**Supplementary Table 2 – Prevalence of Childhood Epilepsy Syndromes**

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| **Epilepsy Syndrome** | **Number of Patients (%)** |
| Neonatal Epileptic Encephalopathies with Burst Suppression\* | 5 (2) |
| Self-limited neonatal/infantile epilepsy | 2 (1) |
| Benign Myoclonic Epilepsy of Infancy | 1 (<1) |
| Infantile Epilepsy with migrating focal seizures | 3 (1) |
| Infantile Spasms | 36 (12) |
| Genetic Epilepsies with Febrile Seizures Plus (GEFS+) | 5 (2) |
| Dravet syndrome | 15 (5) |
| Lennox Gastaut Syndrome | 29 (10) |
| Myoclonic atonic epilepsy (Doose syndrome) | 8 (3) |
| Self-limited Occipital Epilepsy | 20 (7) |
| Self-limited Epilepsy with Centro-temporal Spikes | 82 (28) |
| Childhood Absence Epilepsy | 37 (13) |
| Absence Epilepsy with eyelid myoclonia (Jeavon's syndrome) | 2 (1) |
| Juvenile Absence Epilepsy | 16 (5) |
| Juvenile Myoclonic Epilepsy | 30 (10) |
| Progressive Myoclonic Epilepsy | 1 (<1) |

*\*Early infantile epileptic encephalopathy (EIEE, Ohtahara syndrome), Early myoclonic encephalopathy (EME)*