

Elena Raikina

List of Publications by Year in descending order

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papers

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1684188

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270
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#	ARTICLE	IF	CITATIONS
1	Risk Factors for and the Clinical Impact of Cytomegalovirus and Epstein-Barr Virus Infections in Pediatric Recipients of TCR $\alpha\beta$ ^{hi} /I β ^{hi} and CD19-Depleted Grafts. <i>Biology of Blood and Marrow Transplantation</i> , 2017, 23, 483-490.	2.0	59
2	Relative expansion of CD19 ⁺ negative very early normal B ⁺ cell precursors in children with acute lymphoblastic leukaemia after CD19 targeting by blinatumomab and CAR ⁺ cell therapy: implications for flow cytometric detection of minimal residual disease. <i>British Journal of Haematology</i> , 2021, 193, 602-612.	2.5	30
3	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. <i>International Journal of Hematology</i> , 2021, 114, 725-734.	1.6	17
4	The Clinical and Genetic Spectrum of 82 Patients With RAG Deficiency Including a c.256_257delAA Founder Variant in Slavic Countries. <i>Frontiers in Immunology</i> , 2020, 11, 900.	4.8	16
5	Chimerism evaluation in measurable residual disease ⁺ 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. <i>Cytometry Part B - Clinical Cytometry</i> , 2021, 100, 568-573.	1.5	12
6	Hematopoietic stem cell transplantation in a patient with type 1 mosaic variegated aneuploidy syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 97.	2.7	5
7	Development of flow cytometry assay for Wiskott ⁺ Aldrich syndrome diagnosis by WASP protein evaluation. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 141-151.	0.3	3
8	Update on the genetic landscape of Diamond ⁺ Blackfan anemia in the Russian Federation. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28314.	1.5	2
9	Platelet phenotype in children with ANKRD26-related thrombocytopenia. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 20, 65-73.	0.3	2
10	Clericusio syndrome (poikiloderm with neutropenia). <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2019, 18, 96-103.	0.3	2
11	The use of preventive pleurodesis in patient with Langerhans-cell histiocytosis with lung involvement: the case report and the review. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2018, 17, 74-80.	0.3	2
12	Genetic and Clinical Features of Shwachman-Diamond Syndrome in Russian Population: Prospective Study. <i>Voprosy Sovremennoi Pediatrii - Current Pediatrics</i> , 2020, 18, 393-400.	0.4	2
13	Cryopyrin-associated periodic syndrome assess the efficacy and safety of anakinra therapy: a single center experience. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2022, 21, 88-92.	0.3	2
14	Unique Combination of Diamond ⁺ Blackfan Anemia and Lynch Syndrome in Adult Female: A Case Report. <i>Frontiers in Oncology</i> , 2021, 11, 652696.	2.8	1
15	Mevalonate kinase deficiency syndrome: Single center experience. <i>Nauchno-Prakticheskaya Revmatologiya</i> , 2021, 59, 326-334.	1.0	1
16	Characteristics of a group of patients with WHIM syndrome. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 19, 68-75.	0.3	1
17	The Influence of Natural Killer Cell Alloreactivity on the Outcome of $\alpha\beta$ TCR/CD19 ⁺ depleted Allogeneic Hematopoietic Stem Cell Transplantation in Pediatric Patients with Acute Leukemia. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2018, 17, .	0.3	1
18	EXPERIENCE OF PRENATAL DIAGNOSIS OF PRIMARY IMMUNODEFICIENCY STATES. <i>Pediatriia</i> , 2019, 98, 44-48.	0.2	1

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19	Clinical case of proteasome-associated autoinflammatory syndrome-2 (PRAAS2). <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2019, 18, 108-113.	0.3	1
20	The phenomenon of reverse mutation in a patient with Wiskottâ€Aldrich syndrome. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2019, 18, 104-111.	0.3	1
21	Chromosomal aberrations as the cause of a complex phenotype in children with primary immunodeficiencies. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 19, 62-67.	0.3	0
22	Non-infectious complications in the group of pediatric patients with chronic granulomatous disease. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 19, 50-61.	0.3	0
23	CLINICAL CHARACTERISTICS OF A GROUP OF PATIENTS WITH CTLA4 HAPLOINSUFFICIENCY SYNDROME: EXPERIENCE OF ONE CENTER. <i>Pediatriia</i> , 2021, 100, 22-30.	0.2	0
24	DIFFICULTIES IN VERIFYING THE CAUSES OF IRON-DEFICIENCY ANAEMIA REFRACTORY TO ADEQUATELY ADMINISTERED THERAPY WITH IRON SUPPLEMENTS. <i>Pediatriia</i> , 2021, 100, 279-283.	0.2	0
25	Evans syndrome in children: the results of a retrospective study of 54 patients. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 20, 74-83.	0.3	0
26	A case report of familial dyskeratosis congenital. Case report. <i>Terapevticheskii Arkhiv</i> , 2021, 93, 818-825.	0.8	0
27	The efficacy and safety of romiplostim in the treatment of thrombocytopenia in pediatric patients with Wiskottâ€Aldrich syndrome: the results of a retrospective study. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 19, 39-49.	0.3	0
28	Hemorrhagic thrombocytopathy with defective signal transduction CalDAG-GEFI. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 20, 126-131.	0.3	0
29	Description of the familial case of Wiskottâ€Aldrich syndrome with mild phenotype. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2018, 17, 97-102.	0.3	0
30	The case of rare hereditary thrombocytopenia with a predisposition to the development of acute myeloid leukemia in twin children. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2019, 17, 51-56.	0.3	0
31	MALIGNANT RHABDOID KIDNEY TUMOR IN CHILDREN: RESULTS OF A SINGLE-CENTER STUDY. <i>Pediatriia</i> , 2019, 98, 40-48.	0.2	0
32	ANKRD26-related thrombocytopenia: case report and literature review of inherited thrombocytopenias with predisposition to malignancies. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2019, 18, 54-61.	0.3	0
33	Epsteinâ€Barr virus-associated smooth muscle tumors in patients with primary immunodeficiencies. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 165-177.	0.3	0
34	Immune dysregulation symptoms as a rare manifestation of X-linked lymphoproliferative syndrome type 1. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 178-184.	0.3	0
35	Verification of X-linked lymphoproliferative syndrome type 1 and 2 using a flow cytometry method. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 108-118.	0.3	0
36	The spectrum of genetic variants of the a- and b-globin clusters in patients with hemoglobinopathies living in the Republic of Dagestan. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 50-53.	0.3	0

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37	Diagnostic challenges in pyruvate kinase deficiency. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2020, 19, 90-94.	0.3	0
38	Federal clinical guidelines. Primary immunodeficiency: severe combined immunodeficiency. <i>Russian Journal of Allergy</i> , 2020, 17, 97-114.	0.2	0
39	Analysis of familial cases of primary immunodeficiency in the context of genetic counseling. <i>Pediatric Hematology/Oncology and Immunopathology</i> , 2021, 20, 125-133.	0.3	0