**2022**

**Amlie-Lefond C**. Pediatric Stroke-Are We Asking the Right Questions? The 2022 Sidney Carter Award Lecture. Neurology. 2022 Nov 8:10.1212/WNL.0000000000201487. doi: 10.1212/WNL.0000000000201487. Epub ahead of print. PMID: 36347625.

Benedetti GM, **Amlie-Lefond C**. Caring for Our Sickest Patients: The Expanding Role of the Pediatric Neurohospitalist. Neurology. 2022 Nov 1;99(18):781-782. doi: 10.1212/WNL.0000000000201267. PMID: 36316123.

**Chatterjee JH,** **Blume HK**. Migraine and Mental Health in Pediatrics. JAMA Pediatr. 2022 Oct 31. doi: 10.1001/jamapediatrics.2022.3948. Epub ahead of print. PMID: 36315116.

Vitanza NA, Wilson AL, Huang W, Seidel K, Brown C, Gustafson JA, Yokoyama JK, Johnson AJ, Baxter BA, Koning RW, Reid AN, Meechan M, Biery MC, Myers C, Rawlings-Rhea SD, Albert CM, Browd SR, Hauptman JS, Lee A, Ojemann JG, Berens ME, Dun MD, Foster JB, Crotty EE, Leary SES, Cole BL, Perez FA, Wright JN, Orentas RJ, Chour T, Newell EW, Whiteaker JR, Zhao L, Paulovich AG, Pinto N, **Gust J**, Gardner RA, Jensen MC, Park JR. Intraventricular B7-H3 CAR T cells for diffuse intrinsic pontine glioma: preliminary first-in-human bioactivity and safety. Cancer Discov. 2022 Oct 19:CD-22-0750. doi: 10.1158/2159-8290.CD-22-0750. Epub ahead of print. PMID: 36259971.

Guglieri M, Clemens PR, **Perlman SJ**, Smith EC, Horrocks I, Finkel RS, Mah JK, Deconinck N, Goemans N, Haberlova J, Straub V, Mengle-Gaw LJ, Schwartz BD, Harper AD, Shieh PB, De Waele L, Castro D, Yang ML, Ryan MM, McDonald CM, Tulinius M, Webster R, McMillan HJ, Kuntz NL, Rao VK, Baranello G, Spinty S, Childs AM, Sbrocchi AM, Selby KA, Monduy M, Nevo Y, Vilchez-Padilla JJ, Nascimento-Osorio A, Niks EH, de Groot IJM, Katsalouli M, James MK, van den Anker J, Damsker JM, Ahmet A, Ward LM, Jaros M, Shale P, Dang UJ, Hoffman EP. Efficacy and Safety of Vamorolone vs Placebo and Prednisone Among Boys With Duchenne Muscular Dystrophy: A Randomized Clinical Trial. JAMA Neurol. 2022 Oct 1;79(10):1005-1014. doi: 10.1001/jamaneurol.2022.2480. PMID: 36036925; PMCID: PMC9425287

Mayne EW, Mailo JA, Pabst L, Pulcine E, Harrar DB, Waak M, Rafay MF, Hassanein SM, **Amlie-Lefond C,** Jordan LC. Pediatric Stroke and Cardiac Disease: Challenges in Recognition and Management. Semin Pediatr Neurol. 2022 Oct;43:100992. doi: 10.1016/j.spen.2022.100992. Epub 2022 Aug 31. PMID: 36344023.

Surtees TL, Pearson R, Harrar DB, Lee S, **Amlie-Lefond CM**, Guilliams KP. Acute Hospital Management of Pediatric Stroke. Semin Pediatr Neurol. 2022 Oct;43:100990. doi: 10.1016/j.spen.2022.100990. Epub 2022 Aug 19. PMID: 36344020.

Patel ND, Batra M, Udomphorn Y, **Wainwright M**, Vavilala MS. Cerebral Autoregulation in Healthy Term Newborns: Brief Report. Pediatr Neurol. 2022 Oct;135:4-5. doi: 10.1016/j.pediatrneurol.2022.07.001. Epub 2022 Jul

Persa L, Shaw DW, **Amlie-Lefond C**. Why Would a Child Have a Stroke? J Child Neurol. 2022 Oct 9:8830738221129916. doi: 10.1177/08830738221129916. Epub ahead of print. PMID: 36214173.

Hartford EA, **Blume H**, Barry D, **Hauser Chatterjee J**, Law E. Disparities in the emergency department management of pediatric migraine by race, ethnicity, and language preference. Acad Emerg Med. 2022 Sep;29(9):1057-1066. doi: 10.1111/acem.14550. Epub 2022 Jul 7. PMID: 35726699.

Olson V, Chang IJ, Merritt Nd JL, **Mingbunjerdsuk D**. Refractory Myoclonus as a Presentation of Metabolic Stroke in A Child With Cobalamin B Methylmalonic Acidemia After Liver and Kidney Transplant. J Mov Disord. 2022 Sep;15(3):281-283. doi: 10.14802/jmd.21196. Epub 2022 May 26. PMID: 35614015; PMCID: PMC9536905.

**Natarajan N,** Benedetti G, Perez FA, Wood TR, German KR, **Lockrow JP**, Puia-Dumitrescu M, Myers E, Mietzsch U. Association Between Early EEG Background and Outcomes in Infants With Mild HIE Undergoing Therapeutic Hypothermia. Pediatr Neurol. 2022 Sep;134:52-58. doi: 10.1016/j.pediatrneurol.2022.06.006. Epub 2022 Jun 19. PMID: 35835026.

Hanaford A, **Johnson SC**. The immune system as a driver of mitochondrial disease pathogenesis: a review of evidence. Orphanet J Rare Dis. 2022 Sep 2;17(1):335. doi: 10.1186/s13023-022-02495-3. PMID: 36056365; PMCID: PMC9438277.

**Gust J**, Rawlings-Rhea SD, Wilson AL, Tulberg NM, Sherman AL, Seidel KD, Wu QV, Park JR, Gardner RA, Annesley CE. GFAP and NfL increase during neurotoxicity from high baseline levels in pediatric CD19-CAR T-cell patients. Blood Adv. 2022 Aug 25:bloodadvances.2022008119. doi: 10.1182/bloodadvances.2022008119. Epub ahead of print. PMID: 36006611.

Tran DK, Poliakov AV, Friedman SD, Goldstein HE, Shurtleff HA, **Bowen K**, **Patrick KE**, Warner M, **Novotny EJ,** Ojemann JG, Hauptman JS. Concordance of functional MRI memory task and resting-state functional MRI connectivity used in surgical planning for pediatric temporal lobe epilepsy. J Neurosurg Pediatr. 2022 Jul 29:1-6. doi: 10.3171/2022.6.PEDS221. Epub ahead of print. PMID: 35907201.

Erklauer JC, Thomas AX, Hong SJ, Appavu BL, Carpenter JL, Chiriboga-Salazar NR, Ferrazzano PA, Goldstein Z, Griffith JL, Guilliams KP, Kirschen MP, Lidsky K, Lovett ME, McLaughlin B, Munoz Pareja JC, Murphy S, O'Donnell W, Riviello JJ, Schober ME, Topjian AA, **Wainwright MS**, Simon DW, Pediatric Neurocritical Care Research Group. A Virtual Community of Practice: An International Educational Series in Pediatric Neurocritical Care. Children (Basel). 2022 Jul 20;9(7):1086. doi: 10.3390/children9071086. PMID: 35884070; PMCID: PMC9316633.

Tarazi RA, **Patrick KE**, Iampietro M, Apollonsky N. Hydroxyurea Use Associated with Nonverbal and Executive Skills in Sickle Cell Anemia. J Pediatr Psychol. 2021 Jul 20;46(6):710-718. doi: 10.1093/jpepsy/jsab015. PMID: 33706380.

Ostendorf AP, Axeen ET, Eschbach K, Fedak Romanowski E, **Morgan LA**, Gross P, Narayanan UG, Glader L, Noritz G; CEREBRAL PALSY RESEARCH NETWORK. Epilepsy and proxy-reported health-related quality of life in children and young people with non-ambulatory cerebral palsy. Dev Med Child Neurol. 2022 Jul 12. doi: 10.1111/dmcn.15336. Epub ahead of print. PMID: 35820144.

Faulhaber LD, D'Costa O, Shih AY, **Gust J.** Antibody-based *in vivo* leukocyte label for two-photon brain imaging in mice. Neurophotonics. 2022 Jul;9(3):031917. doi: 10.1117/1.NPh.9.3.031917. Epub 2022 May 24. PMID: 35637871; PMCID: PMC9128835.

Davies OMT, Garzon MC, Frieden IJ, Cottrell CE, Gripp KW, **Saneto RP**, Shwayder T, Mirzaa GM, Drolet BA. Cutaneous vascular anomalies associated with a mosaic variant of AKT3: Genetic analysis continues to refine the diagnosis, nomenclature, and classification of vascular anomalies. J Am Acad Dermatol. 2022 Jul;87(1):162-164. doi: 10.1016/j.jaad.2021.06.877. Epub 2021 Jul 6. PMID: 34237354; PMCID: PMC8733055.

Goldstein HE, Poliakov A, Shaw DW, Barry D, Tran K, **Novotny EJ**, **Saneto RP**, Marashly A, Warner MH, Wright JN, Hauptman JS, Ojemann JG, Shurtleff HA. Precision medicine in pediatric temporal epilepsy surgery: optimization of outcomes through functional MRI memory tasks and tailored surgeries. J Neurosurg Pediatr. 2022 Jul 1:1-12. doi: 10.3171/2022.5.PEDS22148. Epub ahead of print. PMID: 35901731.

Striano P, Auvin S, Collins A, Horvath R, Scheffer IE, Tzadok M, Miller I, Kay Koenig M, Lacy A, Davis R, Garcia-Cazorla A, **Saneto RP**, Brandabur M, Blair S, Koutsoukos T, De Vivo D. A randomized, double-blind trial of triheptanoin for drug-resistant epilepsy in glucose transporter 1 deficiency syndrome. Epilepsia. 2022 Jul;63(7):1748-1760. doi: 10.1111/epi.17263. Epub 2022 May 21. PMID: 35441706.

Hartford EA,  **H**, Barry D, **Hauser Chatterjee J**, Law E. Disparities in the emergency department management of pediatric migraine by race, ethnicity, and language preference. Acad Emerg Med. 2022 Jun 21. doi: 10.1111/acem.14550. Epub ahead of print. PMID: 35726699.

Qin ES, Gold LS, Singh N, Wysham KD, Hough CL, **Patel PB**, Bunnell AE, Andrews JS. Physical function and fatigue recovery at 6 months after hospitalization for COVID-19. PM R. 2022 Jun 21:10.1002/pmrj.12866. doi: 10.1002/pmrj.12866. Epub ahead of print. PMID: 35726518; PMCID: PMC9353339.

**Natarajan N**, **Benedetti G**, Perez FA, Wood TR, German KR, **Lockrow JP**, Puia-Dumitrescu M, Myers E, Mietzsch U. Association Between Early EEG Background and Outcomes in Infants With Mild HIE Undergoing Therapeutic Hypothermia. Pediatr Neurol. 2022 Jun 19;134:52-58. doi: 10.1016/j.pediatrneurol.2022.06.006. Epub ahead of print. PMID: 35835026

**Saneto RP**, Perez FA. Mitochondria-Associated Membrane Scaffolding with Endoplasmic Reticulum: A Dynamic Pathway of Developmental Disease. Front Mol Biosci. 2022 Jun 14;9:908721. doi: 10.3389/fmolb.2022.908721. PMID: 35775081; PMCID: PMC9237565.

Emmanuele V, Ganesh J, Vladutiu G, Haas R, Kerr D, **Saneto RP**, Cohen BH, Van Hove JLK, Scaglia F, Hoppel C, Rosales XQ, Barca E, Buchsbaum R, Thompson JL, DiMauro S, Hirano M; North American Mitochondrial Disease Consortium (NAMDC). Time to harmonize mitochondrial syndrome nomenclature and classification: A consensus from the North American Mitochondrial Disease Consortium (NAMDC). Mol Genet Metab. 2022 Jun;136(2):125-131. doi: 10.1016/j.ymgme.2022.05.001. Epub 2022 May 13. PMID: 35606253; PMCID: PMC9341219.

**Castillo-Pinto C**, Carpenter JL, Donofrio MT, Zhang A, Wernovsky G, Sinha P, Harrar D. Incidence and predictors of epilepsy in children with congenital heart disease. Cardiol Young. 2022 Jun;32(6):918-924. doi: 10.1017/S1047951121003279. Epub 2021 Aug 9. PMID: 34365987.

Crotty E, **Gust J**. Pediatric ependymoma: New perspectives on older trials. Neuro Oncol. 2022 Jun 1;24(6):949-950. doi: 10.1093/neuonc/noac060. PMID: 35325202; PMCID: PMC9159413.

Knight EMP, Amin S, Bahi-Buisson N, Benke TA, Cross JH, Demarest ST, Olson HE, Specchio N, Fleming TR, Aimetti AA, Gasior M, Devinsky O; Marigold Trial Group (**Saneto, RP**). Safety and efficacy of ganaxolone in patients with CDKL5 deficiency disorder: results from the double-blind phase of a randomised, placebo-controlled, phase 3 trial. Lancet Neurol. 2022 May;21(5):417-427. doi: 10.1016/S1474-4422(22)00077-1. Erratum in: Lancet Neurol. 2022 Jul;21(7):e7. PMID: 35429480.

Olson V, Chang IJ, Merritt Nd JL, **Mingbunjerdsuk D**. Refractory Myoclonus as a Presentation of Metabolic Stroke in A Child With Cobalamin B Methylmalonic Acidemia After Liver and Kidney Transplant. J Mov Disord. 2022 May 26. doi: 10.14802/jmd.21196. Epub ahead of print. PMID: 35614015.

Chen WL, Pao E, **Owens J**, Glass I, Pritchard C, Shirts BH, Lockwood C, Mirzaa GM. The utility of cerebrospinal fluid-derived cell-free DNA in molecular diagnostics for the *PIK3CA*-related megalencephaly-capillary malformation (MCAP) syndrome: a case report. Cold Spring Harb Mol Case Stud. 2022 Apr 28;8(3):a006188. doi: 10.1101/mcs.a006188. PMID: 35483878; PMCID: PMC9059787.

Pirozzi F, Berkseth M, Shear R, Gonzalez L, Timms AE, Sulc J, Pao E, Oyama N, Forzano F, Conti V, Guerrini R, Doherty ES, Saitta SC, Lockwood CM, Pritchard CC, Dobyns WB, **Novotny E**, Wright JNN, **Saneto RP**, Friedman S, Hauptman J, Ojemann J, Kapur RP, Mirzaa GM. Profiling PI3K-AKT-MTOR variants in focal brain malformations reveals new insights for diagnostic care. Brain. 2022 Apr 29;145(3):925-938. doi: 10.1093/brain/awab376. Erratum in: Brain. 2022 Oct 19;: PMID: 35355055; PMCID: PMC9630661.

Maio N, **Saneto RP**, Steet R, Sotero de Menezes MA, Skinner C, Rouault TA. Disruption of cellular iron homeostasis by *IREB2* missense variants causes severe neurodevelopmental delay, dystonia and seizures. Brain Commun. 2022 Apr 19;4(3):fcac102. doi: 10.1093/braincomms/fcac102. PMID: 35602653; PMCID: PMC9118103.

Prasad N, **Castillo-Pinto C**, Safadi AL, Osborne B. Neurological Considerations for the Care of Patients With Severe Obesity. Neurohospitalist. 2022 Apr;12(2):264-267. doi: 10.1177/19418744211040697. Epub 2021 Aug 25. PMID: 35419155; PMCID: PMC8995583.

Williams CN, Hall TA, Francoeur C, Kurz J, Rasmussen L, Hartman ME, O'meara AI, Ferguson NM, Fink EL, Walker T, Drury K, Carpenter JL, Erklauer J, Press C, **Wainwright MS**, Lovett M, Dapul H, Murphy S, Risen S, Guerriero RM, Woodruff A, Guilliams KP; PEDIATRIC NEUROCRITICAL CARE RESEARCH GROUP (PNCRG).. Continuing Care For Critically Ill Children Beyond Hospital Discharge: Current State of Follow-up. Hosp Pediatr. 2022 Apr 1;12(4):359-393. doi: 10.1542/hpeds.2021-006464. PMID: 35314865; PMCID: PMC9182716.

Vawter-Lee M, **Natarajan N**, Rang K, Horn PS, Pardo AC, Thomas CW. Topiramate Is Safe for Refractory Neonatal Seizures: A Multicenter Retrospective Cohort Study of Necrotizing Enterocolitis Risk. Pediatr Neurol. 2022 Apr;129:7-13. doi: 10.1016/j.pediatrneurol.2021.12.003. Epub 2021 Dec 30. PMID: 35131568.

Keene JC, **Morgan LA**, Abend NS, Bates SV, Bauer Huang SL, Chang T, Chu CJ, Glass HC, Massey SL, Ostrander B, Pardo AC, Press CA, Soul JS, Shellhaas RA, Thomas C, **Natarajan N**. Treatment of Neonatal Seizures: Comparison of Treatment Pathways From 11 Neonatal Intensive Care Units. Pediatr Neurol. 2022 Mar;128:67-74. doi: 10.1016/j.pediatrneurol.2021.10.004. Epub 2021 Oct 11. PMID: 34750046.

Keene JC, **Wainwright M**, **Morgan LA**, Mietzsch U, Musa N, **Bozarth XL**, **Natarajan N**. Retrospective Evaluation of First-line Levetiracetam use for Neonatal Seizures after Congenital Heart Defect repair with or without Extracorporeal Membrane Oxygenation. J Pediatr Pharmacol Ther. 2022;27(3):254-262. doi: 10.5863/1551-6776-27.3.254. Epub 2022 Mar 21. PMID: 35350164; PMCID: PMC8939278.

Fink EL, Robertson CL, **Wainwright MS**, Roa JD, Lovett ME, Stulce C, Yacoub M, Potera RM, Zivick E, Holloway A, Nagpal A, Wellnitz K, Czech T, Even KM, Brunow de Carvalho W, Rodriguez IS, Schwartz SP, Walker TC, Campos-Miño S, Dervan LA, Geneslaw AS, Sewell TB, Pryce P, Silver WG, Lin JE, Vargas WS, Topjian A, Alcamo AM, McGuire JL, Domínguez Rojas JA, Muñoz JT, Hong SJ, Muller WJ, Doerfler M, Williams CN, Drury K, Bhagat D, Nelson A, Price D, Dapul H, Santos L, Kahoud R, Francoeur C, Appavu B, Guilliams KP, Agner SC, Walson KH, Rasmussen L, Janas A, Ferrazzano P, Farias-Moeller R, Snooks KC, Chang CH, Yun J, Schober ME; Global Consortium Study of Neurologic Dysfunction in COVID-19 (GCS-NeuroCOVID) Investigators. Prevalence and Risk Factors of Neurologic Manifestations in Hospitalized Children Diagnosed with Acute SARS-CoV-2 or MIS-C. Pediatr Neurol. 2022 Mar;128:33-44. doi: 10.1016/j.pediatrneurol.2021.12.010. Epub 2021 Dec 28. PMID: 35066369; PMCID: PMC8713420.

Farias-Moeller R, Jayakar A, Guerriero RM, Carpenter JL, **Wainwright MS**, Harrar DB. Pediatric Critical Care Neurologists in the United States and Canada: A Survey of Clinical Practice Experience. J Child Neurol. 2022 Mar;37(4):288-297. doi: 10.1177/08830738211070099. Epub 2022 Jan 17. PMID: 35037772.

Ko PY, Khalatbari H, Hatt D, Coufal N, Barry D, **Wainwright MS**, Khanna PC, **Amlie-Lefond C.** Hemorrhagic Transformation Following Childhood Cardioembolic Stroke Is Not Increased in Anticoagulated Patients. J Child Neurol. 2022 Mar;37(4):273-280. doi: 10.1177/08830738211025866. Epub 2022 Jan 13. PMID: 35021919.

**Steinman KJ,** Stone WL, Ibañez LV, Attar SM. Reducing Barriers to Autism Screening in Community Primary Care: A Pragmatic Trial Using Web-Based Screening. Acad Pediatr. 2022 Mar;22(2):263-270. doi: 10.1016/j.acap.2021.04.017. Epub 2021 Apr 23. PMID: 33901728; PMCID: PMC8536796.

Price D, Dapul H, Santos L, Kahoud R, Francoeur C, Appavu B, Guilliams KP, Agner SC, Walson KH, Rasmussen L, Janas A, Ferrazzano P, Farias-Moeller R, Snooks KC, Chang CH, Yun J, Schober ME; Global Consortium Study of Neurologic Dysfunction in COVID-19 (GCS-NeuroCOVID) Investigators. Prevalence and Risk Factors of Neurologic Manifestations in Hospitalized Children Diagnosed with Acute SARS-CoV-2 or MIS-C. Pediatr Neurol. 2022 Mar;128:33-44. doi: 10.1016/j.pediatrneurol.2021.12.010. Epub 2021 Dec 28. PMID: 35066369; PMCID: PMC8713420.

Kuo CH, Casimo K, Wu J, Collins K, Rice P, Chen BW, Yang SH, Lo YC, **Novotny EJ,** Weaver KE, Chen YY, Ojemann JG. Electrocorticography to Investigate Age-Related Brain Lateralization on Pediatric Motor Inhibition. Front Neurol. 2022 Mar 7;13:747053. doi: 10.3389/fneur.2022.747053. PMID: 35330804; PMCID: PMC8940229.

Tunc EM, **Otten CE**, Kodish IM, Kim SJ, Kochar A, **Novotny E**, Yoshida H. Seventeen-Year-Old Female With History of Depression Presented With Mania and Enuresis. Pediatrics. 2022 Mar 1;149(3):e2021051176. doi: 10.1542/peds.2021-051176. PMID: 35229119.

Knupp KG, Coryell J, Singh RK, Gaillard WD, Shellhaas RA, Koh S, Mitchell WG, Harini C, Millichap JJ, May A, Dlugos D, Nickels K, Mytinger JR, Keator C, Yozawitz E, Singhal N, **Lockrow J**, Thomas JF, Juarez-Colunga E. Comparison of Cosyntropin, Vigabatrin, and Combination Therapy in New-Onset Infantile Spasms in a Prospective Randomized Trial. J Child Neurol. 2022 Mar;37(3):186-193. doi: 10.1177/08830738211073400. Epub 2022 Jan 19. PMID: 35044272.

Harrar DB, **Benedetti GM**, Jayakar A, Carpenter JL, Mangum TK, Chung M, Appavu B; International Pediatric Stroke Study Group and Pediatric Neurocritical Care Research Group. Pediatric Acute Stroke Protocols in the United States and Canada. J Pediatr. 2022 Mar;242:220-227.e7. doi: 10.1016/j.jpeds.2021.10.048. Epub 2021 Nov 11. PMID: 34774972

Bornstein R, James K, Stokes J, Park KY, Kayser EB, Snell J, Bard A, Chen Y, Kalume F, **Johnson SC**. Differential effects of mTOR inhibition and dietary ketosis in a mouse model of subacute necrotizing encephalomyelopathy. Neurobiol Dis. 2022 Feb;163:105594. doi: 10.1016/j.nbd.2021.105594. Epub 2021 Dec 20. PMID: 34933094; PMCID: PMC8770160

Conway KM, Gedlinske A, Mathews KD, **Perlman S**, Johnson N, Butterfield R, Hung M, Bounsanga J, Matthews D, Oleszek J, Romitti PA. A population-based study of scoliosis among males diagnosed with a dystrophinopathy identified by the Muscular Dystrophy Surveillance, Tracking, and Research Network (MD STARnet). Muscle Nerve. 2022 Feb;65(2):193-202. doi: 10.1002/mus.27464. Epub 2021 Dec 1. PMID: 34787322; PMCID: PMC8752499.

Shalabi H, Nellan A, Shah NN, **Gust J**. Immunotherapy Associated Neurotoxicity in Pediatric Oncology. Front Oncol. 2022 Feb 21;12:836452. doi: 10.3389/fonc.2022.836452. PMID: 35265526; PMCID: PMC8899040.

Madden LK, Rajajee V, Human T, **Wainwright MS**, Guanci M, Mainali S, Rowe S, McLaughlin D, Lunde J, Lele A, Fried H. Correction to: Neurocritical Care Society Guidelines Update: Lessons from a Decade of GRADE Guidelines. Neurocrit Care. 2022 Feb;36(1):333. doi: 10.1007/s12028-021-01411-0. Erratum for: Neurocrit Care. 2022 Feb;36(1):1-10. PMID: 34874505; PMCID: PMC9172880.

Jung S, Kayser EB, **Johnson SC**, Li L, Worstman HM, Sun GX, Sedensky MM, Morgan PG. Tetraethylammonium chloride reduces anaesthetic-induced neurotoxicity in Caenorhabditis elegans and mice. Br J Anaesth. 2022 Jan;128(1):77-88. doi: 10.1016/j.bja.2021.09.036. Epub 2021 Nov 30. PMID: 34857359; PMCID: PMC8787783

Sen K, Whitehead M, **Castillo Pinto C**, Caldovic L, Gropman A. Fifteen years of urea cycle disorders brain research: Looking back, looking forward. Anal Biochem. 2022 Jan 1;636:114343. doi: 10.1016/j.ab.2021.114343. Epub 2021 Oct 9. PMID: 34637785; PMCID: PMC8671367.

**Castillo-Pinto C**, Lamotte G, Mehta A, Sonti R, Di Maria G, Ruiz D, Kumar PN, Stemer AB, Denny MC. Healthcare Worker With Large Vessel Acute Ischemic Stroke Likely Related to Mild SARS-CoV-2 Infection. Neurohospitalist. 2022 Jan;12(1):48-56. doi: 10.1177/1941874420966845. Epub 2020 Nov 10. PMID: 34950386; PMCID: PMC8689530.

Kirschen MP, LaRovere K, Balakrishnan B, Erklauer J, Francoeur C, Ganesan SL, Jayakar A, Lovett M, Luchette M, Press CA, Wolf M, Ferrazzano P, **Wainwright MS**, Appavu B; Pediatric Neurocritical Care Research Group (PNCRG). A Survey of Neuromonitoring Practices in North American Pediatric Intensive Care Units. Pediatr Neurol. 2022 Jan;126:125-130. doi: 10.1016/j.pediatrneurol.2021.11.002. Epub 2021 Nov 12. PMID: 34864306; PMCID: PMC9135309.

Stokes JC, Bornstein RL, James K, Park KY, Spencer K, Vo K, Snell JC, Johnson BM, Morgan PG, Sedensky MM, Baertsch N, **Johnson SC**: Leukocytes mediate disease pathogenesis in the Ndufs4(KO) mouse model of Leigh syndrome. *JCI Insight*. Jan 20, 2022. PMID: 35050903

Keene JC, Woods B, **Wainwright M**, King M, **Morgan LA**. Optimized Benzodiazepine Treatment of Pediatric Status Epilepticus Through a Standardized Emergency Medical Services Resuscitation Tool. Pediatr Neurol. 2022 Jan;126:50-55. doi: 10.1016/j.pediatrneurol.2021.10.001. Epub 2021 Oct 9. PMID: 34736064.

Shurtleff HA, Poliakov A, Barry D, Wright JN, Warner MH, **Novotny EJ**, Marashly A, Buckley R, Goldstein HE, Hauptman JS, Ojemann JG, Shaw DWW. A clinically applicable functional MRI memory paradigm for use with pediatric patients. Epilepsy Behav. 2022 Jan;126:108461. doi: 10.1016/j.yebeh.2021.108461. Epub 2021 Dec 9. PMID: 34896785.

**Wainwright MS**, Guilliams K, Kannan S, Simon DW, Tasker RC, Traube C, Pineda J; Pediatric Organ Dysfunction Information Update Mandate (PODIUM) Collaborative. Acute Neurologic Dysfunction in Critically Ill Children: The PODIUM Consensus Conference. Pediatrics. 2022 Jan 1;149(1 Suppl 1):S32-S38. doi: 10.1542/peds.2021-052888E. PMID: 34970681

Bembea MM, Agus M, Akcan-Arikan A, Alexander P, Basu R, Bennett TD, Bohn D, Brandão LR, Brown AM, Carcillo JA, Checchia P, Cholette J, Cheifetz IM, Cornell T, Doctor A, Eckerle M, Erickson S, Farris RWD, Faustino EVS, Fitzgerald JC, Fuhrman DY, Giuliano JS, Guilliams K, Gaies M, Gorga SM, Hall M, Hanson SJ, Hartman M, Hassinger AB, Irving SY, Jeffries H, Jouvet P, Kannan S, Karam O, Khemani RG, Kissoon N, Lacroix J, Laussen P, Leclerc F, Lee JH, Leteurtre S, Lobner K, McKiernan PJ, Menon K, Monagle P, Muszynski JA, Odetola F, Parker R, Pathan N, Pierce RW, Pineda J, Prince JM, Robinson KA, Rowan CM, Ryerson LM, Sanchez-Pinto LN, Schlapbach LJ, Selewski DT, Shekerdemian LS, Simon D, Smith LS, Squires JE, Squires RH, Sutherland SM, Ouellette Y, Spaeder MC, Srinivasan V, Steiner ME, Tasker RC, Thiagarajan R, Thomas N, Tissieres P, Traube C, Tucci M, Typpo KV, **Wainwright MS**, Ward SL, Watson RS, Weiss S, Whitney J, Willson D, Wynn JL, Yehya N, Zimmerman JJ. Pediatric Organ Dysfunction Information Update Mandate (PODIUM) Contemporary Organ Dysfunction Criteria: Executive Summary. Pediatrics. 2022 Jan 1;149(1 Suppl 1):S1-S12. doi: 10.1542/peds.2021-052888B. PMID:

**Tully HM**, Doherty D, **Wainwright M**. Mortality in pediatric hydrocephalus. Dev Med Child Neurol. 2022 Jan;64(1):112-117. doi: 10.1111/dmcn.14975. Epub 2021 Jul 15. PMID: 34268734; PMCID : PMC8671148

**2021**

Faulhaber LD, Phuong AQ, Hartsuyker KJ, Cho Y, Mand KK, Harper SD, Olson AK, Garden GA, Shih AY, **Gust J**. Brain capillary obstruction during neurotoxicity in a mouse model of anti-CD19 chimeric antigen receptor T-cell therapy. Brain Commun. 2021 Dec 31;4(1):fcab309. doi: 10.1093/braincomms/fcab309. PMID: 35169706; PMCID: PMC8833245.

Bornstein R, James K, Stokes J, Park KY, Snell J, Pan A, Bard A, Kalume F, **Johnson SC**: Differential effects of mTOR inhibition and dietary ketosis in a mouse model of subacute necrotizing encephalomyelopathy. *Neurobiology of Disease.* Dec 20, 2021. PMID: 34933094

Shurtleff HA, Poliakov A, Barry D, Wright JN, Warner MH, **Novotny EJ, Marashly A**, Buckley R, Goldstein HE, Hauptman JS, Ojemann JG, Shaw A clinically applicable functional MRI memory paradigm for use with pediatric patients. DWW.Epilepsy Behav. 2021 Dec 9;126:108461. doi: 10.1016/j.yebeh.2021.108461. Online ahead of print.PMID: 34896785

**Wainwright MS.** Editorial: The path to resilience and recovery: understanding the epidemiology, neuropathology and treatment of neurologic injury due to the SARS-CoV-2 virus in children. Curr Opin Pediatr. 2021 Dec 1;33(6):576-579. doi: 10.1097/MOP.0000000000001079. PMID: 34654051; PMCID: PMC8577292.

Vasquez A, Farias-Moeller R, Sánchez-Fernández I, Abend NS, Amengual-Gual M, Anderson A, Arya R, Brenton JN, Carpenter JL, Chapman K, Clark J, Gaillard WD, Glauser T, Goldstein JL, Goodkin HP, Guerriero RM, Lai YC, McDonough TL, Mikati MA, **Morgan LA**, **Novotny EJ**, Ostendorf AP, Payne ET, Peariso K, Piantino J, Riviello JJ, Sands TT, Sannagowdara K, Tasker RC, Tchapyjnikov D, Topjian A, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; Pediatric Status Epilepticus Research Group (pSERG). Super-Refractory Status Epilepticus in Children: A Retrospective Cohort Study. Pediatr Crit Care Med. 2021 Dec 1;22(12):e613-e625

**Patel PB**, Bearden D. Neuropathogenesis of severe acute respiratory syndrome coronavirus 2. Curr Opin Pediatr. 2021 Dec 1;33(6):597-602. doi: 10.1097/MOP.0000000000001068. PMID: 34734914; PMCID: PMC8577294.

Madden LK, Rajajee V, Human T, **Wainwright MS**, Guanci M, Mainali S, Rowe S, McLaughlin D, Lunde J, Lele A, Fried H. Neurocritical Care Society Guidelines Update: Lessons from a Decade of GRADE Guidelines. Neurocrit Care. 2021 Nov 2:1–10. doi: 10.1007/s12028-021-01375-1. Epub ahead of print. Erratum in: Neurocrit Care. 2021 Dec 7;: PMID: 34729676; PMCID: PMC8562933

Sheehan T, Amengual-Gual M, Vasquez A, Abend NS, Anderson A, Appavu B, Arya R, Barcia Aguilar C, Brenton JN, Carpenter JL, Chapman KE, Clark J, Farias-Moeller R, Gaillard WD, Gaínza-Lein M, Glauser TA, Goldstein JL, Goodkin HP, Guerriero RM, Huh L, Jackson M, Kapur K, Kahoud R, Lai YC, McDonough TL, Mikati MA, **Morgan LA**, **Novotny EJ,** Ostendorf AP, Payne ET, Peariso K, Piantino J, Reece L, Riviello JJ, Sands TT, Sannagowdara K, Shellhaas R, Smith G, Tasker RC, Tchapyjnikov D, Topjian AA, **Wainwright MS**, Wilfong A, Williams K, Zhang B, Loddenkemper T; Pediatric Status Epilepticus Research Group. Benzodiazepine administration patterns before escalation to second-line medications in pediatric refractory convulsive status epilepticus. Epilepsia. 2021 Nov;62(11):2766-2777. doi: 10.1111/epi.17043. Epub 2021 Aug 21. PMID: 34418087.

Conway KM, Gedlinske A, Mathews KD, **Perlman S**, Johnson N, Butterfield R, Hung M, Bounsanga J, Matthews D, Oleszek J, Romitti PA. A population-based study of scoliosis among males diagnosed with a dystrophinopathy identified by the Muscular Dystrophy Surveillance, Tracking, and Research Network (MD STARnet). Muscle Nerve. 2021 Nov 17. doi: 10.1002/mus.27464. Epub ahead of print. PMID: 34787322.

Kirschen MP, LaRovere K, Balakrishnan B, Erklauer J, Francoeur C, Ganesan SL, Jayakar A, Lovett M, Luchette M, Press CA, Wolf M, Ferrazzano P, **Wainwright MS**, Appavu B; Pediatric Neurocritical Care Research Group (PNCRG). A Survey of Neuromonitoring Practices in North American Pediatric Intensive Care Units. Pediatr Neurol. 2021 Nov 12;126:125-130. doi: 10.1016/j.pediatrneurol.2021.11.002. Epub ahead of print. PMID: 34864306

Harrar DB, **Benedetti GM**, Jayakar A, Carpenter JL, Mangum TK, Chung M, Appavu B; Pediatric Acute Stroke Protocols in the United States and Canada. International Pediatric Stroke Study Group and Pediatric Neurocritical Care Research Group.J Pediatr. 2021 Nov 11:S0022-3476(21)01048-9. doi: 10.1016/j.jpeds.2021.10.048. Online ahead of print.PMID: 34774972

Stokes J, Freed A, Pan A, Sun G, Bornstein R, Snell J, Park KY, Morgan P, Sedensky MM, **Johnson SC**: Mechanisms underlying neonate specific metabolic effects of volatile anesthetics. *eLIFE*. Jul 12, 2021. PMID: 34254587.

Cheff DM, Muotri AR, Stockwell BR, Schmidt EE, Ran Q, Kartha RV, **Johnson SC**, Mittel P, Arner ESJ, Wigby KM, Hall MD, Rames SK: Development of Therapies for Rare Genetic Disorders of GPX4: Roadmap and Opportunities. *Orphanet Journal of Rare Diseases*. Epub 2021 Oct 23. PMID: 34688299.

Beatty CW, **Lockrow JP**, Gedela S, Gehred A, Ostendorf AP. The Missed Value of Underutilizing Pediatric Epilepsy Surgery: A Systematic Review. Semin Pediatr Neurol. 2021 Oct;39:100917. doi: 10.1016/j.spen.2021.100917. Epub 2021 Aug 19. PMID: 34620465.

Lee S, **Amlie-Lefond CM.** Target Practice: Aiming for Automated Perfusion in Childhood Stroke.

Stroke. 2021 Oct;52(10):3305-3307. doi: 10.1161/STROKEAHA.121.036218. Epub 2021 Aug 18.PMID: 34404239 No abstract available.

Schober ME, Robertson CL, **Wainwright MS**, Roa JD, Fink EL. COVID-19 and the Pediatric Nervous System: Global Collaboration to Meet a Global Need. Neurocrit Care. 2021 Oct;35(2):283-290. doi: 10.1007/s12028-021-01269-2. Epub 2021 Jun 28. PMID: 34184177; PMCID: PMC8238033.

**Keene JC**, **Morgan LA**, Abend NS, Bates SV, Bauer Huang SL, Chang T, Chu CJ, Glass HC, Massey SL, Ostrander B, Pardo AC, Press CA, Soul JS, Shellhaas RA, Thomas C, **Natarajan N.** Treatment of Neonatal Seizures: Comparison of Treatment Pathways From 11 Neonatal Intensive Care Units. Pediatr Neurol. 2021 Oct 11:S0887-8994(21)00222-8. doi: 10.1016/j.pediatrneurol.2021.10.004. Online ahead of print.PMID: 34750046

**Keene JC**, Woods B, **Wainwright M**, King M, **Morgan LA.** Optimized Benzodiazepine Treatment of Pediatric Status Epilepticus Through a Standardized Emergency Medical Services Resuscitation Tool Pediatr Neurol. 2021 Oct 9;126:50-55. doi: 10.1016/j.pediatrneurol.2021.10.001. Online ahead of print.PMID: 34736064

Johnson JO, Chia R, Miller DE, Li R, Kumaran R, Abramzon Y, Alahmady N, Renton AE, Topp SD, Gibbs JR, Cookson MR, Sabir MS, Dalgard CL, Troakes C, Jones AR, Shatunov A, Iacoangeli A, Al Khleifat A, Ticozzi N, Silani V, Gellera C, Blair IP, Dobson-Stone C, Kwok JB, Bonkowski ES, Palvadeau R, Tienari PJ, Morrison KE, Shaw PJ, Al-Chalabi A, Brown RH Jr, Calvo A, Mora G, Al-Saif H, Gotkine M, Leigh F, Chang IJ**, Perlman SJ,** Glass I, Scott AI, Shaw CE, Basak AN, Landers JE, Chiò A, Crawford TO, Smith BN, Traynor BJ; FALS Sequencing Consortium; American Genome Center; International ALS Genomics Consortium; and ITALSGEN Consortium, Smith BN, Ticozzi N, Fallini C, Gkazi AS, Topp SD, Scotter EL, Kenna KP, Keagle P, Tiloca C, Vance C, Troakes C, Colombrita C, King A, Pensato V, Castellotti B, Baas F, Ten Asbroek ALMA, McKenna-Yasek D, McLaughlin RL, Polak M, Asress S, Esteban-Pérez J, Stevic Z, D'Alfonso S, Mazzini L, Comi GP, Del Bo R, Ceroni M, Gagliardi S, Querin G, Bertolin C, van Rheenen W, Rademakers R, van Blitterswijk M, Lauria G, Duga S, Corti S, Cereda C, Corrado L, Sorarù G, Williams KL, Nicholson GA, Blair IP, Leblond-Manry C, Rouleau GA, Hardiman O, Morrison KE, Veldink JH, van den Berg LH, Al-Chalabi A, Pall H, Shaw PJ, Turner MR, Talbot K, Taroni F, García-Redondo A, Wu Z, Glass JD, Gellera C, Ratti A, Brown RH Jr, Silani V, Shaw CE, Landers JE, Dalgard CL, Adeleye A, Soltis AR, Alba C, Viollet C, Bacikova D, Hupalo DN, Sukumar G, Pollard HB, Wilkerson MD, Martinez EM, Abramzon Y, Ahmed S, Arepalli S, Baloh RH, Bowser R, Brady CB, Brice A, Broach J, Campbell RH, Camu W, Chia R, Cooper-Knock J, Ding J, Drepper C, Drory VE, Dunckley TL, Eicher JD, England BK, Faghri F, Feldman E, Floeter MK, Fratta P, Geiger JT, Gerhard G, Gibbs JR, Gibson SB, Glass JD, Hardy J, Harms MB, Heiman-Patterson TD, Hernandez DG, Jansson L, Kirby J, Kowall NW, Laaksovirta H, Landeck N, Landi F, Le Ber I, Lumbroso S, MacGowan DJL, Maragakis NJ, Mora G, Mouzat K, Murphy NA, Myllykangas L, Nalls MA, Orrell RW, Ostrow LW, Pamphlett R, Pickering-Brown S, Pioro EP, Pletnikova O, Pliner HA, Pulst SM, Ravits JM, Renton AE, Rivera A, Robberecht W, Rogaeva E, Rollinson S, Rothstein JD, Scholz SW, Sendtner M, Shaw PJ, Sidle KC, Simmons Z, Singleton AB, Smith N, Stone DJ, Tienari PJ, Troncoso JC, Valori M, Van Damme P, Van Deerlin VM, Van Den Bosch L, Zinman L, Landers JE, Chiò A, Traynor BJ, Angelocola SM, Ausiello FP, Barberis M, Bartolomei I, Battistini S, Bersano E, Bisogni G, Borghero G, Brunetti M, Cabona C, Calvo A, Canale F, Canosa A, Cantisani TA, Capasso M, Caponnetto C, Cardinali P, Carrera P, Casale F, Chiò A, Colletti T, Conforti FL, Conte A, Conti E, Corbo M, Cuccu S, Dalla Bella E, D'Errico E, DeMarco G, Dubbioso R, Ferrarese C, Ferraro PM, Filippi M, Fini N, Floris G, Fuda G, Gallone S, Gianferrari G, Giannini F, Grassano M, Greco L, Iazzolino B, Introna A, La Bella V, Lattante S, Lauria G, Liguori R, Logroscino G, Logullo FO, Lunetta C, Mandich P, Mandrioli J, Manera U, Manganelli F, Marangi G, Marinou K, Marrosu MG, Martinelli I, Messina S, Moglia C, Mora G, Mosca L, Murru MR, Origone P, Passaniti C, Petrelli C, Petrucci A, Pozzi S, Pugliatti M, Quattrini A, Ricci C, Riolo G, Riva N, Russo M, Sabatelli M, Salamone P, Salivetto M, Salvi F, Santarelli M, Sbaiz L, Sideri R, Simone I, Simonini C, Spataro R, Tanel R, Tedeschi G, Ticca A, Torriello A, Tranquilli S, Tremolizzo L, Trojsi F, Vasta R, Vacchiano V, Vita G, Volanti P, Zollino M, Zucchi E. Association of Variants in the SPTLC1 Gene With Juvenile Amyotrophic Lateral Sclerosis. JAMA Neurol. 2021 Oct 1;78(10):1236-1248. doi: 10.1001/jamaneurol.2021.2598. PMID: 34459874; PMCID: PMC8406220.

Prendergast E, Allen KY, Mills MG, Moran T, Harris ZL, Malakooti M, Smith CM, **Wainwright MS**, McCarthy-Kowols M. Targeted Temperature Management Protocol in a Pediatric Intensive Care Unit: A Quality Improvement Project. Crit Care Nurse. 2021 Oct 1;41(5):41-50. doi: 10.4037/ccn2021554. PMID: 34595494.

Gaínza-Lein M, Barcia Aguilar C, Piantino J, Chapman KE, Sánchez Fernández I, Amengual-Gual M, Anderson A, Appavu B, Arya R, Brenton JN, Carpenter JL, Clark J, Farias-Moeller R, Gaillard WD, Glauser TA, Goldstein JL, Goodkin HP, Huh L, Kahoud R, Kapur K, Lai YC, McDonough TL, Mikati MA, **Morgan LA**, Nayak A, **Novotny E Jr**, Ostendorf AP, Payne ET, Peariso K, Reece L, Riviello J, Sannagowdara K, Sands TT, Sheehan T, Tasker RC, Tchapyjnikov D, Vasquez A, **Wainwright MS**, Wilfong A, Williams K, Zhang B, Loddenkemper T; Factors associated with long-term outcomes in pediatric refractory status epilepticus.Pediatric Status Epilepticus Research Group.Epilepsia. 2021 Sep;62(9):2190-2204. doi: 10.1111/epi.16984. Epub 2021 Jul 12.PMID: 34251039

Park KB, Chapman T, Aldinger KA, Mirzaa GM, Zeiger J, Beck A, Glass IA, Hevner RF, Jansen AC, Marshall DA, Oegema R, Parrini E, **Saneto RP**, Curry CJ, Hall JG, Guerrini R, Leventer RJ, Dobyns WB. The spectrum of brain malformations and disruptions in twins. Am J Med Genet A. 2021 Sep;185(9):2690-2718. doi: 10.1002/ajmg.a.61972. Epub 2020 Nov 18. PMID: 33205886; PMCID: PMC8683564

Vitanza NA, Johnson AJ, Wilson AL, Brown C, Yokoyama JK, Künkele A, Chang CA, Rawlings-Rhea S, Huang W, Seidel K, Albert CM, Pinto N, **Gust J**, Finn LS, Ojemann JG, Wright J, Orentas RJ, Baldwin M, Gardner RA, Jensen MC, Park JR. Locoregional infusion of HER2-specific CAR T cells in children and young adults with recurrent or refractory CNS tumors: an interim analysis. Nat Med. 2021 Sep;27(9):1544-1552. doi: 10.1038/s41591-021-01404-8. Epub 2021 Jul 12.PMID: 34253928 Clinical Trial.

Shurtleff HA, Roberts EA, Young CC, Barry D, Warner MH, **Saneto RP**, Buckley R, Firman T, Poliakov AV, Ellenbogen RG, Hauptman JS, Ojemann JG, **Marashly A**. Pediatric hemispherectomy outcome: Adaptive functioning, intelligence, and memory. Epilepsy Behav. 2021 Sep 16;124:108298. doi: 10.1016/j.yebeh.2021.108298. Epub ahead of print. PMID: 34537627.

Jung S, Kayser EB, **Johnson SC**, Li L, Worstman H, Sun G, Sedensky MM, Morgan PG: Tetraethylammonium chloride reduces anaesthetic-induced neurotoxicity in *Caenorhabditis elegans* and mice. *British Journal of Anaesthesia*. https://doi.org/10.1016/j.bja.2021.09.036, PMID: 34857359.

Miller DE, Sulovari A, Wang T, Loucks H, Hoekzema K, Munson KM, Lewis AP, Fuerte EPA, Paschal CR, Walsh T, Thies J, Bennett JT, Glass I, Dipple KM, Patterson K, Bonkowski ES, Nelson Z, Squire A, Sikes M, Beckman E, Bennett RL, Earl D, Lee W, Allikmets R, **Perlman SJ**, Chow P, Hing AV, Wenger TL, Adam MP, Sun A, Lam C, Chang I, Zou X, Austin SL, Huggins E, Safi A, Iyengar AK, Reddy TE, Majoros WH, Allen AS, Crawford GE, Kishnani PS; University of Washington Center for Mendelian Genomics, King MC, Cherry T, Chong JX, Bamshad MJ, Nickerson DA, Mefford HC, Doherty D, Eichler EE. Targeted long-read sequencing identifies missing disease-causing variation. Am J Hum Genet. 2021 Aug 5;108(8):1436-1449. doi: 10.1016/j.ajhg.2021.06.006. Epub 2021 Jul 2. PMID: 34216551; PMCID: PMC8387463.

Zhang J, Tang W, Bhatia NK, Xu Y, Paudyal N, Liu D, Kim S, Song R, XiangWei W, Shaulsky G, Myers SJ, Dobyns W, Jayaraman V, Traynelis SF, Yuan H, **Bozarth X.** A de novo GRIN1 Variant Associated With Myoclonus and Developmental Delay: From Molecular Mechanism to Rescue Pharmacology. Front Genet. 2021 Aug 3;12:694312. doi: 10.3389/fgene.2021.694312. eCollection 2021.PMID: 34413877 Free PMC article**.**

Engelke UF, van Outersterp RE, Merx J, van Geenen FA, van Rooij A, Berden G, Huigen MC, Kluijtmans LA, Peters TM, Al-Shekaili HH, Leavitt BR, de Vrieze E, Broekman S, van Wijk E, Tseng LA, Kulkarni P, Rutjes FP, Mecinović J, Struys EA, Jansen LA, **Gospe SM Jr**, Mercimek-Andrews S, Hyland K, Willemsen MA, Bok LA, van Karnebeek CD, Wevers RA, Boltje TJ, Oomens J, Martens J, Coene KL. Untargeted metabolomics and infrared ion spectroscopy identify biomarkers for pyridoxine-dependent epilepsy J Clin Invest. 2021 Aug 2;131(15):e148272. doi: 10.1172/JCI148272.PMID: 34138754 Free PMC article**.**

# Jennifer I Koop, Kevin Credille, Yang Wang, Michelle Loman, Ahmad Marashly, Irene Kim, Sean M Lew, Mohit MaheshwariDetermination of language dominance in pediatric patients with epilepsy for clinical decision-making: Correspondence of intracarotid amobarbitol procedure and fMRI modalities

Epilepsy Behav . 2021 Aug;121(Pt A):108041.

**Hannah M Tully**, Dan Doherty, **Mark Wainwright**. Mortality in pediatric hydrocephalus Dev Med Child Neurol. 2021 Jul 15.  doi: 10.1111/dmcn.14975. Online ahead of print.

Stokes J, Freed A, Pan A, Sun G, Bornstein R, Snell J, Park KY, Morgan P, Sedensky MM, **Johnson SC**: Mechanisms underlying neonate specific metabolic effects of volatile anesthetics. *eLIFE*. Jul 12, 2021. PMID: 34254587

**Gospe SM Jr.** In: Adam MP, Ardinger HH, Pagon RA, Wallace SE, Bean LJH, Mirzaa G, Amemiya A, editors. Pyridoxine-Dependent Epilepsy –  ALDH7A1 GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993–2021. 2001 Dec 7 [updated 2021 Jul 29].

Sculier C, Barcia Aguilar C, Gaspard N, Gaínza-Lein M, Sánchez Fernández I, Amengual-Gual M, Anderson A, Arya R, Burrows BT, Brenton JN, Carpenter JL, Chapman KE, Clark J, Gaillard WD, Glauser TA, Goldstein JL, Goodkin HP, Gorman M, Lai YC, McDonough TL, Mikati MA, Nayak A, Peariso K, Riviello J, Rusie A, Sperberg K, Stredny CM, Tasker RC, Tchapyjnikov D, Vasquez A, **Wainwright MS**, Wilfong AA, Williams K, Loddenkemper T; Clinical presentation of new onset refractory status epilepticus in children (the pSERG cohort). pSERG.Epilepsia. 2021 Jul;62(7):1629-1642. doi: 10.1111/epi.16950. Epub 2021 Jun 6.

Barcia Aguilar C, Amengual-Gual M, Sánchez Fernández I, Abend NS, Anderson A, Appavu B, Arya R, Brenton JN, Carpenter JL, Chapman KE, Clark J, Farias-Moeller R, Gaillard WD, Gaínza-Lein M, Glauser T, Goldstein JL, Goodkin HP, Guerriero RM, Huh L, Lai YC, McDonough TL, Mikati MA, **Morgan LA,** **Novotny EJ,** Ostendorf A, Payne ET, Peariso K, Piantino J, Riviello J, Sannagowdara K, Sheehan T, Sands TT, Tasker RC, Tchapyjnikov D, Topjian AA, Vasquez A, **Wainwright MS**, Wilfong AA, Williams K, Loddenkemper T; pSERG. Time to Treatment in Pediatric Convulsive Refractory Status Epilepticus: The Weekend Effect. Pediatr Neurol. 2021 Jul;120:71-79. doi: 10.1016/j.pediatrneurol.2021.03.009. Epub 2021 Mar 26

**Saneto RP**, **Patrick KE**, Perez FA. Homoplasmy of the m. 8993 T>G variant in a patient without MRI findings of Leigh syndrome, ataxia or retinal abnormalities Mitochondrion. 2021 Jul;59:58-62. doi: 10.1016/j.mito.2021.04.010. Epub 2021 Apr 22.

Tarazi RA, **Patrick KE**, Lampietro M, Apollonsky N. Hydroxyurea Use Associated with Nonverbal and Executive Skills in Sickle Cell Anemia.J Pediatr Psychol. 2021 Jul 20;46(6):710-718. doi: 10.1093/jpepsy/jsab015

Vitanza NA, Johnson AJ, Wilson AL, Brown C, Yokoyama JK, Künkele A, Chang CA, Rawlings-Rhea S, Huang W, Seidel K, Albert CM, Pinto N, **Gust J**, Finn LS, Ojemann JG, Wright J, Orentas RJ, Baldwin M, Gardner RA, Jensen MC, Park Locoregional infusion of HER2-specific CAR T cells in children and young adults with recurrent or refractory CNS tumors: an interim analysis. J R.Nat Med. 2021 Jul 12. doi: 10.1038/s41591-021-01404-8. Online ahead of print.

Gaínza-Lein M, Barcia Aguilar C, Piantino J, Chapman KE, Sánchez Fernández I, Amengual-Gual M, Anderson A, Appavu B, Arya R, Brenton JN, Carpenter JL, Clark J, Farias-Moeller R, Gaillard WD, Glauser TA, Goldstein JL, Goodkin HP, Huh L, Kahoud R, Kapur K, Lai YC, McDonough TL, Mikati MA, **Morgan LA**, Nayak A, **Novotny E Jr**, Ostendorf AP, Payne ET, Peariso K, Reece L, Riviello J, Sannagowdara K, Sands TT, Sheehan T, Tasker RC, Tchapyjnikov D, Vasquez A, **Wainwright MS**, Wilfong A, Williams K, Zhang B, Loddenkemper T; Factors associated with long-term outcomes in pediatric refractory status epilepticus. Pediatric Status Epilepticus Research Group.Epilepsia. 2021 Jul 12. doi: 10.1111/epi.16984. Online ahead of print.

Oesch G, Perez FA, **Wainwright MS**, Shaw DWW, **Amlie-Lefond C.** Focal Cerebral Arteriopathy of Childhood: Clinical and Imaging Correlates. Stroke. 2021 Jul;52(7):2258-2265. doi: 10.1161/STROKEAHA.120.031880. Epub 2021 May 27.

Davies OMT, Garzon MC, Frieden IJ, Cottrell CE, Gripp KW, **Saneto RP**, Shwayder T, Mirzaa GM, Drolet BA.J Cutaneous vascular anomalies associated with a mosaic variant of AKT3: Genetic analysis continues to refine the diagnosis, nomenclature, and classification of vascular anomalies. Am Acad Dermatol. 2021 Jul 6:S0190-9622(21)02067-3. doi: 10.1016/j.jaad.2021.06.877. Online ahead of print.

Schober ME, Robertson CL, **Wainwright MS**, Roa JD, Fink EL COVID-19 and the Pediatric Nervous System: Global Collaboration to Meet a Global Need. .Neurocrit Care. 2021 Jun 28:1-8. doi: 10.1007/s12028-021-01269-2. Online ahead of print.

O'Brien NF, Reuter-Rice K, **Wainwright MS**, Kaplan SL, Appavu B, Erklauer JC, Ghosh S, Kirschen M, Kozak B, Lidsky K, Lovett ME, Mehollin-Ray AR, Miles DK, Press CA, Simon DW, Tasker RC, LaRovere KL Practice Recommendations for Transcranial Doppler Ultrasonography in Critically Ill Children in the Pediatric Intensive Care Unit: A Multidisciplinary Expert Consensus Statement. J Pediatr Intensive Care. 2021 Jun;10(2):133-142.

Vasquez A, Farias-Moeller R, Sánchez-Fernández I, Abend NS, Amengual-Gual M, Anderson A, Arya R, Brenton JN, Carpenter JL, Chapman K, Clark J, Gaillard WD, Glauser T, Goldstein JL, Goodkin HP, Guerriero RM, Lai YC, McDonough TL, Mikati MA, **Morgan LA, Novotny EJ,** Ostendorf AP, Payne ET, Peariso K, Piantino J, Riviello JJ, Sands TT, Sannagowdara K, Tasker RC, Tchapyjnikov D, Topjian A, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; Pediatric Status Epilepticus Research Group (pSERG) Super-Refractory Status Epilepticus in Children: A Retrospective Cohort Study. Pediatr Crit Care Med. 2021 Jun 14.

Erickson SL, Killien EY, **Wainwright M**, Mills B, Vavilala MS. Mean Arterial Pressure and Discharge Outcomes in Severe Pediatric Traumatic Brain Injury. Neurocrit Care. 2021 Jun;34(3):1017-1025. doi: 10.1007/s12028-020-01121-z. Epub 2020 Oct 27. PMID: 33108627; PMCID: PMC8076334.

Paczesny S, Pasquini MC, Pavletic SZ, Agarwal A, Spellman S, Kean L, Bernatchez C, **Gust J**, Staedtke V, Perales MA Blueprint for the discovery of biomarkers of toxicity and efficacy for CAR T cells and T-cell engagers. .Blood Adv. 2021 Jun 8;5(11):2519-2522

Shalabi H, **Gust J**, Taraseviciute A, Wolters PL, Leahy AB, Sandi C, Laetsch TW, Wiener L, Gardner RA, Nussenblatt V, Hill JA, Curran KJ, Olson TS, Annesley C, Wang HW, Khan J, Pasquini MC, Duncan CN, Grupp SA, Pulsipher MA, Shah NN. Beyond the storm - subacute toxicities and late effects in children receiving CAR T cells.Nat Rev Clin Oncol. 2021 Jun;18(6):363-378.

Ferrazzano P, Yeske B, Mumford J, Kirk G, Bigler ED, **Bowen K**, O'Brien N, Rosario B, Beers SR, Rathouz P, Bell MJ, Alexander AL. Brain Magnetic Resonance Imaging Volumetric Measures of Functional Outcome after Severe Traumatic Brain Injury in Adolescents. J Neurotrauma. 2021 Jun 1;38(13):1799-1808. doi: 10.1089/neu.2019.6918. Epub 2021 Feb 24

Acsadi G, Crawford TO, Müller-Felber W, Shieh PB, Richardson R, **Natarajan N**, Castro D, Ramirez-Schrempp D, Gambino G, Sun P, Farwell W. Safety and efficacy of nusinersen in spinal muscular atrophy: The EMBRACE study. Muscle Nerve. 2021 May;63(5):668-677. doi: 10.1002/mus.27187. Epub 2021 Feb 16.

Kelly JP, Phillips JO, **Saneto RP**, Khalatbari H, Poliakov A, Tarczy-Hornoch K, Weiss AH. Cerebral Visual Impairment Characterized by Abnormal Visual Orienting Behavior With Preserved Visual Cortical Activation. Invest Ophthalmol Vis Sci. 2021 May 3;62(6):15.

Khalatbari H, Wright JN, Ishak GE, Perez FA, **Amlie-Lefond CM**, Shaw DWW. Deep medullary vein engorgement and superficial medullary vein engorgement: two patterns of perinatal venous stroke. Pediatr Radiol. 2021 May;51(5):675-685.

Ferdinandusse S, McWalter K, Te Brinke H, IJlst L, Mooijer PM, Ruiter JPN, van Lint AEM, Pras-Raves M, Wever E, Millan F, Guillen Sacoto MJ, Begtrup A, Tarnopolsky M, Brady L, Ladda RL, Sell SL, Nowak CB, Douglas J, Tian C, Ulm E, **Perlman S**, Drack AV, Chong K, Martin N, Brault J, Brokamp E, Toro C, Gahl WA, Macnamara EF, Wolfe L; Undiagnosed Diseases Network, Waisfisz Q, Zwijnenburg PJG, Ziegler A, Barth M, Smith R, Ellingwood S, Gaebler-Spira D, Bakhtiari S, Kruer MC, van Kampen AHC, Wanders RJA, Waterham HR, Cassiman D, Vaz FM. An autosomal dominant neurological disorder caused by de novo variants in FAR1 resulting in uncontrolled synthesis of ether lipids. Genet Med. 2021 Apr;23(4):740-750. doi: 10.1038/s41436-020-01027-3. Epub 2020 Nov 26.

**Saneto RP**, **Patrick KE**, Perez FA Homoplasmy of the m. 8993 T>G variant in a patient without MRI findings of Leigh syndrome, ataxia or retinal abnormalities. Mitochondrion. 2021 Apr 22;59:58-62.

Barry M, Barry D, Kansagra AP, Hallam D, Abraham M, **Amlie-Lefond C**; Thrombolysis in Pediatric Stroke (TIPSTER) Investigators\*. Higher-Quality Data Collection Is Critical to Establish the Safety and Efficacy of Pediatric Mechanical Thrombectomy. Stroke. 2021 Apr;52(4):1213-1221.

Lauzier DC, Galardi MM, Guilliams KP, Goyal MS, **Amlie-Lefond C**, Hallam DK, Kansagra AP. Pediatric Thrombectomy: Design and Workflow Lessons From Two Experienced Centers. Stroke. 2021 Apr;52(4):1511-1519.

**Steinman KJ**, Stone WL, Ibañez LV, Attar SM. Reducing Barriers to Autism Screening in Community Primary Care: A Pragmatic Trial using Web-Based Screening. *Acad Pediatr*. 2021 Apr 23:S1876-2859(21)00237-0.

**Novotny EJ** Diverse genetic causes of polymicrogyria with epilepsy. Epilepsy Phenome/Genome Project, Epi4K Consortium. Epilepsia. 2021 Apr;62(4):973-983. doi: 10.1111/epi.16854. Epub 2021 Apr 5.

Simpson TS, Peterson RL, **Patrick KE**, Forster JE, McNally KA. Concussion Symptom Treatment and Education Program: A Feasibility Study J Head Trauma Rehabil. 2021 Mar-Apr 01;36(2):E79-E88

Weiss KE, **Steinman KJ**, Kodish I, Sim L, Yurs S, Steggall C, Fobian AD. Functional Neurological Symptom Disorder in Children and Adolescents within Medical Settings. J Clin Psychol Med Settings. 2021 Mar;28(1):90-101. doi: 10.1007/s10880-020-09736-2.

Barcia Aguilar C, Amengual-Gual M, Sánchez Fernández I, Abend NS, Anderson A, Appavu B, Arya R, Brenton JN, Carpenter JL, Chapman KE, Clark J, Farias-Moeller R, Gaillard WD, Gaínza-Lein M, Glauser T, Goldstein JL, Goodkin HP, Guerriero RM, Huh L, Lai YC, McDonough TL, Mikati MA**, Morgan LA**, **Novotny EJ**, Ostendorf A, Payne ET, Peariso K, Piantino J, Riviello J, Sannagowdara K, Sheehan T, Sands TT, Tasker RC, Tchapyjnikov D, Topjian AA, Vasquez A, **Wainwright MS**, Wilfong AA, Williams K, Loddenkemper T; pSERG Time to Treatment in Pediatric Convulsive Refractory Status Epilepticus: The Weekend Effect. Pediatr Neurol. 2021 Mar 26;120:71-79

**Mingbunjerdsuk D**, **Blume H**, Browd S, Samii A Intraventricular Baclofen Following Deep Brain Stimulation in a Child with Refractory Status Dystonicus. .Mov Disord Clin Pract. 2021 Mar 19;8(3):456-459

Madurski C, Jarvis JM, Beers SR, Houtrow AJ, Wagner AK, Fabio A, Wang C, Smith CM, Doughty L, Janesko-Feldman K, Rubin P, Pollon D, Treble-Barna A, Kochanek PM, Fink EL; PICU-Rehabilitation Study Group (**Mark Wainwright**). Serum Biomarkers of Regeneration and Plasticity are Associated with Functional Outcome in Pediatric Neurocritical Illness: An Exploratory Study Neurocrit Care. 2021 Mar 4.

Kamins J, Richards R, Barney BJ, Locandro C, Pacchia CF, Charles AC, Cook LJ, Gioia G, Giza CC, **Blume HK.** Evaluation of Posttraumatic Headache Phenotype and Recovery Time After Youth Concussion. JAMA Netw Open. 2021 Mar 1;4(3):e211312. doi: 10.1001/jamanetworkopen.2021.1312.

**Gust J**, Annesley CE, Gardner RA, **Bozarth X**. EEG Correlates of Delirium in Children and Young Adults With CD19-Directed CAR T Cell Treatment-Related Neurotoxicity. J Clin Neurophysiol. 2021 Mar 1;38(2):135-142.

# Payal B Patel, Andrew Belden Ryan Handoko, Thanyawee Puthanakit, Stephen Kerr, Pope Kosalaraksa, Pradthana Ounchanum, Suparat Kanjanavanit, Linda Aurpibul Chaiwat Ngampiyasakul Wicharn Luesomboon, Claude A Mellins Kathleen Malee, Jintanat Ananworanich, Robert PaulBehavioral impairment and cognition in Thai adolescents affected by HIV. Glob Ment Health (Camb). 2021 Feb 9;8:e3.  doi: 10.1017/gmh.2021.1. eCollection 2021.

Harini Sarva, Gustavo Patino, Mehmood Rashid, **James WM Owens**, Matthew Robbins, and Stefano Sandrone (2021) The status of neurology fellowships in the United States: clinical needs, educational barriers, and future outlooks. BMC Medical Education. 2021 Feb 17: 21(1)108-117.

**Mingbunjerdsuk D**, Wong M, **Bozarth X**, Sun A. Co-occurrence of Metachromatic Leukodystrophy in Phelan-McDermid Syndrome. J Child Neurol. 2021 Feb;36(2):148-151.

Peter Ferrazzano, Benjamin Yeske, Jeanette Mumford, Gregory Kirk, Erin D Bigler**, Katherine Bowen,**Nicole O'Brien, Bedda Rosario, Sue R Beers, Paul Rathouz, Michael J Bell, Andrew L Alexander, and the ADAPT MRI Biomarkers Investigators. Brain Magnetic Resonance Imaging Volumetric Measures of Functional Outcome after Severe Traumatic Brain Injury in Adolescents**.** J Neurotrauma 2021 Feb 24

**Patel PB**, Belden A, Handoko R, Puthanakit T, Kerr S, Kosalaraksa P, Ounchanum P, Kanjanavanit S, Aurpibul L, Ngampiyasakul C, Luesomboon W, Mellins CA, Malee K, Ananworanich J, Paul R. Behavioral impairment and cognition in Thai adolescents affected by HIV. Glob Ment Health (Camb). 2021 Feb 9;8:e3. doi: 10.1017/gmh.2021.1. PMID: 34026234; PMCID: PMC8127634.

**Amlie-Lefond C**, **Wainwright MS.** Childhood Stroke: Thinking Locally, Acting Globally?

Stroke. 2021 Jan;52(1):162-163.

Murphy OC, Messacar K, Benson L, Bove R, Carpenter JL, Crawford T, Dean J, DeBiasi R, Desai J, Elrick MJ, Farias-Moeller R, Gombolay GY, Greenberg B, Harmelink M, Hong S, Hopkins SE, Oleszek J, **Otten C**, Sadowsky CL, Schreiner TL, Thakur KT, Van Haren K, Carballo CM, Chong PF, Fall A, Gowda VK, Helfferich J, Kira R, Lim M, Lopez EL, Wells EM, Yeh EA, Pardo CA; AFM working group. Acute flaccid myelitis: cause, diagnosis, and management Lancet. 2021 Jan 23;397(10271):334-346.

Acsadi G, Crawford TO, Müller-Felber W, Shieh PB, Richardson R, **Natarajan N**, Castro D, Ramirez-Schrempp D, Gambino G, Sun P, Farwell W. [Safety and efficacy of nusinersen in spinal muscular atrophy: the EMBRACE study.](https://pubmed.ncbi.nlm.nih.gov/33501671/) Muscle Nerve. 2021 Jan 26.

Crow YJ, Marshall H, Rice GI, Seabra L, Jenkinson EM, Baranano K, Battini R, Berger A, Blair E, Blauwblomme T, Bolduc F, Boddaert N, Buckard J, Burnett H, Calvert S, Caumes R, Ng AC, Chiang D, Clifford DB, Cordelli DM, de Burca A, Demic N, Desguerre I, De Waele L, Di Fonzo A, Dunham SR, Dyack S, Elmslie F, Ferrand M, Fisher G, Karimiani EG, Ghoumid J, Gibbon F, Goel H, Hilmarsen HT, Hughes I, Jacob A, Jones EA, Kumar R, Leventer RJ, MacDonald S, Maroofian R, Mehta SG, Metz I, Monfrini E, Neumann D, Noetzel M, O'Driscoll M, Õunap K, Panzer A, Parikh S, Prabhakar P, Ramond F, Sandford R, **Saneto R**, Soh C, Stutterd CA, Subramanian GM, Talbot K, Thomas RH, Toro C, Touraine R, Wakeling E, Wassmer E, Whitney A, Livingston JH, O'Keefe RT, Badrock AP. Leukoencephalopathy with calcifications and cysts: Genetic and phenotypic spectrum. Am J Med Genet A. 2021 Jan;185(1):15-25. doi: 10.1002/ajmg.a.61907. Epub 2020 Oct 7.

Malbari F, Partap S, **Gust J**, Duke E, **Sato A**, Khakoo Y, Ullrich NJ. Neuro-Oncology Training for the Child Neurology Resident.J Child Neurol. 2021 Jan;36(1):79-82

Weaver JJ, Hallam DK, Chick JFB, Vaidya S, Shin DS, **Natarajan N**, Rad N, Reis J, Koo KSH, Shivaram GM, Thibodeau A, Apkon S, Monroe EJ. [Transforaminal intrathecal delivery of nusinersen for older children and adults with spinal muscular atrophy and complex spinal anatomy: an analysis of 200 consecutive injections.](https://pubmed.ncbi.nlm.nih.gov/32471828/) J Neurointerv Surg. 2021 Jan;13(1):75-78.

Grinspan ZM, Patel AD, Shellhaas RA, Berg AT, Axeen ET, Bolton J, Clarke DF, Coryell J, Gaillard WD, Goodkin HP, Koh S, Kukla A, Mbwana JS, **Morgan LA**, Singhal NS, Storey MM, Yozawitz EG, Abend NS, Fitzgerald MP, Fridinger SE, Helbig I, Massey SL, Prelack MS, Buchhalter J; Pediatric Epilepsy Learning Healthcare System. [Design and implementation of electronic health record common data elements for pediatric epilepsy: Foundations for a learning health care system.](https://pubmed.ncbi.nlm.nih.gov/33368200/)  Epilepsia. 2021 Jan;62(1):198-216.

Ko PY, Glass IA, Crandall S, Weiss A, Dorschner MO, Kelly JP, Phillips JO, **Lopez J.** Two Missense CACNA1A Variants in a Single Family with Variable Neurobehavioral, Cerebellar, Epileptic, and Oculomotor Features. Neuropediatrics. 2021 Jan 14.

Coughlin CR 2nd, Tseng LA, Abdenur JE, Ashmore C, Boemer F, Bok LA, Boyer M, Buhas D, Clayton PT, Das A, Dekker H, Evangeliou A, Feillet F, Footitt EJ, **Gospe SM Jr**, Hartmann H, Kara M, Kristensen E, Lee J, Lilje R, Longo N, Lunsing RJ, Mills P, Papadopoulou MT, Pearl PL, Piazzon F, Plecko B, Saini AG, Santra S, Sjarif DR, Stockler-Ipsiroglu S, Striano P, Van Hove JLK, Verhoeven-Duif NM, Wijburg FA, Zuberi SM, van Karnebeek CDM. Consensus guidelines for the diagnosis and management of pyridoxine-dependent epilepsy due to α-aminoadipic semialdehyde dehydrogenase deficiency. J Inherit Metab Dis. 2021 Jan;44(1):178-192.

Dixon SM, Binkley MM, **Gospe SM Jr**, Guerriero RM. Child Neurology Applicants Place Increasing Emphasis on Quality of Life Factors. Pediatr Neurol. 2021 Jan;114:42-46.

**2020**

Lai YC, Muscal E, Wells E, Shukla N, Eschbach K, Hyeong Lee K, Kaliakatsos M, Desai N, Wickström R, Viri M, Freri E, Granata T, Nangia S, Dilena R, Brunklaus A, **Wainwright MS**, Gorman MP, Stredny CM, Asiri A, Hundallah K, Doja A, Payne E, Wirrell E, Koh S, Carpenter JL, Riviello Anakinra usage in febrile infection related epilepsy syndrome: an international cohort.J. Ann Clin Transl Neurol. 2020 Dec;7(12):2467-2474

**Cronin M**, **Wainwright MS.** Plasma Exchange for Treatment of Refractory Demyelination. Pediatr Neurol Briefs. 2020 Dec 9;34:16.

Bass DI, **Shurtleff H**, **Warner M**, Knott D, Poliakov A, Friedman S, Collins MJ, **Lopez J**, **Lockrow JP**, **Novotny EJ**, Ojemann JG, Hauptman JS. Awake Mapping of the Auditory Cortex during Tumor Resection in an Aspiring Musical Performer: A Case Report. Pediatr Neurosurg. 2020;Dec 1 155(6):351-358

**Gust J**, Ponce R, Liles WC, Garden GA, Turtle CJ. [Cytokines in CAR T Cell-Associated Neurotoxicity.](https://pubmed.ncbi.nlm.nih.gov/33391257/)

Front Immunol. 2020 Dec 16;11:577027.

Weisleder P, **Gospe SM Jr** [Withholding Childhood Immunizations: A Parent's Right or a Child's Neglect?](https://pubmed.ncbi.nlm.nih.gov/33190688/)

Pediatr Neurol. 2020 Dec;113:80-81.

Oesch G, **Bozarth XL.** [Rufinamide efficacy and association with phenotype and genotype in children with intractable epilepsy: A retrospective single center study.](https://pubmed.ncbi.nlm.nih.gov/31575436/) Epilepsy Res. 2020 Dec;168:106211.

Vasquez A, Gaínza-Lein M, Abend NS, Amengual-Gual M, Anderson A, Arya R, Brenton JN, Carpenter JL, Chapman K, Clark J, Farias-Moeller R, Gaillard WD, Glauser T, Goldstein JL, Goodkin HP, Guerriero RM, Kapur K, Lai YC, McDonough TL, Mikati MA, **Morgan LA,** **Novotny EJ**, Ostendorf AP, Payne ET, Peariso K, Piantino J, Riviello JJ, Sannagowdara K, Tasker RC, Tchapyjnikov D, Topjian A, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; Pediatric Status Epilepticus Research Group (pSERG). First-line medication dosing in pediatric refractory status epilepticus. Neurology. 2020 Nov 10;95(19):e2683-e2696

Park KB, Chapman T, Aldinger KA, Mirzaa GM, Zeiger J, Beck A, Glass IA, Hevner RF, Jansen AC, Marshall DA, Oegema R, Parrini E, **Saneto RP**, Curry CJ, Hall JG, Guerrini R, Leventer RJ, Dobyns WB. The spectrum of brain malformations and disruptions in twins. Am J Med Genet A. 2020 Nov 18. Online ahead of print. PMID: 33205886

Powell WT, Schaaf CP, Rech ME, **Wrede J.** Polysomnographic characteristics and sleep-disordered breathing in Schaaf-Yang syndrome. Pediatr Pulmonol. 2020 Nov;55(11):3162-3167.

Jeon TY, Poliakov AV, Friedman SD, **Bozarth XL**, **Novotny EJ**, Hauptman JS, Moon SH, Shaw DWW. Structural MRI and tract-based spatial statistical analysis of diffusion tensor imaging in children with hemimegalencephaly Neuroradiology. 2020 Nov;62(11):1467-1474.

Lewis H, Samanta D, Örsell JL, Bosanko KA, Rowell A, Jones M, Dale RC, Taravath S, Hahn CD, Krishnakumar D, Chagnon S, Keller S, Hagebeuk E, Pathak S, Bebin EM, Arndt DH, **Alexander JJ**, Mainali G, Coppola G, Maclean J, Sparagana S, McNamara N, Smith DM, Raggio V, Cruz M, Fernández-Jaén A, Kava MP, Emrick L, Fish JL, Vanderver A, Helman G, Pierson TM, Zarate YA. Epilepsy and Electroencephalographic Abnormalities in SATB2-Associated Syndrome. Pediatr Neurol. 2020 Nov;112:94-100.

Ferdinandusse S, McWalter K, Te Brinke H, IJlst L, Mooijer PM, Ruiter JPN, van Lint AEM, Pras-Raves M, Wever E, Millan F, Guillen Sacoto MJ, Begtrup A, Tarnopolsky M, Brady L, Ladda RL, Sell SL, Nowak CB, Douglas J, Tian C, Ulm E, **Perlman S**, Drack AV, Chong K, Martin N, Brault J, Brokamp E, Toro C, Gahl WA, Macnamara EF, Wolfe L; Undiagnosed Diseases Network, Waisfisz Q, Zwijnenburg PJG, Ziegler A, Barth M, Smith R, Ellingwood S, Gaebler-Spira D, Bakhtiari S, Kruer MC, van Kampen AHC, Wanders RJA, Waterham HR, Cassiman D, Vaz FM. An autosomal dominant neurological disorder caused by de novo variants in FAR1 resulting in uncontrolled synthesis of ether lipids.Genet Med. 2020 Nov 26. Online ahead of print. PMID: 33239752

**Cronin M**, **Wainwright MS.** Underpowered and Too Heterogenous: A Humbling Assessment of the Literature Supporting Neuroprognostication After Pediatric Cardiac Arrest. Pediatr Crit Care Med. 2020 Oct;21(10):915-916.

Weisenberg JLZ, Fitzgerald RT, Constantino JN, Winawer MR, Thio LL; EPGP Investigators (**Novotny EJ**). Familial aggregation of status epilepticus in generalized and focal epilepsies. Neurology. 2020 Oct 13;95(15):e2140-e2149.

Boos MD, **Bozarth XL**, Sidbury R, Cooper AB, Perez F, Chon C, Paras G, **Amlie-Lefond C** [Forehead location and large segmental pattern of facial port-wine stains predict risk of Sturge-Weber syndrome.](https://pubmed.ncbi.nlm.nih.gov/32413446/) J Am Acad Dermatol. 2020 Oct;83(4):1110-1117.

Ebrahimi-Fakhari D, Teinert J, Behne R, Wimmer M, D'Amore A, Eberhardt K, Brechmann B, Ziegler M, Jensen DM, Nagabhyrava P, Geisel G, Carmody E, Shamshad U, Dies KA, Yuskaitis CJ, Salussolia CL, Ebrahimi-Fakhari D, Pearson TS, Saffari A, Ziegler A, Kölker S, Volkmann J, Wiesener A, Bearden DR, Lakhani S, Segal D, Udwadia-Hegde A, Martinuzzi A, Hirst J, **Perlman S**, Takiyama Y, Xiromerisiou G, Vill K, Walker WO, Shukla A, Dubey Gupta R, Dahl N, Aksoy A, Verhelst H, Delgado MR, Kremlikova Pourova R, Sadek AA, Elkhateeb NM, Blumkin L, Brea-Fernández AJ, Dacruz-Álvarez D, Smol T, Ghoumid J, Miguel D, Heine C, Schlump JU, Langen H, Baets J, Bulk S, Darvish H, Bakhtiari S, Kruer MC, Lim-Melia E, Aydinli N, Alanay Y, El-Rashidy O, Nampoothiri S, Patel C, Beetz C, Bauer P, Yoon G, Guillot M, Miller SP, Bourinaris T, Houlden H, Robelin L, Anheim M, Alamri AS, Mahmoud AAH, Inaloo S, Habibzadeh P, Faghihi MA, Jansen AC, Brock S, Roubertie A, Darras BT, Agrawal PB, Santorelli FM, Gleeson J, Zaki MS, Sheikh SI, Bennett JT, Sahin M. Defining the clinical, molecular and imaging spectrum of adaptor protein complex 4-associated hereditary spastic paraplegia. Brain. 2020 Oct 1;143(10):2929-2944.

Sun LR, Harrar D, Drocton G, **Castillo-Pinto C**, Felling R, Carpenter JL, Wernovsky G, McDougall CG, Gailloud P, Pearl MS. Mechanical Thrombectomy for Acute Ischemic Stroke: Considerations in Children. Stroke. 2020 Oct;51(10):3174-3181. doi: 10.1161/STROKEAHA.120.029698. Epub 2020 Sep 11. PMID: 32912096

Kim SH, Green-Snyder L, Lord C, Bishop S, **Steinman KJ**, Bernier R, Hanson E, Goin-Kochel RP, Chung WK. Language characterization in 16p11.2 deletion and duplication syndromes. Am J Med Genet B Neuropsychiatr Genet. 2020 Sep;183(6):380-391.

**Saneto RP.** Mitochondrial diseases: expanding the diagnosis in the era of genetic testing. J Transl Genet Genom. 2020;4:384-428. Epub 2020 Sep 29. PMID: 33426505

Vossler DG, Bainbridge JL, Boggs JG, **Novotny EJ**, Loddenkemper T, Faught E, Amengual-Gual M, Fischer SN, Gloss DS, Olson DM, Towne AR, Naritoku D, Welty TE. Treatment of Refractory Convulsive Status Epilepticus: A Comprehensive Review by the American Epilepsy Society Treatments Committee Epilepsy Curr. 2020 Sep;20(5):245-264.

Bornstein R, Gonzalez B, **Johnson SC**. Mitochondrial pathways in human health and aging. *Mitochondrion*. 2020 Sep 54:72-84. DOI: 10.1016/j.mito.2020.07.007. Epub 2020 Jul 30. PMID: 32738358.

Law EF, **Blume H**, Palermo TM. Longitudinal Impact of Parent Factors in Adolescents With Migraine and Tension-Type Headache. Headache. 2020 Sep;60(8):1722-1733.

Pearl PL, Mink JW, Cohen BH, Bamford N, Bass N, Jordan L, **Wainwright MS**, Larson RB. The President, Past President, Executive Director, and the Board of the Child Neurology Society Denounce Racism and Inequality. Ann Neurol. 2020 Aug;88(2):209-210.

**Patrick KE,** Schultheis MT, Agate FT, McCurdy MD, Daly BP, Tarazi RA, Chute DL, Hurewitz F. Executive function "drives" differences in simulated driving performance between young adults with and without autism spectrum disorder. Child Neuropsychol. 2020 Jul;26(5):649-665.

Simpson TS, Peterson RL, **Patrick KE**, Forster JE, McNally KA. J Concussion Symptom Treatment and Education Program: A Feasibility Study. Head Trauma Rehabil. 2020 Jul 22.

Fernández IS, Abend NS, Amengual-Gual M, Anderson A, Arya R, Aguilar CB, Brenton JN, Carpenter JL, Chapman KE, Clark J, Farias-Moeller R, Gaillard WD, Gaínza-Lein M, Glauser T, Goldstein J, Goodkin HP, Guerriero RM, Lai YC, McDonough T, Mikati MA, **Morgan LA**, **Novotny E Jr**, Payne E, Peariso K, Piantino J, Ostendorf A, Sands TT, Sannagowdara K, Tasker RC, Tchapyjnikov D, Topjian AA, Vasquez A, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; pSERG. Association of guideline publication and delays to treatment in pediatric status epilepticus.Neurology**.** 2020 Jul 1

Crotty EE, Leary SES, Geyer JR, Olson JM, Millard NE, **Sato AA**, Ermoian RP, Cole BL, Lockwood CM, Paulson VA, Browd SR, Ellenbogen RG, Hauptman JS, Lee A, Ojemann JG, Vitanza NA. Children with DIPG and high-grade glioma treated with temozolomide, irinotecan, and bevacizumab: the Seattle Children's Hospital experience. J Neurooncol. 2020 Jul;148(3):607-617.

Nambot S, Faivre L, Mirzaa G, Thevenon J, Bruel AL, Mosca-Boidron AL, Masurel-Paulet A, Goldenberg A, Le Meur N, Charollais A, Mignot C, Petit F, Rossi M, Metreau J, Layet V, Amram D, Boute-Bénéjean O, Bhoj E, Cousin MA, Kruisselbrink TM, Lanpher BC, Klee EW, Fiala E, Grange DK, Meschino WS, Hiatt SM, Cooper GM, Olivié H, Smith WE, Dumas M, Lehman A; CAUSES Study, Inglese C, Nizon M, Guerrini R, Vetro A, Kaplan ES, Miramar D, Van Gils J, Fergelot P, Bodamer O, Herkert JC, Pajusalu S, Õunap K, Filiano JJ, Smol T, Piton A, Gérard B, Chantot-Bastaraud S, Bienvenu T, Li D, Juusola J, Devriendt K, Bilan F, Poé C, Chevarin M, Jouan T, Tisserant E, Rivière JB, Tran Mau-Them F, Philippe C, Duffourd Y, **Dobyns WB**, Hevner R, Thauvin-Robinet C. De novo TBR1 variants cause a neurocognitive phenotype with ID and autistic traits: report of 25 new individuals and review of the literature. Eur J Hum Genet. 2020 Jun;28(6):770-782.

Brock S, Vanderhasselt T, Vermaning S, Keymolen K, Régal L, Romaniello R, Wieczorek D, Storm TM, Schaeferhoff K, Hehr U, Kuechler A, Krägeloh-Mann I, Haack TB, Kasteleijn E, Schot R, Mancini GMS, Webster R, Mohammad S, Leventer RJ, Mirzaa G, **Dobyns WB**, Bahi-Buisson N, Meuwissen M, Jansen AC, Stouffs K. Defining the phenotypical spectrum associated with variants in TUBB2A. J Med Genet. 2020 Jun 22:jmedgenet-2019-106740

**Benedetti GM**, Vartanian RJ, McCaffery H, Shellhaas RA. Early Electroencephalogram Background Could Guide Tailored Duration of Monitoring for Neonatal Encephalopathy Treated with Therapeutic HypothermiaJ Pediatr. 2020 Jun;2 21:81-87.e1

**Marashly A**, Koop J, Loman M, Kim I, Maheshwari M, Lew SM. Multiple hippocampal transections for refractory pediatric mesial temporal lobe epilepsy: seizure and neuropsychological outcomes.

J Neurosurg Pediatr. 2020 Jun 26:1-10.

Rafay MF, Shapiro KA, Surmava AM, deVeber GA, Kirton A, Fullerton HJ, **Amlie-Lefond C**, Weschke B, Dlamini N, Carpenter JL, Mackay MT, Rivkin M, Linds A, Bernard TJ; International Pediatric Stroke Study (IPSS) Group.Spectrum of cerebral arteriopathies in children with arterial ischemic stroke. Neurology. 2020 Jun 9;94(23):e2479-e2490

Holland EL, **Saneto RP**, Knipper EK. Hypermetabolic Syndrome and Dyskinesia After Neurologic Surgery for Labrune Syndrome: A Case Report. A A Pract**.** 2020 May;14**(7**):e01212..

Lennox AL, Hoye ML, Jiang R, Johnson-Kerner BL, Suit LA, Venkataramanan S, Sheehan CJ, Alsina FC, Fregeau B, Aldinger KA, Moey C, Lobach I, Afenjar A, Babovic-Vuksanovic D, Bézieau S, Blackburn PR, Bunt J, Burglen L, Campeau PM, Charles P, Chung BHY, Cogné B, Curry C, D'Agostino MD, Di Donato N, Faivre L, Héron D, Innes AM, Isidor B, Keren B, Kimball A, Klee EW, Kuentz P, Küry S, Martin-Coignard D, Mirzaa G, Mignot C, Miyake N, Matsumoto N, Fujita A, Nava C, Nizon M, Rodriguez D, Blok LS, Thauvin-Robinet C, Thevenon J, Vincent M, Ziegler A, **Dobyns W**, Richards LJ, Barkovich AJ, Floor SN, Silver DL, Sherr EH. Pathogenic DDX3X Mutations Impair RNA Metabolism and Neurogenesis during Fetal Cortical Development. Neuron. 2020 May 6;106(3):404-420.e8

Weaver JJ, Hallam DK, Chick JFB, Vaidya S, Shin DS, **Natarajan N**, Rad N, Reis J, Koo KSH, Shivaram GM, Thibodeau A, Apkon S, Monroe EJ. Transforaminal intrathecal delivery of nusinersen for older children and adults with spinal muscular atrophy and complex spinal anatomy: an analysis of 200 consecutive injections. J Neurointerv Surg. 2020 May

Blumkin L, Leibovitz Z, Krajden-Haratz K, Arad A, Yosovich K, Gindes L, Zerem A, Ben-Sira L, Lev D, Nissenkorn A, Kidron D**, Dobyns WB**, Malinger G, Bahi-Buisson N, Leventer RJ, Lerman-Sagie T. Autosomal dominant TUBB3-related syndrome: Fetal, radiologic, clinical and morphological features Eur J Paediatr Neurol. 2020 May;26:46-60.

Miller I, Scheffer IE, Gunning B, Sanchez-Carpintero R, Gil-Nagel A, Perry MS, **Saneto RP**, Checketts D, Dunayevich E, Knappertz V; GWPCARE2 Study Group. Dose-Ranging Effect of Adjunctive Oral Cannabidiol vs Placebo on Convulsive Seizure Frequency in Dravet Syndrome: A Randomized Clinical Trial. JAMA Neurol. 2020 May 1;77(5):613-621.

Tsai MH, Muir AM, Wang WJ, Kang YN, Yang KC, Chao NH, Wu MF, Chang YC, Porter BE, Jansen LA, Sebire G, Deconinck N, Fan WL, Su SC, Chung WH, Almanza Fuerte EP, Mehaffey MG; University of Washington Center for Mendelian Genomics, Ng CC, Chan CK, Lim KS, Leventer RJ, Lockhart PJ, Riney K, Damiano JA, Hildebrand MS, Mirzaa GM, **Dobyns WB**, Berkovic SF, Scheffer IE, Tsai JW, Mefford HC. Pathogenic Variants in CEP85L Cause Sporadic and Familial Posterior Predominant Lissencephaly. Neuron**.** 2020 Apr 22**;**106(2):237-245.e8.

Sandrone S, Berthaud JV, Carlson C, Cios J, Dixit N, Farheen A, Kraker J, **Owens JWM**, Patino G, Sarva H, Weber D, Schneider LD. Active Learning in Psychiatry Education: Current Practices and Future Perspectives. Front Psychiatry. 2020 Apr 23;11:211.

Nambot S, Hevner RF, **Dobyns WB**. Reply to Hsueh YP et al. Eur J Hum Genet**.** 2020 Apr 9.

Mineyko A, Kirton A, Billinghurst L, Tatishvili NN, Wintermark M, deVeber G, Fox C; SIPS Investigators. Kirton A, Abdalla A, Zafeiriou D, Friedman N, Aprasidze T, Kolk A, Armstrong J, Ichord R, **Amlie-Lefond,C** deVeber G, Kovacevic G, Hernandez Chavez M, Mackay M, Titomanlio L, Guilliams K, Elbers J, Fullerton H, Benedict S, Dowling M, Jordan L, Pergami P Seizures and Outcome One Year After Neonatal and Childhood Cerebral Sinovenous Thrombosis. Pediatr Neurol. 2020 Apr;105:21-26.

**Johnson SC**, Kayser EB, Bornstein R, Stokes J, Bitto A, Park KY, Pan A, Sun G, Raftery D, Kaeberlein M, Sedensky MM, Morgan PG. Regional metabolic signatures in the Ndufs4(KO) mouse brain implicate defective glutamate/α-ketoglutarate metabolism in mitochondrial disease. *Molecular Genetics and Metabolism*. 2020 Jun;130(2):118-132. doi: 10.1016/j.ymgme.2020.03.007. Epub 2020 Apr 3. PMID: 32331968 27.

Perez FA, Oesch G, **Amlie-Lefond CM** MRI Vessel Wall Enhancement and Other Imaging Biomarkers in Pediatric Focal Cerebral Arteriopathy-Inflammatory Subtype. Stroke. 2020 Mar;51(3):853-859.

Friederich MW, Perez FA, Knight KM, Van Hove RA, Yang SP**, Saneto RP**, Van Hove JLK. Pathogenic variants in NUBPL result in failure to assemble the matrix arm of complex I and cause a complex leukoencephalopathy with thalamic involvement. Mol Genet Metab. 2020 Mar;129(3):236-242.

Gonzalez-Perez P, Smith C, Sebetka WL, Gedlinske A**, Perlman S**, Mathews KD. Clinical and electrophysiological evaluation of myasthenic features in an alpha-dystroglycanopathy cohort (FKRP-predominant). Neuromuscul Disord. 2020 Mar;30**(3)**:213-218.

Barca E, Long Y, Cooley V, Schoenaker R, Emmanuele V, DiMauro S, Cohen BH, Karaa A, Vladutiu GD, Haas R, Van Hove JLK, Scaglia F, Parikh S, Bedoyan JK, DeBrosse SD, Gavrilova RH, **Saneto RP**, Enns GM, Stacpoole PW, Ganesh J, Larson A, Zolkipli-Cunningham Z, Falk MJ, Goldstein AC, Tarnopolsky M, Gropman A, Camp K, Krotoski D, Engelstad K, Rosales XQ, Kriger J, Grier J, Buchsbaum R, Thompson JLP, Hirano M. Mitochondrial diseases in North America: An analysis of the NAMDC Registry. Neurol Genet**.** 2020 Mar 2;6(2**):**e402.. eCollection 2020 Apr

Vabres P, Sorlin A, Kholmanskikh SS, Demeer B, St-Onge J, Duffourd Y, Kuentz P, Courcet JB, Carmignac V, Garret P, Bessis D, Boute O, Bron A, Captier G, Carmi E, Devauchelle B, Geneviève D, Gondry-Jouet C, Guibaud L, Lafon A, Mathieu-Dramard M, Thevenon J, **Dobyns W**B, Bernard G, Polubothu S, Faravelli F, Kinsler VA, Thauvin C, Faivre L, Ross ME, Rivière JB. Author Correction: Postzygotic inactivating mutations of RHOA cause a mosaic neuroectodermal syndrome. Nat Genet**.** 2020 Mar;52(3):353.

Lennox AL, Hoye ML, Jiang R, Johnson-Kerner BL, Suit LA, Venkataramanan S, Sheehan CJ, Alsina FC, Fregeau B, Vabres P, Sorlin A, Kholmanskikh SS, Demeer B, St-Onge J, Duffourd Y, Kuentz P, Courcet JB, Carmignac V, Garret P, Bessis D, Boute O, Bron A, Captier G, Carmi E, Devauchelle B, Geneviève D, Gondry-Jouet C, Guibaud L, Lafon A, Mathieu-Dramard M, Thevenon J, **Dobyns WB**, Bernard G, Polubothu S, Faravelli F, Kinsler VA, Thauvin C, Faivre L, Ross ME, Rivière JB. Author Correction: Postzygotic inactivating mutations of RHOA cause a mosaic neuroectodermal syndrome. Nat Genet. 2020 Mar;52(3):353. doi: 10.1038/s41588-019-0565-x.

Blumkin L, Leibovitz Z, Krajden-Haratz K, Arad A, Yosovich K, Gindes L, Zerem A, Ben-Sira L, Lev D, Nissenkorn A, Kidron D, **Dobyns WB**, Malinger G, Bahi-Buisson N, Leventer RJ, Lerman-Sagie T. Autosomal dominant TUBB3-related syndrome: Fetal, radiologic, clinical and morphological features. Eur J Paediatr Neurol. 2020 Mar 4. pii: S1090-3798(20)30044-1.

DelRosso LM, Yi T, **Chan JHM**, **Wrede JE**, **Lockhart CT**, Ferri R. Determinants of ferritin response to oral iron supplementation in children with sleep movement disorders. Sleep. 2020 Mar 12;43(3). pii: zsz234..

Neighbors A, Moss T, Holloway L, Yu SH, Annese F, Skinner S, **Saneto R**, Steet R. Functional analysis of a novel mutation in the TIMM8A gene that causes deafness-dystonia-optic neuronopathy syndrome. Mol Genet Genomic Med. 2020 Mar;8(3):e1121.

Aldinger KA, Moey C, Lobach I, Afenjar A, Babovic-Vuksanovic D, Bézieau S, Blackburn PR, Bunt J, Burglen L, Campeau PM, Charles P, Chung BHY, Cogné B, Curry C, D'Agostino MD, Di Donato N, Faivre L, Héron D, Innes AM, Isidor B, Keren B, Kimball A, Klee EW, Kuentz P, Küry S, Martin-Coignard D, Mirzaa G, Mignot C, Miyake N, Matsumoto N, Fujita A, Nava C, Nizon M, Rodriguez D, Blok LS, Thauvin-Robinet C, Thevenon J, Vincent M, Ziegler A, **Dobyns W**, Richards LJ, Barkovich AJ, Floor SN, Silver DL, Sherr EH Pathogenic DDX3X Mutations Impair RNA Metabolism and Neurogenesis during Fetal Cortical Development. Neuron. 2020 Feb 28.

Delrosso LM, **Lockhart C, Wrede JE**, Chen ML, Samson M, Reed J, Martin-Washo S, Arp M, Ferri R. Comorbidities in children with elevated periodic limb movement index during sleep. Sleep. 2020 Feb 13;43(2).

**Amlie-Lefond C,** Shaw DWW, Cooper A, **Wainwright MS**, Kirton A, Felling RJ, Abraham MG, Mackay MT, Dowling MM, Torres M, Rivkin MJ, Grabowski EF, Lee S, Kurz JE, McMillan HJ, Barry D, **Lee-Eng J**, Ichord RN. Risk of Intracranial Hemorrhage Following Intravenous tPA (Tissue-Type Plasminogen Activator) for Acute Stroke Is Low in Children. Stroke. 2020 Feb;51(2):542-548.

**Amlie-Lefond C, Wainwright MS**. Response by Amlie-Lefond and Wainwright to Letter Regarding Article, "Organizing for Acute Arterial Ischemic Stroke in Children". Stroke. 2020 Feb;51(2):e37.

Tsai MH, Muir AM, Wang WJ, Kang YN, Yang KC, Chao NH, Wu MF, Chang YC, Porter BE, Jansen LA, Sebire G, Deconinck N, Fan WL, Su SC, Chung WH, Almanza Fuerte EP, Mehaffey MG; University of Washington Center for Mendelian Genomics, Ng CC, Chan CK, Lim KS, Leventer RJ, Lockhart PJ, Riney K, Damiano JA, Hildebrand MS, Mirzaa GM**, Dobyns WB**, Berkovic SF, Scheffer IE, Tsai JW, Mefford HC. Pathogenic Variants in CEP85L Cause Sporadic and Familial Posterior Predominant Lissencephaly. Neuron. 2020 Feb 10.

Press CA, Kirschen M, LaRovere K, Risen S, Guilliams KP, Chung M, Griffith J, Erklauer J, Peariso K, Ducharme-Crevier L, Shah SS, Hall M, **Wainwright MS**. Variation in Treatment and Outcomes of Children With Acute Disseminated Encephalomyelitis. Hosp Pediatr. 2020 Feb;10(2):159-165..

**Patel, P**, Belden A, Handoko R, Paul, R, et al.  Behavioral Impairment and Cognition in Thai Adolescents Affected by HIV. *Global Mental Health*. 9;8:e3. Feb 2020. PMID: 34026234

Nelson BR, Roby JA, **Dobyns WB**, Rajagopal L, Gale M Jr, Adams Waldorf KM. Immune Evasion Strategies Used by Zika Virus to Infect the Fetal Eye and Brain. Viral Immunol. 2020 Jan/Feb;33(1):22-37.

**Marashly A**, Koop J, Loman M, Lee YW, Lew SM. Examining the Utility of Resective Epilepsy Surgery in Children With Electrical Status Epilepticus in Sleep: Long Term Clinical and Electrophysiological Outcomes. Front Neurol. 2020 Jan 15;10:1397.

LaRovere KL, Tasker RC, **Wainwright M**, Reuter-Rice K, Appavu B, Miles D, Lidsky K, Vittner P, Gundersen D, O'Brien NF Transcranial Doppler Ultrasound During Critical Illness in Children: Survey of Practices in Pediatric Neurocritical Care Centers. Pediatric Neurocritical Care Research Group (PNCRG).

Pediatr Crit Care Med. 2020 Jan;21(1):67-74.

**Wainwright MS.** PIRSE: Philosophical Investigations in Refractory Status Epilepticus. Pediatr Crit Care Med. 2020 Jan;21(1):98-99.

Mak CCY, Doherty D, Lin AE, Vegas N, Cho MT, Viot G, Dimartino C, Weisfeld-Adams JD, Lessel D, Joss S, Li C, Gonzaga-Jauregui C, Zarate YA, Ehmke N, Horn D, Troyer C, Kant SG, Lee Y, Ishak GE, Leung G, Barone Pritchard A, Yang S, Bend EG, Filippini F, Roadhouse C, Lebrun N, Mehaffey MG, Martin PM, Apple B, Millan F, Puk O, Hoffer MJV, Henderson LB, McGowan R, Wentzensen IM, Pei S, Zahir FR, Yu M, Gibson WT, Seman A, Steeves M, Murrell JR, Luettgen S, Francisco E, Strom TM, Amlie-Wolf L, Kaindl AM, Wilson WG, Halbach S, Basel-Salmon L, Lev-El N, Denecke J, Vissers LELM, Radtke K, Chelly J, Zackai E, Friedman JM, Bamshad MJ, Nickerson DA; University of Washington Center for Mendelian Genomics, Reid RR, Devriendt K, Chae JH, Stolerman E, McDougall C, Powis Z, Bienvenu T, Tan TY, Orenstein N, **Dobyns WB**, Shieh JT, Choi M, Waggoner D, Gripp KW, Parker MJ, Stoler J, Lyonnet S, Cormier-Daire V, Viskochil D, Hoffman TL, Amiel J, Chung BHY, Gordon CT. MN1 C-terminal truncation syndrome is a novel neurodevelopmental and craniofacial disorder with partial rhombencephalosynapsis Brain. 2020 Jan 1;143(1):55-68.

Goeggel Simonetti B, Rafay MF, Chung M, Lo WD, Beslow LA, Billinghurst LL, Fox CK, Pagnamenta A, Steinlin M, Mackay MT; IPSS Study Group. Collaborators (98) Ashwal S, deVeber G, Ferriero D, Fullerton H, Ichord R, Kirkham F, Lynch JK, O’Callaghan F, Pavlakis S, Sebire G, Willan A, deVeber G, Dowling M, Mackay M, Hernandez Chavez M, Bernard T, Nowak-Gottl U, Benedict S, Rafay M, Yager J, Lo W, Parakh M, Kirton A, Ichord R, Steinlin M, Carpenter J, Rivkin M, Wiznitzer M, O’Callaghan F, Kan L, Zafeiriou D, Fullerton H, Bjornson B, Jordan L, Ashwal S, Torres M, Kolk A, Troncoso M, Whelan H, Pavlakis S, Chan A, Kopyta I, Abdalla A, Gerard T, Golomb M, Humphreys P, Noetzel M, Heyer G, **Amlie-Lefond C**, Yeh A, Pergami P, Cardona F, Kovacevic G, Ganesan V, Elbers J, Friedman N, Wong V, Tatishvili N, Sultan S, Abraham L, Gonzalez V, Titomanlio L, Saengpattrachai M, Lerner NB, Crosswell H, Oliviera C, Andrade Alveal L, Felling R, Hunfeld M, Altuna D, Zecavati N, Buckley D, Deray M, Hassanein S, Brankovic-Sreckovic V, Hegde A, Raybagkar D, Sebire G, Miteff C, Condie J, Khoury C, Carpenter S, Holzhauer S, Garcia Soler P, Nassif El-Hakam L, Conto L, Francisco Vilavedra J, Kosofsky B, Guang Y, Taylor JM, Goldenberg N, Toma A, Beslow LA, **Wainwright M**, Xavier F, Grabowski E, Nass R, Wilson J. Comparative study of posterior and anterior circulation stroke in childhood: Results from the International Pediatric Stroke Study. Neurology. 2020 Jan 28;94(4):e337-e344.

Nambot S, Faivre L, Mirzaa G, Thevenon J, Bruel AL, Mosca-Boidron AL, Masurel-Paulet A, Goldenberg A, Le Meur N, Charollais A, Mignot C, Petit F, Rossi M, Metreau J, Layet V, Amram D, Boute-Bénéjean O, Bhoj E, Cousin MA, Kruisselbrink TM, Lanpher BC, Klee EW, Fiala E, Grange DK, Meschino WS, Hiatt SM, Cooper GM, Olivié H, Smith WE, Dumas M, Lehman A; CAUSES Study, Inglese C, Nizon M, Guerrini R, Vetro A, Kaplan ES, Miramar D, Van Gils J, Fergelot P, Bodamer O, Herkert JC, Pajusalu S, Õunap K, Filiano JJ, Smol T, Piton A, Gérard B, Chantot-Bastaraud S, Bienvenu T, Li D, Juusola J, Devriendt K, Bilan F, Poé C, Chevarin M, Jouan T, Tisserant E, Rivière JB, Tran Mau-Them F, Philippe C, Duffourd Y, **Dobyns WB**, Hevner R, Thauvin-Robinet C. De novo TBR1 variants cause a neurocognitive phenotype with ID and autistic traits: report of 25 new individuals and review of the literature. Eur J Hum Genet. 2020 Jan 31.

Monteiro FP, Curry CJ, Hevner R, Elliott S, Fisher JH, Turocy J, **Dobyns WB**, Costa LA, Freitas E, Kitajima JP, Kok F. Biallelic loss of function variants in ATP1A2 cause hydrops fetalis, microcephaly, arthrogryposis and extensive cortical malformations Eur J Med Genet. 2020 Jan;63(1):103624.

Sandrone S, Berthaud JV, Carlson C, Cios J, Dixit N, Farheen A, Kraker J, **Owens JWM**, Patino G, Sarva H, Weber D, Schneider LD. Strategic Considerations for Applying the Flipped Classroom to Neurology Education. Ann Neurol. 2020 Jan;87(1):4-9.

**2019**

Kahn-Kirby AH, Amagata A, Maeder CI, Mei JJ, Sideris S, Kosaka Y, Hinman A, Malone SA, Bruegger JJ, Wang L, Kim V, Shrader WD, Hoff KG, Latham JC, Ashley EA, Wheeler MT, Bertini E, Carrozzo R, Martinelli D, Dionisi- Vici C, Chapman KA, Enns GM, Gahl W, Wolfe L, Saneto RP, **Johnson SC**, Trimmer JK, Klein MB, Holst CR. Targeting ferroptosis: A novel therapeutic strategy for the treatment of mitochondrial disease-related epilepsy. *PLoS One*. 2019. 14(3):e0214250. PMID: 30921410.

**Johnson SC**, Pan A, Sun GX, Freed A, Stokes JC, Bornstein R, Witkowski M, Li L, Ford JM, Howard CRA, Sedensky MM, Morgan PG. Relevance of experimental paradigms of anesthesia induced neurotoxicity in the mouse. PLoS One. 2019. 14(3)e0213543. PMID: 30897103.

Griffiths KK, Morgan PG, **Johnson SC**, Nambyiah P, Soriano SG, Johnson K, Xu J, Garber C, Maxwell L, Saraiya N. A Summary of Preclinical Poster Presentations at the Sixth Biennial Pediatric Anesthesia Neurodevelopment Assessment (PANDA) Symposium. *J Neurosurg Anesthesiol*. 2019. 31(1):163-165. PMID: 30767942

**Johnson, SC**, Pan A, Li L, Sedensky M, Morgan P: Neurotoxicity of Anesthetics: Mechanisms and Meaning from Mouse Intervention Studies. *Neurotoxicology and Teratology*. 2019. 71:22-31. PMID: 30472095*.*

**Johnson, SC**, Martinez F, Bitto A, Gonzalez B, Tazaerslan C, Cohen C, Delaval L, Timsit J, Knebelmann B, Terzi F, Mahal T, Zhu Y, Morgan P, Sedensky M, Kaeberlein M, Legendre C, Suh Y, Canaud G: mTOR inhibitors may benefit kidney transplant recipients with mitochondrial diseases. *Kidney International*. 2019. 95(2):455-466. PMID: 30471880

**Amlie-Lefond C**, **Wainwright lMS**. Organizing for Acute Arterial Ischemic Stroke in Children. Stroke. 2019 Dec;50(12):3662-3668.

**Gust J,** Annesley CE, Gardner RA, **Bozarth X**.EEG Correlates of Delirium in Children and Young Adults With CD19-Directed CAR T Cell Treatment-Related Neurotoxicity. J Clin Neurophysiol. 2019 Dec 13.

**Bozarth XL**, McGuire J, **Novotny E.** Current Status of Continuous Electroencephalographic Monitoring in Critically Ill Children, Pediatr Neurol. 2019 Dec;101**:**11-17

Friederich MW, Perez FA, Knight KM, Van Hove RA, Yang SP, **Saneto RP**, Van Hove JLK Pathogenic variants in NUBPL result in failure to assemble the matrix arm of complex I and cause a complex leukoencephalopathy with thalamic involvement. Mol Genet Metab. 2019 Dec 30.

Farhadian SF, Mistry H, Kirchwey T, Chiarella J, Calvi R, Chintanaphol M**, Patel P**, Landry ML, Robertson K, Spudich Markers of CNS Injury in Adults Living With HIV With CSF HIV Not Detected vs Detected SS.Open Forum Infect Dis. 2019 Dec 14;6(12):ofz528.

RA, Ceppi F, Rivers J, Annesley C, Summers C, Taraseviciute A, **Gust J**, Leger KJ, Tarlock K, Cooper TM, Finney OC, Brakke H, Li DH, Park JR, Jensen MC. Preemptive mitigation of CD19 CAR T-cell cytokine release syndrome without attenuation of antileukemic efficacy.Gardner Blood. 2019 Dec 12;134(24):2149-2158. doi: 10.1182/blood.2019001463.

Ravanpay AC, **Gust J,** Johnson AJ, Rolczynski LS, Cecchini M, Chang CA, Hoglund VJ, Mukherjee R, Vitanza NA, Orentas RJ, Jensen MC. EGFR806-CAR T cells selectively target a tumor-restricted EGFR epitope in glioblastoma Oncotarget. 2019 Dec 17;10(66):7080-7095.

Bearer C, Agostoni C, Anand KJS, Ambalavanan N, Bhandari V, Bliss JM, Bloomfield F, Bonifacio SL, Buhimschi I, Cilio MR, Coppes M, Czinn SJ, El-Khuffash A, Embleton N, Felderhoff-Müser U, Ferriero DM, Florin T, Fuentes-Afflick E, Gardner W, **Gospe SM Jr**, Gunn A, Gressens P, Guissani D, Haiden N, Hauptman M, Kim KS, Klebanoff M, Lachman P, Lanphear B, Ozen S, Roehr C, Roland D, Rosenblum N, Schwarz M, Staiano A, Stroustrup A, Valente EM, Wilson-Costello D, Wynn J, Molloy E. Toward the elimination of bias in Pediatric Research. Pediatr Res. 2019 Dec;86**(**6):680-681.

Gardner RA, Ceppi F, Rivers J, Annesley C, Summers C, Taraseviciute A, **Gust J,** Leger KJ, Tarlock K, Cooper TM, Finney OC, Brakke H, Li DH, Park JR, Jensen MC. Preemptive mitigation of CD19 CAR T-cell cytokine release syndrome without attenuation of antileukemic efficacy.Blood. **2019 Dec 12;134**(24):2149-2158. doi: 10.1182/blood.2019001463

Wilson JL, **Amlie-Lefond C**, Abruzzo T, Orbach DB, Rivkin MJ, deVeber GA, Pergami P. Survey of practice patterns and preparedness for endovascular therapy in acute pediatric stroke. Childs Nerv Syst. 2019 Dec;35(12):2371-2378.

Parikh S, Galioto R, Lapin B, Haas R, Hirano M, Koenig MK, **Saneto RP**, Zolkipli-Cunningham Z, Goldstein A, Karaa A. Fatigue in primary genetic mitochondrial disease: No rest for the weary. Neuromuscul Disord. 2019 Nov;29(11):895-902.

**Lockrow JP**, Wright JN, **Saneto RP**, **Amlie-Lefond C.** Epileptic Spasms Predict Poor Epilepsy Outcomes After Perinatal Stroke.J Child Neurol. 2019 Nov;34(13):830-836.

Kochanek PM, Grenvik AN, Tasker RC, Carney N, Totten AM, Adelson PD, Selden NR, Davis-O'Reilly C, Hart EL, Bell MJ, Bratton SL, Grant GA, Kissoon N, Reuter-Rice KE, Vavilala MS**, Wainwright MS**. The authors reply.Pediatr Crit Care Med. 2019 Nov;20(11):1105-1107. .

Rummey C, Corben LA, Delatycki MB, Subramony SH, Bushara K, Gomez CM, Hoyle JC, Yoon G, Ravina B, Mathews KD, Wilmot G, Zesiewicz T, **Perlman S,** Farmer JM, Lynch DR. Psychometric properties of the Friedreich Ataxia Rating Scale. Neurol Genet. 2019 Oct 29;5(6):371

Wood AM, Geddes GC, **Marashly A**. Language regression, hemichorea and focal subclinical seizures in a 6-year-old girl with GLUT-1 deficiency Epilepsy Behav Rep. **2019 Oct 24**;14:100340.

**Gust J**, Ishak GE. Chimeric Antigen Receptor T-Cell Neurotoxicity Neuroimaging: More Than Meets the Eye. AJNR Am J Neuroradiol. 2019 Oct;40(10):E50-E51.

Oegema R, Barkovich AJ, Mancini GMS, Guerrini R, **Dobyns WB**. Subcortical heterotopic gray matter brain malformations: Classification study of 107 individuals. Neurology. 2019 Oct 1;93(14):e1360-e1373

Law EF, Powers SW, **Blume H**, Palermo TM. Screening Family and Psychosocial Risk in Pediatric Migraine and Tension-Type Headache: Validation of the Psychosocial Assessment Tool (PAT). Headache. 2019 Oct;59(9):1516-1529.

**Chan JHM**, **Owens JW, Wrede JE**. Case of an In-Laboratory Vagal Nerve Stimulator Titration for Vagal Nerve Stimulator-Induced Pediatric Obstructive Sleep Apnea. J Clin Sleep Med. 2019 Oct 15;15(10):1539-1542.

**Patel PB**, Apornpong T, Puthanakit T, Thongpibul K, Kosalaraksa P, Hansudewechakul R, Kanjanavanit S, Ngampiyaskul C, Luesomboon W, Wongsawat J, Sun LP, Chettra K, Saphonn V, Mellins CA, Malee K, Spudich S, Ananworanich J, Kerr SJ, Paul R; PREDICT/Resilience Study Trajectory Analysis of Cognitive Outcomes in Children With Perinatal HIV.

Group.Pediatr Infect Dis J. 2019 Oct;38(10):1038-1044

DelRosso LM, **Lockhart C**, **Wrede JE**, Chen ML, Samson M, Reed J, Martin-Washo S, Arp M, Ferri R. Comorbidities in children with elevated periodic limb movement index during sleep. Sleep. 2019 Sep 26.

DelRosso LM, Yi T, Chan JHM, **Wrede JE, Lockhart CT,** Ferri R. Determinants of ferritin response to oral iron supplementation in children with sleep movement disorders. Sleep. 2019 Sep 29. pii: zsz234.

Dines JN, Liu YJ, Neufeld-Kaiser W, Sawyer T, Ishak GE, **Tully HM**, Racobaldo M, Sanchez-Valle A, Disteche CM, Juusola J, Torti E, McWalter K, Doherty D, Dipple KM. Expanding phenotype with severe midline brain anomalies and missense variant supports a causal role for FOXA2 in 20p11.2 deletion syndrome. Am J Med Genet A. 2019 Sep;179(9):1783-1790

Gripp KW, Morse LA, Axelrad M, Chatfield KC, Chidekel A, **Dobyns W**, Doyle D, Kerr B, Lin AE, Schwartz DD, Sibbles BJ, Siegel D, Shankar SP, Stevenson DA, Thacker MM, Weaver KN, White SM, Rauen KA. Costello syndrome: Clinical phenotype, genotype, and management guidelines Am J Med Genet A. 2019 Sep;179(9):1725-1744.

Sansevere AJ, Arya R, Sánchez Fernández I, Gaillard WD, Tasker RC, Lai YC, Anderson AE, Tchapyjnikov D, Chapman KE, Brenton JN, Carpenter JL, Gaínza-Lein M, Goldstein JL, Goodkin HP, Jackson MC, Kapur K, Mikati MA, Peariso K, Glauser TA, Topjian AA, **Wainwright M,** Wilfong AA, Williams KL, Loddenkemper T, Abend NS; De Novo Variants in WDR37 Are Associated with Epilepsy Pediatric Status Epilepticus Research Group (pSERG). J Clin Neurophysiol. 2019 Sep;36(5):365-370.

Luat AF, Juhász C, Loeb JA, Chugani HT, Falchek SJ, Jain B, Greene-Roethke C, **Amlie-Lefond C**, Ball KL, Davis A, Pinto A. Neurological Complications of Sturge-Weber Syndrome: Current Status and Unmet Needs. Pediatr Neurol. 2019 Sep;98:31-38

Kanca O, Andrews JC, Lee PT, Patel C, Braddock SR, Slavotinek AM, Cohen JS, Gubbels CS, Aldinger KA, Williams J, Indaram M, Fatemi A, Yu TW, Agrawal PB, Vezina G, Simons C, Crawford J, Lau CC; Undiagnosed Diseases Network, Chung WK, Markello TC, **Dobyns WB**, Adams DR, Gahl WA, Wangler MF, Yamamoto S, Bellen HJ, Malicdan MCV. Colobomas, Dysmorphism, Developmental Delay, Intellectual Disability, and Cerebellar Hypoplasia. Am J Hum Genet. 2019 Aug 1;105(2):413-424. doi: 10.1016/j.ajhg.2019.06.014. Epub 2019 Jul 18. Erratum in: Am J Hum Genet. 2019 Sep 5;105(3):672-674.

Berg AT, Wusthoff C, Shellhaas RA, Loddenkemper T, Grinspan ZM, **Saneto RP**, Knupp KG, Patel A, Sullivan JE, Kossoff EH, Chu CJ, Massey S, Valencia I, Keator C, Wirrell EC, Coryell J, Millichap JJ, Gaillard WD. Immediate outcomes in early life epilepsy: A contemporary account.Epilepsy Behav. 2019 Aug;97:44-50.

Topjian AA, de Caen A, **Wainwright MS**, Abella BS, Abend NS, Atkins DL, Bembea MM, Fink EL, Guerguerian AM, Haskell SE, Kilgannon JH, Lasa JJ, Hazinski MF. Pediatric Post-Cardiac Arrest Care: A Scientific Statement From the American Heart Association. 2019 Aug 6;140(6):e194-e233.

**Patel PB**. HIV-Associated Neurocognitive Disorders. Medlink Neurology. 2019 August;

Sánchez Fernández I, Gaínza-Lein M, Abend NS, Amengual-Gual M, Anderson A, Arya R, Brenton JN, Carpenter JL, Chapman KE, Clark J, Farias-Moeller R, Davis Gaillard W, Glauser TA, Goldstein J, Goodkin HP, Guerriero RM, Hecox K, Jackson M, Kapur K, Kelley SA, Kossoff EHW, Lai YC, McDonough TL, Mikati MA, **Morgan LA**, **Novotny EJ**, Ostendorf AP, Payne ET, Peariso K, Piantino J, Riviello JJ Jr, Sannagowdara K, Stafstrom CE, Tasker RC, Tchapyjnikov D, Topjian AA, Vasquez A, **Wainwright MS,** Wilfong A, Williams K, Loddenkemper T; The onset of pediatric refractory status epilepticus is not distributed uniformly during the day.Pediatric Status Epilepticus Research Group. Seizure. 2019 Aug;70:90-96.

Holt RJ, Young RM, Crespo B, Ceroni F, Curry CJ, Bellacchio E, Bax DA, Ciolfi A, Simon M, Fagerberg CR, van Binsbergen E, De Luca A, Memo L, **Dobyns WB**, Mohammed AA, Clokie SJH, Zazo Seco C, Jiang YH, Sørensen KP, Andersen H, Sullivan J, Powis Z, Chassevent A, Smith-Hicks C, Petrovski S, Antoniadi T, Shashi V, Gelb BD, Wilson SW, Gerrelli D, Tartaglia M, Chassaing N, Calvas P, Ragge NK. De Novo Missense Variants in FBXW11 Cause Diverse Developmental Phenotypes Including Brain, Eye, and Digit Anomalies. Am J Hum Genet. 2019 Jul 23. .

Ng BG, Sosicka P, Agadi S, Almannai M, Bacino CA, Barone R, Botto LD, Burton JE, Carlston C, Chung BH, Cohen JS, Coman D, Dipple KM, Dorrani N, **Dobyns WB**, Elias AF, Epstein L, Gahl WA, Garozzo D, Hammer TB, Haven J, Héron D, Herzog M, Hoganson GE, Hunter JM, Jain M, Juusola J, Lakhani S, Lee H, Lee J, Lewis K, Longo N, Lourenço CM, Mak CCY, McKnight D, Mendelsohn BA, Mignot C, Mirzaa G, Mitchell W, Muhle H, Nelson SF, Olczak M, Palmer CGS, Partikian A, Patterson MC, Pierson TM, Quinonez SC, Regan BM, Ross ME, Guillen Sacoto MJ, Scaglia F, Scheffer IE, Segal D, Singhal NS, Striano P, Sturiale L, Symonds JD, Tang S, Vilain E, Willis M, Wolfe LA, Yang H, Yano S, Powis Z, Suchy SF, Rosenfeld JA, Edmondson AC, Grunewald S, Freeze HH. SLC35A2-CDG: Functional characterization, expanded molecular, clinical, and biochemical phenotypes of 30 unreported Individuals. Hum Mutat. 2019 Jul;40(7):908-925.

**Gust J**, Finney OC, Li D, Brakke HM, Hicks RM, Futrell RB, Gamble DN, Rawlings-Rhea SD, Khalatbari HK, Ishak GE, Duncan VE, Hevner RF, Jensen MC, Park JR, Gardner RA. Glial injury in neurotoxicity after pediatric CD19-directed chimeric antigen receptor T cell therapy. Ann Neurol**. 2019 Jul;86(**1):42-54

**Lockrow JP**, Wright JN**, Saneto RP**, **Amlie-Lefond C**. Epileptic Spasms Predict Poor Epilepsy Outcomes After Perinatal Stroke J Child Neurol. 2019 Jul 24:

**Wainwright MS**. Clinician, Heal Thyself": Decisional Conflict, Value Judgments, and Their Role in Practice Variation in Traumatic Brain Injury Pediatr Crit Care Med. 2019 Jul;20(7):679-680.

Sandrone S, Berthaud JV, Carlson C, Cios J, Dixit N, Farheen A, Kraker J, **Owens JWM**, Patino G, Sarva H, Weber D, Schneider LD. Education Research: Flipped classroom in neurology: Principles, practices, and perspectivesNeurology. 2019 Jul 2;93(1):e106-e111.

Kochanek PM, Tasker RC, Carney N, Totten AM, Adelson PD, Selden NR, Davis-O'Reilly C, Hart EL, Bell MJ, Bratton SL, Grant GA, Kissoon N, Reuter-Rice KE, Vavilala MS, **Wainwright MS**. Guidelines for the Management of Pediatric Severe Traumatic Brain Injury, Third Edition: Update of the Brain Trauma Foundation Guidelines, Executive Summary Neurosurgery. 2019 Jun 1;84(6):1169-1178.

Ibañez LV, Stoep AV, Myers K, Zhou C, Dorsey S, **Steinman KJ**, Stone WL. Promoting early autism detection and intervention in underserved communities: study protocol for a pragmatic trial using a stepped-wedge design.Psychiatry. 2019 Jun 7;19(1):169.

Barry M, Hallam DK, Bernard TJ, **Amlie-Lefond C** What is the Role of Mechanical Thrombectomy in Childhood Stroke?

Pediatr Neurol. 2019 Jun;95:19-25.

Chelse AB, Kurz JE, Gorman KM, Epstein LG, Balmert LC, Ciolino JD, **Wainwright MS**. Remote poststroke headache in children: Characteristics and association with stroke recurrence. Neurol Clin Pract. 2019 Jun;9(3):194-200.

McKeever JD, Schultheis MT, Sim T, Goykhman J, **Patrick K,** Ehde DM, Woods SP. Selective reminding of prospective memory in Multiple Sclerosis Neuropsychol Rehabil. 2019 Jun;29(5):675-690.

Karasozen Y, Osbun JW, Parada CA, Busald T, Tatman P, Gonzalez-Cuyar LF, Hale CJ, Alcantara D, O'Driscoll M, **Dobyns WB**, Murray M, Kim LJ, Byers P, Dorschner MO, Ferreira M Jr. Somatic PDGFRB Activating Variants in Fusiform Cerebral Aneurysms Am J Hum Genet. 2019 May 2;104(5):968-976.

Marsillio LE, Manghi T, Carroll MS, Balmert LC, **Wainwright MS** Heart rate variability as a marker of recovery from critical illness in children.PLoS One. 2019 May 17;14(5)

Amengual-Gual M, Sánchez Fernández I, **Wainwright MS**. Novel drugs and early polypharmacotherapy in status epilepticus Seizure. 2019 May;68:79-88. doi: 10.1016/j.seizure.2018.08.004. Epub 2018 Aug 7. Review.

Guilliams KP, Kirkham FJ, Holzhauer S, Pavlakis S, Philbrook B, **Amlie-Lefond C**, Noetzel MJ, Dlamini N, Sharma M, Carpenter JL, Fox CK, Torres M, Ichord RN, Jordan LC, Dowling MM. Arteriopathy Influences Pediatric Ischemic Stroke Presentation, but Sickle Cell Disease Influences Stroke Management. Stroke. 2019 May;50(5):1089-1094

Messacar K, Sillau S, Hopkins SE, **Otten C**, Wilson-Murphy M, Wong B, Santoro JD, Treister A, Bains HK, Torres A, Zabrocki L, Glanternik JR, Hurst AL, Martin JA, Schreiner T, Makhani N, DeBiasi RL, Kruer MC, Tremoulet AH, Van Haren K, Desai J, Benson LA, Gorman MP, Abzug MJ, Tyler KL, Dominguez SR. Safety, tolerability, and efficacy of fluoxetine as an antiviral for acute flaccid myelitis. Neurology. 2019 Apr 30;92(18):e2118-e2126.

Kochanek PM, Tasker RC, Carney N, Totten AM, Adelson PD, Selden NR, Davis-O'Reilly C, Hart EL, Bell MJ, Bratton SL, Grant GA, Kissoon N, Reuter-Rice KE, Vavilala MS, **Wainwright MS**. Guidelines for the Management of Pediatric Severe Traumatic Brain Injury, Third Edition: Update of the Brain Trauma Foundation Guidelines. Pediatr Crit Care Med. 2019 Mar;20(3S Suppl 1):S1-S82. Erratum in: Pediatr Crit Care Med. 2019 Apr;20(4):404.

Vandervore LV, Schot R, Kasteleijn E, Oegema R, Stouffs K, Gheldof A, Grochowska MM, van der Sterre MLT, van Unen LMA, Wilke M, Elfferich P, van der Spek PJ, Heijsman D, Grandone A, Demmers JAA, Dekkers DHW, Slotman JA, Kremers GJ, Schaaf GJ, Masius RG, van Essen AJ, Rump P, van Haeringen A, Peeters E, Altunoglu U, Kalayci T, Poot RA, **Dobyns WB**, Bahi-Buisson N, Verheijen FW, Jansen AC, Mancini GMS. Heterogeneous clinical phenotypes and cerebral malformations reflected by rotatin cellular dynamics.

Brain. 2019 Apr 1;142(4):867-884

Kahn-Kirby AH, Amagata A, Maeder CI, Mei JJ, Sideris S, Kosaka Y, Hinman A, Malone SA, Bruegger JJ, Wang L, Kim V, Shrader WD, Hoff KG, Latham JC, Ashley EA, Wheeler MT, Bertini E, Carrozzo R, Martinelli D, Dionisi-Vici C, Chapman KA, Enns GM, Gahl W, Wolfe L, **Saneto RP**, **Johnson SC,** Trimmer JK, Klein MB, Holst CR. Targeting ferroptosis: A novel therapeutic strategy for the treatment of mitochondrial disease-related epilepsy PLoS One. 2019 Mar 28;14(3):

**Wainwright MS**. Getting Excited About Paroxysms: Why Treating Sympathetic Hyperarousal After Traumatic Brain Injury May Be More Important Than We Appreciated Pediatr Crit Care Med. 2019 Mar;20(3):295-296.

Kochanek PM, Tasker RC, Carney N, Totten AM, Adelson PD, Selden NR, Davis-O'Reilly C, Hart EL, Bell MJ, Bratton SL, Grant GA, Kissoon N, Reuter-Rice KE, Vavilala MS, **Wainwright MS.** Guidelines for the Management of Pediatric Severe Traumatic Brain Injury, Third Edition: Update of the Brain Trauma Foundation Guidelines, Executive Summary.Pediatr Crit Care Med. 2019 Mar;20(3):280-289

Kochanek PM, Tasker RC, Bell MJ, Adelson PD, Carney N, Vavilala MS, Selden NR, Bratton SL, Grant GA, Kissoon N, Reuter-Rice KE, **Wainwright MS**. Management of Pediatric Severe Traumatic Brain Injury: 2019 Consensus and Guidelines-Based Algorithm for First and Second Tier Therapies. Pediatr Crit Care Med. 2019 Mar;20(3):269-279.

Dlamini N, Muthusami P, **Amlie-Lefond C.** Childhood Moyamoya: Looking Back to the Future.

Pediatr Neurol. 2019 Feb;91:11-19.

Recober A, **Patel PB**, Thibault DP, Hill AW, Kaiser EA, Willis AW. Sociodemographic Factors Associated With Hospital Care for Pediatric Migraine: A National Study Using the Kids' Inpatient Dataset. Pediatr Neurol. **2019 Feb;91:34**-40. doi: 10.1016/j.pediatrneurol.2017.10.023.

Wenger TL, Hopper RA, Rosen A, **Tully HM**, Cunningham ML, Lee A. A genotype-specific surgical approach for patients with Pfeiffer syndrome due to W290C pathogenic variant in FGFR2 is associated with improved developmental outcomes and reduced mortality. Genet Med. 2019 Feb;21(2):471-476.

Piao CS, Holloway AL, Hong-Routson S, **Wainwright MS**. Depression following traumatic brain injury in mice is associated with down-regulation of hippocampal astrocyte glutamate transporters by thrombin J Cereb Blood Flow Metab. 2019 Jan;39(1):58-73.

**2018**

|  |
| --- |
| Global, Regional, and Country-Specific Lifetime Risks of Stroke, 1990 and 2016. GBD 2016 Lifetime Risk of Stroke Collaborators, Feigin VL, Nguyen G, Cercy K, Johnson CO, Alam T, Parmar PG, Abajobir AA, Abate KH, Abd-Allah F, Abejie AN, Abyu GY, Ademi Z, Agarwal G, Ahmed MB, Akinyemi RO, Al-Raddadi R, Aminde LN**, Amlie-Lefond C,** Ansari H, Asayesh H, Asgedom SW, Atey TM, Ayele HT, Banach M, Banerjee A, Barac A, Barker-Collo SL, Bärnighausen T, Barregard L, Basu S, Bedi N, Behzadifar M, Béjot Y, Bennett DA, Bensenor IM, Berhe DF, Boneya DJ, Brainin M, Campos-Nonato IR, Caso V, Castañeda-Orjuela CA, Rivas JC, Catalá-López F, Christensen H, Criqui MH, Damasceno A, Dandona L, Dandona R, Davletov K, de Courten B, deVeber G, Dokova K, Edessa D, Endres M, Faraon EJA, Farvid MS, Fischer F, Foreman K, Forouzanfar MH, Gall SL, Gebrehiwot TT, Geleijnse JM, Gillum RF, Giroud M, Goulart AC, Gupta R, Gupta R, Hachinski V, Hamadeh RR, Hankey GJ, Hareri HA, Havmoeller R, Hay SI, Hegazy MI, Hibstu DT, James SL, Jeemon P, John D, Jonas JB, Jóźwiak J, Kalani R, Kandel A, Kasaeian A, Kengne AP, Khader YS, Khan AR, Khang YH, Khubchandani J, Kim D, Kim YJ, Kivimaki M, Kokubo Y, Kolte D, Kopec JA, Kosen S, Kravchenko M, Krishnamurthi R, Kumar GA, Lafranconi A, Lavados PM, Legesse Y, Li Y, Liang X, Lo WD, Lorkowski S, Lotufo PA, Loy CT, Mackay MT, Abd El Razek HM, Mahdavi M, Majeed A, Malekzadeh R, Malta DC, Mamun AA, Mantovani LG, Martins SCO, Mate KK, Mazidi M, Mehata S, Meier T, Melaku YA, Mendoza W, Mensah GA, Meretoja A, Mezgebe HB, Miazgowski T, Miller TR, Ibrahim NM, Mohammed S, Mokdad AH, Moosazadeh M, Moran AE, Musa KI, Negoi RI, Nguyen M, Nguyen QL, Nguyen TH, Tran TT, Nguyen TT, Anggraini Ningrum DN, Norrving B, Noubiap JJ, O’Donnell MJ, Olagunju AT, Onuma OK, Owolabi MO, Parsaeian M, Patton GC, Piradov M, Pletcher MA, Pourmalek F, Prakash V, Qorbani M, Rahman M, Rahman MA, Rai RK, Ranta A, Rawaf D, Rawaf S, Renzaho AM, Robinson SR, Sahathevan R, Sahebkar A, Salomon JA, Santalucia P, Santos IS, Sartorius B, Schutte AE, Sepanlou SG, Shafieesabet A, Shaikh MA, Shamsizadeh M, Sheth KN, Sisay M, Shin MJ, Shiue I, Silva DAS, Sobngwi E, Soljak M, Sorensen RJD, Sposato LA, Stranges S, Suliankatchi RA, Tabarés-Seisdedos R, Tanne D, Nguyen CT, Thakur JS, Thrift AG, Tirschwell DL, Topor-Madry R, Tran BX, Nguyen LT, Truelsen T, Tsilimparis N, Tyrovolas S, Ukwaja KN, Uthman OA, Varakin Y, Vasankari T, Venketasubramanian N, Vlassov VV, Wang W, Werdecker A, Wolfe CDA, Xu G, Yano Y, Yonemoto N, Yu C, Zaidi Z, El Sayed Zaki M, Zhou M, Ziaeian B, Zipkin B, Vos T, Naghavi M, Murray CJL, Roth GA.N Engl J Med. 2018 Dec 20;379(25):2429-2437. |
| Venot Q, Blanc T, Rabia S, Berteloot L, Ladraa S, Duong J, Blanc E, **Johnson SC**, Hoguin C, Boccara O, Sarnacki S, Boddaert N, Pannier S, Martinez F, Magassa S, Yamaguchi J, Knebelmann B, Merville P, Grenier N, Joly D, Cormier-Daire V, Michot C, Bole-Feysot C, Picard A, Soupre V, Lyonnet S, Sadoine J, Slimani L, Chaussain C, Laroche-Raynaud C, Guibaud L, Broissand C, Amiel J, Legendre C, Terzi F, Canaud G: Targeted therapy in patients with PIK3CA-related overgrowth syndrome. *Nature*. 2018. 558:p540–546. PMID: 29899452**Blume HK.** Posttraumatic headache in pediatrics: an update and review**. Curr Opin Pediatr. 2018 Dec;30(6):755-763.** |
| Rhombencephalosynapsis: Fused cerebellum, confused geneticists. Aldinger KA, Dempsey JC**, Tully HM,** Grout ME, Mehaffey MG, **Dobyns WB,** Doherty D. Am J Med Genet C Semin Med Genet. 2018 Dec;178(4):432-439. doi: 10.1002/ajmg.c.31666. Review. |
| An update on oculocerebrocutaneous (Delleman-Oorthuys) syndrome. Moog U, **Dobyns WB.** Am J Med Genet C Semin Med Genet. 2018 Dec;178(4):414-422**Gust J,** Taraseviciute A, Turtle CJ. Neurotoxicity Associated with CD19-Targeted CAR-T Cell Therapies.CNS Drugs. **2018 Dec;32**(12):1091-1101 |
| MACF1 Mutations Encoding Highly Conserved Zinc-Binding Residues of the GAR Domain Cause Defects in Neuronal Migration and Axon Guidance. Dobyns WB, Aldinger KA, Ishak GE, Mirzaa GM, Timms AE, Grout ME, Dremmen MHG, Schot R, Vandervore L, van Slegtenhorst MA, Wilke M, Kasteleijn E, Lee AS, Barry BJ, Chao KR, Szczałuba K, Kobori J, Hanson-Kahn A, Bernstein JA, Carr L, D'Arco F, Miyana K, Okazaki T, Saito Y, Sasaki M, Das S, Wheeler MM, Bamshad MJ, Nickerson DA; University of Washington Center for Mendelian Genomics; Center for Mendelian Genomics at the Broad Institute of MIT and Harvard, Engle EC, Verheijen FW, Doherty D, Mancini GMS. Am J Hum Genet. 2018 Dec 6;103(6):1009-1021. |
| **Bozarth X,** Dines JN, Cong Q, Mirzaa GM, Foss K, Lawrence Merritt J 2nd, Thies J, Mefford HC**, Novotny E.** Expanding clinical phenotype in CACNA1C related disorders: From neonatal onset severe epileptic encephalopathy to late-onset epilepsy**.**Am J Med Genet A. 2018 Dec;176(12):2733-2739. |
| **Benedetti GM**, Silverstein FS. Targeted Temperature Management in Pediatric Neurocritical Care.Pediatr Neurol. 2018 Nov;88:12-24. Messacar K, Sillau S, Hopkins SE**, Otten C,** Wilson-Murphy M, Wong B, Santoro JD, Treister A, Bains HK, Torres A, Zabrocki L, Glanternik JR, Hurst AL, Martin JA, Schreiner T, Makhani N, DeBiasi RL, Kruer MC, Tremoulet AH, Van Haren K, Desai J, Benson LA, Gorman MP, Abzug MJ, Tyler KL, Dominguez SR. Safety, tolerability, and efficacy of fluoxetine as an antiviral for acute flaccid myelitis. Neurology. 2018 Nov 9.  |
| **Amlie-Lefond C**, Flanagan J, Kanter J, **Dobyns WB**. The Genetic Landscape of Cerebral Steno-Occlusive Arteriopathy and Stroke in Sickle Cell Anemia. J Stroke Cerebrovasc Dis. 2018 Nov;27(11):2897-2904.  |
| **Bozarth X,** Foss K, Mefford HC. A de novo in-frame deletion of CASK gene causes early onset infantile spasms and supratentorial cerebral malformation in a female patient Am J Med Genet A. 2018 Nov;176(11):2425-2429. |
| Schanze I, Bunt J, Lim JWC, Schanze D, Dean RJ, Alders M, Blanchet P, Attié-Bitach T, Berland S, Boogert S, Boppudi S, Bridges CJ, Cho MT**, Dobyns WB**, Donnai D, Douglas J, Earl DL, Edwards TJ, Faivre L, Fregeau B, Genevieve D, Gérard M, Gatinois V, Holder-Espinasse M, Huth SF, Izumi K, Kerr B, Lacaze E, Lakeman P, Mahida S, Mirzaa GM, Morgan SM, Nowak C, Peeters H, Petit F, Pilz DT, Puechberty J, Reinstein E, Rivière JB, Santani AB, Schneider A, Sherr EH, Smith-Hicks C, Wieland I, Zackai E, Zhao X, Gronostajski RM, Zenker M, Richards LJ. NFIB Haploinsufficiency Is Associated with Intellectual Disability and Macrocephaly Am J Hum Genet. 2018 Nov 1;103(5):752-768. |
| Piacitelli AM, Jensen DM, Brandling-Bennett H, Gray MM, Batra M**, Gust J**, Thaker A, Paschal C, Tsuchiya K, Pritchard CC, Perkins J, Mirzaa GM, Bennett JT. Characterization of a severe case of PIK3CA-related overgrowth at autopsy by droplet digital polymerase chain reaction and report of PIK3CA sequencing in 22 patients. Am J Med Genet A. 2018 Nov;176(11):2301-2308.**Marashly A**, Loman MM, Lew SM Stereotactic laser ablation for nonlesional cingulate epilepsy: case report. .J Neurosurg Pediatr. 2018 Nov 1;22(5):481-488Fullerton HJ, Stence N, Hills NK, Jiang B, **Amlie-Lefond C,** Bernard TJ, Friedman NR, Ichord R, Mackay MT, Rafay MF, Chabrier S, Steinlin M, Elkind MSV, deVeber GA, Wintermark M; VIPS Investigators. Focal Cerebral Arteriopathy of Childhood: Novel Severity Score and Natural History. Stroke. 2018 Nov;49(11):2590-2596.  |
| **Shurtleff H**, Barry D, Chanprasert S, Firman T, **Warner M, Saneto RP.** Cognitive characteristics of mitochondrial diseases in children. Epilepsy Behav. 2018 Nov;88:235-243 |
| Paganoni S, Nicholson K, **Leigh F,** Swoboda K, Chad D, Drake K, Haley K, Cudkowicz M, Berry JD. Developing multidisciplinary clinics for neuromuscular care and research Muscle Nerve. 2017 Nov;56(5):848-858.  |
| **Leigh F**, Ferlini A, Biggar D, Bushby K, Finkel R, Morgenroth LP, Wagner KR. Neurology Care, Diagnostics, and Emerging Therapies of the Patient With Duchenne Muscular Dystrophy. Pediatrics. 2018 Oct;142(Suppl 2):S5-S16. |
| **Lockrow J, Tully H, Saneto RP.** Epileptic spasms as the presenting seizure type in a patient with a new "O" of TORCH, congenital Zika virus infection Epilepsy Behav Case Rep. 2018 Oct 18;11:1-3. |
| Ramos EM, Carecchio M, Lemos R, Ferreira J, Legati A, Sears RL, Hsu SC, Panteghini C, Magistrelli L, Salsano E, Esposito S, Taroni F, Richard AC, Tranchant C, Anheim M, Ayrignac X, Goizet C, Vidailhet M, Maltete D, Wallon D, Frebourg T, Pimentel L, Geschwind DH, Vanakker O, Galasko D, Fogel BL, Innes AM, Ross A, **Dobyns WB**, Alcantara D, O'Driscoll M, Hannequin D, Campion D; French PFBC study group, Oliveira JR, Garavaglia B, Coppola G, Nicolas G. Primary brain calcification: an international study reporting novel variants and associated phenotypes. Eur J Hum Genet. 2018 Oct;26(10):1462-1477 |
| **Benedetti GM**, Silverstein FS, Rau SM, Lester SG, Benedetti MH, Shellhaas RA. Sedation and Analgesia Influence Electroencephalography Monitoring in Pediatric Neurocritical Care.Pediatr Neurol. 2018 Oct;87:57-64. Coryell J, Gaillard WD, Shellhaas RA, Grinspan ZM, Wirrell EC, Knupp KG, Wusthoff CJ, Keator C, Sullivan JE, Loddenkemper T, Patel A, Chu CJ, Massey S, **Novotny EJ Jr, Saneto RP,** Berg AT. Pediatrics. Neuroimaging of Early Life Epilepsy 2018 Sep;142(3). |
| Riley JS, Quarato G, Cloix C**, Lopez J,** O'Prey J, Pearson M, Chapman J, Sesaki H, Carlin LM, Passos JF, Wheeler AP, Oberst A, Ryan KM, Tait SW. Mitochondrial inner membrane permeabilisation enables mtDNA release during apoptosis. EMBO J. 2018 Sep 3;37(17). |
| Grassia F, Poliakov AV, **Poliachik SL,** Casimo K, Friedman SD, **Shurtleff H,** Giussani C**, Novotny EJ,** Ojemann JG, Hauptman JS. Changes in resting-state connectivity in pediatric temporal lobe epilepsy J Neurosurg Pediatr. 2018 Sep;22(3):270-275. |
| López-Chiriboga AS, Klein C, Zekeridou A, McKeon A, Dubey D, Flanagan EP, Lennon VA, Tillema JM, Wirrell EC, Patterson MC, Gadoth A, Aaen JG, Brenton JN, Bui JD, Moen A**, Otten C,** Piquet A, Pittock SJ. LGI1 and CASPR2 neurological autoimmunity in children. Ann Neurol. 2018 Sep;84(3):473-480. |
| **Wainwright MS.** Exposing and Engaging Unknown Unknowns in Traumatic Brain Injury Crit Care Med. 2018 Sep;46(9):1556-1557. |
| Vasquez A, Gaínza-Lein M, Sánchez Fernández I, Abend NS, Anderson A, Brenton JN, Carpenter JL, Chapman K, Clark J, Gaillard WD, Glauser T, Goldstein J, Goodkin HP, Lai YC, Loddenkemper T, McDonough TL, Mikati MA, Nayak A, Payne E, Riviello J, Tchapyjnikov D, Topjian AA, **Wainwright MS,** Tasker RC; Pediatric Status Epilepticus Research Group (pSERG). Hospital Emergency Treatment of Convulsive Status Epilepticus: Comparison of Pathways From Ten Pediatric Research Centers Pediatr Neurol. 2018 Sep;86:33-41 |
| Vora SB, **Amlie-Lefond C,** Perez FA, Melvin AJ. Varicella-Associated Stroke. J Pediatr. 2018 Aug;199:281-281.e1. |
| Brock S, Stouffs K, Scalais E, D'Hooghe M, Keymolen K, Guerrini R, **Dobyns WB**, Di Donato N, Jansen AC. Tubulinopathies continued: refining the phenotypic spectrum associated with variants in TUBG1. Eur J Hum Genet. 2018 Aug;26(8):1132-1142.  |
| Fat, Pharmacokinetics, and Fosphenytoin: Bigger Doses May Be Better. **Wainwright MS.** Pediatr Crit Care Med. 2018 Aug;19(8):784-785. |
| **Wrede JE**, Parsons EC, Watson NF. A Novel Treatment for Nasolacrimal Air Regurgitation Into the Eye With CPAP: The Total Face Mask J Clin Sleep Med. 2018 Aug 15;14(8):1415-1417 |
| Chan P, **Patel P**, Hellmuth J, Colby DJ, Kroon E, Sacdalan C, Pinyakorn S, Jagodzinski L, Krebs S, Ananworanich J, Valcour V, Spudich S; Distribution of Human Immunodeficiency Virus (HIV) Ribonucleic Acid in Cerebrospinal Fluid and Blood Is Linked to CD4/CD8 Ratio During Acute HIV. RV254/SEARCH 010 Study Team.J Infect Dis. 2018 Aug 14;218(6):937-945. Schaffer AE, Breuss MW, Caglayan AO, Al-Sanaa N, Al-Abdulwahed HY, Kaymakçalan H, Yılmaz C, Zaki MS, Rosti RO, Copeland B, Baek ST, Musaev D, Scott EC, Ben-Omran T, Kariminejad A, Kayserili H, Mojahedi F, Kara M, Cai N, Silhavy JL, Elsharif S, Fenercioglu E, Barshop BA, Kara B, Wang R, Stanley V, James KN, Nachnani R, Kalur A, Megahed H, Incecik F, Danda S, Alanay Y, Faqeih E, Melikishvili G, Mansour L, Miller I, Sukhudyan B, Chelly J, **Dobyns WB**, Bilguvar K, Jamra RA, Gunel M, Gleeson JG. Biallelic loss of human CTNNA2, encoding αN-catenin, leads to ARP2/3 complex overactivity and disordered cortical neuronal migration. Nat Genet. 2018 Aug;50(8):1093-1101 |
| Nickels KC, Zaccariello MJ, **Hamiwka LD**, Wirrell EC. Cognitive and neurodevelopmental comorbidities in paediatric epilepsy Nat Rev Neurol. 2016 Aug;12(8):465-76. |
| Amengual-Gual M, Sánchez Fernández I, **Wainwright MS.** Novel drugs and early polypharmacotherapy in status epilepticus. Seizure. 2018 Aug 7**Patrick KE**, Hurewitz F, McCurdy MD, Agate FT, Daly BP, Tarazi RA, Chute DL, Schultheis MT. Driving Comparisons Between Young Adults with Autism Spectrum Disorder and Typical Development. J Dev Behav Pediatr. **2018 Jul/Aug;39**(6):451-460 |
| The genotypic spectrum of ALDH7A1 mutations resulting in pyridoxine dependent epilepsy: a common epileptic encephalopathy. Coughlin CR 2nd, Swanson MA, Spector E, Meeks NJL, Kronquist KE, Aslamy M, Wempe MF, van Karnebeek CDM**, Gospe SM Jr**, Aziz VG, Tsai BP, Gao H, Nagy PL, Hyland K, van Dooren SJM, Salomons GS, Van Hove JLK. J Inherit Metab Dis. 2018 Jul 24. |
| Piacitelli AM, Jensen DM, Brandling-Bennett H, Gray MM, Batra M**, Gust J,** Thaker A, Paschal C, Tsuchiya K, Pritchard CC, Perkins J, Mirzaa GM, Bennett JT. Characterization of a severe case of PIK3CA-related overgrowth at autopsy by droplet digital polymerase chain reaction and report of PIK3CA sequencing in 22 patients. Am J Med Genet A. 2018 Jul 31  |
| **Gospe SM Jr.** Developmental outcome in pyridoxine-dependent epilepsy: Better late (onset) than early Eur J Paediatr Neurol. 2018 Jul;22(4):575-576. |
| Harms MB, **Shannon Bowen KE**, Hanson JL, Pollak Instrumental learning and cognitive flexibility processes are impaired in children exposed to early life stress. SD.Dev Sci. 2018 Jul;21(4):e12596.  |
| Oesch G, Maga AM, Friedman SD, Poliachik SL, Budech CB, Wright JN, Bok LA, **Gospe SM Jr.** Geometric morphometrics reveal altered corpus callosum shape in pyridoxine-dependent epilepsy Neurology. 2018 Jul 3;91(1):e78-e86. |
| PARD3 dysfunction in conjunction with dynamic HIPPO signaling drives cortical enlargement with massive heterotopia. Liu WA, Chen S, Li Z, Lee CH, Mirzaa G, **Dobyns WB**, Ross ME, Zhang J, Shi SH. Genes Dev. 2018 Jun 1;32(11-12):763-780 |
|  |
| Wenger TL, Hopper RA, Rosen A, **Tully HM**, Cunningham ML, Lee A. A genotype-specific surgical approach for patients with Pfeiffer syndrome due to W290C pathogenic variant in FGFR2 is associated with improved developmental outcomes and reduced mortality Genet Med. 2018 Jun 18. |
| Sánchez Fernández I, Gaínza-Lein M, Abend NS, Anderson AE, Arya R, Brenton JN, Carpenter JL, Chapman KE, Clark J, Gaillard WD, Glauser TA, Goldstein JL, Goodkin HP, Helseth AR, Jackson MC, Kapur K, Lai YC, McDonough TL, Mikati MA, Nayak A, Peariso K, Riviello JJ Jr, Tasker RC, Tchapyjnikov D, Topjian AA, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; Pediatric Status Epilepticus Research Group (pSERG): **Morgan, LA**. Factors associated with treatment delays in pediatric refractory convulsive status epilepticus. Neurology. 2018 May 8;90(19):e1692-e1701. |
| Shukla A**, Saneto RP**, Hebbar M, Mirzaa G, Girisha KM. A neurodegenerative mitochondrial disease phenotype due to biallelic loss-of-function variants in PNPLA8 encoding calcium-independent phospholipase A2γ. Am J Med Genet A. 2018 May;176(5):1232-1237. |
| Kannan N, Wang J, Mink RB, **Wainwright MS**, Groner JI, Bell MJ, Giza CC, Zatzick DF, Ellenbogen RG, Boyle LN, Mitchell PH, Rivara FP, Rowhani-Rahbar A, Vavilala MS; PEGASUS (Pediatric Guideline Adherence Outcomes) Study. Timely Hemodynamic Resuscitation and Outcomes in Severe Pediatric Traumatic Brain Injury: Preliminary Findings. Pediatr Emerg Care. 2018 May;34(5):325-329.  |
| Leibovitz Z, Mandel H, Falik-Zaccai TC, Ben Harouch S, Savitzki D, Krajden-Haratz K, Gindes L, Tamarkin M, Lev D, **Dobyns WB**, Lerman-Sagie T. Walker-Warburg syndrome and tectocerebellar dysraphia: A novel association caused by a homozygous DAG1 mutation. Eur J Paediatr Neurol. 2018 May;22(3):525-531.  |
| Heinzen EL, O'Neill AC, Zhu X, Allen AS, Bahlo M, Chelly J, Chen MH, **Dobyns WB**, Freytag S, Guerrini R, Leventer RJ, Poduri A, Robertson SP, Walsh CA, Zhang M; Epi4K Consortium; Epilepsy Phenome/Genome Project. De novo and inherited private variants in MAP1B in periventricular nodular heterotopia. PLoS Genet.2018 May 8;14(5):. |
| Di Donato N, Timms AE, Aldinger KA, Mirzaa GM, Bennett JT, Collins S, Olds C, Mei D, Chiari S, Carvill G, Myers CT, Rivière JB, Zaki MS; University of Washington Center for Mendelian Genomics, Gleeson JG, Rump A, Conti V, Parrini E,Ross ME, Ledbetter DH, Guerrini R, **Dobyns WB.** Analysis of 17 genes detects mutations in 81% of 811 patients with lissencephaly. Genet Med. 2018 Apr 19.  |
| Grinspan ZM, Shellhaas RA, Coryell J, Sullivan JE, Wirrell EC, Mytinger JR, Gaillard WD, Kossoff EH, Valencia I, Knupp KG, Wusthoff C, Keator C, Ryan N, Loddenkemper T, Chu CJ, **Novotny EJ Jr**, Millichap J, Berg AT. Comparative Effectiveness of Levetiracetam vs Phenobarbital for Infantile Epilepsy. JAMA Pediatr. 2018 Apr 1;172(4):352-360.  |
| Gaínza-Lein M, Sánchez Fernández I, Jackson M, Abend NS, Arya R, Brenton JN, Carpenter JL, Chapman KE, Gaillard WD, Glauser TA, Goldstein JL, Goodkin HP, Kapur K, Mikati MA, Peariso K, Tasker RC, Tchapyjnikov D, Topjian AA, **Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; Pediatric Status Epilepticus Research Group. Association of Time to Treatment With Short-term Outcomes for Pediatric Patients With Refractory Convulsive Status Epilepticus. JAMA Neurol. 2018 Apr 1;75(4):410-418.  |
| **Patel PB**, Spudich, S. Global Health Neurology: HIV/AIDS. Semin Neurol. 2018 Apr;38(2):238-246. |
| Weaver JJ**, Natarajan N**, Shaw DWW, Apkon SD, Koo KSH, Shivaram GM, Monroe EJ. Transforaminal intrathecal delivery of nusinersen using cone-beam computed tomography for children with spinal muscular atrophy and extensive surgical instrumentation: early results of technical success and safety Pediatr Radiol. 2018 Mar;48(3):392-397.**Patel P**, Ross A, Henretig FM, Liu G, Harding B, Panzer J. Clinical Reasoning: A 12-year-old girl with headache and change in mental status. Neurology. **2018 Mar 13**;90(11):524-529. |
| Monroe EJ, **Amlie-Lefond CM** Cone beam computed tomography-guided transpterygoidal aspiration of a carotid space abscess in Lemierre's syndrome.. Radiol Case Rep. 2018 Mar 30;13(3):618-621. doi: 10.1016/j.radcr.2018.03.007. eCollection 2018 Jun. |
| Ruggeri G, Timms AE, Cheng C, Weiss A, **Kollros P**, Chapman T, **Tully H**, Mirzaa GM. Bi-allelic mutations of CCDC88C are a rare cause of severe congenital hydrocephalus. Am J Med Genet A. 2018 Mar;176(3):676-681. |
| Adams Waldorf KM, Nelson BR, Stencel-Baerenwald JE, Studholme C, Kapur RP, Armistead B, Walker CL, Merillat S, Vornhagen J, Tisoncik-Go J, Baldessari A, Coleman M, Dighe MK, Shaw DWW, Roby JA, Santana-Ufret V, Boldenow E, Li J, Gao X,Davis MA, Swanstrom JA, Jensen K, Widman DG, Baric RS, Medwid JT, Hanley KA, Ogle J, Gough GM, Lee W, English C, Durning WM, Thiel J, Gatenby C, Dewey EC, Fairgrieve MR, Hodge RD, Grant RF, Kuller L, **Dobyns WB**, Hevner RF, Gale M Jr, Rajagopal L. Congenital Zika virus infection as a silent pathology with loss of neurogenic output in the fetal brain. Nat Med. 2018 Mar;24(3):368-374.  |
| Berg AT, Chakravorty S, Koh S, Grinspan ZM, Shellhaas RA, **Saneto RP**, Wirrell EC, Coryell J, Chu CJ, Mytinger JR, Gaillard WD, Valencia I, Knupp KG, Loddenkemper T, Sullivan JE, Poduri A, Millichap JJ, Keator C, Wusthoff C, Ryan N, **Dobyns WB**, Hegde M. Why West? Comparisons of clinical, genetic and molecular features of infants with and without spasms. PLoS One. 2018 Mar 8;13(3): |
| **Natarajan N, Beatty CW, Gust J, Hamiwka L.** Provider Practices of Phenobarbital Discontinuation in Neonatal Seizures J Child Neurol. 2018 Feb;33(2):153-157.  |
| Evaluation and Acute Management of Ischemic Stroke in Infants and Children. **Amlie-Lefond C.** Continuum (Minneap Minn). 2018 Feb;24(1, Child Neurology):150-170. doi: 10.1212. Review. |
| PURA syndrome: clinical delineation and genotype-phenotype study in 32 individuals with review of published literature. Reijnders MRF, Janowski R, Alvi M, Self JE, van Essen TJ, Vreeburg M, Rouhl RPW, Stevens SJC, Stegmann APA, Schieving J, Pfundt R, van Dijk K, Smeets E, Stumpel CTRM, Bok LA, Cobben JM, Engelen M, Mansour S, Whiteford M, Chandler KE, Douzgou S, Cooper NS, Tan EC, Foo R, Lai AHM, Rankin J, Green A, Lönnqvist T, Isohanni P, Williams S**, Ruhoy I,** Carvalho KS, Dowling JJ, Lev DL, Sterbova K, Lassuthova P, Neupauerová J, Waugh JL, Keros S, Clayton-Smith J, Smithson SF, Brunner HG, van Hoeckel C, Anderson M, Clowes VE, Siu VM, Ddd Study T, Selber P, Leventer RJ, Nellaker C, Niessing D, Hunt D, Baralle D. J Med Genet. 2018 Feb;55(2):104-113 |
| Weaver KE, Poliakov A, **Novotny EJ,** Olson JD, Grabowski TJ, Ojemann JG.Electrocorticography and the early maturation of high-frequency suppression within the default mode network. J Neurosurg Pediatr. 2018 Feb;21(2):133-140.  |
| Phelps IG, Dempsey JC, Grout ME, Isabella CR, **Tully HM**, Doherty D, Bachmann-Gagescu R. Interpreting the clinical significance of combined variants in multiple recessive disease genes: systematic investigation of Joubert syndrome yields little support for oligogenicity. Genet Med. 2018 Feb;20(2):223-233.  |
| **Wainwright MS.** Neurologic Complications in the Pediatric Intensive Care Unit. Continuum (Minneap Minn). 2018 Feb;24(1, Child Neurology):288-299.  |
| Fry AE, Fawcett KA, Zelnik N, Yuan H, Thompson BAN, Shemer-Meiri L, Cushion TD, Mugalaasi H, Sims D, Stoodley N, Chung SK, Rees MI, Patel CV, Brueton LA,Layet V, Giuliano F, Kerr MP, Banne E, Meiner V, Lerman-Sagie T, Helbig KL,Kofman LH, Knight KM, Chen W, Kannan V, Hu C, Kusumoto H, Zhang J, Swanger SA, Shaulsky GH, Mirzaa GM, Muir AM, Mefford HC, **Dobyns WB**, Mackenzie AB, Mullins JGL, Lemke JR, Bahi-Buisson N, Traynelis SF, Iago HF, Pilz DT. De novo mutations in GRIN1 cause extensive bilateral polymicrogyria. Brain. 2018 Jan 22. doi:10.1093/brain/awx358. |
| Wang AC, Ibrahim GM, Poliakov AV, Wang PI, Fallah A, Mathern GW, Buckley RT, Collins K, Weil AG**, Shurtleff HA, Warner MH,** Perez FA, Shaw DW, Wright JN, **Saneto RP, Novotny EJ,** Lee A, Browd SR, Ojemann JG. Corticospinal tract atrophy and motor fMRI predict motor preservation after functional cerebral hemispherectomy. J Neurosurg Pediatr. 2018 Jan;21(1):81-89 |
| Steiner JE, McCoy GN, Hess CP, **Dobyns WB**, Metry DW, Drolet BA, Maheshwari M, Siegel DH. Structural malformations of the brain, eye, and pituitary gland in PHACE syndrome. Am J Med Genet A. 2018 Jan;176(1):48-55. doi:10.1002/ajmg.a.38523 |

**Patel P**., Ross, A., Lui, G., Harding, B., Henretig, F., Panzer, J. Clinical Reasoning: A 12 year old Female with Headache and Change in Mental Status. Neurology. 90(11): 524-529, 2018

Chan P, Hellmuth J, Colby D, Kroon E, Sacdalan C, Fletcher J, **Patel P**, Pinyakorn S, Valcour V, Ananworanich J, Spudich S.J Safety of lumbar puncture procedure in an international research setting during acute HIV infection.

Virus Erad. 2018 Jan 1;4(1):16-20.

**2017**

**Johnson SC**, Gonzalez B, Zhang Q, Milholland B, Zhang Z, Suh Y: Network analysis of mitonuclear GWAS reveals functional networks and tissue expression profiles of disease-associated genes. *Human Genetics*. 2017. 136: p55-65. PMID: 27704213

**Amlie-Lefond CM**, Pavlakis SG. Preventing cognitive decline in sickle cell disease: A good night's sleep. Neurology. 2017 Dec 12;89(24):2402-2403. doi: 10.1212/WNL.0000000000004754

Wallace AS, Hudac CM, **Steinman KJ**, Peterson JL, DesChamps TD, Duyzend MH, Nuttle X, Eichler EE, Bernier RA. Longitudinal report of child with de novo 16p11.2 triplication. Clin Case Rep. 2017 Dec 6;6(1):147-154. eCollection 2018 Jan. PMID: 29375855

**Gust J**, Hay KA, Hanafi LA, Li D, Myerson D, Gonzalez-Cuyar LF, Yeung C, Liles WC, Wurfel M, Lopez JA, Chen J, Chung D, Harju-Baker S, Özpolat T, Fink KR,Riddell SR, Maloney DG, Turtle CJ. Endothelial Activation and Blood-Brain Barrier Disruption in Neurotoxicity after Adoptive Immunotherapy with CD19 CAR-T Cells.Cancer Discov. 2017 Dec;7(12):1404-1419. doi: 10.1158/2159-8290.CD-17-0698.

Demarest ST, Shellhaas RA, Gaillard WD, Keator C, Nickels KC, Hussain SA, Loddenkemper T, Patel AD**, Saneto RP**, Wirrell E, Sánchez Fernández I, Chu CJ, Grinspan Z, Wusthoff CJ, Joshi S, Mohamed IS, Stafstrom CE, Stack CV, Yozawitz E, Bluvstein JS, Singh RK, Knupp KG; Pediatric Epilepsy Research Consortium. The impact of hypsarrhythmia on infantile spasms treatment response: Observational cohort study from the National Infantile Spasms Consortium. Epilepsia. 2017 Dec;58(12):2098-2103. doi: 10.1111/epi.13937.

Parikh S, Goldstein A, Karaa A, Koenig MK, Anselm I, Brunel-Guitton C,Christodoulou J, Cohen BH, Dimmock D, Enns GM, Falk MJ, Feigenbaum A, Frye RE,Ganesh J, Griesemer D, Haas R, Horvath R, Korson M, Kruer MC, Mancuso M,McCormack S, Raboisson MJ, Reimschisel T, Salvarinova R, **Saneto RP**, Scaglia F,Shoffner J, Stacpoole PW, Sue CM, Tarnopolsky M, Van Karnebeek C, Wolfe LA, Cunningham ZZ, Rahman S, Chinnery PF. Patient care standards for primary mitochondrial disease: a consensus statement from the Mitochondrial Medicine Society. Genet Med. 2017 Dec;19(12). doi: 10.1038/gim.2017.107.

Hay KA, Hanafi LA, Li D**, Gust J**, Liles WC, Wurfel MM, López JA, Chen J, Chung D, Harju-Baker S, Cherian S, Chen X, Riddell SR, Maloney DG, Turtle CJ. Kinetics and biomarkers of severe cytokine release syndrome after CD19 chimeric antigenreceptor-modified T-cell therapy. Blood. 2017 Nov 23;130(21):2295-2306. doi:10.1182/blood-2017-06-793141. Epub 2017 Sep 18. PubMed PMID: 28924019; PubMed

Farhadian S, **Patel P**, Spudich S. Neurological Complications of HIV Infection. Curr Infect Dis Rep. 2017 Nov 21;19(12):50. doi: 10.1007/s11908-017-0606-5.PMID: 29164407 Review. Global Health Neurology: HIV/AIDS

Shellhaas RA, Berg AT, Grinspan ZM, Wusthoff CJ, Millichap JJ, Loddenkemper T,Coryell J, **Saneto RP**, Chu CJ, Joshi SM, Sullivan JE, Knupp KG, Kossoff EH, Keator C, Wirrell EC, Mytinger JR, Valencia I, Massey S, Gaillard WD. Initial Treatment for Nonsyndromic Early-Life Epilepsy: An Unexpected Consensus. Pediatr Neurol.2017 Oct;75:73-79. doi: 10.1016/j.pediatrneurol.2017.06.011.

Kirk V, Baughn J, D'Andrea L, Friedman N, Galion A, Garetz S, Hassan F, **Wrede J,** Harrod CG, Malhotra RK. American Academy of Sleep Medicine Position Paper for the Use of a Home Sleep Apnea Test for the Diagnosis of OSA in Children. J Clin Sleep Med. 2017 Oct 15;13(10):1199-1203. doi: 0.5664/jcsm.6772.

Rilinger JF, Smith CM, deRegnier RAO, Goldstein JL, Mills MG, Reynolds M, Backer CL, Burrowes DM, Mehta P, Piantino J**, Wainwright MS**. Transcranial Doppler Identification of Neurologic Injury during Pediatric Extracorporeal Membrane Oxygenation Therapy. J Stroke Cerebrovasc Dis. 2017 Oct;26(10):2336-2345. doi:10.1016/j.jstrokecerebrovasdis.2017.05.022

De Mori R, Romani M, D'Arrigo S, Zaki MS, Lorefice E, Tardivo S, Biagini T,Stanley V, Musaev D, Fluss J, Micalizzi A, Nuovo S, Illi B, Chiapparini L, Di Marcotullio L, Issa MY, Anello D, Casella A, Ginevrino M, Leggins AS, Roosing S, Alfonsi R, Rosati J, Schot R, Mancini GMS, Bertini E, **Dobyns WB,** Mazza T, Gleeson JG, Valente EM. Hypomorphic Recessive Variants in SUFU Impair the Sonic Hedgehog Pathway and Cause Joubert Syndrome with Cranio-facial and Skeletal Defects. Am J Hum Genet. 2017 Oct 5;101(4):552-563. doi: 10.1016/j.ajhg.2017.08.017. Epub 2017 Sep 28

Alcantara D, Timms AE, Gripp K, Baker L, Park K, Collins S, Cheng C, Stewart F, Mehta SG, Saggar A, Sztriha L, Zombor M, Caluseriu O, Mesterman R, Van Allen MI, Jacquinet A, Ygberg S, Bernstein JA, Wenger AM, Guturu H, Bejerano G, Gomez-Ospina N, Lehman A, Alfei E, Pantaleoni C, Conti V, Guerrini R, Moog U, Graham JM Jr, Hevner R, **Dobyns WB**, O'Driscoll M, Mirzaa GM. Mutations of AKT3 are associated with a wide spectrum of developmental disorders including extreme megalencephaly. Brain. 2017 Oct 1;140(10):2610-2622. doi: 10.1093/brain/awx203

**Saneto RP**. Genetics of Mitochondrial Disease. Adv Genet. 2017;98:63-116. doi: 10.1016/bs.adgen.2017.06.002. Epub 2017 Sep

Berg AT, Coryell J, **Saneto RP**, Grinspan ZM, **Alexander JJ**, Kekis M, Sullivan JE, Wirrell EC, Shellhaas RA, Mytinger JR, Gaillard WD, Kossoff EH, Valencia I, Knupp KG, Wusthoff C, Keator C, **Dobyns WB**, Ryan N, Loddenkemper T, Chu CJ, **Novotny EJ Jr**, Koh S. Early-Life Epilepsies and the Emerging Role of Genetic Testing. JAMA Pediatr. 2017 Sep 1;171(9):863-871. doi:10.1001/jamapediatrics.2017.1743..

Rosenthal ES, Claassen J, **Wainwright MS**, Husain AM, Vaitkevicius H, Raines S, Hoffmann E, Colquhoun H, Doherty JJ, Kanes SJ. Brexanolone as adjunctive therapy in super-refractory status epilepticus. Ann Neurol. 2017 Sep;82(3):342-352. doi: 10.1002/ana.25008.

Syrbe S, Harms FL, Parrini E, Montomoli M, Mütze U, Helbig KL, Polster T, Albrecht B, Bernbeck U, van Binsbergen E, Biskup S, Burglen L, Denecke J, Heron B, Heyne HO, Hoffmann GF, Hornemann F, Matsushige T, Matsuura R, Kato M, Korenke GC, Kuechler A, Lämmer C, Merkenschlager A, Mignot C, Ruf S, Nakashima M, Saitsu H, Stamberger H, Pisano T, Tohyama J, Weckhuysen S, Werckx W, Wickert J, Mari F, Verbeek NE, Møller RS, Koeleman B, Matsumoto N, **Dobyns WB**, Battaglia D, Lemke JR,Kutsche K, Guerrini R. Delineating SPTAN1 associated phenotypes: from isolated epilepsy to encephalopathy with progressive brain atrophy. Brain. 2017 Sep 1;140(9):2322-2336.

Bonwitt J, Poel A, DeBolt C, Gonzales E, Lopez A, Routh J, Rietberg K, Linton N, Reggin J, Sejvar J, Lindquist S, **Otten CE**. Acute Flaccid Myelitis Among Children - Washington, September-November 2016. MMWR Morb Mortal Wkly Rep. 2017 Aug 11;66(31):826-829.

Oegema R, Baillat D, Schot R, van Unen LM, Brooks A, Kia SK, Hoogeboom AJM,Xia Z, Li W, Cesaroni M, Lequin MH, van Slegtenhorst M, **Dobyns WB**, de Coo IFM,van den Berg D, Verheijen FW, Kremer A, van der Spek PJ, Heijsman D, Wagner EJ,Fornerod M, Mancini GMS. Correction: Human mutations in integrator complex subunits link transcriptome integrity to brain development. PLoS Genet. 2017 Aug 1;13(8):e1006923. doi: 10.1371/journal.pgen.1006923. eCollection 2017 Aug. PubMed

McAdams RM, Fleiss B, Traudt C, Schwendimann L, Snyder JM, Haynes RL,**Natarajan N**, Gressens P, Juul SE. Long-Term Neuropathological Changes Associated with Cerebral Palsy in a Nonhuman Primate Model of Hypoxic-IschemicEncephalopathy. Dev Neurosci. 2017;39(1-4):124-140.

**Novotny EJ Jr**. Early genetic testing for neonatal epilepsy: When, why, and how? Neurology. 2017 Aug 29;89(9):880-881.

Platzer K, Yuan H, Schütz H, Winschel A, Chen W, Hu C, Kusumoto H, Heyne HO, Helbig KL, Tang S, Willing MC, Tinkle BT, Adams DJ, Depienne C, Keren B, Mignot C, Frengen E, Strømme P, Biskup S, Döcker D, Strom TM, Mefford HC, Myers CT, MuirAM, LaCroix A, Sadleir L, Scheffer IE, Brilstra E, van Haelst MM, van der Smagt JJ, Bok LA, Møller RS, Jensen UB, Millichap JJ, Berg AT, Goldberg EM, De Bie I,Fox S, Major P, Jones JR, Zackai EH, Abou Jamra R, Rolfs A, Leventer RJ, Lawson JA, Roscioli T, Jansen FE, Ranza E, Korff CM, Lehesjoki AE, Courage C, Linnankivi T, Smith DR, Stanley C, Mintz M, McKnight D, Decker A, Tan WH, Tarnopolsky MA,Brady LI, Wolff M, Dondit L, Pedro HF, Parisotto SE, Jones KL, Patel AD, Franz DN, Vanzo R, Marco E, Ranells JD, Di Donato N**, Dobyns WB**, Laube B, Traynelis SF, Lemke JR. GRIN2B encephalopathy: novel findings on phenotype, variant clustering,functional consequences and treatment aspects. J Med Genet. 2017Jul;54(7):460-470. doi: 10.1136/jmedgenet-2016-104509.

Van De Weghe JC, Rusterholz TDS, Latour B, Grout ME, Aldinger KA, Shaheen R, Dempsey JC, Maddirevula S, Cheng YH, Phelps IG, Gesemann M, Goel H, Birk OS,Alanzi T, Rawashdeh R, Khan AO; University of Washington Center for MendelianGenomics, Bamshad MJ, Nickerson DA, Neuhauss SCF, **Dobyns WB**, Alkuraya FS, RoepmanR, Bachmann-Gagescu R, Doherty D. Mutations in ARMC9, which Encodes a Basal Body Protein, Cause Joubert Syndrome in Humans and Ciliopathy Phenotypes in Zebrafish.Am J Hum Genet. 2017 Jul 6;101(1):23-36. doi: 10.1016/j.ajhg.2017.05.010. Epub

2017 Jun 15. PubMed PMID: 28625504; PubMed Central PMCID: PMC5501774.

AJ, Guerrini R, **Dobyns WB**. Lissencephaly: Expanded imaging and clinical classification. Am J Med Genet A. 2017 Jun;173(6):1473-1488. doi:10.1002/ajmg.a.38245. Epub 2017 Apr 25.

Keros S, Buraniqi E, Alex B, Antonetty A, Fialho H, Hafeez B, Jackson MC, Jawahar R, Kjelleren S, Stewart E, **Morgan LA**, **Wainwright MS**, Sogawa Y, Patel AD, Loddenkemper T, Grinspan ZM. Increasing Ketamine Use for Refractory Status Epilepticus in US Pediatric Hospitals. J Child Neurol. 2017 Jun;32(7):638-646.doi: 10.1177/0883073817698629

Garavelli L, Ivanovski I, Caraffi SG, Santodirocco D, Pollazzon M, Cordelli DM, Abdalla E, Accorsi P, Adam MP, Baldo C, Bayat A, Belligni E, Bonvicini F, Breckpot J, Callewaert B, Cocchi G, Cuturilo G, Devriendt K, Dinulos MB, Djuric O, Epifanio R, Faravelli F, Formisano D, Giordano L, Grasso M, Grønborg S, Iodice A, Iughetti L, Lacombe D, Maggi M, Malbora B, Mammi I, Moutton S, Møller R, Muschke P, Napoli M, Pantaleoni C, Pascarella R, Pellicciari A, Poch-Olive ML,Raviglione F, Rivieri F, Russo C, Savasta S, Scarano G, Selicorni A, Silengo M, Sorge G, Tarani L, Tone LG, Toutain A, Trimouille A, Valera ET, Vergano SS, Zanotta N, Zollino M, **Dobyns WB,** Paciorkowski AR. Neuroimaging findings in Mowat-Wilson syndrome: a study of 54 patients. Genet Med. 2017 Jun;19(6):691-700. doi: 10.1038/gim.2016.176. Epub 2016 Nov 10

Kurz JE, Smith CM, **Wainwright MS.** Thermoregulate, autoregulate and ventilate: brain-directed critical care for pediatric cardiac arrest. Curr Opin Pediatr.2017 Jun;29(3):259-265. Doi. Di Donato N, Chiari S, Mirzaa GM, Aldinger K, Parrini E, Olds C, Barkovich AJ, Guerrini R, **Dobyns WB**. Lissencephaly: Expanded imaging and clinical classification. Am J Med Genet A. 2017 Jun;173(6):1473-1488. doi:10.1002/ajmg.a.38245. Epub 2017 Apr 250.1097/MOP.0000000000000482:.

**Marashly A**, Lew S, Koop J. Successful surgical management of New Onset Refractory Status Epilepticus (NORSE) presenting with gelastic seizures in a 3 year old girl. Epilepsy Behav Case Rep. 2017 May 22;8:18-26. doi: 10.1016/

Ducharme-Crevier L, Press CA, Kurz JE, Mills MG, Goldstein JL, **Wainwright MS.** Early Presence of Sleep Spindles on Electroencephalography Is Associated With Good Outcome After Pediatric Cardiac Arrest. Pediatr Crit Care Med. 2017 May;18(5):452-460. doi: 10.1097/PCC.0000000000001137

Dempsey JC, Phelps IG, Bachmann-Gagescu R, Glass IA, **Tully HM,** Doherty D.Mortality in Joubert syndrome. Am J Med Genet A. 2017 May;173(5):1237-1242. doi: 10.1002/ajmg.a.38158.

Oegema R, Baillat D, Schot R, van Unen LM, Brooks A, Kia SK, Hoogeboom AJM, Xia Z, Li W, Cesaroni M, Lequin MH, van Slegtenhorst M, **Dobyns WB**, de Coo IFM, Verheijen FW, Kremer A, van der Spek PJ, Heijsman D, Wagner EJ, Fornerod M, Mancini GMS. Human mutations in integrator complex subunits link transcriptome integrity to brain development. PLoS Genet. 2017 May 25;13(5):e1006809. doi: 10.1371/journal.pgen.1006809. eCollection 2017 May. Erratum in: PLoS Genet. 2017 Aug 1;13(8):e1006923..

**Carter JC**, **Wrede JE**. Overview of Sleep and Sleep Disorders in Infancy and Childhood. Pediatr Ann. 2017 Apr 1;46(4):e133-e138.

Mahdi J, Shah AC, **Sato A,** Morris SM, McKinstry RC, Listernick R, Packer RJ,Fisher MJ, Gutmann DH. A multi-institutional study of brainstem gliomas in children with neurofibromatosis type 1. Neurology. 2017 Apr 18;88(16):1584-1589.

**Blume HK**. Neurological Disorders in Primary Care Pediatrics. Pediatr Ann. 2017 Apr 1;46(4):e131-e132.

Lee S, Mirsky DM, Beslow LA, **Amlie-Lefond C,** Danehy AR, Lehman L, Stence NV, Vossough A, Wintermark M, Rivkin MJ; International Paediatric Stroke Study Neuroimaging Consortium and the Paediatric Stroke Neuroimaging Consortium. Pathways for Neuroimaging of Neonatal Stroke. Pediatr Neurol. 2017 Apr;69:37-48.doi:10.1016/j.pediatrneurol.2016.12.008.

**Gospe SM Jr**. Pyridoxine-Dependent Epilepsy. 2001 Dec 7 [updated 2017 Apr 13]. In: Adam MP, Ardinger HH, Pagon RA, Wallace SE, Bean LJH, Stephens K, Amemiya A, editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2018.

**Randle SC**. Tuberous Sclerosis Complex: A Review. Pediatr Ann. 2017 Apr 1;46(4):e166-e171.

Moore CA, Staples JE, **Dobyns WB**, Pessoa A, Ventura CV, Fonseca EB, RibeiroEM, Ventura LO, Neto NN, Arena JF, Rasmussen SA. Characterizing the Pattern of Anomalies in Congenital Zika Syndrome for Pediatric Clinicians. JAMA Pediatr. 2017 Mar 1;171(3):288-295. doi: 10.1001/jamapediatrics.2016.3982. Review

**Natarajan N**, Pardo AC. Challenges in neurologic prognostication after neonatal brain injury. Semin Perinatol. 2017 Mar;41(2):117-123. doi:10.1053/j.semperi.2016.11.008. Epub 2017 Jan 28. Review. PubMed PMID: 28139254.

**Amlie-Lefond C**, Ojemann JG. Neonatal Hemorrhagic Stroke. JAMA Pediatr. 2017 Mar 1;171(3):220-221. doi: 10.1001/jamapediatrics.2016.4466

Lardelli RM, Schaffer AE, Eggens VR, Zaki MS, Grainger S, Sathe S, Van Nostrand EL, Schlachetzki Z, Rosti B, Akizu N, Scott E, Silhavy JL, Heckman LD, Rosti RO, Dikoglu E, Gregor A, Guemez-Gamboa A, Musaev D, Mande R, Widjaja A,Shaw TL, Markmiller S, Marin-Valencia I, Davies JH, de Meirleir L, Kayserili H, Altunoglu U, Freckmann ML, Warwick L, Chitayat D, Blaser S, Çağlayan AO, Bilguvar K, Per H, Fagerberg C, Christesen HT, Kibaek M, Aldinger KA, Manchester D, Matsumoto N, Muramatsu K, Saitsu H, Shiina M, Ogata K, Foulds N, **Dobyns WB**, Chi NC, Traver D, Spaccini L, Bova SM, Gabriel SB, Gunel M, Valente EM, Nassogne MC, Bennett EJ, Yeo GW, Baas F, Lykke-Andersen J, Gleeson JG. Biallelic mutations in the 3' exonuclease TOE1 cause pontocerebellar hypoplasia and uncover a role in snRNA processing. Nat Genet. 2017 Mar;49(3):457-464. doi: 10.1038/ng.3762. Epub 2017 Jan 16.

Bordini BJ, **Monrad P.** Differentiating Familial Neuropathies from Guillain-Barré Syndrome.

Pediatr Clin North Am. 2017 Feb;64(1):231-252.

McDonald CL, **Saneto RP**, Carmant L, Sotero de Menezes MA. Focal Seizures in Patients With SCN1A Mutations. J Child Neurol. 2017 Feb;32(2):170-176. doi: 10.1177/0883073816672379

**Beatty CW, Wrede JE, Blume HK.** Diagnosis, treatment, and outcomes of infantile spasms in the Trisomy 21 population. Seizure. 2017 Feb;45:184-188. doi: 10.1016/j.seizure.2016.12.016..

Tuschl K, Clayton PT, **Gospe SM Jr**, Mills PB. Dystonia/Parkinsonism, Hypermanganesemia, Polycythemia, and Chronic Liver Disease. 2012 Aug 30 [updated 2017 Feb 9]. In: Adam MP, Ardinger HH, Pagon RA, Wallace SE, Bean LJH, Stephens K, Amemiya A, editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2018.

Wenger TL, Chow P, **Randle SC**, Rosen A, Birgfeld C, **Wrede J**, Javid P, King D, Manh V, Hing AV, Albers E.Novel findings of left ventricular non-compaction cardiomyopathy, microform cleft lip and poor vision in patient with SMC1A-associated Cornelia de Lange syndrome. Am J Med Genet A. 2017 Feb;173(2):414-420.

Piao CS, Holloway AL, Hong-Routson S, **Wainwright MS**. Depression following traumatic brain injury in mice is associated with down-regulation of hippocampal astrocyte glutamate transporters by thrombin. J Cereb Blood Flow Metab. 2017 Jan 1:271678X17742792. doi: 10.

Brun BN, Mockler SR, Laubscher KM, Stephan CM, Wallace AM, Collison JA,Zimmerman MB, **Dobyns WB**, Mathews KD. Comparison of brain MRI findings with language and motor function in the dystroglycanopathies. Neurology. 2017 Feb 14;88(7):623-629. doi: 10.1212/WNL.0000000000003609. Epub 2017 Jan 13. .

Kannan N, Quistberg A, Wang J, Groner JI, Mink RB, **Wainwright MS**, Bell MJ, Giza CC, Zatzick DF, Ellenbogen RG, Boyle LN, Mitchell PH, Vavilala MS. Frequency of and factors associated with emergency department intracranial pressure monitor placement in severe paediatric traumatic brain injury. Brain Inj.

2017;31(13-14):1745-1752. doi: 10.1080/02699052.2017.

Gorman KM, **Wainwright MS**. Adult Stroke Screening Tool in Childhood IschemicStroke. Pediatr Neurol Briefs. 2017 Jan;31(1):3. doi:10.15844/pedneurbriefs-31-1-3.

Sánchez Fernández I, Jackson MC, Abend NS, Arya R, Brenton JN, Carpenter JL, Chapman KE, Gaillard WD, Gaínza-Lein M, Glauser TA, Goldstein JL, Goodkin HP,Helseth A, Kapur K, McDonough TL, Mikati MA, Peariso K, Riviello J Jr, Tasker RC,Topjian AA**, Wainwright MS**, Wilfong A, Williams K, Loddenkemper T; pediatric Status Epilepticus Research Group (pSERG). Refractory status epilepticus in children with and without prior epilepsy or status epilepticus. Neurology. 2017 Jan 24;88(4):386-394. doi: 10.1212/WNL.0000000000003550.

Press CA, Morgan L, Mills M, Stack CV, Goldstein JL, Alonso EM, **Wainwright MS**. Spectral Electroencephalogram Analysis for the Evaluation of Encephalopathy Grade in Children With Acute Liver Failure. Pediatr Crit Care Med. 2017 Jan;18(1):64-72. doi: 10.1097/PCC.0000000000001016

Haldipur P, Dang D, Aldinger KA, Janson OK, Guimiot F, Adle-Biasette H, **Dobyns WB**, Siebert JR, Russo R, Millen KJ. Phenotypic outcomes in Mouse and HumanFoxc1 dependent Dandy-Walker cerebellar malformation suggest shared mechanisms.Elife. 2017 Jan 16;6. pii: e20898. doi: 10.7554/eLife.20898

Byers HM, Adam MP, LaCroix A, Leary SE, Cole B, **Dobyns W**B, Mefford HC.Description of a new oncogenic mechanism for atypical teratoid rhabdoid tumors in patients with ring chromosome 22. Am J Med Genet A. 2017 Jan;173(1):245-249. doi: 10.1002/ajmg.a.37993. Epub 2016 Oct 12

**Patel, P**., Baier J, Baranov, et al. Health Beliefs Regarding Pedatric Cerebral Palsy Among Caregivers in Botswana: A Qualitative Study. Child Care Health Dev. 43(6):861-868, 2017

Farhadian S., **Patel P.,** Spudich S., Neurological Complications of HIV. Curr Infect Dis Rep. 19(12): 50, 2017

**2016**

Ducharme-Crevier L, Mills MG, Mehta PM, Smith CM, **Wainwright MS**. Use of Transcranial Doppler for Management of Central Nervous System Infections in Critically Ill Children. Pediatr Neurol. 2016 Dec;65:52-58.e2. doi: 10.1016/j.pediatrneurol.2016.08.027.

**Johnson SC**, Kaeberlein M: Rapamycin in aging and disease: maximizing efficacy while minimizing side effects. *Oncotarget.* 2016; 7:p44876-44878. PMID: 27384492.

Teumer A, Qi Q, Nethander M, Aschard H, Bandinelli S, Beekman M, Berndt SI, Bidlingmaier M, Broer L; CHARGE Longevity Working Group, Cappola A, Ceda GP, Chanock S, Chen MH, Chen TC, Chen YI, Chung J, Del Greco Miglianico F, Eriksson J, Ferrucci L, Friedrich N, Gnewuch C, Goodarzi MO, Grarup N, Guo T, Hammer E, Hayes RB, Hicks AA, Hofman A, Houwing-Duistermaat JJ, Hu F, Hunter DJ, Husemoen LL, Isaacs A, Jacobs KB, Janssen JA, Jansson JO, Jehmlich N, **Johnson S**, Juul A, Karlsson M, Kilpelainen TO, Kovacs P, Kraft P, Li C, Linneberg A, Liu Y, Loos RJ; Body Composition Genetics Consortium, Lorentzon M, Lu Y, Maggio M, Magi R, Meigs J, Mellström D, Nauck M, Newman AB, Pollak MN, Pramstaller PP, Prokopenko I, Psaty BM, Reincke M, Rimm EB, Rotter JI, Saint Pierre A, Schurmann C, Seshadri S, Sjögren K, Slagboom PE, Strickler HD, Stumvoll M, Suh Y, Sun Q, Zhang C, Svensson J, Tanaka T, Tare A, Tönjes A, Uh HW, van Duijn CM, van Heemst D, Vandenput L, Vasan R, Völker U, Willems SM, Ohlsson C, Wallaschofski H, Kaplan R: Genomewide meta-analysis identifies loci associated with IGF-I and IGFBP-3 levels with impact on age-related traits. *Aging Cell.* 2016. 15(5):p811-24. PMID: 27329260.

Mohammad S, Wolfe LA, Stöbe P, Biskup S, **Wainwright MS**, Melin-Aldana H, Malladi P, Muenke M, Gahl WA, Whitington PF. Infantile Cirrhosis, Growth Impairment, and Neurodevelopmental Anomalies Associated with Deficiency of PPP1R15B. J Pediatr. 2016 Dec;179:144-149.e2. doi: 0.1016/j.jpeds.2016.08.043

Leto K, Arancillo M, Becker EB, Buffo A, Chiang C, Ding B, **Dobyns WB**, Dusart I, Haldipur P, Hatten ME, Hoshino M, Joyner AL, Kano M, Kilpatrick DL, Koibuchi N, Marino S, Martinez S, Millen KJ, Millner TO, Miyata T, Parmigiani E, Schilling K, Sekerková G, Sillitoe RV, Sotelo C, Uesaka N, Wefers A, Wingate RJ, Hawkes R. Consensus Paper: Cerebellar Development. Cerebellum. 2016 Dec;15(6):789-828.Review

Rivkin MJ, Bernard TJ, Dowling MM, **Amlie-Lefond C**. Corrigendum to 'Guidelines for Urgent Management of Stroke in Children' [Pediatric Neurology 56 (2016)8-17]. Pediatr Neurol. 2016 Nov;64:105. doi: 10.1016/j.pediatrneurol.2016.08.019.

Epub 2016 Sep 1.

**Steinman KJ,** Spence SJ, Ramocki MB, Proud MB, Kessler SK, Marco EJ, Green Snyder L, D'Angelo D, Chen Q, Chung WK, Sherr EH; Simons VIP Consortium. 16p11.2 deletion and duplication: Characterizing neurologic phenotypes in a large clinically ascertained cohort. Am J Med Genet A. 2016 Nov;170(11):2943-2955. doi:10.1002/ajmg.a.37820.

Kim J, Liao YH, **Ionita C**, Bale AE, Darras B, Acsadi G. Mitochondrial Membrane Protein-Associated Neurodegeneration Mimicking Juvenile AmyotrophicLateral Sclerosis. Pediatr Neurol. 2016 Nov;64:83-86. doi:10.1016/j.pediatrneurol.2016.08.013.

Ravenscroft G, Di Donato N, Hahn G, Davis MR, Craven PD, Poke G, Neas KR,Neuhann TM, **Dobyns WB**, Laing NG. Recurrent de novo BICD2 mutation associated witharthrogryposis multiplex congenita and bilateral perisylvian polymicrogyria.Neuromuscul Disord. 2016 Nov;26(11):744-748. doi: 10.1016/j.nmd.2016.09.009. Epub

2016 Sep 19.

Adams Waldorf KM, Stencel-Baerenwald JE, Kapur RP, Studholme C, Boldenow E,Vornhagen J, Baldessari A, Dighe MK, Thiel J, Merillat S, Armistead B,Tisoncik-Go J, Green RR, Davis MA, Dewey EC, Fairgrieve MR, Gatenby JC, Richards T, Garden GA, Diamond MS, Juul SE, Grant RF, Kuller L, Shaw DW, Ogle J, Gough GM,Lee W, English C, Hevner RF, **Dobyns WB,** Gale M Jr, Rajagopal L. Fetal brain lesions after subcutaneous inoculation of Zika virus in a pregnant nonhuman primate. Nat Med. 2016 Nov;22(11):1256-1259. doi: 10.1038/nm.4193. Epub 2016 Sep

Ma M, Adams HR, Seltzer LE, **Dobyns WB**, Paciorkowski AR. Phenotype Differentiation of FOXG1 and MECP2 Disorders: A New Method for Characterization of Developmental Encephalopathies. J Pediatr. 2016 Nov;178:233-240.e10. doi:10.1016/j.jpeds.2016.08.032. Epub 2016 Sep 15.

Di Donato N, Jean YY, Maga AM, Krewson BD, Shupp AB, Avrutsky MI, Roy A, Collins S, Olds C, Willert RA, Czaja AM, Johnson R, Stover JA, Gottlieb S,Bartholdi D, Rauch A, Goldstein A, Boyd-Kyle V, Aldinger KA, Mirzaa GM, Nissen A, Brigatti KW, Puffenberger EG, Millen KJ, Strauss KA, **Dobyns WB**, Troy CM, Jinks RN. Mutations in CRADD Result in Reduced Caspase-2-Mediated Neuronal Apoptosis and Cause Megalencephaly with a Rare Lissencephaly Variant. Am J Hum Genet. 2016 Nov 3;99(5):1117-1129. doi: 10.1016/j.ajhg.2016.09.010. Epub 2016 Oct 20

**Tully HM**, Wenger TL, Kukull WA, Doherty D, **Dobyns WB**. Anatomical configurations associated with posthemorrhagic hydrocephalus among premature infants with intraventricular hemorrhage. Neurosurg Focus. 2016 Nov;41(5):E5.

Garzon MC, Epstein LG, Heyer GL, Frommelt PC, Orbach DB, Baylis AL, Blei F, Burrows PE, Chamlin SL, Chun RH, Hess CP, Joachim S, Johnson K, Kim W, Liang MG, Maheshwari M, McCoy GN, Metry DW, **Monrad PA**, Pope E, Powell J, Shwayder TA, Siegel DH, Tollefson MM, Vadivelu S, Lew SM, Frieden IJ, Drolet BA. PHACE Syndrome: Consensus-Derived Diagnosis and Care Recommendations. J Pediatr. 2016 Nov;178:24-33.e2.

Bernard TJ, Beslow LA, Manco-Johnson MJ, Armstrong-Wells J, Boada R, Weitzenkamp D, Hollatz A, Poisson S, **Amlie-Lefond C**, Lo W, deVeber G, Goldenberg NA, Dowling MM, Roach ES, Fullerton HJ, Benseler SM, Jordan LC, Kirton A, Ichord RN. Inter-Rater Reliability of the CASCADE Criteria: Challenges in Classifying Arteriopathies. Stroke. 2016 Oct;47(10):2443-9. doi:10.1161/STROKEAHA.116.013544. Epub 2016 Sep 15.

Tasker RC, Goodkin HP, Sánchez Fernández I, Chapman KE, Abend NS, Arya R,Brenton JN, Carpenter JL, Gaillard WD, Glauser TA, Goldstein J, Helseth AR,Jackson MC, Kapur K, Mikati MA, Peariso K, **Wainwright MS**, Wilfong AA, Williams K,Loddenkemper T; Pediatric Status Epilepticus Research Group. Refractory Status Epilepticus in Children: Intention to Treat With Continuous Infusions of Midazolam and Pentobarbital. Pediatr Crit Care Med. 2016 Oct;17(10):968-975

Buckley RT, Wang AC, Miller JW, **Novotny EJ**, Ojemann JG. Stereotactic laser ablation for hypothalamic and deep intraventricular lesions. Neurosurg Focus.2016 Oct;41(4):E10.

Ostahowski PJ, Kannan N, **Wainwright MS**, Qiu Q, Mink RB, Groner JI, Bell MJ,Giza CC, Zatzick DF, Ellenbogen RG, Boyle LN, Mitchell PH, Vavilala MS; PEGASUS (Pediatric Guideline Adherence and Outcomes) Study. Variation in seizure prophylaxis in severe pediatric traumatic brain injury. J Neurosurg Pediatr. 2016 Oct;18(4):499-506..

Weisleder P, **Gospe SM Jr**, Ng YT, Sahin M. The Pediatric Neurology Trainee Publication Award for 2015. Pediatr Neurol. 2016 Oct;63:1-2. doi: 10.1016/j.pediatrneurol.2016.07.004.

Gripp KW, Aldinger KA, Bennett JT, Baker L, Tusi J, Powell-Hamilton N,Stabley D, Sol-Church K, Timms AE, **Dobyns WB**. A novel rasopathy caused by recurrent de novo missense mutations in PPP1CB closely resembles Noonan syndrome with loose anagen hair. Am J Med Genet A. 2016 Sep;170(9):2237-47. doi:10.1002/ajmg.a.37781.

Parrini E, Conti V, **Dobyns WB,** Guerrini R. Genetic Basis of Brain Malformations. Mol Syndromol. 2016 Sep;7(4):220-233. Epub 2016 Aug 27. Review.

**Natarajan N, Tully HM**, Chapman T. Prenatal presentation of pyruvate dehydrogenase complex deficiency. Pediatr Radiol. 2016 Aug; 46(9):1354-7.

Tuschl K, Clayton PT, **Gospe SM Jr**, Gulab S, Ibrahim S, Singhi P, Aulakh R, Ribeiro RT, Barsottini OG, Zaki MS, Del Rosario ML, Dyack S, Price V, Rideout A, Gordon K, Wevers RA, Chong WK, Mills PB. Syndrome of Hepatic Cirrhosis, Dystonia,Polycythemia, and Hypermanganesemia Caused by Mutations in SLC30A10, a Manganese Transporter in Man. Am J Hum Genet. 2016 Aug 4;99(2):521. doi: 10.1016/j.ajhg.2016.07.015.

Tsuchida TN, Acharya JN, Halford JJ**, Kuratani JD,** Sinha SR, Stecker MM, Tatum WO 4th, Drislane FW. American Clinical Neurophysiology Society: EEG Guidelines Introduction. J Clin Neurophysiol. 2016 Aug;33(4):301-2. doi: 10.1097/WNP.0000000000000315.

Mossa-Basha M, de Havenon A, Becker KJ, Hallam DK, Levitt MR, Cohen WA, Hippe DS, Alexander MD, Tirschwell DL, Hatsukami T, **Amlie-Lefond C**, Yuan C. Added Value of Vessel Wall Magnetic Resonance Imaging in the Differentiation of Moyamoya Vasculopathies in a Non-Asian Cohort. Stroke. 2016 Jul;47(7):1782-8. doi: 10.1161/STROKEAHA.116.013320

.

**Amlie-Lefond C**, Gilden D. Varicella Zoster Virus: A Common Cause of Stroke in Children and Adults. J Stroke Cerebrovasc Dis. 2016 Jul;25(7):1561-1569. doi:10.1016/j.jstrokecerebrovasdis.2016.03.052. Epub 2016 Apr 29. Review

Mirzaa GM, Campbell CD, Solovieff N, Goold C, Jansen LA, Menon S, Timms AE, Conti V, Biag JD, Adams C, Boyle EA, Collins S, Ishak G, Poliachik S, Girisha KM,Yeung KS, Chung BHY, Rahikkala E, Gunter SA, McDaniel SS, Macmurdo CF, Bernstein JA, Martin B, Leary R, Mahan S, Liu S, Weaver M, Doerschner M, Jhangiani S, MuznyDM, Boerwinkle E, Gibbs RA, Lupski JR, Shendure J, **Saneto RP,** **Novotny EJ**, Wilson CJ, Sellers WR, Morrissey M, Hevner RF, Ojemann JG, Guerrini R, Murphy LO, Winckler W, **Dobyns WB**. Association of MTOR Mutations With Developmental Brain Disorders, Including Megalencephaly, Focal Cortical Dysplasia, and Pigmentary Mosaicism. JAMA Neurol. 2016 Jul 1;73(7):836-845. doi: 10.1001/jamaneurol.2016.0363.

Casimo K, Darvas F, Wander J, Ko A, Grabowski TJ, **Novotny E**, Poliakov A, Ojemann JG, Weaver KE. Regional Patterns of Cortical Phase Synchrony in the Resting State. Brain Connect. 2016 Jul;6(6):470-81. doi: 10.1089/brain.2015.0362. Epub 2016 May 2.

Parikh S, Karaa A, Goldstein A, Ng YS, Gorman G, Feigenbaum A, Christodoulou J, Haas R, Tarnopolsky M, Cohen BK, Dimmock D, Feyma T, Koenig MK, Mundy H,Niyazov D, **Saneto RP, Wainwright MS**, Wusthoff C, McFarland R, Scaglia F. Solid organ transplantation in primary mitochondrial disease: Proceed with caution. Mol Genet Metab. 2016 Jul;118(3):178-84. doi: 10.1016/j.ymgme.2016.04.009

**Saneto RP**. Alpers-Huttenlocher syndrome: the role of a multidisciplinary health care team. J Multidiscip Healthc. 2016 Jul 26;9:323-33. doi:10.2147/JMDH.S84900. eCollection 2016. Review.

Kurz JE, Poloyac SM, Abend NS, Fabio A, Bell MJ, **Wainwright MS**; Investigators for the Approaches and Decisions in Acute Pediatric TBI Trial. Variation in Anticonvulsant Selection and Electroencephalographic Monitoring Following Severe Traumatic Brain Injury in Children-Understanding Resource Availability in Sites Participating in a Comparative Effectiveness Study. Pediatr Crit Care Med. 2016 Jul;17(7):649-57. doi: 10.1097/PCC.0000000000000765.

Byers HM**, Beatty CW**, Hahn SH**, Gospe SM Jr**. Dramatic Response After Lamotrigine in a Patient With Epileptic Encephalopathy and a De NovoCACNA1A Variant. Pediatr Neurol. 2016 Jul;60:79-82. doi: 10.1016/j.pediatrneurol.2016.03.012. Epub 2016 Apr 1.

Twigg SRF, Hufnagel RB, Miller KA, Zhou Y, McGowan SJ, Taylor J, Craft J,Taylor JC, Santoro SL, Huang T, Hopkin RJ, Brady AF, Clayton-Smith J, Clericuzio CL, Grange DK, Groesser L, Hafner C, Horn D, Temple IK, **Dobyns WB,** Curry CJ, Jones MC, Wilkie AOM. A Recurrent Mosaic Mutation in SMO, Encoding the Hedgehog Signal Transducer Smoothened, Is the Major Cause of Curry-Jones Syndrome. Am J Hum Genet. 2016 Jun 2;98(6):1256-1265. doi: 10.1016/j.ajhg.2016.04.007. Epub 2016 May 26.

Mirzaa G, Timms AE, Conti V, Boyle EA, Girisha KM, Martin B, Kircher M, Olds C, Juusola J, Collins S, Park K, Carter M, Glass I, Krägeloh-Mann I, Chitayat D, Parikh AS, Bradshaw R, Torti E, Braddock S, Burke L, Ghedia S, Stephan M, StewartF, Prasad C, Napier M, Saitta S, Straussberg R, Gabbett M, O'Connor BC, Keegan CE, Yin LJ, Lai AH, Martin N, McKinnon M, Addor MC, Boccuto L, Schwartz CE,Lanoel A, Conway RL, Devriendt K, Tatton-Brown K, Pierpont ME, Painter M, Worgan L, Reggin J, Hennekam R, Tsuchiya K, Pritchard CC, Aracena M, Gripp KW, Cordisco M, Van Esch H, Garavelli L, Curry C, Goriely A, Kayserilli H, Shendure J, Graham J Jr, Guerrini R, **Dobyns WB.** PIK3CA-associated developmental disorders exhibit distinct classes of mutations with variable expression and tissue distribution. JCI Insight. 2016 Jun 16;1(9). pii: e87623.

Di Donato N, Neuhann T, Kahlert AK, Klink B, Hackmann K, Neuhann I, Novotna B, Schallner J, Krause C, Glass IA, Parnell SE, Benet-Pages A, Nissen AM, Berger W, Altmüller J, Thiele H, Weber BH, Schrock E, **Dobyns WB**, Bier A, Rump A. Mutations in EXOSC2 are associated with a novel syndrome characterised by retinitis pigmentosa, progressive hearing loss, premature ageing, short stature, mild intellectual disability and distinctive gestalt. J Med Genet. 2016

Jun;53(6):419-25. doi: 10.1136/jmedgenet-2015-103511.

Aldinger KA, Mendelsohn NJ, Chung BH, Zhang W, Cohn DH, Fernandez B, Alkuraya FS, **Dobyns WB**, Curry CJ. Variable brain phenotype primarily affects the brainstem and cerebellum in patients with osteogenesis imperfecta caused by recessive WNT1 mutations. J Med Genet. 2016 Jun;53(6):427-30. doi:10.1136/jmedgenet-2015-103476

**Tully HM**, Kukull WA, Mueller BA. Clinical and Surgical Factors Associated With Increased Epilepsy Risk in Children With Hydrocephalus. Pediatr Neurol. 2016 Jun;59:18-22. doi: 10.1016/j.pediatrneurol.2016.02.011

Chi DL, Momany ET, Mancl LA, Lindgren SD, Zinner SH, **Steinman KJ.** Dental Homes for Children With Autism: A Longitudinal Analysis of Iowa Medicaid's I-Smile Program. Am J Prev Med. 2016 May;50(5):609-615. doi:10.1016/j.amepre.2015.08.022. Epub 2015 Oct 26.

Hong CS, Wang AC, Bonow RH, Abecassis IJ, **Amlie-Lefond C**, Ellenbogen RG. Moyamoya Disease in a Patient with VACTERL Association. World Neurosurg. 2016 May;89:729.e7-729.e10. doi: 10.1016/j.wneu.2016.01.059. Epub 2016 Feb 2.

Bernier R, **Steinman KJ**, Reilly B, Wallace AS, Sherr EH, Pojman N, Mefford HC, Gerdts J, Earl R, Hanson E, Goin-Kochel RP, Berry L, Kanne S, Snyder LG, Spence S, Ramocki MB, Evans DW, Spiro JE, Martin CL, Ledbetter DH, Chung WK; Simons VIP consortium. Clinical phenotype of the recurrent 1q21.1 copy-number variant. Genet Med. 2016 Apr;18(4):341-9. doi: 10.1038/gim.2015.78.

Alqahtani MF, Smith CM, Weiss SL, Dawson S, Ralay Ranaivo H, **Wainwright MS**. Evaluation of New Diagnostic Biomarkers in Pediatric Sepsis: Matrix Metalloproteinase-9, Tissue Inhibitor of Metalloproteinase-1, Mid-Regional Pro-Atrial Natriuretic Peptide, and Adipocyte Fatty-Acid Binding Protein. PLoS One. 2016 Apr 18;11(4):e0153645. doi: 10.1371/journal.pone.0153645. eCollection 2016..

**Lockrow J**, Longstreth W, Davis AP. Intracranial Aneurysms From Presumed Infective Endocarditis: The Dilemma of Persistently Negative Cultures. Neurohospitalist. 2016 Apr;6(2):80-6.. Epub 2015 Sep 21

Bernard TJ, Friedman NR, Stence NV, Jones W, Ichord R, **Amlie-Lefond C**, Dowling MM, Rivkin MJ. Preparing for a "Pediatric Stroke Alert". Pediatr Neurol. 2016 Mar;56:18-24. doi: 10.1016/j.pediatrneurol.2015.10.012. Epub 2015 Dec 11.Review.

Bennett JT, Tan TY, Alcantara D, Tétrault M, Timms AE, Jensen D, Collins S, Nowaczyk MJM, Lindhurst MJ, Christensen KM, Braddock SR, Brandling-Bennett H, Hennekam RCM, Chung B, Lehman A, Su J, Ng S, Amor DJ; University of Washington Center for Mendelian Genomics; Care4Rare Canada Consortium, Majewski J, Biesecker LG, Boycott KM, **Dobyns WB**, O'Driscoll M, Moog U, McDonell LM. Mosaic Activating Mutations in FGFR1 Cause cephalocraniocutaneous Lipomatosis. Am J Hum Genet.2016 Mar 3;98(3):579-587. doi: 10.1016/j.ajhg.2016.02.006

Rivkin MJ, Bernard TJ, Dowling MM, **Amlie-Lefond C**. Guidelines for Urgent Management of Stroke in Children. Pediatr Neurol. 2016 Mar;56:8-17. doi:10.1016/j.pediatrneurol.2016.01.016. Epub 2016 Jan 21. Review. Erratum in:Pediatr Neurol. 2016 Nov;64:105.

**Tully HM**, Ishak GE, Rue TC, Dempsey JC, Browd SR, Millen KJ, Doherty D, **Dobyns WB**. Two Hundred Thirty-Six Children With Developmental Hydrocephalus: Causes and Clinical Consequences. J Child Neurol. 2016 Mar;31(3):309-20. doi:10.1177/0883073815592222

Cohen AS, Yap DB, Lewis ME, Chijiwa C, Ramos-Arroyo MA, Tkachenko N, Milano V, Fradin M, McKinnon ML, Townsend KN, Xu J, Van Allen MI, Ross CJ, **Dobyns WB,**Weaver DD, Gibson WT. Weaver Syndrome-Associated EZH2 Protein Variants Show Impaired Histone Methyltransferase Function In Vitro. Hum Mutat. 2016 Mar;37(3):301-7. doi: 10.1002/humu.22946. Epub 2016 Jan 12.

.

Di Donato N, Rump A, Mirzaa GM, Alcantara D, Oliver A, Schrock E, **Dobyns WB**, O'Driscoll M. Identification and Characterization of a Novel Constitutional PIK3CA Mutation in a Child Lacking the Typical Segmental Overgrowth of

"PIK3CA-Related Overgrowth Spectrum". Hum Mutat. 2016 Mar;37(3):242-5. doi:10.1002/humu.22933. Epub 2015 Dec 15.

Elkind MS, Hills NK, Glaser CA, Lo WD, **Amlie-Lefond C**, Dlamini N, Kneen R, Hod EA, Wintermark M, deVeber GA, Fullerton HJ; VIPS Investigators\*. Herpesvirus Infections and Childhood Arterial Ischemic Stroke: Results of the VIPS Study.Circulation. 2016 Feb 23;133(8):732-41. doi: 10.1161/CIRCULATIONAHA.115.018595.Epub 2016 Jan 26.

Marin SE, **Saneto RP**. Neuropsychiatric Features in Primary Mitochondrial Disease. Neurol Clin. 2016 Feb;34(1):247-94. doi: 10.1016/j.ncl.2015.08.011. Review. PubMed PMID: 26614002.

Storey GP, Gonzalez-Fernandez G, Bamford IJ, Hur M, McKinley JW, Heimbigner L, Minasyan A, Walwyn WM, **Bamford NS.** Nicotine Modifies Corticostriatal Plasticity and Amphetamine Rewarding Behaviors in Mice(1,2,3). eNeuro. 2016 Feb 2;3(1). pii:ENEURO.0095-15.2015. doi: 10.1523/ENEURO.0095-15.2015. eCollection 2016 Jan-Feb

Burns TM, Smith GA, Allen JA, Amato AA, Arnold WD, Barohn R, Benatar M, Bird SJ, Bromberg M, Chahin N, Ciafaloni E, Cohen JA, Corse A, Crum BA, David WS, Dimberg E, Sousa EA, Donofrio PD, Dyck PJ, Engel AG, Ensrud ER, Ferrante M,Freimer M, Gable KL, Gibson S, Gilchrist JM, Goldstein JM, Gooch CL, Goodman BP, Gorelov D, **Gospe SM Jr,** Goyal NA, Guidon AC, Guptill JT, Gutmann L, Gutmann L,Gwathmey K, Harati Y, Harper CM Jr, Hehir MK, Hobson-Webb LD, Howard JF Jr,Jackson CE, Johnson N, Jones SM, Juel VC, Kaminski HJ, Karam C, Kennelly KD, Khella S, Khoury J, Kincaid JC, Kissel JT, Kolb N, Lacomis D, Ladha S, Larriviere D, Lewis RA, Li Y, Litchy WJ, Logigian E, Lou JS, MacGowen DJ, Maselli R, Massey JM, Mauermann ML, Mathews KD, Meriggioli MN, Miller RG, Moon JS, Mozaffar T, Nations SP, Nowak RJ, Ostrow LW, Pascuzzi RM, Peltier A, Ruzhansky K, Richman DP, Ross MA, Rubin DI, Russell JA, Sachs GM, Salajegheh MK, Saperstein DS, Scelsa S, Selcen D, Shaibani A, Shieh PB, Silvestri NJ, Singleton JR, Smith BE, So YT, Solorzano G, Sorenson EJ, Srinivasen J, Tavee J, Tawil R, Thaisetthawatkul P, Thornton C, Trivedi J, Vernino S, Wang AK, Webb TA, Weiss MD, Windebank AJ, Wolfe GI. Editorial by concerned physicians: Unintended effect of the orphan drug act on the potential cost of 3,4-diaminopyridine. Muscle Nerve. 2016 Feb;53(2):165-8.

doi: 10.1002/mus.25009..

Berg AT, **Dobyns WB**. Progress in autism research and postgenomic studies -Authors' reply. Lancet Neurol. 2016 Feb;15(2):136-137. doi:10.1016/S1474-4422(15)00403-2. Epub 2016 Jan 12

Huang L, Vanstone MR, Hartley T, Osmond M, Barrowman N, Allanson J, Baker L,Dabir TA, Dipple KM, **Dobyns WB**, Estrella J, Faghfoury H, Favaro FP, Goel H,Gregersen PA, Gripp KW, Grix A, Guion-Almeida ML, Harr MH, Hudson C, Hunter AG,Johnson J, Joss SK, Kimball A, Kini U, Kline AD, Lauzon J, Lildballe DL,López-González V, Martinezmoles J, Meldrum C, Mirzaa GM, Morel CF, Morton JE,Pyle LC, Quintero-Rivera F, Richer J, Scheuerle AE, Schönewolf-Greulich B, Shears DJ, Silver J, Smith AC, Temple IK; UCLA Clinical Genomics Center, van de Kamp JM,van Dijk FS, Vandersteen AM, White SM, Zackai EH, Zou R; Care4Rare Canada Consortium, Bulman DE, Boycott KM, Lines MA. Mandibulofacial Dysostosis with Microcephaly: Mutation and Database Update. Hum Mutat. 2016 Feb;37(2):148-54.

doi: 10.1002/humu.22924. Epub 2015 Nov 19. Review

**Amlie-Lefond C**, Shaw D. Vascular disease. Handb Clin Neurol. 2016;136:1159-71. doi: 10.1016/B978-0-444-53486-6.00060-0. Review.

**Amlie-Lefond C**. Dissecting etiologies of posterior circulation stroke. Dev Med Child Neurol. 2016 Jan;58(1):10-1. doi: 10.1111/dmcn.12969

Olson JD, Wander JD, Johnson L, Sarma D, Weaver K, **Novotny EJ**, Ojemann JG,Darvas F. Comparison of subdural and subgaleal recordings of cortical high-gamma activity in humans. Clin Neurophysiol. 2016 Jan;127(1):277-284. doi:10.1016/j.clinph.2015.03.014.

D'Angelo D, Lebon S, Chen Q, Martin-Brevet S, Snyder LG, Hippolyte L, Hanson E, Maillard AM, Faucett WA, Macé A, Pain A, Bernier R, Chawner SJ, David A,Andrieux J, Aylward E, Baujat G, Caldeira I, Conus P, Ferrari C, Forzano F,Gérard M, Goin-Kochel RP, Grant E, Hunter JV, Isidor B, Jacquette A, Jønch AE, Keren B, Lacombe D, Le Caignec C, Martin CL, Männik K, Metspalu A, Mignot C,Mukherjee P, Owen MJ, Passeggeri M, Rooryck-Thambo C, Rosenfeld JA, Spence SJ, **Steinman KJ,** Tjernagel J, Van Haelst M, Shen Y, Draganski B, Sherr EH, Ledbetter DH, van den Bree MB, Beckmann JS, Spiro JE, Reymond A, Jacquemont S, Chung WK;Cardiff University Experiences of Children With Copy Number Variants (ECHO)Study; 16p11.2 European Consortium; Simons Variation in Individuals Project (VIP)Consortium. Defining the Effect of the 16p11.2 Duplication on Cognition,Behavior,and Medical Comorbidities. JAMA Psychiatry. 2016 Jan;73(1):20-30. doi: 10.1001/jamapsychiatry.2015.2123

Poliachik SL, Friedman SD, Poliakov AV, Budech CB, Ishak GE, Shaw DW, **Gospe SM Jr.** Corpus Callosum Diffusion and Connectivity Features in High Functioning Subjects With Pyridoxine-Dependent Epilepsy. Pediatr Neurol. 2016 Jan;54:43-8. doi: 10.1016/j.pediatrneurol.2015.09.012.

Tsuchida TN, Acharya JN, Halford JJ**, Kuratani JD**, Sinha SR, Stecker MM, Tatum WO, Drislane FW. American Clinical Neurophysiology Society: EEG Guidelines Introduction. Neurodiagn J. 2016;56(4):231-234. doi:10.1080/21646821.2016.1245513.

**2015**

McCormick MA, Delaney JR, Tsuchiya M, Tsuchiyama S, Shemorry A, Sim S, Chou AC, Ahmed U, Carr D, Murakami CJ, Schleit J, Sutphin GL, Wasko BM, Bennett CF, Wang AM, Olsen B, Beyer RP, Bammler TK, Prunkard D, **Johnson SC**, Pennypacker JK, An E, Anies A, Castanza AS, Choi E, Dang N, Enerio S, Fletcher M, Fox L, Goswami S, Higgins SA, Holmberg MA, Hu D, Hui J, Jelic M, Jeong KS, Johnston E, Kerr EO, Kim J, Kim D, Kirkland K, Klum S, Kotireddy S, Liao E, Lim M, Lin MS, Lo WC, Lockshon D, Miller HA, Moller RM, Muller B, Oakes J, Pak DN, Peng ZJ, Pham KM, Pollard TG, Pradeep P, Pruett D, Rai D, Robison B, Rodriguez AA, Ros B, Sage M, Singh MK, Smith ED, Snead K, Solanky A, Spector BL, Steffen KK, Tchao BN, Ting MK, Vander Wende H, Wang D, Welton KL, Westman EA, Brem RB, Liu XG, Suh Y, Zhou Z, Kaeberlein M, Kennedy BK: A Comprehensive Analysis of Replicative Lifespan in 4,698 Single-Gene Deletion Strains Uncovers Conserved Mechanisms of Aging. *Cell Metab*. 2015. 22(5):p895-906. PMID: 26456335.

**Johnson SC**, Yanos ME, Bitto A, Castanza A, Gagnidze A, Gonzalez B, Gupta K, Hui J, Jarvie C, Johnson BM, Letexier N, McCanta L, Sangesland M, Tamis O, Uhde L, Van Den Ende A, Rabinovitch PS, Suh Y, Kaeberlein M: Dose-dependent Effects of mTOR Inhibition on Weight and Mitochondrial Disease in Mice. *Front. Genet.* 2015. 6:p247. PMID: 26257774.

**Amlie-Lefond C**, Rivkin MJ, Friedman NR, Bernard TJ, Dowling MM, deVeber G.The Way Forward: Challenges and Opportunities in Pediatric Stroke. Pediatr Neurol. 2016 Mar;56:3-7. doi: 10.1016/j.pediatrneurol.2015.10.021. Epub 2015 Dec 10. PubMed PMID: 26803334.

Schwarz AC, **Amlie-Lefond C**. A 15-Year-Old Boy With Trisomy 21 and Postoperative Weakness. JAMA Pediatr. 2016 Jan;170(1):85-6. doi:10.1001/jamapediatrics.2015.1677.

Crealey M, Allen NM, Webb D, **Bouldin A,** Mc Sweeney N, Peake D, Tirupathi S, Butler K, King MD. Sydenham's chorea: not gone but perhaps forgotten. Arch Dis Child. 2015 Dec;100(12):1160-2. doi: 10.1136/archdischild-2015-308693. Epub 2015 Sep 15

Roy A, Skibo J, Kalume F, Ni J, Rankin S, Lu Y, **Dobyns WB**, Mills GB, Zhao JJ, Baker SJ, Millen KJ. Mouse models of human PIK3CA-related brain overgrowth have acutely treatable epilepsy. Elife. 2015 Dec 3;4. pii: e12703. doi:10.7554/eLife.12703..

Mirzaa GM, Conti V, Timms AE, Smyser CD, Ahmed S, Carter M, Barnett S, Hufnagel RB, Goldstein A, Narumi-Kishimoto Y, Olds C, Collins S, Johnston K,Deleuze JF, Nitschké P, Friend K, Harris C, Goetsch A, Martin B, Boyle EA, Parrini E, Mei D, Tattini L, Slavotinek A, Blair E, Barnett C, Shendure J, Chelly J, **Dobyns WB**, Guerrini R. haracterisation of mutations of the phosphoinositide-3-kinase regulatory subunit, PIK3R2, in perisylvian polymicrogyria: a next-generation sequencing study. Lancet Neurol. 2015 Dec;14(12):1182-95. doi: 10.1016/S1474-4422(15)00278-1. Epub 2015 Oct 29.

McMahon KQ, Papandreou A, Ma M, Barry BJ, Mirzaa GM, **Dobyns WB**, Scott RH,Trump N, Kurian MA, Paciorkowski AR. Familial recurrences of FOXG1-related disorder: Evidence for mosaicism. Am J Med Genet A. 2015 Dec;167A(12):3096-102.doi: 10.1002/ajmg.a.37353. Epub 2015 Sep 14.

Miller DT, Chung W, Nasir R, Shen Y, **Steinman KJ,** Wu BL, Hanson E. 16p11.2 Recurrent Microdeletion. 2009 Sep 22 [updated 2015 Dec 10]. In: Adam MP, Ardinger HH, Pagon RA, Wallace SE, Bean LJH, Stephens K, Amemiya A, editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2018.

Mirzaa GM, Collins S, **Dobyns WB**. Corrigendum to "Congenital microcephaly and chorioretinopathy due to de novo heterozygous KIF11 mutations: Five novel mutations and review of the literature. Am J Med Genet Part A 2014 164A:2879-86". Am J Med Genet A. 2015 Nov 14. doi: 10.1002/ajmg.a.37449. [

**Shurtleff HA**, Barry D, Firman T, **Warner MH**, Aguilar-Estrada RL, **Saneto RP**, **Kuratani JD**, Ellenbogen RG, **Novotny EJ**, Ojemann JG. Impact of epilepsy surgery on development of preschool children: identification of a cohort likely to benefit from early intervention. J Neurosurg Pediatr. 2015 Oct;16(4):383-92. doi:10.3171/2015.3.PEDS14359

**Wrede JE,** Mengel-From J, Buchwald D, Vitiello MV, Bamshad M, Noonan C, Christiansen L, Christensen K, Watson NF. Mitochondrial DNA Copy Number in Sleep Duration Discordant Monozygotic Twins. Sleep. 2015 Oct 1;38(10):1655-8. doi:10.5665/sleep.5068

Feyissa EA, Cornell E, Chandhok L, Wang D, **Ionita C**, Schwab J, Kostyun R,Wilion F, Rubin K. Impact of Co-management at the Primary-Subspecialty Care Interface on Follow-up and Referral Patterns for Patients With Concussion. Clin Pediatr (Phila). 2015 Sep;54(10):969-75. doi: 0.1177/0009922814566929

Mefford HC, Zemel M, Geraghty E, Cook J, Clayton PT, Paul K, Plecko B, Mills PB, Nordli DR Jr, **Gospe SM Jr**. Intragenic deletions of ALDH7A1 in pyridoxine-dependent epilepsy caused by Alu-Alu recombination. Neurology. 2015 Sep 1;85(9):756-62.

Arredondo J, Lara M, **Gospe SM Jr**, Mazia CG, Vaccarezza M, Garcia-Erro M, Bowe CM, Chang CH, Mezei MM, Maselli RA. Choline Acetyltransferase Mutations Causing Congenital Myasthenic Syndrome: Molecular Findings and Genotype-Phenotype Correlations. Hum Mutat. 2015 Sep;36(9):881-93. doi: 10.1002/humu.22823

**Monrad P,** Sannagowdara K, Bozarth X, Bhosrekar S, Hecox K, Nwosu M, Schwabe M, Meyer M, Szabo A, Prigge J, Lemke R, Horn B, Whelan HT. Haemodynamic response associated with both ictal and interictal epileptiform activity using simultaneous video electroencephalography/near infrared spectroscopy in a within-subject study. J Near Infrared Spectrosc. 2015;23(4):209-218. doi: 10.1255/jnirs.1170. Epub 2015 Sep 24. PMID: 26538840; PMCID: PMC4629858

Oegema R, Cushion TD, Phelps IG, Chung SK, Dempsey JC, Collins S, Mullins JG,Dudding T, Gill H, Green AJ, **Dobyns WB**, Ishak GE, Rees MI, Doherty D.Recognizable cerebellar dysplasia associated with mutations in multiple tubulin genes. Hum Mol Genet. 2015 Sep 15;24(18):5313-25. doi: 10.1093/hmg/ddv250. Epub 2015 Jun 30.

**Blume HK**. Headaches after Concussion in Pediatrics: a Review. Curr Pain Headache Rep. 2015 Sep;19(9):42. doi: 10.1007/s11916-015-0516-x. Review.

Ng YT, **Gospe SM Jr**, Sahin M. Pediatric Neurology 2014 Trainee Publication Award Winner: Dr. Mitchel T. Williams. Pediatr Neurol. 2015 Aug;53(2):103-4. doi:10.1016/j.pediatrneurol.2015.04.010.

**Morgan LA**, Buchhalter J. Psychogenic Paroxysmal Nonepileptic Events in Children: A Review. Pediatr Neurol. 2015 Jul;53(1):13-22.

Elbers J, **Wainwright MS**, **Amlie-Lefond C**. The Pediatric Stroke Code: Early Management of the Child with Stroke. J Pediatr. 2015 Jul;167(1):19-24.e1-4. doi:10.1016/j.jpeds.2015.03.051. Epub 2015 May 1. PubMed PMID: 25937428. 21: Amlie-Lefond C, Ellenbogen RG. Factors associated with the presentation of moyamoya in childhood. J Stroke Cerebrovasc Dis. 2015 Jun;24(6):1204-10. doi:10.1016/j.jstrokecerebrovasdis.2015.01.018. Epub 2015 Apr 10.

Roth CL, Eslamy H, Werny D, Elfers C, Shaffer ML, Pihoker C, Ojemann J, **Dobyns WB.** Semiquantitative analysis of hypothalamic damage on MRI predicts risk for hypothalamic obesity. Obesity (Silver Spring). 2015 Jun;23(6):1226-33. doi:10.1002/oby.21067. Epub 2015 Apr 17

Jansen LA, Mirzaa GM, Ishak GE, O'Roak BJ, Hiatt JB, Roden WH, Gunter SA,Christian SL, Collins S, Adams C, Rivière JB, St-Onge J, Ojemann JG, Shendure J, Hevner RF, **Dobyns WB.** PI3K/AKT pathway mutations cause a spectrum of brain malformations from megalencephaly to focal cortical dysplasia. Brain. 2015 Jun;138(Pt 6):1613-28. doi: 10.1093/brain/awv045. Epub 2015 Feb 25.

Roosing S, Hofree M, Kim S, Scott E, Copeland B, Romani M, Silhavy JL, Rosti RO, Schroth J, Mazza T, Miccinilli E, Zaki MS, Swoboda KJ, Milisa-Drautz J, **Dobyns WB**, Mikati MA, İncecik F, Azam M, Borgatti R, Romaniello R, Boustany RM, Clericuzio CL, D'Arrigo S, Strømme P, Boltshauser E, Stanzial F,Mirabelli-Badenier M, Moroni I, Bertini E, Emma F, Steinlin M, Hildebrandt F,Johnson CA, Freilinger M, Vaux KK, Gabriel SB, Aza-Blanc P, Heynen-Genel S,Ideker T, Dynlacht BD, Lee JE, Valente EM, Kim J, Gleeson JG. Functional genome-wide siRNA screen identifies KIAA0586 as mutated in Joubert syndrome. Elife. 2015 May 30;4:e06602. doi: 10.7554/eLife.06602.

Barkovich AJ**, Dobyns WB**, Guerrini R. Malformations of cortical development and epilepsy. Cold Spring Harb Perspect Med. 2015 May 1;5(5):a022392. doi:10.1101/cshperspect.a022392. Review.

Berg AT, **Dobyns WB**. Progress in autism and related disorders of brain development. Lancet Neurol. 2015 Nov;14(11):1069-70. doi:10.1016/S1474-4422(15)00048-4. Epub 2015 Apr 16

Saini A, Emke AR, Silva MC, **Perlman SJ**. Response to Eculizumab in Escherichia coli O157: H7-induced hemolytic uremic syndrome with severe neurological manifestations. 24817079 Clinical pediatrics, 2015 April : 54(4)387-9

Hansen J, Snow C, Tuttle E, Ghoneim DH, Yang CS, Spencer A, Gunter SA, Smyser CD, Gurnett CA, Shinawi M, **Dobyns WB**, Wheless J, Halterman MW, Jansen LA, Paschal BM, Paciorkowski AR. De novo mutations in SIK1 cause a spectrum of developmental epilepsies. Am J Hum Genet. 2015 Apr 2;96(4):682-90. doi:10.1016/j.ajhg.2015.02.013. Erratum in: Am J Hum Genet. 2015 Jun 4;96(6):1009.

Luks VL, Kamitaki N, Vivero MP, Uller W, Rab R, Bovée JV, Rialon KL, Guevara CJ, Alomari AI, Greene AK, Fishman SJ, Kozakewich HP, Maclellan RA, Mulliken JB, Rahbar R, Spencer SA, Trenor CC 3rd, Upton J, Zurakowski D, Perkins JA, Kirsh A, Bennett JT, **Dobyns WB**, Kurek KC, Warman ML, McCarroll SA, Murillo R. Lymphatic and other vascular malformative/overgrowth disorders are caused by somatic mutations in PIK3CA. J Pediatr. 2015 Apr;166(4):1048-54.e1-5. doi:10.1016/j.jpeds.2014.12.069. Epub 2015 Feb 11

Lerche H, **Novotny EJ Jr**. Microscopic brain structure revisited in genetic epilepsy. Neurology. 2015 Mar 31;84(13):1290-1.

Szabo A, Hill MD, Scholz K, **Amlie-Lefond C.** Thrombolysis in pediatric stroke study. Stroke. 2015 Mar;46(3):880-5. doi: 10.1161/STROKEAHA.114.008210. Epub 2015 Jan 22.

**Tully HM,** Capote RT, Saltzman BS. Maternal and infant factors associated with infancy-onset hydrocephalus in Washington State. Pediatr Neurol. 2015 Mar;52(3):320-5. doi: 0.1016/j.pediatrneurol.2014.10.030.

Paciorkowski AR, McDaniel SS, Jansen LA, **Tully H**, Tuttle E, Ghoneim DH, Tupal S, Gunter SA, Vasta V, Zhang Q, Tran T, Liu YB, Ozelius LJ, Brashear A, Sweadner KJ, **Dobyns WB**, Hahn S. Novel mutations in ATP1A3 associated with catastrophic early life epilepsy, episodic prolonged apnea, and postnatal microcephaly.Epilepsia. 2015 Mar;56(3):422-30. doi: 10.1111/epi.12914. Epub 2015 Feb 5..

Choe MC, **Blume HK**. Pediatric Posttraumatic Headache: A Review. J Child Neurol. 2016 Jan;31(1):76-85.

**Simard-Tremblay E**, Berry P, Owens A, Cook WB, **Sittner HR**, **Mazzanti M**, Huber J, **Warner M, Shurtleff H**, **Saneto RP**. High-fat diets and seizure control in myoclonic-astatic epilepsy: a single center's experience. Seizure. 2015 Feb;25:184-6. doi: 10.1016/j.seizure.2014.10.009

Mishra-Gorur K, Çağlayan AO, Schaffer AE, Chabu C, Henegariu O, Vonhoff F,Akgümüş GT, Nishimura S, Han W, Tu S, Baran B, Gümüş H, Dilber C, Zaki MS, HossniHAA, Rivière JB, Kayserili H, Spencer EG, Rosti RÖ, Schroth J, Per H, Çağlar C,Çağlar Ç, Dölen D, Baranoski JF, Kumandaş S, Minja FJ, Erson-Omay EZ, Mane SM,Lifton RP, Xu T, Keshishian H, **Dobyns WB**, Chi NC, Šestan N, Louvi A, Bilgüvar K, Yasuno K, Gleeson JG, Günel M. Mutations in KATNB1 Cause Complex Cerebral Malformations by Disrupting Asymmetrically Dividing Neural Progenitors. Neuron.

2015 Jan 7;85(1):228. doi: 10.1016/j.neuron.2014.12.046.

**Carapetian S (Randle**), Hageman J, Lyons E, Leonard D, Janies K, Kelley K, Fuchs S. Emergency Department Evaluation and Management of Children With Simple Febrile Seizures. Clin Pediatr (Phila). 2015;54:992-8.

**2014**

Carbonell A, Takeda A, Fahlgren N, **Johnson SC**, Cuperus JT, Carrington JC: New Generation of Artificial MicroRNA and Synthetic Trans-Acting Small Interfering RNA Vectors for Efficient Gene Silencing in Arabidopsis. *Plant Physiology*. 165(1):p15-29. 2014. PMID: 24647477.

Mishra-Gorur K, Çağlayan AO, Schaffer AE, Chabu C, Henegariu O, Vonhoff F, Akgümüş GT, Nishimura S, Han W, Tu S, Baran B, Gümüş H, Dilber C, Zaki MS, Hossni HA, Rivière JB, Kayserili H, Spencer EG, Rosti RÖ, Schroth J, Per H, Çağlar C,Çağlar Ç, Dölen D, Baranoski JF, Kumandaş S, Minja FJ, Erson-Omay EZ, Mane SM,Lifton RP, Xu T, Keshishian H, **Dobyns WB,** Chi NC, Šestan N, Louvi A, Bilgüvar K, Yasuno K, Gleeson JG, Günel M. Mutations in KATNB1 cause complex cerebral malformations by disrupting asymmetrically dividing neural progenitors. Neuron.2014 Dec 17;84(6):1226-39. doi: 10.1016/j.neuron.2014.12.014. Erratum in: Neuron.2015 Jan 7;85(1):228. Neuron. 2015 Jan 7;85(1):228

**Johnson SC**: Translational Medicine. A target for pharmacological intervention in an untreatable human disease. *Science*. 2014 Dec 5;346(6214). PMID: 25477449.

Mirzaa GM, Enyedi L, Parsons G, Collins S, Medne L, Adams C, Ward T, Davitt B, Bicknese A, Zackai E, Toriello H, **Dobyns WB**, Christian S. Congenital microcephaly and chorioretinopathy due to de novo heterozygous KIF11 mutations: five novel mutations and review of the literature. Am J Med Genet A. 2014 Nov;164A(11):2879-86. doi: 10.1002/ajmg.a.36707. Epub 2014 Aug 12. Review. PubMed

**Wainwright MS**, Grimason M, Goldstein J, Smith CM, **Amlie-Lefond** C, Revivo G,Noah ZL, Harris ZL, Epstein LG. Building a pediatric neurocritical care program: a multidisciplinary approach to clinical practice and education from the intensive care unit to the outpatient clinic. Semin Pediatr Neurol. 2014 Dec;21(4):248-54. doi: 0.1016/j.spen.2014.10.006. Epub 2014 Nov 1.

**Ruhoy IS,** **Saneto RP**. The genetics of Leigh syndrome and its implications for clinical practice and risk management. Appl Clin Genet. 2014 Nov 13;7:221-34. doi: 10.2147/TACG.S46176. eCollection 2014. Review..

Friedman SD, Ishak GE, Poliachik SL, Poliakov AV, Otto RK, Shaw DW, Willemsen MA, Bok LA, **Gospe** **SM Jr**. Callosal alterations in pyridoxine-dependent epilepsy.Dev Med Child Neurol. 2014 Nov;56(11):1106-10. doi: 10.1111/dmcn.12511. Epu

**Johnson SC**, Dong X, Vijg J, Suh Y: Genetic evidence for common pathways in human age-related diseases. *Aging Cell*. 2014 Oct;14(5):p809-17. PMID: 26077337.

Yamamoto S, Jaiswal M, Charng WL, Gambin T, Karaca E, Mirzaa G, Wiszniewski W, Sandoval H, Haelterman NA, Xiong B, Zhang K, Bayat V, David G, Li T, Chen K,Gala U, Harel T, Pehlivan D, Penney S, Vissers LELM, de Ligt J, Jhangiani SN, Xie Y, Tsang SH, Parman Y, Sivaci M, Battaloglu E, Muzny D, Wan YW, Liu Z, Lin-Moore AT, Clark RD, Curry CJ, Link N, Schulze KL, Boerwinkle E, **Dobyns WB**, Allikmets R, Gibbs RA, Chen R, Lupski JR, Wangler MF, Bellen HJ. A drosophila genetic resource of mutants to study mechanisms underlying human genetic diseases. Cell. 2014 Sep 25;159(1):200-214. doi: 10.1016/j.cell.2014.09.002

French CR, Seshadri S, Destefano AL, Fornage M, Arnold CR, Gage PJ, Skarie JM, **Dobyns WB,** Millen KJ, Liu T, Dietz W, Kume T, Hofker M, Emery DJ, Childs SJ, Waskiewicz AJ, Lehmann OJ. Mutation of FOXC1 and PITX2 induces cerebral small-vessel disease. J Clin Invest. 2014 Nov;124(11):4877-81. doi:10.1172/JCI75109. Epub 2014 Sep 24.

Kakar N, Ahmad J, Morris-Rosendahl DJ, Altmüller J, Friedrich K, Barbi G, Nürnberg P, Kubisch C, **Dobyns WB,** Borck G. STIL mutation causes autosomal recessive microcephalic lobar holoprosencephaly. Hum Genet. 2015 Jan;134(1):45-51. doi: 10.1007/s00439-014-1487-4. Epub 2014 Sep 14.

**Tully HM, Dobyns WB.** Infantile hydrocephalus: a review of epidemiology, classification and causes. Eur J Med Genet. 2014 Aug;5(8):359-68. doi:10.1016/j.ejmg.2014.06.002.

Apkon SD, Grady R, Hart S, Lee A, McNalley T, Niswander L, Petersen J, Remley S, Rotenstein D, **Shurtleff H, Warner M**, Walker WO Jr. Advances in the care of children with spina bifida. Adv Pediatr. 2014 Aug;61(1):33-74. doi:10.1016/j.yapd.2014.03.007

**Dobyns WB**, Das S. LIS1-Associated Lissencephaly/Subcortical Band Heterotopia.2009 Mar 3 [updated 2014 Aug 14]. In: Adam MP, Ardinger HH, Pagon RA, Wallace SE,Bean LJH, Stephens K, Amemiya A, editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2018. Available from http://www.ncbi.nlm.nih.gov/books/NBK5189/.

Seltzer LE, Ma M, Ahmed S, Bertrand M, **Dobyns WB**, Wheless J, Paciorkowski AR Epilepsy and outcome in FOXG1-related disorders. Epilepsia. 2014 Aug;55(8):1292-300. doi: 10.1111/epi.12648. Epub WB. De novo mutations in the beta-tubulin gene TUBB2A cause simplified gyral patterning and infantile-onset epilepsy. Am J Hum Genet. 2014 Apr 3;94(4):634-41. doi: 10.1016/j.ajhg.2014.03.009.:

Mirzaa GM, Millen KJ, Barkovich AJ, **Dobyns WB,** Paciorkowski AR. The Developmental Brain Disorders Database (DBDB): a curated neurogenetics knowledge base with clinical and research applications. Am J Med Genet A. 2014 Jun;164A(6):1503-11. doi: 10.1002/ajmg.a.36517. Epub 2014 Apr 3. PubMed PMID:

**Otten CE**, Creutzfeldt CJ. Fulminant acute disseminated encephalomyelitis presenting in an adult. JAMA Neurol. 2014 71:648-9.

Nerloes A, Di Donato N, Masliah-Planchon J, Jongmans M, Abdul-Raman OA,Albrecht B, Allanson J, Brunner H, Bertola D, Chassaing N, David A, Devriendt K, Eftekhari P, Drouin-Garraud V, Faravelli F, Faivre L, Giuliano F, Guion Almeida L, Juncos J, Kempers M, Eker HK, Lacombe D, Lin A, Mancini G, Melis D, Lourenço, CM, Siu VM, Morin G, Nezarati M, Nowaczyk MJ, Ramer JC, Osimani S, Philip N, Pierpont ME, Procaccio V, Roseli ZS, Rossi M, Rusu C, Sznajer Y, Templin L,Uliana V, Klaus M, Van Bon B, Van Ravenswaaij C, Wainer B, Fry AE, Rump A,Hoischen A, Drunat S, Rivière JB, **Dobyns WB**, Pilz DT. Baraitser-Winter cerebrofrontofacial syndrome: delineation of the spectrum in 42 cases. Eur J Hum Genet. 2015 Mar;23(3):292-301. doi: 10.1038/ejhg.2014.95. Epub 2014 Jul 23.

Rivkin MJ, deVeber G, Ichord RN, Kirton A, Chan AK, Hovinga CA, Gill JC, Bernard TJ, Rivkin MJ, Scholz K, deVeber G, Kirton A, Gill JC, Chan AK,Hovinga CA, Ichord RN, Grotta JC, Jordan LC, Benedict S, Friedman NR, Dowling MM, Elbers J, Torres M, Sultan S, Cummings DD, Grabowski EF, McMillan HJ, Beslow LA, **Amlie-Lefond C**; Thrombolysis in Pediatric Stroke Study. Emergence of the primary pediatric stroke center: impact of the thrombolysis in pediatric stroke trial. Stroke. 2014 Jul;45(7):2018-23. doi: 10.1161/STROKEAHA.114.004919. Epub 2014 Jun 10

**Beatty CW,** Ko PR, Nixon J**, Gospe SM Jr**. Delayed-onset movement disorder and encephalopathy after oxycodone ingestion. Semin Pediatr Neurol. 2014 Jun;21(2):160-5. doi: 10.1016/j.spen.2014.06.009. Epub 2014 Jun 18. PubMed

Fernández-López D, **Natarajan N,** Ashwal S, Vexler ZS. Mechanisms of perinatal arterial ischemic stroke. J Cereb Blood Flow Metab. 2014 Jun;34(6):921-32. doi:10.1038/jcbfm.2014.41. Epub 2014 Mar 26. Review.

**Tully HM, Dobyns WB**. Infantile hydrocephalus: a review of epidemiology, classification and causes. Eur J Med Genet. 2014 Aug;57(8):359-68. doi:10.1016/j.ejmg.2014.06.002. Epub 2014 Jun 13. Review. PubMed PMID: 24932902;

Guerrini R, **Dobyns WB**. Malformations of cortical development: clinical features and genetic causes. Lancet Neurol. 2014 Jul;13(7):710-26. doi:10.1016/S1474-4422(14)70040-7. Epub 2014 Jun 2. Review.

Paciorkowski AR, Weisenberg J, Kelley JB, Spencer A, Tuttle E, Ghoneim D, Thio LL, Christian SL, **Dobyns WB**, Paschal BM. Autosomal recessive mutations in nuclear transport factor KPNA7 are associated with infantile spasms and cerebellar malformation. Eur J Hum Genet. 2014 May;22(5):587-93. doi:10.1038/ejhg.2013.196. Epub 2013 Sep 18.

Segal MM, Williams MS, Gropman AL, Torres AR, Forsyth R, Connolly AM, El-Hattab AW, **Perlman SJ**, Samanta D, Parikh S, Pavlakis SG, Feldman LK, Betensky RA, **Gospe SM Jr** Evidence-based decision support for neurological diagnosis reduces errors and unnecessary workup. 23576414 Journal of child neurology, 2014 April : 29(4)487-92

Pearl PL, **Gospe SM Jr**. Pyridoxine or pyridoxal-5'-phosphate for neonatal epilepsy: the distinction just got murkier. Neurology. 2014 Apr 22;82(16):1392-4.doi: 10.1212/WNL.0000000000000351

Wallace SE, Conta JH, Winder TL, Willer T, Eskuri JM, Haas R, Patterson K, Campbell KP, Moore SA, **Gospe SM Jr**. A novel missense mutation in POMT1 modulates the severe congenital muscular dystrophy phenotype associated with POMT1 nonsense mutations. Neuromuscul Disord. 2014 Apr;24(4):312-20. doi:10.1016/j.nmd.2014.01.001. Epub 2014 Jan 11.

Cacciagli P, Desvignes JP, Girard N, Delepine M, Zelenika D, Lathrop M, Lévy N, Ledbetter DH, **Dobyns WB**, Villard L. AP1S2 is mutated in X-linked Dandy-Walker malformation with intellectual disability, basal ganglia disease and seizures (Pettigrew syndrome). Eur J Hum Genet. 2014 Mar;22(3):363-8. doi:10.1038/ejhg.2013.135. Epub 2013 Jun 12.

**Ruhoy IS,** Merritt JL 2nd, **Amlie-Lefond** C. Cystathionine beta-synthase deficiency heralded by cerebral sinus venous thrombosis and stroke. Pediatr Neurol. 2014 Jan;50(1):108-11. doi: d10.1016/j.pediatrneurol.2013.08.021. Epub 2013 Oct 15.

**Amlie-Lefond C**, Gill JC. Approach to acute ischemic stroke in childhood. Curr Treat Options Cardiovasc Med. 2014 Jan;16(1):276. doi: 10.1007/s11936-013-0276-z.

van Karnebeek CD, Stockler-Ipsiroglu S, Jaggumantri S, Assmann B, Baxter P,Buhas D, Bok LA, Cheng B, Coughlin CR 2nd, Das AM, Giezen A, Al-Hertani W, Ho G, Meyer U, Mills P, Plecko B, Struys E, Ueda K, Albersen M, Verhoeven N, **Gospe SM Jr,** Gallagher RC, Van Hove JK, Hartmann H. Lysine-Restricted Diet as Adjunct Therapy for Pyridoxine-Dependent Epilepsy: The PDE Consortium Consensus Recommendations. JIMD Rep. 2014;15:1-11. doi: 10.1007/8904\_2014\_296.

Johnston JJ, Sapp JC, Curry C, Horton M, Leon E, Cusmano-Ozog K, **Dobyns WB**, Hudgins L, Zackai E, Biesecker LG. Expansion of the TARP syndrome phenotype associated with de novo mutations and mosaicism. Am J Med Genet A. 2014Jan;164A(1):120-8. doi: 10.1002/ajmg.a.36212.

**Beatty CW**, Creutzfeldt CJ, Davis AP, Hoffer Z, Khot SP. The diagnostic conundrum and treatment dilemma of a patient with a rapidly progressive encephalopathy. Neurohospitalist. 2014 Jan;4(1):34-41. doi:10.1177/1941874413496792.

**Carapetian S (Randle),** Hesselink J, Nass R, Trauner D, Stiles J. Face and location processing in children with early unilateral brain injury. Brain Cogn. 2014;88:6-13.

Wei Z, Chigurupati S, Bagsiyao P, **Henriquez A**, Chan SL. The brain uncoupling protein UCP4 attenuates mitochondrial toxin-induced cell death: role of extracellular signal-regulated kinases in bioenergetics adaptation and cell survival.

Neurotox Res. 2009 Jul;16(1):14-29. doi: 10.1007/s12640-009-9039-8. Epub 2009 Mar 25.