Meri Kaustio

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4094342/publications.pdf

Version: 2024-02-01

| | | 1163117 | 1372567 | |
|----------|----------------|--------------|----------------|--|
| 14 | 524 | 8 | 10 | |
| papers | citations | h-index | g-index | |
| | | | | |
| | | | | |
| 15 | 15 | 15 | 1324 | |
| all docs | docs citations | times ranked | citing authors | |
| | | | | |

| # | Article | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Autoimmunity, hypogammaglobulinemia, lymphoproliferation, and mycobacterial disease in patients with activating mutations in STAT3. Blood, 2015, 125, 639-648. | 1.4 | 229 |
| 2 | Damaging heterozygous mutations in NFKB1 lead to diverse immunologic phenotypes. Journal of Allergy and Clinical Immunology, 2017, 140, 782-796. | 2.9 | 113 |
| 3 | ADA2 deficiency: Clonal lymphoproliferation in a subset of patients. Journal of Allergy and Clinical Immunology, 2018, 141, 1534-1537.e8. | 2.9 | 71 |
| 4 | Loss-of-function mutation in <i>IKZF2</i> leads to immunodeficiency with dysregulated germinal center reactions and reduction of MAIT cells. Science Immunology, 2021, 6, eabe3454. | 11.9 | 30 |
| 5 | Loss of DIAPH1 causes SCBMS, combined immunodeficiency, and mitochondrial dysfunction. Journal of Allergy and Clinical Immunology, 2021, 148, 599-611. | 2.9 | 23 |
| 6 | Enrichment of rare variants in population isolates: single AICDA mutation responsible for hyper-IgM syndrome type 2 in Finland. European Journal of Human Genetics, 2016, 24, 1473-1478. | 2.8 | 22 |
| 7 | Constant B cell lymphocytosis since early age in a patient with CARD11 mutation: A 20-year follow-up. Clinical Immunology, 2016, 165, 19-20. | 3.2 | 17 |
| 8 | Identification of novel regulators of STAT3 activity. PLoS ONE, 2020, 15, e0230819. | 2.5 | 12 |
| 9 | Primary Immunodeficiency, a Possible Cause of Neutrophilic Necrotizing Dermatosis. JAMA Dermatology, 2019, 155, 863. | 4.1 | 5 |
| 10 | Dominant NFKB1 Mutations Cause Antibody Deficiency and Autoinflammatory Episodes. Blood, 2015, 126, 206-206. | 1.4 | 1 |
| 11 | Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819. | | O |
| 12 | Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819. | | 0 |
| 13 | Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819. | | 0 |
| 14 | Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819. | | 0 |