

# Fady M Mikhail

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8864511/publications.pdf>

Version: 2024-02-01

38  
papers

1,871  
citations

430754

18  
h-index

345118

36  
g-index

38  
all docs

38  
docs citations

38  
times ranked

3008  
citing authors

#	ARTICLE	IF	CITATIONS
1	Chromosomal microarray analysis, including constitutional and neoplastic disease applications, 2021 revision: a technical standard of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021, 23, 1818-1829.	1.1	18
2	Outcomes of high-risk acute promyelocytic leukemia patients treated with arsenic trioxide (ATO)/all trans retinoic acid (ATRA) based induction and consolidation without maintenance phase: A case Series. <i>Hematology/ Oncology and Stem Cell Therapy</i> , 2020, 13, 143-146.	0.6	1
3	Recurrent microdeletions at chromosome 2p11.2 are associated with thymic hypoplasia and features resembling DiGeorge syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 358-367.e2.	1.5	24
4	Fragile Bones Secondary to SMURF1 Gene Duplication. <i>Calcified Tissue International</i> , 2020, 106, 567-573.	1.5	4
5	Evidence-based review of genomic aberrations in B-lymphoblastic leukemia/lymphoma: Report from the cancer genomics consortium working group for lymphoblastic leukemia. <i>Cancer Genetics</i> , 2020, 243, 52-72.	0.2	14
6	Masked hypodiploidy: Hypodiploid acute lymphoblastic leukemia (ALL) mimicking hyperdiploid ALL in children: A report from the Children's Oncology Group. <i>Cancer Genetics</i> , 2019, 238, 62-68.	0.2	32
7	Menkes disease complicated by concurrent Koolenâ€de Vries syndrome (17q21.31 deletion). <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e829.	0.6	3
8	Technical laboratory standards for interpretation and reporting of acquired copy-number abnormalities and copy-neutral loss of heterozygosity in neoplastic disorders: a joint consensus recommendation from the American College of Medical Genetics and Genomics (ACMG) and the Cancer Genomics Consortium (CGC). <i>Genetics in Medicine</i> , 2019, 21, 1903-1916.	1.1	39
9	Relapse after Prolonged Remission in Philadelphia-Like Acute Lymphoblastic Leukemia. <i>Case Reports in Hematology</i> , 2019, 2019, 1-3.	0.3	1
10	Central Nervous System Double Relapse of Acute Promyelocytic Leukemia and Acute Myelomonocytic Leukemia. <i>Case Reports in Hematology</i> , 2019, 2019, 1-6.	0.3	1
11	Therapyâ€associated myelodysplastic syndrome with monosomy 7 arising in a Muirâ€Torre Syndrome patient carrying SETBP1 mutation. <i>Molecular and Clinical Oncology</i> , 2018, 8, 306-309.	0.4	0
12	Maternal uniparental isodisomy for chromosome 6 discovered by paternity testing: a case report. <i>Molecular Cytogenetics</i> , 2018, 11, 60.	0.4	8
13	A previously unrecognized 22q13.2 microdeletion syndrome that encompasses <i>TCF20</i> and <i>TNFRSF13C</i> . <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 2791-2797.	0.7	22
14	Omission of Maintenance in Patients with High-Risk Acute Promyelocytic Leukemia (APL) in the Era of ATRA/Arsenic Consolidation. <i>Blood</i> , 2018, 132, 5192-5192.	0.6	1
15	Overview of Genetic Diagnosis in Cancer. <i>Current Protocols in Human Genetics</i> , 2017, 93, 10.1.1-10.1.9.	3.5	2
16	Enhanced targeting of CML stem and progenitor cells by inhibition of porcupine acyltransferase in combination with TKI. <i>Blood</i> , 2017, 129, 1008-1020.	0.6	58
17	Diagnostic and Prognostic Utility of Fluorescence In situ Hybridization (FISH) Analysis in Acute Myeloid Leukemia. <i>Current Hematologic Malignancy Reports</i> , 2017, 12, 568-573.	1.2	12
18	Clinical relevance of small copy-number variants in chromosomal microarray clinical testing. <i>Genetics in Medicine</i> , 2017, 19, 377-385.	1.1	24

#	ARTICLE	IF	CITATIONS
19	Section E6.5â€“6.8 of the ACMG technical standards and guidelines: chromosome studies of lymph node and solid tumorâ€“acquired chromosomal abnormalities. <i>Genetics in Medicine</i> , 2016, 18, 643-648.	1.1	17
20	Section E6.1â€“6.4 of the ACMG technical standards and guidelines: chromosome studies of neoplastic blood and bone marrowâ€“acquired chromosomal abnormalities. <i>Genetics in Medicine</i> , 2016, 18, 635-642.	1.1	35
21	Cleft palate in a patient with the nested 22q11.2 LCR C to D deletion. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 260-262.	0.7	2
22	Family history of hematologic malignancies and risk of multiple myeloma: differences by race and clinical features. <i>Cancer Causes and Control</i> , 2016, 27, 81-91.	0.8	35
23	Array comparative genomic hybridisation testing in CHD. <i>Cardiology in the Young</i> , 2015, 25, 1155-1172.	0.4	7
24	Copy number variations and human genetic disease. <i>Current Opinion in Pediatrics</i> , 2014, 26, 646-652.	1.0	36
25	The recurrent distal 22q11.2 microdeletions are often de novo and do not represent a single clinical entity: a proposed categorization system. <i>Genetics in Medicine</i> , 2014, 16, 92-100.	1.1	49
26	Incidental Detection of Cancer Predisposition Gene Copy Number Variations by Array Comparative Genomic Hybridization. <i>Journal of Pediatrics</i> , 2014, 165, 1057-1059.e4.	0.9	9
27	Quality Assurance and Quality Control in Clinical Cytogenetics. <i>Current Protocols in Human Genetics</i> , 2014, 82, 8.2.1-10.	3.5	1
28	Clinically relevant single gene or intragenic deletions encompassing critical neurodevelopmental genes in patients with developmental delay, mental retardation, and/or autism spectrum disorders. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 2386-2396.	0.7	159
29	Rearrangement of CRLF2 is associated with mutation of JAK kinases, alteration of IKZF1, Hispanic/Latino ethnicity, and a poor outcome in pediatric B-progenitor acute lymphoblastic leukemia. <i>Blood</i> , 2010, 115, 5312-5321.	0.6	503
30	Rearrangement of CRLF2 in B-progenitorâ€“ and Down syndromeâ€“associated acute lymphoblastic leukemia. <i>Nature Genetics</i> , 2009, 41, 1243-1246.	9.4	559
31	Rearrangement of CRLF2 in B-Progenitor and Down Syndrome Associated Acute Lymphoblastic Leukemia.. <i>Blood</i> , 2009, 114, 182-182.	0.6	0
32	Distal 22q11.2 microduplication encompassing the <i>BCR</i> gene. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 3075-3081.	0.7	30
33	Overlapping phenotype of Wolfâ€“Hirschhorn and Beckwithâ€“Wiedemann syndromes in a girl with der(4)t(4;11)(pter;pter). <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 1760-1766.	0.7	5
34	A previously unrecognized microdeletion syndrome on chromosome 22 band q11.2 encompassing the <i>BCR</i> gene. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 2178-2184.	0.7	42
35	Normal and transforming functions of RUNX1: A perspective. <i>Journal of Cellular Physiology</i> , 2006, 207, 582-593.	2.0	60
36	Complete trisomy 17p syndrome in a girl with der(14)t(14;17)(p11.2;p11.2). <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 1647-1654.	0.7	11

#	ARTICLE	IF	CITATIONS
37	A novel gene, FGA7, is fused to RUNX1/AML1 in a t(4;21)(q28;q22) in a patient with T-cell acute lymphoblastic leukemia. <i>Genes Chromosomes and Cancer</i> , 2004, 39, 110-118.	1.5	26
38	A new translocation that rearranges the AML1 gene in a patient with T-cell acute lymphoblastic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2002, 135, 96-100.	1.0	21