

# Susanne Gerit Kircher

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2455586/publications.pdf>

Version: 2024-02-01

12  
papers

61  
citations

1937685

4  
h-index

1720034

7  
g-index

12  
all docs

12  
docs citations

12  
times ranked

100  
citing authors

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