Elena Raikina

List of Publications by Year
in descending order

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39 papers	161 citations	1684188 5 h-index	1199594 12 g-index
39 all docs	39 docs citations	39 times ranked	270 citing authors

#	Article	IF	Citations
1	Cryopyrin-associated periodic syndrome assess the efficacy and safety of anakinra therapy: a single center experience. Pediatric Hematology/Oncology and Immunopathology, 2022, 21, 88-92.	0.3	2
2	Chromosomal aberrations as the cause of a complex phenotype in children with primary immunodeficiencies. Pediatric Hematology/Oncology and Immunopathology, 2021, 19, 62-67.	0.3	0
3	Non-infectious complications in the group of pediatric patients with chronic granulomatous disease. Pediatric Hematology/Oncology and Immunopathology, 2021, 19, 50-61.	0.3	0
4	Relative expansion of CD19â€negative veryâ€early normal Bâ€cell precursors in children with acute lymphoblastic leukaemia after CD19 targeting by blinatumomab and CARâ€T cell therapy: implications for flow cytometric detection of minimal residual disease. British Journal of Haematology, 2021, 193, 602-612.	2.5	30
5	CLINICAL CHARACTERISTICS OF A GROUP OF PATIENTS WITH CTLA4 HAPLOINSUFFICIENCY SYNDROME: EXPERIENCE OF ONE CENTER. Pediatriia, 2021, 100, 22-30.	0.2	0
6	Unique Combination of Diamondâ€“Blackfan Anemia and Lynch Syndrome in Adult Female: A Case Report. Frontiers in Oncology, 2021, 11, 652696.	2.8	1
7	DIFFICULTIES IN VERIFYING THE CAUSES OF IRON-DEFICIENCY ANAEMIA REFRACTORY TO ADEQUATELY ADMINISTERED THERAPY WITH IRON SUPPLEMENTS. Pediatriia, 2021, 100, 279-283.	0.2	0
8	Platelet phenotype in children with ANKRD26-related thrombocytopenia. Pediatric Hematology/Oncology and Immunopathology, 2021, 20, 65-73.	0.3	2
9	Evans syndrome in children: the results of a retrospective study of 54 patients. Pediatric Hematology/Oncology and Immunopathology, 2021, 20, 74-83.	0.3	0
10	Mevalonate kinase deficiency syndrome: Single center experience. Nauchno-Prakticheskaya Revmatologiya, 2021, 59, 326-334.	1.0	1
11	A case report of familial dyskeratosis congenital. Case report. Terapevticheskii Arkhiv, 2021, 93, 818-825.	0.8	0
12	Vemurafenib provides a rapid and robust clinical response in pediatric Langerhans cell histiocytosis with the BRAF V600E mutation but does not eliminate low-level minimal residual disease per ddPCR using cell-free circulating DNA. International Journal of Hematology, 2021, 114, 725-734.	1.6	17
13	Characteristics of a group of patients with WHIM syndrome. Pediatric Hematology/Oncology and Immunopathology, 2021, 19, 68-75.	0.3	1
14	The efficacy and safety of romiplostim in the treatment of thrombocytopenia in pediatric patients with Wiskottâ€“Aldrich syndrome: the results of a retrospective study. Pediatric Hematology/Oncology and Immunopathology, 2021, 19, 39-49.	0.3	0
15	Chimerism evaluation in measurable residual diseaseâ€suspected cells isolated by flow cell sorting as a reliable tool for measurable residual disease verification in acute leukemia patients after allogeneic hematopoietic stem cell transplantation. Cytometry Part B - Clinical Cytometry, 2021, 100, 568-573.	1.5	12
16	Hemorrhagic thrombocytopathy with defective signal transduction CalDAG-GEFI. Pediatric Hematology/Oncology and Immunopathology, 2021, 20, 126-131.	0.3	0
17	Analysis of familial cases of primary immunodeficiency in the context of genetic counseling. Pediatric Hematology/Oncology and Immunopathology, 2021, 20, 125-133.	0.3	0
18	Update on the genetic landscape of Diamondâ€Blackfan anemia in the Russian Federation. Pediatric Blood and Cancer, 2020, 67, e28314.	1.5	2

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19	The Clinical and Genetic Spectrum of 82 Patients With RAG Deficiency Including a c.256_257delAA Founder Variant in Slavic Countries. Frontiers in Immunology, 2020, 11, 900.	4.8	16
20	Development of flow cytometry assay for Wiskottâ€“Aldrich syndrome diagnosis by WASP protein evaluation. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 141-151.	0.3	3
21	Epsteinâ€“Barr virus-associated smooth muscle tumors in patients with primary immunodeficiencies. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 165-177.	0.3	0
22	Immune dysregulation symptoms as a rare manifestation of X-linked lymphoproliferative syndrome type 1. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 178-184.	0.3	0
23	Verification of X-linked lymphoproliferative syndrome type 1 and 2 using a flow cytometry method. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 108-118.	0.3	0
24	Genetic and Clinical Features of Shwachman-Diamond Syndrome in Russian Population: Prospective Study. Voprosy Sovremennoi Pediatrii - Current Pediatrics, 2020, 18, 393-400.	0.4	2
25	The spectrum of genetic variants of the a- and b-globin clusters in patients with hemoglobinopathies living in the Republic of Dagestan. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 50-53.	0.3	0
26	Diagnostic challenges in pyruvate kinase deficiency. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 90-94.	0.3	0
27	Federal clinical guidelines. Primary immunodeficiency: severe combined immunodeficiency. Russian Journal of Allergy, 2020, 17, 97-114.	0.2	0
28	Hematopoietic stem cell transplantation in a patient with type 1 mosaic variegated aneuploidy syndrome. Orphanet Journal of Rare Diseases, 2019, 14, 97.	2.7	5
29	Clericusio syndrome (poikiloderm with neutropenia). Pediatric Hematology/Oncology and Immunopathology, 2019, 18, 96-103.	0.3	2
30	The case of rare hereditary thrombocytopenia with a predisposition to the development of acute myeloid leukemia in twin children. Pediatric Hematology/Oncology and Immunopathology, 2019, 17, 51-56.	0.3	0
31	EXPERIENCE OF PRENATAL DIAGNOSIS OF PRIMARY IMMUNODEFICIENCY STATES. Pediatriia, 2019, 98, 44-48.	0.2	1
32	Clinical case of proteasome-associated autoinflammatory syndrome-2 (PRAAS2). Pediatric Hematology/Oncology and Immunopathology, 2019, 18, 108-113.	0.3	1
33	MALIGNANT RHABDOID KIDNEY TUMOR IN CHILDREN: RESULTS OF A SINGLE-CENTER STUDY. Pediatriia, 2019, 98, 40-48.	0.2	0
34	The phenomenon of reverse mutation in a patient with Wiskottâ€“Aldrich syndrome. Pediatric Hematology/Oncology and Immunopathology, 2019, 18, 104-111.	0.3	1
35	ANKRD26-related thrombocytopenia: case report and literature review of inherited thrombocytopenias with predisposition to malignancies. Pediatric Hematology/Oncology and Immunopathology, 2019, 18, 54-61.	0.3	0
36	The use of preventive pleurodesis in patient with Langerhans-cell histiocytosis with lung involvement: the case report and the review. Pediatric Hematology/Oncology and Immunopathology, 2018, 17, 74-80.	0.3	2

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37	Description of the familial case of Wiskottâ€“Aldrich syndrome with mild phenotype. Pediatric Hematology/Oncology and Immunopathology, 2018, 17, 97-102.	0.3	0
38	The Influence of Natural Killer Cell Alloreactivity on the Outcome of Î±/Î²TCR/CD19+ depleted Allogeneic Hematopoietic Stem Cell Transplantation in Pediatric Patients with Acute Leukemia. Pediatric Hematology/Oncology and Immunopathology, 2018, 17, .	0.3	1
39	Risk Factors for and the Clinical Impact of Cytomegalovirus and Epstein-Barr Virus Infections in Pediatric Recipients of TCR-Î±/Î²â€“ and CD19-Depleted Grafts. Biology of Blood and Marrow Transplantation, 2017, 23, 483-490.	2.0	59