

# Davide Comoletti

## List of Publications by Year in descending order

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Version: 2024-02-01

36  
papers

2,601  
citations

331670

21  
h-index

361022

35  
g-index

37  
all docs

37  
docs citations

37  
times ranked

3208  
citing authors

#	ARTICLE	IF	CITATIONS
1	Shed CNTNAP2 ectodomain is detectable in CSF and regulates Ca <sup>2+</sup> homeostasis and network synchrony via PMCA2/ATP2B2. <i>Neuron</i> , 2022, 110, 627-643.e9.	8.1	17
2	Structure of Reelin repeat 8 and the adjacent C-terminal region. <i>Biophysical Journal</i> , 2022, 121, 2526-2537.	0.5	0
3	Case Report: Is Catatonia a Clinical Feature of the Natural Progression of NLGN2-Related Neurodevelopmental Disorder?. <i>Journal of Autism and Developmental Disorders</i> , 2021, 51, 371-376.	2.7	5
4	Comparative mapping of selected structural determinants on the extracellular domains of cholinesterase-like cell-adhesion molecules. <i>Neuropharmacology</i> , 2021, 184, 108381.	4.1	4
5	The structure-function relationship of a signaling-competent, dimeric Reelin fragment. <i>Structure</i> , 2021, 29, 1156-1170.e6.	3.3	6
6	In trans neuregulin3-Caspr3 interaction controls DA axonal bassoon cluster development. <i>Current Biology</i> , 2021, 31, 3330-3342.e7.	3.9	2
7	Purification of a heterodimeric Reelin construct to investigate binding stoichiometry. <i>European Biophysics Journal</i> , 2020, 49, 773-779.	2.2	3
8	Synapse type-specific proteomic dissection identifies IgSF8 as a hippocampal CA3 microcircuit organizer. <i>Nature Communications</i> , 2020, 11, 5171.	12.8	35
9	In utero exposure to endogenous maternal polyclonal anti-Caspr2 antibody leads to behavioral abnormalities resembling autism spectrum disorder in male mice. <i>Scientific Reports</i> , 2020, 10, 14446.	3.3	12
10	A Proteomic Screen of Neuronal Cell-Surface Molecules Reveals IgLONs as Structurally Conserved Interaction Modules at the Synapse. <i>Structure</i> , 2019, 27, 893-906.e9.	3.3	44
11	Secreted amyloid- $\beta$ precursor protein functions as a GABA <sub>B</sub> R1a ligand to modulate synaptic transmission. <i>Science</i> , 2019, 363, .	12.6	205
12	An ELISA-Based Screening Platform for Ligand-Receptor Discovery. <i>Methods in Enzymology</i> , 2019, 615, 453-475.	1.0	18
13	P1 $\beta$ : SECRETED AMYLOID PRECURSOR PROTEIN IS A GABABR1A LIGAND THAT SUPPRESSES SYNAPTIC VESICLE RELEASE. <i>Alzheimer's and Dementia</i> , 2018, 14, P349.	0.8	1
14	Structural Insights into Reelin Function: Present and Future. <i>Frontiers in Cellular Neuroscience</i> , 2016, 10, 137.	3.7	27
15	Super-complexes of adhesion GPCRs and neural guidance receptors. <i>Nature Communications</i> , 2016, 7, 11184.	12.8	84
16	Structural Characterization of the Extracellular Domain of CASPR2 and Insights into Its Association with the Novel Ligand Contactin1. <i>Journal of Biological Chemistry</i> , 2016, 291, 5788-5802.	3.4	29
17	Autism-associated R451C mutation in neuroligin3 leads to activation of the unfolded protein response in a PC12 Tet-On inducible system. <i>Biochemical Journal</i> , 2016, 473, 423-434.	3.7	37
18	Structural and Mechanistic Insights into the Latrophilin3-FLRT3 Complex that Mediates Glutamatergic Synapse Development. <i>Structure</i> , 2015, 23, 1665-1677.	3.3	42

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19	LRP8-Reelin-Regulated Neuronal Enhancer Signature Underlying Learning and Memory Formation. <i>Neuron</i> , 2015, 86, 696-710.	8.1	130
20	Reelin Induces Erk1/2 Signaling in Cortical Neurons Through a Non-canonical Pathway. <i>Journal of Biological Chemistry</i> , 2014, 289, 20307-20317.	3.4	49
21	Ecto-Fc MS identifies ligand-receptor interactions through extracellular domain Fc fusion protein baits and shotgun proteomic analysis. <i>Nature Protocols</i> , 2014, 9, 2061-2074.	12.0	21
22	Processing of Cholinesterase-like &#945;/&#946;-Hydrolase Fold Proteins: Alterations Associated with Congenital Disorders. <i>Protein and Peptide Letters</i> , 2012, 19, 173-179.	0.9	8
23	FLRT Proteins Are Endogenous Latrophilin Ligands and Regulate Excitatory Synapse Development. <i>Neuron</i> , 2012, 73, 903-910.	8.1	221
24	The Crystal Structure of the Î±-Neurexin-1 Extracellular Region Reveals a Hinge Point for Mediating Synaptic Adhesion and Function. <i>Structure</i> , 2011, 19, 767-778.	3.3	56
25	The Macromolecular Architecture of Extracellular Domain of Î±NRXN1: Domain Organization, Flexibility, and Insights into Trans-Synaptic Disposition. <i>Structure</i> , 2010, 18, 1044-1053.	3.3	30
26	Structureâ€“function relationships of the Î±/Î²-hydrolase fold domain of neuroligin: A comparison with acetylcholinesterase. <i>Chemico-Biological Interactions</i> , 2010, 187, 49-55.	4.0	10
27	LRRTM2 Interacts with Neurexin1 and Regulates Excitatory Synapse Formation. <i>Neuron</i> , 2009, 64, 799-806.	8.1	338
28	Characterization of the solution structure of a neuroligin/Î²-neurexin complex. <i>Chemico-Biological Interactions</i> , 2008, 175, 150-155.	4.0	8
29	Structural Analysis of the Synaptic Protein Neuroligin and Its Î²-Neurexin Complex: Determinants for Folding and Cell Adhesion. <i>Neuron</i> , 2007, 56, 979-991.	8.1	142
30	Synaptic Arrangement of the Neuroligin/Î²-Neurexin Complex Revealed by X-Ray and Neutron Scattering. <i>Structure</i> , 2007, 15, 693-705.	3.3	64
31	Gene Selection, Alternative Splicing, and Post-translational Processing Regulate Neuroligin Selectivity for Î²-Neurexinsâ€™. <i>Biochemistry</i> , 2006, 45, 12816-12827.	2.5	117
32	A Mutation Linked with Autism Reveals a Common Mechanism of Endoplasmic Reticulum Retention for the Î±,Î²-Hydrolase Fold Protein Family. <i>Journal of Biological Chemistry</i> , 2006, 281, 9667-9676.	3.4	53
33	A Splice Code for trans-Synaptic Cell Adhesion Mediated by Binding of Neuroligin 1 to Î±- and Î²-Neurexins. <i>Neuron</i> , 2005, 48, 229-236.	8.1	416
34	The Arg451Cys-Neuroligin-3 Mutation Associated with Autism Reveals a Defect in Protein Processing. <i>Journal of Neuroscience</i> , 2004, 24, 4889-4893.	3.6	214
35	Structural Characterization of Recombinant Soluble Rat Neuroligin 1:â€™ Mapping of Secondary Structure and Glycosylation by Mass Spectrometry. <i>Biochemistry</i> , 2004, 43, 1496-1506.	2.5	41
36	Characterization of the Interaction of a Recombinant Soluble Neuroligin-1 with Neurexin-1Î². <i>Journal of Biological Chemistry</i> , 2003, 278, 50497-50505.	3.4	111