Kantonsspital St.Gallen

André Schaller

Kontakt

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Publikationen (7)

Sanlialp A, Escher P, Schaller A, Todorova M. Clinical Heterogeneity in Two Siblings Harbouring a Heterozygous PRPH2 Pathogenic Variant. Klin Monbl Augenheilkd 2023; 240:536-543.

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Lazdinyte S, Schorderet D, Schaller A, Valmaggia C, Todorova M. Analysis of Inherited Optic Neuropathies. Klin Monbl Augenheilkd 2019; 236:451-461.

Schreglmann S, Bhatia K, Baumann C, Michels L, Krasemann E, Hidding U, Schaller A, Jaunmuktane Z, Waldvogel D, Kägi G, Ganos C, Galovic M, Riederer F, Jung H. Movement disorders in genetically confirmed mitochondrial disease and the putative role of the cerebellum. Mov Disord 2017

Jackson C, Neuwirth C, Hahn D, Nuoffer J, Frank S, Gallati S, Schaller A. Novel mitochondrial tRNA(Ile) m.4282A>G gene mutation leads to chronic progressive external ophthalmoplegia plus phenotype. Br J Ophthalmol 2014; 98:1453-9.

Zürcher S, Schaller A, Gorgievski-Hrisoho M, Garzoni C, Mohacsi P, Barbani M, Mühlemann K, Lüthi A, Mooser C, Flatz L. Sensitive and rapid detection of ganciclovir resistance by PCR based MALDI-TOF analysis. J Clin Virol 2012; 54:359-63.

Projekte (0)

Keine Resultate gefunden.

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