

Bodo Grimbacher

List of Publications by Year in descending order

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Version: 2024-02-01

192
papers

24,090
citations

9264

74
h-index

7745

150
g-index

200
all docs

200
docs citations

200
times ranked

18218
citing authors

#	ARTICLE	IF	CITATIONS
1	Therapeutic options for CTLA-4 insufficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 149, 736-746.	2.9	68
2	Pembrolizumab for treatment of progressive multifocal leukoencephalopathy in primary immunodeficiency and/or hematologic malignancy: a case series of five patients. <i>Journal of Neurology</i> , 2022, 269, 973-981.	3.6	9
3	Therapeutic targeting of endoplasmic reticulum stress in acute graft-&i>versus&/i>-host disease. <i>Haematologica</i> , 2022, 107, 1538-1554.	3.5	3
4	Bowel Histology of CVID Patients Reveals Distinct Patterns of Mucosal Inflammation. <i>Journal of Clinical Immunology</i> , 2022, 42, 46-59.	3.8	10
5	Diagnostic Yield and Therapeutic Consequences of Targeted Next-Generation Sequencing in Sporadic Primary Immunodeficiency. <i>International Archives of Allergy and Immunology</i> , 2022, 183, 337-349.	2.1	6
6	Dysregulated PI3K Signaling in B Cells of CVID Patients. <i>Cells</i> , 2022, 11, 464.	4.1	6
7	Predictive Factors for and Complications of Bronchiectasis in Common Variable Immunodeficiency Disorders. <i>Journal of Clinical Immunology</i> , 2022, 42, 572-581.	3.8	5
8	Resolving the polygenic aetiology of a late onset combined immune deficiency caused by NFKB1 haploinsufficiency and modified by PIK3R1 and TNFRSF13B variants. <i>Clinical Immunology</i> , 2022, 234, 108910.	3.2	3
9	Treatment of STAT3-deficient hyperâimmunoglobulin E syndrome with monoclonal antibodies targeting allergic inflammation. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2022, 10, 1367-1370.e1.	3.8	8
10	Phenotype, genotype, treatment, and survival outcomes in patients with X-linked inhibitor of apoptosis deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 150, 456-466.	2.9	15
11	Copy Number Analysis in a Large Cohort Suggestive of Inborn Errors of Immunity Indicates a Wide Spectrum of Relevant Chromosomal Losses and Gains. <i>Journal of Clinical Immunology</i> , 2022, 42, 1083-1092.	3.8	6
12	Genomic characterization of lymphomas in patients with inborn errors of immunity. <i>Blood Advances</i> , 2022, 6, 5403-5414.	5.2	12
13	Bile acids regulate intestinal antigen presentation and reduce graft-versus-host disease without impairing the graft-versus-leukemia effect. <i>Haematologica</i> , 2021, 106, 2131-2146.	3.5	26
14	Hematopoietic Stem Cell Transplantation Resolves the Immune Deficit Associated with STAT3-Dominant-Negative Hyper-IgE Syndrome. <i>Journal of Clinical Immunology</i> , 2021, 41, 934-943.	3.8	21
15	Gain-of-function variants in SYK cause immune dysregulation and systemic inflammation in humans and mice. <i>Nature Genetics</i> , 2021, 53, 500-510.	21.4	56
16	A Pathogenic Missense Variant in NFKB1 Causes Common Variable Immunodeficiency Due to Detrimental Protein Damage. <i>Frontiers in Immunology</i> , 2021, 12, 621503.	4.8	12
17	Altered Spectrum of Lymphoid Neoplasms in a Single-Center Cohort of Common Variable Immunodeficiency with Immune Dysregulation. <i>Journal of Clinical Immunology</i> , 2021, 41, 1250-1265.	3.8	15
18	Immune checkpoint deficiencies and autoimmune lymphoproliferative syndromes. <i>Biomedical Journal</i> , 2021, 44, 400-411.	3.1	23

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19	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 1332-1341.e5.	2.9	75
20	Activation-induced deaminase is critical for the establishment of DNA methylation patterns prior to the germinal center reaction. <i>Nucleic Acids Research</i> , 2021, 49, 5057-5073.	14.5	5
21	What can clinical immunology learn from inborn errors of epigenetic regulators?. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 1602-1618.	2.9	8
22	Dysregulated immunity in PID patients with low GARP expression on Tregs due to mutations in LRRC32. <i>Cellular and Molecular Immunology</i> , 2021, 18, 1677-1691.	10.5	11
23	Genetic Analysis of a Cohort of 275 Patients with Hyper-IgE Syndromes and/or Chronic Mucocutaneous Candidiasis. <i>Journal of Clinical Immunology</i> , 2021, 41, 1804-1838.	3.8	12
24	TAC1 deficiency "a complex system out of balance. <i>Current Opinion in Immunology</i> , 2021, 71, 81-88.	5.5	21
25	Biochemically deleterious human <i>NFKB1</i> variants underlie an autosomal dominant form of common variable immunodeficiency. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	32
26	Simple Measurement of IgA Predicts Immunity and Mortality in Ataxia-Telangiectasia. <i>Journal of Clinical Immunology</i> , 2021, 41, 1878-1892.	3.8	9
27	BTK operates a phospho-tyrosine switch to regulate NLRP3 inflammasome activity. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	33
28	There is no gene for CVID "novel monogenetic causes for primary antibody deficiency. <i>Current Opinion in Immunology</i> , 2021, 72, 176-185.	5.5	26
29	A distinct CD38+CD45RA+ population of CD4+, CD8+, and double-negative T cells is controlled by FAS. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	25
30	The expansion of human T-bet ^{high} CD21 ^{low} B cells is T cell dependent. <i>Science Immunology</i> , 2021, 6, eabh0891.	11.9	82
31	Establishing the Molecular Diagnoses in a Cohort of 291 Patients With Predominantly Antibody Deficiency by Targeted Next-Generation Sequencing: Experience From a Monocentric Study. <i>Frontiers in Immunology</i> , 2021, 12, 786516.	4.8	19
32	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1452-1463.	2.9	112
33	Distinct molecular response patterns of activating STAT3 mutations associate with penetrance of lymphoproliferation and autoimmunity. <i>Clinical Immunology</i> , 2020, 210, 108316.	3.2	40
34	Structural Noninfectious Manifestations of the Central Nervous System in Common Variable Immunodeficiency Disorders. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020, 8, 1047-1062.e6.	3.8	10
35	Altered Microbiota, Impaired Quality of Life, Malabsorption, Infection, and Inflammation in CVID Patients With Diarrhoea. <i>Frontiers in Immunology</i> , 2020, 11, 1654.	4.8	17
36	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	64

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37	Nonpermissive bone marrow environment impairs early B-cell development in common variable immunodeficiency. <i>Blood</i> , 2020, 135, 1452-1457.	1.4	7
38	Incidence of SCID in Germany from 2014 to 2015 an ESPED* Survey on Behalf of the API*** Erhebungseinheit für Seltene Pädiatrische Erkrankungen in Deutschland (German Paediatric) <i>Trends in Pediatrics</i> 2020, 40, 708-717.	3.8	9
39	Glucagon-like peptide 2 for intestinal stem cell and Paneth cell repair during graft-versus-host disease in mice and humans. <i>Blood</i> , 2020, 136, 1442-1455.	1.4	60
40	Clinical Phenotypes and Immunological Characteristics of 18 Egyptian LRBA Deficiency Patients. <i>Journal of Clinical Immunology</i> , 2020, 40, 820-832.	3.8	17
41	Safety and efficacy of abatacept in patients with treatment-resistant Sarcoidosis (ABASARC) – protocol for a multi-center, single-arm phase IIa trial. <i>Contemporary Clinical Trials Communications</i> , 2020, 19, 100575.	1.1	10
42	Dynamics in protein translation sustaining T cell preparedness. <i>Nature Immunology</i> , 2020, 21, 927-937.	14.5	120
43	Cancer Tendency in a Patient with ZNF341 Deficiency. <i>Journal of Clinical Immunology</i> , 2020, 40, 534-538.	3.8	3
44	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 901-911.	2.9	78
45	Enabling External Inquiries to an Existing Patient Registry by Using the Open Source Registry System for Rare Diseases: Demonstration of the System Using the European Society for Immunodeficiencies Registry. <i>JMIR Medical Informatics</i> , 2020, 8, e17420.	2.6	2
46	The German National Registry of Primary Immunodeficiencies (2012–2017). <i>Frontiers in Immunology</i> , 2019, 10, 1272.	4.8	71
47	The architecture of the IgG anti-carbohydrate repertoire in primary antibody deficiencies. <i>Blood</i> , 2019, 134, 1941-1950.	1.4	19
48	FAS mutations are an uncommon cause of immune thrombocytopenia in children and adults without additional features of immunodeficiency. <i>British Journal of Haematology</i> , 2019, 186, e163-e165.	2.5	6
49	Signaling mechanisms inducing hyporesponsiveness of phagocytes during systemic inflammation. <i>Blood</i> , 2019, 134, 134-146.	1.4	39
50	Evaluating laboratory criteria for combined immunodeficiency in adult patients diagnosed with common variable immunodeficiency. <i>Clinical Immunology</i> , 2019, 203, 59-62.	3.2	36
51	Assessing the Functional Relevance of Variants in the IKAROS Family Zinc Finger Protein 1 (IKZF1) in a Cohort of Patients With Primary Immunodeficiency. <i>Frontiers in Immunology</i> , 2019, 10, 568.	4.8	37
52	The European Society for Immunodeficiencies (ESID) Registry Working Definitions for the Clinical Diagnosis of Inborn Errors of Immunity. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 1763-1770.	3.8	381
53	Clinical and Immunological Phenotype of Patients With Primary Immunodeficiency Due to Damaging Mutations in NFKB2. <i>Frontiers in Immunology</i> , 2019, 10, 297.	4.8	117
54	Late-Onset Antibody Deficiency Due to Monoallelic Alterations in NFKB1. <i>Frontiers in Immunology</i> , 2019, 10, 2618.	4.8	29

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55	What did we learn from <sc>CTLA</sc> insufficiency on the human immune system?. Immunological Reviews, 2019, 287, 33-49.	6.0	121
56	Imaging of Bronchial Pathology in Antibody Deficiency: Data from the European Chest CT Group. Journal of Clinical Immunology, 2019, 39, 45-54.	3.8	32
57	A novel monoallelic gain of function mutation in p110 $\hat{\nu}$ causing atypical activated phosphoinositide 3-kinase $\hat{\nu}$ syndrome (APDS-1). Clinical Immunology, 2019, 200, 31-34.	3.2	11
58	Hypomorphic caspase activation and recruitment domain 11 (CARD11) mutations associated with diverse immunologic phenotypes with or without atopic disease. Journal of Allergy and Clinical Immunology, 2019, 143, 1482-1495.	2.9	116
59	Autoimmunity and primary immunodeficiency: two sides of the same coin?. Nature Reviews Rheumatology, 2018, 14, 7-18.	8.0	103
60	Treatment of severe forms of LPS-responsive beige-like anchor protein deficiency with allogeneic hematopoietic stem cell transplantation. Journal of Allergy and Clinical Immunology, 2018, 141, 770-775.e1.	2.9	52
61	The TH1 phenotype of follicular helper T cells indicates an IFN- $\hat{\nu}$ associated immune dysregulation in patients with CD21 ^{low} common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 730-740.	2.9	109
62	Respiratory Infections and Antibiotic Usage in Common Variable Immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 159-168.e3.	3.8	46
63	Plasma cell deficiency in human subjects with heterozygous mutations in Sec61 translocon alpha 1 subunit (SEC61A1). Journal of Allergy and Clinical Immunology, 2018, 141, 1427-1438.	2.9	63
64	The burden of common variable immunodeficiency disorders: a retrospective analysis of the European Society for Immunodeficiency (ESID) registry data. Orphanet Journal of Rare Diseases, 2018, 13, 201.	2.7	119
65	A novel LPS-responsive beige-like anchor protein (LRBA) mutation presents with normal cytotoxic T lymphocyte-associated protein 4 (CTLA-4) and overactive TH17 immunity. Journal of Allergy and Clinical Immunology, 2018, 142, 1968-1971.	2.9	13
66	Increased Risk for Malignancies in 131 Affected CTLA4 Mutation Carriers. Frontiers in Immunology, 2018, 9, 2012.	4.8	79
67	Nuclear factor $\hat{\nu}$ B mutations in human subjects: The devil is in the details. Journal of Allergy and Clinical Immunology, 2018, 142, 1062-1065.	2.9	22
68	Novel LRBA Mutation and Possible Germinal Mosaicism in a Slavic Family. Journal of Clinical Immunology, 2018, 38, 471-474.	3.8	5
69	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4 ^{insufficient} subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	2.9	344
70	CTLA-4 regulates human Natural Killer cell effector functions. Clinical Immunology, 2018, 194, 43-45.	3.2	30
71	Diagnosis of DOCK8 deficiency using Flow cytometry Biomarkers: an Egyptian Center experience. Clinical Immunology, 2018, 195, 36-44.	3.2	9
72	Disease Evolution and Response to Rapamycin in Activated Phosphoinositide 3-Kinase $\hat{\nu}$ Syndrome: The European Society for Immunodeficiencies-Activated Phosphoinositide 3-Kinase $\hat{\nu}$ Syndrome Registry. Frontiers in Immunology, 2018, 9, 543.	4.8	137

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73	Rapid Flow Cytometry-Based Test for the Diagnosis of Lipopolysaccharide Responsive Beige-Like Anchor (LRBA) Deficiency. <i>Frontiers in Immunology</i> , 2018, 9, 720.	4.8	24
74	Immune TORopathies, a Novel Disease Entity in Clinical Immunology. <i>Frontiers in Immunology</i> , 2018, 9, 966.	4.8	35
75	Is It Safe to Switch From Intravenous Immunoglobulin to Subcutaneous Immunoglobulin in Patients With Common Variable Immunodeficiency and Autoimmune Thrombocytopenia?. <i>Frontiers in Immunology</i> , 2018, 9, 1656.	4.8	12
76	A recessive form of hyper-IgE syndrome by disruption of ZNF341-dependent STAT3 transcription and activity. <i>Science Immunology</i> , 2018, 3, .	11.9	132
77	ZNF341 controls STAT3 expression and thereby immunocompetence. <i>Science Immunology</i> , 2018, 3, .	11.9	113
78	Disturbed canonical nuclear factor of κ B light chain signaling in B cells of patients with common variable immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 220-231.e8.	2.9	39
79	Evaluation of RAG1 mutations in an adult with combined immunodeficiency and progressive multifocal leukoencephalopathy. <i>Clinical Immunology</i> , 2017, 179, 1-7.	3.2	24
80	Human NACHT, LRR, and PYD domain-containing protein 3 (NLRP3) inflammasome activity is regulated by and potentially targetable through Bruton tyrosine kinase. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1054-1067.e10.	2.9	105
81	Rescue of Cytokine Storm Due to HLH by Hemoadsorption in a CTLA4-Deficient Patient. <i>Journal of Clinical Immunology</i> , 2017, 37, 273-276.	3.8	45
82	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
83	NFKB1 regulates human NK cell maturation and effector functions. <i>Clinical Immunology</i> , 2017, 175, 99-108.	3.2	38
84	Targeted Gene Panel Sequencing for Early-onset Inflammatory Bowel Disease and Chronic Diarrhea. <i>Inflammatory Bowel Diseases</i> , 2017, 23, 2109-2120.	1.9	33
85	A biallelic mutation in <i>IL6ST</i> encoding the GP130 co-receptor causes immunodeficiency and craniosynostosis. <i>Journal of Experimental Medicine</i> , 2017, 214, 2547-2562.	8.5	158
86	Immunological phenotype of the murine <i>Lrb</i> knockout. <i>Immunology and Cell Biology</i> , 2017, 95, 789-802.	2.3	24
87	Autoantibodies against BAFF, APRIL or IL21 - an alternative pathogenesis for antibody-deficiencies?. <i>BMC Immunology</i> , 2017, 18, 34.	2.2	10
88	Clinical spectrum and features of activated phosphoinositide 3-kinase γ syndrome: A large patient cohort study. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 597-606.e4.	2.9	377
89	Vedolizumab as a successful treatment of CTLA-4-associated autoimmune enterocolitis. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 1043-1046.e5.	2.9	24
90	Treatment of Infantile Inflammatory Bowel Disease and Autoimmunity by Allogeneic Stem Cell Transplantation in LPS-Responsive Beige-Like Anchor Deficiency. <i>Frontiers in Immunology</i> , 2017, 8, 52.	4.8	24

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91	14 Years after Discovery: Clinical Follow-up on 15 Patients with Inducible Co-Stimulator Deficiency. <i>Frontiers in Immunology</i> , 2017, 8, 964.	4.8	57
92	Other Well-Defined Immunodeficiencies. , 2017, , 461-517.		1
93	Unique and shared signaling pathways cooperate to regulate the differentiation of human CD4+ T cells into distinct effector subsets. <i>Journal of Experimental Medicine</i> , 2016, 213, 1589-1608.	8.5	77
94	Successful Granulocyte Colony-stimulating Factor Treatment of Relapsing <i>Candida albicans</i> Meningoencephalitis Caused by CARD9 Deficiency. <i>Pediatric Infectious Disease Journal</i> , 2016, 35, 428-431.	2.0	70
95	<i>DOCK8</i> deficiency in six Iranian patients. <i>Clinical Case Reports (discontinued)</i> , 2016, 4, 593-600.	0.5	7
96	Common variable immunodeficiency, impaired neurological development and reduced numbers of T regulatory cells in a 10-year-old boy with a STAT1 gain-of-function mutation. <i>Gene</i> , 2016, 586, 234-238.	2.2	31
97	Heterozygous STAT1 gain-of-function mutations underlie an unexpectedly broad clinical phenotype. <i>Blood</i> , 2016, 127, 3154-3164.	1.4	465
98	Activating PI3K γ mutations in a cohort of 669 patients with primary immunodeficiency. <i>Clinical and Experimental Immunology</i> , 2016, 183, 221-229.	2.6	82
99	Health-Related Quality of Life and Health Resource Utilization in Patients with Primary Immunodeficiency Disease Prior to and Following 12 Months of Immunoglobulin G Treatment. <i>Journal of Clinical Immunology</i> , 2016, 36, 450-461.	3.8	24
100	The extended phenotype of LPS-responsive beige-like anchor protein (LRBA) deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 223-230.	2.9	247
101	Ruxolitinib Induces Interleukin 17 and Ameliorates Chronic Mucocutaneous Candidiasis Caused by STAT1 Gain-of-Function Mutation. <i>Clinical Infectious Diseases</i> , 2016, 62, 951.2-953.	5.8	73
102	Deficiency of Adenosine Deaminase 2 Causes Antibody Deficiency. <i>Journal of Clinical Immunology</i> , 2016, 36, 179-186.	3.8	78
103	Infancy-Onset T1DM, Short Stature, and Severe Immunodysregulation in Two Siblings With a Homozygous LRBA Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 898-904.	3.6	43
104	The crossroads of autoimmunity and immunodeficiency: Lessons from polygenic traits and monogenic defects. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 3-17.	2.9	100
105	The Extended Clinical Phenotype of 26 Patients with Chronic Mucocutaneous Candidiasis due to Gain-of-Function Mutations in STAT1. <i>Journal of Clinical Immunology</i> , 2016, 36, 73-84.	3.8	124
106	Defects in B Cell Survival and Activation. , 2016, , 466-478.		2
107	Atypical Manifestation of LRBA Deficiency with Predominant IBD-like Phenotype. <i>Inflammatory Bowel Diseases</i> , 2015, 21, 40-47.	1.9	81
108	Pathogenic Fungi Regulate Immunity by Inducing Neutrophilic Myeloid-Derived Suppressor Cells. <i>Cell Host and Microbe</i> , 2015, 17, 507-514.	11.0	99

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109	The extended clinical phenotype of 64 patients with deficiency of cytokines 8. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 402-412.	2.9	163
110	DOCK8 Deficiency: Clinical and Immunological Phenotype and Treatment Options - a Review of 136 Patients. <i>Journal of Clinical Immunology</i> , 2015, 35, 189-198.	3.8	284
111	Multicenter experience in hematopoietic stem cell transplantation for serious complications of common variable immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 988-997.e6.	2.9	123
112	Long-term remission after allogeneic hematopoietic stem cell transplantation in LPS-responsive beige-like anchor (LRBA) deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 1384-1390.e8.	2.9	65
113	Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 993-1006.e1.	2.9	181
114	Lung disease in primary antibody deficiency. <i>Lancet Respiratory Medicine</i> , 2015, 3, 651-660.	10.7	92
115	Symptomatic Males and Female Carriers in a Large Caucasian Kindred with XIAP Deficiency. <i>Journal of Clinical Immunology</i> , 2015, 35, 439-444.	3.8	29
116	Autoimmune lymphoproliferative syndrome-like disease in patients with LRBA mutation. <i>Clinical Immunology</i> , 2015, 159, 84-92.	3.2	96
117	Association of CLEC16A with human common variable immunodeficiency disorder and role in murine B cells. <i>Nature Communications</i> , 2015, 6, 6804.	12.8	63
118	DCLRE1C (ARTEMIS) mutations causing phenotypes ranging from atypical severe combined immunodeficiency to mere antibody deficiency. <i>Human Molecular Genetics</i> , 2015, 24, 7361-7372.	2.9	72
119	Haploinsufficiency of the NF- κ B1 Subunit p50 in Common Variable Immunodeficiency. <i>American Journal of Human Genetics</i> , 2015, 97, 389-403.	6.2	232
120	An inherited immunoglobulin class-switch recombination deficiency associated with a defect in the INO80 chromatin remodeling complex. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 998-1007.e6.	2.9	37
121	The diagnosis of hyper immunoglobulin e syndrome based on project management. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2015, 14, 126-32.	0.4	2
122	Gain-of-function mutations in signal transducer and activator of transcription 1 (STAT1): Chronic mucocutaneous candidiasis accompanied by enamel defects and delayed dental shedding. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 1209-1213.e6.	2.9	41
123	Ten-Year Follow-Up of a DOCK8-Deficient Child With Features of Systemic Lupus Erythematosus. <i>Pediatrics</i> , 2014, 134, e1458-e1463.	2.1	29
124	The Many Faces of the Hyper-IgE Syndrome. , 2014, , 241-254.		1
125	Hyper-IgE syndromes. <i>Current Opinion in Pediatrics</i> , 2014, 26, 697-703.	2.0	40
126	The use of databases in primary immunodeficiencies. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2014, 14, 501-508.	2.3	21

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127	Common variable immunodeficiency is associated with a functional deficiency of invariant natural killer T cells. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1420-1428.e1.	2.9	19
128	Activity, Severity and Impact of Respiratory Disease in Primary Antibody Deficiency Syndromes. <i>Journal of Clinical Immunology</i> , 2014, 34, 68-75.	3.8	34
129	Clinical picture and treatment of 2212 patients with common variable immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 116-126.e11.	2.9	512
130	Autosomal dominant immune dysregulation syndrome in humans with CTLA4 mutations. <i>Nature Medicine</i> , 2014, 20, 1410-1416.	30.7	723
131	Reply. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 990.	2.9	0
132	Evidence for non-neutralizing autoantibodies against IL-10 signalling components in patients with inflammatory bowel disease. <i>BMC Immunology</i> , 2014, 15, 10.	2.2	9
133	Hypomorphic homozygous mutations in phosphoglucomutase 3 (PGM3) impair immunity and increase serum IgE levels. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1410-1419.e13.	2.9	160
134	CD161 expression characterizes a subpopulation of human regulatory T cells that produces IL-17 in a STAT3-dependent manner. <i>European Journal of Immunology</i> , 2013, 43, 2043-2054.	2.9	114
135	Deep Dermatophytosis and Inherited CARD9 Deficiency. <i>New England Journal of Medicine</i> , 2013, 369, 1704-1714.	27.0	362
136	Deficiency of caspase recruitment domain family, member 11 (CARD11), causes profound combined immunodeficiency in human subjects. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 131, 477-485.e1.	2.9	166
137	B-cell biology and development. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 131, 959-971.	2.9	376
138	Autosomal Dominant Cases of Chronic Mucocutaneous Candidiasis Segregates with Mutations of Signal Transducer and Activator of Transcription 1, But Not of Toll-Like Receptor 3. <i>Journal of Pediatrics</i> , 2013, 163, 277-279.	1.8	22
139	CVID-associated TACI mutations affect autoreactive B cell selection and activation. <i>Journal of Clinical Investigation</i> , 2013, 123, 4283-4293.	8.2	153
140	The phenotype of human STK4 deficiency. <i>Blood</i> , 2012, 119, 3450-3457.	1.4	286
141	Mendelian traits causing susceptibility to mucocutaneous fungal infections in human subjects. <i>Journal of Allergy and Clinical Immunology</i> , 2012, 129, 294-305.	2.9	74
142	An update on the hyper-IgE syndromes. <i>Arthritis Research and Therapy</i> , 2012, 14, 228.	3.5	126
143	Soluble BAFF Levels Inversely Correlate with Peripheral B Cell Numbers and the Expression of BAFF Receptors. <i>Journal of Immunology</i> , 2012, 188, 497-503.	0.8	155
144	Deleterious Mutations in LRBA Are Associated with a Syndrome of Immune Deficiency and Autoimmunity. <i>American Journal of Human Genetics</i> , 2012, 90, 986-1001.	6.2	452

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145	Clinical, immunologic and genetic profiles of DOCK8-deficient patients in Kuwait. <i>Clinical Immunology</i> , 2012, 143, 266-272.	3.2	60
146	Inflammatory bowel disease: is it a primary immunodeficiency?. <i>Cellular and Molecular Life Sciences</i> , 2012, 69, 41-48.	5.4	99
147	“A Rose is a Rose is a Rose,” but CVID is Not CVID. <i>Advances in Immunology</i> , 2011, 111, 47-107.	2.2	155
148	IL10 and IL10 receptor defects in humans. <i>Annals of the New York Academy of Sciences</i> , 2011, 1246, 102-107.	3.8	223
149	The Phenotype of Human STK4 Deficiency. <i>Blood</i> , 2011, 118, 692-692.	1.4	2
150	Chronic mucocutaneous candidiasis and congenital susceptibility to <i>Candida</i> . <i>Current Opinion in Allergy and Clinical Immunology</i> , 2010, 10, 542-550.	2.3	59
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