Maja Agnieszka Klaudel-Dreszler

List of Publications by Year
in descending order

13 papers	598 citations	1040056 9 h-index	1058476 14 g-index
15 all docs	15 docs citations	15 times ranked	1549 citing authors

#	Article	IF	Citations
1	Heterozygous TBK1 mutations impair TLR3 immunity and underlie herpes simplex encephalitis of childhood. Journal of Experimental Medicine, 2012, 209, 1567-1582.	8.5	231
2	Human RIPK1 deficiency causes combined immunodeficiency and inflammatory bowel diseases. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 970-975.	7.1	130
3	Nijmegen Breakage Syndrome: Clinical and Immunological Features, Long-Term Outcome and Treatment Options â€“ a Retrospective Analysis. Journal of Clinical Immunology, 2015, 35, 538-549.	3.8	73
4	Disseminated Bacillus Calmette-GuÃ©rin Infection and Immunodeficiency. Emerging Infectious Diseases, 2007, 13, 799-801.	4.3	61
5	Giant Cell Hepatitis With Autoimmune Hemolytic Anemia in Children. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 669-673.	1.8	23
6	Common Variable Immune Deficiency in Childrenâ€”Clinical Characteristics Varies Depending on Defect in Peripheral B Cell Maturation. Journal of Clinical Immunology, 2013, 33, 731-741.	3.8	20
7	Early onset lysosomal acid lipase deficiency presenting as secondary hemophagocytic lymphohistiocytosis: Two infants treated with sebelipase alfa. Clinics and Research in Hepatology and Gastroenterology, 2018, 42, e77-e82.	1.5	16
8	Treosulfan-based conditioning regimen in a second matched unrelated peripheral blood stem cell transplantation for a pediatric patient with CGD and invasive aspergillosis, who experienced initial graft failure after RIC. International Journal of Hematology, 2009, 90, 571-575.	1.6	14
9	Pulmonary Lymphomatoid Granulomatosis in Griscelli Syndrome Type 2. Viral Immunology, 2011, 24, 471-473.	1.3	10
10	Severe congenital neutropeniaâ€associated JAGN1 mutations unleash a calpainâ€dependent cell death programme in myeloid cells. British Journal of Haematology, 2021, 192, 200-211.	2.5	7
11	Endocrine dysfunction in children with Shwachman-Diamond syndrome. Endokrynologia Polska, 2021, 72, 211-216.	1.0	6
12	Clinical immunology Disseminated Mycobacterium tuberculosis complex infection in a girl with partial dominant IFN-Î³ receptor 1 deficiency. Central-European Journal of Immunology, 2012, 4, 378-381.	1.2	4
13	Case Report: Liver as a Source of Hematopoietic Stem Cells After Liver Transplantation Following Hematopoietic Stem Cell Transplantation. Frontiers in Pediatrics, 2022, 10, 861692.	1.9	1