

Pediatric-onset Evans syndrome: Heterogeneous presentation of monogenic disorders including LRBA and CTLA4 mutations

Clinical Immunology

188, 52-57

DOI: [10.1016/j.clim.2017.12.009](https://doi.org/10.1016/j.clim.2017.12.009)

Citation Report

#	ARTICLE	IF	CITATIONS
1	TPP2 mutation associated with sterile brain inflammation mimicking MS. <i>Neurology: Genetics</i> , 2018, 4, e285.	0.9	6
2	Tregopathies: Monogenic diseases resulting in regulatory T-cell deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 1679-1695.	1.5	106
3	Evans syndrome: clinical perspectives, biological insights and treatment modalities. <i>Journal of Blood Medicine</i> , 2018, Volume 9, 171-184.	0.7	54
4	Novel LRBA Mutation and Possible Germinal Mosaicism in a Slavic Family. <i>Journal of Clinical Immunology</i> , 2018, 38, 471-474.	2.0	5
5	Bacille Calmette-Guérin Complications in Newly Described Primary Immunodeficiency Diseases: 2010-2017. <i>Frontiers in Immunology</i> , 2018, 9, 1423.	2.2	20
6	Monogenic polyautoimmunity in primary immunodeficiency diseases. <i>Autoimmunity Reviews</i> , 2018, 17, 1028-1039.	2.5	24
7	Evans Syndrome in Childhood: Long Term Follow-Up and the Evolution in Primary Immunodeficiency or Rheumatological Disease. <i>Frontiers in Pediatrics</i> , 2019, 7, 304.	0.9	19
8	Human DEF6 deficiency underlies an immunodeficiency syndrome with systemic autoimmunity and aberrant CTLA-4 homeostasis. <i>Nature Communications</i> , 2019, 10, 3106.	5.8	48
9	Clinical Aspects of STAT3 Gain-of-Function Germline Mutations: A Systematic Review. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 1958-1969.e9.	2.0	144
10	Clinical, Immunologic, and Molecular Spectrum of Patients with LPS-Responsive Beige-Like Anchor Protein Deficiency: A Systematic Review. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 2379-2386.e5.	2.0	88
11	Neurological Involvement in Childhood Evans Syndrome. <i>Journal of Clinical Immunology</i> , 2019, 39, 171-181.	2.0	6
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13	Paradoxical CD4 Lymphopenia in Autoimmune Lymphoproliferative Syndrome (ALPS). <i>Frontiers in Immunology</i> , 2019, 10, 1193.	2.2	18
14	Pediatric Evans syndrome is associated with a high frequency of potentially damaging variants in immune genes. <i>Blood</i> , 2019, 134, 9-21.	0.6	102
15	Papilledema from gain-of-function mutations in the <i>STAT3</i> gene. <i>Ophthalmic Genetics</i> , 2019, 40, 165-169.	0.5	4
16	Autoimmunity as a continuum in primary immunodeficiency. <i>Current Opinion in Pediatrics</i> , 2019, 31, 851-862.	1.0	46
17	Bortezomib treatment of steroid-refractory Evans syndrome in children. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28725.	0.8	3
18	Activating mutations of STAT3: Impact on human growth. <i>Molecular and Cellular Endocrinology</i> , 2020, 518, 110979.	1.6	14

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19	Evansâ€™ Syndrome: From Diagnosis to Treatment. <i>Journal of Clinical Medicine</i> , 2020, 9, 3851.	1.0	50
20	How to evaluate for immunodeficiency in patients with autoimmune cytopenias: laboratory evaluation for the diagnosis of inborn errors of immunity associated with immune dysregulation. <i>Hematology American Society of Hematology Education Program</i> , 2020, 2020, 661-672.	0.9	10
21	Single Nucleotide Polymorphisms in PPAR α Associated with Systemic Lupus Erythematosus in Chinese Populations. <i>Journal of Immunology Research</i> , 2020, 2020, 1-7.	0.9	1
22	Secondâ€line treatment trends and longâ€term outcomes of 392 children with chronic immune thrombocytopenic purpura: the French experience over the past 25 years. <i>British Journal of Haematology</i> , 2020, 189, 931-942.	1.2	12
23	Pediatric immune thrombocytopenia (ITP) treatment. <i>Annals of Blood</i> , 0, 6, 4-4.	0.4	2
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25	Refractory autoimmune cytopenias in pediatric Evans syndrome with underlying systemic immune dysregulation. <i>European Journal of Haematology</i> , 2021, 106, 783-787.	1.1	9
26	Primary Immunodeficiency in Children With Autoimmune Cytopenias: Retrospective 154-Patient Cohort. <i>Frontiers in Immunology</i> , 2021, 12, 649182.	2.2	12
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28	Potential proteinâ€phenotype correlation in three lipopolysaccharide-responsive beige-like anchor protein-deficient patients. <i>World Journal of Clinical Cases</i> , 2021, 9, 5873-5888.	0.3	0
29	Primary Immune Regulatory Disorders With an Autoimmune Lymphoproliferative Syndrome-Like Phenotype: Immunologic Evaluation, Early Diagnosis and Management. <i>Frontiers in Immunology</i> , 2021, 12, 671755.	2.2	35
30	Response to rituximab in children and adults with immune thrombocytopenia (ITP). <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12587.	1.0	4
31	Long term follow-up of pediatric-onset Evans syndrome: broad immunopathological manifestations and high treatment burden. <i>Haematologica</i> , 2022, 107, 457-466.	1.7	9
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37	Autoimmune Cytopenias and Dysregulated Immunophenotype Act as Warning Signs of Inborn Errors of Immunity: Results From a Prospective Study. <i>Frontiers in Immunology</i> , 2021, 12, 790455.	2.2	11
38	Adult Evans' Syndrome. <i>Hematology/Oncology Clinics of North America</i> , 2022, 36, 381-392.	0.9	6

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41	Paraneoplastic Evans Syndrome in a Patient With Prostate Cancer With Small Cell Transformation. Cureus, 2022, , .	0.2	2
42	Underlying Inborn Errors of Immunity in Patients With Evans Syndrome and Multilineage Cytopenias: A Single-Centre Analysis. Frontiers in Immunology, 2022, 13, .	2.2	7
43	Autoimmune Cytopenias in Common Variable Immunodeficiency Are a Diagnostic and Therapeutic Conundrum: An Update. Frontiers in Immunology, 0, 13, .	2.2	7
44	Genetic Diagnosis Guides Treatment of Autoimmune Enteropathy. Clinical Gastroenterology and Hepatology, 2023, 21, 1368-1371.e2.	2.4	2
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