# Amit Nahum

List of Publications by Year
in descending order



| 16 | 187 | 1478505 6 | 1125743 13 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 17 | 17 | 17 | 276 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | First Year of Israeli Newborn Screening for Severe Combined Immunodeficiencyâ€"Clinical Achievements and Insights. Frontiers in Immunology, 2017, 8, 1448. | 4.8 | 67 |
| 2 | Defining the biological responses of IL-6 by the study of a novel IL-6 receptor chain immunodeficiency. Journal of Allergy and Clinical Immunology, 2020, 145, 1011-1015.e6. | 2.9 | 25 |
| 3 | Inhaled nitric oxide therapy in acute bronchiolitis: A multicenter randomized clinical trial. Scientific Reports, 2020, 10, 9605. | 3.3 | 20 |
| 4 | Lessons Learned From Five Years of Newborn Screening for Severe Combined Immunodeficiency in Israel. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 2722-2731.e9. | 3.8 | 15 |
| 5 | The clinical and laboratory spectrum of dedicator of cytokinesis 8 immunodeficiency syndrome in patients with a unique mutation. Immunologic Research, 2017, 65, 651-657. | 2.9 | 12 |
| 6 | Chronic Mucocutaneous Candidiasis in Early Life: Insights Into Immune Mechanisms and Novel Targeted Therapies. Frontiers in Immunology, 2020, 11, 593289. | 4.8 | 11 |
| 7 | Cytokine profile of foodâ€allergic postâ€liver transplant children is identified by high levels of ILâ€5 and low ILâ€10 secretion from patients' peripheral blood mononuclear cells. Pediatric Transplantation, 2015, 19, 716-721. | 1.0 | 9 |
| 8 | Unusual phenotype in patients with a hypomorphic mutation in the DCLRE1C gene: IgG hypergammaglobulinemia with IgA and IgE deficiency. Clinical Immunology, 2020, 213, 108366. | 3.2 | 7 |
| 9 | Sesame: An unrecognized trigger of food protein-induced enterocolitis syndrome. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 305-306. | 3.8 | 6 |
| 10 | Toll-like receptor 3 (TLR3) variant and NLRP12 mutation confer susceptibility to a complex clinical presentation. Clinical Immunology, 2020, 212, 108249. | 3.2 | 6 |
| 11 | Phenotypic variability in patients with unique double homozygous mutations causing variant ataxia telangiectasia. European Journal of Paediatric Neurology, 2021, 32, 36-39. | 1.6 | 2 |
| 12 | Chronic mucocutaneous candidiasis: a spectrum of genetic disorders. LymphoSign Journal, 0, , . | 0.2 | 2 |
| 13 | B cell repertoire in patients with a novel BTK mutation: expanding the spectrum of atypical X-linked agammaglobulinemia. Immunologic Research, 2022, 70, 216-223. | 2.9 | 2 |
| 14 | Hodgkin's lymphoma, nephrotic syndrome, and echinococcosis cysts: an unusual association and literature review. Pediatric Hematology and Oncology, 2019, 36, 40-45. | 0.8 | 1 |
| 15 | Fast fluorometric method for measuring circulating cellâ€free DNA could aid the diagnosis of febrile children. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 1577-1578. | 1.5 | 1 |
| 16 | Trough Concentrations of Specific Antibodies in Primary Immunodeficiency Patients Receiving Intravenous Immunoglobulin Replacement Therapy. Journal of Clinical Medicine, 2021, 10, 592. | 2.4 | 1 |