

Test screening for leading infectious cause of hearing loss in babies feasible and wellreceived

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A saliva test taken by parents to screen for the leading infectious cause of hearing loss in babies was feasible and well-received, according to a new study. Credit: Filip Mroz

A saliva test taken by parents to screen for the leading infectious cause of hearing loss in babies was feasible and well-received, according to a new study.

The research led by the Murdoch Children's Research Institute (MCRI) and published in the *Journal of Paediatrics and Child Health*, could pave the way for testing of congenital cytomegalovirus (cCMV) to be added to infant <u>hearing</u> screening programs across Australia.

cCMV can be harmless, but in some babies it leads to hearing loss and neurodevelopmental disorders such as cerebral palsy or vision loss. About 1 in every 200 babies is born with cCMV.

MCRI researcher Emma Webb said this was the first study to show that saliva swabs taken by parents to check for cCMV in their babies was achievable and acceptable to Australian families, even when the swabs were taken at home after discharge from hospital.

The study involved 96 families across four maternity hospitals in Melbourne whose babies did not pass their second Victorian Infant Hearing Screening Program (VIHSP) tests. Accurate diagnosis of cCMV requires a <u>saliva test</u> from an infant in the first 21 days after birth, with antiviral treatment recommended, if clinically indicated, within the first month of life.



During the study, 26 per cent of parents did the saliva test at home, 60 per cent at the hospital and 10 per cent as outpatients. The research found a high uptake by parents with 76 per cent agreeing to do the screen and 100 per cent of infant saliva swabs taken within the required timeframe. One baby was found to have cCMV and was immediately treated.

The findings also showed more than 90 per cent of parents thought the screening was easy to do, was a good idea, and were glad their baby had the test.

Ms Webb said CMV was present in 80 per cent of Australian adults and while it's mostly harmless, it could affect a developing fetus.

"Once a person becomes infected, the virus remains present but usually dormant for life," she said. "Infection or reactivation can occur during pregnancy, with the small risk that the unborn baby may contract the virus."

International guidelines recommend targeted cCMV screening of newborns who do not pass their hearing checks.

MCRI Associate Professor Valerie Sung said because cCMV was not routinely screened for in Australia, detecting affected infants in time to offer potential antiviral treatment was less likely.

"Given this test allows for an early and accurate cCMV diagnosis, could reduce unnecessary parental guilt, and help prevent lifelong disabilities it should be rolled out nation-wide through newborn hearing screening programs," she said.

Associate Professor Sung said timely screening for cCMV became more difficult after women and babies were discharged.



"Early discharge, as we're seeing more now of during COVID-19, can be a big problem because it means we might miss cases of cCMV," she said. "But our study has shown parents can do the swab themselves even at home after leaving the hospital."

Associate Professor Sung said training of hearing screeners, midwives and nurses to complete swabs in hospital would likely reduce false positive rates and further improve the uptake and turnaround time for results.

Victoria Cottrell, whose son Teddy was the only one to test positive for cCMV in the trial, said the test was quick and simple and taking part was the best decision she had ever made.

"I had never heard of cCMV before and thought the cause of the hearing loss most likely was genetic," she said. "During my pregnancy check-ups many infant conditions and foods to steer clear from were mentioned but cCMV was never talked about. The diagnosis came as a shock and was a lot for our family to process."

Profoundly deaf, Teddy, now aged 2 and wears cochlear implants, underwent six months of antiviral treatments and currently has physiotherapy to help with walking.

"If his cCMV diagnosis wasn't picked up so soon he would be further behind in his gross motor skills," Victoria said. "Having the awareness early we have received support quickly and our specialists have been on top everything."

Victoria said she would like to see a test for cCMV added to infant hearing screening programs across Australia.

"If the trial hadn't happened, we would never have known Teddy had



cCMV," she said. "I'm just so thankful and relieved that this trial existed to get the best outcomes for Teddy."

More information: Emma Webb et al, Feasibility and acceptability of targeted salivary cytomegalovirus screening through universal newborn hearing screening, *Journal of Paediatrics and Child Health* (2021). DOI: 10.1111/jpc.15705

Provided by Murdoch Children's Research Institute

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