

# Michael Hershfield

## List of Publications by Year in descending order

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Version: 2024-02-01

32  
papers

1,100  
citations

430874

18  
h-index

454955

30  
g-index

33  
all docs

33  
docs citations

33  
times ranked

1247  
citing authors

#	ARTICLE	IF	CITATIONS
1	Hematopoietic stem cell transplantation rescues the hematological, immunological, and vascular phenotype in DADA2. <i>Blood</i> , 2017, 130, 2682-2688.	1.4	140
2	Screening of 181 Patients With Antibody Deficiency for Deficiency of Adenosine Deaminase 2 Sheds New Light on the Disease in Adulthood. <i>Arthritis and Rheumatology</i> , 2017, 69, 1689-1700.	5.6	103
3	Mitochondrial Basis for Immune Deficiency. <i>Journal of Experimental Medicine</i> , 2000, 191, 2197-2208.	8.5	100
4	Deficiency of Adenosine Deaminase 2 Causes Antibody Deficiency. <i>Journal of Clinical Immunology</i> , 2016, 36, 179-186.	3.8	78
5	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
6	Tandem mass spectrometry, but not T-cell receptor excision circle analysis, identifies newborns with late-onset adenosine deaminase deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 131, 1604-1610.	2.9	65
7	Autoimmune phenotype with type I interferon signature in two brothers with ADA2 deficiency carrying a novel CECR1 mutation. <i>Pediatric Rheumatology</i> , 2017, 15, 67.	2.1	58
8	Diagnosis of immunodeficiency caused by a purine nucleoside phosphorylase defect by using tandem mass spectrometry on dried blood spots. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 155-159.e3.	2.9	56
9	Human adenosine deaminase 2 deficiency: A multifaceted inborn error of immunity. <i>Immunological Reviews</i> , 2019, 287, 62-72.	6.0	54
10	Hematopoietic Cell Transplantation Cures Adenosine Deaminase 2 Deficiency: Report on 30 Patients. <i>Journal of Clinical Immunology</i> , 2021, 41, 1633-1647.	3.8	43
11	Deficiency of Adenosine Deaminase 2 (DADA2): Hidden Variants, Reduced Penetrance, and Unusual Inheritance. <i>Journal of Clinical Immunology</i> , 2020, 40, 917-926.	3.8	32
12	Clinical, Immunological, and Molecular Features of Severe Combined Immune Deficiency: A Multi-Institutional Experience From India. <i>Frontiers in Immunology</i> , 2020, 11, 619146.	4.8	31
13	Long-term outcomes after gene therapy for adenosine deaminase severe combined immune deficiency. <i>Blood</i> , 2021, 138, 1304-1316.	1.4	28
14	ADA Deficiency: Evaluation of the Clinical and Laboratory Features and the Outcome. <i>Journal of Clinical Immunology</i> , 2018, 38, 484-493.	3.8	26
15	ADA2 Deficiency Mimicking Idiopathic Multicentric Castleman Disease. <i>Pediatrics</i> , 2018, 142, .	2.1	26
16	Outcomes following treatment for ADA-deficient severe combined immunodeficiency: a report from the PIDTC. <i>Blood</i> , 2022, 140, 685-705.	1.4	26
17	Renal Amyloidosis in Deficiency of Adenosine Deaminase 2: Successful Experience With Canakinumab. <i>Pediatrics</i> , 2018, 142, .	2.1	23
18	Intrinsic Defects in B Cell Development and Differentiation, T Cell Exhaustion and Altered Unconventional T Cell Generation Characterize Human Adenosine Deaminase Type 2 Deficiency. <i>Journal of Clinical Immunology</i> , 2021, 41, 1915-1935.	3.8	23

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19	Novel compound heterozygous variants in <i>CECR1</i> gene associated with childhood onset polyarteritis nodosa and deficiency of ADA2. <i>Rheumatology</i> , 2016, 55, 1145-1147.	1.9	22
20	Combined immunodeficiencies: twenty years experience from a single center in Turkey. <i>Central-European Journal of Immunology</i> , 2016, 1, 107-115.	1.2	20
21	Childhood Hodgkin Lymphoma: Think DADA2. <i>Journal of Clinical Immunology</i> , 2019, 39, 26-29.	3.8	20
22	A Case with Purine Nucleoside Phosphorylase Deficiency Suffering from Late-Onset Systemic Lupus Erythematosus and Lymphoma. <i>Journal of Clinical Immunology</i> , 2020, 40, 833-839.	3.8	16
23	Complexities of genetic diagnosis illustrated by an atypical case of congenital hypoplastic anemia. <i>Journal of Physical Education and Sports Management</i> , 2018, 4, a003384.	1.2	12
24	Adenosine Deaminase 2 Deficiency As a Cause of Pure Red Cell Aplasia Mimicking Diamond Blackfan Anemia. <i>Blood</i> , 2015, 126, 3615-3615.	1.4	9
25	Adenosine Deaminase-Deficient Severe Combined Immunodeficiency and Diffuse Large B-Cell Lymphoma. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2015, 28, 138-142.	0.8	4
26	Deficiency of adenosine deaminase 2: a case series revealing clinical manifestations, genotypes and treatment outcomes from Turkey. <i>Rheumatology</i> , 2020, 59, 254-256.	1.9	4
27	Diamond-Blackfan Anemia Phenotype Caused By Deficiency of Adenosine Deaminase 2. <i>Blood</i> , 2017, 130, 874-874.	1.4	4
28	Two patients with novel missense mutation in the purine nucleoside phosphorylase gene without serious or recurrent infections. <i>Clinical and Experimental Neuroimmunology</i> , 2016, 7, 79-82.	1.0	3
29	Comparative Results of Gene Therapy for Adenosine Deaminase Deficiency with or without PEG-ADA Withdrawal and Myelosuppressive Chemotherapy. <i>Blood</i> , 2007, 110, 501-501.	1.4	2
30	Deficiency of Adenosine Deaminase 2 (DADA2) Presenting As Familial Hodgkin Lymphoma. <i>Blood</i> , 2018, 132, 5373-5373.	1.4	1
31	Normal IgH Repertoire Diversity in an Infant with ADA Deficiency After Gene Therapy. <i>Journal of Clinical Immunology</i> , 2021, 41, 1597-1606.	3.8	0
32	Deficiency of Adenosine Deaminase 2 a Monogenic Cause of Wunderlich Syndrome. <i>Journal of Clinical Immunology</i> , 2021, 41, 1693-1695.	3.8	0