# DISEASE:
# Benign familial neonatal epilepsy

| | |
|---|---|
| **NAME:** | Benign familial neonatal epilepsy |
| **DESCRIPTION:** | Benign familial neonatal epilepsy (BFNE) is a rare genetic epilepsy syndrome characterized by the occurrence of afebrile seizures in otherwise healthy newborns with onset in the first few days of life. |
| **ORPHACODE:** | 1949 |
| **SYNONYMS:** | BFNS<br>Benign familial neonatal convulsions<br>Benign familial neonatal seizures |
| **XREF(S):** | Orphanet<br>MeSH<br>MeSH<br>MedDRA<br>ICD-10<br>OMIM<br>OMIM<br>OMIM<br>OMIM |
| **ANALYTE(S):** | KCNQ2<br>KCNQ3 |

| | |
|---|---|
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

---

# RELATED CONTENT

## Related Genetic Tests

- Epilepsy (gene panel)

## Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

## Related Analytes

- potassium voltage-gated channel subfamily Q member 2
- potassium voltage-gated channel subfamily Q member 3

## Related Gene Panels

- Rare epilepsy with developmental delay (> 240 genes) - UZA