Victoria Y Voinova

List of Publications by Year
in descending order

20	158	1307594 7	1199594 12
papers	citations	h-index	g-index

20	20	20	224
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Cytogenetic, Molecular-Cytogenetic, and Clinical-Genealogical Studies of the Mothers of Children with Autism: A Search for Familial Genetic Markers for Autistic Disorders. Neuroscience and Behavioral Physiology, 2010, 40, 745-756.	0.4	28
2	3p22.1p21.31 microdeletion identifies CCK as Asperger syndrome candidate gene and shows the way for therapeutic strategies in chromosome imbalances. Molecular Cytogenetics, 2015, 8, 82.	0.9	27
3	Mutations of the Transcriptional Corepressor ZMYM2 Cause Syndromic Urinary Tract Malformations. American Journal of Human Genetics, 2020, 107, 727-742.	6.2	25
4	Xq28 (MECP2) microdeletions are common in mutation-negative females with Rett syndrome and cause mild subtypes of the disease. Molecular Cytogenetics, 2013, 6, 53.	0.9	24
5	Immune tolerance induction for laronidase treatment in mucopolysaccharidosis I. Molecular Genetics and Metabolism Reports, 2017, 10, 61-66.	1.1	12
6	Dental manifestations of hypophosphatasia in children and the effects of enzyme replacement therapy on dental status: A series of clinical cases. Clinical Case Reports (discontinued), 2020, 8, 911-918.	0.5	10
7	4q21.2q21.3 Duplication: Molecular and Neuropsychological Aspects. Current Genomics, 2018, 19, 173-178.	1.6	10
8	40-Hz Auditory Steady-State Response (ASSR) as a Biomarker of Genetic Defects in the SHANK3 Gene: A Case Report of 15-Year-Old Girl with a Rare Partial SHANK3 Duplication. International Journal of Molecular Sciences, 2021, 22, 1898.	4.1	8
9	High-performance DNA sequencing to identify genetically determined diseases in pediatric practice. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2019, 64, 103-109.	0.3	4
10	GENETICS AND THE HEART: THE BASIS FOR INTRODUCING GENETIC TESTING INTO CLINICAL PRACTICE. Pediatriia, 2020, 99, 8-15.	0.2	3
11	The relationship of bone mineral density with the of the intima-media thickness in premenopausal women. Osteoporosis and Bone Diseases, 2021, 23, 13-18.	1.4	2
12	Replicative cell ageing: the role of insulin resistance in patients with arterial hypertension. Arterial Hypertension (Russian Federation), 2019, 25, 225-231.	0.4	2
13	Phenotypic variability and modifier variants in children with hereditary heart diseases. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2021, 66, 12-19.	0.3	1
14	Rett syndrome in Russia and abroad: a scientific historical review. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2020, 65, 25-31.	0.3	1
15	CLINICAL MANIFESTATIONS AND STRUCTURE OF CHROMOSOMAL PATHOLOGY IN CHILDREN WITH IMPAIRED COGNITIVE DEVELOPMENT IN A SPECIALIZED GENETIC CLINIC. Pediatriia, 2020, 99, 102-108.	0.2	1
16	A RARE FORM OF HYPOPHOSPHATEMIC RICKETS DUE TO MUTATIONS IN THE SLC34A3 GENE: DIAGNOSIS AND TREATMENT PECULIARITIES. Pediatriia, 2018, 97, 69-73.	0.2	0
17	ANOMALY OF THE URINARY SYSTEM IN A CHILD WITH RARE MONOGENIC STAR SYNDROME. Pediatriia, 2019, 98, 263-268.	0.2	0
18	Biomarkers of early cardiovascular aging. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2019, 64, 11-18.	0.3	0

#	Article	IF	Citations
19	USE OF THE NEXT -GENERATION SEQUENCING TECHNOLOGY FOR HEREDITARY DISEASES IN CHILDREN DIAGNOSTICS IN A SPECIALIZED CLINIC. Pediatriia, 2019, 98, 74-78.	0.2	0
20	New genome editing technologies in the treatment of X-linked adrenoleukodystrophy. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2020, 65, 104-107.	0.3	0