

Raphael Carapito

List of Publications by Year in descending order

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Version: 2024-02-01

71
papers

2,024
citations

279487

23
h-index

288905

40
g-index

81
all docs

81
docs citations

81
times ranked

3959
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of driver genes for critical forms of COVID-19 in a deeply phenotyped young patient cohort. <i>Science Translational Medicine</i> , 2022, 14, eabj7521.	5.8	71
2	A Homozygous Missense Variant in <i>PPP1R1B/DARPP32</i> Is Associated With Generalized Complex Dystonia. <i>Movement Disorders</i> , 2022, 37, 365-374.	2.2	7
3	The MHC class I MICA gene is a histocompatibility antigen in kidney transplantation. <i>Nature Medicine</i> , 2022, 28, 989-998.	15.2	20
4	A gain-of-function variant in the Wiskott-Aldrich syndrome gene is associated with a MYH9-related disease-like syndrome. <i>Blood Advances</i> , 2022, 6, 5279-5284.	2.5	2
5	Fc receptors and the diversity of antibody responses to HIV infection and vaccination. <i>Genes and Immunity</i> , 2022, 23, 149-156.	2.2	1
6	selectBoost: a general algorithm to enhance the performance of variable selection methods. <i>Bioinformatics</i> , 2021, 37, 659-668.	1.8	1
7	Structural and Functional Impact of SRP54 Mutations Causing Severe Congenital Neutropenia. <i>Structure</i> , 2021, 29, 15-28.e7.	1.6	12
8	<i>SRP54</i> mutations induce congenital neutropenia via dominant-negative effects on <i>XBP1</i> splicing. <i>Blood</i> , 2021, 137, 1340-1352.	0.6	15
9	Atypical focal segmental glomerulosclerosis associated with a new <i>PODXL</i> nonsense variant. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1658.	0.6	2
10	Temporal multiomic modeling reveals a B-cell receptor proliferative program in chronic lymphocytic leukemia. <i>Leukemia</i> , 2021, 35, 1463-1474.	3.3	6
11	Tenascin ϵ immobilizes infiltrating T lymphocytes through CXCL12 promoting breast cancer progression. <i>EMBO Molecular Medicine</i> , 2021, 13, e13270.	3.3	27
12	Impairing flow-mediated endothelial remodeling reduces extravasation of tumor cells. <i>Scientific Reports</i> , 2021, 11, 13144.	1.6	12
13	Impact of Tenascin-C on Radiotherapy in a Novel Syngeneic Oral Squamous Cell Carcinoma Model With Spontaneous Dissemination to the Lymph Nodes. <i>Frontiers in Immunology</i> , 2021, 12, 636108.	2.2	6
14	Ral GTPases promote breast cancer metastasis by controlling biogenesis and organ targeting of exosomes. <i>ELife</i> , 2021, 10, .	2.8	70
15	Contrasting role of <i>NLRP12</i> in autoinflammation: evidence from a case report and mouse models. <i>RMD Open</i> , 2021, 7, e001824.	1.8	5
16	A de novo synonymous variant in <i>EFTUD2</i> disrupts normal splicing and causes mandibulofacial dysostosis with microcephaly: case report. <i>BMC Medical Genetics</i> , 2020, 21, 182.	2.1	8
17	Tenascin-C Orchestrates an Immune-Suppressive Tumor Microenvironment in Oral Squamous Cell Carcinoma. <i>Cancer Immunology Research</i> , 2020, 8, 1122-1138.	1.6	40
18	A <i>FCER1a</i> polymorphism has a HLA-B57 and HLA-B27 independent effect on HIV disease outcome. <i>Genes and Immunity</i> , 2020, 21, 263-268.	2.2	5

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19	Progressive multifocal leukoencephalopathy and sarcoidosis under interleukin 7. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020, 7, e862.	3.1	6
20	<i>NCKAP1L</i> defects lead to a novel syndrome combining immunodeficiency, lymphoproliferation, and hyperinflammation. <i>Journal of Experimental Medicine</i> , 2020, 217, .	4.2	48
21	Improvement of therapy-induced myelodysplastic syndrome by infusion of autologous CD34-positive hematopoietic progenitor cells without chemotherapy. <i>Leukemia and Lymphoma</i> , 2020, 61, 3259-3262.	0.6	1
22	Two Cases of Recessive Intellectual Disability Caused by NDST1 and METTL23 Variants. <i>Genes</i> , 2020, 11, 1021.	1.0	9
23	Protocol for TRAUMADORNASE: a prospective, randomized, multicentre, double-blinded, placebo-controlled clinical trial of aerosolized dornase alfa to reduce the incidence of moderate-to-severe hypoxaemia in ventilated trauma patients. <i>Trials</i> , 2020, 21, 274.	0.7	12
24	Matrix-Targeting Immunotherapy Controls Tumor Growth and Spread by Switching Macrophage Phenotype. <i>Cancer Immunology Research</i> , 2020, 8, 368-382.	1.6	42
25	High-Throughput MICA/B Genotyping of Over Two Million Samples: Workflow and Allele Frequencies. <i>Frontiers in Immunology</i> , 2020, 11, 314.	2.2	28
26	A mouse model of MSU-induced acute inflammation <i>in vivo</i> suggests imiquimod-dependent targeting of <i>IL-1β</i> as relevant therapy for gout patients. <i>Theranostics</i> , 2020, 10, 2158-2171.	4.6	28
27	Compatibility at amino acid position 98 of MICB reduces the incidence of graft-versus-host disease in conjunction with the CMV status. <i>Bone Marrow Transplantation</i> , 2020, 55, 1367-1378.	1.3	9
28	Multi-omics dataset to decipher the complexity of drug resistance in diffuse large B-cell lymphoma. <i>Scientific Reports</i> , 2019, 9, 895.	1.6	38
29	<i>IKZF1</i> Loss-of-Function Variant Causes Autoimmunity and Severe Familial Antiphospholipid Syndrome. <i>Journal of Clinical Immunology</i> , 2019, 39, 353-357.	2.0	15
30	Exome sequencing identifies a novel missense variant in <i>CTSC</i> causing nonsyndromic aggressive periodontitis. <i>Journal of Human Genetics</i> , 2019, 64, 689-694.	1.1	8
31	HLA*LA HLA typing from linearly projected graph alignments. <i>Bioinformatics</i> , 2019, 35, 4394-4396.	1.8	88
32	Update on Fc-Mediated Antibody Functions Against HIV-1 Beyond Neutralization. <i>Frontiers in Immunology</i> , 2019, 10, 2968.	2.2	44
33	<i>ZMIZ1</i> Variants Cause a Syndromic Neurodevelopmental Disorder. <i>American Journal of Human Genetics</i> , 2019, 104, 319-330.	2.6	30
34	Comprehensive characterization of pseudomyxoma peritonei.. <i>Journal of Clinical Oncology</i> , 2019, 37, e15701-e15701.	0.8	1
35	A Core Proliferative Program Induced By B-Cell Receptor Stimulation in Chronic Lymphocytic Leukemia Cells. <i>Blood</i> , 2019, 134, 3777-3777.	0.6	0
36	Mutation-Specific Dominant Negative Effects Determine the Phenotype in <i>SRP54</i> Deficiency. <i>Blood</i> , 2019, 134, 437-437.	0.6	0

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37	Hemodynamic Forces Tune the Arrest, Adhesion, and Extravasation of Circulating Tumor Cells. <i>Developmental Cell</i> , 2018, 45, 33-52.e12.	3.1	219
38	An unusually high substitution rate in transplant-associated BK polyomavirus in vivo is further concentrated in HLA-C-bound viral peptides. <i>PLoS Pathogens</i> , 2018, 14, e1007368.	2.1	22
39	Multi-OMICS analyses unveil <i>STAT1</i> as a potential modifier gene in mevalonate kinase deficiency. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 1675-1687.	0.5	19
40	Mutation-Specific Dose Reduction in Functional SRP54 Protein Causes Isolated or Syndromic Neutropenia with Shwachman-Diamond-like Features. <i>Blood</i> , 2018, 132, 1300-1300.	0.6	1
41	MicroRNA-146a governs fibroblast activation and joint pathology in arthritis. <i>Journal of Autoimmunity</i> , 2017, 82, 74-84.	3.0	43
42	A new MHC-linked susceptibility locus for primary Sjögren's syndrome: MICA. <i>Human Molecular Genetics</i> , 2017, 26, 2565-2576.	1.4	22
43	A novel zebrafish model of congenital neutropenia. <i>Experimental Hematology</i> , 2017, 53, S88.	0.2	0
44	Natural Killer Group 2, Member D/NKG2D Ligands in Hematopoietic Cell Transplantation. <i>Frontiers in Immunology</i> , 2017, 8, 368.	2.2	21
45	Mutations in signal recognition particle SRP54 cause syndromic neutropenia with Shwachman-Diamond-like features. <i>Journal of Clinical Investigation</i> , 2017, 127, 4090-4103.	3.9	126
46	Protein-altering MYH3 variants are associated with a spectrum of phenotypes extending to spondylocarpotarsal synostosis syndrome. <i>European Journal of Human Genetics</i> , 2016, 24, 1746-1751.	1.4	21
47	Next-Generation Sequencing of the HLA locus: Methods and impacts on HLA typing, population genetics and disease association studies. <i>Human Immunology</i> , 2016, 77, 1016-1023.	1.2	66
48	The TRANSPLANTEX initiative. <i>Human Immunology</i> , 2016, 77, 1005-1007.	1.2	1
49	Matching for the nonconventional MHC-I MICA gene significantly reduces the incidence of acute and chronic GVHD. <i>Blood</i> , 2016, 128, 1979-1986.	0.6	66
50	Homozygosity for the V377I mutation in mevalonate kinase causes distinct clinical phenotypes in two sibs with hyperimmunoglobulinaemia D and periodic fever syndrome (HIDS). <i>RMD Open</i> , 2016, 2, e000196.	1.8	20
51	Germline variation of TNFAIP3 in primary Sjögren's syndrome-associated lymphoma. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 780-783.	0.5	40
52	Genetics, genomics, and evolutionary biology of NKG2D ligands. <i>Immunological Reviews</i> , 2015, 267, 88-116.	2.8	99
53	A <i>de novo</i> <i>ADCY5</i> mutation causes early-onset autosomal dominant chorea and dystonia. <i>Movement Disorders</i> , 2015, 30, 423-427.	2.2	131
54	Eosinophilic myositis as first manifestation in a patient with type 2 myotonic dystrophy CCTG expansion mutation and rheumatoid arthritis. <i>Neuromuscular Disorders</i> , 2015, 25, 149-152.	0.3	11

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55	On the genetics of the Silk Route: association analysis of HLA, IL10, and IL23R-IL12RB2 regions with Behçet's disease in an Iranian population. <i>Immunogenetics</i> , 2015, 67, 289-293.	1.2	21
56	Polymorphisms in EGFR and IL28B are associated with spontaneous clearance in an HCV-infected Iranian population. <i>Genes and Immunity</i> , 2015, 16, 514-518.	2.2	8
57	Homozygous <i>IL36RN</i> mutation and <i>NSD1</i> duplication in a patient with severe pustular psoriasis and symptoms unrelated to deficiency of interleukin-36 receptor antagonist. <i>British Journal of Dermatology</i> , 2015, 172, 302-305.	1.4	10
58	A new mutation in the C-SH2 domain of PTPN11 causes Noonan syndrome with multiple giant cell lesions. <i>Journal of Human Genetics</i> , 2014, 59, 57-59.	1.1	13
59	High diversity of MIC genes in non-human primates. <i>Immunogenetics</i> , 2014, 66, 581-587.	1.2	13
60	Matching of MHC Class I Chain-Related Genes a and B Is Associated with Reduced Incidence of Severe Acute Graft-Versus-Host Disease after Unrelated Hematopoietic Stem Cell Transplantation. <i>Blood</i> , 2014, 124, 664-664.	0.6	3
61	Genome-wide transcriptional responses of <i>Fusarium graminearum</i> to plant cell wall substrates. <i>FEMS Microbiology Letters</i> , 2013, 340, 129-134.	0.7	15
62	±-l-Arabinofuranosylated pyrrolidines as arabinanase inhibitors. <i>Chemical Communications</i> , 2011, 47, 9684.	2.2	6
63	TACKLING HUMULUS LUPULUS FUNGAL DISEASES BY STUDYING A HOP CELL WALL / FUSARIUM GRAMINEARUM MODEL. <i>Acta Horticulturae</i> , 2009, , 103-114.	0.1	0
64	Efficient hydrolysis of hemicellulose by a <i>Fusarium graminearum</i> xylanase blend produced at high levels in <i>Escherichia coli</i> . <i>Bioresource Technology</i> , 2009, 100, 845-850.	4.8	27
65	Molecular Basis of Arabinobio-hydrolase Activity in Phytopathogenic Fungi. <i>Journal of Biological Chemistry</i> , 2009, 284, 12285-12296.	1.6	42
66	Plant Cell Wall Degradation with a Powerful <i>Fusarium graminearum</i> Enzymatic Arsenal. <i>Journal of Microbiology and Biotechnology</i> , 2009, 19, 573-81.	0.9	15
67	Gene expression in <i>Fusarium graminearum</i> grown on plant cell wall. <i>Fungal Genetics and Biology</i> , 2008, 45, 738-748.	0.9	39
68	Common Alterations in PBP1a from Resistant <i>Streptococcus pneumoniae</i> Decrease Its Reactivity toward β-Lactams. <i>Journal of Biological Chemistry</i> , 2008, 283, 4886-4894.	1.6	44
69	Automated high-throughput process for site-directed mutagenesis, production, purification, and kinetic characterization of enzymes. <i>Analytical Biochemistry</i> , 2006, 355, 110-116.	1.1	11
70	Pneumococcal β-Lactam Resistance Due to a Conformational Change in Penicillin-binding Protein 2x. <i>Journal of Biological Chemistry</i> , 2006, 281, 1771-1777.	1.6	55
71	Identical Penicillin-Binding Domains in Penicillin-Binding Proteins of <i>Streptococcus pneumoniae</i> Clinical Isolates with Different Levels of β-Lactam Resistance. <i>Antimicrobial Agents and Chemotherapy</i> , 2005, 49, 2895-2902.	1.4	44