

Mohammad R Ahmadian

List of Publications by Year in descending order

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papers

5,274
citations

147801

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times ranked

6846
citing authors

#	ARTICLE	IF	CITATIONS
1	The Ras-RasGAP Complex: Structural Basis for GTPase Activation and Its Loss in Oncogenic Ras Mutants. <i>Science</i> , 1997, 277, 333-338.	12.6	1,378
2	GTPase-activating proteins: helping hands to complement an active site. <i>Trends in Biochemical Sciences</i> , 1998, 23, 257-262.	7.5	395
3	Confirmation of the arginine-finger hypothesis for the GAP-stimulated GTP-hydrolysis reaction of Ras. <i>Nature Structural Biology</i> , 1997, 4, 686-689.	9.7	333
4	A restricted spectrum of NRAS mutations causes Noonan syndrome. <i>Nature Genetics</i> , 2010, 42, 27-29.	21.4	271
5	Juvenile myelomonocytic leukemia displays mutations in components of the RAS pathway and the PRC2 network. <i>Nature Genetics</i> , 2015, 47, 1334-1340.	21.4	152
6	Always look on the bright site of Rho: structural implications for a conserved intermolecular interface. <i>EMBO Reports</i> , 2004, 5, 1130-1136.	4.5	141
7	Functional Dysregulation of CDC42 Causes Diverse Developmental Phenotypes. <i>American Journal of Human Genetics</i> , 2018, 102, 309-320.	6.2	138
8	Individual Rate Constants for the Interaction of Ras Proteins with GTPase-Activating Proteins Determined by Fluorescence Spectroscopy. <i>Biochemistry</i> , 1997, 36, 4535-4541.	2.5	135
9	A novel disorder involving dyshematopoiesis, inflammation, and HLH due to aberrant CDC42 function. <i>Journal of Experimental Medicine</i> , 2019, 216, 2778-2799.	8.5	132
10	Alternative Splicing of Rac1 Generates Rac1b, a Self-activating GTPase. <i>Journal of Biological Chemistry</i> , 2004, 279, 4743-4749.	3.4	127
11	Germline KRAS mutations cause aberrant biochemical and physical properties leading to developmental disorders. <i>Human Mutation</i> , 2011, 32, 33-43.	2.5	126
12	An Electrostatic Steering Mechanism of Cdc42 Recognition by Wiskott-Aldrich Syndrome Proteins. <i>Molecular Cell</i> , 2005, 20, 313-324.	9.7	117
13	Structural Insights into the Interaction of ROCK1 with the Switch Regions of RhoA. <i>Journal of Biological Chemistry</i> , 2004, 279, 7098-7104.	3.4	116
14	Activating mutations in RRAS underlie a phenotype within the RASopathy spectrum and contribute to leukaemogenesis. <i>Human Molecular Genetics</i> , 2014, 23, 4315-4327.	2.9	114
15	The RHO Family GTPases: Mechanisms of Regulation and Signaling. <i>Cells</i> , 2021, 10, 1831.	4.1	113
16	Deciphering the Molecular and Functional Basis of Dbl Family Proteins. <i>Journal of Biological Chemistry</i> , 2013, 288, 4486-4500.	3.4	91
17	Deciphering the Molecular and Functional Basis of RHOGAP Family Proteins. <i>Journal of Biological Chemistry</i> , 2016, 291, 20353-20371.	3.4	87
18	The C2 domain of SynGAP is essential for stimulation of the Rap GTPase reaction. <i>EMBO Reports</i> , 2008, 9, 350-355.	4.5	82

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19	Comparative functional analysis of the Rac GTPases. <i>FEBS Letters</i> , 2003, 555, 556-560.	2.8	65
20	Galectin-1 dimers can scaffold Raf-effectors to increase H-ras nanoclustering. <i>Scientific Reports</i> , 2016, 6, 24165.	3.3	65
21	Structural Fingerprints of the Ras-GTPase Activating Proteins Neurofibromin and p120GAP. <i>Journal of Molecular Biology</i> , 2003, 329, 699-710.	4.2	58
22	Mechanistic Insights into Specificity, Activity, and Regulatory Elements of the Regulator of G-protein Signaling (RGS)-containing Rho-specific Guanine Nucleotide Exchange Factors (GEFs) p115, PDZ-RhoGEF (PRG), and Leukemia-associated RhoGEF (LARG). <i>Journal of Biological Chemistry</i> , 2011, 286, 18202-18212.	3.4	58
23	The RAS-Effector Interface: Isoform-Specific Differences in the Effector Binding Regions. <i>PLoS ONE</i> , 2016, 11, e0167145.	2.5	55
24	Subcellular Fractionation and Localization Studies Reveal a Direct Interaction of the Fragile X Mental Retardation Protein (FMRP) with Nucleolin. <i>PLoS ONE</i> , 2014, 9, e91465.	2.5	51
25	Guanine Nucleotide Exchange Factors Operate by a Simple Allosteric Competitive Mechanism. <i>Biochemistry</i> , 2005, 44, 15423-15429.	2.5	49
26	Activating Mutations of RRAS2 Are a Rare Cause of Noonan Syndrome. <i>American Journal of Human Genetics</i> , 2019, 104, 1223-1232.	6.2	43
27	Fluorescence approaches for monitoring interactions of Rho GTPases with nucleotides, regulators, and effectors. <i>Methods</i> , 2005, 37, 173-182.	3.8	42
28	bFGF-mediated pluripotency maintenance in human induced pluripotent stem cells is associated with NRAS-MAPK signaling. <i>Cell Communication and Signaling</i> , 2018, 16, 96.	6.5	38
29	New insight into the molecular switch mechanism of human Rho family proteins: shifting a paradigm. <i>Biological Chemistry</i> , 2013, 394, 89-95.	2.5	35
30	In Vitro GEF and GAP Assays. <i>Current Protocols in Cell Biology</i> , 2009, 43, Unit 14.9.	2.3	34
31	Structural fingerprints, interactions, and signaling networks of RAS family proteins beyond RAS isoforms. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2018, 53, 130-156.	5.2	34
32	Rigosertib potently protects against colitis-associated intestinal fibrosis and inflammation by regulating PI3K/AKT and NF- κ B signaling pathways. <i>Life Sciences</i> , 2020, 249, 117470.	4.3	34
33	Crystal structure of Rnd3/RhoE: functional implications1. <i>FEBS Letters</i> , 2002, 525, 100-104.	2.8	32
34	Accessory proteins of the RAS-MAPK pathway: moving from the side line to the front line. <i>Communications Biology</i> , 2021, 4, 696.	4.4	32
35	Functional Cross-talk between Ras and Rho Pathways. <i>Journal of Biological Chemistry</i> , 2014, 289, 6839-6849.	3.4	31
36	Novel FMRP interaction networks linked to cellular stress. <i>FEBS Journal</i> , 2021, 288, 837-860.	4.7	31

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37	Activating MRAS mutations cause Noonan syndrome associated with hypertrophic cardiomyopathy. <i>Human Molecular Genetics</i> , 2020, 29, 1772-1783.	2.9	30
38	Structural snapshots of RAF kinase interactions. <i>Biochemical Society Transactions</i> , 2018, 46, 1393-1406.	3.4	28
39	Biophysical Characterization of Nucleophosmin Interactions with Human Immunodeficiency Virus Rev and Herpes Simplex Virus US11. <i>PLoS ONE</i> , 2015, 10, e0143634.	2.5	27
40	IQGAP1 Interaction with RHO Family Proteins Revisited. <i>Journal of Biological Chemistry</i> , 2016, 291, 26364-26376.	3.4	26
41	The Role of Embryonic Stem Cell-expressed RAS (ERAS) in the Maintenance of Quiescent Hepatic Stellate Cells. <i>Journal of Biological Chemistry</i> , 2016, 291, 8399-8413.	3.4	26
42	Biochemical Assays to Characterize Rho GTPases. <i>Methods in Molecular Biology</i> , 2012, 827, 37-58.	0.9	24
43	Rho inhibition by lovastatin affects apoptosis and DSB repair of primary human lung cells in vitro and lung tissue in vivo following fractionated irradiation. <i>Cell Death and Disease</i> , 2017, 8, e2978-e2978.	6.3	24
44	Purification and Biochemical Properties of Rac1, 2, 3 and the Splice Variant Rac1b. <i>Methods in Enzymology</i> , 2006, 406, 1-11.	1.0	22
45	Subcellular Localization and Mitotic Interactome Analyses Identify SIRT4 as a Centrosomally Localized and Microtubule Associated Protein. <i>Cells</i> , 2020, 9, 1950.	4.1	19
46	New model for the interaction of IQGAP1 with CDC42 and RAC1. <i>Small GTPases</i> , 2020, 11, 16-22.	1.6	17
47	Interaction characteristics of Plexin-B1 with Rho family proteins. <i>Biochemical and Biophysical Research Communications</i> , 2013, 434, 785-790.	2.1	16
48	The Function of Embryonic Stem Cell-expressed RAS (E-RAS), a Unique RAS Family Member, Correlates with Its Additional Motifs and Its Structural Properties. <i>Journal of Biological Chemistry</i> , 2015, 290, 15892-15903.	3.4	15
49	The intramolecular allostery of GRB2 governing its interaction with SOS1 is modulated by phosphotyrosine ligands. <i>Biochemical Journal</i> , 2021, 478, 2793-2809.	3.7	15
50	From basic researches to new achievements in therapeutic strategies of KRAS-driven cancers. <i>Cancer Biology and Medicine</i> , 2019, 16, 435-461.	3.0	15
51	Aberrant <i>HRAS</i> transcript processing underlies a distinctive phenotype within the RASopathy clinical spectrum. <i>Human Mutation</i> , 2017, 38, 798-804.	2.5	14
52	Fragile X mental retardation protein protects against tumour necrosis factor-mediated cell death and liver injury. <i>Gut</i> , 2020, 69, 133-145.	12.1	14
53	A comprehensive analysis of RAS-effector interactions reveals interaction hotspots and new binding partners. <i>Journal of Biological Chemistry</i> , 2021, 296, 100626.	3.4	14
54	Selective inhibition of IL-6 trans-signaling by a miniaturized, optimized chimeric soluble gp130 inhibits T _H 17 cell expansion. <i>Science Signaling</i> , 2021, 14, .	3.6	13

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55	Liposome Reconstitution and Modulation of Recombinant Prenylated Human Rac1 by GEFs, GDI1 and Pak1. PLoS ONE, 2014, 9, e102425.	2.5	13
56	The Pseudo-Natural Product Rhonin Targets RHO GDI. Angewandte Chemie - International Edition, 2022, 61, .	13.8	11
57	KRAS-related long noncoding RNAs in human cancers. Cancer Gene Therapy, 2022, 29, 418-427.	4.6	8
58	IL-2 Inducible Kinase ITK is Critical for HIV-1 Infection of Jurkat T-cells. Scientific Reports, 2018, 8, 3217.	3.3	7
59	RHO GTPase-Related Long Noncoding RNAs in Human Cancers. Cancers, 2021, 13, 5386.	3.7	7
60	Electrostatic Forces Mediate the Specificity of RHO GTPase-GDI Interactions. International Journal of Molecular Sciences, 2021, 22, 12493.	4.1	6
61	Selectivity Determinants of RHO GTPase Binding to IQGAPs. International Journal of Molecular Sciences, 2021, 22, 12596.	4.1	6
62	Inhibition of the RacGEF VAV3 by the small molecule IODVA1 impedes RAC signaling and overcomes resistance to tyrosine kinase inhibition in acute lymphoblastic leukemia. Leukemia, 2022, 36, 637-647.	7.2	5
63	MRI-based molecular imaging of epicardium-derived stromal cells (EpiSC) by peptide-mediated active targeting. Scientific Reports, 2020, 10, 21669.	3.3	4
64	Spotlight on Accessory Proteins: RTK-RAS-MAPK Modulators as New Therapeutic Targets. Biomolecules, 2021, 11, 895.	4.0	4
65	Allosteric regulation of GRB2 modulates RAS activation. Small GTPases, 2022, 13, 282-286.	1.6	3
66	Physical Interaction between Embryonic Stem Cell-Expressed Ras (ERas) and Arginase-1 in Quiescent Hepatic Stellate Cells. Cells, 2022, 11, 508.	4.1	2