



Matthias Jürgens

Contact

Matthias Jürgens

Publications (9)

Glas J, Czamara D, Diegelmann J, Lohse P, Ochsenkühn T, Göke B, Müller-Myhsok B, Weidinger M, Laubender R, Olszak T, Jürgens M, Beigel F, Pfennig S, Tillack C, Seiderer J, Brand S. The NOD2 single nucleotide polymorphisms rs2066843 and rs2076756 are novel and common Crohn's disease susceptibility gene variants. *PLoS one* 2010; 5:e14466.

Jürgens M, Ochsenkühn T, Göke B, Tillack C, Beigel F, Hasbargen U, Hübener C, Filik L, Brand S, Seiderer J. Safety of adalimumab in Crohn's disease during pregnancy: case report and review of the literature. *Inflamm Bowel Dis* 2010; 16:1634-6.

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.

Beigel F, Jürgens M, Filik L, Bader L, Lück C, Göke B, Ochsenkühn T, Brand S, Seiderer J. Severe *Legionella pneumophila* pneumonia following infliximab therapy in a patient with Crohn's disease. *Inflamm Bowel Dis* 2009; 15:1240-4.

Beigel F, Jürgens M, Tillack C, Subklewe M, Mayr D, Göke B, Brand S, Ochsenkühn T. Hepatosplenic T-cell lymphoma in a patient with Crohn's disease. *Nat Rev Gastroenterol Hepatol* 2009; 6:433-6.

Glas J, Klein W, Epplen J, Schiemann U, Mussack T, Lohse P, Göke B, Ochsenkühn T, Folwaczny M, Müller-Myhsok B, Griga T, Jürgens M, Stallhofer J, Seiderer J, Pasciuto G, Tillack C, Diegelmann J, Pfennig S, Konrad A, Schmechel S, Wetzke M, Török H, Brand S. rs224136 on chromosome 10q21.1 and variants in PHOX2B, NCF4, and FAM92B are not major genetic risk factors for susceptibility to Crohn's disease in the German population. *Am J Gastroenterol* 2009; 104:665-72.

Seiderer J, Lohse P, Müller-Myhsok B, Ochsenkühn T, Göke B, Konrad A, Schmechel S, Jürgens M, Pfennig S, Tillack C, Stallhofer J, Glas J, Diegelmann J, Elben I, Brand S. Role of the novel Th17 cytokine IL-17F in inflammatory bowel disease (IBD): upregulated colonic IL-17F expression in active Crohn's disease and analysis of the IL17F p.His161Arg polymorphism in IBD. *Inflamm Bowel Dis* 2008; 14:437-45.

Seiderer J, Reinecker H, Lohse P, Ochsenkühn T, Göke B, Müller-Myhsok B, Jürgens M, Pfennig S, Niess J, Glas J, Tillack C, Leistner D, Dambacher J, Brand S. Genotype-phenotype analysis of the CXCL16 p.Ala181Val polymorphism in inflammatory bowel disease. *Clin Immunol* 2008; 127:49-55.

Projects (0)

No results found.

Kantonsspital St.Gallen

Rorschacher Strasse 95

CH-9007 St.Gallen

T: +41 71 494 11 11

support.forschung@kssg.ch