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	Newborn Screening in the Diagnosis of Primary Immunodeficiency. Clinical Reviews in Allergy and Immunology, 2022, 63, 9-21.	6.5	5
2	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
3	HSCT using carrier donors for CD40L deficiency results in excellent immune function and higher CD40L expression in cTfh. Blood Advances, 2022, , .	5.2	1
4	Infections in Infants with SCID: Isolation, Infection Screening, and Prophylaxis in PIDTC Centers. Journal of Clinical Immunology, 2021, 41, 38-50.	3.8	36
5	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
6	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
7	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
8	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
9	Management of ADA-Deficient SCID Patient on Adagen During Pregnancy. Journal of Clinical Immunology, 2019, 39, 846-848.	3.8	1
10	Practical approach to genetic testing for primary immunodeficiencies. Annals of Allergy, Asthma and Immunology, 2019, 123, 433-439.	1.0	11
11	Primary Immunodeficiency Disorders. Immunology and Allergy Clinics of North America, 2019, 39, xi-xii.	1.9	0
12	Use of Genetic Testing for Primary Immunodeficiency Patients. Journal of Clinical Immunology, 2018, 38, 320-329.	3.8	88
13	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
14	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
15	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
16	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
17	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
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	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
20	ICON: The Early Diagnosis of Congenital Immunodeficiencies. Journal of Clinical Immunology, 2014, 34, 398-424.	3.8	34
21	Prevalence and Morbidity of Primary Immunodeficiency Diseases, United States 2001â€“2007. Journal of Clinical Immunology, 2014, 34, 954-961.	3.8	115
22	Malakoplakia and Primary Immunodeficiency. Journal of Pediatrics, 2014, 165, 1053-1056.	1.8	5
23	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
24	Immunoglobulin Deficiencies: The B-Lymphocyte Side of DiGeorge Syndrome. Journal of Pediatrics, 2012, 161, 950-953.e1.	1.8	63
25	Subcutaneous immunoglobulin therapy: a new option for patients with primary immunodeficiency diseases. Biologics: Targets and Therapy, 2012, 6, 277.	3.2	56
26	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
27	Primary immunodeficiencies presenting in adolescence. Adolescent Medicine: State of the Art Reviews, 2009, 20, 121-48, ix-x.	0.2	0
28	Velocardiofacial syndrome, DiGeorge syndrome: the chromosome 22q11.2 deletion syndromes. Lancet, The, 2007, 370, 1443-1452.	13.7	513
29	Combined Immune Deficiencies in Children. Journal of Infusion Nursing, 2006, 29, 206-213.	2.3	5
30	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
31	Cutting Edge: Antibody Production to Pneumococcal Polysaccharides Requires CD1 Molecules and CD8+T Cells. Journal of Immunology, 2005, 174, 1787-1790.	0.8	60
32	Practice parameter for the diagnosis and management of primary immunodeficiency. Annals of Allergy, Asthma and Immunology, 2005, 94, S1-S63.	1.0	452
33	Primary Immunodeficiencies Presenting in Adolescence. , 2005, , 121-148.		1
34	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