# Alberto Tommasini

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5718892/publications.pdf

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|---|---|---|---|
| 176 papers | 5,488 citations | 87843 38 h-index | 95218 68 g-index |
| 179 all docs | 179 docs citations | 179 times ranked | 7281 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | GNE-related thrombocytopenia: evidence for a mutational hotspot in the ADP/substrate domain of the GNE bifunctional enzyme. Haematologica, 2022, 107, 750-754. | 1.7 | 11 |
| 2 | Cytofluorimetric assay to investigate variability in blinatumomab in vitro response. Frontiers in Bioscience, 2022, 27, 039. | 0.8 | 0 |
| 3 | Evolution of SARS-CoV-2 IgG Seroprevalence in Children and Factors Associated with Seroconversion: Results from a Multiple Time-Points Study in Friuli-Venezia Giulia Region, Italy. Children, 2022, 9, 246. | 0.6 | 6 |
| 4 | Druggable monogenic immune defects hidden in diverse medical specialties: Focus on overlap syndromes. World Journal of Clinical Pediatrics, 2022, 11, 136-150. | 0.6 | 3 |
| 5 | Ultrasonographic Assessment for Tenosynovitis in Juvenile Idiopathic Arthritis with Ankle Involvement: Diagnostic and Therapeutic Significance. Children, 2022, 9, 509. | 0.6 | 1 |
| 6 | Case Report: Refractory Autoimmune Gastritis Responsive to Abatacept in LRBA Deficiency. Frontiers in Immunology, 2021, 12, 619246. | 2.2 | 8 |
| 7 | SARS-CoV-2 Infection and Inflammatory Response in a Twin Pregnancy. International Journal of Environmental Research and Public Health, 2021, 18, 3075. | 1.2 | 2 |
| 8 | Hydroxychloroquine modulates immunological pathways activated by RNA:DNA hybrids in Aicardiâ€“GoutiÃ¨res syndrome patients carrying RNASEH2 mutations. Cellular and Molecular Immunology, 2021, 18, 1593-1595. | 4.8 | 3 |
| 9 | Plasmacytoid Dendritic Cells Depletion and Elevation of IFN-Î³ Dependent Chemokines CXCL9 and CXCL10 in Children With Multisystem Inflammatory Syndrome. Frontiers in Immunology, 2021, 12, 654587. | 2.2 | 39 |
| 10 | Immunity and Genetics at the Revolving Doors of Diagnostics in Primary Immunodeficiencies. Diagnostics, 2021, 11, 532. | 1.3 | 2 |
| 11 | Topical clobetasol: an overlooked cause of Cushing syndrome. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2021, 21, . | 0.6 | 1 |
| 12 | Priming of the cGAS-STING-TBK1 Pathway Enhances LPS-Induced Release of Type I Interferons. Cells, 2021, 10, 785. | 1.8 | 16 |
| 13 | Le sindromi autoinfiammatorie: quando non Ã¨ solo PFAPA. Medico E Bambino, 2021, 40, 221-225. | 0.1 | 2 |
| 14 | Vasculitis, Autoimmunity, and Cytokines: How the Immune System Can Harm the Brain. International Journal of Environmental Research and Public Health, 2021, 18, 5585. | 1.2 | 1 |
| 15 | Genetic and immunologic findings in children with recurrent aphthous stomatitis with systemic inflammation. Pediatric Rheumatology, 2021, 19, 70. | 0.9 | 12 |
| 16 | In Vitro Effects of Sulforaphane on Interferon-Driven Inflammation and Exploratory Evaluation in Two Healthy Volunteers. Molecules, 2021, 26, 3602. | 1.7 | 2 |
| 17 | Post-COVID-19 symptoms 6Âmonths after acute infection among hospitalized and non-hospitalized patients. Clinical Microbiology and Infection, 2021, 27, 1507-1513. | 2.8 | 197 |
| 18 | Case Report: EBV Chronic Infection and Lymphoproliferation in Four APDS Patients: The Challenge of Proper Characterization, Therapy, and Follow-Up. Frontiers in Pediatrics, 2021, 9, 703853. | 0.9 | 8 |

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|---|---|---|---|
| 19 | Clinical and Cytometric Study of Immune Involvement in a Heterogeneous Cohort of Subjects With RASopathies and mTORopathies. Frontiers in Pediatrics, 2021, 9, 703613. | 0.9 | 5 |
| 20 | Swollen Ankle with a Hole: Brodie Abscess. Journal of Pediatrics, 2021, 236, 319-320. | 0.9 | 0 |
| 21 | Mevalonate Kinase Deficiency and Squalene Synthase Inhibitor (TAK-475): The Balance to Extinguish the Inflammation. Biomolecules, 2021, 11, 1438. | 1.8 | 1 |
| 22 | Le interferonopatie di tipo I. Medico E Bambino, 2021, 40, 509-514. | 0.1 | 0 |
| 23 | A Novel ELISA-Based Peptide Biosensor Assay for Screening ABL1 Activity in vitro: A Challenge for Precision Therapy in BCR-ABL1 and BCR-ABL1 Like Leukemias. Frontiers in Pharmacology, 2021, 12, 749361. | 1.6 | 4 |
| 24 | Genetic Variants Assessing Crohnâ€™s Disease Pattern in Pediatric Inflammatory Bowel Disease Patients by a Clinical Exome Survey. Bioinformatics and Biology Insights, 2021, 15, 117793222110552. | 1.0 | 1 |
| 25 | Complex regional pain syndrome of the foot in a girl with post-SARS-CoV-2 chilblains. Clinical and Experimental Rheumatology, 2021, , . | 0.4 | 0 |
| 26 | B-cell depletion induces prolonged remission in patients with giant cell hepatitis and autoimmune hemolytic anemia. Clinics and Research in Hepatology and Gastroenterology, 2020, 44, 66-72. | 0.7 | 6 |
| 27 | Diagnostic Approach to Monogenic Inflammatory Bowel Disease in Clinical Practice: A Ten-Year Multicentric Experience. Inflammatory Bowel Diseases, 2020, 26, 720-727. | 0.9 | 32 |
| 28 | Biological and Clinical Changes in a Pediatric Series Treated with Off-Label JAK Inhibitors. International Journal of Molecular Sciences, 2020, 21, 7767. | 1.8 | 18 |
| 29 | High prevalence of rare FBLIM1 gene variants in an Italian cohort of patients with Chronic Non-bacterial Osteomyelitis (CNO). Pediatric Rheumatology, 2020, 18, 55. | 0.9 | 9 |
| 30 | The Italian Registry for Primary Immunodeficiencies (Italian Primary Immunodeficiency Network;) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 3 | 2.0 | 15 |
| 31 | From bone to skin. Journal of Paediatrics and Child Health, 2020, 56, 1481-1481. | 0.4 | 0 |
| 32 | A Case of Uveitis in a Patient With Juvenile Myelomonocytic Leukemia Successfully Treated With Adalimumab. Journal of Pediatric Hematology/Oncology, 2020, 42, e373-e376. | 0.3 | 5 |
| 33 | Consensus of the Italian Primary Immunodeficiency Network on transition management from pediatric to adult care in patients affected with childhood-onset inborn errors of immunity. Journal of Allergy and Clinical Immunology, 2020, 146, 967-983. | 1.5 | 12 |
| 34 | Recent Insight into SARS-CoV2 Immunopathology and Rationale for Potential Treatment and Preventive Strategies in COVID-19. Vaccines, 2020, 8, 224. | 2.1 | 47 |
| 35 | Biomarkers and Precision Therapy for Primary Immunodeficiencies: An In Vitro Study Based on Induced Pluripotent Stem Cells From Patients. Clinical Pharmacology and Therapeutics, 2020, 108, 358-367. | 2.3 | 8 |
| 36 | Higher interferon score and normal complement levels may identify a distinct clinical subset in children with systemic lupus erythematosus. Arthritis Research and Therapy, 2020, 22, 91. | 1.6 | 22 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Case Report: Use of Anakinra in Multisystem Inflammatory Syndrome During COVID-19 Pandemic. Frontiers in Pediatrics, 2020, 8, 624248. | 0.9 | 28 |
| 38 | Acute rheumatic fever prophylaxis in high-income countries: clinical observations from an Italian multicentre, retrospective study. Clinical and Experimental Rheumatology, 2020, 38, 1016-1020. | 0.4 | 2 |
| 39 | Cutaneous manifestations in mevalonate kinase deficient patients treated with canakinumab. Clinical and Experimental Rheumatology, 2020, 38, 760-762. | 0.4 | 1 |
| 40 | Pulmonary arterial hypertension in interferonophaties: a case report and a review of the literature. Pulmonary Circulation, 2019, 9, 1-4. | 0.8 | 9 |
| 41 | Combined Analysis of Methylation and Gene Expression Profiles in Separate Compartments of Small Bowel Mucosa Identified Celiac Disease Patientsâ€™ Signatures. Scientific Reports, 2019, 9, 10020. | 1.6 | 15 |
| 42 | Clinical, Immunological, and Molecular Features of Typical and Atypical Severe Combined Immunodeficiency: Report of the Italian Primary Immunodeficiency Network. Frontiers in Immunology, 2019, 10, 1908. | 2.2 | 41 |
| 43 | Tregs and Th17 lymphocytes in human DYRK1A haploinsufficiency. Immunology Letters, 2019, 214, 52-54. | 1.1 | 1 |
| 44 | Familial hypogammaglobulinemia with high RTE and naÃ¯ve T lymphocytes. Inflammation Research, 2019, 68, 901-904. | 1.6 | 2 |
| 45 | An Easy and Reliable Strategy for Making Type I Interferon Signature Analysis Comparable among Research Centers. Diagnostics, 2019, 9, 113. | 1.3 | 14 |
| 46 | Monocyteâ€predominant engraftment, cytokine levels and early transplantâ€related complications in pediatric hematopoietic stem cell recipients. Cancer Medicine, 2019, 8, 890-901. | 1.3 | 4 |
| 47 | Off-Label Use of Sirolimus and Everolimus in a Pediatric Center: A Case Series and Review of the Literature. Paediatric Drugs, 2019, 21, 185-193. | 1.3 | 37 |
| 48 | Hughes Stovin: Sustained remission and regression of pulmonary aneurysms with antiâ€"tumor necrosis factor treatment. Pediatric Pulmonology, 2019, 54, E13-E15. | 1.0 | 11 |
| 49 | AB1036â€...AN UNSOLVED CASE: IS THIS A CANDLE-LIKE SYNDROME?. , 2019, , . | | 0 |
| 50 | AB1062â€...INTER-LABORATORY COMPARISON OF TYPE I INTERFERON SIGNATURE ANALYSES: PAVING THE WAY TO SHARE RECOMMENDATIONS.. , 2019, , . | | 0 |
| 51 | AB1105â€...A NOVEL AUTOINFLAMMATORY AND LYMPHOPROLIFERATIVE SYNDROME ASSOCIATED WITH PIM1 MUTATIONS. , 2019, , . | | 0 |
| 52 | FRI0579â€...PREDNISONE VERSUS STANDARD THERAPY IN SYDENHAMâ€™S CHOREA: RESULTS FROM A RETROSPECTIVE STUDY. , 2019, , . | | 0 |
| 53 | Antibodies reacting to mimotopes of Simian virus 40 large T antigen, the viral oncoprotein, in sera from children. Journal of Cellular Physiology, 2019, 234, 3170-3179. | 2.0 | 4 |
| 54 | Is autophagy an elective strategy to protect neurons from dysregulated cholesterol metabolism?. Neural Regeneration Research, 2019, 14, 582. | 1.6 | 4 |

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|---|---|---|---|
| 55 | Lowâ€dose sirolimus in two cousins with autoimmune lymphoproliferative syndromeâ€associated infection. Pediatrics International, 2018, 60, 315-317. | 0.2 | 6 |
| 56 | Theophylline as a precision therapy in a young girl with PIK3R1 immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 2165-2167. | 2.0 | 19 |
| 57 | Genetic profile of patients with early onset inflammatory bowel disease. Gene, 2018, 645, 18-29. | 1.0 | 29 |
| 58 | Novel NOD2 Mutation in Early-Onset Inflammatory Bowel Phenotype. Inflammatory Bowel Diseases, 2018, 24, 1204-1212. | 0.9 | 13 |
| 59 | Long term outcome of eight patients with type 1 Leukocyte Adhesion Deficiency (LAD-1): Not only infections, but high risk of autoimmune complications. Clinical Immunology, 2018, 191, 75-80. | 1.4 | 33 |
| 60 | The Challenge of Next Generation Sequencing in a Boy With Severe Mononucleosis and EBV-related Lymphoma. Journal of Pediatric Hematology/Oncology, 2018, 40, e323-e326. | 0.3 | 2 |
| 61 | The Complex Interplay between Lipids, Immune System and Interleukins in Cardio-Metabolic Diseases. International Journal of Molecular Sciences, 2018, 19, 4058. | 1.8 | 46 |
| 62 | Neuronal Dysfunction Associated with Cholesterol Deregulation. International Journal of Molecular Sciences, 2018, 19, 1523. | 1.8 | 9 |
| 63 | Fecal Calprotectin to Detect Inflammatory Bowel Disease in Juvenile Idiopathic Arthritis. Journal of Rheumatology, 2018, 45, 1418-1421. | 1.0 | 4 |
| 64 | Role of the Long Nonâ€Coding RNA Growth Arrestâ€Specific 5 in Glucocorticoid Response in Children with Inflammatory Bowel Disease. Basic and Clinical Pharmacology and Toxicology, 2018, 122, 87-93. | 1.2 | 41 |
| 65 | Repositioning Drugs for Rare Immune Diseases: Hopes and Challenges for a Precision Medicine. Current Medicinal Chemistry, 2018, 25, 2764-2782. | 1.2 | 7 |
| 66 | Reappraisal of Antimalarials in Interferonopathies: New Perspectives for Old Drugs. Current Medicinal Chemistry, 2018, 25, 2797-2810. | 1.2 | 13 |
| 67 | Neither hereditary periodic fever nor periodic fever, aphthae, pharingitis, adenitis: Undifferentiated periodic fever in a tertiary pediatric center. World Journal of Clinical Pediatrics, 2018, 7, 49-55. | 0.6 | 9 |
| 68 | Innovation for rare diseases and bioethical concerns: A thin thread between medical progress and suffering. World Journal of Clinical Pediatrics, 2018, 7, 75-82. | 0.6 | 4 |
| 69 | Describing Kawasaki shock syndrome: results from a retrospective study and literature review. Clinical Rheumatology, 2017, 36, 223-228. | 1.0 | 68 |
| 70 | Mevalonate kinase deficiency: therapeutic targets, treatments, and outcomes. Expert Opinion on Orphan Drugs, 2017, 5, 515-524. | 0.5 | 1 |
| 71 | ADA2 deficiency (DADA2) as an unrecognised cause of early onset polyarteritis nodosa and stroke: a multicentre national study. Annals of the Rheumatic Diseases, 2017, 76, 1648-1656. | 0.5 | 199 |
| 72 | Cryopyrin-associated Periodic Syndromes in Italian Patients: Evaluation of the Rate of Somatic NLRP3 Mosaicism and Phenotypic Characterization. Journal of Rheumatology, 2017, 44, 1667-1673. | 1.0 | 28 |

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|---|---|---|---|
| 73 | Type I interferon-mediated autoinflammation due to DNase II deficiency. Nature Communications, 2017, 8, 2176. | 5.8 | 164 |
| 74 | Curcumin Anti-Apoptotic Action in a Model of Intestinal Epithelial Inflammatory Damage. Nutrients, 2017, 9, 578. | 1.7 | 27 |
| 75 | Dealing with Chronic Non-Bacterial Osteomyelitis: a practical approach. Pediatric Rheumatology, 2017, 15, 87. | 0.9 | 54 |
| 76 | Ex vivo response to mucosal bacteria and muramyl dipeptide in inflammatory bowel disease. World Journal of Gastroenterology, 2016, 22, 9734. | 1.4 | 2 |
| 77 | Innovative Target Therapies Are Able to Block the Inflammation Associated with Dysfunction of the Cholesterol Biosynthesis Pathway. International Journal of Molecular Sciences, 2016, 17, 47. | 1.8 | 8 |
| 78 | Differential expression of <scp>GAS</scp>5 in rapamycinâ€induced reversion of glucocorticoid resistance. Clinical and Experimental Pharmacology and Physiology, 2016, 43, 602-605. | 0.9 | 15 |
| 79 | Putative modifier genes in mevalonate kinase deficiency. Molecular Medicine Reports, 2016, 13, 3181-3189. | 1.1 | 4 |
| 80 | Carbamazepine-induced thrombocytopenic purpura in a child: Insights from a genomic analysis. Blood Cells, Molecules, and Diseases, 2016, 59, 97-99. | 0.6 | 2 |
| 81 | Clinical Characteristics of Patients Carrying the Q703K Variant of the NLRP3 Gene: A 10-year Multicentric National Study. Journal of Rheumatology, 2016, 43, 1093-1100. | 1.0 | 31 |
| 82 | Action of methotrexate and tofacitinib on directly stimulated and bystander-activated lymphocytes. Molecular Medicine Reports, 2016, 14, 574-582. | 1.1 | 1 |
| 83 | Next-generation sequencing and its initial applications for molecular diagnosis of systemic auto-inflammatory diseases. Annals of the Rheumatic Diseases, 2016, 75, 1550-1557. | 0.5 | 57 |
| 84 | Altered pattern of tumor necrosis factor-alpha production in peripheral blood monocytes from Crohn's disease. World Journal of Gastroenterology, 2016, 22, 9117. | 1.4 | 7 |
| 85 | The diagnostic challenge of very early-onset enterocolitis in an infant with XIAP deficiency. BMC Pediatrics, 2015, 15, 208. | 0.7 | 25 |
| 86 | Two-gene mutation in a single patient: Biochemical and functional analysis for a correct interpretation of exome results. Molecular Medicine Reports, 2015, 12, 6128-6132. | 1.1 | 2 |
| 87 | Genetics of inflammatory bowel disease from multifactorial to monogenic forms. World Journal of Gastroenterology, 2015, 21, 12296. | 1.4 | 113 |
| 88 | Patients' Induced Pluripotent Stem Cells to Model Drug Induced Adverse Events: A Role in Predicting Thiopurine Induced Pancreatitis?. Current Drug Metabolism, 2015, 17, 91-98. | 0.7 | 7 |
| 89 | To Extinguish the Fire from Outside the Cell or to Shutdown the Gas Valve Inside? Novel Trends in Anti-Inflammatory Therapies. International Journal of Molecular Sciences, 2015, 16, 21277-21293. | 1.8 | 5 |
| 90 | Long Noncoding RNA GAS5: A Novel Marker Involved in Glucocorticoid Response. Current Molecular Medicine, 2015, 15, 94-99. | 0.6 | 42 |

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|---|---|---|---|
| 91 | Altered germinal center reaction and abnormal B cell peripheral maturation in PI3KR1-mutated patients presenting with HIGM-like phenotype. Clinical Immunology, 2015, 159, 33-36. | 1.4 | 51 |
| 92 | Wharton's jelly derived mesenchymal stromal cells: Biological properties, induction of neuronal phenotype and current applications in neurodegeneration research. Acta Histochemica, 2015, 117, 329-338. | 0.9 | 53 |
| 93 | Genetic profiling of autoinflammatory disorders in patients with periodic fever: a prospective study. Pediatric Rheumatology, 2015, 13, 11. | 0.9 | 22 |
| 94 | Failure of interferon-Î³ pre-treated mesenchymal stem cell treatment in a patient with Crohnâ€™s disease. World Journal of Gastroenterology, 2015, 21, 4379. | 1.4 | 21 |
| 95 | Fever tree revisited: From malaria to autoinflammatory diseases. World Journal of Clinical Pediatrics, 2015, 4, 106. | 0.6 | 4 |
| 96 | Different presentations of mevalonate kinase deficiency: a case series. Clinical and Experimental Rheumatology, 2015, 33, 437-42. | 0.4 | 10 |
| 97 | Clinical significance of hyper-IgA in a paediatric laboratory series. Archives of Disease in Childhood, 2014, 99, 1114-1116. | 1.0 | 4 |
| 98 | Novel Missense Mutation in the NOD2 Gene in a Patient with Early Onset Ulcerative Colitis: Causal or Chance Association?. International Journal of Molecular Sciences, 2014, 15, 3834-3841. | 1.8 | 9 |
| 99 | TNF-Î± SNP rs1800629 and risk of relapse in childhood acute lymphoblastic leukemia: relation to immunophenotype. Pharmacogenomics, 2014, 15, 619-627. | 0.6 | 2 |
| 100 | In vivo detection of polyomaviruses JCV and SV40 in mesenchymal stem cells from human umbilical cords. Pediatric Blood and Cancer, 2014, 61, 1347-1349. | 0.8 | 10 |
| 101 | â€œHyper-IgD syndromeâ€•or â€œmevalonate kinase deficiencyâ€• an old syndrome needing a new name?. Rheumatology International, 2014, 34, 423-424. | 1.5 | 5 |
| 102 | Inhibition of mesenchymal stromal cells by pre-activated lymphocytes and their culture media. Stem Cell Research and Therapy, 2014, 5, 3. | 2.4 | 16 |
| 103 | Very-early onset IBD in male as expression of XIAP mutation. Digestive and Liver Disease, 2014, 46, e91. | 0.4 | 0 |
| 104 | Severe inflammatory bowel disease associated with congenital alteration of transforming growth factor beta signaling. Journal of Crohn's and Colitis, 2014, 8, 770-774. | 0.6 | 28 |
| 105 | Hemophagocytic Lymphohistiocytosis in Total Parenteral Nutrition Dependent Children. Journal of Pediatric Hematology/Oncology, 2014, 36, e440-e442. | 0.3 | 5 |
| 106 | Fate of Lymphocytes after Withdrawal of Tofacitinib Treatment. PLoS ONE, 2014, 9, e85463. | 1.1 | 16 |
| 107 | F402L variant in NLRP12 in subjects with undiagnosed periodic fevers and in healthy controls. Clinical and Experimental Rheumatology, 2014, 32, 993-4. | 0.4 | 16 |
| 108 | Selective IgA Deficiency: Ruling out Coeliac Disease and Selective Antibody Deficiency to Polysaccharides. Journal of Clinical Immunology, 2013, 33, 1149-1149. | 2.0 | 3 |

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|---|---|---|---|
| 109 | Human IL2RA null mutation mediates immunodeficiency with lymphoproliferation and autoimmunity. Clinical Immunology, 2013, 146, 248-261. | 1.4 | 186 |
| 110 | Evolutionary hypothesis of the Mevalonate Kinase Deficiency. Medical Hypotheses, 2013, 80, 67-69. | 0.8 | 10 |
| 111 | Clinical Genetic Testing of Periodic Fever Syndromes. BioMed Research International, 2013, 2013, 1-8. | 0.9 | 10 |
| 112 | Genetic and Functional Profiling of Crohn's Disease: Autophagy Mechanism and Susceptibility to Infectious Diseases. BioMed Research International, 2013, 2013, 1-11. | 0.9 | 10 |
| 113 | Clinical impact ofMEFVmutations in children with periodic fever in a prevalent western European Caucasian population. Annals of the Rheumatic Diseases, 2012, 71, 1961-1965. | 0.5 | 65 |
| 114 | Phylloid Pattern of Hypomelanosis Closely Related to Chromosomal Abnormalities in the 13q Detected by SNP Array Analysis. Dermatology, 2012, 225, 294-297. | 0.9 | 15 |
| 115 | Defective and Excessive Immunities in Pediatric Diseases. Current Pharmaceutical Design, 2012, 18, 5729-5734. | 0.9 | 4 |
| 116 | A common genetic background could explain early-onset Crohnâ€™s disease. Medical Hypotheses, 2012, 78, 520-522. | 0.8 | 15 |
| 117 | The MDM2 inhibitor Nutlin-3 modulates dendritic cellâ€“induced T cell proliferation. Human Immunology, 2012, 73, 342-345. | 1.2 | 28 |
| 118 | Inflammation profile of four early onset Crohn patients. Gene, 2012, 493, 282-285. | 1.0 | 12 |
| 119 | A red baby should not be taken too lightly. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e573-7. | 0.7 | 3 |
| 120 | Clinical immunology Primary Immunodeficiency Diseases in two neighboring pediatric centers: registry data bring out a wide spectrum of diseases with complex clinical presentations. Central-European Journal of Immunology, 2012, 4, 365-370. | 0.4 | 2 |
| 121 | Immunomodulatory drugs in autoimmune lymphoproliferative syndrome (ALPS). Pediatric Blood and Cancer, 2012, 58, 310-310. | 0.8 | 7 |
| 122 | JCV+ Patients with Inflammatory Bowel Disease show elevated plasma levels of MIG and SCF. Inflammatory Bowel Diseases, 2012, 18, 1194-1196. | 0.9 | 5 |
| 123 | Forkhead box protein 3 (FOXP3) mutations lead to increased TH17 cell numbers and regulatory T-cell instability. Journal of Allergy and Clinical Immunology, 2011, 128, 1376-1379.e1. | 1.5 | 54 |
| 124 | Differential action of 3-hydroxyanthranilic acid on viability and activation of stimulated lymphocytes. International Immunopharmacology, 2011, 11, 2242-2245. | 1.7 | 8 |
| 125 | Gastrointestinal Foxp3 expression in normal, inflammatory and neoplastic conditions. Pathology, 2011, 43, 465-471. | 0.3 | 3 |
| 126 | Functional type 1 regulatory T cells develop regardless of FOXP3 mutations in patients with IPEX syndrome. European Journal of Immunology, 2011, 41, 1120-1131. | 1.6 | 72 |

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|---|---|---|---|
| 127 | Longâ€term clinical profile of children with the lowâ€penetrance R92Q mutation of the TNFRSF1A gene. Arthritis and Rheumatism, 2011, 63, 1141-1150. | 6.7 | 99 |
| 128 | The Farnesyltransferase Inhibitors Tipifarnib and Lonafarnib Inhibit Cytokines Secretion in a Cellular Model of Mevalonate Kinase Deficiency. Pediatric Research, 2011, 70, 78-82. | 1.1 | 20 |
| 129 | Ages of celiac disease: From changing environment to improved diagnostics. World Journal of Gastroenterology, 2011, 17, 3665. | 1.4 | 18 |
| 130 | Decreased cholesterol levels reflect a consumption of anti-inflammatory isoprenoids associated with an impaired control of inflammation in a mouse model of mevalonate kinase deficiency. Inflammation Research, 2010, 59, 335-338. | 1.6 | 14 |
| 131 | Follow-Up and Quality of Life of Patients with Cryopyrin-Associated Periodic Syndromes Treated with Anakinra. Journal of Pediatrics, 2010, 157, 310-315.e1. | 0.9 | 105 |
| 132 | Selective resistance to different glucocorticoids in severe autoimmune disorders. Clinical Immunology, 2010, 134, 313-319. | 1.4 | 8 |
| 133 | Therapeutic strategy in p47-phox deficient chronic granulomatous disease presenting as inflammatory bowel disease. Journal of Allergy and Clinical Immunology, 2010, 125, 943-946.e1. | 1.5 | 40 |
| 134 | Targeting farnesyl-transferase as a novel therapeutic strategy for mevalonate kinase deficiency: In vitro and in vivo approaches. Pharmacological Research, 2010, 61, 506-510. | 3.1 | 17 |
| 135 | Natural isoprenoids inhibit LPS-induced-production of cytokines and nitric oxide in aminobisphosphonate-treated monocytes. International Immunopharmacology, 2010, 10, 639-642. | 1.7 | 37 |
| 136 | The immunosuppressive effect of Wharton's jelly stromal cells depends on the timing of their licensing and on lymphocyte activation. Cytotherapy, 2010, 12, 154-160. | 0.3 | 37 |
| 137 | The universe of immune deficiencies in Crohn's disease: a new viewpoint for an old disease?. Scandinavian Journal of Gastroenterology, 2010, 45, 1141-1149. | 0.6 | 17 |
| 138 | Regulatory T-Cell Function Is Impaired in Celiac Disease. Digestive Diseases and Sciences, 2009, 54, 1513-1519. | 1.1 | 59 |
| 139 | Differentiating PFAPA Syndrome From Monogenic Periodic Fevers. Pediatrics, 2009, 124, e721-e728. | 1.0 | 138 |
| 140 | Wild-type FOXP3 is selectively active in CD4+CD25hi regulatory T cells of healthy female carriers of different FOXP3 mutations. Blood, 2009, 114, 4138-4141. | 0.6 | 49 |
| 141 | The efficacy of anakinra in an adolescent with colchicine-resistant familial Mediterranean fever. European Journal of Pediatrics, 2008, 167, 695-696. | 1.3 | 119 |
| 142 | Clinical and molecular profile of a new series of patients with immune dysregulation, polyendocrinopathy, enteropathy, X-linked syndrome: Inconsistent correlation between forkhead box protein 3 expression and disease severity. Journal of Allergy and Clinical Immunology, 2008, 122, 1105-1112.e1. | 1.5 | 199 |
| 143 | Natural Isoprenoids are Able to Reduce Inflammation in a Mouse Model of Mevalonate Kinase Deficiency. Pediatric Research, 2008, 64, 177-182. | 1.1 | 54 |
| 144 | Diagnostics and Therapeutic Insights in a Severe Case of Mevalonate Kinase Deficiency. Pediatrics, 2007, 119, e523-e527. | 1.0 | 34 |

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|---|---|---|---|
| 145 | Autoinflammatory syndromes and coeliac disease: One observation and two hypotheses. Digestive and Liver Disease, 2007, 39, A83-A84. | 0.4 | 0 |
| 146 | T cells stimulated in vitro have a suppressive function but do not contain only regulatory T cells. Clinical and Experimental Immunology, 2007, 150, 561-566. | 1.1 | 8 |
| 147 | Medium-term survival without haematopoietic stem cell transplantation in a case of IPEX: insights into nutritional and immunosuppressive therapy. European Journal of Pediatrics, 2007, 166, 1195-1197. | 1.3 | 25 |
| 148 | Heterozygous nucleotide-binding oligomerization domain-2 mutations affect monocyte maturation in Crohnâ€™s disease. World Journal of Gastroenterology, 2007, 13, 6191. | 1.4 | 1 |
| 149 | Diagnostic challenge of hyper-IgD syndrome in four children with inflammatory gastrointestinal complaints. Scandinavian Journal of Gastroenterology, 2006, 41, 430-436. | 0.6 | 15 |
| 150 | CARD15/NOD2 mutations are not related to abdominal PFAPA. Journal of Pediatrics, 2006, 149, 427. | 0.9 | 10 |
| 151 | Cryptic gluten intolerance in type 1 diabetes: identifying suitable candidates for a gluten free diet. Gut, 2006, 55, 133-134. | 6.1 | 14 |
| 152 | Looking for Celiac Disease: Diagnostic Accuracy of Two Rapid Commercial Assays. American Journal of Gastroenterology, 2006, 101, 1597-1600. | 0.2 | 50 |
| 153 | MVK mutations and associated clinical features in Italian patients affected with autoinflammatory disorders and recurrent fever. European Journal of Human Genetics, 2005, 13, 314-320. | 1.4 | 71 |
| 154 | AMANTADINE IN CHRONIC GRANULOMATOUS DISEASE. Pediatric Hematology and Oncology, 2005, 22, 147-151. | 0.3 | 6 |
| 155 | Characterization of the Anti-Tissue Transglutaminase Antibody Response in Nonobese Diabetic Mice. Journal of Immunology, 2005, 174, 5830-5836. | 0.4 | 23 |
| 156 | Dual sugar gut-permeability testing on blood drop in animal models. Clinica Chimica Acta, 2005, 352, 191-197. | 0.5 | 18 |
| 157 | Testing for Anti-Human Transglutaminase Antibodies in Saliva Is Not Useful for Diagnosis of Celiac Disease. Clinical Chemistry, 2004, 50, 216-219. | 1.5 | 16 |
| 158 | Mass screening for coeliac disease using antihuman transglutaminase antibody assay. Archives of Disease in Childhood, 2004, 89, 512-515. | 1.0 | 185 |
| 159 | Anti-Transglutaminase Antibodies and Age. Clinical Chemistry, 2004, 50, 1856-1860. | 1.5 | 10 |
| 160 | Human tissue transglutaminase ELISA and an old study: a revision of the blood donor screening study for coeliac disease in the USA. Scandinavian Journal of Gastroenterology, 2004, 39, 195-197. | 0.6 | 9 |
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