Lisa J Kobrynski

List of Publications by Year
in descending order

34	1,767	623734 14	414414 32
papers	citations	h-index	g-index

36	36	36	2522
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Velocardiofacial syndrome, DiGeorge syndrome: the chromosome 22q11.2 deletion syndromes. Lancet, The, 2007, 370, 1443-1452.	13.7	513
2	Practice parameter for the diagnosis and management of primary immunodeficiency. Annals of Allergy, Asthma and Immunology, 2005, 94, S1-S63.	1.0	452
3	Prevalence and Morbidity of Primary Immunodeficiency Diseases, United States 2001â€“2007. Journal of Clinical Immunology, 2014, 34, 954-961.	3.8	115
4	Use of Genetic Testing for Primary Immunodeficiency Patients. Journal of Clinical Immunology, 2018, 38, 320-329.	3.8	88
5	Immunoglobulin Deficiencies: The B-Lymphocyte Side of DiGeorge Syndrome. Journal of Pediatrics, 2012, 161, 950-953.e1.	1.8	63
6	Cutting Edge: Antibody Production to Pneumococcal Polysaccharides Requires CD1 Molecules and CD8+T Cells. Journal of Immunology, 2005, 174, 1787-1790.	0.8	60
7	Subcutaneous immunoglobulin therapy: a new option for patients with primary immunodeficiency diseases. Biologics: Targets and Therapy, 2012, 6, 277.	3.2	56
8	Diagnostic interpretation of genetic studies in patients with primary immunodeficiency diseases: AÂworking group report of the Primary Immunodeficiency Diseases Committee of the American Academy of Allergy, Asthma & Immunology. Journal of Allergy and Clinical Immunology, 2020, 145, 46-69.	2.9	54
9	Exome Sequencing Identifies a Novel FOXP3 Mutation in a 2â€Generation Family With Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 561-568.	1.8	47
10	Applying public health strategies to primary immunodeficiency diseases: a potential approach to genetic disorders. MMWR Recommendations and Reports, 2004, 53, 1-29.	61.1	43
11	Cost-Effectiveness/Cost-Benefit Analysis of Newborn Screening for Severe Combined Immune Deficiency in Washington State. Journal of Pediatrics, 2016, 172, 127-135.	1.8	40
12	Infections in Infants with SCID: Isolation, Infection Screening, and Prophylaxis in PIDTC Centers. Journal of Clinical Immunology, 2021, 41, 38-50.	3.8	36
13	ICON: The Early Diagnosis of Congenital Immunodeficiencies. Journal of Clinical Immunology, 2014, 34, 398-424.	3.8	34
14	Primary immunodeficiency diseases: Practice among primary care providers and awareness among the general public, United States, 2008. Genetics in Medicine, 2010, 12, 792-800.	2.4	26
15	Rubella Virus Infected Macrophages and Neutrophils Define Patterns of Granulomatous Inflammation in Inborn and Acquired Errors of Immunity. Frontiers in Immunology, 2021, 12, 796065.	4.8	19
16	Hypogammaglobulinemia with decreased class-switched B-cells and dysregulated T-follicular-helper cells in IPEX syndrome. Clinical Immunology, 2018, 197, 219-223.	3.2	15
17	T-follicular helper cell expansion and chronic T-cell activation are characteristic immune anomalies in Evans syndrome. Blood, 2022, 139, 369-383.	1.4	14
18	Subcutaneous Immunoglobulin Replacement Therapy with HizentraÂ® is Safe and Effective in Children Less Than 5 Years of Age. Journal of Clinical Immunology, 2015, 35, 558-565.	3.8	13

#	Article	IF	Citations
19	Practical approach to genetic testing for primary immunodeficiencies. Annals of Allergy, Asthma and Immunology, 2019, 123, 433-439.	1.0	11
20	Newborn screening for severe combined immune deficiency (technical and political aspects). Current Opinion in Allergy and Clinical Immunology, 2015, 15, 539-546.	2.3	10
21	Severe Congenital Neutropenia associated with SRP54 mutation in 22q11.2 Deletion Syndrome: Hematopoietic Stem Cell Transplantation Results in Correction of Neutropenia with Adequate Immune Reconstitution. Journal of Clinical Immunology, 2018, 38, 546-549.	3.8	10
22	Identification of nonâ€“severe combined immune deficiency T-cell lymphopenia at newborn screening for severe combined immune deficiency. Annals of Allergy, Asthma and Immunology, 2019, 123, 424-427.	1.0	9
23	NAPDH Oxidase-Specific Flow Cytometry Allows for Rapid Genetic Triage and Classification of Novel Variants in Chronic Granulomatous Disease. Journal of Clinical Immunology, 2020, 40, 191-202.	3.8	8
24	Combined Immune Deficiencies in Children. Journal of Infusion Nursing, 2006, 29, 206-213.	2.3	5
25	Malakoplakia and Primary Immunodeficiency. Journal of Pediatrics, 2014, 165, 1053-1056.	1.8	5
26	A Pyrosequencing-Based Assay for the Rapid Detection of the 22q11.2 Deletion in DNA from Buccal and Dried Blood Spot Samples. Journal of Molecular Diagnostics, 2014, 16, 533-540.	2.8	5
27	MALDI-TOF-MS Assay to Detect the Hemizygous 22q11.2 Deletion in DNA from Dried Blood Spots. Clinical Chemistry, 2016, 62, 287-292.	3.2	5
28	Newborn Screening in the Diagnosis of Primary Immunodeficiency. Clinical Reviews in Allergy and Immunology, 2022, 63, 9-21.	6.5	5
29	Monoclonal IgA gammopathy due to maternal B cells in an infant with severe combined immunodeficiency (SCID) prior to hematopoietic stem cell transplantation. Journal of Pediatric Hematology/Oncology, 2006, 28, 53-6.	0.6	2
30	Management of ADA-Deficient SCID Patient on Adagen During Pregnancy. Journal of Clinical Immunology, 2019, 39, 846-848.	3.8	1
31	Primary Immunodeficiencies Presenting in Adolescence. , 2005, , 121-148.		1
32	HSCT using carrier donors for CD40L deficiency results in excellent immune function and higher CD40L expression in cTfh. Blood Advances, 2022, , .	5.2	1
33	Primary Immunodeficiency Disorders. Immunology and Allergy Clinics of North America, 2019, 39, xi-xii.	1.9	0
34	Primary immunodeficiencies presenting in adolescence. Adolescent Medicine: State of the Art Reviews, 2009, 20, 121-48, ix-x.	0.2	0