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Cardiovascular response to 8-hydroxy-2-(di-n-propylamino) tetralin (8-OH-DPAT) in the rat: site of action and pharmacological analysis

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#	Paper	IF	Citations
202	Comparison of effects of some 5-HT1 agonists on blood pressure and heart rate of normotensive anaesthetized rats. **1987**, 140, 259-66		38
201	Evidence that the putative 5-HT1A receptor agonists, 8-OH-DPAT and ipsapirone, have a central hypotensive action that differs from that of clonidine in anaesthetised cats. **1987**, 138, 179-91		107
200	Communications. *British Journal of Pharmacology*, **1987**, 91, 293P-510P	8.6	0
199	COMMUNICATION. *British Journal of Pharmacology*, **1987**, 92, 511P-792p	8.6	1
198	Urapidil and some analogues with hypotensive properties show high affinities for 5-hydroxytryptamine (5-HT) binding sites of the 5-HT1A subtype and for alpha 1-adrenoceptor binding sites. **1987**, 336, 597-601		66
197	Further characterization of 5-hydroxytryptamine1-like receptors mediating tachycardia in the cat: No apparent relationship to known subtypes of the 5-hydroxytryptamine1 binding site. **1988**, 13, 245-258		9
196	Comparison between the cardiovascular effects of 8-hydroxy-2-(di-n-propylamino) tetralin (8-OH-DPAT) and clonidine in the conscious sino-aortic denervated rat. **1988**, 8, 267-76		6
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194	MDL 72832: a potent and stereoselective ligand at central and peripheral 5-HT1A receptors. **1988**, 149, 107-20		44
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187	Feeding responses to a high dose of 8-OH-DPAT in young and adult rats: influence of food texture. **1988**, 151, 267-73		17
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180	Antagonism by Ketanserin of 8-OH-DPAT-Induced Vasoconstriction. **1989**, 9, 43-44		
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