

Gabriel T Mindler

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6478811/publications.pdf>

Version: 2024-02-01

9
papers

132
citations

1684188

5
h-index

1588992

8
g-index

11
all docs

11
docs citations

11
times ranked

98
citing authors

#	ARTICLE	IF	CITATIONS
1	Interdisciplinary management of FGF23-related phosphate wasting syndromes: a Consensus Statement on the evaluation, diagnosis and care of patients with X-linked hypophosphataemia. <i>Nature Reviews Endocrinology</i> , 2022, 18, 366-384.	9.6	42
2	Multidisciplinary patient care in X-linked hypophosphatemic rickets: one challenge, many perspectives. <i>Wiener Medizinische Wochenschrift</i> , 2020, 170, 116-123.	1.1	32
3	Disease-specific gait deviations in pediatric patients with X-linked hypophosphatemia. <i>Gait and Posture</i> , 2020, 81, 78-84.	1.4	17
4	Lower Limb Deformity and Gait Deviations Among Adolescents and Adults With X-Linked Hypophosphatemia. <i>Frontiers in Endocrinology</i> , 2021, 12, 754084.	3.5	15
5	The unstable knee in congenital limb deficiency. <i>Journal of Children's Orthopaedics</i> , 2016, 10, 521-528.	1.1	10
6	Persistent Lower Limb Deformities Despite Amelioration of Rickets in X-Linked Hypophosphatemia (XLH) - A Prospective Observational Study. <i>Frontiers in Endocrinology</i> , 2022, 13, 866170.	3.5	10
7	Leg lengthening and deformity correction in rare bone diseases: a multidisciplinary approach. <i>Wiener Medizinische Wochenschrift</i> , 2021, 171, 126-132.	1.1	4
8	Expanding the Phenotype of the FAM149B1-Related Ciliopathy and Identification of Three Neurogenetic Disorders in a Single Family. <i>Genes</i> , 2021, 12, 1648.	2.4	2
9	Clinical Phenotype and Bone Biopsy Characteristics in a Child with Proteus Syndrome. <i>Calcified Tissue International</i> , 2021, 109, 586-595.	3.1	0