Susan Akbaroghli

List of Publications by Year in descending order

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2258059 2053705 6 43 3 5 citations h-index g-index papers 6 6 6 128 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Next-generation sequencing identified novel truncating mutations in BBS9 causing Bardet Biedl syndrome in two Iranian consanguineous families Iranian Journal of Child Neurology, 2022, 16, 123-133.	0.3	O
2	Investigation of Chromosomal Abnormalities and Microdeletion/ Microduplication(s) in Fifty Iranian Patients with Multiple Congenital Anomalies. Cell Journal, 2019, 21, 337-349.	0.2	1
3	Van Maldergem syndrome and Hennekam syndrome: Further delineation of allelic phenotypes. American Journal of Medical Genetics, Part A, 2018, 176, 1166-1174.	1.2	14
4	Discriminative Features in Three Autosomal Recessive Cutis Laxa Syndromes: Cutis Laxa IIA, Cutis Laxa IIB, and Geroderma Osteoplastica. International Journal of Molecular Sciences, 2017, 18, 635.	4.1	18
5	Identification of a new mutation in an Iranian family with hereditary multiple osteochondromas. Therapeutics and Clinical Risk Management, 2016, Volume 13, 15-19.	2.0	1
6	A Case Series: Congenital Hyperinsulinism. International Journal of Endocrinology and Metabolism, 2016, 14, e37311.	1.0	9