CMV RESEARCH STUDY

We want to hear your opinion: what are the cytomegalovirus (CMV) research priorities?

This study aims to find out the most important research priorities for preventing and reducing the impacts of CMV infection for women and babies in Australia and New Zealand.

Knowing what is important to those most affected will help direct research funding and build the most relevant research agenda.

Who can take part?

1. People with lived experience of CMV
2. Clinicians, researchers and health policy experts with an interest and/or knowledge in CMV

To participate scan the code

or click HERE

What is involved?

3 short surveys over a 4-month period.

People with lived experience will also be invited to attend a one-hour online meeting.

In Australia, more than 400 babies every year are born with disabilities such as hearing loss, cerebral palsy and intellectual impairment as a result of CMV.

To find out more about this study, please contact: **Ms Kath Swinburn**

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This study has ethical approval from the University of Sydney HREC: 2024HE001863

Cerebral Palsy ALLIANCE

THE UNIVERSITY OF SYDNEY