I DON'T WANT TO BE INVOLVED?

It is 100% okay if you don't want to be involved! Your doctor will be happy to talk to you about this and answer any questions.



REACH US

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Chronic Recurrent Multifocal Osteomyelitis/ Chronic Nonbacterial Osteomyelitis

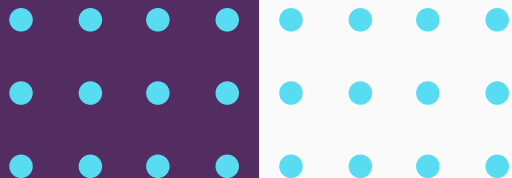
A PUBLIC INFORMATION LEAFLET
FOR KIDS & TEENS



WHAT IS CRMO/CNO?

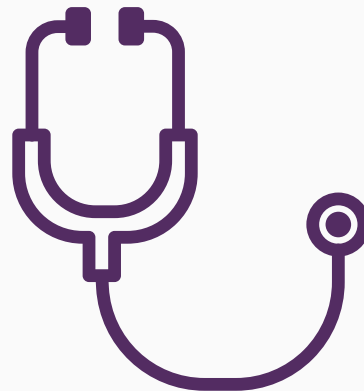
Chronic Recurrent Multifocal Osteomyelitis/Chronic Nonbacterial Osteomyelitis (CRMO/CNO) is a rare disease affecting the bones. The body's immune system attacks the bones by mistake, causing pain and swelling.

Usually, painkillers can really help children with CRMO/CNO. Unfortunately, CRMO/CNO can sometimes cause more serious problems like breaks in bones.



WHO IS BPSU?

The British Paediatric Surveillance Unit (BPSU) studies rare conditions in children. It allows doctors and researchers to find out how many children in the UK & Ireland are affected by these conditions every year. This information helps us become better at treating children with these conditions in the future.



HOW DOES THIS STUDY WORK?

HOW?

We want to find out how common CRMO/CNO is, the people involved in the care of children with CRMO/CNO, and the different treatments that are currently given to these children.

WHO?

Specialist doctors who treat children are sent online report cards every month. They use these report cards to let us know if they have seen any children with CRMO/CNO. We then send them a short survey, for them to tell us more about the patient's medical journey.

WHEN?

We want to identify new cases of CRMO/CNO over 13 months. The follow up study will last 12 months.