

# GATA2 deficiency: a protean disorder of hematopoiesis,

Blood

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Collaborating constitutive and somatic genetic events in myeloid malignancies: ASXL1 mutations in patients with germline GATA2 mutations. <i>Haematologica</i> , 2014, 99, 201-203.	1.7	39
2	Mendelian Genetics of Human Susceptibility to Fungal Infection. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2014, 4, a019638-a019638.	2.9	81
3	Inflammatory Monocytes Orchestrate Innate Antifungal Immunity in the Lung. <i>PLoS Pathogens</i> , 2014, 10, e1003940.	2.1	154
4	DOCK8 regulates lymphocyte shape integrity for skin antiviral immunity. <i>Journal of Experimental Medicine</i> , 2014, 211, 2549-2566.	4.2	150
5	Secondary pulmonary alveolar proteinosis in hematologic malignancies. <i>Hematology/ Oncology and Stem Cell Therapy</i> , 2014, 7, 127-135.	0.6	36
6	Host genetic factors in susceptibility to mycobacterial disease. <i>Clinical Medicine</i> , 2014, 14, s17-s21.	0.8	4
7	Infections Caused by Non-Tuberculous Mycobacteria in Recipients of Hematopoietic Stem Cell Transplantation. <i>Frontiers in Oncology</i> , 2014, 4, 311.	1.3	48
8	Cellular immune controls over Epstein-Barr virus infection: new lessons from the clinic and the laboratory. <i>Trends in Immunology</i> , 2014, 35, 159-169.	2.9	121
9	Generalized verrucosis in a patient with GATA2 deficiency. <i>British Journal of Dermatology</i> , 2014, 170, 1182-1186.	1.4	21
10	Mendelian susceptibility to mycobacterial disease: Genetic, immunological, and clinical features of inborn errors of IFN- $\gamma$ immunity. <i>Seminars in Immunology</i> , 2014, 26, 454-470.	2.7	582
11	GATA2 deficiency: flesh and blood. <i>Blood</i> , 2014, 123, 799-800.	0.6	8
12	Nonmyeloablative Allogeneic Hematopoietic Stem Cell Transplantation for GATA2 Deficiency. <i>Biology of Blood and Marrow Transplantation</i> , 2014, 20, 1940-1948.	2.0	84
14	<i>Mycobacterium simiae</i> Infection in Two Unrelated Patients with Different Forms of Inherited IFN- $\gamma$ R2 Deficiency. <i>Journal of Clinical Immunology</i> , 2014, 34, 904-909.	2.0	20
15	Inherited Predisposition to Acute Myeloid Leukemia. <i>Seminars in Hematology</i> , 2014, 51, 306-321.	1.8	85
16	A woman with warts, leg swelling, and deafness. <i>Journal of the American Academy of Dermatology</i> , 2014, 71, 577-580.	0.6	7
17	Inherited bone marrow failure syndromes in adolescents and young adults. <i>Annals of Medicine</i> , 2014, 46, 353-363.	1.5	53
18	Human Invasive Mycoses: Immunogenetics on the Rise. <i>Journal of Infectious Diseases</i> , 2015, 211, 1205-7.	1.9	6
19	Synergistic contribution of SMAD signaling blockade and high localized cell density in the differentiation of neuroectoderm from H9 cells. <i>Biochemical and Biophysical Research Communications</i> , 2014, 452, 895-900.	1.0	7

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20	A Single Oncogenic Enhancer Rearrangement Causes Concomitant EVI1 and GATA2 Deregulation in Leukemia. <i>Cell</i> , 2014, 157, 369-381.	13.5	571
21	Cavitary pulmonary disease in a patient treated with natalizumab. <i>Presse Medicale</i> , 2014, 43, 1009-1012.	0.8	7
22	Discovery of single-gene inborn errors of immunity by next generation sequencing. <i>Current Opinion in Immunology</i> , 2014, 30, 17-23.	2.4	83
23	Old variables, new value: a refined IPI for DLBCL. <i>Blood</i> , 2014, 123, 800-801.	0.6	4
24	Functional characterization of the human dendritic cell immunodeficiency associated with the IRF8K108E mutation. <i>Blood</i> , 2014, 124, 1894-1904.	0.6	65
25	Spectrum of myeloid neoplasms and immune deficiency associated with germline <i>GATA2</i> mutations. <i>Cancer Medicine</i> , 2015, 4, 490-499.	1.3	43
26	Primary immunodeficiencies and the control of Epstein-Barr virus infection. <i>Annals of the New York Academy of Sciences</i> , 2015, 1356, 22-44.	1.8	42
27	Genomic analysis of bone marrow failure and myelodysplastic syndromes reveals phenotypic and diagnostic complexity. <i>Haematologica</i> , 2015, 100, 42-48.	1.7	108
28	T-cell-restricted T-bet overexpression induces aberrant hematopoiesis of myeloid cells and impairs function of macrophages in the lung. <i>Blood</i> , 2015, 125, 370-382.	0.6	19
29	GATA2 deficiency-associated bone marrow disorder differs from idiopathic aplastic anemia. <i>Blood</i> , 2015, 125, 56-70.	0.6	131
30	Young woman with mild bone marrow dysplasia, GATA2 and ASXL1 mutation treated with allogeneic hematopoietic stem cell transplantation. <i>Leukemia Research Reports</i> , 2015, 4, 72-75.	0.2	10
31	Successful hematopoietic cell transplantation in a patient with X-linked agammaglobulinemia and acute myeloid leukemia. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1674-1676.	0.8	30
32	GATA family transcriptional factors: emerging suspects in hematologic disorders. <i>Experimental Hematology and Oncology</i> , 2015, 4, 28.	2.0	57
33	Severe combined immunodeficiency (SCID) presenting with neonatal aplastic anemia. <i>Pediatric Blood and Cancer</i> , 2015, 62, 2047-2049.	0.8	1
34	Lymphatic vessel development: fluid flow and valve-forming cells. <i>Journal of Clinical Investigation</i> , 2015, 125, 2924-2926.	3.9	17
35	Host susceptibility to non-tuberculous mycobacterial infections. <i>Lancet Infectious Diseases</i> , The, 2015, 15, 968-980.	4.6	195
37	Dexamethasone targeted directly to macrophages induces macrophage niches that promote erythroid expansion. <i>Haematologica</i> , 2015, 100, 178-187.	1.7	59
38	The Immunology of Epstein-Barr Virus-Induced Disease. <i>Annual Review of Immunology</i> , 2015, 33, 787-821.	9.5	416

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39	Genetic predisposition syndromes: When should they be considered in the work-up of MDS?. Best Practice and Research in Clinical Haematology, 2015, 28, 55-68.	0.7	52
40	GATA2 Germline Mutations Impair <i>GATA2</i> Transcription, Causing Haploinsufficiency: Functional Analysis of the p.Arg396Gln Mutation. Journal of Immunology, 2015, 194, 2190-2198.	0.4	29
41	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. Immunological Reviews, 2015, 264, 103-120.	2.8	180
42	Disseminated Mycobacterial Infection and Scabies Infestation. American Journal of Medicine, 2015, 128, e41-e42.	0.6	3
43	Transcriptional Control of NK Cells. Current Topics in Microbiology and Immunology, 2015, 395, 1-36.	0.7	23
44	Pathology of bone marrow failure syndromes. Diagnostic Histopathology, 2015, 21, 174-180.	0.2	0
45	Hematopoietic Signaling Mechanism Revealed from a Stem/Progenitor Cell Cistrome. Molecular Cell, 2015, 59, 62-74.	4.5	40
46	Pulmonary alveolar proteinosis: time to shift?. Expert Review of Respiratory Medicine, 2015, 9, 337-349.	1.0	22
47	Rheumatologic manifestations of the "MonoMAC" syndrome. a systematic review. Clinical Rheumatology, 2015, 34, 1643-1645.	1.0	5
48	Inherited and Somatic Defects in DDX41 in Myeloid Neoplasms. Cancer Cell, 2015, 27, 658-670.	7.7	341
49	Diffuse parenchymal lung disease as first clinical manifestation of GATA-2 deficiency in childhood. BMC Pulmonary Medicine, 2015, 15, 8.	0.8	20
50	Mononuclear phagocyte-mediated antifungal immunity: the role of chemotactic receptors and ligands. Cellular and Molecular Life Sciences, 2015, 72, 2157-2175.	2.4	14
52	Recent advances in understanding the pathophysiology of primary T cell immunodeficiencies. Trends in Molecular Medicine, 2015, 21, 408-416.	3.5	18
53	GATA2 deficiency. Current Opinion in Allergy and Clinical Immunology, 2015, 15, 104-109.	1.1	113
54	Anemia of Central Origin. Seminars in Hematology, 2015, 52, 321-338.	1.8	9
55	GATA2 deficiency in children and adults with severe pulmonary alveolar proteinosis and hematologic disorders. BMC Pulmonary Medicine, 2015, 15, 87.	0.8	63
56	Regulation of GATA-binding Protein 2 Levels via Ubiquitin-dependent Degradation by Fbw7. Journal of Biological Chemistry, 2015, 290, 10368-10381.	1.6	27
57	Haematopoietic and immune defects associated with <i>GATA2</i> mutation. British Journal of Haematology, 2015, 169, 173-187.	1.2	197

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58	Low-level GATA2 overexpression promotes myeloid progenitor self-renewal and blocks lymphoid differentiation in mice. <i>Experimental Hematology</i> , 2015, 43, 565-577.e10.	0.2	43
59	Diversification and Functional Specialization of Human NK Cell Subsets. <i>Current Topics in Microbiology and Immunology</i> , 2015, 395, 63-93.	0.7	56
60	Case 32-2015. <i>New England Journal of Medicine</i> , 2015, 373, 1554-1564.	13.9	7
61	Primary Immunodeficiencies Associated with EBV Disease. <i>Current Topics in Microbiology and Immunology</i> , 2015, 390, 241-265.	0.7	109
62	Primary immunodeficiency update. <i>Journal of the American Academy of Dermatology</i> , 2015, 73, 367-381.	0.6	26
63	Pediatric myelodysplastic syndromes. <i>Journal of Hematopathology</i> , 2015, 8, 127-141.	0.2	8
64	Practice parameter for the diagnosis and management of primary immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 1186-1205.e78.	1.5	564
65	Cis-regulatory mechanisms governing stem and progenitor cell transitions. <i>Science Advances</i> , 2015, 1, e1500503.	4.7	57
66	Human Genetic Defects Resulting in Increased Susceptibility to Viral Infections. , 2016, , 375-388.		1
67	Post-infectious inflammatory response syndrome (PIIRS): Dissociation of T-cell-macrophage signaling in previously healthy individuals with cryptococcal fungal meningoencephalitis. <i>Macrophage</i> , 2015, 2, .	1.0	13
68	NK Cell Influence on the Outcome of Primary Epstein-Barr Virus Infection. <i>Frontiers in Immunology</i> , 2016, 7, 323.	2.2	48
69	Cancers Related to Immunodeficiencies: Update and Perspectives. <i>Frontiers in Immunology</i> , 2016, 7, 365.	2.2	137
70	Genetic Causes of Human NK Cell Deficiency and Their Effect on NK Cell Subsets. <i>Frontiers in Immunology</i> , 2016, 7, 545.	2.2	69
71	Hereditary Predispositions to Myelodysplastic Syndrome. <i>International Journal of Molecular Sciences</i> , 2016, 17, 838.	1.8	58
72	Acute lymphoblastic leukemia in a patient with MonoMAC syndrome/ <i>GATA2</i> haploinsufficiency. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1844-1847.	0.8	22
73	Severe disseminated primary herpes simplex infection as skin manifestation of <i>GATA2</i> deficiency. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016, 30, 1248-1250.	1.3	6
74	Successful reduced-intensity stem cell transplantation for <i>GATA2</i> deficiency before progression of advanced <i>MDS</i> . <i>Pediatric Transplantation</i> , 2016, 20, 333-336.	0.5	20
75	Monocyte, Macrophage, and Dendritic Cell Development: the Human Perspective. <i>Microbiology Spectrum</i> , 2016, 4, .	1.2	24

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76	Megakaryocytic Transcription Factors in Disease and Leukemia. , 2016, , 61-91.		1
77	Association of pulmonary alveolar proteinosis and fibrosis: patient with GATA2 deficiency. European Respiratory Journal, 2016, 48, 1510-1514.	3.1	23
78	Mutations in AML: prognostic and therapeutic implications. Hematology American Society of Hematology Education Program, 2016, 2016, 348-355.	0.9	136
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82	Gata2 Is a Rheostat for Mesenchymal Stem Cell Fate in Male Mice. Endocrinology, 2016, 157, 1021-1028.	1.4	10
83	Persistent nodal histoplasmosis in nuclear factor kappa B essential modulator deficiency: Report of a case and review of infection in primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2016, 138, 903-905.	1.5	14
84	Association of GATA2 Deficiency With Severe Primary Epstein-Barr Virus (EBV) Infection and EBV-associated Cancers. Clinical Infectious Diseases, 2016, 63, 41-47.	2.9	56
85	Epstein Barr virus "a tumor virus that needs cytotoxic lymphocytes to persist asymptotically. Current Opinion in Virology, 2016, 20, 34-39.	2.6	14
86	GATA Factor-Dependent Positive-Feedback Circuit in Acute Myeloid Leukemia Cells. Cell Reports, 2016, 16, 2428-2441.	2.9	59
87	Prevalence, clinical characteristics, and prognosis of GATA2-related myelodysplastic syndromes in children and adolescents. Blood, 2016, 127, 1387-1397.	0.6	304
88	Timely follow-up of a GATA2 deficiency patient allows successful treatment. Journal of Allergy and Clinical Immunology, 2016, 138, 1480-1483.e4.	1.5	7
89	The Hematopoietic Stem and Progenitor Cell Cistrome. Current Topics in Developmental Biology, 2016, 118, 45-76.	1.0	21
90	Loss of B cells and their precursors is the most constant feature of GATA-2 deficiency in childhood myelodysplastic syndrome. Haematologica, 2016, 101, 707-716.	1.7	51
91	Guidelines for the diagnosis and management of adult aplastic anaemia. British Journal of Haematology, 2016, 172, 187-207.	1.2	539
92	Merkel cell carcinoma in a patient with GATA 2 deficiency: a novel association with primary immunodeficiency. British Journal of Dermatology, 2016, 174, 169-171.	1.4	16
93	Mediastinal and Disseminated Mycobacterium kansasii Disease in GATA2 Deficiency. Annals of the American Thoracic Society, 2016, 13, 2169-2173.	1.5	11
94	Successful umbilical cord blood hematopoietic stem cell transplantation in pediatric patients with <scp>MDS</scp>/<scp>AML</scp> associated with underlying <scp>GATA</scp>2 mutations: two case reports and review of literature. Pediatric Transplantation, 2016, 20, 1004-1007.	0.5	6
95	Navigating Transcriptional Coregulator Ensembles to Establish Genetic Networks. Current Topics in Developmental Biology, 2016, 118, 205-244.	1.0	19

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96	Severe Epstein-Barr virus infection in primary immunodeficiency and the normal host. <i>British Journal of Haematology</i> , 2016, 175, 559-576.	1.2	47
97	How I diagnose and manage individuals at risk for inherited myeloid malignancies. <i>Blood</i> , 2016, 128, 1800-1813.	0.6	149
98	Primary Immune Deficiencies in the Adult: A Previously Underrecognized Common Condition. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2016, 4, 1101-1107.	2.0	28
99	Genetic predisposition to leukemia and other hematologic malignancies. <i>Seminars in Oncology</i> , 2016, 43, 598-608.	0.8	58
100	Myelodysplastic syndromes and acute leukemia with genetic predispositions: a new challenge for hematologists. <i>Expert Review of Hematology</i> , 2016, 9, 1189-1202.	1.0	19
101	Requirements for human natural killer cell development informed by primary immunodeficiency. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2016, 16, 541-548.	1.1	7
102	Single-gene association between GATA-2 and autoimmune hepatitis: A novel genetic insight highlighting immunologic pathways to disease. <i>Journal of Hepatology</i> , 2016, 64, 1190-1193.	1.8	23
103	Evaluation of Patients and Families With Concern for Predispositions to Hematologic Malignancies Within the Hereditary Hematologic Malignancy Clinic (HHMC). <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2016, 16, 417-428.e2.	0.2	74
104	“Why me, why now?” Using clinical immunology and epidemiology to explain who gets nontuberculous mycobacterial infection. <i>BMC Medicine</i> , 2016, 14, 54.	2.3	98
105	<i>Gata3</i> Hypomorphic Mutant Mice Rescued with a Yeast Artificial Chromosome Transgene Suffer a Glomerular Mesangial Cell Defect. <i>Molecular and Cellular Biology</i> , 2016, 36, 2272-2281.	1.1	11
106	X-linked Hyper IgM Syndrome Presenting as Pulmonary Alveolar Proteinosis. <i>Journal of Clinical Immunology</i> , 2016, 36, 564-570.	2.0	12
107	GATA factors in endocrine neoplasia. <i>Molecular and Cellular Endocrinology</i> , 2016, 421, 2-17.	1.6	19
108	Altered chemotactic response to CXCL12 in patients carrying <i>GATA2</i> mutations. <i>Journal of Leukocyte Biology</i> , 2016, 99, 1065-1076.	1.5	32
109	Loss of B Cells in Patients with Heterozygous Mutations in IKAROS. <i>New England Journal of Medicine</i> , 2016, 374, 1032-1043.	13.9	217
110	GATA2 deficiency underlying severe blastomycosis and fatal herpes simplex virus-associated hemophagocytic lymphohistiocytosis. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 638-640.	1.5	36
111	Genetic predisposition and hematopoietic malignancies in children: Primary immunodeficiency. <i>European Journal of Medical Genetics</i> , 2016, 59, 647-653.	0.7	15
112	Down-regulation of EZH2 expression in myelodysplastic syndromes. <i>Leukemia Research</i> , 2016, 44, 1-7.	0.4	13
113	Familial skin cancer syndromes. <i>Journal of the American Academy of Dermatology</i> , 2016, 74, 437-451.	0.6	46

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114	Flow Cytometry, a Versatile Tool for Diagnosis and Monitoring of Primary Immunodeficiencies. <i>Vaccine Journal</i> , 2016, 23, 254-271.	3.2	76
115	NK cells and cancer: you can teach innate cells new tricks. <i>Nature Reviews Cancer</i> , 2016, 16, 7-19.	12.8	903
116	Genetic predisposition to myelodysplastic syndrome and acute myeloid leukemia in children and young adults. <i>Leukemia and Lymphoma</i> , 2016, 57, 520-536.	0.6	96
117	A genetic perspective on granulomatous diseases with an emphasis on mycobacterial infections. <i>Seminars in Immunopathology</i> , 2016, 38, 199-212.	2.8	16
118	Defects of Innate Immunity. , 2016, , 101-111.e3.		1
119	Pulmonary Complications of Primary Immunodeficiencies. , 2016, , 1624-1638.e4.		0
120	Pulmonary Alveolar Proteinosis Syndrome. , 2016, , 1260-1274.e12.		4
121	Transcriptional and post-transcriptional regulation of NK cell development and function. <i>Clinical Immunology</i> , 2017, 177, 60-69.	1.4	23
122	Natural killer cell biology illuminated by primary immunodeficiency syndromes in humans. <i>Clinical Immunology</i> , 2017, 177, 29-42.	1.4	26
123	Exome sequencing identifies highly recurrent somatic GATA2 and CEBPA mutations in acute erythroid leukemia. <i>Leukemia</i> , 2017, 31, 195-202.	3.3	37
124	<i>Pneumocystis jiroveci</i> pneumonia and GATA2 deficiency: Expanding the spectrum of the disease. <i>Journal of Infection</i> , 2017, 74, 425-427.	1.7	9
125	Gain-of-function SAMD9L mutations cause a syndrome of cytopenia, immunodeficiency, MDS, and neurological symptoms. <i>Blood</i> , 2017, 129, 2266-2279.	0.6	152
126	Adaptive NK cells can persist in patients with GATA2 mutation depleted of stem and progenitor cells. <i>Blood</i> , 2017, 129, 1927-1939.	0.6	89
127	Germline Genetic Predisposition to Hematologic Malignancy. <i>Journal of Clinical Oncology</i> , 2017, 35, 1018-1028.	0.8	80
128	GATA2 null mutation associated with incomplete penetrance in a family with Emberger syndrome. <i>Hematology</i> , 2017, 22, 1-5.	0.7	8
129	The GATA factor revolution in hematology. <i>Blood</i> , 2017, 129, 2092-2102.	0.6	115
130	GATA factor mutations in hematologic disease. <i>Blood</i> , 2017, 129, 2103-2110.	0.6	149
131	GATA2 deficiency and related myeloid neoplasms. <i>Seminars in Hematology</i> , 2017, 54, 81-86.	1.8	125



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132	Haemodynamically proven pulmonary hypertension in a patient with GATA2 deficiency-associated pulmonary alveolar proteinosis and fibrosis. <i>European Respiratory Journal</i> , 2017, 49, 1700178.	3.1	9
133	Haemodynamically proven pulmonary hypertension in a patient with GATA2 deficiency-associated pulmonary alveolar proteinosis and fibrosis. <i>European Respiratory Journal</i> , 2017, 49, 1700407.	3.1	8
134	Practical considerations for diagnosis and management of patients and carriers. <i>Seminars in Hematology</i> , 2017, 54, 69-74.	1.8	15
135	Primary Immunodeficiency Diseases. , 2017, , .		22
137	Turning the tide in myelodysplastic/myeloproliferative neoplasms. <i>Nature Reviews Cancer</i> , 2017, 17, 425-440.	12.8	117
138	Predispositions to Leukemia in Down Syndrome and Other Hereditary Disorders. <i>Current Treatment Options in Oncology</i> , 2017, 18, 41.	1.3	15
139	Recommendations for Surveillance for Children with Leukemia-Predisposing Conditions. <i>Clinical Cancer Research</i> , 2017, 23, e14-e22.	3.2	80
141	Allogeneic Hematopoietic Cell Transplantation Using Treosulfan-Based Conditioning for Treatment of Marrow Failure Disorders. <i>Biology of Blood and Marrow Transplantation</i> , 2017, 23, 1669-1677.	2.0	45
142	Immunogenetics of Fungal Diseases. , 2017, , .		2
143	WILD syndrome is GATA2 deficiency: A novel deletion in the GATA2 gene. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2017, 5, 1149-1152.e1.	2.0	16
144	Fungal Infections in Primary and Acquired Immunodeficiencies. , 2017, , 1-34.		0
145	Resolution of Multifocal Epstein-Barr Virus-Related Smooth Muscle Tumor in a Patient with GATA2 Deficiency Following Hematopoietic Stem Cell Transplantation. <i>Journal of Clinical Immunology</i> , 2017, 37, 61-66.	2.0	20
146	Multiple Opportunistic Infections in a Woman with GATA2 Mutation. <i>International Journal of Infectious Diseases</i> , 2017, 54, 89-91.	1.5	11
147	Introduction on Primary Immunodeficiency Diseases. , 2017, , 1-81.		3
148	Vulvar lymphedema and refractory VIN-III heralding GATA2 deficiency syndrome. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2017, 218, 138-140.	0.5	5
149	Familial myelodysplastic syndrome/acute myeloid leukemia. <i>Best Practice and Research in Clinical Haematology</i> , 2017, 30, 287-289.	0.7	25
150	Long-Term Survival After Hematopoietic Stem Cell Transplantation for Complete STAT1 Deficiency. <i>Journal of Clinical Immunology</i> , 2017, 37, 701-706.	2.0	21
151	Integrating Enhancer Mechanisms to Establish a Hierarchical Blood Development Program. <i>Cell Reports</i> , 2017, 20, 2966-2979.	2.9	46

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152	Natural killer cell-mediated immunosurveillance of human cancer. <i>Seminars in Immunology</i> , 2017, 31, 20-29.	2.7	240
153	Case 28-2017. <i>New England Journal of Medicine</i> , 2017, 377, 1077-1091.	13.9	3
154	Transplant for NEMO: this and much, much more. <i>Blood</i> , 2017, 130, 1391-1393.	0.6	2
155	Congenital neutropenia in the era of genomics: classification, diagnosis, and natural history. <i>British Journal of Haematology</i> , 2017, 179, 557-574.	1.2	115
156	Successful Myeloablative Matched Unrelated Donor Hematopoietic Stem Cell Transplantation in a Young Girl With GATA2 Deficiency and Emberger Syndrome. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, 230-232.	0.3	8
157	Pulmonary Alveolar Proteinosis: A Comprehensive Clinical Perspective. <i>Pediatrics</i> , 2017, 140, e20170610.	1.0	45
158	Proposed Terminology and Classification of Pre-Malignant Neoplastic Conditions: A Consensus Proposal. <i>EBioMedicine</i> , 2017, 26, 17-24.	2.7	24
159	Intraluminal valves: development, function and disease. <i>DMM Disease Models and Mechanisms</i> , 2017, 10, 1273-1287.	1.2	48
160	Autoimmunity/inflammation in a monogenic primary immunodeficiency cohort. <i>Clinical and Translational Immunology</i> , 2017, 6, e155.	1.7	27
161	Heterogeneity of GATA2-related myeloid neoplasms. <i>International Journal of Hematology</i> , 2017, 106, 175-182.	0.7	44
162	Germline Mutations Associated with Leukemia in Childhood: New Discoveries and Emerging Phenotypes. <i>Current Genetic Medicine Reports</i> , 2017, 5, 59-65.	1.9	4
163	Modern management of phagocyte defects. <i>Pediatric Allergy and Immunology</i> , 2017, 28, 124-134.	1.1	9
164	Mutational profiling of a MonoMAC syndrome family with GATA2 deficiency. <i>Leukemia</i> , 2017, 31, 244-245.	3.3	22
165	Immunodeficiencies. , 2017, , 705-722.e2.		2
166	Evaluation of the HIV-Uninfected Adult with Suspected Immunodeficiency. , 2017, , 808-811.e1.		0
167	The role of GATA2 in lethal prostate cancer aggressiveness. <i>Nature Reviews Urology</i> , 2017, 14, 38-48.	1.9	71
168	Cryptococcal meningitis: epidemiology, immunology, diagnosis and therapy. <i>Nature Reviews Neurology</i> , 2017, 13, 13-24.	4.9	344
169	Case Report of an Adolescent Male With Unexplained Pancytopenia: <i>GATA2</i>-Associated Bone Marrow Failure and Genetic Testing. <i>Global Pediatric Health</i> , 2017, 4, 2333794X1774494.	0.3	2

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170	Chronic neutropenia in adults. <i>Hematologie</i> , 2017, 23, 333-337.	0.0	0
171	New monogenic disorders identify more pathways to neutropenia: from the clinic to next-generation sequencing. <i>Hematology American Society of Hematology Education Program</i> , 2017, 2017, 172-180.	0.9	7
172	Somatic mutations in children with GATA2-associated myelodysplastic syndrome who lack other features of GATA2 deficiency. <i>Blood Advances</i> , 2017, 1, 443-448.	2.5	23
173	Monocyte, Macrophage, and Dendritic Cell Development: the Human Perspective. , 2017, , 79-97.		1
174	Recurrent and Sustained Viral Infections in Primary Immunodeficiencies. <i>Frontiers in Immunology</i> , 2017, 8, 665.	2.2	37
175	Cellular and Molecular Defects Underlying Invasive Fungal Infections—Revelations from Endemic Mycoses. <i>Frontiers in Immunology</i> , 2017, 8, 735.	2.2	57
176	Acquired Senescent T-Cell Phenotype Correlates with Clinical Severity in GATA Binding Protein 2-Deficient Patients. <i>Frontiers in Immunology</i> , 2017, 8, 802.	2.2	18
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