Andrew Gennery

List of Publications by Year
in descending order

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311 papers	15,872 citations	12330 69 h-index	22166 113 g-index
405 all docs	405 docs citations	405 times ranked	13658 citing authors

#	Article	IF	Citations
1	International retrospective study of allogeneic hematopoietic cell transplantation for activated PI3K-delta syndrome. Journal of Allergy and Clinical Immunology, 2022, 149, 410-421.e7.	2.9	34
2	Comparison of hematopoietic cell transplant conditioning regimens for hemophagocytic lymphohistiocytosis disorders. Journal of Allergy and Clinical Immunology, 2022, 149, 1097-1104.e2.	2.9	16
3	Recommendations for uniform definitions used in newborn screening for severe combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2022, 149, 1428-1436.	2.9	19
4	Hematopoietic cell transplantation in severe combined immunodeficiency: The SCETIDE 2006-2014 European cohort. Journal of Allergy and Clinical Immunology, 2022, 149, 1744-1754.e8.	2.9	51
5	Hematopoietic stem cell transplantation for Wiskott-Aldrich syndrome: an EBMT Inborn ErrorsÂWorking Party analysis. Blood, 2022, 139, 2066-2079.	1.4	33
6	Hematopoietic Cell Transplant for CD40 Ligand Deficiencyâ€"Comparing Busulfan Versus Treosulfan. Journal of Clinical Immunology, 2022, 42, 703-705.	3.8	6
7	Outcomes following SARS-CoV-2 infection in patients with primary and secondary immunodeficiency in the UK. Clinical and Experimental Immunology, 2022, 209, 247-258.	2.6	25
8	A Prospective, Multicenter Study of Closed System Extracorporeal Photopheresis for Children With Steroid-Refractory Acute Graft-Versus-Host Disease. Transplantation and Cellular Therapy, 2022, , .	1.2	3
9	T-replete HLA-matched grafts vs T-depleted HLA-mismatched grafts in inborn errors of immunity. Blood Advances, 2022, 6, 1319-1328.	5.2	8
10	Management of Chronic Graft-vs.-Host Disease in Children and Adolescents With ALL: Present Status and Model for a Personalised Management Plan. Frontiers in Pediatrics, 2022, 10, 808103.	1.9	8
11	Primary immune regulatory disorders: Undiagnosed needles in the haystack?. Orphanet Journal of Rare Diseases, 2022, 17, 99.	2.7	9
12	Hematopoietic stem cell transplantation for adolescents and adults with inborn errors of immunity: an EBMT IEWP study. Blood, 2022, 140, 1635-1649.	1.4	20
13	Sodium stibogluconate and CD47-SIRPÎ± blockade overcome resistance of anti-CD20â€"opsonized B cells to neutrophil killing. Blood Advances, 2022, 6, 2156-2166.	5.2	12
14	TCRÎ±Î²-Depleted Haploidentical Grafts Are a Safe Alternative to HLA-Matched Unrelated Donor Stem Cell Transplants for Infants with Severe Combined Immunodeficiency. Journal of Clinical Immunology, 2022, 42, 851-858.	3.8	13
15	22q11.2 Deletion and Duplication Syndromes and COVID-19. Journal of Clinical Immunology, 2022, 42, 746-748.	3.8	2
16	Allogeneic hematopoietic stem cell transplantation for adult HLH: a retrospective study by the chronic malignancies and inborn errors working parties of EBMT. Bone Marrow Transplantation, 2022, 57, 817-823.	2.4	4
17	Hematopoietic Cell Transplantation for Adenosine Deaminase Severe Combined Immunodeficiencyâ€"Improved Outcomes in the Modern Era. Journal of Clinical Immunology, 2022, 42, 819-826.	3.8	8
18	Biomarkers for Diagnosing Febrile Illness in Immunocompromised Children: A Systematic Review of the Literature. Frontiers in Pediatrics, 2022, 10, 828569.	1.9	3

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19	Preemptive hematopoietic cell transplantation for asymptomatic patients with X-linked lymphoproliferative syndrome type 1. Clinical Immunology, 2022, 237, 108993.	3.2	1
20	Phenotype, genotype, treatment, and survival outcomes in patients with X-linked inhibitor of apoptosis deficiency. Journal of Allergy and Clinical Immunology, 2022, 150, 456-466.	2.9	15
21	Autoimmunity in combined immunodeficiency. , 2022, , 97-127.		1
22	Consensus Recommendations for the Clinical Management of Hematological Malignancies in Patients with DNA Double Stranded Break Disorders. Cancers, 2022, 14, 2000.	3.7	5
23	Granulocyte Transfusions in Patients with Chronic Granulomatous Disease Undergoing Hematopoietic Cell Transplantation or Gene Therapy. Journal of Clinical Immunology, 2022, 42, 1026-1035.	3.8	3
24	Seek and you shall find: immune lymphoid cells in holobiont health. Blood, 2022, 139, 2577-2578.	1.4	1
25	HSCT is effective in patients with PSTPIP1-associated myeloid-related proteinemia inflammatory (PAMI) syndrome. Journal of Allergy and Clinical Immunology, 2021, 148, 250-255.e1.	2.9	18
26	Progress in treating chronic granulomatous disease. British Journal of Haematology, 2021, 192, 251-264.	2.5	17
27	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	2.9	278
28	Long-term outcomes for adults with chronic granulomatous disease in the United Kingdom. Journal of Allergy and Clinical Immunology, 2021, 147, 1104-1107.	2.9	10
29	Hematopoietic Stem Cell Transplantation Positively Affects the Natural History of Cancer in Nijmegen Breakage Syndrome. Clinical Cancer Research, 2021, 27, 575-584.	7.0	13
30	Improved survival and graft function in ex vivo T-cell depleted haploidentical hematopoietic cell transplantation for primary immunodeficiency. Bone Marrow Transplantation, 2021, 56, 1200-1204.	2.4	11
31	COVID-19 in patients with primary and secondary immunodeficiency: The United Kingdom experience. Journal of Allergy and Clinical Immunology, 2021, 147, 870-875.e1.	2.9	188
32	Transplantation for congenital heart disease is associated with an increased risk of Epstein-Barr virusâ€“related post-transplant lymphoproliferative disorder in children. Journal of Heart and Lung Transplantation, 2021, 40, 24-32.	0.6	7
33	Outcome of Non-hematological Autoimmunity After Hematopoietic Cell Transplantation in Children with Primary Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 171-184.	3.8	5
34	European dermatology forum: Updated guidelines on the use of extracorporeal photopheresis 2020 â€“ Part 2. Journal of the European Academy of Dermatology and Venereology, 2021, 35, 27-49.	2.4	28
35	Treosulfan-based conditioning for inborn errors of immunity. Therapeutic Advances in Hematology, 2021, 12, 204062072110139.	2.5	6
36	Hematopoietic Stem Cell Transplantation Resolves the Immune Deficit Associated with STAT3-Dominant-Negative Hyper-IgE Syndrome. Journal of Clinical Immunology, 2021, 41, 934-943.	3.8	21

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37	Thymopoiesis, Alterations in Dendritic Cells and Tregs, and Reduced T Cell Activation in Successful Extracorporeal Photopheresis Treatment of GVHD. Journal of Clinical Immunology, 2021, 41, 1016-1030.	3.8	9
38	Adenosine Deaminase Deficient SCID with Myocardial Hypertrophy. Journal of Clinical Immunology, 2021, 41, 1128-1130.	3.8	3
39	STAT3 Hyper-IgE Syndromeâ€"an Update and Unanswered Questions. Journal of Clinical Immunology, 2021, 41, 864-880.	3.8	63
40	Haematopoietic Stem Cell Transplant for Norovirus-Induced Intestinal Failure in X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2021, 41, 1574-1581.	3.8	4
41	Thymic Epithelial Cell Alterations and Defective Thymopoiesis Lead to Central and Peripheral Tolerance Perturbation in MHCII Deficiency. Frontiers in Immunology, 2021, 12, 669943.	4.8	8
42	Congenital nephrotic syndrome in IL7RÎ±-SCID: A rare feature of maternofetal graft-versus-host disease. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 4151-4153.e1.	3.8	2
43	EBMT/ESID inborn errors working party guidelines for hematopoietic stem cell transplantation for inborn errors of immunity. Bone Marrow Transplantation, 2021, 56, 2052-2062.	2.4	95
44	Newborn screening for severe combined immunodeficiencyâ€"Coming to a region near you soon. Clinical and Experimental Immunology, 2021, 205, 343-345.	2.6	2
45	Stem cell transplantation as treatment for major histocompatibility class I deficiency. Clinical Immunology, 2021, 229, 108801.	3.2	3
46	HSCT in two brothers with CGD arising from mutations in CYBC1 corrects the defect in neutrophil function. Clinical Immunology, 2021, 229, 108799.	3.2	3
47	Outcome of Hematopoietic Stem Cell Transplantation in patients with Mendelian Susceptibility to Mycobacterial Diseases. Journal of Clinical Immunology, 2021, 41, 1774-1780.	3.8	3
48	Hematopoietic Cell Transplantation Ameliorates Autoinflammation in A20 Haploinsufficiency. Journal of Clinical Immunology, 2021, 41, 1954-1956.	3.8	9
49	COVID-19 and X-linked agammaglobulinemia (XLA) â€“ insights from a monogenic antibody deficiency. Current Opinion in Allergy and Clinical Immunology, 2021, 21, 525-534.	2.3	22
50	Management of inborn errors of immunity. , 2021, , 345-361.		0
51	Evaluation of newborn screening for severe combined immunodeficiency (SCID). British Journal of General Practice, 2021, 71, 456-457.	1.4	0
52	Establishing Newborn Screening for SCID in the USA: Experience in California. International Journal of Neonatal Screening, 2021, 7, 72.	3.2	7
53	Allogeneic hematopoietic stem cell transplantation in leukocyte adhesion deficiency type I and III. Blood Advances, 2021, 5, 262-273.	5.2	9
54	UK national survey of anticoagulation in extraâ€corporeal photopheresisâ€"Is it time for a UK consensus statement?. Transfusion Medicine, 2021, , .	1.1	0

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55	Granulomatousâ€“lymphocytic interstitial lung disease: an international research prioritisation. ERJ Open Research, 2021, 7, 00467-2021.	2.6	6
56	The EHA Research Roadmap: Infections in Hematology. HemaSphere, 2021, 5, e662.	2.7	5
57	Combined T and B Lymphocyte Deficiencies. , 2021, , .		0
58	Proposed Therapeutic Range of Treosulfan in Reduced Toxicity Pediatric Allogeneic Hematopoietic Stem Cell Transplant Conditioning: Results From a Prospective Trial. Clinical Pharmacology and Therapeutics, 2020, 108, 264-273.	4.7	22
59	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. Journal of Allergy and Clinical Immunology, 2020, 145, 1452-1463.	2.9	112
60	Developing a career in academic paediatrics. Paediatrics and Child Health (United Kingdom), 2020, 30, 70-73.	0.4	0
61	European dermatology forum â€“ updated guidelines on the use of extracorporeal photopheresis 2020 â€“ part 1. Journal of the European Academy of Dermatology and Venereology, 2020, 34, 2693-2716.	2.4	49
62	Hematopoietic Stem Cell Transplantation and Vasculopathy Associated With STAT3-Dominant-Negative Hyper-IgE Syndrome. Frontiers in Pediatrics, 2020, 8, 575.	1.9	7
63	Managing Granulomatousâ€“Lymphocytic Interstitial Lung Disease in Common Variable Immunodeficiency Disorders: e-GLILDnet International Clinicians Survey. Frontiers in Immunology, 2020, 11, 606333.	4.8	10
64	CD27-CD70 defects: a wolf in wolfâ€™s clothing?. Blood, 2020, 136, 2600-2602.	1.4	2
65	Systematic Review of Primary Immunodeficiency Diseases in Malaysia: 1979â€“2020. Frontiers in Immunology, 2020, 11, 1923.	4.8	11
66	The challenges presented by haematopoietic stem cell transplantation in children with primary immunodeficiency. British Medical Bulletin, 2020, 135, 4-15.	6.9	3
67	Diagnosis and management of severe combined immunodeficiency in Australia and New Zealand. Journal of Paediatrics and Child Health, 2020, 56, 1508-1513.	0.8	4
68	UK national audit of extracorporeal photopheresis (ECP) in chronic graft versus host disease. Leukemia and Lymphoma, 2020, 61, 3511-3514.	1.3	1
69	A survey of extracorporeal photopheresis treatment in pediatric patients in the United Kingdom. EJHaem, 2020, 1, 293-296.	1.0	1
70	COVIDâ€19 â€“ Impact on Childhood Haematology Patients. HemaSphere, 2020, 4, e465.	2.7	9
71	Guidelines on the use of irradiated blood components. British Journal of Haematology, 2020, 191, 704-724.	2.5	61
72	BCG lymphadenitis: a potential complication of immune reconstitution following haematopoietic stem cell transplant. Archives of Disease in Childhood: Education and Practice Edition, 2020, , edpract-2020-320883.	0.5	1

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73	HSCT provides effective treatment for lymphoproliferative disorders in children with primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2020, 146, 447-450.	2.9	8
74	Targeted busulfan-based reduced-intensity conditioning and HLA-matched HSCT cure hemophagocytic lymphohistiocytosis. Blood Advances, 2020, 4, 1998-2010.	5.2	30
75	Impact of Different Ex Vivo T-Cell Depletion Strategies on Outcomes Following Hematopoietic Cell Transplantation for Children with Primary Immunodeficiency. Biology of Blood and Marrow Transplantation, 2020, 26, S17-S18.	2.0	0
76	New insights into risk factors for transplant-associated thrombotic microangiopathy in pediatric HSCT. Blood Advances, 2020, 4, 2418-2429.	5.2	24
77	Hematopoietic Cell Transplantation in Patients With Primary Immune Regulatory Disorders (PIRD): A Primary Immune Deficiency Treatment Consortium (PIDTC) Survey. Frontiers in Immunology, 2020, 11, 239.	4.8	57
78	Neonatal thymectomy in childrenâ€"accelerating the immunologic clock?. Journal of Allergy and Clinical Immunology, 2020, 146, 236-243.	2.9	27
79	Update on DNA-Double Strand Break Repair Defects in Combined Primary Immunodeficiency. Current Allergy and Asthma Reports, 2020, 20, 57.	5.3	15
80	Hematopoietic cell transplantation in chronic granulomatous disease: a study of 712 children and adults. Blood, 2020, 136, 1201-1211.	1.4	97
81	Hematopoietic stem cell transplantation for primary immune deficiencies. , 2020, , 1175-1214.		0
82	Outcome and Risk Factors of Autoimmune Cytopenia after Hematopoietic Cell Transplantation for Children with Primary Immunodeficiency. Biology of Blood and Marrow Transplantation, 2020, 26, S208.	2.0	0
83	Primary Immunodeficiency Diseases and Bacillus Calmette-GuÃ©rin (BCG)-Vaccineâ€"Derived Complications: A Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1371-1386.	3.8	51
84	International Retrospective Study of Allogeneic Hematopoietic Cell Transplantation (HCT) for Activated Phosphoinositide 3-Kinase Delta (PI3K) Syndrome. Biology of Blood and Marrow Transplantation, 2020, 26, S14-S15.	2.0	4
85	Cutaneous barrier leakage and gut inflammation drive skin disease in Omenn syndrome. Journal of Allergy and Clinical Immunology, 2020, 146, 1165-1179.e11.	2.9	13
86	Improved transplant survival and long-term disease outcome in children with MHC class II deficiency. Blood, 2020, 135, 954-973.	1.4	23
87	Outcome of autoimmune cytopenia after hematopoietic cell transplantation in primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2020, 146, 406-416.	2.9	18
88	Conditioning Regimens and Outcomes after Allogeneic Hematopoietic Cell Transplant for Hyperinflammatory Inborn Errors of Immunity. Blood, 2020, 136, 36-37.	1.4	0
89	Hematopoietic stem cell transplantation for cytidine triphosphate synthase 1 (CTPS1) deficiency. Bone Marrow Transplantation, 2019, 54, 130-133.	2.4	13
90	Recombination activity of human recombination-activating gene 2 (RAG2) mutations and correlation with clinical phenotype. Journal of Allergy and Clinical Immunology, 2019, 143, 726-735.	2.9	39

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91	Editorial: The Relationship Between Cancer Predisposition and Primary Immunodeficiency. Frontiers in Immunology, 2019, 10, 1781.	4.8	15
92	Allogeneic HSCT for Autoimmune Diseases: A Retrospective Study From the EBMT ADWP, IEWP, and PDWP Working Parties. Frontiers in Immunology, 2019, 10, 1570.	4.8	48
93	Late relapse of primary hemophagocytic lymphohistiocytosis after hematopoietic stem cell transplantation: a consequence of low-level chimerism from a carrier donor?. Immunologic Research, 2019, 67, 261-264.	2.9	0
94	Long Term Outcome and Immune Function After Hematopoietic Stem Cell Transplantation for Primary Immunodeficiency. Frontiers in Pediatrics, 2019, 7, 381.	1.9	31
95	Adoptive T Cell Therapy Strategies for Viral Infections in Patients Receiving Haematopoietic Stem Cell Transplantation. Cells, 2019, 8, 47.	4.1	32
96	Gonadal Function after Busulfan Compared with Treosulfan in Children and Adolescents Undergoing Allogeneic Hematopoietic Stem Cell Transplant. Biology of Blood and Marrow Transplantation, 2019, 25, 1786-1791.	2.0	42
97	Recent advances in understanding RAG deficiencies. F1000Research, 2019, 8, 148.	1.6	27
98	New graft manipulation strategies improve the outcome of mismatched stem cell transplantation in children with primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2019, 144, 280-293.	2.9	35
99	Two decades of excellent transplant survival for chronic granulomatous disease: a supraregional immunology transplant center report. Blood, 2019, 133, 2546-2549.	1.4	26
100	Health-Related Quality of Life and Emotional Health in X-Linked Carriers of Chronic Granulomatous Disease in the United Kingdom. Journal of Clinical Immunology, 2019, 39, 195-199.	3.8	9
101	Indications for haematopoietic stem cell transplantation for haematological diseases, solid tumours and immune disorders: current practice in Europe, 2019. Bone Marrow Transplantation, 2019, 54, 1525-1552.	2.4	218
102	Combined liver and hematopoietic stem cell transplantation in patients with X-linked hyper-IgM syndrome. Journal of Allergy and Clinical Immunology, 2019, 143, 1952-1956.e6.	2.9	10
103	Hematopoietic Cell Transplantation for MHC Class II Deficiency. Frontiers in Pediatrics, 2019, 7, 516.	1.9	13
104	Universal donor strategy for primary immunodeficiency. Blood, 2019, 134, 1688-1689.	1.4	0
105	Choice of conditioning regimens for bone marrow transplantation in severe aplastic anemia. Blood Advances, 2019, 3, 3123-3131.	5.2	37
106	Conditioning Regimens for Hematopoietic Cell Transplantation in Primary Immunodeficiency. Current Allergy and Asthma Reports, 2019, 19, 52.	5.3	19
107	Hematopoietic Stem Cell Transplantation for Primary Immunodeficiencies. Frontiers in Pediatrics, 2019, 7, 445.	1.9	24
108	An update on X-Linked agammaglobulinaemia: clinical manifestations and management. Current Opinion in Allergy and Clinical Immunology, 2019, 19, 571-577.	2.3	11

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109	Solid organ transplantation after hematopoietic stem cell transplantation in childhood: A multicentric retrospective survey. American Journal of Transplantation, 2019, 19, 1798-1805.	4.7	9
110	Hematopoietic Stem Cell Transplantation as Treatment for Patients with DOCK8 Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 848-855.	3.8	67
111	Pulmonary Manifestations of Combined T- and B-Cell Immunodeficiencies. , 2019, , 37-75.		0
112	Hematopoietic Stem Cell Transplantation for DNA Double Strand Breakage Repair Disorders. Frontiers in Pediatrics, 2019, 7, 557.	1.9	14
113	Primary Immunodeficiencies. , 2019, , 663-670.		2
114	Class-Switch Recombination Defects. Rare Diseases of the Immune System, 2019, , 179-199.	0.1	0
115	Erythrodermic Rash and Seizures. , 2019, , 695-699.		0
116	Thymopoiesis following HSCT; a retrospective review comparing interventions for aGVHD in a pediatric cohort. Clinical Immunology, 2018, 193, 33-37.	3.2	9
117	Endothelial cell damage in idiopathic pneumonia syndrome. Bone Marrow Transplantation, 2018, 53, 515-518.	2.4	13
118	In vivo T-depleted reduced-intensity transplantation for GATA2-related immune dysfunction. Blood, 2018, 131, 1383-1387.	1.4	32
119	Hematopoietic cell transplantation in primary immunodeficiency â€“ conventional and emerging indications. Expert Review of Clinical Immunology, 2018, 14, 103-114.	3.0	42
120	Adenosine deaminase deficiency: a review. Orphanet Journal of Rare Diseases, 2018, 13, 65.	2.7	144
121	Recommendations from the European Society for Blood and Marrow Transplantation (EBMT) for a curriculum in hematopoietic cell transplantation. Bone Marrow Transplantation, 2018, 53, 1548-1552.	2.4	6
122	Non-posttransplant lymphoproliferative disorder malignancy after hematopoietic stem cell transplantation in patients with primary immunodeficiency: UK experience. Journal of Allergy and Clinical Immunology, 2018, 141, 2319-2321.e1.	2.9	7
123	Ikaros family zinc finger 1 regulates dendritic cell development and function in humans. Nature Communications, 2018, 9, 1239.	12.8	62
124	Clinical considerations in the hematopoietic stem cell transplant management of primary immunodeficiencies. Expert Review of Clinical Immunology, 2018, 14, 297-306.	3.0	24
125	Outcome of hematopoietic cell transplantation for DNA double-strand break repair disorders. Journal of Allergy and Clinical Immunology, 2018, 141, 322-328.e10.	2.9	79
126	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

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127	Hematopoietic stem cell transplantation in patients with gain-of-function signal transducer and activator of transcription 1 mutations. Journal of Allergy and Clinical Immunology, 2018, 141, 704-717.e5.	2.9	128
128	Treosulfan and Fludarabine Conditioning for Hematopoietic Stem Cell Transplantation in Children with Primary Immunodeficiency: UK Experience. Biology of Blood and Marrow Transplantation, 2018, 24, 529-536.	2.0	75
129	Biallelic interferon regulatory factor 8 mutation: AÂcomplex immunodeficiency syndrome with dendritic cell deficiency, monocytopenia, and immune dysregulation. Journal of Allergy and Clinical Immunology, 2018, 141, 2234-2248.	2.9	63
130	T-cell receptor Î±Î²+ and CD19+ cellâ€“depleted haploidentical and mismatched hematopoietic stem cell transplantation in primary immune deficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1417-1426.e1.	2.9	119
131	Transplantation of Hematopoietic Stem Cells for Primary Immunodeficiencies in Brazil: Challenges in Treating Rare Diseases in Developing Countries. Journal of Clinical Immunology, 2018, 38, 917-926.	3.8	13
132	Current Understanding and Future Research Priorities in Malignancy Associated With Inborn Errors of Immunity and DNA Repair Disorders: The Perspective of an Interdisciplinary Working Group. Frontiers in Immunology, 2018, 9, 2912.	4.8	48
133	Predicting the future with TRECs. Blood, 2018, 132, 1731-1733.	1.4	2
134	Jakinibs for the treatment of immune dysregulation in patients with gain-of-function signal transducer and activator of transcription 1 (STAT1) or STAT3 mutations. Journal of Allergy and Clinical Immunology, 2018, 142, 1665-1669.	2.9	196
135	Allogeneic hematopoietic stem cell transplantation for severe, refractory juvenile idiopathic arthritis. Blood Advances, 2018, 2, 777-786.	5.2	37
136	Early and late outcomes after cord blood transplantation for pediatric patients with inherited leukodystrophies. Blood Advances, 2018, 2, 49-60.	5.2	45
137	Multicenter phase 1/2 application of adenovirus-specific T cells in high-risk pediatric patients after allogeneic stem cell transplantation. Cytotherapy, 2018, 20, 830-838.	0.7	20
138	Outcome of domino hematopoietic stem cell transplantation in human subjects: An international case series. Journal of Allergy and Clinical Immunology, 2018, 142, 1628-1631.e4.	2.9	1
139	Interleukin-2-Inducible T-Cell Kinase Deficiencyâ€"New Patients, New Insight?. Frontiers in Immunology, 2018, 9, 979.	4.8	31
140	Long-Term Health Outcome and Quality of Life Post-HSCT for IL7RÎ±-, Artemis-, RAG1- and RAG2-Deficient Severe Combined Immunodeficiency: a Single Center Report. Journal of Clinical Immunology, 2018, 38, 727-732.	3.8	32
141	The International Alliance of Primary Immune Deficiency Societies. Journal of Clinical Immunology, 2018, 38, 447-449.	3.8	2
142	Advances in genetic and molecular understanding of Omenn syndrome - implications for the future. Expert Opinion on Orphan Drugs, 2018, 6, 351-359.	0.8	2
143	Busulfan/Fludarabine- or Treosulfan/Fludarabine-Based Conditioning Regimen in Patients with Wiskott-Aldrich Syndrome Given Allogeneic Hematopoietic Cell Transplantation â€" an EBMT Inborn Errors Working Party and Scetide Retrospective Analysis. Blood, 2018, 132, 2175-2175.	1.4	4
144	Allogeneic Hematopoietic Stem Cell Transplantation in Children and Adults with Chronic Granulomatous Disease (CGD): A Study of the Inborn Errors Working Party (IEWP) of the EBMT. Blood, 2018, 132, 970-970.	1.4	2

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145	Current Knowledge and Priorities for Future Research in Late Effects after Hematopoietic Stem Cell Transplantation (HCT) for Severe Combined Immunodeficiency Patients: A Consensus Statement from the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects after Pediatric HCT. Biology of Blood and Marrow Transplantation, 2017, 23, 379-387.	2.0	49
146	Use of defibrotide to treat transplant-associated thrombotic microangiopathy: a retrospective study of the Paediatric Diseases and Inborn Errors Working Parties of the European Society of Blood and Marrow Transplantation. Bone Marrow Transplantation, 2017, 52, 762-764.	2.4	39
147	Long-term outcome of hematopoietic stem cell transplantation for IL2RG/JAK3 SCID: a cohort report. Blood, 2017, 129, 2198-2201.	1.4	54
148	The role of extracorporeal photopheresis in the management of cutaneous Tâ€cell lymphoma, graftâ€versusâ€host disease and organ transplant rejection: a consensus statement update from the UK Photopheresis Society. British Journal of Haematology, 2017, 177, 287-310.	2.5	109
149	A risk factor analysis of outcomes after unrelated cord blood transplantation for children with Wiskott-Aldrich syndrome. Haematologica, 2017, 102, 1112-1119.	3.5	30
150	Raised Serum IL-8 Levels Are Associated with Excessive Fatigue in Female Carriers of X-Linked Chronic Granulomatous Disease in the UK. Journal of Clinical Immunology, 2017, 37, 279-281.	3.8	6
151	Recommendations for Screening and Management of Late Effects in Patients with Severe Combined Immunodeficiency after Allogenic Hematopoietic Cell Transplantation: A Consensus Statement from the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects after Pediatric HCT. Biology of Blood and Marrow Transplantation, 2017, 23, 1229-1240.	2.0	44
152	Enterovirus-Related Immune Reconstitution Inflammatory Syndrome (IRIS) Following Haploidentical Stem Cell Transplantation in an MHC Class II-Deficient Child. Journal of Clinical Immunology, 2017, 37, 419-421.	3.8	7
153	Reticular dysgenesis: international survey on clinical presentation, transplantation, and outcome. Blood, 2017, 129, 2928-2938.	1.4	31
154	Inflammatory and autoimmune manifestations in X-linked carriers of chronic granulomatous disease in the United Kingdom. Journal of Allergy and Clinical Immunology, 2017, 140, 628-630.e6.	2.9	48
155	Approaches to the removal of T-lymphocytes to minimize graft-versus-host disease in patients with primary immunodeficiencies who do not have a matched sibling donor. Current Opinion in Allergy and Clinical Immunology, 2017, 17, 414-420.	2.3	11
156	Preclinical modeling highlights the therapeutic potential of hematopoietic stem cell gene editing for correction of SCID-X1. Science Translational Medicine, 2017, 9, .	12.4	176
157	X-Linked Agammaglobulinaemia: Outcomes in the modern era. Clinical Immunology, 2017, 183, 54-62.	3.2	72
158	Hematopoietic stem cell transplantation in 29 patients hemizygous for hypomorphic IKBKG/NEMO mutations. Blood, 2017, 130, 1456-1467.	1.4	95
159	Identification of Heterozygous Single- and Multi-exon Deletions in IL7R by Whole Exome Sequencing. Journal of Clinical Immunology, 2017, 37, 42-50.	3.8	20
160	Hematopoietic stem cell transplant in patients with activated PI3K delta syndrome. Journal of Allergy and Clinical Immunology, 2017, 139, 1046-1049.	2.9	90
161	A prospective study on the natural history of patients with profound combined immunodeficiency: An interim analysis. Journal of Allergy and Clinical Immunology, 2017, 139, 1302-1310.e4.	2.9	71
162	Long-term outcomes of 176 patients with X-linked hyper-IgM syndrome treated with or without hematopoietic cell transplantation. Journal of Allergy and Clinical Immunology, 2017, 139, 1282-1292.	2.9	107

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