

Nadezda Savenkova

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

68
citations

2258059

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1872680

6
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21
citing authors

#	ARTICLE	IF	CITATIONS
1	Development of hormone dependence in children with hormone-sensitive nephrotic syndrome at the onset of the disease. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2022, 66, 77-82.	0.3	1
2	Nephronophthisis due to mutation in the TMEM67 gene. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2022, 67, 121-126.	0.3	0
3	Prediction of cardiovascular complications and progression of renal failure in pediatric patients with chronic kidney disease according to NKF-K/DOQI (2002) and KDIGO (2012) classifications. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2022, 67, 12-19.	0.3	0
4	World Kidney Day: 2006–2022. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2022, 67, 153-157.	0.3	1
5	Hereditary Congenital and Infantile Nephrotic Syndrome in Children: Strategy of Management with New Possibilities for Genetic Diagnosis and Therapy. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2021, 65, 12-21.	0.3	1
6	Renal oligohydramnios and Potter sequence with cystic kidney disease. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2021, 66, 47-51.	0.3	4
7	Orpha disease – FRASER syndrome (ORPHA:2052) in children: phenotype and genotype characteristics. Nephrology (Saint-Petersburg), 2021, 25, 28-35.	0.4	0
8	Pediatric problems of stratification of the severity of stages, cardiovascular complications and renal forecast of chronic kidney disease by NKF-K / DOQI (2002) and KDIGO (2012) classifications. Nephrology (Saint-Petersburg), 2021, 25, 9-19.	0.4	2
9	Prenatal renal oligohydramnion and renal function in newborns and infants with cystic kidney diseases. Nephrology (Saint-Petersburg), 2021, 25, 68-74.	0.4	0
10	Orphanic hereditary hypophosphatemic rachit with hypercalciuria, nephrocalcinosis on account of mutation gene SLC34A3 (Review and case report). Nephrology (Saint-Petersburg), 2021, 25, 52-60.	0.4	0
11	Albert vazgenovich papayan – outstanding scientist, pediatrician, teacher (towards 85th anniversary). Pediatrician (St Petersburg), 2021, 12, 93-100.	0.3	0
12	Clinical phenotype, diagnostics, strategy of hypophosphatasia therapy due to <i>ALPL</i> gene mutations in pediatric and adult patients. Nephrology (Saint-Petersburg), 2021, 25, 16-26.	0.4	1
13	CHARGE syndrome. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2020, 65, 116-121.	0.3	3
14	Research of the level of erythropoietin and hypoxia-inducible factor 1-alpha in the blood of children and adolescents with anemia at stage C1–5 of chronic kidney disease. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2020, 65, 77-85.	0.3	2
15	The hereditary nephrotic syndrome in children and adults. Nephrology (Saint-Petersburg), 2020, 24, 15-27.	0.4	3
16	Feature of distribution of polymorphic options of genes associated with thrombophilia and arterial hypertension in children with hemolytic-uremic syndrome. Nephrology (Saint-Petersburg), 2020, 24, 90-94.	0.4	3
17	Professor Alexandra Antonovna Valentinovich (to the 110th anniversary of birth). Nephrology (Saint-Petersburg), 2020, 24, 105-106.	0.4	1
18	Clinico-genetic specifications of Bartter and Gitelman syndrome in children. Nephrology (Saint-Petersburg), 2020, 24, 42-53.	0.4	1

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19	Clinical and genetic features of glomerulocystic kidney in childhood. Nephrology (Saint-Petersburg), 2020, 24, 54-63.	0.4	0
20	Comparison of first choice cytostatic therapy with calcineurin inhibitors and nucleotides synthesis inhibitors in children with steroid-sensitive, steroid-dependent nephrotic syndrome with steroid toxicity. Nephrology (Saint-Petersburg), 2020, 24, 72-78.	0.4	3
21	Treatment of autosomal recessive and autosomal dominant polycystic kidney disease. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2019, 64, 22-29.	0.3	2
22	Clinical-genetic features and therapy strategy of hereditary congenital and infantile nephrotic syndrome in children (literature review). Nephrology (Saint-Petersburg), 2019, 23, 17-28.	0.4	3
23	Risk factors of acute cardiac surgery-associated kidney injury in newborns and infants with congenital heart defects. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2019, 64, 63-67.	0.3	0
24	Course of autosomal dominant and autosomal recessive polycystic kidney disease (ADPKD and ARPKD) wich detected in prenatal, neonatal and infant periods in children.. Nephrology (Saint-Petersburg), 2019, 23, 77-87.	0.4	2
25	Acute kidney injury in the newborn infant: classification, causes and epidemiology. Nephrology (Saint-Petersburg), 2019, 23, 9-16.	0.4	1
26	Treatment strategy ANCA-associated renal vasculitides in children and adolescents. Nephrology (Saint-Petersburg), 2019, 23, 107-115.	0.4	3
27	PEDIATRIC CLASSIFICATION AND EPIDEMIOLOGY OF ACUTE RENAL INJURY. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2018, 63, 36-42.	0.3	3
28	CLASSIFICATION PERFECTION OF ACUTE KIDNEY INJURY AND CHRONIC KIDNEY DIEASE IN ØEDIATRIC NEPHROLOGY. Nephrology (Saint-Petersburg), 2018, 22, 11-17.	0.4	3
29	Kidney cystosis inÂtuberous sclerosis inÂinfants. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2018, 63, 100-105.	0.3	0
30	The atypical form ofÂGitelman syndrome with cerebral calcifications. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2018, 63, 90-95.	0.3	1
31	MYH9-ASSOCIATED SYNDROME EPSTEIN: MACROTHROMBOCYTOPENIA, SENSORINEURAL HEARING LOSS, NEPHROPATHY IN CHILDREN (ACCORDING TO THE LITERATURE). Nephrology (Saint-Petersburg), 2018, 22, 88-94.	0.4	1
32	RENAL ARTERIAL HYPERTENSION IN CHILDREN AND ADOLESCENTS: CAUSES, CLASSIFICATION, DIAGNOSIS. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2017, 62, 43-48.	0.3	8
33	IRON STATUS, ERYTHROPOIETIN, HYPOXIA INDUCATOR FACTORS IN CHILDREN WITH ANEMIA WITH CHRONIC KIDNEY DISEASE. Nephrology (Saint-Petersburg), 2017, 21, 68-77.	0.4	2
34	ACUTE KIDNEY INJURY IN NEONATES AND INFANTS WITH CONGENITAL HEART DISORDERS AFTER CARDIAC SURGERY. Nephrology (Saint-Petersburg), 2017, 21, 54-60.	0.4	5
35	CAKUT-SYNDROME IN THE ETIOLOGICAL STRUCTURE OF CHRONIC KIDNEY DISEASE IN CHILDREN AND ADOLESCENTS. Nephrology (Saint-Petersburg), 2017, 21, 69-74.	0.4	8