siham Chafai Elalaoui

List of Publications by Year
in descending order

13	159	1163117 8	1199594 12
papers	citations	h-index	g-index

13	13	13	269
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	Natural history of non-lethal Raine syndrome during childhood. Orphanet Journal of Rare Diseases, 2020, 15, 93.	2.7	17
3	High frequency of the recurrent c.1310_1313delAAGA BRCA2 mutation in the North-East of Morocco and implication for hereditary breastâ€“ovarian cancer prevention and control. BMC Research Notes, 2017, 10, 188.	1.4	24
4	Non lethal Raine syndrome and differential diagnosis. European Journal of Medical Genetics, 2016, 59, 577-583.	1.3	33
5	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
6	Novel DDR2 mutation identified by whole exome sequencing in a Moroccan patient with spondyloâ€metaâ€epiphyseal dysplasia, short limbâ€abnormal calcification type. American Journal of Medical Genetics, Part A, 2016, 170, 460-465.	1.2	12
7	Distribution of allelic and genotypic frequencies of NAT2 and CYP2E1variants in Moroccan population. BMC Genetics, 2014, 15, 156.	2.7	23
8	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
9	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
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	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
12	Severe form of hypoglossiaâ€“hypodactylia syndrome associated with complex cardiopathy: A case report. International Journal of Pediatric Otorhinolaryngology, 2010, 74, 1092-1094.	1.0	7
13	MicrodÃ©lÃ©tion 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