

Brian F Thomas

List of Publications by Year in descending order

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75
papers

3,504
citations

159358

30
h-index

143772

57
g-index

77
all docs

77
docs citations

77
times ranked

3358
citing authors

#	ARTICLE	IF	CITATIONS
1	Bisphenol A Exposure Causes Meiotic Aneuploidy in the Female Mouse. <i>Current Biology</i> , 2003, 13, 546-553.	1.8	575
2	Behavioral, biochemical, and molecular modeling evaluations of cannabinoid analogs. <i>Pharmacology Biochemistry and Behavior</i> , 1991, 40, 471-478.	1.3	384
3	Safety and pharmacokinetics of purified soy isoflavones: single-dose administration to postmenopausal women,,,. <i>American Journal of Clinical Nutrition</i> , 2002, 76, 1126-1137.	2.2	162
4	Potent rewarding and reinforcing effects of the synthetic cathinone 3,4- ϵ -methylenedioxypropylvalerone (<sc>MDPV</sc>). <i>Addiction Biology</i> , 2014, 19, 165-174.	1.4	156
5	AB-CHMINACA, AB-PINACA, and FUBIMINA: Affinity and Potency of Novel Synthetic Cannabinoids in Producing δ^9 -Tetrahydrocannabinol-Like Effects in Mice. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2015, 354, 328-339.	1.3	110
6	Structure-Activity Analysis of Anandamide Analogs: A Relationship to a Cannabinoid Pharmacophore. <i>Journal of Medicinal Chemistry</i> , 1996, 39, 471-479.	2.9	101
7	Synthesis and Structure-Activity Relationships of Amide and Hydrazone Analogues of the Cannabinoid CB1 Receptor Antagonist N-(Piperidinyl)-5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-4-methyl-1H-pyrazole-3-carboxamide (SR141716). <i>Journal of Medicinal Chemistry</i> , 2002, 45, 2708-2719.	2.9	94
8	Cannabinoids in disguise: δ^9 -Tetrahydrocannabinol-like effects of tetramethylcyclopropyl ketone indoles. <i>Neuropharmacology</i> , 2013, 75, 145-154.	2.0	94
9	Analysis of Synthetic Cannabinoids Using High-Resolution Mass Spectrometry and Mass Defect Filtering: Implications for Nontargeted Screening of Designer Drugs. <i>Analytical Chemistry</i> , 2012, 84, 5574-5581.	3.2	91
10	Allosteric Modulation: An Alternate Approach Targeting the Cannabinoid CB1 Receptor. <i>Medicinal Research Reviews</i> , 2017, 37, 441-474.	5.0	76
11	Quantitative analysis of the principle soy isoflavones genistein, daidzein and glycitein, and their primary conjugated metabolites in human plasma and urine using reversed-phase high-performance liquid chromatography with ultraviolet detection. <i>Biomedical Applications</i> , 2001, 760, 191-205.	1.7	70
12	Dose-Response Effects of Spectrum Research Cigarettes. <i>Nicotine and Tobacco Research</i> , 2013, 15, 1113-1121.	1.4	69
13	Molecular and Behavioral Pharmacological Characterization of Abused Synthetic Cannabinoids MMB- and MDMB-FUBINACA, MN-18, NNE1, CUMYL-PICA, and 5-Fluoro-CUMYL-PICA. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2018, 365, 437-446.	1.3	69
14	Overcoming the Psychiatric Side Effects of the Cannabinoid CB1 Receptor Antagonists: Current Approaches for Therapeutics Development. <i>Current Topics in Medicinal Chemistry</i> , 2019, 19, 1418-1435.	1.0	69
15	Synthesis and Pharmacological Comparison of Dimethylheptyl and Pentyl Analogs of Anandamide. <i>Journal of Medicinal Chemistry</i> , 1997, 40, 3626-3634.	2.9	63
16	Synthesis and Biological Evaluation of Bivalent Ligands for the Cannabinoid 1 Receptor. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 7048-7060.	2.9	62
17	The Bioactive Conformation of Aminoalkylindoles at the Cannabinoid CB1 and CB2 Receptors: Insights Gained from (E)- and (Z)-Naphthylidene Indenes. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 5177-5187.	2.9	60
18	Peripherally Selective Cannabinoid 1 Receptor (CB1R) Agonists for the Treatment of Neuropathic Pain. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7525-7543.	2.9	53

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19	Relationship between the biodisposition of [3H]soman and its pharmacological effects in mice. Toxicology and Applied Pharmacology, 1985, 80, 409-420.	1.3	49
20	Evaluation of first generation synthetic cannabinoids on binding at non-cannabinoid receptors and in a battery of in vivo assays in mice. Neuropharmacology, 2016, 110, 143-153.	2.0	49
21	Hijacking of Basic Research: The Case of Synthetic Cannabinoids. , 2011, 2011, .		45
22	Thermolytic Degradation of Synthetic Cannabinoids: Chemical Exposures and Pharmacological Consequences. Journal of Pharmacology and Experimental Therapeutics, 2017, 361, 162-171.	1.3	41
23	Diarylureas as Allosteric Modulators of the Cannabinoid CB1 Receptor: Structure-Activity Relationship Studies on 1-(4-Chlorophenyl)-3-{3-[6-(pyrrolidin-1-yl)pyridin-2-yl]phenyl}urea (PSNCBAM-1). Journal of Medicinal Chemistry, 2014, 57, 7758-7769.	2.9	40
24	Structure-activity relationships for 1 ¹ ,1 ² -dimethylalkyl- δ 8 -tetrahydrocannabinols. Bioorganic and Medicinal Chemistry, 2003, 11, 1397-1410.	1.4	38
25	Kappa opioid mediation of cannabinoid effects of the potent hallucinogen, salvinorin A, in rodents. Psychopharmacology, 2010, 210, 275-284.	1.5	37
26	Substituted Tetrahydroisoquinolines as Selective Antagonists for the Orexin 1 Receptor. Journal of Medicinal Chemistry, 2013, 56, 6901-6916.	2.9	36
27	In vitro and in vivo pharmacokinetics and metabolism of synthetic cannabinoids CUMYL-PICA and 5F-CUMYL-PICA. Forensic Toxicology, 2017, 35, 333-347.	1.4	35
28	Cannabinoid CB1 receptor antagonists as potential pharmacotherapies for drug abuse disorders. International Review of Psychiatry, 2009, 21, 134-142.	1.4	33
29	Vaping Synthetic Cannabinoids: A Novel Preclinical Model of E-Cigarette Use in Mice. Substance Abuse: Research and Treatment, 2017, 11, 117822181770173.	0.5	33
30	Synthetic Cannabinoid Hydroxypentyl Metabolites Retain Efficacy at Human Cannabinoid Receptors. Journal of Pharmacology and Experimental Therapeutics, 2019, 368, 414-422.	1.3	33
31	Use of SPME-HS-GC-MS for the Analysis of Herbal Products Containing Synthetic Cannabinoids. Journal of Analytical Toxicology, 2012, 36, 293-302.	1.7	32
32	Structure-activity relationships of substituted 1H-indole-2-carboxamides as CB1 receptor allosteric modulators. Bioorganic and Medicinal Chemistry, 2015, 23, 2195-2203.	1.4	31
33	QSAR Analysis of δ -THC Analogues: Relationship of Side-Chain Conformation to Cannabinoid Receptor Affinity and Pharmacological Potency. Journal of Medicinal Chemistry, 2000, 43, 59-70.	2.9	30
34	Finding order in chemical chaos - Continuing characterization of synthetic cannabinoid receptor agonists. Neuropharmacology, 2018, 134, 73-81.	2.0	29
35	Combination Chemistry: Structure-Activity Relationships of Novel Psychoactive Cannabinoids. Current Topics in Behavioral Neurosciences, 2016, 32, 231-248.	0.8	28
36	Toxic by design? Formation of thermal degradants and cyanide from carboxamide-type synthetic cannabinoids CUMYL-PICA, 5F-CUMYL-PICA, AMB-FUBINACA, MDMB-FUBINACA, NNEI, and MN-18 during exposure to high temperatures. Forensic Toxicology, 2019, 37, 17-26.	1.4	28

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37	Synthesis of long-chain amide analogs of the cannabinoid CB1 receptor antagonist N-(piperidinyl)-5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-4-methyl-1H-pyrazole-3-carboxamide (SR141716) with unique binding selectivities and pharmacological activities. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 5463-5474.	1.4	27
38	Toward the Development of Bivalent Ligand Probes of Cannabinoid CB1 and Orexin OX1 Receptor Heterodimers. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 634-638.	1.3	25
39	Synthesis and in vivo studies of a selective ligand for the dopamine transporter: 3 ^Î -(4-[125I]iodophenyl) tropan-2 ^Î -carboxylic acid isopropyl ester ([125I]RTM-21). <i>Nuclear Medicine and Biology</i> , 1996, 23, 277-284.	0.3	24
40	The great divide: Separation between inÂvitro and inÂvivo effects of PSNCBAM-based CB 1 receptor allosteric modulators. <i>Neuropharmacology</i> , 2017, 125, 365-375.	2.0	23
41	Novel Diarylurea Based Allosteric Modulators of the Cannabinoid CB1 Receptor: Evaluation of Importance of 6-Pyrrolidinylpyridinyl Substitution. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 7410-7424.	2.9	21
42	Identification of Eight Synthetic Cannabinoids, Including 5Fâ€AKB48 in Seized Herbal Products Using DARTâ€TOFâ€MS and LCâ€QTOFâ€MS as Nontargeted Screening Methods. <i>Journal of Forensic Sciences</i> , 2017, 62, 1151-1158.	0.9	20
43	Determination of ibogaine in plasma by gas chromatography-chemical ionization mass spectrometry. <i>Journal of Chromatography A</i> , 1996, 723, 101-109.	1.8	19
44	The Importance of Hydrogen Bonding and Aromatic Stacking to the Affinity and Efficacy of Cannabinoid Receptor CB ₂ Antagonist, 5-(4-chloro-3-methylphenyl)-1-[(4-methylphenyl)methyl]-N-[(1 <i>S</i> ,2 <i>S</i> ,4 <i>R</i>)-1,3,3-trimethylbicyclo[2.2.1]hept-2-yl] (SR144528). <i>Journal of Medicinal Chemistry</i> , 2013, 56, 6593-6612.	2.9	19
45	Physical design analysis and mainstream smoke constituent yields of the new potential reduced exposure product, Marlboro UltraSmooth. <i>Nicotine and Tobacco Research</i> , 2007, 9, 1197-1206.	1.4	18
46	Conformationally Constrained Analogues of N-(Piperidinyl)-5-(4-Chlorophenyl)-1-(2,4-) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 387 Td (Dic Analysis, And Biological Evaluations. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 3526-3539.	2.9	18
47	Truncated Orexin Peptides: Structureâ€Activity Relationship Studies. <i>ACS Medicinal Chemistry Letters</i> , 2013, 4, 1224-1227.	1.3	18
48	Indenopyridine derivative RTI-4587-073(l): A candidate for male contraception in stallions. <i>Theriogenology</i> , 2013, 80, 1006-1016.	0.9	17
49	The importance of the 6- and 7-positions of tetrahydroisoquinolines as selective antagonists for the orexin 1 receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5709-5724.	1.4	17
50	Structure elucidation of a novel ring-constrained biaryl pyrazole CB1 cannabinoid receptor antagonist. <i>Magnetic Resonance in Chemistry</i> , 2003, 41, 265-268.	1.1	16
51	Novel, potent THC/anandamide (hybrid) analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 7850-7864.	1.4	16
52	Diaryl urea analogues of SB-334867 as orexin-1 receptor antagonists. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 2980-2985.	1.0	16
53	Comparison of cigarette, little cigar, and waterpipe tobacco smoke condensate and e-cigarette aerosol condensate in a self-administration model. <i>Behavioural Brain Research</i> , 2019, 372, 112061.	1.2	16
54	Conformational characteristics of the interaction of SR141716A with the CB1 cannabinoid receptor as determined through the use of conformationally constrained analogs. <i>AAPS Journal</i> , 2006, 8, E665-E671.	2.2	15

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55	Effect of 1-Substitution on Tetrahydroisoquinolines as Selective Antagonists for the Orexin-1 Receptor. <i>ACS Chemical Neuroscience</i> , 2015, 6, 599-614.	1.7	14
56	The Spicy Story of Cannabimimetic Indoles. <i>Molecules</i> , 2021, 26, 6190.	1.7	13
57	Tricyclic Pyrazoles. Part 5. Novel 1,4-Dihydroindeno[1,2-]pyrazole CB2 Ligands Using Molecular Hybridization Based on Scaffold Hopping. <i>Open Medicinal Chemistry Journal</i> , 2012, 6, 1-14.	0.9	12
58	Synthesis and Pharmacological Evaluation of 1-Phenyl-3-Thiophenylurea Derivatives as Cannabinoid Type-1 Receptor Allosteric Modulators. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 9806-9823.	2.9	12
59	Assessment of structural commonality between tetrahydrocannabinol and anandamide. <i>European Journal of Pharmacology</i> , 2002, 435, 35-42.	1.7	11
60	Structural analogs of pyrazole and sulfonamide cannabinoids: Effects on acute food intake in mice. <i>European Journal of Pharmacology</i> , 2012, 695, 62-70.	1.7	11
61	Kinetic and metabolic profiles of synthetic cannabinoids NNE1 and MN-18. <i>Drug Testing and Analysis</i> , 2018, 10, 137-147.	1.6	11
62	In vitro and in vivo pharmacological evaluation of the synthetic cannabinoid receptor agonist EG-018. <i>Pharmacology Biochemistry and Behavior</i> , 2020, 193, 172918.	1.3	11
63	Analytical surveillance of emerging drugs of abuse and drug formulations. <i>Life Sciences</i> , 2013, 92, 512-519.	2.0	10
64	Medullary Endocannabinoids Contribute to the Differential Resting Baroreflex Sensitivity in Rats with Altered Brain Renin-Angiotensin System Expression. <i>Frontiers in Physiology</i> , 2016, 7, 207.	1.3	9
65	Diarylureas Containing 5-Membered Heterocycles as CB ₁ Receptor Allosteric Modulators: Design, Synthesis, and Pharmacological Evaluation. <i>ACS Chemical Neuroscience</i> , 2019, 10, 518-527.	1.7	8
66	Modification of phencyclidine intoxication and biodisposition by charcoal and other treatments. <i>Pharmacology Biochemistry and Behavior</i> , 1988, 30, 371-377.	1.3	6
67	Interactions of Cannabinoids With Biochemical Substrates. <i>Substance Abuse: Research and Treatment</i> , 2017, 11, 117822181771141.	0.5	6
68	Complete ¹ H and ¹³ C assignments of fluorinated analogs of dehydroepiandrosterone. <i>Magnetic Resonance in Chemistry</i> , 2006, 44, 1051-1053.	1.1	5
69	Neuroanatomical basis for therapeutic applications of cannabinoid receptor 1 antagonists. <i>Drug Development Research</i> , 2009, 70, 527-554.	1.4	5
70	Pharmacokinetic and tissue distribution study of [¹⁴ C]fluasterone in male Beagle dogs following intravenous, oral and subcutaneous dosing routes. <i>Chemico-Biological Interactions</i> , 2010, 183, 317-326.	1.7	5
71	Identification of [¹⁴ C]Fluasterone Metabolites in Urine and Feces Collected from Dogs after Subcutaneous and Oral Administration of [¹⁴ C]Fluasterone. <i>Drug Metabolism and Disposition</i> , 2009, 37, 1089-1097.	1.7	3
72	Cannabinoid CB1 receptor antagonists. <i>Drug Development Research</i> , 2009, 70, 525-526.	1.4	1

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73	Structure-Activity Relationships and Conformational Freedom of CB1 Receptor Antagonists and Inverse Agonists. , 2009, , 95-119.		1
74	Determination of l- \pm -acetylmethadol, l- \pm -noracetylmethadol and l- \pm -dinoracetylmethadol in plasma by gas chromatography-mass spectrometry. Biomedical Applications, 1994, 655, 201-211.	1.7	0
75	Conformational Characteristics of the Interaction of SR141716A with the CB1 Cannabinoid Receptor as Determined Through the Use of Conformationally Constrained Analogs. , 2008, , 707-718.		0