Nebal Waill Saadi

List of Publications by Year
in descending order

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6 papers	46 citations	2258059 3 h-index	2550090 3 g-index
6 all docs	6 docs citations	6 times ranked	150 citing authors

#	Article	IF	Citations
1	HTRA2 Defect: A Recognizable Inborn Error of Metabolism with 3-Methylglutaconic Aciduria as Discriminating Feature Characterized by Neonatal Movement Disorder and Epilepsyâ€"Report of 11 Patients. Neuropediatrics, 2018, 49, 373-378.	0.6	21
2	A novel homozygous <scp>SLC13A5</scp> wholeâ€gene deletion generated by <scp>Alu/Alu</scp>â€mediated rearrangement in an Iraqi family with epileptic encephalopathy. American Journal of Medical Genetics, Part A, 2021, 185, 1972-1980.	1.2	16
3	Haploinsufficiency of ARFGEF1 is associated with developmental delay, intellectual disability, and epilepsy with variable expressivity. Genetics in Medicine, 2021, 23, 1901-1911.	2.4	9
4	13-year diagnostic delay as cerebral palsy of an Iraqi patient with NBIA type 4. Neurology: Clinical Practice, 2019, 9, e22-e24.	1.6	0
5	Pitfalls in the diagnosis of Gaucher disease in Iraq:ÂAÂdiagnostic experience from a developing country. Pakistan Journal of Medical Sciences, 2021, 37, 782-787.	0.6	0
6	Molecular Genetic Testing in Pediatric and Adult Neurology in Iraq: New Experience and Challenges from a Developing Country. Journal of Pediatric Neurology, 2021, 19, 395-401.	0.2	0