F Ivy Carroll

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

9,363 82 289 52 h-index g-index citations papers 10,064 295 5.2 5.71 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
289	Interactions between 2Qfluoro-(carbamoylpyridinyl)deschloroepibatidine analogues and acetylcholine-binding protein inform on potent antagonist activity against nicotinic receptors <i>Acta Crystallographica Section D: Structural Biology</i> , 2022 , 78, 353-362	5.5	O
288	Blockade of kappa-opioid receptors amplifies microglia-mediated inflammatory responses. <i>Pharmacology Biochemistry and Behavior</i> , 2021 , 173301	3.9	O
287	Designer drugs: a medicinal chemistry perspective (II). <i>Annals of the New York Academy of Sciences</i> , 2021 , 1489, 48-77	6.5	1
286	Nanobody-enabled monitoring of kappa opioid receptor states. <i>Nature Communications</i> , 2020 , 11, 1145	17.4	50
285	Effects of chronic treatment with bupropion on self-administration of nicotine + cocaine mixtures in nonhuman primates. <i>Experimental and Clinical Psychopharmacology</i> , 2020 , 28, 517-526	3.2	2
284	Nicotinic Acetylcholine Receptor Accessory Subunits Determine the Activity Profile of Epibatidine Derivatives. <i>Molecular Pharmacology</i> , 2020 , 98, 328-342	4.3	2
283	Formulation and Characterization of Conjugate Vaccines to Reduce Opioid Use Disorders Suitable for Pharmaceutical Manufacturing and Clinical Evaluation. <i>Molecular Pharmaceutics</i> , 2019 , 16, 2364-237	5 ^{5.6}	10
282	Kappa Opioid Receptors Drive a Tonic Aversive Component of Chronic Pain. <i>Journal of Neuroscience</i> , 2019 , 39, 4162-4178	6.6	53
281	Behavioral Pharmacology of Novel Kappa Opioid Receptor Antagonists in Rats. <i>International Journal of Neuropsychopharmacology</i> , 2019 , 22, 735-745	5.8	12
280	The selective Eppioid receptor antagonist JDTic attenuates the alcohol deprivation effect in rats. European Neuropsychopharmacology, 2019 , 29, 1386-1396	1.2	3
279	Synthesis and Characterization of the Selective, Reversible PKC Inhibitor (9 S)-9-[(Dimethylamino)methyl]-6,7,10,11-tetrahydro-9 H,18 H-5,21:12,17-dimethenodibenzo[e,k]pyrrolo[3,4-h][1,4,13]oxadiazacyclohexadecine-18,20(19 H)-dione, Ruboxistaurin (LY333531).	5.7	
278	Blockade of nicotinic acetylcholine receptor enhances the responsiveness to bupropion in the mouse forced swim test. <i>Behavioural Brain Research</i> , 2019 , 360, 262-269	3.4	4
277	The Eppioid receptor antagonist JDTic decreases ethanol intake in alcohol-preferring AA rats. <i>Psychopharmacology</i> , 2018 , 235, 1581-1591	4.7	5
276	Structure of the Nanobody-Stabilized Active State of the Kappa Opioid Receptor. Cell, 2018, 172, 55-67	. e 5652	205
275	Opioid Dose- and Route-Dependent Efficacy of Oxycodone and Heroin Vaccines in Rats. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2018 , 365, 346-353	4.7	27
274	Dissociable effects of the kappa opioid receptor agonist nalfurafine on pain/itch-stimulated and pain/itch-depressed behaviors in male rats. <i>Psychopharmacology</i> , 2018 , 235, 203-213	4.7	30
273	Potent and Selective Tetrahydroisoquinoline Kappa Opioid Receptor Antagonists of Lead Compound (3 R)- N-[1 R)-1-(Cyclohexylmethyl)-2-methylpropyl]-7-hydroxy-1,2,3,4-tetrahydroisoquinoline-3-carboxamide	8.3	3

(2015-2018)

272	Potent and Selective Tetrahydroisoquinoline Kappa Opioid Receptor Antagonists of Lead Compound (3 R)-7-Hydroxy- N-[(1 S)-2-methyl-1-(piperidin-1-ylmethyl)propyl]-1,2,3,4-tetrahydroisoquinoline-3-carboxamide (PDTic).	8.3	6	
271	Caged Naloxone: Synthesis, Characterization, and Stability of 3-O-(4,5-Dimethoxy-2-nitrophenyl)carboxymethyl Naloxone (CNV-NLX). ACS Chemical Neuroscience, 2018, 9, 563-567	5.7	3	
270	A Double-Blind, Placebo-Controlled Trial Demonstrating the Safety, Tolerability, and Pharmacokinetics of Single, Escalating Oral Doses of RTI-336. <i>Frontiers in Pharmacology</i> , 2018 , 9, 712	5.6	1	
269	New insights on the effects of varenicline on nicotine reward, withdrawal and hyperalgesia in mice. <i>Neuropharmacology</i> , 2018 , 138, 72-79	5.5	14	
268	Probing the Allosteric Role of the \$\textrm{\text{\text{B}}}\$ Subunit of \$\text{\text{\text{\text{\text{B}}}}\$ Nicotinic Acetylcholine Receptors by Functionally Selective Modulators and Ligands. \$ACS Chemical Biology, \$\text{2017}\$, 12, 702-714	4.9	8	
267	In⊡ivo interactions between ⊞7 nicotinic acetylcholine receptor and nuclear peroxisome proliferator-activated receptor-⊞: Implication for nicotine dependence. <i>Neuropharmacology</i> , 2017 , 118, 38-45	5.5	26	
266	Simple Tetrahydroisoquinolines Are Potent and Selective Kappa Opioid Receptor Antagonists. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 742-745	4.3	5	
265	Safety and efficacy of an oxycodone vaccine: Addressing some of the unique considerations posed by opioid abuse. <i>PLoS ONE</i> , 2017 , 12, e0184876	3.7	35	
264	Effects of Chronic Social Defeat Stress on Sleep and Circadian Rhythms Are Mitigated by Kappa-Opioid Receptor Antagonism. <i>Journal of Neuroscience</i> , 2017 , 37, 7656-7668	6.6	53	
263	Synthesis, Nicotinic Acetylcholine Receptor Binding, and in Vitro and in Vivo Pharmacological Properties of 2@Fluoro-(substituted thiophenyl)deschloroepibatidine Analogues. <i>ACS Chemical Neuroscience</i> , 2017 , 8, 115-127	5.7	1	
262	Sex Differences and Drug Dose Influence the Role of the ± 7 Nicotinic Acetylcholine Receptor in the Mouse Dextran Sodium Sulfate-Induced Colitis Model. <i>Nicotine and Tobacco Research</i> , 2017 , 19, 460-468	34.9	12	
261	Design, synthesis, and pharmacological evaluation of JDTic analogs to examine the significance of replacement of the 3-hydroxyphenyl group with pyridine or thiophene bioisosteres. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 3842-8	3.4	5	
260	Synthesis, Nicotinic Acetylcholine Binding, and in Vitro and in Vivo Pharmacological Properties of 2QFluoro-(carbamoylpyridinyl)deschloroepibatidine Analogues. <i>ACS Chemical Neuroscience</i> , 2016 , 7, 100	o4÷12	4	
259	Attenuated nicotine-like effects of varenicline but not other nicotinic ACh receptor agonists in monkeys receiving nicotine daily. <i>British Journal of Pharmacology</i> , 2016 , 173, 3454-3466	8.6	3	
258	Pharmacodynamic Relationships between Duration of Action of JDTic-like Kappa-Opioid Receptor Antagonists and Their Brain and Plasma Pharmacokinetics in Rats. <i>ACS Chemical Neuroscience</i> , 2016 , 7, 1737-1745	5.7	7	
257	Design, Synthesis, and Biological Evaluation of Structurally Rigid Analogues of 4-(3-Hydroxyphenyl)piperidine Opioid Receptor Antagonists. <i>Journal of Organic Chemistry</i> , 2016 , 81, 10383-10391	4.2	6	
256	Nicotine Enhances the Hypnotic and Hypothermic Effects of Alcohol in the Mouse. <i>Alcoholism:</i> Clinical and Experimental Research, 2016 , 40, 62-72	3.7	5	
255	In vitro and in vivo neuronal nicotinic receptor properties of (+)- and (-)-pyrido[3,4]homotropane [(+)- and (-)-PHT]: (+)-PHT is a potent and selective full agonist at ⊞6½ containing neuronal nicotinic acetylcholine receptors. <i>ACS Chemical Neuroscience</i> , 2015 , 6, 920-6	5.7	7	

254	Examination of the metabolite hydroxybupropion in the reinforcing and aversive stimulus effects of nicotine in rats. <i>Psychopharmacology</i> , 2015 , 232, 2763-71	4.7	8
253	Design, synthesis, and pharmacological evaluation of JDTic analogs to examine the significance of the 3- and 4-methyl substituents. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 6379-88	3.4	12
252	A Double-Blind, Placebo-Controlled Trial to Evaluate the Safety, Tolerability, and Pharmacokinetics of Single, Escalating Oral Doses of JDTic. <i>Neuropsychopharmacology</i> , 2015 , 40, 2059-65	8.7	37
251	Quantitative Signaling and Structure-Activity Analyses Demonstrate Functional Selectivity at the Nociceptin/Orphanin FQ Opioid Receptor. <i>Molecular Pharmacology</i> , 2015 , 88, 502-11	4.3	28
250	Synthesis, nicotinic acetylcholine receptor binding, in vitro and in vivo pharmacology properties of 3Q(substituted pyridinyl)-deschloroepibatidine analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 5693-701	3.4	1
249	Novel Synthesis and Pharmacological Characterization of NOP Receptor Agonist 8-[(1S,3aS)-2,3,3a,4,5,6-Hexahydro-1H-phenalen-1-yl]-1-phenyl-1,3,8-triazaspiro[4.5]decan-4-one (Ro 64-6198). ACS Chemical Neuroscience, 2015 , 6, 1956-64	5.7	13
248	Effects of orally-bioavailable short-acting kappa opioid receptor-selective antagonist LY2456302 on nicotine withdrawal in mice. <i>Neuropharmacology</i> , 2015 , 97, 270-4	5.5	23
247	Abnormal error processing in depressive states: a translational examination in humans and rats. <i>Translational Psychiatry</i> , 2015 , 5, e564	8.6	15
246	Anti-nociception mediated by a lippioid receptor agonist is blocked by a lipeceptor agonist. <i>British Journal of Pharmacology</i> , 2015 , 172, 691-703	8.6	15
245	Combining Active Immunization with Monoclonal Antibody Therapy To Facilitate Early Initiation of a Long-Acting Anti-Methamphetamine Antibody Response. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 4665-77	8.3	23
244	Effects of ketoprofen, morphine, and kappa opioids on pain-related depression of nesting in mice. <i>Pain</i> , 2015 , 156, 1153-1160	8	50
243	Effect of the 3- and 4-methyl groups on the opioid receptor properties of N-substituted trans-3,4-dimethyl-4-(3-hydroxyphenyl)piperidines. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 3140-7	8.3	4
242	Emergence and properties of spice and bath salts: a medicinal chemistry perspective. <i>Life Sciences</i> , 2014 , 97, 9-19	6.8	60
241	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 2@fluoro-3@(substituted pyridinyl)-7-deschloroepibatidine analogues. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 836-48	8.3	12
240	Design, synthesis, and biological evaluation of (3R)-1,2,3,4-tetrahydro-7-hydroxy-N-[(1S)-1-[[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]m (JDTic) analogues: in vitro pharmacology and ADME profile. <i>Journal of Medicinal Chemistry</i> , 2014 ,	ietki¥l]-:	2- <u>ഩ</u> ethylp
239	57, 7367-81 Discriminative stimulus and hypothermic effects of some derivatives of the nAChR agonist epibatidine in mice. <i>Psychopharmacology</i> , 2014 , 231, 4455-66	4.7	13
238	Bupropion and bupropion analogs as treatments for CNS disorders. <i>Advances in Pharmacology</i> , 2014 , 69, 177-216	5.7	41
237	Effects of chronic varenicline treatment on nicotine, cocaine, and concurrent nicotine+cocaine self-administration. <i>Neuropsychopharmacology</i> , 2014 , 39, 1222-31	8.7	17

236	The discovery and development of the N-substituted trans-3,4-dimethyl-4-(3@hydroxyphenyl)piperidine class of pure opioid receptor antagonists. <i>ChemMedChem</i> , 2014 , 9, 1638-54	3.7	10
235	Simple radiometric method for accurately quantitating epitope densities of hapten-protein conjugates with sulfhydryl linkages. <i>Bioconjugate Chemistry</i> , 2014 , 25, 2112-5	6.3	7
234	Effects of the kappa opioid receptor antagonist, norbinaltorphimine, on stress and drug-induced reinstatement of nicotine-conditioned place preference in mice. <i>Psychopharmacology</i> , 2013 , 226, 763-8	4.7	47
233	4EMethyl-5-(3-hydroxyphenyl)morphan opioid agonist and partial agonist derived from a 4Emethyl-5-(3-hydroxyphenyl)morphan pure antagonist. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 8826-	3 ⁸ .3	6
232	Vaccination protects rats from methamphetamine-induced impairment of behavioral responding for food. <i>Vaccine</i> , 2013 , 31, 4596-602	4.1	35
231	Patterns of nicotinic receptor antagonism II: cardiovascular effects in rats. <i>Drug and Alcohol Dependence</i> , 2013 , 131, 284-97	4.9	19
230	Development of Epioid receptor antagonists. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 2178-95	8.3	120
229	Discovery of N-{4-[(3-hydroxyphenyl)-3-methylpiperazin-1-yl]methyl-2-methylpropyl}-4-phenoxybenzamide analogues as selective kappa opioid receptor antagonists. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 455	8.3 1-67	10
228	Structurally distinct nicotine immunogens elicit antibodies with non-overlapping specificities. <i>Biochemical Pharmacology</i> , 2012 , 83, 543-50	6	55
227	The long-lasting effects of JDTic, a kappa opioid receptor antagonist, on the expression of ethanol-seeking behavior and the relapse drinking of female alcohol-preferring (P) rats. <i>Pharmacology Biochemistry and Behavior</i> , 2012 , 101, 581-7	3.9	36
226	The kappa opioid receptor antagonist JDTic attenuates alcohol seeking and withdrawal anxiety. <i>Addiction Biology</i> , 2012 , 17, 634-47	4.6	79
225	Influence of chronic dopamine transporter inhibition by RTI-336 on motor behavior, sleep, and hormone levels in rhesus monkeys. <i>Experimental and Clinical Psychopharmacology</i> , 2012 , 20, 77-83	3.2	6
224	Corticotropin-releasing factor (CRF)-induced disruption of attention in rats is blocked by the Ebpioid receptor antagonist JDTic. <i>Neuropsychopharmacology</i> , 2012 , 37, 2809-16	8.7	42
223	Effects of the specific ⊞4½ nAChR antagonist, 2-fluoro-3-(4-nitrophenyl) deschloroepibatidine, on nicotine reward-related behaviors in rats and mice. <i>Psychopharmacology</i> , 2012 , 223, 159-68	4.7	19
222	Structure of the human Eppioid receptor in complex with JDTic. <i>Nature</i> , 2012 , 485, 327-32	50.4	695
221	Synthesis and nicotinic acetylcholine receptor in vitro and in vivo pharmacological properties of 2@fluoro-3@(substituted phenyl)deschloroepibatidine analogues of 2@fluoro-3@(4-nitrophenyl)deschloroepibatidine. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 6512-22	8.3	24
220	Designer drugs: a medicinal chemistry perspective. <i>Annals of the New York Academy of Sciences</i> , 2012 , 1248, 18-38	6.5	104
219	Bupropion and its main metabolite reverse nicotine chronic tolerance in the mouse. <i>Nicotine and Tobacco Research</i> , 2012 , 14, 1356-61	4.9	11

218	Antagonism of the hypothermic effects of nicotinic receptor ligands in mice. <i>FASEB Journal</i> , 2012 , 26, 1048.5	0.9	
217	Kappa opioid receptor signaling in the basolateral amygdala regulates conditioned fear and anxiety in rats. <i>Biological Psychiatry</i> , 2011 , 70, 425-33	7.9	103
216	Synthesis of mercapto-(+)-methamphetamine haptens and their use for obtaining improved epitope density on (+)-methamphetamine conjugate vaccines. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 5221-8	8.3	40
215	Duration of action of a broad range of selective Eppioid receptor antagonists is positively correlated with c-Jun N-terminal kinase-1 activation. <i>Molecular Pharmacology</i> , 2011 , 80, 920-9	4.3	88
214	Interaction of tyrosine 151 in norepinephrine transporter with the 2lgroup of cocaine analog RTI-113. <i>Neuropharmacology</i> , 2011 , 61, 112-20	5.5	13
213	Synthesis and evaluation of 1,2,4-methyltriazines as mGluR5 antagonists. <i>Organic and Biomolecular Chemistry</i> , 2011 , 9, 4276-86	3.9	5
212	Synthesis and Evaluation of Metabotropic Glutamate Receptor Subtype 5 Antagonists Based on Fenobam(). ACS Medicinal Chemistry Letters, 2011, 2, 882-884	4.3	7
211	Synthesis of 2-(substituted phenyl)-3,5,5-trimethylmorpholine analogues and their effects on monoamine uptake, nicotinic acetylcholine receptor function, and behavioral effects of nicotine. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1441-8	8.3	7
210	Preparation of a Series of 5-Methyl-3-(substituted)-[1,2,4]triazines. <i>Tetrahedron Letters</i> , 2011 , 52, 3345-	-3 <u>2</u> 346	2
209	Acute administration of cocaine decreases cell surface expression of DAT in the squirrel monkey caudate. <i>FASEB Journal</i> , 2011 , 25, 1083.3	0.9	
208	Effects of hydroxymetabolites of bupropion on nicotine dependence behavior in mice. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010 , 334, 1087-95	4.7	36
207	Synthesis and characterization of in vitro and in vivo profiles of hydroxybupropion analogues: aids to smoking cessation. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 4731-48	8.3	36
206	Synthesis and biological evaluation of bupropion analogues as potential pharmacotherapies for smoking cessation. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 2204-14	8.3	35
205	Analogues of (3R)-7-hydroxy-N-[(1S)-1-{[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl}-2-methylproperidical (JDTic). Synthesis and in vitro and in vivo opioid receptor antagonist activity. <i>Journal of Medicinal</i>	oyd); 1,2	2, <u>3</u> ,4-tetra
204	1-Substituted 4-(3-Hydroxyphenyl)piperazines Are Pure Opioid Receptor Antagonists. <i>ACS Medicinal Chemistry Letters</i> , 2010 , 1, 365-369	4.3	6
203	Nicotinic acetylcholine receptor efficacy and pharmacological properties of 3-(substituted phenyl)-2Bubstituted tropanes. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 8345-53	8.3	5
202	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 3Q(substituted phenyl)epibatidine analogues. Nicotinic partial agonists. <i>Journal of Natural Products</i> , 2010 , 73, 306-12	4.9	11
201	Kappa opioid mediation of cannabinoid effects of the potent hallucinogen, salvinorin A, in rodents. <i>Psychopharmacology</i> , 2010 , 210, 275-84	4.7	35

200	Role of kappa-opioid receptors in the effects of salvinorin A and ketamine on attention in rats. <i>Psychopharmacology</i> , 2010 , 210, 263-74	4.7	67
199	Effectiveness of analogs of the kappa opioid receptor antagonist (3R)-7-hydroxy-N-((1S)-1-{[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl}-2-methylpro (JDTic) to reduce U50,488-induced diuresis and stress-induced cocaine reinstatement in rats.	р <u>у</u> l) ₇ 1,	2,3 <i>5</i> 4-tetra
198	Lower reinforcing strength of the phenyltropane cocaine analogs RTI-336 and RTI-177 compared to cocaine in nonhuman primates. <i>Pharmacology Biochemistry and Behavior</i> , 2010 , 96, 274-8	3.9	10
197	From rapid in vitro screening to rapid in vivo screening in the drug discovery process. <i>Neuropsychopharmacology</i> , 2009 , 34, 251-2	8.7	9
196	High specific activity (+)-amphetamine and (+)-methamphetamine. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2009 , 52, 457-462	1.9	3
195	Neutral antagonist activity of naltrexone and 6beta-naltrexol in naWe and opioid-dependent C6 cells expressing a mu-opioid receptor. <i>British Journal of Pharmacology</i> , 2009 , 156, 1044-53	8.6	22
194	Positive allosteric modulation of the human cannabinoid (CB) receptor by RTI-371, a selective inhibitor of the dopamine transporter. <i>British Journal of Pharmacology</i> , 2009 , 156, 1178-84	8.6	70
193	Synthesis and structure-activity relationship of 3beta-(4-alkylthio, -methylsulfinyl, and -methylsulfonylphenyl)tropane and 3beta-(4-alkylthiophenyl)nortropane derivatives for monoamine transporters. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 5126-32	3.4	5
192	Synthesis and in vitro opioid receptor functional antagonism of methyl-substituted analogues of (3R)-7-hydroxy-N-[(1S)-1-{[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl}-2-methylpro (JDTic). <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 7463-72	pyվ <u></u> ե1,	2, 3 ,84-tetra
191	Functional and biological determinants affecting the duration of action and efficacy of anti-(+)-methamphetamine monoclonal antibodies in rats. <i>Vaccine</i> , 2009 , 27, 7011-20	4.1	27
190	Synthesis and biological evaluation of bupropion analogues as potential pharmacotherapies for cocaine addiction. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 6768-81	8.3	100
189	The synthesis of haptens and their use for the development of monoclonal antibodies for treating methamphetamine abuse. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 7301-9	8.3	27
188	EPIBATIDINE ANALOGS SYNTHESIZED FOR CHARACTERIZATION OF NICOTINIC PHARMACOPHORES-A REVIEW. <i>Heterocycles</i> , 2009 , 79, 99-120	0.8	18
187	mGluR5 antagonists that block calcium mobilization in vitro also reverse (S)-3,5-DHPG-induced hyperalgesia and morphine antinociceptive tolerance in vivo. <i>Brain Research</i> , 2008 , 1187, 58-66	3.7	17
186	Relationship between rate of drug uptake in brain and behavioral pharmacology of monoamine transporter inhibitors in rhesus monkeys. <i>Pharmacology Biochemistry and Behavior</i> , 2008 , 90, 453-62	3.9	38
185	Development of 3-phenyltropane analogues with high affinity for the dopamine and serotonin transporters and low affinity for the norepinephrine transporter. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 8048-56	8.3	9
184	Improved Synthesis of the ORL Antagonist 1-[(3R,4R)-1-Cyclooctylmethyl-3-ethoxycarbonyl-4-piperidinyl]-3-ethyl-1,3-dihydro-2H-benzimidazol-2-c (J-113397). Synthetic Communications, 2008 , 38, 1926-1930	one ₇	1
183	Preparation of carbon-14 labeled (3R)-7-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl}-2-methylprop (JDTic). <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2008 , 51, 440-443	oy l :.0,2,	3, 4 -tetral

182	A new synthesis of the ORL-1 antagonist 1-[(3R,4R)-1-cyclooctylmethyl-3-hydroxymethyl-4-piperidinyl]-3-ethyl-1,3-dihydro-2H-benzimidazol-2-or (J-113397) and activity in a calcium mobilization assay. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 82		9
181	Synthesis, nicotinic acetylcholine receptor binding, and pharmacological properties of 3Q(substituted phenyl)deschloroepibatidine analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 746	-5 ³ 4 ⁴	15
180	Synthesis and receptor binding properties of 2beta-alkynyl and 2beta-(1,2,3-triazol)substituted 3beta-(substituted phenyl)tropane derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 5529-35	3.4	5
179	Synthesis and monoamine transporter binding properties of 2beta-[3@substituted benzyl)isoxazol-5-yl]- and 2beta-[3@methyl-4@substituted phenyl)isoxazol-5-yl]-3beta-(substituted phenyl)tropanes. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 6682-8	3.4	6
178	3@Fluoro substitution in the pyridine ring of epibatidine improves selectivity and efficacy for alpha4beta2 versus alpha3beta4 nAChRs. <i>Neuropharmacology</i> , 2008 , 55, 1287-92	5.5	5
177	Synthesis and in vitro opioid receptor functional antagonism of analogues of the selective kappa opioid receptor antagonist (3R)-7-hydroxy-N-((1S)-1-{[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl}-2-methylpro	8.3 oyl)-1,2	21 2 ,3,4-te tra
176	Antagonists at metabotropic glutamate receptor subtype 5: structure activity relationships and therapeutic potential for addiction. <i>Annals of the New York Academy of Sciences</i> , 2008 , 1141, 221-32	6.5	48
175	Development of the Dopamine Transporter Selective RTI-336 as a Pharmacotherapy for Cocaine Abuse 2008 , 179		2
174	Using hapten design to discover therapeutic monoclonal antibodies for treating methamphetamine abuse. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 322, 30-9	4.7	54
173	Synthesis, monoamine transporter binding, properties, and functional monoamine uptake activity of 3beta-[4-methylphenyl and 4-chlorophenyl]-2 beta-[5-(substituted phenyl)thiazol-2-yl]tropanes. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3686-95	8.3	9
172	Synthesis, nicotinic acetylcholine receptor binding, antinociceptive and seizure properties of methyllycaconitine analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 678-85	3.4	15
171	Faster onset and dopamine transporter selectivity predict stimulant and reinforcing effects of cocaine analogs in squirrel monkeys. <i>Pharmacology Biochemistry and Behavior</i> , 2007 , 86, 45-54	3.9	47
170	Effects of combined dopamine and serotonin transporter inhibitors on cocaine self-administration in rhesus monkeys. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 320, 757-65	4.7	64
169	Improved Procedure for the Synthesis of DAMGO. <i>Synthetic Communications</i> , 2007 , 37, 2345-2348	1.7	O
168	Synthesis and nicotinic acetylcholine receptor binding properties of bridged and fused ring analogues of epibatidine. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 6383-91	8.3	29
167	Anxiolytic-like effects of kappa-opioid receptor antagonists in models of unlearned and learned fear in rats. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 323, 838-45	4.7	196
166	Synthesis and pharmacological evaluation of phenylethynyl[1,2,4]methyltriazines as analogues of 3-methyl-6-(phenylethynyl)pyridine. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3388-91	8.3	19
165	Effects of dopamine transporter selective 3-phenyltropane analogs on locomotor activity, drug discrimination, and cocaine self-administration after oral administration. <i>European Journal of Pharmacology</i> , 2006 , 553, 149-56	5.3	18

(2005-2006)

164	Dopamine transporter ligands: recent developments and therapeutic potential. <i>Current Topics in Medicinal Chemistry</i> , 2006 , 6, 1825-43	3	63
163	2-Fluoro-3-(4-nitro-phenyl)deschloroepibatidine is a novel potent competitive antagonist of human neuronal alpha4beta2 nAChRs. <i>Molecular Pharmacology</i> , 2006 , 69, 1945-52	4.3	17
162	Development of the dopamine transporter selective RTI-336 as a pharmacotherapy for cocaine abuse. <i>AAPS Journal</i> , 2006 , 8, E196-203	3.7	32
161	Synthesis and monoamine transporter binding properties of 2,3-cyclo analogues of 3beta-(4@minophenyl)-2beta-tropanemethanol. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 4589-94	8.3	2
160	N-substituted 4beta-methyl-5-(3-hydroxyphenyl)-7alpha-amidomorphans are potent, selective kappa opioid receptor antagonists. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 1781-91	8.3	19
159	Highly potent and selective phenylmorphan-based inverse agonists of the opioid delta receptor. Journal of Medicinal Chemistry, 2006 , 49, 5597-609	8.3	15
158	A reduced rate of in vivo dopamine transporter binding is associated with lower relative reinforcing efficacy of stimulants. <i>Neuropsychopharmacology</i> , 2006 , 31, 351-62	8.7	28
157	Synthesis and pharmacological characterization of nicotinic acetylcholine receptor properties of (+)- and (-)-pyrido-[3,4-b]homotropanes. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 3244-50	8.3	18
156	Synthesis and pharmacological characterization of exo-2-(2Qchloro-5-pyridinyl)-7-(endo and exo)-aminobicyclo[2.2.1]heptanes as novel epibatidine analogues. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 7491-5	8.3	12
155	Synthesis and monoamine transporter binding properties of 3alpha-(substituted phenyl)nortropane-2beta-carboxylic acid methyl esters. Norepinephrine transporter selective compounds. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 3852-7	8.3	11
154	Synthesis and monoamine transporter binding properties of 2,3-diaryltropanes. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 7437-44	8.3	5
153	N-substituted cis-4a-(3-hydroxyphenyl)-8a-methyloctahydroisoquinolines are opioid receptor pure antagonists. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 8182-93	8.3	15
152	Bupropion increases striatal vesicular monoamine transport. <i>Neuropharmacology</i> , 2005 , 49, 820-30	5.5	60
151	Synthesis and monoamine transporter binding properties of 3beta-(3Q4Qdisubstituted phenyl)tropane-2beta-carboxylic acid methyl esters. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 2767-71	8.3	14
150	Synthesis, structural identification, and ligand binding of tropane ring analogs of paroxetine and an unexpected aza-bicyclo[3.2.2]nonane rearrangement product. <i>Bioorganic and Medicinal Chemistry</i> , 2005 , 13, 2439-49	3.4	4
149	Structure-activity relationship of trihexyphenidyl analogs with respect to the dopamine transporter in the on going search for a cocaine inhibitor. <i>European Journal of Medicinal Chemistry</i> , 2005 , 40, 1013-2	16.8	6
148	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 3@substituted deschloroepibatidine analogues. Novel nicotinic antagonists. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 1221-8	8.3	28
147	Effects of JDTic, a selective kappa-opioid receptor antagonist, on the development and expression of physical dependence on morphine using a rat continuous-infusion model. <i>European Journal of Pharmacology</i> , 2005 , 524, 89-94	5.3	32

146	Cocaine self-administration and locomotor activity are altered in Lewis and F344 inbred rats by RTI 336, a 3-phenyltropane analog that binds to the dopamine transporter. <i>Brain Research</i> , 2005 , 1055, 186	- 3 5	22
145	Tritiation of the cannabinoid receptor antagonist SR144528 involving lithium aluminum tritide reduction; assessment of the kinetic isotope effect by 3H-NMR. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2005 , 48, 589-596	1.9	5
144	Differential effects of the novel kappa opioid receptor antagonist, JDTic, on reinstatement of cocaine-seeking induced by footshock stressors vs cocaine primes and its antidepressant-like effects in rats. <i>Psychopharmacology</i> , 2005 , 183, 118-26	4.7	236
143	Effects of murine-derived anti-methamphetamine monoclonal antibodies on (+)-methamphetamine self-administration in the rat. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2004 , 309, 1248-5	5 ∮ ∙7	64
142	Enantioselective effects of hydroxy metabolites of bupropion on behavior and on function of monoamine transporters and nicotinic receptors. <i>Molecular Pharmacology</i> , 2004 , 66, 675-82	4.3	175
141	Effects of dopamine transporter inhibitors on cocaine self-administration in rhesus monkeys: relationship to transporter occupancy determined by positron emission tomography neuroimaging. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2004 , 309, 959-69	4.7	55
140	Epibatidine structure-activity relationships. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 1889-9	6 .9	65
139	Importance of phenolic address groups in opioid kappa receptor selective antagonists. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 1070-3	8.3	25
138	Monoamine transporter binding, locomotor activity, and drug discrimination properties of 3-(4-substituted-phenyl)tropane-2-carboxylic acid methyl ester isomers. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 6401-9	8.3	34
137	Synthesis, monoamine transporter binding properties, and behavioral pharmacology of a series of 3beta-(substituted phenyl)-2beta-(3@substituted isoxazol-5-yl)tropanes. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 296-302	8.3	37
136	Discovery of the first N-substituted 4beta-methyl-5-(3-hydroxyphenyl)morphan to possess highly potent and selective opioid delta receptor antagonist activity. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 281-4	8.3	16
135	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 2Qfluoro-3Q(substituted phenyl)deschloroepibatidine analogues. Novel nicotinic antagonist. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 4588-94	8.3	39
134	2002 Medicinal Chemistry Division Award address: monoamine transporters and opioid receptors. Targets for addiction therapy. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 1775-94	8.3	77
133	Identification of (3R)-7-hydroxy-N-((1S)-1-[[(3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl]methyl]-2-methylpropyl)-1,2,3,4-tetrahydro- 3-isoquinolinecarboxamide as a novel potent and selective opioid kappa receptor antagonist. <i>Journal of Medicinal Chemistry</i> ,	8.3	72
132	Tritiation of SR141716 by metallation Eduction: tritium-proton nOe study. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2002 , 45, 59-70	1.9	13
131	Self-administration of cocaine and the cocaine analog RTI-113: relationship to dopamine transporter occupancy determined by PET neuroimaging in rhesus monkeys. <i>Synapse</i> , 2002 , 43, 78-85	2.4	55
130	In vitro and in vivo characterization of [1251]iodomethyllycaconitine in the rat. Synapse, 2002, 44, 117-23	32.4	19
129	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 2-exo-2-(2@Qdisubstituted 5Qpyridinyl)-7-azabicyclo[2.2.1]heptanes: epibatidine analogues. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 4755-61	8.3	47

128	Effects of pyridine ring substitutions on affinity, efficacy, and subtype selectivity of neuronal nicotinic receptor agonist epibatidine. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2002 , 302, 1246-52	4.7	34
127	Discovery of an opioid kappa receptor selective pure antagonist from a library of N-substituted 4beta-methyl-5-(3-hydroxyphenyl)morphans. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 3524-30	8.3	25
126	Synthesis and transporter binding properties of 3beta-[4Qphenylalkyl, -phenylalkenyl, and -phenylalkynyl)phenyl]tropane-2beta-carboxylic acid methyl esters: evidence of a remote phenyl binding domain on the dopamine transporter. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 4029-37	8.3	13
125	Synthesis of bridged analogs of epibatidine. 3-Chloro-5,7,8,9,9a,10-hexahydro-7,10-methanopyrrolo[1,2-b]-2,6-naphthyridine and 2-chloro-5,5a,6,7,8,10-hexahydro-5,8-methanopyrrolo[2,1-b]-1,7-naphthyridine. <i>Tetrahedron Letters</i> ,	2	18
124	Opioid peptide receptor studies. 15. Relative efficacy of 4-[(N-allyl-3-methyl-4-piperidinyl)phenylamino]-N,N-diethylbenzamide and related compounds at the cloned human delta-opioid receptor. <i>Synapse</i> , 2001 , 40, 269-74	2.4	4
123	Anticocaine catalytic antibodies have no affinity for RTI compounds: implications for treatment. <i>Synapse</i> , 2001 , 41, 176-8	2.4	5
122	[(125)I]3beta-(4-ethyl-3-iodophenyl)nortropane-2beta-carboxylic acid methyl ester ([(125)I]EINT): a potent and selective radioligand for the brain serotonin transporter. <i>Synapse</i> , 2001 , 41, 241-7	2.4	3
121	Opioid peptide receptor studies. 14. Stereochemistry determines agonist efficacy and intrinsic efficacy in the [(35)S]GTP-gamma-S functional binding assay. <i>Synapse</i> , 2001 , 39, 64-9	2.4	30
120	Factors influencing agonist potency and selectivity for the opioid delta receptor are revealed in structure-activity relationship studies of the 4-[(N-substituted-4-piperidinyl)arylamino]-N,N-diethylbenzamides. <i>Journal of Medicinal Chemistry</i> ,	8.3	16
119	Identification of the first trans-(3R,4R)- dimethyl-4-(3-hydroxyphenyl)piperidine derivative to possess highly potent and selective opioid kappa receptor antagonist activity. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 2687-90	8.3	100
118	Locomotor stimulant effects of novel phenyltropanes in the mouse. <i>Drug and Alcohol Dependence</i> , 2001 , 65, 25-36	4.9	18
117	Generation of anti-(+)methamphetamine antibodies is not impeded by (+)methamphetamine administration during active immunization of rats. <i>International Immunopharmacology</i> , 2001 , 1, 329-38	5.8	58
116	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 2-exo-2-(2&ubstituted 5&pyridinyl)-7-azabicyclo[2.2.1]heptanes. Epibatidine analogues. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 2229-37	8.3	76
115	Synthesis, nicotinic acetylcholine receptor binding, and antinociceptive properties of 2-exo-2-(2&ubstituted-3&phenyl-5&pyridinyl)-7-azabicyclo[2.2.1]heptanes. Novel nicotinic antagonist. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 4039-41	8.3	40
114	Agonist-, antagonist-, and inverse agonist-regulated trafficking of the delta-opioid receptor correlates with, but does not require, G protein activation. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2001 , 298, 1015-20	4.7	30
113	Synthesis of [125I]-3E(4-ethyl-3-iodophenyl)nortropane-2Etarboxylic acid methyl ester ([125I]EINT). <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2000 , 43, 137-146	1.9	3
112	4-[(8-Alkyl-8-azabicyclo[3.2.1]octyl-3-yl)-3-arylanilino]-N,N-diethylbenzamides: high affinity, selective ligands for the delta opioid receptor illustrate factors important to antagonist activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000 , 10, 1281-4	2.9	8
111	3alpha-(4-Substituted phenyl)nortropane-2beta-carboxylic acid methyl esters show selective binding at the norepinephrine transporter. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000 , 10, 2445-7	2.9	11

110	RTI-76, an isothiocyanate derivative of a phenyltropane cocaine analog, as a tool for irreversibly inactivating dopamine transporter function in vitro. <i>Naunyn-Schmiedeberg Archives of Pharmacology</i> , 2000 , 362, 238-47	3.4	7
109	Dopamine transporter synthesis and degradation rate in rat striatum and nucleus accumbens using RTI-76. <i>Neuropharmacology</i> , 2000 , 39, 578-85	5.5	35
108	Synthesis and pharmacological characterization of [(125)I]iodomethyllycaconitine ([(125)I]iodo-MLA). A new ligand for the alpha(7) nicotinic acetylcholine receptor. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 142-5	8.3	28
107	(+/-)-4-[(N-allyl-cis-3-methyl-4-piperidinyl)phenylamino]-N,N-diethylbenzamide displays selective binding for the delta opioid receptor. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999 , 9, 3053-6	2.9	14
106	Optically pure (-)-4-[(N-allyl-3-methyl-4-piperidinyl)phenyl-amino]-N,N-diethylbenzami de displays selective binding and full agonist activity for the delta opioid receptor. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999 , 9, 3347-50	2.9	13
105	Serotonin transporter production and degradation rates: studies with RTI-76. <i>Brain Research</i> , 1999 , 841, 1-10	3.7	19
104	Opioid peptide receptor studies, 11: involvement of Tyr148, Trp318 and His319 of the rat mu-opioid receptor in binding of mu-selective ligands. <i>Synapse</i> , 1999 , 32, 23-8	2.4	32
103	Synthesis of 3E(4-[125I]iodophenyl)tropane-2-Epyrrolidine carboxamide ([125I]RTI-229). <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 1999 , 42, 281-286	1.9	3
102	Synthesis of [1-3H]morphine-6-ED-glucuronide. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 1999 , 42, 851-857	1.9	5
101	Asymmetric synthesis of 9-alkyl-2-benzyl-6,7-benzomorphans: characterization as novel sigma receptor ligands. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 4621-9	8.3	7
100	Comparative studies of epibatidine derivatives [18F]NFEP and [18F]N-methyl-NFEP: kinetics, nicotine effect, and toxicity. <i>Nuclear Medicine and Biology</i> , 1999 , 26, 139-48	2.1	48
99	Pharmacotherapies for treatment of cocaine abuse: preclinical aspects. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 2721-36	8.3	219
98	Pharmacological characterization of nicotine@interaction with cocaine and cocaine analogs. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 1999 , 289, 1229-36	4.7	27
97	Protection of the allylic alcohol double bond from catalytic reduction in the preparation of [1-3H]morphine and [1-3H]codeine 1998 , 41, 811-821		7
96	Opioid peptide receptor studies. 7. The methylfentanyl congener RTI-4614-4 and its four enantiomers bind to different domains of the rat mu opioid receptor. <i>Synapse</i> , 1998 , 28, 117-24	2.4	12
95	RTI-113 administration reduces cocaine self-administration at high occupancy of dopamine transporter. <i>Synapse</i> , 1998 , 30, 49-55	2.4	29
94	New synthesis of 7-(tert-butoxycarbonyl)-7-azabicyclo[2.2.1]hept-2-ene. A key intermediate in the synthesis of epibatidine and analogs. <i>Tetrahedron Letters</i> , 1998 , 39, 5321-5322	2	18
93	N-substituted octahydro-4a-(3-hydroxyphenyl)-10a-methyl-benzo[g]isoquinolines are opioid receptor pure antagonists. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1998 , 8, 3149-52	2.9	8

92	Synthesis and transporter binding properties of (R)-2 beta, 3 beta- and (R)-2 alpha, 3 alpha-diaryltropanes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1998 , 8, 3689-92	2.9	8
91	In vitro and ex vivo autoradiographic studies of nicotinic acetylcholine receptors using [18F]fluoronochloroepibatidine in rodent and human brain. <i>Nuclear Medicine and Biology</i> , 1998 , 25, 449	-54 ¹	27
90	Identification of an opioid kappa receptor subtype-selective N-substituent for (+)-(3R,4R)-dimethyl-4-(3-hydroxyphenyl)piperidine. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 5188-97	8.3	58
89	N-Substituted 9beta-methyl-5-(3-hydroxyphenyl)morphans are opioid receptor pure antagonists. Journal of Medicinal Chemistry, 1998 , 41, 4143-9	8.3	30
88	Synthesis and ligand binding of tropane ring analogues of paroxetine. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 247-57	8.3	21
87	Investigation of the N-substituent conformation governing potency and mu receptor subtype-selectivity in (+)-(3R, 4R)-dimethyl-4-(3-hydroxyphenyl)piperidine opioid antagonists. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 1980-90	8.3	38
86	3 Beta-(4-ethyl-3-iodophenyl)nortropane-2 beta-carboxylic acid methyl ester as a high-affinity selective ligand for the serotonin transporter. <i>Journal of Medicinal Chemistry</i> , 1997 , 40, 3861-4	8.3	43
85	Synthesis and nicotinic acetylcholine receptor binding properties of exo-2-(2@fluoro-5@pyridinyl)-7-azabicyclo-[2.2.1]heptane: a new positron emission tomography ligand for nicotinic receptors. <i>Journal of Medicinal Chemistry</i> , 1997 , 40, 2293-5	8.3	55
84	AN IMPROVED PROCEDURE FOR THE SYNTHESIS OF ANHYDROECGONINE METHYL ESTER. <i>Organic Preparations and Procedures International</i> , 1997 , 29, 308-311	1.1	1
83	Fluoro-norchloroepibatidine: preclinical assessment of acute toxicity. <i>Nuclear Medicine and Biology</i> , 1997 , 24, 743-7	2.1	64
82	Synthesis of [18F]norchlorofluoroepibatidine and its N-methyl derivative: new PET ligands for mapping nicotinic acetylcholine receptors. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 1997 , 39, 827-832	1.9	28
81	N-substituted phenyltropanes as in vivo binding ligands for rapid imaging studies of the dopamine transporter. <i>Synapse</i> , 1997 , 25, 345-9	2.4	36
80	RTI-352: a 3 alpha analogue of RTI-55 as an in vivo dopamine transporter binding ligand. <i>Synapse</i> , 1997 , 25, 389-92	2.4	1
79	Imaging Transporters for Dopamine and Other Neurotransmitters in Brain 1997 , 297-313		4
78	Dopamine Transporter Uptake Blockers 1997 , 263-295		13
77	Synthesis of [18F]norchlorofluoroepibatidine and its N-methyl derivative: new PET ligands for mapping nicotinic acetylcholine receptors 1997 , 39, 827		2
76	Ohmefentanyl and Its Stereoisomers: Chemistry and Pharmacology. <i>Current Medicinal Chemistry</i> , 1997 , 4, 247-270	4.3	9
75	Synthesis, ligand binding, and quantitative structure-activity relationship study of 3 beta-(4@substituted phenyl)-2 beta-heterocyclic tropanes: evidence for an electrostatic interaction at the 2 beta-position. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 2753-63	8.3	65

74	3 alpha-(4@substituted phenyl)tropane-2 beta-carboxylic acid methyl esters: novel ligands with high affinity and selectivity at the dopamine transporter. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 4139-41	8.3	33
73	Synthesis and in vivo studies of a selective ligand for the dopamine transporter: 3 beta-(4-[1251]iodophenyl) tropan-2 beta-carboxylic acid isopropyl ester ([1251]RTI-121). <i>Nuclear Medicine and Biology</i> , 1996 , 23, 277-84	2.1	23
72	Highly potent cocaine analogs cause long-lasting increases in locomotor activity. <i>European Journal of Pharmacology</i> , 1996 , 311, 109-14	5.3	33
71	Synthesis and transporter binding properties of 3 beta-(4Qalkyl-, 4Qalkenyl-, and 4Qalkynylphenyl)nortropane-2 beta-carboxylic acid methyl esters: serotonin transporter selective analogs. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 4027-35	8.3	51
70	Benzyne addition to N-alkyl-4-hydroxy-1-methylisoquinolinium salts; a new and convenient synthesis of (日)-5-methyl-10,11-dihydro-5H-dibenzo-[a,d]cyclohepten-5,10-imine (MK801). <i>Chemical Communications</i> , 1996 , 717-718	5.8	13
69	Mapping nicotinic acetylcholine receptors with PET. <i>Synapse</i> , 1996 , 24, 403-7	2.4	64
68	Mapping dopamine transporters in the human brain with novel selective cocaine analog [125I]RTI-121. <i>Synapse</i> , 1995 , 21, 364-72	2.4	62
67	Facile synthesis of (⊞)-, (+)-, and (-)-galanthamine. <i>Journal of Heterocyclic Chemistry</i> , 1995 , 32, 195-199	1.9	28
66	Evaluation of retinoid lactones as topical therapeutic agents in dermatology. <i>Pharmaceutical Research</i> , 1995 , 12, 983-92	4.5	2
65	Synthesis and ligand binding study of 3 beta-(4@substituted phenyl)-2 beta-(heterocyclic)tropanes. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 3451-3	8.3	23
64	Cocaine and 3 beta-(4@ubstituted phenyl)tropane-2 beta-carboxylic acid ester and amide analogues. New high-affinity and selective compounds for the dopamine transporter. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 379-88	8.3	187
63	PRACTICAL SYNTHESIS AND CHARACTERIZATION OF 7-BENZYLIDENENALTREXONE (BNTX). Organic Preparations and Procedures International, 1995, 27, 621-624	1.1	3
62	Synthesis of (+) and (DEpibatidine. Synthetic Communications, 1995, 25, 63-71	1.7	28
61	Enantiomers of diastereomeric cis-N-[1-(2-hydroxy-2-phenylethyl)-3-methyl-4-piperidyl]-N-phenylpropanamides: synthesis, X-ray analysis, and biological activities. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 1547-57	8.3	30
60	Synthesis and sigma binding properties of 2@substituted 5,9 alpha-dimethyl-6,7-benzomorphans. Journal of Medicinal Chemistry, 1995 , 38, 2978-85	8.3	24
59	In vivo labeling of neuronal nicotinic acetylcholine receptors with radiolabeled isomers of norchloroepibatidine. <i>NeuroReport</i> , 1995 , 6, 2483-8	1.7	24
58	Development of imaging agents for the dopamine transporter. <i>Medicinal Research Reviews</i> , 1995 , 15, 419-44	14.4	30
57	Rate of binding of various inhibitors at the dopamine transporter in vivo. <i>Psychopharmacology</i> , 1995 , 119, 376-84	4.7	66

56	RTI-4793-14, a new ligand with high affinity and selectivity for the (+)-MK801-insensitive [3H]1-]1-(2-thienyl)cyclohexyl]piperidine binding site (PCP site 2) of guinea pig brain. <i>Synapse</i> , 1994 , 16, 59-65	2.4	5
55	Metallation/reduction as a new approach to tritium labeling. The synthesis of [3H]ibogaine. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 1994 , 34, 367-375	1.9	4
54	Synthesis and structural determination of 5H-benzocyclohepten-5,8-imines. <i>Journal of Heterocyclic Chemistry</i> , 1994 , 31, 187-192	1.9	15
53	Synthesis of 4,4-disubstituted piperidine analogs of (日)-cis-N-[1-(2-hydroxy-2-phenylethyl)-3-methyl-4-piperidyl]-N-phenylpropanamide. <i>Journal of Heterocyclic Chemistry</i> , 1994 , 31, 513-520	1.9	3
52	Secondary amine analogues of 3 beta-(4@substituted phenyl)tropane-2 beta-carboxylic acid esters and N-norcocaine exhibit enhanced affinity for serotonin and norepinephrine transporters. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1220-3	8.3	85
51	Evaluation of retinoids as therapeutic agents in dermatology. <i>Pharmaceutical Research</i> , 1994 , 11, 192-20)φ .5	13
50	The requirement for a retinoic acid lactone of 11-cis, 13-cis-stereochemistry for topical dermatologic activity. <i>Pharmaceutical Research</i> , 1994 , 11, 1065-7	4.5	1
49	Synthesis, ligand binding, and QSAR (CoMFA and classical) study of 3 beta-(3@ubstituted phenyl)-, 3 beta-(4@ubstituted phenyl)-, and 3 beta-(3@4@disubstituted phenyl)tropane-2 beta-carboxylic acid methyl esters. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 2865-73	8.3	103
48	3-Aryl-2-(3@substituted-1@Q4@oxadiazol-5@yl)tropane analogues of cocaine: affinities at the cocaine binding site at the dopamine, serotonin, and norepinephrine transporters. <i>Journal of Medicinal Chemistry</i> , 1993 , 36, 2886-90	8.3	106
47	Radiosynthesis of a photoaffinity probe for the cocaine receptor of the dopamine transporter: 3E(p-chlorophenyl)tropan-2Ecarboxylic acid m-([125I]-iodo)-p-azidophenethyl ester ([125I]-RTI-82). <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 1993 , 33, 1131-1137	1.9	19
46	[125I]RTI-55 binding to cocaine-sensitive dopaminergic and serotonergic uptake sites in the human brain. <i>Journal of Neurochemistry</i> , 1993 , 61, 1996-2006	6	45
45	Stimulus generalization from cocaine to analogs with high in vitro affinity for dopamine uptake sites. <i>Behavioural Pharmacology</i> , 1992 , 3, 113???116	2.4	17
44	Dopamine transporter imaging with novel, selective cocaine analogs. <i>NeuroReport</i> , 1992 , 3, 969-72	1.7	24
43	Enantiomeric N-substituted N-normetazocines: a comparative study of affinities at sigma, PCP, and mu opioid receptors. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 2812-8	8.3	68
42	Isopropyl and phenyl esters of 3 beta-(4-substituted phenyl)tropan-2 beta-carboxylic acids. Potent and selective compounds for the dopamine transporter. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 2497-5	5 <mark>0</mark> 0	75
41	High potency cocaine analogs: neurochemical, imaging, and behavioral studies. <i>Annals of the New York Academy of Sciences</i> , 1992 , 654, 282-91	6.5	28
40	2 beta-substituted analogues of cocaine. Synthesis and inhibition of binding to the cocaine receptor. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 135-40	8.3	73
39	Probes for the cocaine receptor. Potentially irreversible ligands for the dopamine transporter. Journal of Medicinal Chemistry, 1992 , 35, 1813-7	8.3	55

38	A cocaine analog and a GBR analog label the same protein in rat striatal membranes. <i>Brain Research</i> , 1992 , 576, 173-4	3.7	14
37	In vivo imaging of dopamine reuptake sites in the primate brain using single photon emission computed tomography (SPECT) and iodine-123 labeled RTI-55. <i>Synapse</i> , 1992 , 10, 169-72	2.4	78
36	High-affinity binding of [125I]RTI-55 to dopamine and serotonin transporters in rat brain. <i>Synapse</i> , 1992 , 12, 27-36	2.4	195
35	In vivo binding of [125I]RTI-55 to dopamine transporters: pharmacology and regional distribution with autoradiography. <i>Synapse</i> , 1992 , 12, 37-46	2.4	60
34	Synthesis of 2-(3-substituted-1,2,4-oxadiazol-5-yl)-8-methyl-8-azabicyclo [3.2.1]octanes and 2 alpha-(3-substituted-1,2,4-oxadiazol-5-yl)-8-methyl-8- azabicyclo[3.2.1]oct-2-enes as potential muscarinic agonists. <i>Pharmaceutical Research</i> , 1992 , 9, 1474-9	4.5	7
33	Cocaine receptor: biochemical characterization and structure-activity relationships of cocaine analogues at the dopamine transporter. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 969-81	8.3	275
32	Synthesis and spectral analysis of (H)-cis-N-[1-(2-hydroxy-2-phenyl-ethyl)-3-methyl-4-piperidyl]-N-phenylpropanamide and (H)-cis-N-[1-(2-hydroxy-1-phenylethyl)-3-methyl-4-piperidyl]-N-phenylpropanamide. <i>Journal of Heterocyclic Chemistry</i> , 1992 , 29, 1773-1779	1.9	2
31	Synthesis and ligand binding of cocaine isomers at the cocaine receptor. <i>Journal of Medicinal Chemistry</i> , 1991 , 34, 883-6	8.3	72
30	Potent substituted-3 beta-phenyltropane analogs of cocaine have cocaine-like discriminative stimulus effects. <i>Drug and Alcohol Dependence</i> , 1991 , 29, 145-51	4.9	20
29	Synthesis, ligand binding, QSAR, and CoMFA study of 3 beta-(p-substituted phenyl)tropane-2 beta-carboxylic acid methyl esters. <i>Journal of Medicinal Chemistry</i> , 1991 , 34, 2719-25	8.3	181
28	Isothiocyanate derivatives of cocaine: irreversible inhibition of ligand binding at the dopamine transporter. <i>Molecular Pharmacology</i> , 1991 , 39, 339-45	4.3	25
27	An improved resolution of (日)-cis-N-normetazocine. <i>Journal of Heterocyclic Chemistry</i> , 1990 , 27, 2139-21	439	18
26	SYNTHESIS OF POTENTIAL 1-[1-(2-THIENYL)CYCLOHEXYL]PIPERIDINE METABOLITES. <i>Organic Preparations and Procedures International</i> , 1990 , 22, 124-128	1.1	0
25	Carbon-13 nuclear magnetic resonance spectra of fentanyl analogs. <i>Journal of Heterocyclic Chemistry</i> , 1989 , 26, 677-686	1.9	17
24	Solid-state and solution conformations of methadone hydrochloride and related compounds. <i>Magnetic Resonance in Chemistry</i> , 1989 , 27, 311-317	2.1	2
23	Synthesis of fentanyl analogs. <i>NIDA Research Monograph</i> , 1989 , 95, 497-8		
22	An improved synthesis of galanthamine. <i>Journal of Heterocyclic Chemistry</i> , 1988 , 25, 1809-1811	1.9	28
21	AN IMPROVED SYNTHESIS OF 1,4-BIS(2?-METHYL-6?-KIHYLANILINO)ANTHRAQUINONE. <i>Organic Preparations and Procedures International</i> , 1987 , 19, 57-60	1.1	O

20	A practical synthesis of (+)-cocaine. Journal of Heterocyclic Chemistry, 1987, 24, 19-21	1.9	24
19	Synthesis and spectral properties of optically active 2-ethylidene-1,5-dimethyl-3,3-diphenylpyrrolidine: Primary methadone metabolite. <i>Journal of Heterocyclic Chemistry</i> , 1986 , 23, 369-374	1.9	O
18	Carbon-13 nuclear magnetic resonance spectra of phencyclidine analogs substituted in the piperidine and aromatic rings. <i>Journal of Heterocyclic Chemistry</i> , 1984 , 21, 71-75	1.9	5
17	SYNTHESIS OF 4-(4?-HYDROXYPIPERIDINO)-4-PHENYLCYCLOHEXANOL, A DIHYDROXY PHENCYCLIDINE METABOLITE. <i>Organic Preparations and Procedures International</i> , 1983 , 15, 371-377	1.1	2
16	Synthesis of naphthyridinone derivatives as potential antimalarials. <i>Journal of Heterocyclic Chemistry</i> , 1981 , 18, 941-946	1.9	10
15	Carbon-13 nuclear magnetic resonance spectra of methaqualone metabolites. <i>Journal of Heterocyclic Chemistry</i> , 1979 , 16, 25-28	1.9	2
14	Carbon-13 nuclear magnetic resonance spectra of phenycyclidine analogs. <i>Journal of Heterocyclic Chemistry</i> , 1979 , 16, 1425-1429	1.9	15
13	SYNTHESIS OF (R)-5-ALKYL-5-(1@METHYL-3@CARBOXYPROPYL) BARBITURIC ACIDS AND (R)-ALKYL-5-(1@METHYL-3-CARBOXYPROPYL)-2-THIOBARBITURIC ACIDS. <i>Organic Preparations and Procedures International</i> , 1978 , 10, 21-27	1.1	1
12	FORMYLATION OF ARENES BY ⊞,⊞-DICHLOROMETHYL METHYL ETHER. AN IMPROVED EXPERIMENTAL PROCEDURE. <i>Organic Preparations and Procedures International</i> , 1978 , 10, 201-204	1.1	11
11	Synthesis of Morphine-3-Glucuronide. <i>Synthetic Communications</i> , 1975 , 5, 231-236		
	Synthesis of Morphine 5 didedroniae. Synthetic Communications, 1213, 5, 251 250	1.7	18
10	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. Organic Preparations and Procedures International, 1975 , 7, 117-122	1.7	2
10	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. Organic Preparations		
	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. <i>Organic Preparations</i> and <i>Procedures International</i> , 1975 , 7, 117-122	1.1	2
9	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. Organic Preparations and Procedures International, 1975, 7, 117-122 4-PYRIDYLETHYLENE OXIDE. Organic Preparations and Procedures International, 1971, 3, 121-124 Synthesis of (3S,5R)-cis- and (3S,5S)-trans-3,5-Dimethylvalerolactones and Their Conversion to Biotransformation Products of (S)-5-Ethyl-5-(2?-pentyl)barbituric Acid. Synthetic Communications,	1.1	2
9	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. <i>Organic Preparations and Procedures International</i> , 1975 , 7, 117-122 4-PYRIDYLETHYLENE OXIDE. <i>Organic Preparations and Procedures International</i> , 1971 , 3, 121-124 Synthesis of (3S,5R)-cis- and (3S,5S)-trans-3,5-Dimethylvalerolactones and Their Conversion to Biotransformation Products of (S)-5-Ethyl-5-(2?-pentyl)barbituric Acid. <i>Synthetic Communications</i> , 1971 , 1, 169-174 Synthesis of 1,2-disubstituted naphth [1,2-d]imidazole-4,5-diones. <i>Journal of Heterocyclic Chemistry</i> ,	1.1	2 2 3
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9 8 7 6	SYNTHESIS OF METABOLITES OF PHENOBARBITAL AND MEPHOBARBITAL. <i>Organic Preparations and Procedures International</i> , 1975 , 7, 117-122 4-PYRIDYLETHYLENE OXIDE. <i>Organic Preparations and Procedures International</i> , 1971 , 3, 121-124 Synthesis of (3S,5R)-cis- and (3S,5S)-trans-3,5-Dimethylvalerolactones and Their Conversion to Biotransformation Products of (S)-5-Ethyl-5-(2?-pentyl)barbituric Acid. <i>Synthetic Communications</i> , 1971 , 1, 169-174 Synthesis of 1,2-disubstituted naphth [1,2-d]imidazole-4,5-diones. <i>Journal of Heterocyclic Chemistry</i> , 1970 , 7, 297-306 THE REACTION OF 2,3-DICHLORO-1,4-NAPHTHOQUINONE WITH p-NITROBENZHYDRAZIDE1. <i>Organic Preparations and Procedures</i> , 1970 , 2, 259-263	1.1	2 2 3 1

The synthesis of some 1-(Ediethylaminoethyl)-2-(p-ethoxybenzyl)-5-substituted benzimidazoles.

Journal of Heterocyclic Chemistry, 1967, 4, 262-267

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