

# Atsuko Noguchi

## List of Publications by Year in descending order

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Version: 2024-02-01

15  
papers

245  
citations

1040056

9  
h-index

1199594

12  
g-index

15  
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15  
docs citations

15  
times ranked

326  
citing authors

#	ARTICLE	IF	CITATIONS
1	Diagnosis of SLC25A46-related pontocerebellar hypoplasia in two siblings with fulminant neonatal course: role of postmortem CT and whole genomic analysis: a case report. <i>BMC Neurology</i> , 2022, 22, 20.	1.8	0
2	Precocious puberty in a case of Simpsonâ€“Golabiâ€“Behmel syndrome with a de novo 240-kb deletion including GPC3. <i>Human Genome Variation</i> , 2022, 9, .	0.7	0
3	Phenotypic characteristics of pediatric inflammatory bowel disease in Japan: results from a multicenter registry. <i>Intestinal Research</i> , 2020, 18, 412-420.	2.6	20
4	Detection in Japan of an equine-like G3P[8] reassortant rotavirus A strain that is highly homologous to European strains across all genome segments. <i>Archives of Virology</i> , 2018, 163, 791-794.	2.1	21
5	Barth Syndrome: Different Approaches to Diagnosis. <i>Journal of Pediatrics</i> , 2018, 193, 256-260.	1.8	14
6	Familial episodic limb pain in kindreds with novel Nav1.9 mutations. <i>PLoS ONE</i> , 2018, 13, e0208516.	2.5	12
7	Biopsyâ€“proven acute interstitial nephritis due to fosfomycin in a child. <i>Nephrology</i> , 2018, 23, 890-890.	1.6	2
8	Regional Variations in the Incidence of Rotavirus Hospitalization in Children Living in Defined Regions of Akita and Kyoto Prefectures, Japan. <i>Japanese Journal of Infectious Diseases</i> , 2017, 70, 167-170.	1.2	1
9	Infantile Pain Episodes Associated with Novel Nav1.9 Mutations in Familial Episodic Pain Syndrome in Japanese Families. <i>PLoS ONE</i> , 2016, 11, e0154827.	2.5	38
10	Spread and predominance in Japan of novel G1P[8] double-reassortant rotavirus strains possessing a DS-1-like genotype constellation typical of G2P[4] strains. <i>Infection, Genetics and Evolution</i> , 2014, 28, 426-433.	2.3	59
11	Stroke-like Episode Involving a Cerebral Artery in a Patient With MELAS. <i>Pediatric Neurology</i> , 2005, 33, 70-71.	2.1	19
12	Psychosocial Dwarfism: A Case Report. <i>Clinical Pediatric Endocrinology</i> , 2005, 14, S24_43-S24_46.	0.8	0
13	SLC7A7 genomic structure and novel variants in three Japanese lysinuric protein intolerance families. <i>Human Mutation</i> , 2000, 15, 367-372.	2.5	29
14	A cluster of lysinuric protein intolerance (LPI) patients in a northern part of Iwate, Japan due to a founder effect. <i>Human Mutation</i> , 2000, 16, 270-271.	2.5	27
15	Chylothorax in a polysplenia infant with cor triatriatum, pulmonary stenosis and sick sinus syndrome. <i>Heart and Vessels</i> , 1997, 12, 247-249.	1.2	3