

# Chantal Farra

## List of Publications by Year in descending order

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Version: 2024-02-01

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	FISH analyses for 1p and 19q status on gliomas: Reporting an 8 years' experience from a tertiary care center in the Middle East. <i>Annals of Diagnostic Pathology</i> , 2022, 57, 151899.	1.3	3
2	CFTR mutational screening by next-generation sequencing reveals novel variants and a high carrier rate in a Middle Eastern population. <i>Annals of Human Genetics</i> , 2022, 86, 80-86.	0.8	0
3	A synonymous variant in MYO15A enriched in the Ashkenazi Jewish population causes autosomal recessive hearing loss due to abnormal splicing. <i>European Journal of Human Genetics</i> , 2021, 29, 988-997.	2.8	8
4	The Spectrum of $\beta^2$ -Thalassemia Mutations in the Population Migration in Lebanon: A 6-Year Retrospective Study. <i>Hemoglobin</i> , 2021, , 1-6.	0.8	1
5	17p13.3 Microduplication Syndrome: Further Delineating the Clinical Spectrum. <i>Journal of Pediatric Genetics</i> , 2021, 10, 239-244.	0.7	2
6	BACs-on-Beads assay, a rapid aneuploidy test, improves the diagnostic yield of conventional karyotyping. <i>Molecular Biology Reports</i> , 2020, 47, 169-177.	2.3	4
7	The Impact of Partial Weak Staining in Normal Breast Epithelium on the Reliability of Immunohistochemistry Results in Herceptin-positive Breast Cancer. <i>Clinical Breast Cancer</i> , 2019, 19, 340-344.	2.4	4
8	BRCA mutations in a cohort of Iraqi patients presenting to a tertiary referral center. <i>BMC Medical Genetics</i> , 2019, 20, 154.	2.1	0
9	Acquired centromeric heteromorphism of chromosome 7 yields discordant results between fluorescent in situ hybridization and karyotype analysis in a child with severe congenital neutropenia. <i>Pediatric Hematology and Oncology</i> , 2019, 36, 432-437.	0.8	0
10	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. <i>Molecular Biology Reports</i> , 2019, 46, 2003-2011.	2.3	2
11	BRCA mutation screening and patterns among high-risk Lebanese subjects. <i>Hereditary Cancer in Clinical Practice</i> , 2019, 17, 4.	1.5	13
12	Large scale multifactorial likelihood quantitative analysis of BRCA1 and BRCA2 variants: An ENIGMA resource to support clinical variant classification. <i>Human Mutation</i> , 2019, 40, 1557-1578.	2.5	102
13	Novel pleiotropic BRCA2 pathogenic variants in Lebanese families. <i>Cancer Genetics</i> , 2019, 231-232, 32-35.	0.4	0
14	A novel cystic fibrosis gene mutation c.2490insT in a Palestinian patient: A case report and review of the literature. <i>Annals of Thoracic Medicine</i> , 2017, 12, 290.	1.8	1
15	Genotype/Phenotype Correlation in Primary Congenital Glaucoma Patients in the Lebanese Population: A Pilot Study. <i>Ophthalmic Genetics</i> , 2016, 37, 1-6.	1.2	16
16	Incidence of Alpha-Globin Gene Defect in the Lebanese Population: A Pilot Study. <i>BioMed Research International</i> , 2015, 2015, 1-3.	1.9	5
17	Ellis-van Creveld Syndrome: Mutations Uncovered in Lebanese Families. <i>Case Reports in Genetics</i> , 2015, 2015, 1-4.	0.2	4
18	Alpha thalassemia allelic frequency in Lebanon. <i>Pediatric Blood and Cancer</i> , 2015, 62, 120-122.	1.5	9

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19	De novo exceptional complex chromosomal rearrangement in a healthy fertile male: case report and review of the literature. <i>Fertility and Sterility</i> , 2011, 96, 1160-1164.	1.0	7
20	A Lebanese family with autosomal recessive oculo-auriculo-vertebral (OAV) spectrum and review of the literature: is OAV a genetically heterogeneous disorder. <i>The Application of Clinical Genetics</i> , 2011, 4, 93.	3.0	5
21	Implementation of An Aggressive Risk Stratified Treatment Protocol for Children and Adolescents with Acute Lymphoblastic Leukemia in a Developing Country. <i>Blood</i> , 2011, 118, 876-876.	1.4	1
22	Goldenhar syndrome associated with prenatal maternal Fluoxetine ingestion: Cause or coincidence?. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2010, 88, 582-585.	1.6	14
23	Mutational spectrum of cystic fibrosis in the Lebanese population. <i>Journal of Cystic Fibrosis</i> , 2010, 9, 406-410.	0.7	32
24	Validation of a reverse-hybridization StripAssay for the simultaneous analysis of common $\alpha$ -thalassemia point mutations and deletions. <i>Clinical Chemistry and Laboratory Medicine</i> , 2007, 45, 605-10.	2.3	33
25	Novel human pathological mutations. Gene symbol: CFTR. Disease: cystic fibrosis. <i>Human Genetics</i> , 2007, 122, 553.	3.8	0