PITCHES

PITCHES Trial Results

Phase III trial in IntrahepaTic CHolestasis of pregnancy (ICP) to Evaluate urSodeoxycholic acid (UDCA) in improving perinatal outcomes.



Why did we do this trial?

Intrahepatic cholestasis of pregnancy (ICP) is the commonest pregnancyspecific liver disorder in the UK. It affects around 5,500 women a year, causing premature birth and, in extreme cases, stillbirth.

The most popular current drug used to treat ICP is called ursodeoxycholic acid (commonly known as 'urso'), but it hasn't been tested in any large clinical trials to show whether it prevents premature birth and stillbirth. Our trial asked:

"If a woman has ICP, what are the effects on the baby if she is treated with ursodeoxycholic acid (or placebo)?"

What did we do?

Between December 2015 and September 2018, we recruited 605 pregnant women with ICP. Half of the women received urso and half a placebo (a 'dummy' tablet that contained no active ingredients). This is the most reliable way to test a drug.

During the trial we also:

- Collected blood test results
- Measured the women's level of itching
- Recorded information about the births
- Collected blood samples from some women to use for future research

What did we find?

We found that urso is not a drug that helps women with ICP. It did not reduce stillbirths or the chances of a baby needing to be admitted to a neonatal unit. It did not show any meaningful improvement in itch for most women, nor did it reduce the woman's bile acid levels.

What does this mean for women with ICP?

It means that most women don't need to take urso because it won't help their itching or protect their baby from stillbirth. Further research is needed to identify whether there may still be a group of women who would benefit from taking urso, and whether other drugs could reduce the itch in women with ICP or prevent their babies from being stillborn.

PITCHES Trial Coordinating Centre

NPEU Clinical Trials Unit (NPEU CTU) National Perinatal Epidemiology Unit, Nuffield Department of Population Health, University of Oxford, Old Road Campus, Headington, Oxford, OX3 7LF Tel: +44 (0)1865 289728 Fax: +44 (0)1865 289740 Email: ctu@npeu.ox.ac.uk Web: www.npeu.ox.ac.uk/ctu

This study was organised by the National Perinatal Epidemiology Unit (NPEU) CTU at the University of Oxford. The Unit is dedicated to improving the care provided to women and their families during pregnancy, childbirth and the period after birth, as well as the care provided to the newborn.





