Kantonsspital St.Gallen

Andreas Klein-Franke

Kontakt

Andreas Klein-Franke

Publikationen (5)

Steiner L, Tscherter A, Henzi B, Branca M, Carda S, Enzmann C, Fluss J, Jacquier D, Neuwirth C, Ripellino P, Scheidegger O, Schlaeger R, Schreiner B, Stettner G, Klein-Franke A, Swiss-Reg-NMD Group. Chronic Pain in Patients with Spinal Muscular Atrophy in Switzerland: A Query to the Spinal Muscular Atrophy Registry. J Clin Med 2024; 13

Baumdick M, Niehrs A, Degenhardt F, Schwerk M, Hinrichs O, Jordan-Paiz A, Padoan B, Wegner L, Schloer S, Zecher B, Malsy J, Joshi V, Illig C, Schröder-Schwarz J, Möller K, Hamburg Intestinal Tissue Study Group, Martin M, Yuki Y, Ozawa M, Sauter J, Schmidt A, Perez D, Giannou A, Carrington M, Davis R, Schumacher U, Sauter G, Huber S, Puelles V, Melling N, Klein-Franke A, International Inflammatory Bowel Disease Genetics Consortium, Altfeld M, Bunders M, Brand S. HLA-DP on Epithelial Cells Enables Tissue Damage by NKp44 Natural Killer Cells in Ulcerative Colitis. Gastroenterology 2023; 165:946-962.e13.

Tscherter A, Rüsch C, Baumann D, Enzmann C, Hasselmann O, Jacquier D, Jung H, Kruijshaar M, Kuehni C, Neuwirth C, Stettner G, Klein-Franke A. Evaluation of real-life outcome data of patients with spinal muscular atrophy treated with nusinersen in Switzerland. Neuromuscul Disord 2022; 32:399-409.

van Rheenen W, van der Spek R, Bakker M, Farei-Campagna J, Hop P, Zwamborn R, de Klein N, Westra H, Bakker O, Deelen P, Shireby G, Hannon E, Moisse M, Baird D, Restuadi R, Dolzhenko E, Dekker A, Gawor K, Westeneng H, Tazelaar G, van Eijk K, Kooyman M, Byrne R, Doherty M, Heverin M, Al Khleifat A, Iacoangeli A, Shatunov A, Ticozzi N, Cooper-Knock J, Smith B, Gromicho M, Chandran S, Pal S, Morrison K, Openshaw P, Hardy J, Orrell R, Sendtner M, Meyer T, Başak N, van der Kooi A, Ratti A, Fogh I, Gellera C, Lauria G, Corti S, Cereda C, Sproviero D, D'Alfonso S, Sorarù G, Siciliano G, Filosto M, Padovani A, Chio A, Calvo A, Moglia C, Brunetti M, Canosa A, Grassano M, Beghi E, Pupillo E, Logroscino G, Nefussy B, Osmanovic A, Nordin A, Lerner Y, Zabari M, Gotkine M, Baloh R, Bell S, Vourc'h P, Corcia P, Couratier P, Millecamps S, Meininger V, Salachas F, Mora J, Assialioui A, Rojas-García R, Dion P, Ross J, Ludolph A, Weishaupt J, Brenner D, Freischmidt A, Bensimon G, Brice A, Dürr A, Payan C, Saker-Delye S, Wood N, Topp S, Rademakers R, Tittmann L, Lieb W, Klein-Franke A, Ripke S, Braun A, Kraft J, Whiteman D, Olsen C, Uitterlinden A, Hofman A, Rietschel M, Cichon S, Nöthen M, Amouyel P, SLALOM Consortium, PARALS Consortium, SLAGEN Consortium, SLAP Consortium, Traynor B, Singleton A, Mitne Neto M, Cauchi R, Ophoff R, Wiedau-Pazos M, Lomen-Hoerth C, Van Deerlin V, Grosskreutz J, Roediger A, Gaur N, Jörk A, Barthel T, Theele E, Ilse B, Stubendorff B, Witte O, Steinbach R, Hübner C, Graff C, Brylev L, Fominykh V, Demeshonok V, Ataulina A, Rogelj B, Koritnik B, Zidar J, Ravnik-Glavač M, Glavač D, Stević Z, Drory V, Povedano M, Blair I, Kiernan M, Benyamin B, Henderson R, Furlong S, Mathers S, McCombe P, Needham M, Ngo S, Nicholson G, Pamphlett R, Rowe D, Steyn F, Williams K, Mather K, Sachdev P, Henders A, Wallace L, de Carvalho M, Pinto S, Petri S, Weber M, Rouleau G, Silani V, Curtis C, Breen G, Glass J, Brown R, Landers J, Shaw C, Andersen P, Groen E, van Es M, Pasterkamp R, Fan D, Garton F, McRae A, Davey Smith G, Gaunt T, Eberle M, Mill J, McLaughlin R, Hardiman O, Kenna K, Wray N, Tsai E, Runz H, Franke L, Al-Chalabi A, Van Damme P, van den Berg L, Veldink J. Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. Nat Genet 2021; 53:1636-1648.

Neumann H, Apel T, Treier M, Reineke M, Walz M, Hoang-Vu C, Brauckhoff M, Klein-Franke A, Klose P, Schmidt H, Maier M, Peczkowska M, Szmigielski C, Glaesker S, Manz T, Munk R, Bausch B, McWhinney S, Bender B, Gimm O, Franke G, Schipper J, Klisch J, Altehoefer C, Zerres K, Januszewicz A, Eng C, Smith W, Freiburg-Warsaw-Columbus Pheochromocytoma Study Group. Germ-line mutations in nonsyndromic pheochromocytoma. N Engl J Med 2002; 346:1459-66.

Projekte (0)

Keine Resultate gefunden.

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