Susanne Gerit Kircher

List of Publications by Year in descending order

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1937685 1720034 12 61 4 7 citations g-index h-index papers 12 12 12 100 docs citations times ranked citing authors all docs

#	Article	lF	CITATIONS
1	Infantile systemic hyalinosis: Variable grades of severity. African Journal of Paediatric Surgery, 2021, 18, 224-230.	0.6	0
2	Skeletal phenotype/genotype in progressive pseudorheumatoid chondrodysplasia. Clinical Rheumatology, 2020, 39, 553-560.	2.2	7
3	Progressive Deformity of the Lower Limbs in a Patient with KID (Keratitis-Ichthyosis-Deafness) Syndrome. Case Reports in Orthopedics, 2020, 2020, 1-5.	0.3	1
4	Tomographic Study of the Malformation Complex in Correlation With the Genotype in Patients With Robinow Syndrome: Review Article. Journal of Investigative Medicine High Impact Case Reports, 2020, 8, 232470962091177.	0.6	4
5	The articular and the craniocervical abnormalities are of confusing age of onset in patients with Maroteaux-Lamy disease (MPS VI). Minerva Pediatrics, 2020, , .	0.4	0
6	The Managment of cervical spine abnormalities in children with spondyloepiphyseal dysplasia congenita. Medicine (United States), 2019, 98, e13780.	1.0	17
7	Leri-Weill Dyschondrosteosis Syndrome: Analysis via 3DCT Scan. Medicines (Basel, Switzerland), 2019, 6, 60.	1.4	0
8	Massive Axial and Appendicular Skeletal Deformities in Connection with Gorham-Stout Syndrome. Medicines (Basel, Switzerland), 2019, 6, 54.	1.4	1
9	A constellation of orthopaedic deformities in connection with cartilage oligomeric matrix protein mutation. African Journal of Paediatric Surgery, 2019, 16, 23.	0.6	1
10	Muscle Weakness. Journal of Investigative Medicine High Impact Case Reports, 2017, 5, 232470961668958.	0.6	3
11	The Diversity of the Clinical Phenotypes in Patients With Fibrodysplasia Ossificans Progressiva. Journal of Clinical Medicine Research, 2016, 8, 246-253.	1.2	22
12	Can Multiple Hereditary Exostoses Overlap With Mesomelic Dysplasia?. Journal of Clinical Medicine Research, 2016, 8, 605-609.	1,2	5