Presenter	Poster No	Abstract Title
F Abdel-Salam	14	SEPT9 Gene Duplication in a Child with Bilateral Neuralgic Amyotrophy
R Aljeaid*	116	Clinical audit of oesophageal pH and impedance monitoring in children with neurodisabilities
NM Allen*	33	Exome Sequencing in Early Onset Epileptic Encephalopathy: A Single Centre Study of 42 Patients
G P Ambegaonkar	45	Seizure cessation in a breast fed infant with medically refractory epilepsy following commencement of Modifies Atkins diet by mother
T Antoniadi	17	Gene panel testing reveals causative mutation in the alanyl-tRNA synthetase gene (AARS) in two patients with inherited peripheral neuropathy
F Ardestani*	94	Neurologically asymptomatic case of Adrenoleukodystrophy, potential for Bone Marrow Transplant
L Arkush*	38	Neuroimaging in acute admissions with seizures
MOE Babiker*	44	De novo gain-of-function mutation in SCN11A: No pain, More pain, or a bit of both?
B Bansagi*	8	Autosomal dominant distal hereditary motor neuropathy with neuromuscular transmission defect in a large British pedigree due to a mutation in the Synaptotagmin 2 (SYT2) gene
B Bansagi*	15	STAT5B mutation as a cause of growth retardation associated with neuromuscular symptoms
N Bhangu*	92	A case of transient pseduobulbar palsy in severe diabetic ketoacidosis
NK Bhangu*	109	Neuro-developmental Assessment in Children Pre Cardio-Pulmonary Bypass
R Boal*	75	Neurosarcoidosis with ? concurrent CNS Herpes Simplex viral reactivation
J Borbone	90	Tyrosine hydroxylase deficiency: a new patient with unusual clinical and neuroimaging
CM Brand	107	Bones and brains: a prospective case-control study of seasonal vitamin D in childhood epilepsy

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Presenter	Poster No	Abstract Title
LC Brennan**	64	Pegboard performance at altitude in normal children: results from Young Everest 2
A Brunklaus*	35	Genotype phenotype associations in SCN2A related epilepsies
S Byrne*	70	Screening for catatonic features in children with NMDA receptor antibody encephalitis: a review of published cases
DI Campbell	119	Enteral Feed Induced Dystonic Spasms: The role of PN
G Chow	11	A Novel Mutation causing Congenital Myasthenia
JH Cross	3	Sleep benefits memory consolidation in children with focal epilepsy
E Crossley**	115	Parental and child experiences from a service evaluation of the TANDeM MDT clinic
KCS Dalpatadu	113	Recurrent apnoeic episodes with cyanosis responding to acetazolamide in child with
AZ Demetriou*	59	Are We Considering Neonatal Stroke Early Enough in our Differential Diagnosis? A Case Report
S Deoraj**	117	Posterior reversible encephalopathy syndrome in Children: Towards an improved clinico-radiological definition
I Dey*	83	â€˜Brain on fireâ€™ with months of smoke- Addressing Ongoing dilemmas in management
A Donald*	66	Syndrome of the Trephined â€“ A Rare Intrusion to Neurorehabilitation
A Donald*	80	An unusual case of subcortical white matter lesions in a young adult with transverse myelitis
BM Dunning-Davies*	4	Investigating Isolated Intellectual Disability.
BM Dunning-Davies*	96	A pragmatic guideline for annual review in a child with NF1
BM Dunning-Davies*	103	UK90 OFC charts: a head above WHO?
NT EL Tantawi	20	The role of a controlled Propofol infusion in the rehabilitation of functional dystonia in a teenage girl

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Presenter	Poster No	Abstract Title
NT EL Tantawi*	57	Starting Antieplipeltic medications by non specialists : what are the hazards?
S Egger*	76	Vomiting and Singultus as presenting signs and symptoms of Neuromyelitis Optica spectrum disorder in a 14 years old boy.
O F El-Rashidy	56	Non -convulsive status epilepticus
CM Eltze	48	Ketogenic diet in refractory status epilepticus: North London experience
MR Eyre*	73	Anticipating intensive care requirement in paediatric Guillain-BarrÃ© syndrome
A Fadilah*	104	Consider Spinal Cord Arteriovenous Malformation in the Differential Diagnosis of Spinal Cord Swelling with Haemorrhage
T Fosi*	49	Pyridoxine-dependent epilepsy presenting as early neonatal metabolic encephalopathy with hyperlactaemia
S Ghani*	82	Paroxysmal autonomic instability with dystonia syndrome (PAID) with enteroviral encephalitis
K Gorman*	72	Subacute encephalopathy as presentation of horizontal HIV-1/EBV acquisition in a preadolescent girl
K Gorman*	88	Mucolipidosis Type IV : The first Irish case.
R Grattan*	68	Early Thrombectomy following Basilar artery thrombosis
H Griffin*	108	Should we recommend hip surveillance radiographs in children with bilateral cerebral palsy earlier than current local guidance of 30 months?
M Guglieri	21	FOR-DMD: Double-blind randomised trial to optimise corticosteroid regime in Duchenne Muscular Dystrophy (DMD)
Y Hacohen	78	Traumatic head injury N-methyl-D-aspartate receptor (NMDA) antibody mediated leukoencephalopathy
Y Hacohen*	79	N-methyl-D-aspartate receptor (NMDA) antibody encephalitis mimicking an autistic regression in a toddler.
A Hilley	28	Appropriate Phenytoin Level Monitoring in a Paediatric Population
S Hummaida*	114	Kernohanâ€™s notch: False Localising Hemiparesis in a child

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Presenter	Poster No	Abstract Title
MB Hunter**	99	NEUROdevelopment in PReschool Children Of FIfe and Lothian Epilepsy Study: NEUROPROFILES â€“ A population-based study
MA Iro	85	A phase III multicenter randomized controlled double blind trial to assess the role of intravenous immunoglobulin (IVIG) in the treatment of childhood encephalitis (The IgNiTE study)
N Ismayilova*	87	Reversible cerebral and brainstem white matter abnormalities associated with classical homocystinuria
TJ Jackson	106	Longitudinal analysis of the neurological features of Ataxia Telangiectasia
S Jain*	51	Interesting cases of Pyridoxine dependent epilepsy in late childhood
SS Jose	120	A Paediatricians guide to setting up an Epilepsy Database
S Joseph*	77	Seronegative neuromyelitis optica(NMO) with longuituidinally extensive transverse myelitis (LETM). Implication of myelin-oligodendrocyte glycoprotein (MOG-Abs) on clinical practice.
KS Kallambella*	102	Gorlin Syndrome
V Kalra*	18	A case of acute rhabdomyolysis with intermittent myositis and persistently low levels of Parvovirus B19 infection
V Kalra*	55	Ring chromosome 20 as a rare cause of refractory epilepsy in children: Case report and literature
SD Karayiannis*	62	Paediatric Head Injuries: Improving quality of assessment in a District General Hospital.
DST Kariyawasam*	71	An unusual case of Infantile Botulism: Clostridium botulinum Bf infection in UK
OO Kehinde*	31	Prevalence of symptoms of disordered sleep in children with epilepsy
JE Kisler	1	Can Taping improve hand posture and use in infants with asymmetric cerebral palsy?
R Kumar	112	A pilot high-definition video-enabled tele-clinic service in tertiary paediatric neurology

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Presenter	Poster No	Abstract Title
L Lagae	42	Impact of adjunctive perampanel on behaviour in adolescents with refractory partial-onset seizures
B Mahendran**	84	Inflammatory childhood cranial neuropathies: clinical features, management and outcome.
A McTague*	26	The first UK case of GABRBR3 mutation causing early onset epilepsy and developmental delay.
A McWilliams*	100	A study of Psychogenic Non-Epileptic Seizures in children and adolescents; characteristics and experiences of a case series
LC Mellish*	39	Treatment of rolandic epilepsy and Panayiotopoulos syndrome - a survey of clinical practice and clinical trial feasibility
Leena Mewasingh	98	Visual snow in a young child
SR Mordekar	54	A comparison of ambulatory EEG followed by video telemetry in children
F Motaleb*	50	Convulsive status epilepticus in the children of South Yorkshire and Humber: an audit of patients requiring intensive care admission
F Motaleb*	110	It may not be as serious as it looks!: An Unexpected result on brain biopsy
V Mundada*	16	Hopkins Syndrome Can be caused by Enterovirus-68: A Common Pathogen
V Nesbitt*	86	Benign Intracranial Hypertension & Visual Loss Secondary to Vitamin A Deficiency in a Child with Autism
V Nesbitt*	91	General anaesthesia in children with mitochondrial disease
V Nesbitt*	97	Cardiac transplantation in children with mitochondrial disease
MT Ong*	22	Apnoeas as manifestation of left temporal lobe seizure
MT Ong*	36	KCTD7 mutation as a cause of infantile neuronal ceroid lipofucinosis
MT Ong*	65	Post-anoxic reticular reflex myoclonus â€“ a form of Lance-Adams myoclonus?
APJ Parker	95	Adrenoleukodystrophy - An Avoidable and Treatable Illness

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Presenter	Poster No	Abstract Title
D Pathak*	58	Red cell indices at presentation of paroxysmal disorders: local results from Epilepsy 12
D Ram*	9	Novel Use of Eculizumab in Guillain-BarrÃ© Syndrome
D Ram*	10	TK2 Mutation: An Expanding Clinical Phenotype
D Ram*	13	Intraneural Perineurioma: A Rare Cause of Monomelic Weakness
D Ram*	61	Value of Cervical Spine Imaging after Vertebral Artery Dissection: A Case Report
D Ram*	67	Down Syndrome and Moyamoya Disease: An Autoimmune Link?
D Ram*	101	A Rare Case of Childhood Gliomatosis Cerebri Presenting as Encephalitis
M Ramphul*	2	Isolated intellectual disability: utility of investigations for inborn errors of metabolism
M Ramphul*	52	Macrocytosis without anaemia in children on the ketogenic diet â€“ a newly described complication of treatment
RR Rattihalli*	32	Another phenotype of GLUT1 deficiency syndrome, with different phenotypes in dominant transmission.
B Renfroe	30	Interim efficacy and safety analysis of adjunctive perampanel in the adolescent population from the extension phase of three double-blind, placebo-controlled, Phase III (core) studies in patients with refractory partial-onset seizures
RE Rosch*	40	Copy-number variations in absence epilepsy
A Salonikiou*	74	Association of systemic disability and optical coherence tomography (OCT) findings in children with demyelinating disorders
L Scotson	6	Chest size in children with cerebral palsy
A Shahin*	24	Audit of Epilepsy Care in Children with Epilepsy attending Special Schools
H Shekhar	5	Stock Car racing related cranio-cervical junction injury presenting with bilateral hypoglossal and abducens nerve palsy

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Presenter	Poster No	Abstract Title
A Silwal*	89	Cystic leukoencephalopathy due to NDUFV1 mutation - complex I deficiency in a mitochondrial disorder - a report of the phenotype and its rare association with Primary Hyperoxaluria
J Singh*	63	Laboratory predictors of death and poor functional outcome following paediatric traumatic brain injury
T Smallbone*	46	Vigabatrin for Refractory Epilepsy ...out because of sight and out of mind?
M Smith*	111	Clinical audit of video-EEG head-up tilt tests and ocular compression tests in children and young people
B Soliman**	53	The role of cerebral haemodynamics, breathing patterns, iron deficiency and autoimmunity in acute paediatric seizure activity
M Spyridon	34	Effects of adjunctive zonisamide treatment on weight and body mass index in paediatric patients with partial epilepsy
HE Steele*	105	The Eye of the Tiger with Night Blindness: Retinitis Pigmentosa as a presenting feature of PKAN
N Stewart*	19	A new variant of Dynein Cytoplasmic 1 Heavy Chain 1 (DYNC1H1) associated with lower limb wasting and contractures and global developmental delay with cortical abnormalities on MRI
N Swiderska*	27	Provision of Ketogenic Dietary Therapy across the UK â€“ a national survey of Paediatric Neurologists.
JD Symonds	47	Genetic Heterogeneity in Dravet Syndrome: the role of SCN2A, SCN1B, and GABRG2 variants
LK Tapper**	93	X-linked Adrenoleukodystrophy- How to improve presymptomatic identification
MJ Taylor*	29	Narcolepsy in Children
HK Terry*	25	Use of Pyridoxine in Drug-resistant Epilepsy with Dialeptic and Myoclonic Semiology
JT Wiggett	43	Can the QTc interval be calculated accurately by consultants and clinical physiologists using a single lead electrocardiogram during the electroencephalogram?
K Wood*	81	Periodic Lateralised Epileptiform Discharges (PLEDS) in TB Meningitis

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Presenter	Poster No	Abstract Title
I Woodcock*	12	The prevalence of neuromuscular disease in the paediatric population in Yorkshire
SK Wright*	23	Automated seizure detection in a mouse model of autoimmune epilepsy
Z Yusuf**	60	Vitamin B12 Deficiency - A Poorly Recognised and Preventable Childhood Stroke Risk Factor
Z Yusuf**	69	A case series on childhood acquired hypoxic-ischemic encephalopathy
Y de Alwis*	37	Temporal lobe non convulsive status epilepticus in Ornithine transcarbamylase (OTC) deficiency
HJA van Ruiten*	7	Lowering the age of diagnosis of Duchenne muscular dystrophy

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