Susan Akbaroghli

List of Publications by Year
in descending order

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

#	Article	IF	Citations
1	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
2	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
3	A Case Series: Congenital Hyperinsulinism. International Journal of Endocrinology and Metabolism, 2016, 14, e37311.	1.0	9
4	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
5	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
6	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	0