Muriel de la Dure Molla

List of Publications by Year
in descending order

12	557	933447 10	1281871 11
papers	citations	h-index	g-index

12	12	12	739
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	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
3	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
4	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
5	Amelogenesis imperfecta: therapeutic strategy from primary to permanent dentition across case reports. BMC Oral Health, 2018, 18, 108.	2.3	15
6	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
7	Isolated dentinogenesis imperfecta and dentin dysplasia: revision of the classification. European Journal of Human Genetics, 2015, 23, 445-451.	2.8	90
8	Pathognomonic oral profile of Enamel Renal Syndrome (ERS) caused by recessive FAM20A mutations. Orphanet Journal of Rare Diseases, 2014, 9, 84.	2.7	63
9	Tracking Endogenous Amelogenin and Ameloblastin In Vivo. PLoS ONE, 2014, 9, e99626.	2.5	23
10	Nephrocalcinosis (Enamel Renal Syndrome) Caused by Autosomal Recessive FAM20A Mutations. Nephron Physiology, 2013, 122, 1-6.	1.2	84
11	Enamel Defects Reflect Perinatal Exposure to Bisphenol A. American Journal of Pathology, 2013, 183, 108-118.	3.8	106
12	Oral Phenotype of Singletonâ€“Merten Syndrome: A Systematic Review Illustrated With a Case Report. Frontiers in Genetics, 0, 13, .	2.3	0