

PUBLICACIONES 2020-2021

Año 2020

- **"Nueva mutación en el gen CASK en un niño con síndrome de microcefalia e hipoplasia pontocerebelosa [New mutation in the CASK gene in a child with microcephaly syndrome and pontocerebellar hypoplasia]"** González-Roca I, Alonso-Rivero P, Sánchez-Soblechero A, Vázquez-López M.. Rev Neurol. 2020 Aug 16;71(4):161-162. Spanish. doi: 10.33588/rn.7104.2020120. PMID: 32700313
- **"Associations of paediatric demyelinating and encephalitic syndromes with myelin oligodendrocyte glycoprotein antibodies: a multicentre observational study."** Armangue T, Olivé-Cirera G, Martínez-Hernandez E, Sepulveda M, Ruiz-García R, Muñoz-Batista M, Ariño H, González-Álvarez V, Felipe-Rucián A, Jesús Martínez-González M, Cantarín-Extremera V, Concepción Miranda-Herrero MC, Monge-Galindo L, Tomás-Vila M, Miravet E, Málaga I, Arrambide G, Auger C, Tintoré M, Montalban X, Vanderver A, Graus F, Saiz A, Dalmau J; **Spanish Pediatric anti-MOG Study Group**. Lancet Neurol. 2020 Mar;19(3):234-246. doi: 10.1016/S1474-4422(19)30488-0. Epub 2020 Feb 10. Erratum in: Lancet Neurol. 2020 Apr;19(4):e4. PMID: 32057303
- **"Parálisis súbita de la supravversión de la mirada en una niña de 9 años" ["Sudden paralysis of upward gaze in a 9-year-old girl"]**. de la Mata-Navazo S, Toledo-Del Castillo B, Rodríguez-Jiménez C, Miranda-Herrero MC, Aguado-Del Hoyo A, González-Martínez F Rev Neurol. 2020 Nov 16;71(10):387-388. Spanish. doi:10.33588/rn.7110.2020275. PMID: 33145750
- **"Neurologic Complications in Children Hospitalized With Influenza Infections: Prevalence, Risk Factors and Impact on Disease Severity"**. Solís-García G, Chacón-Pascual A, González Martínez F, Miranda Herrero MC, Hernández-Sampelayo T, Catalán Alonso P, Rodríguez-Fernández R Pediatr Infect Dis J. 2020 Sep;39(9):789-793. doi: 10.1097/INF.0000000000002686. PMID: 32282657
- **"Endocrine and Growth Abnormalities in 4H Leukodystrophy Caused by Variants in POLR3A, POLR3B, and POLR1C"**. Pelletier F, Perrier S, Cayami FK, Mirchi A, Saikali S, Tran LT, Ulrick N, Guerrero K, Rampakakis E, van Spaendonk RML, Naidu S, Pohl D, Gibson WT, Demos M, Goizet C, Tejera-Martin I, Potic A, Fogel BL, Brais B, Sylvain M, Sebire G, Lourenço CM, Bonkowsky JL, Catsman-Berrevoets C, Pinto PS, Tirupathi S, Strømme P, de Grauw T, Gieruszczak-Bialek D, Krägeloh-Mann I, Mierzewska H, Philippi H, Rankin J, Atik T, Banwell B, Benko WS, Blaschek A, Bley A, Boltshauser E, Bratkovic D, Brozova K, Cimas I, Clough C, Corenblum B, Dinopoulos A, Dolan G, Faletra F, Fernandez R, Fletcher J, Garcia Garcia ME, Gasparini P, Gburek-Augustat J, Gonzalez Moron D, Hamati A, Harting

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- **"Biallelic mutations in *NDUFA8* cause complex I deficiency in two siblings with favorable clinical evolution"**. Tort F, Barredo E, Parthasarathy R, Ugarteburu O, Ferrer-Cortès X, García-Villoria J, Gort L, González-Quintana A, Martín MA, Fernández-Vizarra E, Zeviani M, Ribes A.. *Mol Genet Metab.* 2020 Nov;131(3):349-357. doi: 10.1016/j.ymgme.2020.10.005. Epub 2020 Oct 13. PMID: 33153867.
- **"Microbiota y dieta cetogénica"**. Alvarez-Calatayud G, Tolín-Hernani M, Barredo-Valderrama E, Meneses D, Morencos-Pinedo C. *An Microbiota Probióticos Prebióticos* 2020; 1(2).

Año 2021

- **"Visuospatial functions in preterm schoolchildren without cognitive delay: Using Pascual's Graphomotor test as a screening method"**. Miranda-Herrero MC, Vázquez-López M, Barredo-Valderrama E, de Castro de Castro P, Chacón-Pascual A, Pascual-Pascual SI.. *Early Hum Dev.* 2021 Aug 30;161:105454. doi: 10.1016/j.earlhumdev.2021.105454. Epub ahead of print. PMID: 34496347.
- **"Afectación del VI y el VII par craneal secundaria a hipertensión intracraneal idiopática en un paciente pediátrico [VI and VII cranial nerve injury secondary to intracranial idiopathic hypertension in a pediatric patient]"**. Fernández-Monteaigudo B, Miranda-Herrero MC, Chacón-Pascual A, Vázquez-López M. *Rev Neurol* 2021 Jun 16;72(12):443. Spanish. doi: 10.33588/rn.7212.2021111. PMID: 34110000
- **"Spanish Study Group NeuroimmunoPed-Covid. Impact of COVID-19 in Immunosuppressed Children With Neuroimmunologic Disorders"**. Olivé-Cirera G, Fonseca E, Cantarín-Extremera V, Vázquez-López M, Jiménez-Legido M, González-Álvarez V, Ribeiro-Constante J, Camacho-Salas A, Martí I, Cancho-Candela R, Martínez-González MJ, Saiz A, Armangué T; *Neurol Neuroimmunol Neuroinflamm.* 2021 Nov 10;9(1):e1101

- ***“Quality of life and psychiatric comorbidities in pediatric patients with Gilles de la Tourette syndrome”***. Solís-García G, Jové-Blanco A, Chacón-Pascual A, Vázquez-López M, Castro-De Castro P, Carballo JJ, Pina-Camacho L, Miranda-Herrero MC. Rev Neurol. 2021 Nov 16;73(10):339-344.
- ***“The Genetic Landscape of Mitochondrial Diseases in Spain: A Nationwide Call”*** Bellusci M, Paredes-Fuentes AJ, Ruiz-Pesini E, Gómez B; MITOSPAIN Working Group, Martín MA, Montoya J, Artuch R. Genes (Basel). 2021 Oct 9;12(10):1590.
- ***“COVID-19 in children with neuromuscular disorders”***. Natera-de Benito D, Aguilera-Albesa S, Costa-Comellas L, García-Romero M, Miranda-Herrero MC, Rúbies Olives J, García-Campos Ó, Martínez Del Val E, Martínez Garcia MJ, Medina Martínez I, Cancho-Candela R, Fernández-García MA, Pascual-Pascual SI, Gómez-Andrés D, Nascimento A; Neuromuscular Working Group of Spanish Pediatric Neurology Society. J Neurol. 2021 Sep;268(9):3081-3085. doi: 10.1007/s00415-020-10339-y. Epub 2021 Jan 2. PMID: 33387010; PMCID: PMC7775833.