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List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5977735/publications.pdf>

Version: 2024-02-01

13
papers

159
citations

1163117

8
h-index

1199594

12
g-index

13
all docs

13
docs citations

13
times ranked

269
citing authors

#	ARTICLE	IF	CITATIONS
1	Non lethal Raine syndrome and differential diagnosis. European Journal of Medical Genetics, 2016, 59, 577-583.	1.3	33
2	High frequency of the recurrent c.1310_1313delAAGA BRCA2 mutation in the North-East of Morocco and implication for hereditary breast-ovarian cancer prevention and control. BMC Research Notes, 2017, 10, 188.	1.4	24
3	Distribution of allelic and genotypic frequencies of NAT2 and CYP2E1 variants in Moroccan population. BMC Genetics, 2014, 15, 156.	2.7	23
4	First application of next-generation sequencing in Moroccan breast/ovarian cancer families and report of a novel frameshift mutation of the BRCA1 gene. Oncology Letters, 2016, 12, 1192-1196.	1.8	18
5	Natural history of non-lethal Raine syndrome during childhood. Orphanet Journal of Rare Diseases, 2020, 15, 93.	2.7	17
6	Novel <i>DDR2</i> mutation identified by whole exome sequencing in a Moroccan patient with spondylo-epiphyseal dysplasia, short limb abnormal calcification type. American Journal of Medical Genetics, Part A, 2016, 170, 460-465.	1.2	12
7	Germinal mosaicism in Noonan syndrome: A family with two affected siblings of normal parents. American Journal of Medical Genetics, Part A, 2010, 152A, 2850-2853.	1.2	9
8	Low Level of Consanguinity in Moroccan Families at High Risk of Breast Cancer. Asian Pacific Journal of Cancer Prevention, 2013, 14, 723-726.	1.2	8
9	Severe form of hypoglossia-hypodactylia syndrome associated with complex cardiopathy: A case report. International Journal of Pediatric Otorhinolaryngology, 2010, 74, 1092-1094.	1.0	7
10	A recurrent mutation in Moroccan patients with Dyggve-Melchior-Clausen syndrome: Report of a new case and review. Indian Journal of Human Genetics, 2011, 17, 97.	0.7	5
11	Mutational spectrum of BRCA1/2 genes in Moroccan patients with hereditary breast and/or ovarian cancer, and review of BRCA mutations in the MENA region. Breast Cancer Research and Treatment, 2022, 194, 187-198.	2.5	2
12	Characterization of a rare short arm heteromorphism of chromosome 22 in a girl with down-syndrome like facies. Indian Journal of Human Genetics, 2014, 20, 89.	0.7	1
13	Microdélétion 22q11.2 rÃ©vÃ©lÃ©e par une hypocalcÃ©mie nÃ©onatale avec une dysmorphie faciale mineure. Revue Francophone Des Laboratoires, 2009, 2009, 83-85.	0.0	0