Chantal Farra

List of Publications by Year
in descending order

25 papers	266 citations	1163117 8 h-index	940533 16 g-index
26 all docs	26 docs citations	26 times ranked	765 citing authors

#	Article	IF	Citations
1	Large scale multifactorial likelihood quantitative analysis of BRCA1 and BRCA2 variants: An ENIGMA resource to support clinical variant classification. Human Mutation, 2019, 40, 1557-1578.	2.5	102
2	Validation of a reverse-hybridization StripAssay for the simultaneous analysis of common Î±-thalassemia point mutations and deletions. Clinical Chemistry and Laboratory Medicine, 2007, 45, 605-10.	2.3	33
3	Mutational spectrum of cystic fibrosis in the Lebanese population. Journal of Cystic Fibrosis, 2010, 9, 406-410.	0.7	32
4	Genotype/Phenotype Correlation in Primary Congenital Glaucoma Patients in the Lebanese Population: A Pilot Study. Ophthalmic Genetics, 2016, 37, 1-6.	1.2	16
5	Goldenhar syndrome associated with prenatal maternal Fluoxetine ingestion: Cause or coincidence?. Birth Defects Research Part A: Clinical and Molecular Teratology, 2010, 88, 582-585.	1.6	14
6	BRCA mutation screening and patterns among high-risk Lebanese subjects. Hereditary Cancer in Clinical Practice, 2019, 17, 4.	1.5	13
7	Alpha thalassemia allelic frequency in Lebanon. Pediatric Blood and Cancer, 2015, 62, 120-122.	1.5	9
8	A synonymous variant in MYO15A enriched in the Ashkenazi Jewish population causes autosomal recessive hearing loss due to abnormal splicing. European Journal of Human Genetics, 2021, 29, 988-997.	2.8	8
9	De novo exceptional complex chromosomal rearrangement in a healthy fertile male: case report and review of the literature. Fertility and Sterility, 2011, 96, 1160-1164.	1.0	7
10	A Lebanese family with autosomal recessive oculo-auriculo-vertebral (OAV) spectrum and review of the literature: is OAV a genetically heterogeneous disorder. The Application of Clinical Genetics, 2011, 4, 93.	3.0	5
11	Incidence of Alpha-Globin Gene Defect in the Lebanese Population: A Pilot Study. BioMed Research International, 2015, 2015, 1-3.	1.9	5
12	Ellis-van Creveld Syndrome: Mutations Uncovered in Lebanese Families. Case Reports in Genetics, 2015, 2015, 1-4.	0.2	4
13	The Impact of Partial Weak Staining in Normal Breast Epithelium on the Reliability of Immunohistochemistry Results in HercepTest-positive Breast Cancer. Clinical Breast Cancer, 2019, 19, 340-344.	2.4	4
14	BACs-on-Beadsâ„¢ assay, a rapid aneuploidy test, improves the diagnostic yield of conventional karyotyping. Molecular Biology Reports, 2020, 47, 169-177.	2.3	4
15	FISH analyses for 1p and 19q status on gliomas: Reporting an 8Âyears' experience from a tertiary care center in the Middle East. Annals of Diagnostic Pathology, 2022, 57, 151899.	1.3	3
16	Molecular profiling of adult acute myeloid and lymphoid leukemia in a major referral center in Lebanon: a 10-year experience report and review of the literature. Molecular Biology Reports, 2019, 46, 2003-2011.	2.3	2
17	17p13.3 Microduplication Syndrome: Further Delineating the Clinical Spectrum. Journal of Pediatric Genetics, 2021, 10, 239-244.	0.7	2
18	The Spectrum of Î²-Thalassemia Mutations in the Population Migration in Lebanon: A 6-Year Retrospective Study. Hemoglobin, 2021, , 1-6.	0.8	1

#	Article	IF	Citations
19	Implementation of An Aggressive Risk Stratified Treatment Protocol for Children and Adolescents with Acute Lymphoblastic Leukemia in a Developing Country. Blood, 2011, 118, 876-876.	1.4	1
20	A novel cystic fibrosis gene mutation c.2490insT in a Palestinian patient: A case report and review of the literature. Annals of Thoracic Medicine, 2017, 12, 290.	1.8	1
21	BRCA mutations in a cohort of Iraqi patients presenting to a tertiary referral center. BMC Medical Genetics, 2019, 20, 154.	2.1	0
22	Acquired centromeric heteromorphism of chromosome 7 yields discordant results between fluorescent in situ hybridization and karyotype analysis in a child with severe congenital neutropenia. Pediatric Hematology and Oncology, 2019, 36, 432-437.	0.8	0
23	Novel pleiotropic BRCA2 pathogenic variants in Lebanese families. Cancer Genetics, 2019, 231-232, 32-35.	0.4	0
24	Novel human pathological mutations. Gene symbol: CFTR. Disease: cystic fibrosis. Human Genetics, 2007, 122, 553.	3.8	0
25	CFTR mutational screening by nextâ€generation sequencing reveals novel variants and a high carrier rate in a Middle Eastern population. Annals of Human Genetics, 2022, 86, 80-86.	0.8	0