

## **Seizures - Investigations**

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**Approved by:** Neonatal Guidelines Group 17/02/2015.  
Reviewed Neonatal Clinical Guidelines 24.04.2018

**Review date:** 01.05.2021

**Version:** 3

## Investigations for neonatal seizures

Seizures occur most commonly in the neonatal period than in any other period of life and suggest a serious neurological abnormality. Furthermore, they are more likely to occur in the preterm than the term infant and the list of aetiologies are vast.<sup>1,2</sup> An incidence of 0.15 – 3.5 per 1000 births has been reported with higher rates in preterm infants.<sup>3,4,5</sup> The majority of seizures are reactive, secondary to an underlying condition.<sup>2</sup> The commonest cause of seizures in the term infant is Hypoxic Ischaemic Encephalopathy while in preterm infants it is intracranial haemorrhage. Other aetiologies include:<sup>1,6,7</sup>

- Infection
- Intracranial pathology
- Electrolyte imbalance
- Hypoglycaemia
- Inborn errors of metabolism
- Drug withdrawal
- Familial
- Idiopathic

When considering the underlying aetiology of seizures and appropriate investigations it is important that we are guided by the individual case. A structured approach should be taken as shown below.

1. **History** – antenatal, perinatal, family (including family tree with demographics and consanguinity) and feeding history. Consider risk factors including.<sup>2,8</sup>
  - **Maternal**
    - Advancing maternal age
    - Nulliparity
    - Pre existing/gestational diabetes
    - Illicit drug use
  - **Intrapartum**
    - Fetal distress
    - Placental abruption, cord prolapse, prolonged second stage
    - Maternal pyrexia/chorioamnionitis
  - **Neonatal**
    - Lower gestational age
    - Post term (>42 weeks)
    - Low birth weight
    - Male
2. **Clinical** – detailed neurological and general examination<sup>8</sup>
  - **Neurological** – level of consciousness, tone, gaze, posture, cranial nerves (pupillary reaction, facial movements, suck, gag reflex) and reflexes. Documentation and description of seizures should be charted.
  - **General** - looking for underlying disease
    - Skin                    bleeding, bruising, neurocutaneous lesions,                    birth marks.
    - Cardiac                to include auscultation for murmurs and possible bruits over fontanelle (AV malformation).
    - Abdominal            looking for intra-abdominal masses.

**3. 1<sup>st</sup> line investigations<sup>2,8,9,10,11</sup>**

- Blood gas – pH, pCO<sub>2</sub>, BE, lactate, glucose, ionised calcium
- FBC
- U&E – renal function, Na, K, Mg, Ca
- LFT's
- Ammonia
- Septic screen – Blood culture, CRP, LP (consider viral PCR's if appropriate), urine culture

***Neuroimaging***

- Cranial USS – to identify intraventricular haemorrhage, arterial stroke, malformations and infections. Should be carried out as early as possible.
- MRI – to identify changes of HIE, arterial and venous stroke, meningitis/encephalitis, inborn errors of metabolism and congenital malformations. Can be carried out later into investigations.

***Electrophysiology***

- aEEG – all babies being investigated for seizures require EEG monitoring.
- Multichannel EEG – contact Neurophysiology Department (SLH)

If abnormalities are found in first line investigations, further investigations may be required to determine the underlying cause for the abnormal result. **Discussion with Neurology, Genetics and the Metabolic Team would be appropriate for advice on further management and investigations.**

**4. 2<sup>nd</sup> line investigations**

- Metabolic screen – serum amino acids, acylcarnitine, TORCH, urine organic and amino acids, urine reducing substances
- Genetic testing
- LP – neurotransmitters, paired glucose/lactate/amino acids

The National Metabolic Biochemistry Website has some useful guidelines -

<http://www.metbio.net/metbioGuidelines.asp>

including presentation and investigations for suspected metabolic disorders.

**SEIZURE INVESTIGATIONS LIST**

Name

Hospital number

Date of birth

INVESTIGATION	DATE SENT	DATE EXPECTED	RESULT
<b>Blood gas</b>			
<b>FBC</b>			
<b>U&amp;E's</b>			
<b>LFT's</b>			
<b>Ammonia</b>			
<b>Septic screen</b> <ul style="list-style-type: none"> <li>• Blood cultures</li> <li>• Urine culture</li> <li>• LP</li> <li>• CRP</li> </ul>			
<b>Metabolic screen</b> <ul style="list-style-type: none"> <li>• Acylcarnitine</li> <li>• Serum AA's</li> <li>• Urine OA's</li> <li>• Urine AA's</li> <li>• Urine reducing substances</li> </ul>			
<b>Genetic testing (discuss with genetics)</b>			
<b>TORCH</b>			
<b>CSF</b>			
<b>Cr USS</b>			
<b>MRI</b>			
<b>EEG</b>			

**INVESTIGATION SHEET TO BE FILED BEHIND DAILY BLOOD TESTS  
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