List of Publications by Year in descending order

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FELLY HALLSCH

#	Article	IF	CITATIONS
1	FK506 Binding Protein 5 Shapes Stress Responsiveness: Modulation of Neuroendocrine Reactivity and Coping Behavior. Biological Psychiatry, 2011, 70, 928-936.	1.3	235
2	The involvement of FK506-binding protein 51 (FKBP5) in the behavioral and neuroendocrine effects of chronic social defeat stress. Neuropharmacology, 2012, 62, 332-339.	4.1	195
3	Epigenetic upregulation of FKBP5 by aging and stress contributes to NF-ήB–driven inflammation and cardiovascular risk. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 11370-11379.	7.1	193
4	Selective inhibitors of the FK506-binding protein 51 by induced fit. Nature Chemical Biology, 2015, 11, 33-37.	8.0	188
5	FKBP Ligands—Where We Are and Where to Go?. Frontiers in Pharmacology, 2018, 9, 1425.	3.5	110
6	Large FK506-Binding Proteins Shape the Pharmacology of Rapamycin. Molecular and Cellular Biology, 2013, 33, 1357-1367.	2.3	106
7	A C-terminal HSP90 inhibitor restores glucocorticoid sensitivity and relieves a mouse allograft model of Cushing disease. Nature Medicine, 2015, 21, 276-280.	30.7	92
8	Pharmacological Inhibition of the Psychiatric Risk Factor FKBP51 Has Anxiolytic Properties. Journal of Neuroscience, 2015, 35, 9007-9016.	3.6	90
9	The Prospect of FKBP51 as a Drug Target. ChemMedChem, 2012, 7, 1351-1359.	3.2	86
10	The stress regulator FKBP51 drives chronic pain by modulating spinal glucocorticoid signaling. Science Translational Medicine, 2016, 8, 325ra19.	12.4	82
11	Stress-responsive FKBP51 regulates AKT2-AS160 signaling and metabolic function. Nature Communications, 2017, 8, 1725.	12.8	82
12	The Many Faces of FKBP51. Biomolecules, 2019, 9, 35.	4.0	79
13	FKBPs and the Akt/mTOR pathway. Cell Cycle, 2013, 12, 2366-2370.	2.6	75
14	Fluorescent Probes to Characterise FK506â€Binding Proteins. ChemBioChem, 2009, 10, 1402-1410.	2.6	73
15	A regulatory role for the co-chaperone FKBP51s in PD-L1 expression in glioma. Oncotarget, 2017, 8, 68291-68304.	1.8	71
16	FKBP51 employs both scaffold and isomerase functions to promote NF-κB activation in melanoma. Nucleic Acids Research, 2015, 43, 6983-6993.	14.5	68
17	Orexin–Corticotropin-Releasing Factor Receptor Heteromers in the Ventral Tegmental Area as Targets for Cocaine. Journal of Neuroscience, 2015, 35, 6639-6653.	3.6	66
18	Hippocampal neuroligin-2 links early-life stress with impaired social recognition and increased aggression in adult mice. Psychoneuroendocrinology, 2015, 55, 128-143.	2.7	63

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19	FKBP5 expression in human adipose tissue: potential role in glucose and lipid metabolism, adipogenesis and type 2 diabetes. Endocrine, 2018, 62, 116-128.	2.3	63
20	Evaluation of Synthetic FK506 Analogues as Ligands for the FK506-Binding Proteins 51 and 52. Journal of Medicinal Chemistry, 2012, 55, 4114-4122.	6.4	59
21	Structural characterization of the PPIase domain of FKBP51, a cochaperone of human Hsp90. Acta Crystallographica Section D: Biological Crystallography, 2011, 67, 549-559.	2.5	57
22	Increasing the Efficiency of Ligands for FK506-Binding Protein 51 by Conformational Control. Journal of Medicinal Chemistry, 2013, 56, 3922-3935.	6.4	53
23	Stress-primed secretory autophagy promotes extracellular BDNF maturation by enhancing MMP9 secretion. Nature Communications, 2021, 12, 4643.	12.8	50
24	Exploration of Pipecolate Sulfonamides as Binders of the FK506-Binding Proteins 51 and 52. Journal of Medicinal Chemistry, 2012, 55, 4123-4131.	6.4	46
25	The stress regulator FKBP51: a novel and promising druggable target for the treatment of persistent pain states across sexes. Pain, 2018, 159, 1224-1234.	4.2	46
26	Chemogenomic Profiling of Human and Microbial FK506-Binding Proteins. Journal of Medicinal Chemistry, 2018, 61, 3660-3673.	6.4	42
27	Crystal Structures of the Free and Ligand-Bound FK1–FK2 Domain Segment of FKBP52 Reveal a Flexible Inter-Domain Hinge. Journal of Molecular Biology, 2013, 425, 4134-4144.	4.2	41
28	Deficiency of <scp>FK</scp> 506â€binding protein ( <scp>FKBP</scp> ) 51 alters sleep architecture and recovery sleep responses to stress in mice. Journal of Sleep Research, 2014, 23, 176-185.	3.2	41
29	InterAKTions with FKBPs - Mutational and Pharmacological Exploration. PLoS ONE, 2013, 8, e57508.	2.5	39
30	The Hsp90 machinery facilitates the transport of diphtheria toxin into human cells. Scientific Reports, 2017, 7, 613.	3.3	36
31	Structure–Affinity Relationship Analysis of Selective FKBP51 Ligands. Journal of Medicinal Chemistry, 2015, 58, 7796-7806.	6.4	32
32	FKBPs and their role in neuronal signaling. Biochimica Et Biophysica Acta - General Subjects, 2015, 1850, 2035-2040.	2.4	31
33	Rapid, Structure-Based Exploration of Pipecolic Acid Amides as Novel Selective Antagonists of the FK506-Binding Protein 51. Journal of Medicinal Chemistry, 2016, 59, 2410-2422.	6.4	31
34	Pharmacological Modulation of the Psychiatric Risk Factor FKBP51 Alters Efficiency of Common Antidepressant Drugs. Frontiers in Behavioral Neuroscience, 2018, 12, 262.	2.0	29
35	Structure-Based Design of High-Affinity Macrocyclic FKBP51 Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 3320-3349.	6.4	28
36	Rational Design and Asymmetric Synthesis of Potent and Neurotrophic Ligands for FK506â€Binding Proteins (FKBPs). Angewandte Chemie - International Edition, 2015, 54, 345-348.	13.8	27

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37	Modulating FKBP5/FKBP51 and autophagy lowers HTT (huntingtin) levels. Autophagy, 2021, 17, 4119-4140.	9.1	27
38	Clues to molecular glues. Current Research in Chemical Biology, 2022, 2, 100018.	2.9	27
39	Stereoselective Construction of the 5-Hydroxy Diazabicyclo[4.3.1]decane-2-one Scaffold, a Privileged Motif for FK506-Binding Proteins. Organic Letters, 2014, 16, 5254-5257.	4.6	26
40	Targeting the Glucocorticoid Receptor Reduces Bingeâ€Like Drinking in High Drinking in the Dark (HDIDâ€1) Mice. Alcoholism: Clinical and Experimental Research, 2020, 44, 1025-1036.	2.4	26
41	Recent Progress in FKBP Ligand Development. Current Molecular Pharmacology, 2015, 9, 27-36.	1.5	22
42	Structural and Dynamical Basis of G Protein Inhibition by YM-254890 and FR900359: An Inhibitor in Action. Journal of Chemical Information and Modeling, 2019, 59, 4361-4373.	5.4	22
43	The selective FKBP51 inhibitor SAFit2 reduces alcohol consumption and reinstatement of conditioned alcohol effects in mice. Addiction Biology, 2020, 25, e12758.	2.6	21
44	Identification of phenothiazine derivatives as UHM-binding inhibitors of early spliceosome assembly. Nature Communications, 2020, 11, 5621.	12.8	20
45	Picomolar FKBP inhibitors enabled by a single water-displacing methyl group in bicyclic [4.3.1] aza-amides. Chemical Science, 2021, 12, 14758-14765.	7.4	19
46	Synthesis and Neurotrophic Activity Studies of <i>Illicium</i> Sesquiterpene Natural Product Analogues. Chemistry - A European Journal, 2017, 23, 3178-3183.	3.3	18
47	Selective Inhibitors of FKBP51 Employ Conformational Selection of Dynamic Invisible States. Angewandte Chemie - International Edition, 2019, 58, 9429-9433.	13.8	18
48	Betablockers at Work: The Crystal Structure of the β <sub>2</sub> â€Adrenergic Receptor. Angewandte Chemie - International Edition, 2008, 47, 3314-3316.	13.8	16
49	The Seven Pillars of Molecular Pharmacology: GPCR Research Honored with Nobel Prize for Chemistry. Angewandte Chemie - International Edition, 2012, 51, 12172-12175.	13.8	16
50	The splicing FK506-binding protein-51 isoform plays a role in glioblastoma resistance through programmed cell death ligand-1 expression regulation. Cell Death Discovery, 2019, 5, 137.	4.7	14
51	Initial Metabolic Step of a Novel Ethanolamine Utilization Pathway and Its Regulation in <i>Streptomyces coelicolor</i> M145. MBio, 2019, 10, .	4.1	13
52	Sex differences in the effect of the FKBP5 inhibitor SAFit2 on anxiety and stress-induced reinstatement following cocaine self-administration. Neurobiology of Stress, 2020, 13, 100232.	4.0	13
53	Macrocyclic FKBP51 Ligands Define a Transient Binding Mode with Enhanced Selectivity. Angewandte Chemie - International Edition, 2021, 60, 13257-13263.	13.8	13
54	Development of NanoBRETâ€Binding Assays for FKBP‣igand Profiling in Living Cells. ChemBioChem, 2021, 22, 2257-2261.	2.6	12

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55	Facile Synthesis of a Fluorescent Cyclosporin A Analogue To Study Cyclophilin 40 and Cyclophilin 18 Ligands. ACS Medicinal Chemistry Letters, 2010, 1, 536-539.	2.8	11
56	Structures of Classâ€B G Proteinâ€Coupled Receptors: Prospects for Drug Discovery. Angewandte Chemie - International Edition, 2013, 52, 12783-12785.	13.8	9
57	A Novel Decalin-Based Bicyclic Scaffold for FKBP51-Selective Ligands. Journal of Medicinal Chemistry, 2020, 63, 231-240.	6.4	9
58	Duration of Reduction in Enduring Stress-Induced Hyperalgesia Via FKBP51 Inhibition Depends on Timing of Administration Relative to Traumatic Stress Exposure. Journal of Pain, 2022, 23, 1256-1267.	1.4	7
59	Fenton-Chemistry-Based Oxidative Modification of Proteins Reflects Their Conformation. International Journal of Molecular Sciences, 2021, 22, 9927.	4.1	6
60	Azidobupramine, an Antidepressant-Derived Bifunctional Neurotransmitter Transporter Ligand Allowing Covalent Labeling and Attachment of Fluorophores. PLoS ONE, 2016, 11, e0148608.	2.5	5
61	FKBP51 and FKBP12.6—Novel and tight interactors of Glomulin. PLoS ONE, 2019, 14, e0221926.	2.5	5
62	Selective Inhibitors of FKBP51 Employ Conformational Selection of Dynamic Invisible States. Angewandte Chemie, 2019, 131, 9529-9533.	2.0	5
63	Enantioselective Synthesis of a Tricyclic, sp 3 â€Rich Diazatetradecanedione: an Amino Acidâ€Based Natural Productâ€Like Scaffold. Chemistry - A European Journal, 2020, 26, 4677-4681.	3.3	5
64	Cryoâ€EM Structures of Class B GPCR Reveal the Activation Mechanism. Angewandte Chemie - International Edition, 2017, 56, 12412-12414.	13.8	3
65	Med Chem Remote: The Frontiers in Medicinal Chemistry 2021. ChemMedChem, 2021, 16, 2411-2416.	3.2	1
66	Break Away: FKBP12 sequestration as a target for increasing BMP activity. Cell Chemical Biology, 2021, 28, 1253-1255.	5.2	1
67	Typâ€Bâ€GPCRâ€&trukturen verdeutlichen Aktivierungsmechanismus. Angewandte Chemie, 2017, 129, 12584-12586.	2.0	0
68	Makrozyklische FKBP51â€Liganden enthüllen einen transienten Bindungsmodus mit erhöhter Selektivitä Angewandte Chemie, 2021, 133, 13366-13372.	2.0	0