

# Ahmed Sharaf

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3684408/publications.pdf>

Version: 2024-02-01

11  
papers

191  
citations

1163117

8  
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1199594

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g-index

13  
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13  
docs citations

13  
times ranked

357  
citing authors

#	ARTICLE	IF	CITATIONS
1	Distribution of microglia in the postnatal murine nigrostriatal system. <i>Cell and Tissue Research</i> , 2013, 351, 373-382.	2.9	47
2	ApoER2 and VLDLr Are Required for Mediating Reelin Signalling Pathway for Normal Migration and Positioning of Mesencephalic Dopaminergic Neurons. <i>PLoS ONE</i> , 2013, 8, e71091.	2.5	28
3	Proteolytic cleavage of transmembrane cell adhesion molecule L1 by extracellular matrix molecule Reelin is important for mouse brain development. <i>Scientific Reports</i> , 2017, 7, 15268.	3.3	21
4	A Fragment of Adhesion Molecule L1 Binds to Nuclear Receptors to Regulate Synaptic Plasticity and Motor Coordination. <i>Molecular Neurobiology</i> , 2018, 55, 7164-7178.	4.0	19
5	Localization of reelin signaling pathway components in murine midbrain and striatum. <i>Cell and Tissue Research</i> , 2015, 359, 393-407.	2.9	18
6	Systematic Affinity Purification Coupled to Mass Spectrometry Identified p62 as Part of the Cannabinoid Receptor CB2 Interactome. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 224.	2.9	15
7	The WNT1G177C mutation specifically affects skeletal integrity in a mouse model of osteogenesis imperfecta type XV. <i>Bone Research</i> , 2021, 9, 48.	11.4	13
8	Cardiovascular Complications in Diabetic Patients Undergoing Regular Hemodialysis. <i>Angiology</i> , 2015, 66, 225-230.	1.8	10
9	Mice Carrying a Ubiquitous <sc>R235W</sc> Mutation of <i>Wnt1</i> Display a Boneâ€Œspecific Phenotype. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 1726-1737.	2.8	8
10	Reelin controls the positioning of brainstem serotonergic raphe neurons. <i>PLoS ONE</i> , 2018, 13, e0200268.	2.5	6
11	A rare heterozygous <i>TREM2</i> coding variant identified in familial clustering of dementia affects an intrinsically disordered protein region and function of TREM2. <i>Human Mutation</i> , 2020, 41, 169-181.	2.5	4