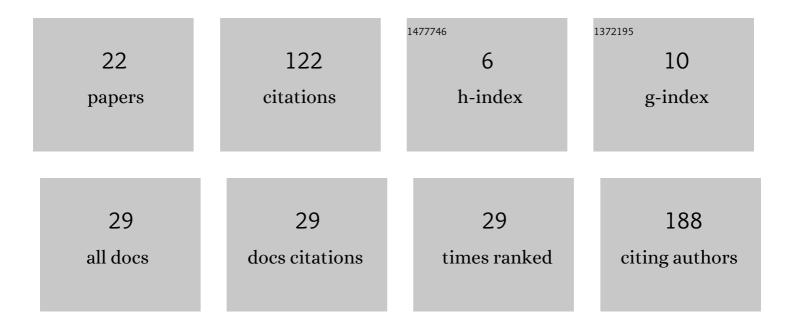
## Jong-Hee Chae

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6779811/publications.pdf Version: 2024-02-01



#	ARTICLE	IF	CITATION
1	Diagnostic Challenges Associated with GLUT1 Deficiency: Phenotypic Variabilities and Evolving Clinical Features. Yonsei Medical Journal, 2019, 60, 1209.	0.9	16
2	The Korean undiagnosed diseases program: lessons from a one-year pilot project. Orphanet Journal of Rare Diseases, 2019, 14, 68.	1.2	14
3	Urological Problems in Patients with Menkes Disease. Journal of Korean Medical Science, 2019, 34, e4.	1.1	13
4	Spectrum of movement disorders in GNAO1 encephalopathy: in-depth phenotyping and case-by-case analysis. Orphanet Journal of Rare Diseases, 2020, 15, 343.	1.2	12
5	Association Analysis of Interleukin-1β, Interleukin-6, and HMGB1 Variants with Postictal Serum Cytokine Levels in Children with Febrile Seizure and Generalized Epilepsy with Febrile Seizure Plus. Journal of		

Jong-Hee Chae

#	Article	IF	CITATIONS
19	Variable Phenotypes of ZC4H2-Associated Rare Disease in Six Patients. Annals of Child Neurology, 2022, 30, 120-126.	0.0	1
20	Clinical and Genetic Spectrum of STXBP1 Encephalopathy in the Korean Pediatric Population. Annals of Child Neurology, 2021, 29, 68-74.	0.0	0
21	Heterogeneous Clinical Characteristics of Allan-Herndon-Dudley Syndrome with SLC16A2 Mutations. Annals of Child Neurology, 2021, 29, 149-158.	0.0	0
22	Clinical Characteristics and Neurologic Outcomes of X-Linked Myotubular Myopathy. Annals of Child Neurology, 0, , .	0.0	0