Shunsuke Maehara

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Unique Antipsychotic Activities of the Selective Metabotropic Glutamate Receptor 1 Allosteric Antagonist 2-Cyclopropyl-5-[1-(2-fluoro-3-pyridinyl)-5-methyl-1 <i>H</i> -1,2,3-triazol-4-yl]-2,3-dihydro-1 <i>H</i> -isoindol-1-one Journal of Pharmacology and Experimental Therapeutics, 2009, 330, 179-190.	2.5	47
2	Antipsychotic property of a muscarinic receptor agonist in animal models for schizophrenia. Pharmacology Biochemistry and Behavior, 2008, 91, 140-149.	2.9	28
3	A Selective Phosphodiesterase 10A Inhibitor Reduces <scp>L</scp> â€Dopaâ€Induced Dyskinesias in Parkinsonian Monkeys. Movement Disorders, 2018, 33, 805-814.	3.9	19
4	Pharmacological characterization of a novel potent, selective, and orally active phosphodiesterase 10A inhibitor, PDMâ€042 [(E)â€4â€{2â€{2â€{5,8â€dimethylâ€{1,2,4]triazolo[1,5―a) Tj ETQq0 0 0 rgBT /Overle	ock 10 Tf	50 622 Td (]
	schizophrenia. Pharmacology Research and Perspectives, 2016, 4, e00241.		
5	Ameliorative effect of N-desmethylclozapine in animal models of social deficits and cognitive functions. Brain Research Bulletin, 2011, 86, 146-151.	3.0	10
6	Antipsychotic effects of N-desmethylclozapine on sensorimotor gating function in rats — Possible involvement of activation of M1 muscarinic receptors. European Journal of Pharmacology, 2011, 667, 242-249.	3.5	9
7	Behavioral effects of N-desmethylclozapine on locomotor activity and sensorimotor gating function in mice—Possible involvement of muscarinic receptors. Brain Research, 2011, 1418, 111-119.	2.2	9
8	Ameliorative effects of a phosphodiesterase 10A inhibitor, MR1916 on l-DOPA-induced dyskinesia in parkinsonian rats. Pharmacological Reports, 2020, 72, 443-448.	3.3	8
9	Dopamine D 1 signaling involvement in the effects of the phosphodiesterase 10A inhibitor, PDM-042 on cognitive function and extrapyramidal side effect in rats. Behavioural Brain Research, 2017, 317, 204-209.	2.2	5
10	Combination of the phosphodiesterase 10A inhibitor, MR1916 with risperidone shows additive antipsychoticâ€like effects without affecting cognitive enhancement and cataleptic effects in rats. Neuropsychopharmacology Reports, 2020, 40, 190-195.	2.3	5
11	Pharmacological characterization of a novel potent, selective, and orally active orexin 2 receptor antagonist, SDMâ€878. Neuropsychopharmacology Reports, 2020, 40, 182-189.	2.3	4
12	Pharmacological characterization of a novel, potent, selective, and orally active fatty acid amide hydrolase inhibitor, PKMâ€833 [(<i>R</i>)â€ <i>N</i> àê€(pyridazinâ€3â€yl)â€4â€(7â€(trifluoromethyl)chromanâ€4â€yl)piperazineâ€1â€carboxar Potential for the treatment of inflammatory pain. Pharmacology Research and Perspectives, 2020, 8, e00569.	nûdae]in ra	it s:
13	Pharmacological characterization of a novel potent, selective, and orally active phosphodiesterase 2A inhibitor, PDM-631. European Journal of Pharmacology, 2017, 811, 110-116.	3.5	2
14	Orexin 2 receptor is involved in orexin A-induced hyperlocomotion in rats. Pharmacological Reports, 2019, 71, 1147-1150.	3.3	1