

Muriel de la Dure Molla

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8883856/publications.pdf>

Version: 2024-02-01

12
papers

557
citations

933447

10
h-index

1281871

11
g-index

12
all docs

12
docs citations

12
times ranked

739
citing authors

#	ARTICLE	IF	CITATIONS
1	Enamel Defects Reflect Perinatal Exposure to Bisphenol A. American Journal of Pathology, 2013, 183, 108-118.	3.8	106
2	A targeted next-generation sequencing assay for the molecular diagnosis of genetic disorders with orodental involvement. Journal of Medical Genetics, 2016, 53, 98-110.	3.2	100
3	Isolated dentinogenesis imperfecta and dentin dysplasia: revision of the classification. European Journal of Human Genetics, 2015, 23, 445-451.	2.8	90
4	Nephrocalcinosis (Enamel Renal Syndrome) Caused by Autosomal Recessive FAM20A Mutations. Nephron Physiology, 2013, 122, 1-6.	1.2	84
5	Pathognomonic oral profile of Enamel Renal Syndrome (ERS) caused by recessive FAM20A mutations. Orphanet Journal of Rare Diseases, 2014, 9, 84.	2.7	63
6	Elements of morphology: Standard terminology for the teeth and classifying genetic dental disorders. American Journal of Medical Genetics, Part A, 2019, 179, 1913-1981.	1.2	41
7	Tracking Endogenous Amelogenin and Ameloblastin In Vivo. PLoS ONE, 2014, 9, e99626.	2.5	23
8	Oral health related quality of life of children and adolescents affected by rare orofacial diseases: a questionnaire-based cohort study. Orphanet Journal of Rare Diseases, 2019, 14, 124.	2.7	16
9	Amelogenesis imperfecta: therapeutic strategy from primary to permanent dentition across case reports. BMC Oral Health, 2018, 18, 108.	2.3	15
10	Translation and cross-cultural validation of the French version of the Sleep-Related Breathing Disorder scale of the Pediatric Sleep Questionnaire. Sleep Medicine, 2019, 58, 123-129.	1.6	14
11	Gingival inflammation, enamel defects, and tooth sensitivity in children with amelogenesis imperfecta: a case-control study. Journal of Applied Oral Science, 2020, 28, e20200170.	1.8	5
12	Oral Phenotype of Singletonâ€™Merten Syndrome: A Systematic Review Illustrated With a Case Report. Frontiers in Genetics, 0, 13, .	2.3	0