Table S1. Main and secondary diagnose.

Main diagnoses	***n* = 20**
Hypoxic-ischemic encephalopathy	2 (10)
Complex developmental disorder of unclear origin	1 (5)
De-Lange Syndrome	1 (5)
Edwards syndrome	1 (5)
Epileptic encephalopathy	1 (5)
Epileptic encephalopathy (ARV1 mutation)	1 (5)
Epileptic encephalopathy (CACNA1E mutation)	1 (5)
Epileptic encephalopathy (SPTAN1 duplication)	1 (5)
Holoprosencephaly	1 (5)
Krabbe disease	1 (5)
Leukencephalopathy	1 (5)
Metachromatic leukodystrophy	1 (5)
Nemaline myopathy (ACTA 1 deficiency)	1 (5)
Neonatal cerebral leukomalacia following preterm birth	1 (5)
Nonketotic hyperglycinemia	1 (5)
Opitz trigonocephaly (C) syndrome	1 (5)
Perinatal asphyxia and hypoxic-ischemic encephalopathy	1 (5)
Pontocerebellar hypoplasia	1 (5)
Severe neurodevelopmental disorder of unknown origin	1 (5)
Secondary diagnoses	
Symptomatic focal epilepsy	10 (50)
Bilateral spastic cerebral palsy	8 (40)
Hypotonia	3 (15)
Bilateral spastic tetraparesis	3 (15)
Structural focal epilepsy	2 (10)
Obstructive sleep apnea	2 (10)
Symptomatic epilepsy	2 (10)
High-flow nasal canula	1 (5)
Bilateral hippocampal sclerosis	1 (5)
Dyskinesia	1 (5)
Epilepsy	1 (5)
Epilepsy and infantile spasms	1 (5)
Hypertrophic cardiomyopathy	1 (5)
Metabolic epilepsy	1 (5)
Chronic respiratory failure	1 (5)
Tracheostomy	1 (5)
Ventricular septal defect	1 (5)

Counts as *n* (% of included cases)