Miriam Erandi Reyna-FabiÃ¡n

List of Publications by Year
in descending order

14 papers	188 citations	1307594 7 h-index	1058476 14 g-index
15 all docs	15 docs citations	15 times ranked	318 citing authors

#	Article	IF	Citations
1	Detecting a Complex of Cryptic Species within Neoechinorhynchus golvani (Acanthocephala:) Tj ETQq1 1 0.784314 rgBT /Overlock 10 T 2009, 95, 1040-1047.	0.7	63
2	Analysis of the Bacterial Diversity in Liver Abscess: Differences Between Pyogenic and Amebic Abscesses. American Journal of Tropical Medicine and Hygiene, 2016, 94, 147-155.	1.4	21
3	5,10â€Methylenetetrahydrofolate reductase single nucleotide polymorphisms and geneâ€“environment interaction analysis in nonâ€syndromic cleft lip/palate. European Journal of Oral Sciences, 2014, 122, 109-113.	1.5	17
4	First comprehensive TSC1/TSC2 mutational analysis in Mexican patients with Tuberous Sclerosis Complex reveals numerous novel pathogenic variants. Scientific Reports, 2020, 10, 6589.	3.3	16
5	Association of interactions among the IRF6 gene, the 8q24 region, and maternal folic acid intake with nonâ€syndromic cleft lip/palate in Mexican Mestizos. American Journal of Medical Genetics, Part A, 2012, 158A, 3207-3210.	1.2	15
6	Gene Interactions Provide Evidence for Signaling Pathways Involved in Cleft Lip/Palate in Humans. Journal of Dental Research, 2016, 95, 1257-1264.	5.2	14
7	Germline Mutations in NKX2-5, GATA4, and CRELD1 are Rare in a Mexican Sample of Down Syndrome Patients with Endocardial Cushion and Septal Heart Defects. Pediatric Cardiology, 2015, 36, 802-808.	1.3	9
8	Validating the systematic position of Plationus Segers, Murugan & Dumont, 1993 (Rotifera:) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 C oxidase. Hydrobiologia, 2010, 644, 361-370.	2.0	7
9	Deletion of Exon 1 of the SLC16A2 Gene: A Common Occurrence in Patients with Allan-Herndon-Dudley Syndrome. Thyroid, 2015, 25, 361-367.	4.5	7
10	Predominance of Dystrophinopathy Genotypes in Mexican Male Patients Presenting as Muscular Dystrophy with A Normal Multiplex Polymerase Chain Reaction DMD Gene Result: A Study Including Targeted Next-Generation Sequencing. Genes, 2019, 10, 856.	2.4	6
11	TSC2/PKD1 contiguous gene syndrome, with emphasis on a case with an atypical mild polycystic kidney phenotype and a novel genetic variant. Nefrologia, 2020, 40, 91-98.	0.4	6
12	Screening of IRF6 Variants in Patients Subjected to Genetic Association Studies for Nonsyndromic Cleft Lip/Palate. Cleft Palate-Craniofacial Journal, 2021, 58, 1128-1134.	0.9	3
13	TSC2/PKD1 contiguous gene syndrome, with emphasis on a case with an atypical mild polycystic kidney phenotype and a novel genetic variant. Nefrologia, 2020, 40, 91-98.	0.4	2
14	Unusual Clinical Manifestations in a Mexican Patient with Sanfilippo B Syndrome. Diagnostics, 2022, 12, 1268.	2.6	1