

# Yuji Ishichi

## List of Publications by Year in descending order

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18  
papers

551  
citations

840776

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18  
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18  
docs citations

18  
times ranked

310  
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery of 4-chloro-2-(2,4-dichloro-6-methylphenoxy)-1-methyl-7-(pentan-3-yl)-1H-benzimidazole, a novel CRF1 receptor antagonist. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 1556-1570.	3.0	4
2	Design, synthesis, and biological evaluation of a novel series of peripheral-selective noradrenaline reuptake inhibitors – Part 3. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3716-3726.	3.0	8
3	Design, synthesis, and biological evaluation of a novel series of peripheral-selective noradrenaline reuptake inhibitors – Part 2. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3207-3217.	3.0	4
4	Design, synthesis and biological evaluation of a novel series of peripheral-selective noradrenaline reuptake inhibitor. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5000-5014.	3.0	7
5	Preparation of (S)-4-(1-(3,4-dichlorophenyl)-2-methoxyethyl)piperidine. <i>Tetrahedron: Asymmetry</i> , 2015, 26, 935-942.	1.8	3
6	Design, synthesis, and biological activities of 1-aryl-1,4-diazepan-2-one derivatives as novel triple reuptake inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3898-3902.	2.2	6
7	Novel triple reuptake inhibitors with low risk of CAD associated liabilities: Design, synthesis and biological activities of 4-[(1S)-1-(3,4-dichlorophenyl)-2-methoxyethyl]piperidine and related compounds. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 4600-4613.	3.0	11
8	Novel acetylcholinesterase inhibitor as increasing agent on rhythmic bladder contractions: SAR of 8-{3-[1-(3-fluorobenzyl)piperidin-4-yl]propanoyl}-1,2,5,6-tetrahydro-4H-pyrrolo[3,2,1-ij]quinolin-4-one (TAK-802) and related compounds. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 1901-1911.	3.0	13
9	Effects of TAK-802, a novel acetylcholinesterase inhibitor, on distension-induced rhythmic bladder contractions in rats and guinea pigs. <i>European Journal of Pharmacology</i> , 2004, 485, 299-305.	3.5	12
10	Amide-based atropisomers in tachykinin NK1-receptor antagonists: synthesis and antagonistic activity of axially chiral N-benzylcarboxamide derivatives of 2,3,4,5-tetrahydro-6H-pyrido[2,3-b][1,5]oxazocin-6-one. <i>Tetrahedron</i> , 2004, 60, 4481-4490.	1.9	35
11	Axially Chiral 1,7-Naphthyridine-6-carboxamide Derivatives as Orally Active Tachykinin NK1 Receptor Antagonists: A Synthesis, Antagonistic Activity, and Effects on Bladder Functions. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 3982-3993.	6.4	94
12	Axially Chiral N-Benzyl-N,7-dimethyl-5-phenyl-1,7-naphthyridine-6-carboxamide Derivatives as Tachykinin NK1 Receptor Antagonists: Determination of the Absolute Stereochemical Requirements. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 4232-4239.	6.4	54
13	Novel, Potent, and Orally Active Substance P Antagonists: Synthesis and Antagonist Activity of N-Benzylcarboxamide Derivatives of Pyrido[3,4-b]pyridine. <i>Journal of Medicinal Chemistry</i> , 1995, 38, 3106-3120.	6.4	77
14	Anodic oxidation of $\hat{\iota}$ -alkoxycarbonyloxy and $\hat{\iota}$ -acyloxy organotin compounds. <i>Tetrahedron Letters</i> , 1994, 35, 5247-5250.	1.4	17
15	Anodic cyclization of unsaturated $\hat{\iota}$ -stannyl ethers. Termination by bromide derived from dibromomethane. <i>Journal of the Chemical Society Chemical Communications</i> , 1994, , 2361-2362.	2.0	35
16	Intramolecular carbon-carbon bond formation by the anodic oxidation of unsaturated $\alpha$ -stannyl heteroatom compounds. Synthesis of fluorine containing heterocyclic compounds. <i>Journal of the American Chemical Society</i> , 1992, 114, 7594-7595.	13.7	113
17	Electrochemical oxidation of $\hat{\iota}$ -alkoxystannanes. <i>Tetrahedron Letters</i> , 1992, 33, 2599-2602.	1.4	36
18	A new method for the synthesis of acylsilanes via one-carbon homologation of aldehydes. <i>Journal of Organic Chemistry</i> , 1991, 56, 1307-1309.	3.2	22