Cobblestone Lissencephaly in Schinzel-Giedion Syndro

Journal of Child Neurology
28, 259-263
DOI: 10.1177/0883073812441250

Citation Report

#	Article	IF	Citations
1	Transformation of Merkel cell carcinoma to ganglioneuroblastoma in intracranial metastasis. Human Pathology, 2014, 45, 1978-1981.	1.1	5
2	Congenital generalized hypertrichosis: the skin as a clue to complex malformation syndromes. Italian Journal of Pediatrics, 2015, 41, 55.	1.0	38
4	Schinzelâ€“Giedion syndrome in two Brazilian patients: Report of a novel mutation in SETBP1 and literature review of the clinical features. American Journal of Medical Genetics, Part A, 2015, 167, 1039-1046.	0.7	25
5	Schinzel-Giedion Syndrome with Congenital Megacalycosis in a Turkish Patient: Report of SETBP1 Mutation and Literature Review of the Clinical Features. Case Reports in Genetics, 2017, 2017, 1-4.	0.1	0
6	Schinzelâ€“Giedion syndrome: a novel case, review and revised diagnostic criteria. Journal of Genetics, 2018, 97, 35-46.	0.4	18
7	Targeted deletion of RIC8A in mouse neural precursor cells interferes with the development of the brain, eyes, and muscles. Developmental Neurobiology, 2018, 78, 374-390.	1.5	0
8	SETBP1 induces transcription of a network of development genes by acting as an epigenetic hub. Nature Communications, 2018, 9, 2192.	5.8	66
9	Schinzel-Giedion syndrome: a novel case, review and revised diagnostic criteria. Journal of Genetics, 2018, 97, 35-46.	0.4	2