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Publikationen (20)

Stallhofer J, Veith L, Diegelmann J, Probst P, Brand S, Schnitzler F, Olszak T, Török H, Mayerle J, Stallmach A, Beigel F. Iron Deficiency in Inflammatory Bowel Disease Is Associated With Low Levels of Vitamin D Modulating Serum Hepcidin and Intestinal Ceruloplasmin Expression. Clin Transl Gastroenterol 2022; 13:e00450.

Schnitzler F, Friedrich M, Angelberger M, Diegelmann J, Stallhofer J, Wolf C, Dütschler J, Truniger S, Olszak T, Beigel F, Tillack C, Lohse P, Brand S. Development of a uniform, very aggressive disease phenotype in all homozygous carriers of the NOD2 mutation p.Leu1007fsX1008 with Crohn's disease and active smoking status resulting in ileal stenosis requiring surgery. PloS one 2020; 15:e0236421.

Stallhofer J, Beigel F, Schnitzler F, Tillack-Schreiber C, Glas J, Lohse P, Wetzke M, Konrad-Zerna A, Friedrich M, Brand S. Lipocalin-2 Is a Disease Activity Marker in Inflammatory Bowel Disease Regulated by IL-17A, IL-22, and TNF-α and Modulated by IL23R Genotype Status. Inflamm Bowel Dis 2015; 21:2327-40.

Schnitzler F, Grüner N, Rust C, Guba M, Denk G, Zachoval R, Göke B, Tillack C, Beigel F, Olszak T, Angelberger M, Wolf C, Karbalai N, Habicht A, Fischereder M, Schönermarck U, Stallhofer J, Friedrich M, Brand S. Solid Organ Transplantation in Patients with Inflammatory Bowel Diseases (IBD): Analysis of Transplantation Outcome and IBD Activity in a Large Single Center Cohort. PloS one 2015; 10:e0135807.

Schnitzler F, Lohse P, Glas J, Göke B, Beigel F, Tillack C, Olszak T, Diegelmann J, Angelberger M, Stallhofer J, Wolf C, Friedrich M, Brand S. The NOD2 Single Nucleotide Polymorphism rs72796353 (IVS4+10 A>C) Is a Predictor for Perianal Fistulas in Patients with Crohn's Disease in the Absence of Other NOD2 Mutations. PloS one 2015; 10:e0116044.

Schnitzler F, Lohse P, Glas J, Göke B, Stallhofer J, Tillack C, Beigel F, Olszak T, Diegelmann J, Angelberger M, Wolf C, Friedrich M, Brand S. The NOD2 p.Leu1007fsX1008 mutation (rs2066847) is a stronger predictor of the clinical course of Crohn's disease than the FOXO3A intron variant rs12212067. PloS one 2014; 9:e108503.

Beigel F, Deml M, Schnitzler F, Breiteneicher S, Göke B, Ochsenkühn T, Brand S. Rate and predictors of mucosal healing in patients with inflammatory bowel disease treated with anti-TNF-alpha antibodies. PloS one 2014; 9:e99293.

Beigel F, Brand S, Seiderer J, Göke B, Laubender R, Van Steen K, John J, Breiteneicher S, Tillack C, Schnitzler F, Steinborn A, Ochsenkühn T. Risk of malignancies in patients with inflammatory bowel disease treated with thiopurines or anti-TNF alpha antibodies. Pharmacoepidemiol Drug Saf 2014; 23:735-44.

Beigel F, Ochsenkühn T, Seiderer J, Göke B, Weidinger M, Breiteneicher S, Schnitzler F, Tillack C, Laubender R, Löhr B, Brand S. Iron status and analysis of efficacy and safety of ferric carboxymaltose treatment in patients with inflammatory bowel disease. Digestion 2011; 85:47-54.

Schnitzler F, Seiderer J, Stallhofer J, Brand S. Dominant disease-causing effect of NOD2 mutations in a family with all family members affected by Crohn's disease. Inflamm Bowel Dis 2011; 18:395-6.

Beigel F, Schnitzler F, Paul Laubender R, Pfennig S, Weidinger M, Göke B, Seiderer J, Ochsenkühn T, Brand S. Formation of antinuclear and double-strand DNA antibodies and frequency of lupus-like syndrome in anti-TNF-α antibody-treated patients with inflammatory bowel disease. Inflamm Bowel Dis 2011; 17:91-8.

Jürgens M, Herrmann K, Lohse P, Göke B, Kreis M, Schnitzler F, Weidinger M, Beigel F, Tillack C, Pfennig S, Wagner J, Wetzke M, Glas J, Seiderer J, Laubender R, Brand S, Ochsenkühn T. The presence of fistulas and NOD2 homozygosity strongly predict intestinal stenosis in Crohn's disease independent of the IL23R genotype. J Gastroenterol 2010; 45:721-31.

Jürgens M, Ochsenkühn T, Glas J, Göke B, Lohse P, Tillack C, Schnitzler F, Stallhofer J, Pfennig S, Beigel F, Wetzke M, Wagner J, Seiderer J, Weidinger M, Hartl F, Laubender R, Brand S. Disease activity, ANCA, and IL23R genotype status determine early response to infliximab in patients with ulcerative colitis. Am J Gastroenterol 2010; 105:1811-9.

Dambacher J, Lohse P, Ochsenkühn T, Göke B, Diebold J, Otte J, Tillack C, Konrad A, Hofbauer K, Pfennig S, Schnitzler F, Sisic Z, Seiderer J, Staudinger T, Brand S. Macrophage migration inhibitory factor (MIF) -173G/C promoter polymorphism influences upper gastrointestinal tract involvement and disease activity in patients with Crohn's disease. Inflamm Bowel Dis 2007; 13:71-82.

Seiderer J, Lohse P, Göke B, Sackmann M, Tillack C, Dambacher J, Hofbauer K, Herrmann K, Pfennig S, Staudinger T, Brand S, Schnitzler F, Ochsenkühn T. Homozygosity for the CARD15 frameshift mutation 1007fs is predictive of early onset of Crohn's disease with ileal stenosis, entero-enteral fistulas, and frequent need for surgical intervention with high risk of re-stenosis. Scand J Gastroenterol 2006; 41:1421-32.

Seiderer J, Lohse P, Göke B, Schoenberg S, Pfennig S, Crispin A, Hatz R, Schnitzler F, Herrmann K, Brand S, Ochsenkühn T. Predictive value of the CARD15 variant 1007fs for the diagnosis of intestinal stenoses and the need for surgery in Crohn's disease in clinical practice: results of a prospective study. Inflamm Bowel Dis 2006; 12:1114-21.

Glas J, Folwaczny M, Ochsenkühn T, Brand S, Schnitzler F, Seiderer J, Lohse P, Schiemann U, Griga T, Epplen J, Klein W, Wetzke M, Mussack T, Müller-Myhsok B, Tonenchi L, Török H, Folwaczny C. Role of the NFKB1 -94ins/delATTG promoter polymorphism in IBD and potential interactions with polymorphisms in the CARD15/NOD2, IKBL, and IL-1RN genes. Inflamm Bowel Dis 2006; 12:606-11.

Schnitzler F, Ochsenkühn T, Göke B, Tillack C, Seiderer J, Hofbauer K, Pfennig S, Staudinger T, Brand S, Lohse P. Eight novel CARD15 variants detected by DNA sequence analysis of the CARD15 gene in 111 patients with inflammatory bowel disease. Immunogenetics 2006; 58:99-106.

Brand S, Ochsenkühn T, Göke B, Konrad A, Tillack C, Seiderer J, Pfennig S, Staudinger T, Schnitzler F, Dambacher J, Hofbauer K, Lohse P. Increased expression of the chemokine fractalkine in Crohn's disease and association of the fractalkine receptor T280M polymorphism with a fibrostenosing disease Phenotype. Am J Gastroenterol 2006; 101:99-106.

Brand S, Lohse P, Göke B, Crispin A, Konrad A, Tillack C, Seiderer J, Dambacher J, Hofbauer K, Pfennig S, Schnitzler F, Staudinger T, Ochsenkühn T. The role of Toll-like receptor 4 Asp299Gly and Thr399Ile polymorphisms and CARD15/NOD2 mutations in the susceptibility and phenotype of Crohn's disease. Inflamm Bowel Dis 2005; 11:645-52.

Projekte (0)

Keine Resultate gefunden.

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