

Timo J Hautala

List of Publications by Year in descending order

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Version: 2024-02-01

48
papers

1,488
citations

361413

20
h-index

330143

37
g-index

50
all docs

50
docs citations

50
times ranked

2080
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Central Nervous System and Ocular Manifestations in Puumala Hantavirus Infection. <i>Viruses</i> , 2021, 13, 1040. | 3.3 | 9 |
| 2 | Clinical characteristics and evaluation of the incidence of cryptococcosis in Finland 2004–2018. <i>Infectious Diseases</i> , 2021, 53, 684-690. | 2.8 | 3 |
| 3 | Loss of DIAPH1 causes SCBMS, combined immunodeficiency, and mitochondrial dysfunction. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 599-611. | 2.9 | 23 |
| 4 | Effect of first-line antifungal treatment on ocular complication risk in <i>Candida</i> or yeast blood stream infection. <i>BMJ Open Ophthalmology</i> , 2021, 6, e000837. | 1.6 | 4 |
| 5 | A Family With A20 Haploinsufficiency Presenting With Novel Clinical Manifestations and Challenges for Treatment. <i>Journal of Clinical Rheumatology</i> , 2021, 27, e583-e587. | 0.9 | 9 |
| 6 | Aetiology of posterior uveitis in a tertiary centre in Finland. <i>Acta Ophthalmologica</i> , 2020, 98, e135-e136. | 1.1 | 1 |
| 7 | Herpes simplex virus 2 encephalitis in a patient heterozygous for a TLR3 mutation. <i>Neurology: Genetics</i> , 2020, 6, e532. | 1.9 | 6 |
| 8 | Heterozygous TLR3 Mutation in Patients with Hantavirus Encephalitis. <i>Journal of Clinical Immunology</i> , 2020, 40, 1156-1162. | 3.8 | 12 |
| 9 | Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 901-911. | 2.9 | 78 |
| 10 | Outcomes for Nitazoxanide Treatment in a Case Series of Patients with Primary Immunodeficiencies and Rubella Virus-Associated Granuloma. <i>Journal of Clinical Immunology</i> , 2019, 39, 112-117. | 3.8 | 19 |
| 11 | Primary Immunodeficiency, a Possible Cause of Neutrophilic Necrotizing Dermatitis. <i>JAMA Dermatology</i> , 2019, 155, 863. | 4.1 | 5 |
| 12 | ADA2 deficiency: Clonal lymphoproliferation in a subset of patients. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1534-1537.e8. | 2.9 | 71 |
| 13 | Nitazoxanide May Modify the Course of Progressive Multifocal Leukoencephalopathy. <i>Journal of Clinical Immunology</i> , 2018, 38, 4-6. | 3.8 | 5 |
| 14 | Haploinsufficiency of A20 impairs protein–protein interactome and leads into caspase-8-dependent enhancement of NLRP3 inflammasome activation. <i>RMD Open</i> , 2018, 4, e000740. | 3.8 | 26 |
| 15 | Damaging heterozygous mutations in NFKB1 lead to diverse immunologic phenotypes. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 782-796. | 2.9 | 113 |
| 16 | Combined immunodeficiency and hypoglycemia associated with mutations in hypoxia upregulated 1. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 1391-1393.e11. | 2.9 | 14 |
| 17 | Fatal Puumala Hantavirus Disease: Involvement of Complement Activation and Vascular Leakage in the Pathobiology. <i>Open Forum Infectious Diseases</i> , 2017, 4, ofx229. | 0.9 | 15 |
| 18 | Inhibition of rubella virus replication by the broad-spectrum drug nitazoxanide in cell culture and in a patient with a primary immune deficiency. <i>Antiviral Research</i> , 2017, 147, 58-66. | 4.1 | 36 |

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|----|---|-----|-----------|
| 19 | Long-term hormonal follow-up after human Puumala hantavirus infection. <i>Clinical Endocrinology</i> , 2016, 84, 85-91. | 2.4 | 18 |
| 20 | Rubella persistence in epidermal keratinocytes and granuloma M2 macrophages in patients with primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1436-1439.e11. | 2.9 | 73 |
| 21 | Enrichment of rare variants in population isolates: single AICDA mutation responsible for hyper-IgM syndrome type 2 in Finland. <i>European Journal of Human Genetics</i> , 2016, 24, 1473-1478. | 2.8 | 22 |
| 22 | Large Granular Lymphocyte Infiltration in the Bone Marrow in Children and Young Adults May Suggest Primary Immune Deficiency. <i>Blood</i> , 2015, 126, 1024-1024. | 1.4 | 1 |
| 23 | Plasma B-type natriuretic peptide (BNP) in acute Puumala hantavirus infection. <i>Annals of Medicine</i> , 2014, 46, 38-43. | 3.8 | 5 |
| 24 | TLR3 deficiency in herpes simplex encephalitis. <i>Neurology</i> , 2014, 83, 1888-1897. | 1.1 | 128 |
| 25 | Elevated Cerebrospinal Fluid Neopterin Concentration Is Associated with Disease Severity in Acute Puumala Hantavirus Infection. <i>Clinical and Developmental Immunology</i> , 2013, 2013, 1-4. | 3.3 | 12 |
| 26 | Signs of general inflammation and kidney function are associated with the ocular features of acute Puumala hantavirus infection. <i>Scandinavian Journal of Infectious Diseases</i> , 2012, 44, 956-962. | 1.5 | 9 |
| 27 | True bacteremias caused by coagulase negative Staphylococcus are difficult to distinguish from blood culture contaminants. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 2012, 31, 2639-2644. | 2.9 | 24 |
| 28 | Late-presenting vascular graft infection caused by Aspergillus in an immunocompetent patient. <i>Mycoses</i> , 2012, 55, 95-98. | 4.0 | 6 |
| 29 | IL-2 may possess neuroprotective properties in glaucomatous optic neuropathy. <i>Acta Ophthalmologica</i> , 2012, 90, e246-7. | 1.1 | 5 |
| 30 | Young male patients are at elevated risk of developing serious central nervous system complications during acute Puumala hantavirus infection. <i>BMC Infectious Diseases</i> , 2011, 11, 217. | 2.9 | 22 |
| 31 | Prospective study on ocular findings in acute Puumala hantavirus infection in hospitalised patients. <i>British Journal of Ophthalmology</i> , 2011, 95, 559-562. | 3.9 | 29 |
| 32 | Clinical Candida krusei isolates remain susceptible during extensive exposure to antifungal drugs. <i>Medical Mycology</i> , 2010, 48, 79-84. | 0.7 | 3 |
| 33 | Central nervous system-related symptoms and findings are common in acute Puumala hantavirus infection. <i>Annals of Medicine</i> , 2010, 42, 344-351. | 3.8 | 43 |
| 34 | Adaptive or maladaptive response to adenoviral adrenomedullin gene transfer is context-dependent in the heart. <i>Journal of Gene Medicine</i> , 2008, 10, 867-877. | 2.8 | 6 |
| 35 | Blood culture Gram stain and simple clinical categorisation can be used to identify patients at risk for delay in appropriate antibiotic treatment. <i>International Journal of Antimicrobial Agents</i> , 2008, 32, 546-547. | 2.5 | 1 |
| 36 | Major Age Group-Specific Differences in Conjunctival Bacteria and Evolution of Antimicrobial Resistance Revealed by Laboratory Data Surveillance. <i>Current Eye Research</i> , 2008, 33, 907-911. | 1.5 | 16 |

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|----|--|-----|-----------|
| 37 | Puumala virus RNA in cerebrospinal fluid in a patient with uncomplicated nephropathia epidemica. <i>Journal of Clinical Virology</i> , 2007, 40, 248-251. | 3.1 | 27 |
| 38 | A cluster of <i>Candida krusei</i> infections in a haematological unit. <i>BMC Infectious Diseases</i> , 2007, 7, 97. | 2.9 | 25 |
| 39 | Endostatin inhibits VEGF-A induced osteoclastic bone resorption in vitro. <i>BMC Musculoskeletal Disorders</i> , 2006, 7, 56. | 1.9 | 31 |
| 40 | Adrenomedullin gene transfer induces neointimal apoptosis and inhibits neointimal hyperplasia in injured rat artery. <i>Journal of Gene Medicine</i> , 2006, 8, 452-458. | 2.8 | 7 |
| 41 | Blood culture Gram stain and clinical categorization based empirical antimicrobial therapy of bloodstream infection. <i>International Journal of Antimicrobial Agents</i> , 2005, 25, 329-333. | 2.5 | 31 |
| 42 | Adenoviral VEGF gene transfer induces angiogenesis and promotes bone formation in healing osseous tissues. <i>Journal of Gene Medicine</i> , 2003, 5, 560-566. | 2.8 | 125 |
| 43 | Hypophyseal Hemorrhage and Panhypopituitarism during Puumala Virus Infection: Magnetic Resonance Imaging and Detection of Viral Antigen in the Hypophysis. <i>Clinical Infectious Diseases</i> , 2002, 35, 96-101. | 5.8 | 104 |
| 44 | Adenoviral Gene Transfer Restores Lysyl Hydroxylase Activity in Type VI Ehlers-Danlos Syndrome. <i>Journal of Investigative Dermatology</i> , 2001, 116, 602-605. | 0.7 | 2 |
| 45 | A compound heterozygote patient with Ehlers-Danlos syndrome type VI has a deletion in one allele and a splicing defect in the other allele of the lysyl hydroxylase gene. <i>Human Mutation</i> , 1998, 11, 55-61. | 2.5 | 22 |
| 46 | Structure and Expression of the Human Lysyl Hydroxylase Gene (PLOD): Introns 9 and 16 Contain Alu Sequences at the Sites of Recombination in Ehlers-Danlos Syndrome Type VI Patients. <i>Genomics</i> , 1994, 24, 464-471. | 2.9 | 48 |
| 47 | A Large Duplication in the Gene for Lysyl Hydroxylase Accounts for the Type VI Variant of Ehlers-Danlos Syndrome in Two Siblings. <i>Genomics</i> , 1993, 15, 399-404. | 2.9 | 70 |
| 48 | Cloning of human lysyl hydroxylase: Complete cDNA-derived amino acid sequence and assignment of the gene (PLOD) to chromosome 1p36.3-p36.2. <i>Genomics</i> , 1992, 13, 62-69. | 2.9 | 124 |