

Helen Cross

List of Publications by Year in descending order

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Version: 2024-02-01

80
papers

9,665
citations

212478

28
h-index

73587

79
g-index

85
all docs

85
docs citations

85
times ranked

10472
citing authors

#	ARTICLE	IF	CITATIONS
1	Epilepsy-related stigma and attitudes: Systematic review of screening instruments and interventions – Report by the International League Against Epilepsy Task Force on Stigma in Epilepsy. <i>Epilepsia</i> , 2022, 63, 598-628.	2.6	12
2	Systematic review of frequency of felt and enacted stigma in epilepsy and determining factors and attitudes toward persons living with epilepsy – Report from the International League Against Epilepsy Task Force on Stigma in Epilepsy. <i>Epilepsia</i> , 2022, 63, 573-597.	2.6	35
3	Broadband-NIRS System Identifies Epileptic Focus in a Child with Focal Cortical Dysplasia – A Case Study. <i>Metabolites</i> , 2022, 12, 260.	1.3	7
4	Guidance on Dravet syndrome from infant to adult care: Road map for treatment planning in Europe. <i>Epilepsia Open</i> , 2022, 7, 11-26.	1.3	32
5	International League Against Epilepsy classification and definition of epilepsy syndromes with onset in childhood: Position paper by the ILAE Task Force on Nosology and Definitions. <i>Epilepsia</i> , 2022, 63, 1398-1442.	2.6	263
6	Methodology for classification and definition of epilepsy syndromes with list of syndromes: Report of the ILAE Task Force on Nosology and Definitions. <i>Epilepsia</i> , 2022, 63, 1333-1348.	2.6	84
7	Core outcome set development for childhood epilepsy treated with ketogenic diet therapy: Results of a scoping review and parent interviews. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2022, 99, 54-67.	0.9	3
8	ILAE classification and definition of epilepsy syndromes with onset in neonates and infants: Position statement by the ILAE Task Force on Nosology and Definitions. <i>Epilepsia</i> , 2022, 63, 1349-1397.	2.6	237
9	Immunomodulation With Azathioprine Therapy in Rasmussen Syndrome. <i>Neurology</i> , 2021, 96, e267-e279.	1.5	8
10	Epilepsy in 2020 – a new dawn. <i>Lancet Neurology</i> , The, 2021, 20, 8-10.	4.9	4
11	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. <i>Epilepsia Open</i> , 2021, 6, 160-170.	1.3	3
12	The initial impact of the SARS-CoV-2 pandemic on epilepsy research. <i>Epilepsia Open</i> , 2021, 6, 255-265.	1.3	2
13	Fenfluramine as antiseizure medication for epilepsy. <i>Developmental Medicine and Child Neurology</i> , 2021, 63, 899-907.	1.1	20
14	Newer versus older antiseizure medications: further forward?. <i>Lancet</i> , The, 2021, 397, 1327-1329.	6.3	2
15	Time to onset of cannabidiol (CBD) treatment effect in Lennox-Gastaut syndrome: Analysis from two randomized controlled trials. <i>Epilepsia</i> , 2021, 62, 1130-1140.	2.6	20
16	Classification of complications of epilepsy surgery and invasive diagnostic procedures: A proposed protocol and feasibility study. <i>Epilepsia</i> , 2021, 62, 2685-2696.	2.6	5
17	Epilepsy care during the COVID-19 pandemic. <i>Epilepsia</i> , 2021, 62, 2322-2332.	2.6	48
18	Quantitative MRI susceptibility mapping reveals cortical signatures of changes in iron, calcium and zinc in malformations of cortical development in children with drug-resistant epilepsy. <i>NeuroImage</i> , 2021, 238, 118102.	2.1	11

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19	The impact of SARS-CoV-2 vaccination in Dravet syndrome: A UK survey. <i>Epilepsy and Behavior</i> , 2021, 124, 108258.	0.9	15
20	Hypothalamic Hamartomas. <i>Neurology</i> , 2021, 97, 864-873.	1.5	12
21	Optimising epilepsy care throughout the Afghan refugee crisis. <i>Lancet, The</i> , 2021, 398, 1563.	6.3	2
22	Mapping degeneration of the visual system in long-term follow-up after childhood hemispherectomy "A series of four cases. <i>Epilepsy Research</i> , 2021, 178, 106808.	0.8	2
23	Meeting report: EpiXchange II brings together European epilepsy research projects to discuss latest advances. <i>Epilepsy Research</i> , 2021, 178, 106811.	0.8	1
24	Optimal clinical management of children receiving ketogenic parenteral nutrition: a clinical practice guide. <i>Developmental Medicine and Child Neurology</i> , 2020, 62, 48-56.	1.1	23
25	Trends in pediatric epilepsy surgery in Europe between 2008 and 2015: Country-, center-, and age-specific variation. <i>Epilepsia</i> , 2020, 61, 216-227.	2.6	44
26	The concept of disease modification. <i>European Journal of Paediatric Neurology</i> , 2020, 24, 43-46.	0.7	6
27	Ketogenic diet therapy in infants with epilepsy. <i>Paediatrics and Child Health (United Kingdom)</i> , 2020, 30, 356-360.	0.2	1
28	An accelerated shift in the use of remote systems in epilepsy due to the COVID-19 pandemic. <i>Epilepsy and Behavior</i> , 2020, 112, 107376.	0.9	29
29	Genetics in the epilepsies "A broadening concept. <i>European Journal of Paediatric Neurology</i> , 2020, 24, 8.	0.7	0
30	Optimising Evidence-Based Psychological Treatment for the Mental Health Needs of Children with Epilepsy: Principles and Methods. <i>Clinical Child and Family Psychology Review</i> , 2020, 23, 284-295.	2.3	17
31	Association of quality of paediatric epilepsy care with mortality and unplanned hospital admissions among children and young people with epilepsy in England: a national longitudinal data linkage study. <i>The Lancet Child and Adolescent Health</i> , 2019, 3, 627-635.	2.7	16
32	Advancing research toward faster diagnosis, better treatment, and end of stigma in epilepsy. <i>Epilepsia</i> , 2019, 60, 1281-1292.	2.6	17
33	Determinants of IQ outcome after focal epilepsy surgery in childhood: A longitudinal case-control neuroimaging study. <i>Epilepsia</i> , 2019, 60, 872-884.	2.6	32
34	Drug Development for Rare Paediatric Epilepsies: Current State and Future Directions. <i>Drugs</i> , 2019, 79, 1917-1935.	4.9	13
35	Fenfluramine hydrochloride for the treatment of seizures in Dravet syndrome: a randomised, double-blind, placebo-controlled trial. <i>Lancet, The</i> , 2019, 394, 2243-2254.	6.3	227
36	Dravet syndrome: Treatment options and management of prolonged seizures. <i>Epilepsia</i> , 2019, 60, S39-S48.	2.6	56

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37	A new multichannel broadband NIRS system for quantitative monitoring of brain hemodynamics and metabolism during seizures. , 2019, , .		1
38	The ketogenic diet is effective for refractory epilepsy associated with acquired structural epileptic encephalopathy. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 718-723.	1.1	8
39	Mechanisms of action for the medium-chain triglyceride ketogenic diet in neurological and metabolic disorders. <i>Lancet Neurology</i> , The, 2018, 17, 84-93.	4.9	296
40	A web-based diagnostic reference centre for the European Reference Network "EpiCare" recommendations of the eNeuropathology working group. <i>Epileptic Disorders</i> , 2018, 20, 339-345.	0.7	2
41	Optimal clinical management of children receiving dietary therapies for epilepsy: Updated recommendations of the International Ketogenic Diet Study Group. <i>Epilepsia Open</i> , 2018, 3, 175-192.	1.3	412
42	Genome-wide association study: Exploring the genetic basis for responsiveness to ketogenic dietary therapies for drug-resistant epilepsy. <i>Epilepsia</i> , 2018, 59, 1557-1566.	2.6	23
43	Pre- and postsurgical cognitive trajectories and quantitative sMRI changes in Rasmussen syndrome. <i>Epilepsia</i> , 2018, 59, 1210-1219.	2.6	10
44	Current standards of neuropsychological assessment in epilepsy surgery centers across Europe. <i>Epilepsia</i> , 2017, 58, 343-355.	2.6	69
45	Operational classification of seizure types by the International League Against Epilepsy: Position Paper of the ILAE Commission for Classification and Terminology. <i>Epilepsia</i> , 2017, 58, 522-530.	2.6	2,191
46	Management of Dravet syndrome and emerging therapy options. <i>Expert Opinion on Orphan Drugs</i> , 2017, , .	0.5	0
47	Practice guideline summary: Sudden unexpected death in epilepsy incidence rates and risk factors. <i>Neurology</i> , 2017, 88, 1674-1680.	1.5	384
48	Ketogenic diet in the treatment of epilepsy in children under the age of 2 years: study protocol for a randomised controlled trial. <i>Trials</i> , 2017, 18, 195.	0.7	17
49	Medical management and antiepileptic drugs in hypothalamic hamartoma. <i>Epilepsia</i> , 2017, 58, 16-21.	2.6	27
50	Trial of Cannabidiol for Drug-Resistant Seizures in the Dravet Syndrome. <i>New England Journal of Medicine</i> , 2017, 376, 2011-2020.	13.9	1,148
51	Professor Brian George Richard Neville. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 243-244.	0.7	0
52	An examination of biochemical parameters and their association with response to ketogenic dietary therapies. <i>Epilepsia</i> , 2017, 58, 893-900.	2.6	13
53	Histopathological Findings in Brain Tissue Obtained during Epilepsy Surgery. <i>New England Journal of Medicine</i> , 2017, 377, 1648-1656.	13.9	621
54	Practice Guideline Summary: Sudden Unexpected Death in Epilepsy Incidence Rates and Risk Factors: Report of the Guideline Development, Dissemination, and Implementation Subcommittee of the American Academy of Neurology and the American Epilepsy Society. <i>Epilepsy Currents</i> , 2017, 17, 180-187.	0.4	87

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55	Integrated care for childhood epilepsy: ongoing challenges and lessons for other long-term conditions. <i>Archives of Disease in Childhood</i> , 2016, 101, 1057-1062.	1.0	14
56	Ketogenic dietary therapies in adults with epilepsy: a practical guide. <i>Practical Neurology</i> , 2016, 16, 208-214.	0.5	38
57	Current practices in long-term video-EEG monitoring services: A survey among partners of the E-PILEPSY pilot network of reference for refractory epilepsy and epilepsy surgery. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2016, 38, 38-45.	0.9	67
58	Targeted Treatment in Childhood Epilepsy Syndromes. <i>Current Treatment Options in Neurology</i> , 2016, 18, 29.	0.7	7
59	Ketogenic diet guidelines for infants with refractory epilepsy. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 798-809.	0.7	134
60	The genetic landscape of the epileptic encephalopathies of infancy and childhood. <i>Lancet Neurology</i> , The, 2016, 15, 304-316.	4.9	474
61	Cognitive consequences of early versus late antiepileptic drug withdrawal after pediatric epilepsy surgery, the TimeToStop (TTS) trial: study protocol for a randomized controlled trial. <i>Trials</i> , 2015, 16, 482.	0.7	18
62	Favourable response to ketogenic dietary therapies: undiagnosed glucose 1 transporter deficiency syndrome is only one factor. <i>Developmental Medicine and Child Neurology</i> , 2015, 57, 969-976.	1.1	8
63	Summary of recommendations for the management of infantile seizures: Task Force Report for the International League Against Epilepsy of Paediatrics. <i>Epilepsia</i> , 2015, 56, 1185-1197.	2.6	323
64	Transition in lesional focal epilepsy, and following epilepsy surgery. <i>Epilepsia</i> , 2014, 55, 34-36.	2.6	9
65	The ketogenic diet component decanoic acid increases mitochondrial citrate synthase and complex I activity in neuronal cells. <i>Journal of Neurochemistry</i> , 2014, 129, 426-433.	2.1	153
66	CARDIAC FEATURES IN ADULTS WITH ALTERNATING HEMIPLEGIA. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, e4.214-e4.	0.9	0
67	Diagnostic test utilization in evaluation for resective epilepsy surgery in children. <i>Epilepsia</i> , 2014, 55, 507-518.	2.6	174
68	Safety and tolerability of zonisamide in paediatric patients with epilepsy. <i>European Journal of Paediatric Neurology</i> , 2014, 18, 747-758.	0.7	16
69	Differential diagnosis of epileptic seizures in infancy including the neonatal period. <i>Seminars in Fetal and Neonatal Medicine</i> , 2013, 18, 192-195.	1.1	26
70	Nodding syndrome—a challenge for African public health. <i>Lancet Neurology</i> , The, 2013, 12, 125-126.	4.9	5
71	Advancing the management of childhood epilepsies. <i>European Journal of Paediatric Neurology</i> , 2013, 17, 334-347.	0.7	27
72	The epileptic encephalopathies. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2013, 111, 619-626.	1.0	21

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73	Neurological features of epilepsy, ataxia, sensorineural deafness, tubulopathy syndrome. <i>Developmental Medicine and Child Neurology</i> , 2013, 55, 846-856.	1.1	53
74	New Research With Diets and Epilepsy. <i>Journal of Child Neurology</i> , 2013, 28, 970-974.	0.7	14
75	Fever and fever-related epilepsies. <i>Epilepsia</i> , 2012, 53, 3-8.	2.6	31
76	Surgical Approaches to Treating Epilepsy in Children. <i>Current Treatment Options in Neurology</i> , 2012, 14, 620-629.	0.7	13
77	The effect of the classical and medium chain triglyceride ketogenic diet on vitamin and mineral levels. <i>Journal of Human Nutrition and Dietetics</i> , 2012, 25, 16-26.	1.3	54
78	A randomized trial of classical and medium-chain triglyceride ketogenic diets in the treatment of childhood epilepsy. <i>Epilepsia</i> , 2009, 50, 1109-1117.	2.6	343
79	The ketogenic diet for the treatment of childhood epilepsy: a randomised controlled trial. <i>Lancet Neurology</i> , The, 2008, 7, 500-506.	4.9	966
80	Eye rolling as a manifestation of clobazam toxicity in a child with epilepsy. <i>Developmental Medicine and Child Neurology</i> , 2007, 48, 612-615.	1.1	4