Atsuko Noguchi

List of Publications by Year
in descending order

15	245	1040056 9	1199594 12
papers	citations	h-index	g-index

15	15	15	326
all docs	docs citations	times ranked	citing authors

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