Roula A Farah

List of Publications by Year
in descending order

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63	2,968	393982	182168
papers	citations	19 h-index	51 g-index

64	64	64	5976
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Immune Checkpoint Inhibition for Hypermutant Glioblastoma Multiforme Resulting From Germline Biallelic Mismatch Repair Deficiency. Journal of Clinical Oncology, 2016, 34, 2206-2211.	0.8	692
2	Comprehensive Analysis of Hypermutation in Human Cancer. Cell, 2017, 171, 1042-1056.e10.	13.5	596
3	Combined hereditary and somatic mutations of replication error repair genes result in rapid onset of ultra-hypermutated cancers. Nature Genetics, 2015, 47, 257-262.	9.4	306
4	Defective cytotoxic lymphocyte degranulation in syntaxin-11â€“deficient familial hemophagocytic lymphohistiocytosis 4 (FHL4) patients. Blood, 2007, 110, 1906-1915.	0.6	272
5	Genetic and clinical determinants of constitutional mismatch repair deficiency syndrome: Report from the constitutional mismatch repair deficiency consortium. European Journal of Cancer, 2014, 50, 987-996.	1.3	180
6	Flash survey on severe acute respiratory syndrome coronavirus-2 infections in paediatric patients on anticancer treatment. European Journal of Cancer, 2020, 132, 11-16.	1.3	155
7	Identification of new Wilms tumour predisposition genes: an exome sequencing study. The Lancet Child and Adolescent Health, 2019, 3, 322-331.	2.7	82
8	Comparison of laparoscopic and open splenectomy in children with hematologic disorders. Journal of Pediatrics, 1997, 131, 41-46.	0.9	80
9	The Development of Monoclonal Antibodies for the Therapy of Cancer. Critical Reviews in Eukaryotic Gene Expression, 1998, 8, 321-356.	0.4	66
10	Immunotoxins against CD19 and CD22 are effective in killing precursor-B acute lymphoblastic leukemia cells in vitro. Leukemia, 2000, 14, 853-858.	3.3	65
11	Lentiviral gene therapy corrects platelet phenotype and function in patients with Wiskott-Aldrich syndrome. Journal of Allergy and Clinical Immunology, 2019, 144, 825-838.	1.5	50
12	Survival Benefit for Individuals With Constitutional Mismatch Repair Deficiency Undergoing Surveillance. Journal of Clinical Oncology, 2021, 39, 2779-2790.	0.8	40
13	Gastrointestinal Findings in the Largest Series of Patients With Hereditary Biallelic Mismatch Repair Deficiency Syndrome: Report from the International Consortium. American Journal of Gastroenterology, 2016, 111, 275-284.	0.2	33
14	Functional Repair Assay for the Diagnosis of Constitutional Mismatch Repair Deficiency From Non-Neoplastic Tissue. Journal of Clinical Oncology, 2019, 37, 461-470.	0.8	23
15	Germline-driven replication repair-deficient high-grade gliomas exhibit unique hypomethylation patterns. Acta Neuropathologica, 2020, 140, 765-776.	3.9	23
16	Limb salvage surgery for children and adolescents with malignant bone tumors in a developing country. Pediatric Blood and Cancer, 2008, 51, 787-791.	0.8	22
17	Contribution of next generation sequencing in pediatric practice in Lebanon. A Study on 213 cases. Molecular Genetics & Genomic Medicine, 2018, 6, 1041-1052.	0.6	22
18	Successful prophylaxis against intracranial hemorrhage using weekly administration of activated recombinant factor VII in a newborn with severe factor VII deficiency. Journal of Thrombosis and Haemostasis, 2007, 5, 433-434.	1.9	20

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19	Current recommendations for clinical surveillance and genetic testing in rhabdoid tumor predisposition: a report from the SIOPE Host Genome Working Group. Familial Cancer, 2021, 20, 305-316.	0.9	20
20	The phenotypic spectrum of germline YARS2 variants: from isolated sideroblastic anemia to mitochondrial myopathy, lactic acidosis and sideroblastic anemia 2. Haematologica, 2018, 103, 2008-2015.	1.7	19
21	Disseminated Coxsackie A9 Infection Complicating Bone Marrow Transplantation. Pediatric Infectious Disease Journal, 1996, 15, 1053-1054.	1.1	19
22	Lifeâ€threatening bleeding in factor <scp>VII</scp> deficiency: the role of prenatal diagnosis and primary prophylaxis. British Journal of Haematology, 2015, 168, 452-455.	1.2	17
23	Clinical and genetic characteristics of children with acute lymphoblastic leukemia and Liâ€“Fraumeni syndrome. Leukemia, 2021, 35, 1475-1479.	3.3	17
24	Next-generation sequencing and recombinant expression characterized aberrant splicing mechanisms and provided correction strategies in factor VII deficiency. Haematologica, 2020, 105, 829-837.	1.7	16
25	Relapsed Childhood Acute Lymphoblastic Leukemia Presenting as an Isolated Breast Mass. Clinical Pediatrics, 1999, 38, 545-546.	0.4	13
26	Homozygous mutation in ELMO2 may cause Ramon syndrome. Clinical Genetics, 2018, 93, 703-706.	1.0	13
27	SLC25A38 congenital sideroblastic anemia: Phenotypes and genotypes of 31 individuals from 24 families, including 11 novel mutations, and a review of the literature. Human Mutation, 2021, 42, 1367-1383.	1.1	11
28	Safety and Cost-Effectiveness of Outpatient Total Body Irradiation in Pediatric Patients Undergoing Stem Cell Transplantation. Journal of Pediatric Hematology/Oncology, 1998, 20, 319-321.	0.3	10
29	PARAMENINGEAL ALVEOLAR RHABDOMYOSARCOMA WITH AN ISOLATED PANCREATIC METASTASIS. Pediatric Hematology and Oncology, 1999, 16, 463-467.	0.3	10
30	A MULTICENTER EXPERIENCE FROM LEBANON IN CHILDHOOD AND ADOLESCENT ACUTE MYELOID LEUKEMIA:HIGH RATE OF EARLY DEATH IN CHILDHOOD ACUTE PROMYELOCYTIC LEUKEMIA.. Mediterranean Journal of Hematology and Infectious Diseases, 2014, 7, e2015012.	0.5	8
31	Combined factor V â€“ factor VIII deficiency (F5F8D): Compound heterozygosity for two novel truncating mutations in LMAN1 in a consanguineous patient. Thrombosis and Haemostasis, 2006, 95, 893-895.	1.8	8
32	Ongoing issues with the management of children with Constitutional Mismatch Repair Deficiency syndrome. European Journal of Medical Genetics, 2019, 62, 103706.	0.7	7
33	Acquired protein C deficiency in a child with acute myelogenous leukemia, splenic, renal, and intestinal infarction. Blood Coagulation and Fibrinolysis, 2011, 22, 140-143.	0.5	6
34	Establishment of a formal program for retinoblastoma: Feasibility of clinical coordination across borders and impact on outcome. Pediatric Blood and Cancer, 2019, 66, e27959.	0.8	6
35	Immune thrombocytopenic purpura associated with brucellosis. Case report and review of the literature. Journal Medical Libanais, 2010, 58, 241-3.	0.0	6
36	Spontaneous Epidural Hematoma in a Child With Inherited Factor XIII Deficiency. Journal of Pediatric Hematology/Oncology, 2014, 36, 62-65.	0.3	5

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37	Association of CYP3A4/5 genotypes and expression with the survival of patients with neuroblastoma. Molecular Medicine Reports, 2015, 11, 1462-1468.	1.1	4
38	Collaborative Pediatric Bone Tumor Program to Improve Access to Specialized Care: An Initiative by the Lebanese Childrenâ€™s Oncology Group. Journal of Global Oncology, 2017, 3, 23-30.	0.5	4
39	Treatment of pediatric Hodgkin's disease with chemotherapy alone or combined modality therapy. Radiation Oncology Investigations, 1999, 7, 365-373.	1.3	3
40	Double Heterozygosity for Hb G-San JosÃ© [Î²7(A4)Gluâ†'Gly] and Hb Fukuoka [Î²2(NA2)Hisâ†'Tyr] in a 2¹/₂-Year-Old Girl. Hemoglobin, 1999, 23, 383-387.	0.4	3
41	Accurate characterization of the IVS7 repeat polymorphism of FVII gene and identification of three novel allelic forms. Thrombosis and Haemostasis, 2006, 96, 95-97.	1.8	3
42	COVID-19 associated respiratory failure complicating a pericardial effusion in a patient with sideroblastic anemia. Respiratory Medicine Case Reports, 2021, 34, 101543.	0.2	3
43	Extragonadal germ cell tumor presenting with spinal cord compression: a case report and literature review. Annals of Pediatric Surgery, 2022, 18, .	0.1	2
44	Correlation of Serum Cholyiglycine Level with Hepatic Dysfunction in Children with Sickle Cell Anemia. Clinical Pediatrics, 1999, 38, 293-296.	0.4	1
45	Pandemic (H1N1) 2009 Influenza in Pediatric Hematology/Oncology Units in Lebanon. Journal of Pediatric Hematology/Oncology, 2013, 35, 144-147.	0.3	1
46	Refractory autoimmune disease: an overview of when first-line therapy is not enough. Seminars in Hematology, 2016, 53, S35-S38.	1.8	1
47	Histiocytoses initiatives in Asia: The Asian-Middle Eastern (AME) Histiocytoses Network. Pediatric Hematology Oncology Journal, 2020, 5, 137-139.	0.1	1
48	MBRS-54. POOR SURVIVAL IN REPLICATION REPAIR DEFICIENT HYPERMUTANT MEDULLOBLASTOMA AND CNS EMBRYONAL TUMORS: A REPORT FROM THE INTERNATIONAL RRD CONSORTIUM. Neuro-Oncology, 2020, 22, iii407-iii407.	0.6	1
49	Clinical and Genetic Features of Patients With Fanconi Anemia in Lebanon and Report on Novel Mutations in the FANCA and FANCG Genes. Journal of Pediatric Hematology/Oncology, 2021, 43, e727-e735.	0.3	1
50	Experience of the use of trabectedin in pretreated unresectable, advanced, or metastatic soft tissue sarcoma in nine centers in Lebanon on a compassionate-use basis.. Journal of Clinical Oncology, 2011, 29, e20533-e20533.	0.8	1
51	Palliative Tumor Control by Trabectedin in Pediatric Advanced Sarcoma. Current Drug Therapy, 2011, 6, 97-99.	0.2	1
52	SP-057 RARE BLEEDING DISORDERS: UPDATE ON DIAGNOSIS AND MANAGEMENT, ROLE OF PRENATAL DIAGNOSIS AND PROPHYLAXIS. Leukemia Research, 2014, 38, S23-S24.	0.4	0
53	Sa2045 Tumor Spectrum, Diagnostic Tools and Survival in Patients With Biallelic Mismatch Repair Gene Deficiency (BMMRD) Syndrome: Report From the International BMMRD Consortium. Gastroenterology, 2016, 150, S438-S439.	0.6	0
54	HGG-20. DNA METHYLATION ANALYSIS OF HIGH-GRADE GLIOMA IN PATIENTS WITH MISMATCH REPAIR DEFICIENCIES. Neuro-Oncology, 2018, 20, i92-i93.	0.6	0

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55	A Novel Deletion in the RPL5 Gene in a Lebanese Child With Diamond Blackfan Anemia Unresponsive to Steroid Treatment. Journal of Pediatric Hematology/Oncology, 2020, 42, e235-e237.	0.3	0
56	Updated results of a phase II trial of therapy with cyclophosphamide (CYP)/pentostatin/rituximab for indolent NHL and CLL. Journal of Clinical Oncology, 2005, 23, 6713-6713.	0.8	0
57	SERUM CHOLYLGLYCINE LEVEL AS A MEASURE OF HEPATIC DYSFUNCTION IN CHILDREN WITH SICKLE CELL ANEMIA. Journal of Pediatric Gastroenterology and Nutrition, 1997, 25, 476.	0.9	0
58	Abstract 35: Novel genetic and clinical determinants of Constitutional Mismatch Repair Deficiency syndrome: Report from the CMMRD consortium. , 2014, , .		0
59	A multi-institutional collaborative pediatric bone tumor program for improving access to specialized care.. Journal of Clinical Oncology, 2015, 33, e21020-e21020.	0.8	0
60	Successful international collaboration establishing a regional retinoblastoma program.. Journal of Clinical Oncology, 2015, 33, e21024-e21024.	0.8	0
61	Abstract B09: DNA polymerase mutations trigger rapid onset of ultra-hypermutant malignant brain tumors in children with biallelic mismatch repair deficiency. , 2015, , .		0
62	RARE-17. SURVIVAL BENEFIT FOR INDIVIDUALS WITH CONSTITUTIONAL MISMATCH REPAIR DEFICIENCY SYNDROME AND BRAIN TUMORS WHO UNDERGO SURVEILLANCE PROTOCOL. A REPORT FROM THE INTERNATIONAL REPLICATION REPAIR CONSORTIUM. Neuro-Oncology, 2020, 22, iii445-iii446.	0.6	0
63	HGG-20. DIAGNOSTIC AND BIOLOGICAL ROLE OF METHYLATION PATTERNS IN REPLICATION REPAIR DEFICIENT HIGH GRADE GLIOMAS. Neuro-Oncology, 2020, 22, iii347-iii348.	0.6	0