# Assessing the effects of maternal seizures during pregnancy on the brain development of children

Submission date	Recruitment status  No longer recruiting	[X] Prospectively registered		
14/12/2018		☐ Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
09/01/2019		Results		
Last Edited	<b>Condition category</b> Nervous System Diseases	] Individual participant data		
09/07/2020		<ul><li>Record updated in last year</li></ul>		

#### Plain English summary of protocol

Background and study aims

The treatment of epilepsy during pregnancy is a balance between treating the seizures and reducing, as far as possible, the risks presented to the developing child by medication. This study will investigate whether seizures which the mother has during pregnancy alters the brain development of the developing child.

#### Who can participate?

Women with epilepsy who are either pregnant (and not yet 21 weeks) or women who are trying for a baby currently. Women must be taking the medications lamotrigine (lamictal) or levetiracetam (keppra) alone or alongside other antiepileptic medications (but not if they are taking sodium valproate (epilim) or carbamazepine (tegtretol)). The study is looking for volunteers both who are having seizures and those who are not having seizures.

#### What does the study involve?

All participants will be asked to sign a consent form indicating that they understand the study and are happy for themselves and their baby to participate. They will then be asked to provide some information about their health, education, occupation and to record on a daily basis whether or not they have had seizure. Women can do this either in a paper seizure diary or using an app. When the women's pregnancy has passed 32 weeks they will be asked about their general health and the medications they are taking again. When the baby is born researchers at the local hospital will record information about the babies birth from medical records. Shortly after the child's first birthday a home visit will be completed where a play-based assessment will be completed with the child. This involves playing a number of games with the child, which in turn provides information on their development in areas such as reasoning, language and motor abilities. Their mothers will also be asked to complete a questionnaire about their early social development and about the mothers own mood. Mother's will also be asked to complete two brief tasks (e.g. one puzzle and one language task) to inform on their reasoning abilities.

What are the possible benefits and risks of participating?

There may be no direct benefits personally to taking part. However, following the assessment of

the child feedback would be provided to parents and a letter be sent the child's GP and kept their medical record. If the person who completed the assessment had concerns about specific areas of the child's development they would discuss it with the parents and then send a letter to the child's GP. In the longer term, the findings of this study will be of benefit to both women with epilepsy and their doctors by providing information about how seizures in pregnancy may or may not impact on how the child develops in their first year of life. This may be directly relevant to participants in the study if they were planning another pregnancy in the future.

If the participant does not already record her seizures it might seem a big task to record them everyday, however only a few details are needed and on most days the participant may simply be ticking 'no' if they did not have any seizures. A further possible disadvantage of this study is that the developmental assessment may reveal that the child is experiencing some difficulties in one or more areas of their development. In this situation, with parental permission, we would share this information with the child's GP to make sure that they can arrange the necessary support for the child.

#### Where is the study run from?

The study is being led by researchers at the University of Manchester and Manchester University Hospitals NHS Foundation Trust but involves doctors and nurses from 21 hospitals around the UK.

When is the study starting and how long is it expected to run for? Set up for the study started in May 2018 and it will open to recruitment on the 14th January 2018. Women will be able to opt into the study until May 2020 and after this time no new participants will be able to opt into the study. The follow-ups after the child's first birthday will run until August 2021. The study will close on 30th October 2021 with the results being available shortly after that.

Who is funding the study? Epilepsy Research UK

Who is the main contact?
Myfanwy Rawson
Myfanwy.rawson@mft.nhs.uk

## Contact information

Type(s)
Public

#### Contact name

Ms Myfanwy Rawson

#### Contact details

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#### Type(s)

Scientific

#### Contact name

Dr Rebecca Bromley

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## Additional identifiers

Integrated Research Application System (IRAS)

232507

Protocol serial number

**IRAS 232507** 

# Study information

#### Scientific Title

Neurodevelopment After Prenatal Exposure to Seizures study: a cohort study

#### Acronym

**NAPES** 

#### Study objectives

Children exposed to frequent seizures in the womb will have poorer development at 12 months of age.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Greater Manchester Central Regional Ethics Committee, 17/05/2018, ref. 18/NW/0261.

#### Study design

Multi-centre prospective observational cohort study.

#### Primary study design

Observational

#### Study type(s)

Other

#### Health condition(s) or problem(s) studied

**Epilepsy** 

#### **Interventions**

Participants are recruited from their local participating hospital. Following recruitment information will be taken from participants regarding their background, such as their education, health and occupation. From this point they record their seizures on a daily basis using either a paper seizure diary or an app. Each month their local research team will contact them to collect that months seizure information. In the third trimester (after 32 weeks) the participant will be contacted to provide information about their current health and medications which they are taking. Following the birth of the child information is recorded about the delivery method and the health of the child at birth. After the child's first birthday, an appointment will be arranged to see the child and parent at home where a play-based assessment will be completed. Whilst this is fun for the child it provides the assessor with important information about where the child is up to in terms of their cognitive (reasoning), language and motor development. Following the assessment of development a questionnaire will be posted to participants which will ask them about their experience of taking part in the study.

#### **Intervention Type**

Other

#### Primary outcome(s)

- 1. Cognitive development is measured using Bayley Scales of Infant and Toddler Development at 12-15 months of age.
- 2. Language development is measured using Bayley Scales of Infant and Toddler Development at 12-15 months of age.
- 3. Motor development is measured using Bayley Scales of Infant and Toddler Development at 12-15 months of age.

#### Key secondary outcome(s))

- 1. Social and behavioural development is measured using Bayley Scales of Infant and Toddler Development at 12- 15 months of age.
- 2. Maternal experiences of participating in the study is measured using questionnaires when the child is 12- 15 months of age.

#### Completion date

30/04/2022

## **Eligibility**

#### Key inclusion criteria

- 1. They have a diagnosis of epilepsy and are either:
- 1.1. Trying to conceive

- 1.2. They are pregnant and before 21 weeks gestation
- 1.2.1. They have an expected date of delivery prior to the recruitment closure date (30th November 2019).
- 2. They are taking either:
- 2.1. Monotherapy lamotrigine or levetiracetam
- 2.2. Polytherapy combinations including lamotrigine and levetiracetam and other antiepileptic drugs; with the exception of valproate and carbamazepine.
- 3. They are receiving their care through one of the participating centres
- 4. They are willing to complete prospectively recorded seizure diaries

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

Adult

#### Sex

Female

#### Key exclusion criteria

- 1. Significant learning disability, requiring support to live independently
- 2. Experience or history of non-epileptic attacks (also known as NEAD or psychogenic seizures)
- 3. Taking non-antiepileptic medications which are known to be teratogenic (e.g. warfarin, mycophenolate)
- 4. English is not the first language in the home

#### Date of first enrolment

14/01/2019

#### Date of final enrolment

30/03/2021

## Locations

#### Countries of recruitment

United Kingdom

England

Northern Ireland

Study participating centre
Manchester University NHS Foundation Trust
Oxford Road

Manchester United Kingdom M13 9WL

# Study participating centre Belfast Health and Social Care Trust

Royal Hospitals Grosvenor Road Belfast United Kingdom BT12 6BA

#### Study participating centre Lancashire Teaching Hospitals NHS Foundation Trust

Royal Preston Hospital Sharoe Green Lane Preston United Kingdom PR2 9HT

#### Study participating centre Liverpool Women's Hospital NHS Foundation Trust

Crown Street Liverpool United Kingdom L8 7SS

#### Study participating centre Salford Royal NHS Foundation Trust

Stott Lane Salford United Kingdom M6 8HD

# Study participating centre City Hospitals Sunderland NHS Foundation Trust

Sunderland Royal Hospital Kayla Road Sunderland United Kingdom SR4 7TR

#### Study participating centre Newcastle Upon Tyne Hospitals NHS Foundation Trust

Royal Victoria Hospital Queen Victoria Road Newcastle Upon Tyne United Kingdom NE1 4LP

#### Study participating centre South Tees Hospitals NHS Foundation Trust

Marton Road Middlesbrough United Kingdom TS4 3BW

# Study participating centre Walton Centre for Neurology and Neurosurgery NHS Foundation Trust

Lower Lane Liverpool United Kingdom L9 7LJ

# Study participating centre University Hospitals of Morecambe Bay NHS Foundation Trust

Lancaster Royal Infirmary Ashton Road Lancaster United Kingdom LA1 4RP

#### Study participating centre Warrington and Halton Hospitals NHS Foundation Trust

Lovely Lane Warrington United Kingdom WA5 1QG

#### Study participating centre

#### Countess of Chester Hospital NHS Foundation Trust

The Countess Of Chester Health Park Liverpool Road Chester United Kingdom CH2 1UL

#### Study participating centre East Lancashire Hospitals NHS Foundation Trust

Royal Blackburn Hospital Haslingden Road Blackburn United Kingdom BB2 3HH

#### Study participating centre Mid Cheshire Hospitals NHS Foundation Trust

Leighton Hospital Crewe United Kingdom CW1 4QJ

#### Study participating centre York Teaching Hospitals NHS Foundation Trust

Wigginton Road York United Kingdom YO31 8HE

#### Study participating centre Leeds Teaching Hospitals NHS Foundation Trust

St James Hospital Beckett Street Leeds United Kingdom LS9 7TF

#### Study participating centre

University Hospitals Birmingham NHS Foundation Trust

Queen Elizabeth Hospital Birmingham Mindelsohn Way Birmingham United Kingdom B15 2GW

#### Study participating centre County Durham and Darlington NHS Foundation Trust

North Road Durham United Kingdom DH1 5TW

#### Study participating centre Blackpool Teaching Hospitals NHS Foundation Trust

Blackpool Victoria Hospital Whinney Heys Road Blackpool United Kingdom FY3 8NR

#### Study participating centre Stockport NHS Foundation Trust

Stepping Hill Hospital Stockport United Kingdom SK2 7JE

# Study participating centre The Mid Yorks Hospital NHS Foundation Trust

Pinderfields Hospital Wakefield United Kingdom WF1 4DG

# Sponsor information

#### Organisation

University of Manchester

**ROR** 

# Funder(s)

#### Funder type

Charity

#### **Funder Name**

Epilepsy Research UK

#### Alternative Name(s)

Epilepsy Research UK, The Epilepsy Research Institute, ERUK

#### **Funding Body Type**

Private sector organisation

#### **Funding Body Subtype**

Other non-profit organizations

#### Location

United Kingdom

### **Results and Publications**

#### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Dr Rebecca Bromley, rebecca.bromley@manchester.ac.uk, in electronic format, within a year of the study end for utilisation in studies with similar aims and where the participant has consented.

#### IPD sharing plan summary

Available on request

#### **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
HRA research summary			28/06/2023	No	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes