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(54) **COMPOUNDS COMPRISING
4-BENZOYLPIPERIDINE AS A
SIGMA-1-SELECTIVE LIGAND**

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A61M 36/14 (2006.01)

(52) **U.S. Cl.**
USPC **424/1.89**; 424/1.81; 546/187; 546/189

(58) **Field of Classification Search**
None
See application file for complete search history.

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(57) **ABSTRACT**

Bipiperidiny compounds and salts thereof are disclosed. The compounds include high affinity ligands for σ_1 receptors. Some compounds are also highly selective for σ_1 receptor compared to σ_2 receptor. Compounds can comprise radioisotopes, including ^{18}F or ^{11}C . Radiolabeled compounds can be used as probes for imaging distribution of σ_1 receptor in a subject such as a human using positron emission tomography (PET) scanning.

17 Claims, 4 Drawing Sheets

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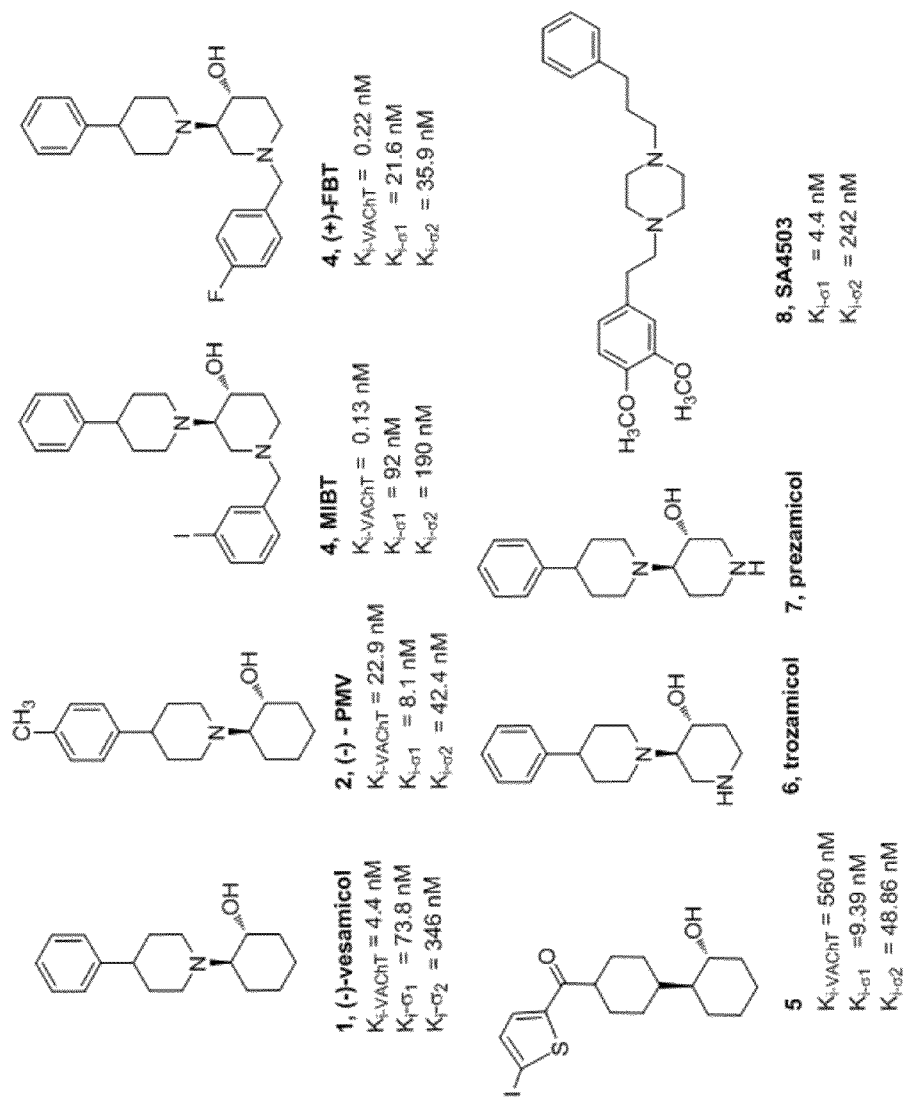


FIG. 1

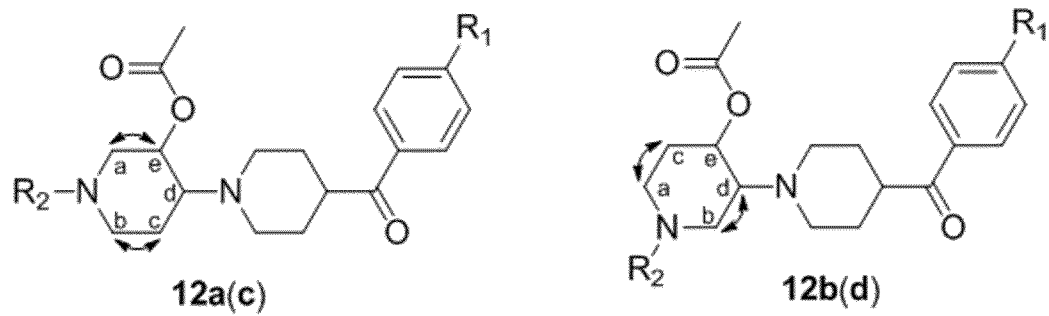


FIG. 2

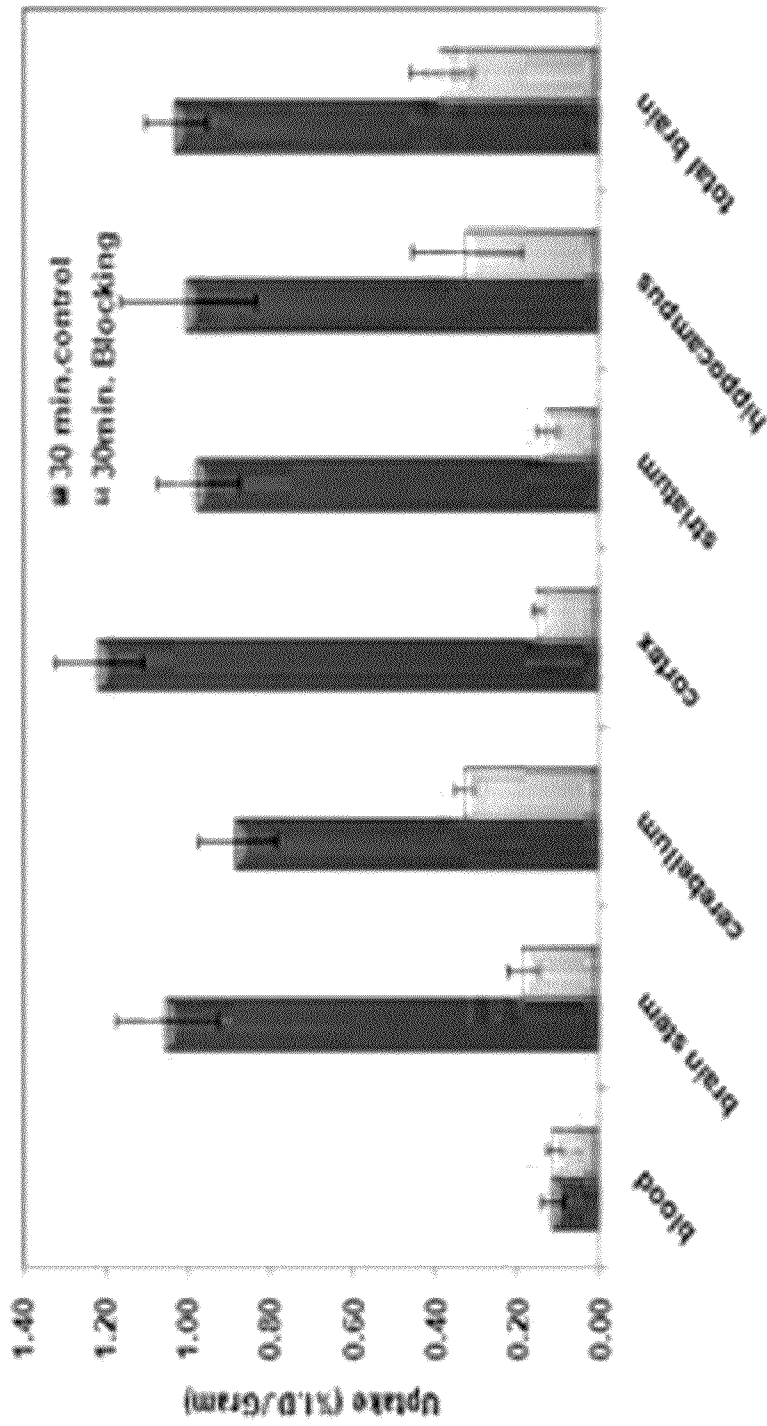


FIG. 3

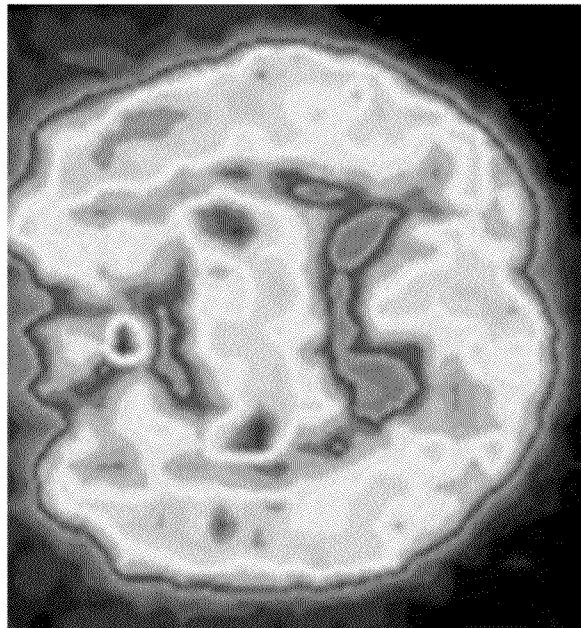
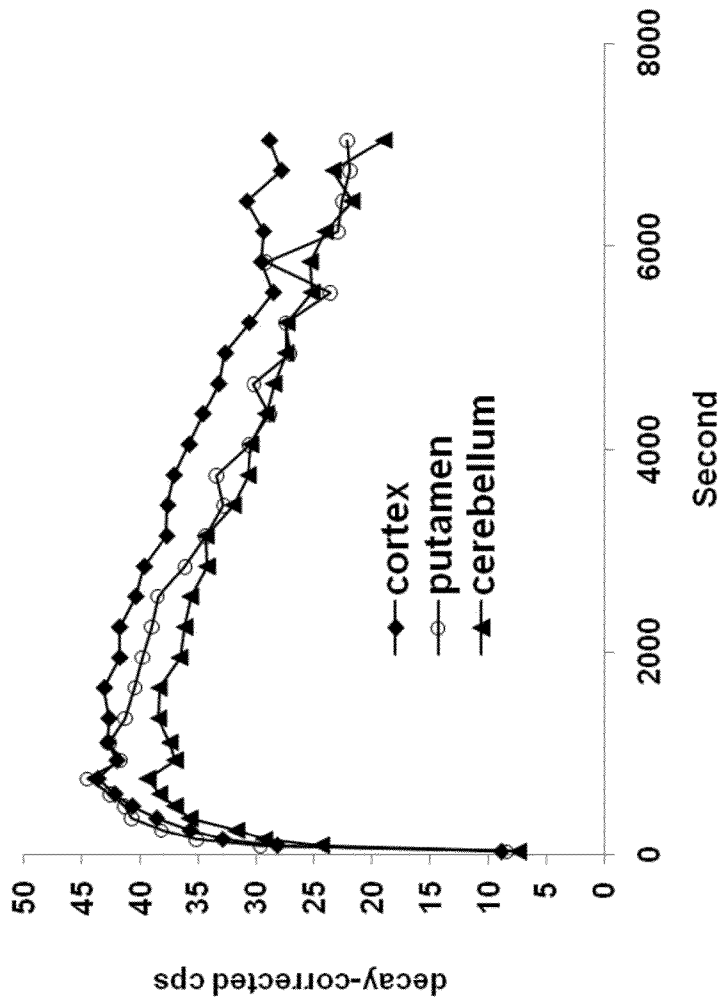


FIG. 4

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**COMPOUNDS COMPRISING
4-BENZOYLPIPERIDINE AS A
SIGMA-1-SELECTIVE LIGAND**

CROSS-REFERENCE TO RELATED
APPLICATIONS

This application claims the benefit of and priority to U.S. Provisional Application No. 61/356,997, filed on Jun. 21, 2010, which is incorporated herein by reference in its entirety.

GOVERNMENT SUPPORT

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INTRODUCTION

Sigma receptors are a class of receptors that are expressed in many normal tissues, including liver, kidneys, endocrine glands, and the central nervous system (CNS) (Walker, J. M., et al. *Pharmacol Rev* 42: 355-402 1990). It has been well established that there are at least two types of sigma receptors, sigma-1 (σ_1) and sigma-2 (σ_2) (Walker, J. M., et al. *Pharmacol Rev* 42, 355-402, 1990). Overexpression of σ_2 receptors has been reported in a variety of human and murine tumors (Bern, W. T., et al., *Cancer Res* 51: 6558-6562, 1991; Vilner, B. J., et al., In: *Multiple sigma and PCP receptor ligands: mechanisms for neuromodulation and neuroprotection*, Kamenka, J. M., and Domino, E. F., ed, Ann Arbor (Mich.), 7 NPP Books, p. 341-353, 1992; Mach, R. H., et al., *Cancer Res.* 57: 156-161, 1997). The human σ_1 receptor is a protein of 223 amino acids. σ_1 receptor is expressed in limbic and motor structures in the brain, and it is believed to modulate various neurotransmitter systems via Ca^{2+} -dependent cell signaling cascades. It has been reported that activation of σ_1 receptors is antidepressive, anxiolytic, neuroprotective, and cognition-enabling.

In the search of highly selective and clinically suitable σ_1 ligands, investigations have revealed that high affinity σ_1 ligands must contain a basic amine having two hydrophobic appendages.^{6,40} Structure-activity relationship analysis of vesamicol ligands (FIG. 1), which bind to the vesicular acetylcholine transporter (VACHT) in presynaptic cholinergic nerve terminals, found that many vesamicol ligands contain a pharmacophore that also has high affinity for σ_1 receptors.^{41, 42} In fact, some of the VACHT ligands show moderate to high binding affinities for σ_1 receptor. As shown in FIG. 1, (-)-vasamicol (1) has moderate σ_1 receptor binding affinity values of 73.5 nM, and its methyl substituted derivative (-)-p-methylvesamicol (2) has high σ_1 receptor binding affinity values of 8.10 nM; the trozamicol analogues meta-iodobenzyltrozamicol (3) and (+)-4-fluorobenzyltrozamicol (4) have σ_1 receptor binding affinity value of 92 and 21.6 nM respectively.⁴³ Several carbonyl containing VACHT analogues also bind to σ_1 very well; for example, (1S,2S)-2-(4-(5-iodothiophen-2-yl)piperidin-1-yl)cyclohexanol (5) has 9.39 nM affinity for K_i value of σ_1 receptors.⁴²⁻⁴⁹ To identify new, highly selective ligands for σ_1 receptors, our strategies are to (1) replace the 4-phenylpiperidiny group in presamicol and trozamicol structures with a 4-substituted benzoylpiperidiny group which is already known to favor a receptors,⁵⁰ (2) alkylate or acylate the secondary amine in (3'-hydroxy-1,4'-bipiperidin-4-yl)(4-substituted phenyl)methanone structurally similar to the presamicol scaffold,⁴³ and (3) alkylate the secondary amine in (4'-hydroxy-1,3'-bipiperidin-4-yl)(4-

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substituted phenyl)methanone, which is structurally similar to the trozamicol scaffold.⁴³ The present inventors identify the ligands that have high affinity for σ_1 receptors and selectivity for σ_1 versus σ_2 receptors. This investigation was inspired by 1) the observation that σ_1 receptor ligands may be potential therapeutic drugs for the treatment of neurological disorders³ and cancer³³ and 2) the need for highly selective and potent σ_1 receptor ligands that can be labeled with F-18 or C-11. Moreover, a novel clinical PET probe for imaging the σ_1 receptor provides a unique tool to assess the relationship between changes in σ_1 receptors in the brain during the progression of CNS disorders and provide a useful tool to monitor the treatment efficacy of the CNS disorders and cancer.

SUMMARY

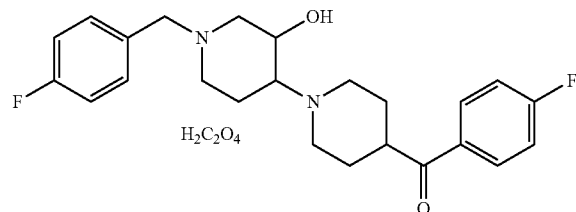
Imaging distribution of σ_1 receptors, e.g., by Positron Emission Tomography (PET) scanning, can be used diagnostically, e.g., to monitor progression of brain diseases or cancers or therapeutic efficacy of treatments. Furthermore, inhibitors of σ_1 receptors can be used clinically, e.g., for treatment of brain diseases and cancer in a subject in need of therapy.

The present inventors disclose various bipiperidiny compounds, and salts thereof. In some aspects, a compound or salt thereof of the present teachings can have activity as a Sigma receptor antagonist or inhibitor, and can show specificity in binding and/or inhibition for Sigma-1 (σ_1) receptors compared to Sigma-2 (σ_2) receptors. In some aspects, a compound or salt thereof of the present teachings can have activity as a vesicular acetylcholine transporter (VACHT) antagonist or inhibitor. In various aspects, a compound or salt thereof of the present teachings can serve as a probe for PET scanning.

In various embodiments, a bipiperidiny compound or salt thereof of the present teachings can comprise a 1,4'-bipiperidin-3'-ol, or a 1,3'-bipiperidin-4'-ol. In various embodiments, a compound or salt thereof of the present teachings can exhibit a K_i for a σ_1 receptor of $K_i < 5$ nM. In some embodiments, a compound or salt thereof of the present teachings can exhibit selectivity for σ_1 versus σ_2 receptors, e.g., $\sigma_1/\sigma_2 > 1000$ -fold. In some embodiments, a compound or salt thereof can exhibit low potency for VACHT, e.g. $K_i > 1000$ nM. In some embodiments, a compound or salt thereof can exhibit selectivity for σ_1 versus VACHT, e.g., $\sigma_1/VACHT > 1000$ -fold.

In some aspects, a compound or salt thereof of the present teachings can comprise a positron-emitting isotope, such as, for example, a positron-emitting isotope of fluorine, such as ¹⁸F, or a positron-emitting isotope of carbon, such as ¹¹C. In some aspects, a compound or salt thereof of the present teachings can be used as a tracer for PET scanning.

In an embodiment, a compound or salt thereof of the present teachings can be 14a of structure

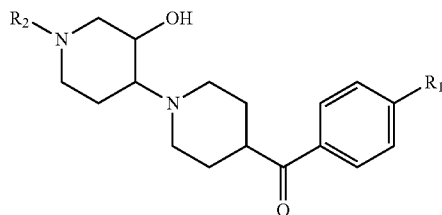


In some aspects, $H_2C_2O_4$ can be oxalic acid. In some aspects, 14a can exhibit subnanomolar σ_1 affinity, such as a K_i of 0.48

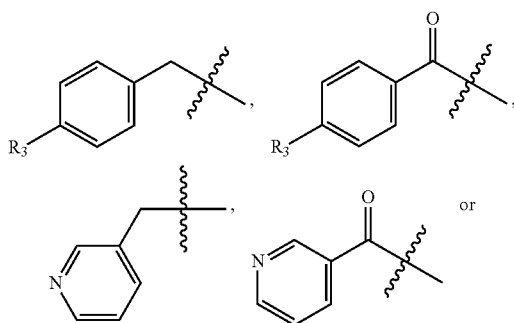
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nM. In various aspects, 14a can exhibit a selectivity for σ_1 versus σ_2 of about 1360-fold, and selectivity of about 3600-fold for σ_1 versus VACHT.

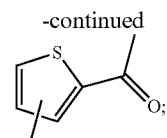
In some aspects, a compound of the present teachings can have a structure



wherein R_1 can be F or methoxy, R_2 can be



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R_3 can be F or methoxy. In various configurations, an F can be an ^{18}F , and a methoxy can comprise a ^{11}C . In various configurations, in a structure of the present teachings comprising more than one fluorine atom, one or more fluorine atoms can be an ^{18}F . In various configurations, in a structure of the present teachings comprising more than one methoxy group, one or more carbon atoms of a methoxy group can be a ^{11}C .

The moderate to high affinity of vesamicol and many of its analogues for σ_1 and σ_2 receptors reduces the selectivity of these compounds for the VACHT and potentially compromises their utility as VACHT ligands. Consequently, we also screened the target compounds for sigma binding to identify selective VACHT ligands from this new structural class of agents. Among the compounds tested, there was wide variability in affinity for σ_1 or σ_2 receptors; only one compound (TZ-3-156) displayed high affinity for σ_1 receptors ($K_i=13.23\pm 0.48$ nM), while all other compounds exhibited poor affinity for σ receptors.

In various aspects, a structure of the present teachings can be a structure set forth herein. Table 1 sets forth K_i and Log P values for disclosed structures with respect to σ_1 , σ_2 and VACHT. Table 2 sets forth elemental analysis of analogues of disclosed structures. Table 3 sets forth structures of VACHT specific compounds. Table 4 sets forth K_i values of VACHT specific compounds.

TABLE 1

Affinities of new analogues for σ_1 receptor, σ_2 receptor, and VACHT. ^a						
Compound	σ_1 (K_i , nM) ^b	σ_2 (K_i , nM) ^c	VACHT (K_i , nM) ^d	σ_1/σ_2 selectivity	σ_1/VACHT selectivity	LogP ^e
14a	0.48 ± 0.14	1741 ± 286	1360 ± 295	3627	2833	2.83
14b	4.03 ± 0.47	5521 ± 1352	3310 ± 907	1370	821	2.61
14c	1.36 ± 0.28	2301 ± 249	401 ± 42.0	1692	295	2.73
14d	22.8 ± 2.32	4208 ± 115	2030 ± 385	184	89	1.48
14e	2.51 ± 0.34	2788 ± 718	294 ± 16.1	1111	117	2.82
14f	25.9 ± 0.96	5157 ± 202	14800 ± 3460	199	569	2.61
14g	4.05 ± 0.88	3033 ± 248	44.2 ± 3.03	749	11	2.72
14h	59.64 ± 2.22	4540 ± 1606	137 ± 14.3	76	2.2	1.46
15a	3144 ± 140	8642 ± 812	3600 ± 499	2.7	1.1	1.92
15b	2238 ± 271	>10000	30900 ± 6400	>4.5	14	2.25
15c	2833 ± 374	20450 ± 4887	683 ± 90	7.2	0.24	1.67
15d	7739 ± 662	13565 ± 1435	3340 ± 706	1.7	0.43	0.57
15g	2088 ± 154	32850 ± 443	3460 ± 476	15.7	1.66	1.90
15h	2061 ± 113	15862 ± 3045	555 ± 65.4	7.7	0.27	1.65
15i	26646 ± 8738	17041 ± 8626	372 ± 65.7	0.64	0.01	0.56
16a	50.0 ± 7.9	3443 ± 928	136 ± 13.8	69	2.72	1.98
16b	91.1 ± 19.9	4979 ± 507	1970 ± 196	54	22	1.73
16c	106 ± 28	832 ± 147	149 ± 22.5	7.8	1.4	1.88
16d	1159 ± 128	13018 ± 2626	237 ± 41.4	11	0.20	0.79
16e	137 ± 21	5598 ± 1033	48.6 ± 8.37	41	0.35	2.00
16f	297 ± 27	4208 ± 439	1080 ± 296	14	3.6	1.75
16g	208 ± 53	8539 ± 1900	35.5 ± 11.1	41	0.17	1.89
16h	2225 ± 168	17135 ± 3863	107 ± 11	7.7	0.40	0.80

^a K_i values (mean ± SEM) were determined in at least three experiments.

^b The σ_1 binding assay used membrane preparations of guinea pig brain.

^c The σ_2 binding assay used homogenates of rat liver.

^d The VACHT binding assay used expressed human VACHT.

^e Calculated value at pH 7.4 by ACD/L- Lab, version 7.0. (Advanced Chemistry Development, Inc., Canada).

TABLE 2

Elemental Analysis of Analogues							
Compound	Molecular Formula	Calculated			Found		
		C	H	N	C	H	N
14a	C ₂₄ H ₂₈ F ₂ N ₂ O ₂ •H ₂ C ₂ O ₄ •0.25H ₂ O	61.35	6.04	5.50	61.43	5.91	5.54
14b	C ₂₅ H ₃₁ FN ₂ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	61.70	6.52	5.33	62.09	6.41	5.30
14c	C ₂₄ H ₂₉ FN ₂ O ₂ •H ₂ C ₂ O ₄ •0.25H ₂ O	63.60	6.47	5.70	63.52	6.47	5.67
14d	C ₂₃ H ₂₈ FN ₃ O ₂ •1.5H ₂ C ₂ O ₄ •H ₂ O	56.72	6.04	7.63	56.88	5.93	7.31
14e	C ₂₅ H ₃₁ FN ₂ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	61.70	6.52	5.33	61.56	6.28	5.24
14f	C ₂₅ H ₃₁ FN ₂ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	62.56	6.94	5.21	62.42	6.78	5.19
14g	C ₂₅ H ₃₂ N ₂ O ₃ •2H ₂ C ₂ O ₄	65.04	6.87	5.62	64.58	6.72	5.56
14h	C ₂₄ H ₃₁ N ₃ O ₃ •2H ₂ C ₂ O ₄ •0.5H ₂ O	56.18	6.06	7.02	56.37	5.98	6.82
15a	C ₂₄ H ₂₆ F ₂ N ₂ O ₅ •H ₂ C ₂ O ₄	60.23	5.44	5.40	59.96	5.43	5.37
15b	C ₂₄ H ₂₇ FN ₂ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	60.10	5.98	5.19	59.99	6.07	5.14
15c	C ₂₄ H ₂₇ FN ₂ O ₃ •H ₂ C ₂ O ₄ •H ₂ O	60.22	6.03	5.40	60.72	5.98	5.50
15d	C ₂₃ H ₂₆ FN ₃ O ₃ •H ₂ C ₂ O ₄ •0.25H ₂ O	59.34	5.61	8.30	59.25	5.55	8.26
15e	C ₂₅ H ₂₉ FN ₂ O ₄ •H ₂ C ₂ O ₄	61.12	5.89	5.28	61.03	5.92	5.26
15f	C ₂₅ H ₃₀ N ₂ O ₈ •H ₂ C ₂ O ₄ •1.5H ₂ O	60.10	6.54	5.19	60.47	6.39	5.41
15g	C ₂₄ H ₂₉ N ₃ O ₄ •H ₂ C ₂ O ₄	60.81	6.08	8.18	60.57	6.06	7.96
16a	C ₂₄ H ₂₈ F ₂ N ₂ O ₂ •H ₂ C ₂ O ₄	61.90	5.99	5.55	61.78	5.97	5.52
16b	C ₂₅ H ₃₁ FN ₂ O ₃ •H ₂ C ₂ O ₄	62.78	6.44	5.42	62.61	6.32	5.40
16c	C ₂₄ H ₂₉ FN ₂ O ₃ •H ₂ C ₂ O ₄	64.18	6.42	5.76	63.95	6.40	5.71
16d	C ₂₃ H ₂₈ FN ₃ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	60.47	6.29	8.46	60.59	6.29	8.41
16e	C ₂₅ H ₃₁ FN ₂ O ₃ •H ₂ C ₂ O ₄	52.78	6.44	5.42	62.55	6.49	5.46
16f	C ₂₆ H ₃₄ N ₂ O ₄ •H ₂ C ₂ O ₄	63.62	6.86	5.30	63.33	6.87	5.27
16g	C ₂₅ H ₃₂ N ₂ O ₃ •H ₂ C ₂ O ₄ •0.5H ₂ O	63.89	6.95	5.52	63.93	6.85	5.51
16h	C ₂₄ H ₃₁ N ₃ O ₃ •H ₂ C ₂ O ₄	62.51	6.66	8.41	62.14	6.72	8.27

30

TABLE 3

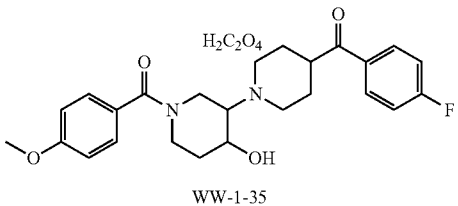
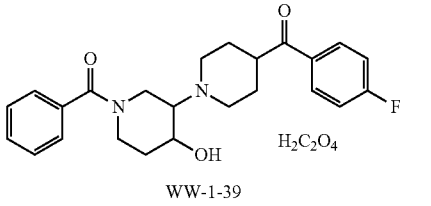
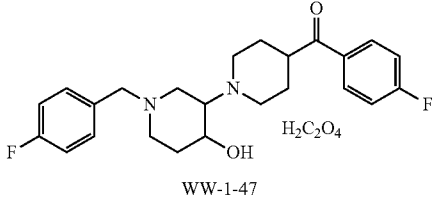
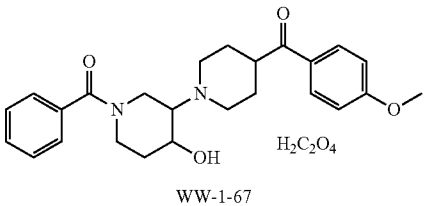
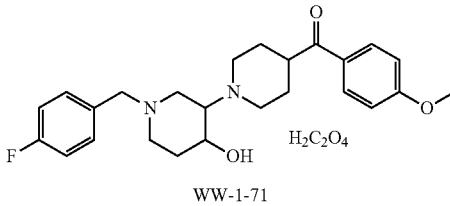
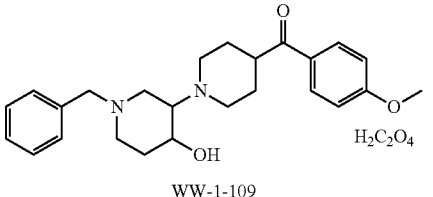
Structures of VACHT specific compounds		
Name	#	Structure
WW-1-35	1	
WW-1-39	2	
WW-1-47	3	

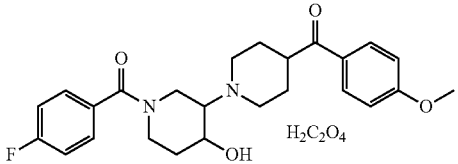
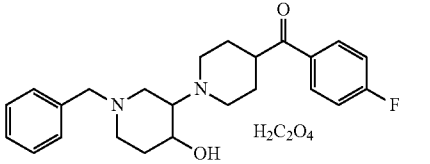
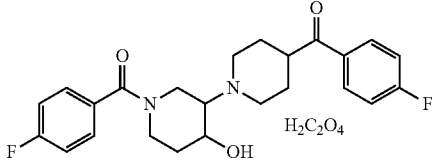
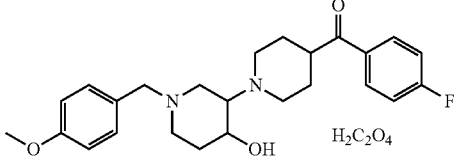
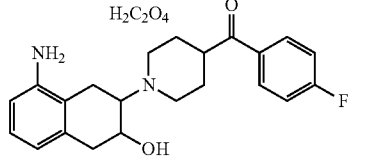
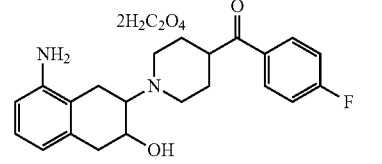
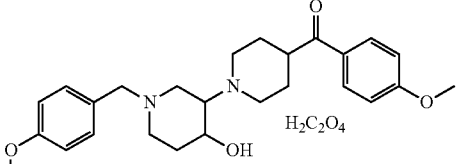
TABLE 3-continued

Structures of VACHT specific compounds		
Name	#	Structure
WW-1-67	4	
WW-1-71	5	
WW-1-109	6	

65

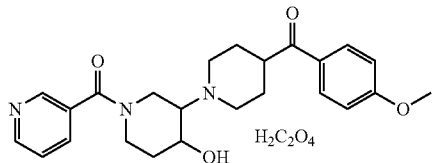
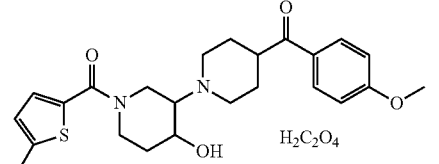
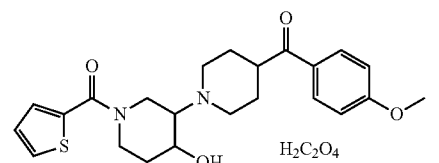
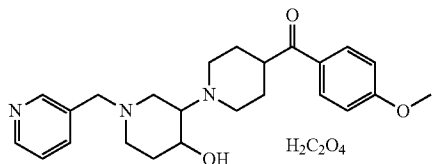
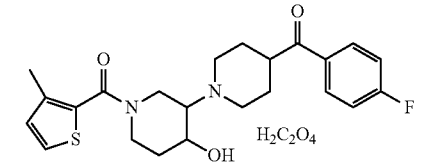
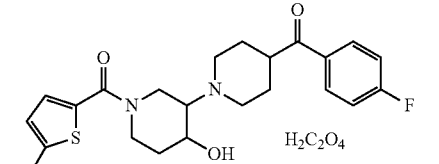
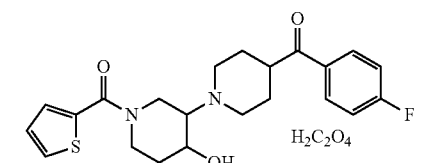
7

TABLE 3-continued

Name #		Structure
Structures of VACHT specific compounds		
WW-1-107	7	 WW-1-107
WW-1-117	8	 WW-1-117
WW-1-123	9	 WW-1-123
WW-1-127	10	 WW-1-127
WW-1-88-1	11	 WW-1-88-1
isomer	(+)	
WW-1-88-2	12	 WW-1-88-2
isomer	(-)	
WW-111-10	13	 WW-111-10

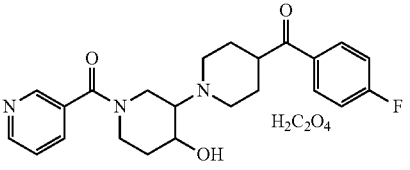
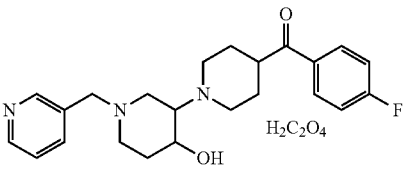
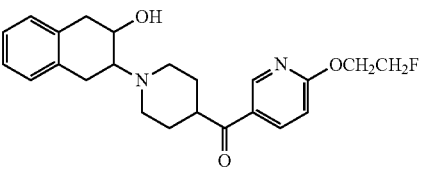
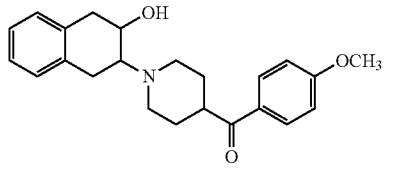
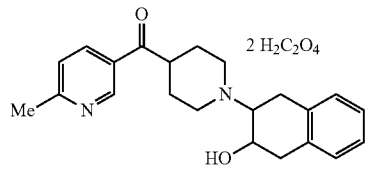
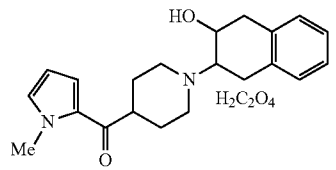
8

TABLE 3-continued

Name #		Structure
Structures of VACHT specific compounds		
WW-111-13	14	 WW-111-13
WW-III-17	15	 WW-III-17
WW-III-21	15	 WW-III-21
WW-III-25	16	 WW-III-25
WW-III-39	17	 WW-III-39
WW-III-41	18	 WW-III-41
WW-III-43	19	 WW-III-43

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TABLE 3-continued

Structures of VACHT specific compounds		
Name	#	Structure
WW-III-45	20	 WW-III-45
WW-III-47	21	 WW-III-47
DZT-I-137	22	 0.5H ₂ C ₂ O ₄
DZT-I-107	23	 0.5H ₂ C ₂ O ₄
OS-5-40	24	 OS-5-40
OS-5-60	25	 OS-5-60

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TABLE 3-continued

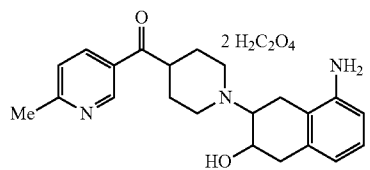
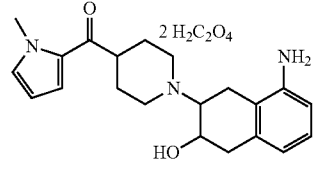
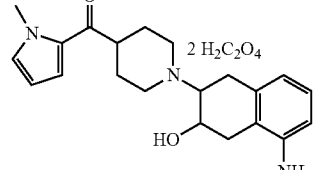
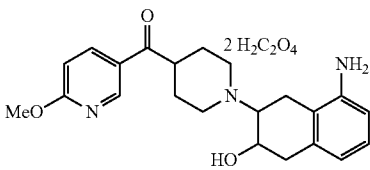
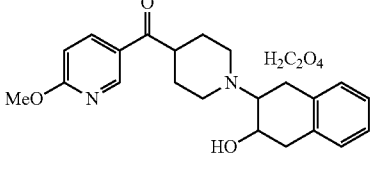
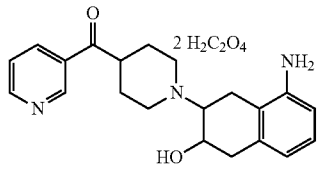
Structures of VACHT specific compounds		
Name	#	Structure
OS-5-70	26	 OS-5-70
OS-5-72-1	27	 OS-5-72-1
OS-5-72-2	28	 OS-5-72-2 C ₂₅ H ₃₁ N ₃ O ₁₀ MW: 533.53
OS-5-82	29	 OS-5-82
OS-5-88	30	 OS-5-88
OS-5-110	31	 OS-5-110

TABLE 4

Ki values of VAcHT specific compounds					
Name	Formula # (MW)	Sigma-1 σ_1 (nM)	Sigma-2 σ_2 (nM)	VAcHT (nM)	Log P
WW-1-35	1 C ₂₇ H ₃₁ FN ₂ O ₈ 530.54	661 ± 72	>10000 nM	13000 ± 1900	2.25
WW-1-39	2 C ₂₆ H ₂₉ FN ₂ O ₇ + H ₂ O 518.53	37885 ± 271	3946 ± 278	62.7 ± 14.3	1.75
WW-1-47	3 C ₂₆ H ₃₀ F ₂ N ₂ O ₆ 504.52	50 ± 7.9	3443 ± 928	136 ± 13.8	1.98
WW-1-67	4 C ₂₇ H ₃₂ N ₂ O ₈ + H ₂ O 530.54	5760 ± 477	4847 ± 649	15.4 ± 0.94	1.73
WW-1-71	5 C ₂₇ H ₃₃ FN ₂ O ₇ 516.56	137 ± 21	5598 ± 1033	48.6 ± 8.37	2.00
WW-1-109	6 C ₂₇ H ₃₄ N ₂ O ₇ + 0.5H ₂ O 507.58	208 ± 53	8539 ± 1900	35.5 ± 11.1	1.89
WW-1-107	7 C ₂₇ H ₃₁ FN ₂ O ₈ + 1.5H ₂ O 557.56	11932 ± 5613	2597 ± 363	233 ± 36.8	1.98
WW-1-117	8 C ₂₆ H ₃₁ FN ₂ O ₆ 486.53	106 ± 28	832 ± 147	149 ± 22.5	1.88
WW-1-123	9 C ₂₆ H ₂₈ F ₂ N ₂ O ₇ + 0.5H ₂ O 527.51	3119 ± 360	1375 ± 135	8700 ± 1620	2.00
WW-1-127	10 C ₂₇ H ₃₃ FN ₂ O ₇ 516.56	91.1 ± 19.9	4979 ± 507	1970 ± 196	1.73
WW-1-88-1	11 C ₂₄ H ₂₇ FN ₂ O ₆ + H ₂ O 476.49	3693 ± 537	14717 ± 1936	366 ± 36.0	2.06
WW-1-88-2	12 C ₂₆ H ₂₉ FN ₂ O ₁₀ + 1.25H ₂ O 571.03	3553 ± 438	6889 ± 546	20.9 ± 4.98	2.06
WW-III-10	13 C ₂₈ H ₃₆ N ₂ O ₈	297.41 ± 27.22	4207.71 ± 439.10	1080 ± 296	1.75
WW-III-13	14 C ₂₆ H ₃₁ N ₃ O ₈ + 1.5H ₂ O	8601.17 ± 375.62	24270.32 ± 1910.12	112 ± 12.9	0.65
WW-III-17	C ₂₆ H ₃₁ N ₂ O ₈ S + 0.5H ₂ O	39396.39 ± 4099.55	24439.19 ± 3627.47	142 ± 16.4	2.51
WW-III-21	15 C ₂₅ H ₃₀ N ₂ O ₈ S + 0.5H ₂ O	33960.81 ± 6118.62	6377.96 ± 1002.53	19.0 ± 2.12	2.05
WW-III-25	16 C ₂₆ H ₃₃ N ₃ O ₇	2225.99 ± 167.98	17135.24 ± 3863.84	107 ± 11.1	0.8
WW-III-39	17 C ₂₅ H ₂₉ FN ₂ O ₇ S + 0.25H ₂ O	12125.27 ± 2447.43	4219.28 ± 204.70	11.4 ± 3.67	2.53
WW-III-41	18 C ₂₅ H ₂₉ FN ₂ O ₇ S	2800.43 ± 210.38	16418.19 ± 1479.77	1190 ± 137	2.53
WW-III-43	19 C ₂₄ H ₂₇ FN ₂ O ₇ S + H ₂ O	22509.06 ± 5877.28	4046.14 ± 244.35	123 ± 14.9	2.07
WW-III-45	20 C ₂₅ H ₂₈ FN ₃ O ₇ + H ₂ O	13796.39 ± 2628.81	15496.51 ± 2172.13	807 ± 139	0.66
WW-III-47	21 C ₂₅ H ₃₀ FN ₃ O ₆ + 0.5H ₂ O	1159.95 ± 128.12	13018.08 ± 2626.50	237 ± 41.4	0.79
DZT-I-137	22 C ₂₅ H ₂₈ FN ₂ O ₅ 443.20	2122.26 ± 49.01	1859.79 ± 144.36	38 ± 3.84	2.87
DZT-I-107	23 C ₂₄ H ₂₈ NO ₅ 410.20	132.00 ± 10.10	1952.95 ± 13.12	4.64 ± 0.47	2.94
OS-5-40	24 C ₂₆ H ₃₀ N ₂ O ₁₀ MW: 530.52	2115 ± 289	3771 ± 886	27.9 ± 7.99	2.29
OS-5-60	25 C ₂₃ H ₂₈ N ₂ O ₆ MW: 428.48	1302 ± 93	2553 ± 339	18.4 ± 2.54	1.42
OS-5-70	26 C ₂₆ H ₃₁ N ₃ O ₁₀ MW: 545.54	11362 ± 2040	27525 ± 6943	48.0 ± 11.4	1.33
OS-5-72-1	27 C ₂₅ H ₃₁ N ₃ O ₁₀ MW: 533.53	9237 ± 793	11396 ± 2500	38.7 ± 6.95	0.53
OS-5-72-2	28 C ₂₅ H ₃₁ N ₃ O ₁₀ MW: 533.53	7664 ± 233	6987 ± 1482	2310 ± 393	0.48
OS-5-82	29 C ₂₆ H ₃₁ N ₃ O ₁₁ MW: 561.54	5682 ± 26	10813 ± 1282	23.3 ± 3.43	1.66
OS-5-88	30 C ₂₄ H ₂₈ N ₂ O ₇ MW: 456.49	986 ± 160	2572 ± 139	8.36 ± 0.68	2.64
OS-5-110	31 C ₂₅ H ₂₉ N ₃ O ₁₀ MW: 531.51	20346 ± 8133	5078 ± 221	2.2	0.87

Structures of the present teachings can be synthesized using standard methods of organic synthesis following well known principles such as set forth in Tu, Z., et al., *J. Med. Chem.* 52: 1358-1369, 2009; Efang, S. M. N., et al., *J. Med. Chem.* 53: 2825-2835, 2006; Hedrickson et al., *Organic Chemistry* 3rd edition, McGraw Hill, New York, 1970; Caruthers, W., and Coldham, I., *Modern Methods of Organic Synthesis* (4th Edition), Cambridge University Press, Cambridge, U.K., 2004. Methods of imaging using PET scanning are set forth in references such as Curati, W. L., *Imaging in Oncology*, Cambridge University Press, Cambridge, U.K., 1998; and Welch, M. J., and Redvanly, C. S., eds. *Handbook of Radiopharmaceuticals: Radiochemistry and Applications*, J. Wiley, N.Y., 2003.

The present inventors synthesized ((3'R,4'R)-1'-(4-fluorobenzyl)-3'-hydroxy-1,4'-bipiperidin-4-yl)(4-[¹¹C]methoxy-

phenyl)methanone (14e), which has high affinity for σ_1 receptors ($K_i=2.51\pm 0.34$ nM, $\sigma_1/\sigma_2>1100$ -fold, log P=2.82) and can be labeled with either C-11 or F-18.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates prior art vesamicol ligands which bind to the vesicular acetylcholine transporter (VAcHT) in presynaptic cholinergic nerve terminals.

FIG. 2 illustrates assignment of regioisomers of 12a(c) and 12b(d) by ¹H-¹H COSY.

FIG. 3 illustrates the uptake of [¹¹C]11e in control rats.

FIG. 4 illustrates transverse image of representative MicroPET images (summed from 0-120 min) of 2 hr dynamic scans

in cynologous Monkeys (left); and time tissue-activity curves for cortex, putamen and cerebellum (right).

DETAILED DESCRIPTION

The present inventors have synthesized a new class of compounds targeting σ_1 receptors. The inventors explored carbonyl-containing VAcHT inhibitors by introducing substituted benzyl or benzoyl groups on the nitrogen of the terminal piperidine ring.

The present inventors have developed five racemic compounds, 14a (0.48 nM), 14b (4.03 nM), 14c (1.36 nM), 14e (2.51 nM), and 14g (4.05 nM), that have very high binding affinities for σ_1 receptors. The first four compounds also display high selectivity for σ_1 vs. σ_2 receptors and do not bind well to VAcHT. They can be useful pharmacological agents targeting σ_1 receptors. Fluorine-18 or carbon-11 isotopes can be introduced into these compounds and can be used as PET probes PET probes for imaging σ_1 receptors in human beings and animals. Particularly for 14a and 14c, the σ_1/σ_2 ratios are greater than 3600 and 1600 fold and the $\sigma_1/VAcHT$ ratios are greater than 2800 fold and 290 fold, respectively. The fluorine-18 or carbon-11 versions of 14a and 14c can be used to quantify the density of σ_1 receptors in human beings or animals. In addition, the SAR information from the current study provides new insight into σ_1 receptor ligands.

Abbreviations List:

Anal., Analysis; BOP-Cl, bis(2-oxo-3-oxazolidinyl)phosphonic chloride; Calcd., calculated; CIMS., Chemical ionization mass spectrometry; CNS, central nervous system; SAR, structure-activity relationship; DCC, N,N'-dicyclohexylcarbodiimide; DMF, N,N-dimethylformamide; DMSO, dimethyl sulfoxide; DTG, 1,3-ditolyguanidine; ND, not determined; PET, positron emission tomography; prezmicol: 3-hydroxy-4-(4-phenylpiperidinyl)piperidine; SPECT, single photon emission computed tomography; THF, tetrahydrofuran; TLC, thin layer chromatography; trozamicol, 4-hydroxy-3-(4-phenylpiperidinyl)piperidine; VAcHT, vesicular acetylcholine transporter; vesamicol, (-)-trans-2-(4-phenylpiperidino)cyclohexanol.

Experimental Section

General.

All reagents and chemicals were purchased from commercial suppliers and used without further purification unless otherwise stated. All anhydrous reactions were carried out in an oven-dried glassware under nitrogen unless otherwise stated. Flash column chromatography was conducted using silica gel 60A, "40 Micron Flash" [32-63 μ m] (Scientific Absorbents, Inc.); the mobile phase used is reported in the experimental procedure for each compound. Melting points were determined using the MEL-TEMP 3.0 apparatus (Laboratory Devices Inc.) and are uncorrected. 1H NMR spectra were recorded at 300 MHz on a Mercury-VX spectrometer (Varian, Inc.) with $CDCl_3$ as solvent and tetramethylsilane (TMS) as the internal standard unless otherwise stated. All chemical shift values are reported in parts per million (PPM) (δ). Peak multiplicities are singlet, s; doublet, d; triplet, t; multiplet, m; broad, br. Elemental analyses (C, H, N) were determined by Atlantic Microlab, Inc. Elemental analysis was used to determine the purity of the target compounds that were assessed with biological data. All the compounds reported in the manuscript have a purity of $\geq 95\%$.

(5,6-Dihydropyridin-1(2H)-yl)(phenyl)methanone
(10)

Into a solution of 1,2,3,6-tetrahydropyridine (1.00 g, 12.0 mmol) and triethylamine (3 mL) in CH_2Cl_2 (30 mL), benzoyl

chloride (1.69 g, 12.0 mmol) was added dropwise at 0° C. The reaction mixture was stirred at room temperature for 2 hrs and washed with saturated Na_2CO_3 (20 mL \times 3) and brine solution (20 mL), dried over Na_2SO_4 and concentrated under vacuum to afford crude product. The crude product was purified by silica gel column chromatography using hexane/ethyl acetate (10/3, v/v) to afford target product as a colorless oil (2.27 g, 100%). 1H NMR ($CDCl_3$): δ 2.16-2.25 (m, 2H), 3.46-4.20 (m, 4H), 5.53-5.91 (m, 2H), 7.29-7.53 (m, 5H).

7-Oxa-3-azabicyclo[4.1.0]heptan-3-yl(phenyl)
methanone (11)

Into a solution of 10 (2.27 g, 12.1 mmol) in CH_2Cl_2 (20 mL), a solution of 3-chloroperoxybenzoic acid (5.43 g, 77% pure, 24.3 mmol) in CH_2Cl_2 (40 mL) was added dropwise at 0° C. The reaction mixture was stirred at room temperature for 5 hrs. After the reaction was complete as determined by TLC (hexane/ethyl acetate, 1/1, v/v), saturated Na_2CO_3 solution (50 mL) was added into the reaction vial slowly with stirring. The reaction mixture was stirred for 30 min. The organic layer was washed with saturated Na_2CO_3 solution (50 mL \times 3), brine solution (50 mL), dried over Na_2SO_4 and concentrated under reduced pressure. The crude product was purified by silica gel column chromatograph using hexane/ethyl acetate (3/1, v/v) as the mobile phase to afford the product as a sticky oil (1.54 g, 62%). 1H NMR ($CDCl_3$): δ 2.04-2.18 (m, 2H), 3.09-4.40 (m, 6H), 7.28-7.47 (m, 5H).

Procedure A: General Method for Preparing 1'-Substituted Benzoyl-4-acyl-[1,4'-bipiperidin]-3'-yl Acetate (12a and 12c) or 1'-Substituted Benzoyl-4-acyl-[1,3'-bipiperidin]-4'-yl Acetate (12b and 12d)

1'-Benzoyl-4-(4-fluorobenzoyl)-[1,4'-bipiperidin]-3'-yl acetate (12a) and 1'-benzoyl-4-(4-methoxybenzoyl)-1,3'-bipiperidin-4'-yl acetate (12b)

A mixture of 11 (0.50 g, 1.97 mmol), 4-(4-fluorobenzoyl) piperidine hydrochloride (1.20 g, 4.92 mmol), triethylamine (1 mL) and sodium carbonate (2.12 g, 20.0 mmol) in ethanol (50 mL) was heated to 70° C. with stirring for over 24 hrs. The reaction mixture was cooled, filtered and concentrated under reduced pressure. The crude product was dissolved in CH_2Cl_2 (50 mL) and washed with water (50 mL), saturated Na_2CO_3 (50 mL) and brine solution. The organic layer was dried over Na_2SO_4 and concentrated under reduced pressure to afford crude product as an oil. Acetic anhydride (1.11 g, 10.92 mmol) was added into a solution of this crude product in CH_2Cl_2 (30 mL). The reaction mixture was stirred at room temperature overnight. The mixture was washed with saturated Na_2CO_3 (30 mL \times 3) and brine solution. The organic layer was dried over Na_2SO_4 and concentrated under reduced pressure. The crude product was purified by silica gel column chromatography using ethyl acetate/hexane (2/1, v/v) as the mobile phase to afford 12a (0.41 g, 37%), the first eluted compound as a colorless oil. 1H NMR ($CDCl_3$): δ 1.40-1.74 (m, 3H), 2.04 (s, 3H), 2.30-2.37 (m, 1H), 2.51-2.69 (m, 2H), 2.86-3.21 (m, 6H), 4.30-4.70 (m, 1H), 4.80-5.10 (br s, 1H), 7.10-7.16 (t, J=5.7 Hz, 2H), 7.40-7.41 (m, 5H), 7.93-7.97 (m, 2H). At the same time, 1'-benzoyl-4-(4-fluorobenzoyl)-[1,3'-bipiperidin]-4'-yl acetate (12b) (0.50 g, 45%) was eluted as the second compound and 12b was a colorless oil. 1H NMR ($CDCl_3$): δ 1.50-1.90 (m, 3H), 2.07 (s, 3H), 2.81-3.29 (m, 6H), 2.20-2.80 (m, 4H), 3.50-4.00 (m, 2H), 4.64 (br s, 1H), 5.10-5.18 (m, 1H), 7.12 (t, J=5.7 Hz, 2H), 7.41-7.42 (m, 5H), 7.93 (m, 2H).

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1'-Benzoyl-4-(4-methoxybenzoyl)-[1,4'-bipiperidin]-3'-yl acetate (12c) and 1'-benzoyl-4-(4-methoxybenzoyl)-1,3'-bipiperidin-4'-yl acetate (12d)

Procedure A was followed to prepare 12c (0.42 g, 37%) as a colorless oil that was eluted first and 12d (0.34 g, 37%) as a colorless oil that was eluted second. The ¹H NMR of 12c (CDCl₃): δ 1.40-1.90 (m, 8H), 2.04 (s, 3H), 2.22-2.40 (m, 1H), 2.24-2.70 (m, 2), 2.80-3.22 (m, 4H), 3.87 (s, 3H), 4.00-4.60 (m, 1H), 6.94 (d, J=8.7 Hz, 2H), 7.41 (s, 5H), 7.91 (d, J=8.7 Hz, 2H). The ¹H NMR of 12d (CDCl₃): δ 1.45-1.90 (m, 7H), 2.04 (s, 3H), 2.22-2.75 (m, 2H), 2.80-3.30 (m, 5H), 3.64 (br s, 1H), 3.86 (s, 3H), 4.64 (br s, 1H), 5.10-5.20 (m, 1H), 6.92 (d, J=8.1 Hz, 2H), 7.64 (br s, 5H), 7.89 (m, 2H).

Procedure B. General Method for Preparing (3'-Hydroxy-[1,4'-bipiperidin]-4-yl)(4-Substituted-phenyl)Methanone (13a and 13c) and (4'-Hydroxy-[1,3'-bipiperidin]-4-yl)(4-Substituted-phenyl)Methanone (13b and 13d)

(4-Fluorophenyl)(3'-hydroxy-[1,4'-bipiperidin]-4-yl)methanone (13a)

To a solution of 12a (0.27 g, 0.66 mmol) in ethanol (10 mL), 6N HCl (4 mL, 2.40 mmol) was added. The reaction mixture was heated to reflux for 14 hours. The mixture was cooled and concentrated under vacuum. 1N NaOH solution (20 mL) was added to the residue, the aqueous phase was extracted with CH₂Cl₂ (20 mL×3). The combined organic phases were washed with saturated sodium carbonate (20 mL×3) and brine solution, dried over Na₂SO₄ and concentrated under vacuum. The crude product was purified by silica gel column chromatography using methanol/ethyl acetate/triethylamine (50/50/2, v/v/v) as the mobile phase to afford 13a as a colorless oil (132 mg, 66%). ¹H NMR (CDCl₃): δ 1.37-1.89 (m, 8H), 2.24-2.77 (m, 6H), 2.98-3.01 (m, 1H), 3.12-3.24 (m, 2H), 3.35-3.51 (m, 2H), 7.11-7.17 (t, J=5.7 Hz, 2H), 7.94-7.99 (m, 2H).

(3'-Hydroxy-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (13b)

Procedure B was used to prepare 13b as a colorless oil (173 mg, 76%). ¹H NMR (CDCl₃+CD₃OD): δ 1.56-1.92 (m, 7H), 2.41-3.47 (m, 11H), 3.62-3.70 (m, 1H), 3.87 (s, 3H), 6.99-7.02 (m, 2H), 7.94-7.98 (m, 2H).

(1'-Benzoyl-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (13c)

Procedure B was followed to afford 13c as a colorless oil (0.22 g, 67%). NMR (CDCl₃+CD₃OD): δ 1.43-1.51 (m, 1H), 1.74-1.90 (m, 5H), 2.10-2.12 (m, 1H), 2.29-2.40 (m, 2H), 2.49-2.61 (m, 2H), 2.77-2.88 (m, 2H), 3.03-3.06 (m, 2H), 3.17-3.26 (m, 2H), 3.51-3.59 (m, 1H), 7.14 (t, J=5.4 Hz, 2H), 7.94-7.98 (m, 2H).

(1'-Benzoyl-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (13d)

Procedure B was followed to afford 13d as a colorless oil (0.18 g, 82%). ¹H NMR (CDCl₃+CD₃OD): δ 1.56-1.92 (m, 6H), 2.07-2.11 (m, 1H), 2.41-2.80 (m, 5H), 2.80-3.00 (m, 1H), 3.00-3.18 (m, 2H), 3.20-3.40 (m, 2H), 3.62-3.70 (m, 1H), 3.87 (s, 3H), 6.99-7.02 (m, 2H), 7.94-7.98 (m, 2H).

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Procedure C. General Method of Preparation of Benzyl Compounds (14a-h)

(1'-(4-Fluorobenzyl)-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (14a)

Into a solution of 13a (132 mg, 0.42 mmol) and triethylamine (17.1 mg, 1.69 mmol) in CH₂Cl₂ (15 mL), a solution of 4-fluorobenzyl bromide (80 mg, 0.42 mmol) in CH₂Cl₂ (5 mL) was slowly added. The reaction mixture was stirred at room temperature overnight. The reaction mixture was washed with water (20 mL×2) and brine solution (20 mL). The organic layer was dried over Na₂SO₄ and concentrated under reduced pressure. The crude product was purified by silica gel column chromatograph using triethylamine/ethyl acetate (1/50, v/v) as mobile phase to afford free base 14a as pale yellow oil (78.9 mg, 45%). ¹H NMR (CDCl₃): δ 1.53-2.04 (m, 9H), 2.20-2.32 (m, 2H), 2.71-2.76 (m, 2H), 2.91-3.01 (m, 2H), 3.17-3.22 (m, 2H), 3.51-3.63 (m, 3H), 6.96-7.02 (m, 2H), 7.10-7.17 (m, 2H), 7.23-7.27 (m, 2H), 7.93-7.98 (m, 2H). Free base was converted to the oxalate salt by dissolving it in acetone and mixing with 1 equivalent of oxalic acid in acetone. mp: 234° C. (decomposed). Anal. (C₂₄H₂₈F₂N₂O₂·H₂C₂O₄·0.25H₂O) C, H, N.

(4-Fluorophenyl)(3'-hydroxy-1'-(4-methoxybenzyl)-[1,4'-bipiperidin]-4-yl)methanone (14b)

Procedure C was followed to afford 14b (54 mg, 78%). ¹H NMR (CDCl₃): δ 1.55-1.87 (m, 7H), 2.23-2.27 (m, 2H), 2.70-2.75 (m, 2H), 2.92-3.00 (m, 2H), 3.19-3.24 (m, 2H), 3.48-3.60 (m, 4H), 3.80 (s, 3H), 6.83-6.86 (m, 2H), 7.10-7.21 (m, 4H), 7.93-7.98 (m, 2H). Free base was converted to the oxalate salt. mp: 237° C. (decomposed). Anal. (C₂₅H₃₁FN₂O₃·H₂C₂O₄·0.5H₂O) C, H, N.

(1'-Benzyl-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (14c). Procedure C was followed to afford 14c (54 mg, 84%). ¹H NMR (CDCl₃): δ 1.57-2.03 (m, 8H), 2.17-2.28 (m, 2H), 2.71-2.76 (m, 2H), 2.93-3.01 (m, 2H), 3.20-3.25 (m, 2H), 3.50-3.63 (m, 4H), 7.10-7.16 (m, 2H), 7.24-7.31 (m, 5H), 7.93-7.98 (m, 2H). The free base was converted to the oxalate salt. mp: 243° C. (decomposed). Anal. (C₂₄H₂₉FN₂O₂·H₂C₂O₄·0.25H₂O) C, H, N.

(4-Fluorophenyl)(3'-hydroxy-1'-(pyridin-4-ylmethyl)-[1,4'-bipiperidin]-4-yl)methanone (14d)

Procedure C was followed to yield 14d (19 mg, 50%). The mobile phase used for column chromatographic separation was triethylamine/ethyl acetate/methanol (1/9/1, v/v/v). ¹H NMR (CDCl₃): δ 1.71-2.02 (m, 8H), 2.24-2.28 (m, 2H), 2.74-2.76 (m, 2H), 2.91-3.00 (m, 2H), 3.17-3.21 (m, 2H), 3.50-3.60 (m, 4H), 7.10-7.17 (m, 2H), 7.23-7.27 (m, 1H), 7.62-7.65 (m, 1H), 7.94-7.98 (m, 2H), 8.50-8.52 (m, 2H). The free base was converted to the oxalate salt. mp: 178° C. (decomposed). Anal. (C₂₃H₂₈FN₃O₂·1.5H₂C₂O₄·H₂O) C, H, N. The structure of 14d has two tertiary amines and a pyridine ring and each of them neutralized with one proton. Each oxalic acid can generate two protons. Totally, each equivalent of base needs one and half oxalic acid.

(1'-(4-Fluorobenzyl)-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (14e)

Procedure C was followed to prepare 14e (24.5 mg, 63%). ¹H NMR (CDCl₃): δ 1.56-2.28 (m, 8H), 2.18-2.30 (m, 2H), 2.70-2.74 (m, 2H), 2.92-2.99 (m, 2H), 3.19-3.23 (m, 2H),

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3.48-3.62 (m, 4H), 3.87 (s, 3H), 6.83-6.95 (m, 4H), 7.18-7.21 (m, 2H), 7.90-7.94 (m, 2H). The free base was converted to the oxalate salt. mp: 231° C. (decomposed). Anal. (C₂₅H₃₁FN₂O₃·H₂C₂O₄·0.5H₂O) C, H, N.

(3'-Hydroxy-1'-(4-methoxybenzyl)-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (14f)

Procedure C was followed to prepare 14f (30 mg, 73%). ¹H NMR (CDCl₃): δ 1.51-2.00 (m, 8H), 2.20-2.32 (m, 2H), 2.71-2.76 (m, 2H), 2.91-3.01 (m, 2H), 3.17-3.22 (m, 2H), 3.51-3.63 (m, 4H), 3.79 (s, 3H), 3.86 (s, 3H), 6.96-7.02 (m, 2H), 7.10-7.17 (m, 2H), 7.23-7.27 (m, 2H), 7.93-7.98 (m, 2H). The free base was converted to the oxalate salt. mp: 225° C. (decomposed). Anal. (C₂₅H₃₁FN₂O₃·H₂C₂O₄·0.5H₂O) C, H, N.

(1'-Benzyl-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (14g)

Procedure C was followed to prepare 14g (32 mg, 34%). ¹H NMR (CDCl₃): δ 1.54-2.04 (m, 8H), 2.22-2.27 (m, 2H), 2.70-2.75 (m, 2H), 2.93-3.00 (m, 2H), 3.20-3.25 (m, 2H), 3.54-3.62 (m, 4H), 3.86 (s, 3H), 6.92-6.95 (d, J=9 Hz, 2H), 7.26-7.31 (m, 5H), 7.91-7.94 (d, J=9 Hz, 2H). The free base was converted to the oxalate salt. mp: 234° C. (decomposed). Anal. (C₂₅H₃₂N₂O₃·2H₂C₂O₄) C, H, N.

(3'-Hydroxy-1'-(pyridin-4-ylmethyl)-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (14h)

Procedure C was followed to prepare 14h (20 mg, 52%). The mobile phase used for column chromatographic separation was triethylamine/ethyl acetate/methanol (1/9/1, v/v/v). ¹H NMR (CDCl₃): δ 1.72-2.03 (m, 8H), 2.25-2.29 (m, 2H), 2.74-2.76 (m, 2H), 2.91-3.00 (m, 2H), 3.19-3.21 (m, 2H), 3.55-3.61 (m, 4H), 3.87 (s, 3H), 6.93-6.96 (d, J=9 Hz, 2H), 7.23-7.27 (m, 1H), 7.62-7.65 (m, 1H), 7.91-7.94 (d, J=9 Hz, 2H), 8.50-8.52 (m, 2H). The free base was converted to the oxalate salt. mp: 188° C. (decomposed). Anal. (C₂₄H₃₁N₃O₃·2H₂C₂O₄·0.5H₂O) C, H, N.

Procedure D. General Method of Preparing Benzoic Compounds (15a-g)

(3'-Hydroxy-[1,4'-bipiperidine]-1',4'-diyl)bis(4-fluorophenyl)methanone (15a)

Into a solution of 13a (50 mg, 0.163 mmol), BOP-Cl (100 mg, 0.40 mmol), and triethylamine (1 mL) in methylene chloride (30 mL) was added 4-fluorobenzoic acid (40 mg, 0.285 mmol). The reaction mixture was stirred overnight at room temperature. The reaction mixture was washed with saturated sodium carbonate (20 mL×5) and brine solution (20 mL). The organic layer was dried over Na₂SO₄ and concentrated under reduced pressure. The crude product was purified by silica gel column chromatography using triethylamine/ethyl acetate (1/50, v/v) as mobile phase to afford target product (62 mg, 89%). ¹H NMR (CDCl₃): δ 1.54-1.93 (m, 7H), 2.27-2.34 (m, 1H), 2.44-2.52 (m, 1H), 2.80-2.98 (m, 5H), 3.23-3.26 (m, 1H), 3.30-3.60 (m, 1H), 4.80 (br s, 1H), 7.07-7.18 (m, 4H), 7.40-7.44 (m, 2H), 7.94-7.99 (m, 2H). The free base was converted to the oxalate salt. mp: 213° C. (decomposed). Anal. (C₂₄H₂₆F₂N₂O₅·H₂C₂O₄) C, H, N.

(4-(4-Fluorobenzoyl)-3'-hydroxy-[1,4'-bipiperidin]-1'-yl)(4-methoxyphenyl)methanone (15b)

Procedure D was followed to prepare 15b (48 mg, 22%). ¹H NMR (CDCl₃): δ 1.54-1.93 (m, 7H), 2.31-2.34 (m, 1H),

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2.44-2.51 (m, 1H), 2.79-2.98 (m, 5H), 3.23-3.26 (m, 1H), 3.47 (m, 2H), 3.82 (s, 3H), 6.91 (d, J=9 Hz, 2H), 7.14 (t, J=6 Hz, 2H), 7.38 (d, J=9 Hz, 2H), 7.94-7.99 (m, 2H). The free base was converted to the oxalate salt. mp: 206° C. (decomposed). Anal. (C₂₄H₂₇FN₂O₃·H₂C₂O₄·0.5H₂O) C, H, N.

(1'-Benzoyl-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (15c)

Procedure D was followed to prepare 15c (0.34 g, 78%). ¹H NMR (CDCl₃): δ 1.75-1.80 (m, 2H), 1.80-1.95 (m, 4H), 2.20-2.40 (m, 1H), 2.40-2.60 (m, 1H), 2.60-3.00 (m, 5H), 3.10-3.30 (m, 1H), 4.10 (s br s, 1H), 4.86 (br s, 1H), 7.12-7.18 (m, 2H), 7.28-7.42 (m, 5H), 7.95-7.99 (m, 2H). The free base was converted to the oxalate salt. mp: 152° C. (decomposed). Anal. (C₂₄H₂₇FN₂O₃·H₂C₂O₄·H₂O) C, H, N.

(4-(4-Fluorobenzoyl)-3'-hydroxy-[1,4'-bipiperidin]-1'-yl)(pyridin-4-yl)methanone (15d)

Procedure D was followed to prepare 15d (33 mg, 83%). The mobile phase used for column chromatographic separation was triethylamine/ethyl acetate/methanol (1/9/1, v/v/v). ¹H NMR (CDCl₃): δ 1.44-2.11 (m, 6H), 2.20-2.40 (m, 1H), 2.40-2.60 (m, 1H), 2.60-3.10 (m, 5H), 3.10-3.30 (m, 1H), 3.38-3.85 (m, 2H), 3.90-4.30 (m, 1H), 4.80-5.10 (m, 1H), 7.12-7.18 (m, 2H), 7.34-7.38 (m, 1H), 7.73-7.75 (m, 1H), 7.95-7.99 (m, 2H), 8.66-8.68 (m, 2H). The free base was converted to the oxalate salt. mp: 215° C. (decomposed). Anal. (C₂₃H₂₆FN₃O₃·H₂C₂O₄·0.25H₂O) C, H, N.

(1'-(4-Fluorobenzoyl)-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (15e)

Procedure D was followed to prepare 15e (84 mg, 67%). ¹H NMR (CDCl₃): 1.30-1.98 (m, 5H), 2.16-2.33 (m, 1H), 2.34-2.41 (m, 1H), 2.45-3.65 (m, 8H), 3.77 (s, 3H), 4.03 (br s, 1H), 4.72 (br s, 1H), 6.83-6.87 (m, 2H), 6.97-7.03 (m, 2H), 7.29-7.35 (m, 2H), 7.81-7.85 (m, 2H).

The free base was converted to the oxalate salt. mp: 216° C. (decomposed). Anal. (C₂₅H₂₉FN₂O₄·H₂C₂O₄) C, H, N.

(1'-Benzoyl-3'-hydroxy-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (15f)

Procedure D was used to prepare 15f (0.14 g, 80%). ¹H NMR (CDCl₃): δ 1.60-1.90 (m, 5H), 2.10-2.50 (m, 2H), 2.60-3.00 (m, 5H), 3.20-3.50 (m, 2H), 3.65 (s, 1H), 3.87 (s, 3H), 4.10 (s br, 1H), 5.30 (br s, 1H), 6.94 (d, J=8.7 Hz, 2H), 7.38-7.40 (m, 5H), 7.92 (d, J=9 Hz, 2H). The free base was converted to the oxalate salt. mp: 188° C. (decomposed). Anal. (C₂₅H₃₀N₂O₈·H₂C₂O₄·1.5H₂O) C, H, N.

(3'-Hydroxy-1'-isonicotinoyl-[1,4'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (15g)

Procedure D was followed to prepare 15g (36 mg, 90%). The mobile phase used for column chromatographic separation was triethylamine/ethyl acetate/methanol (1/9/1, v/v/v). ¹H NMR (CDCl₃): δ 1.61-1.96 (m, 6H), 2.04-2.08 (m, 1H), 2.27-2.56 (m, 2H), 2.80-2.09 (m, 4H), 2.09-3.27 (m, 2H), 3.44-3.69 (m, 2H), 3.87 (s, 3H), 4.85-5.15 (m, 1H), 6.93-6.97 (m, 2H), 7.28-7.38 (m, 1H), 7.73-7.80 (m, 1H), 7.85-7.94 (m, 2H), 8.67-8.68 (m, 2H). The free base was converted to the oxalate salt. mp: 220° C. (decomposed). EIMS: Calcd, 429.1843. Found, 429.1843. Anal. (C₂₄H₂₉N₃O₄·H₂C₂O₄) C, H, N.

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(1'-(4-Fluorobenzyl)-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (16a)

Procedure C was followed to yield 16a as a pale yellow oil (28.7 mg, 43%). ¹H NMR (CDCl₃): δ 1.55-2.05 (m, 9H), 2.24-2.31 (m, 1H), 2.49-2.57 (m, 1H), 2.71-2.83 (m, 3H), 2.94-3.04 (m, 2H), 3.17-3.20 (m, 1H), 3.40-3.49 (m, 3H), 6.98-7.05 (m, 2H), 7.10-7.17 (m, 2H), 7.25-7.30 (m, 2H), 7.92-7.97 (m, 2H). The free base was converted to the oxalate salt. mp: 209° C. (decomposed). Anal. (C₂₄H₂₈F₂N₂O₂·H₂C₂O₄) C, H, N.

(4-Fluorophenyl)(4'-hydroxy-1'-(4-methoxybenzyl)-[1,3'-bipiperidin]-4-yl)methanone (16b)

Procedure C was followed to prepare 16b (28 mg, 45%). ¹H NMR (CDCl₃): δ 1.56-2.06 (m, 8H), 2.26-2.30 (m, 1H), 2.53-2.56 (m, 1H), 2.72-2.86 (m, 3H), 3.00-3.03 (m, 2H), 3.17-3.20 (m, 1H), 3.40-3.50 (m, 4H), 3.81 (s, 3H), 6.86-6.89 (m, 2H), 7.10-7.27 (m, 4H), 7.92-7.97 (m, 2H). The free base was converted to the oxalate salt. mp: 204° C. (decomposed). Anal. (C₂₅H₃₁FN₂O₃·H₂C₂O₄) C, H, N.

(1'-Benzyl-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-fluorophenyl)methanone (16c)

Procedure C was followed to prepare 16c (40 mg, 62%). ¹H NMR (CDCl₃): δ 1.56-2.06 (m, 8H), 2.24-2.3 (m, 1H), 2.51-2.59 (m, 1H), 2.72-2.86 (m, 3H), 3.00-3.04 (m, 2H), 3.17-3.20 (m, 1H), 3.40-3.54 (m, 4H), 7.10-7.15 (m, 2H), 7.26-7.33 (m, 5H), 7.92-7.97 (m, 2H). The free base was converted to the oxalate salt. mp: 200° C. (decomposed). Anal. (C₂₄H₂₉FN₂O₂·H₂C₂O₄) C, H, N.

(4-Fluorophenyl)(4'-hydroxy-1'-(pyridin-4-ylmethyl)-[1,3'-bipiperidin]-4-yl)methanone (16d)

Procedure C was followed to prepare 16d (22 mg, 56%). ¹H NMR (CDCl₃): δ 1.56-2.08 (m, 8H), 2.24-2.32 (m, 1H), 2.48-2.53 (m, 1H), 2.70-3.04 (m, 5H), 3.19-3.59 (m, 5H), 7.13 (t, J=5.7 Hz, 2H), 7.26-7.31 (m, 1H), 7.66-7.69 (m, 1H), 7.92-7.97 (m, 2H), 8.52-8.54 (m, 2H). The free base was converted to the oxalate salt. mp: 182° C. Anal. (C₂₃H₂₈FN₃O₂·H₂C₂O₄·0.5H₂O) C, H, N.

(1'-(4-Fluorobenzyl)-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (16e)

Procedure C was followed to prepare 16e (38 mg, 55%). ¹H NMR (CDCl₃): δ 1.55-2.06 (m, 8H), 2.28 (td, J=13.8, 2.1 Hz, 1H), 2.52 (td, J=10.5, 3.3 Hz, 1H), 2.71-2.84 (m, 3H), 2.90-3.10 (m, 2H), 3.04-3.20 (m, 1H), 3.40-3.49 (m, 4H), 3.86 (s, 3H), 6.92-7.04 (m, 4H), 7.25-7.30 (m, 2H), 7.89-7.92 (m, 2H). The free base was converted to the oxalate salt. mp: 221° C. (decomposed). Anal. (C₂₅H₃₁FN₂O₃·H₂C₂O₄) C, H, N.

(4'-Hydroxy-1'-(4-methoxybenzyl)-[1,3'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (16f)

Procedure C was followed to prepare 16f (32 mg, 78%). ¹H NMR (CDCl₃): δ 1.54-2.06 (m, 8H), 2.24-2.28 (m, 1H), 2.50-2.53 (m, 1H), 2.72-2.86 (m, 3H), 2.98-3.02 (m, 2H), 3.17-3.20 (m, 1H), 3.41-3.48 (m, 3H), 3.62 (br s, 1H), 3.81 (s, 3H), 3.86 (s, 3H), 6.85-6.95 (m, 4H), 7.22 (dd, J=2.1, 6.9 Hz, 2H), 7.90 (dd, J=2.1, 6.9 Hz, 2H). The free base was con-

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verted to the oxalate salt. mp: 182° C. (decomposed). Anal. (C₂₆H₃₄N₂O₄·H₂C₂O₄) C, H, N

(1'-Benzyl-4'-hydroxy-[1,3'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (16g)

Procedure C was followed to prepare 16g (90 mg, 71%). ¹H NMR (CDCl₃): δ 1.57-2.04 (m, 8H), 2.25-2.32 (t, J=7 Hz, 1H), 2.51-2.60 (m, 1H), 2.70-2.86 (m, 3H), 3.00-3.04 (m, 2H), 3.17-3.20 (m, 1H), 3.40-3.59 (m, 4H), 3.86 (s, 3H), 6.91-6.94 (d, J=9 Hz, 2H), 7.26-7.33 (m, 5H), 7.89-7.92 (d, J=9 Hz, 2H). The free base was converted to the oxalate salt. mp: 220.3° C. (decomposed). Anal. (C₂₅H₃₂N₂O₃·H₂C₂O₄·0.5H₂O) C, H, N.

(4'-Hydroxy-1'-(pyridin-4-ylmethyl)-[1,3'-bipiperidin]-4-yl)(4-methoxyphenyl)methanone (16h)

Procedure C was followed to prepare 16h (13 mg, 34%). ¹H NMR (CDCl₃): δ 1.58-2.08 (m, 8H), 2.22-2.29 (m, 1H), 2.52-2.53 (m, 1H), 2.70-3.04 (m, 5H), 3.18-3.20 (m, 1H), 3.42-3.54 (m, 4H), 3.87 (s, 3H), 6.91-6.96 (m, 2H), 7.26-7.30 (m, 1H), 7.63-7.70 (m, 1H), 7.89-7.94 (m, 2H), 8.52-8.54 (m, 2H). The free base was converted to the oxalate salt. mp: 202° C. Anal. (C₂₄H₃₁N₃O₃·H₂C₂O₄) C, H, N.

Procedure E. Method of preparing 1a,2,7,7a-tetrahydronaphtho[2,3-b]oxirene 2

To a solution of 1,4-dihydronaphthalene (8.6 g, 86% purity, 56.8 mmol) in dichloromethane (100 mL) at 0° C., m-chloroperoxybenzoic acid (17.0 g, 77% Purity, 75.8 mmol) in dichloromethane was added. After stirring for 5 h at 0° C., the solution was filtered to separate out the precipitated m-chlorobenzoic acid. The organic layer was washed with saturated Na₂CO₃, water, and brine (2×50 mL each). The organic layer was dried and concentrated under reduced pressure, the residue was purified by chromatography to give compound 2 (6.1 g, 73%). ¹H NMR (CDCl₃): δ 3.17 (d, J=17.4 Hz, 2H), 3.31 (d, J=16.8 Hz, 2H), 3.46 (s, 2H), 7.02-7.15 (m, 4H).

5,8-Dihydronaphthalen-1-amine hydrochloride 4

A solution of 3 (10 g, 69.8 mmol) in oxylene (100 mL) and ethanol (30 mL) was stirred, and small pieces sodium (7.5 g, 326.0 mmol) was added during 1.5 h. The mixture was heated at 80° C. for 2 h, and poured onto ice. The organic layer was separated, washed with water. Hydrochloric acid (1:3, 50 mL) was added, the solid was collected by filtration, dried to give the crude product (7.0 g, 55%). The crude product was used in next step without further purification. ¹H NMR (CDCl₃—CD₃OD): δ 3.45 (m, 7H), 5.90 (br, 2H), 7.12-7.35 (m, 3H).

N-(5,8-dihydronaphthalen-1-yl)-2,2,2-trifluoroacetamide 5

To a solution of 4 (7.0 g, 38.5 mmol) and NEt₃ (15 mL) in dichloromethane (120 mL) at 0° C., trifluoroacetic anhydride (13.0 g, 61.9 mmol) was added slowly. The reaction mixture was allowed to warm to room temperature and stirred for 2 h until complete by TLC, and then the mixture was washed by water, aqueous Na₂CO₃ solution, dried over Na₂SO₄, and concentrated under reduced pressure. The residue was purified by column chromatography to give 5 (8.0 g, 86%). ¹H NMR (CDCl₃): δ 3.23 (m, 2H), 3.44 (m, 2H), 5.91 (m, 2H), 7.08 (d, J=7.8 Hz, 1H), 7.22 (t, J=8.1 Hz, 1H), 7.59 (d, J=7.8 Hz, 1H), 8.67 (br, 1H).

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2,2,2-Trifluoro-N-(1a,2,7,7a-tetrahydronaphtho[2,3-b]oxiren-3-yl)acetamide 6

Compound 6 was prepared from 5 as describe in procedure A. Yield, 53%. ¹H NMR (CDCl₃): δ 2.80-2.97 (m, 2H), 3.21-3.41 (m, 2H), 3.51-3.57 (m, 2H), 7.03-7.23 (m, 3H), 8.25 (br, 1H).

tert-Butyl 4-(methoxy(methyl)carbamoyl)piperidine-1-carboxylate 8

To a solution of 7 (2.5 g, 10.9 mmol) in dichloromethane (30 mL) at room temperature, carbonyldiimidazole (1.77 g, 11.0 mmol) was added. The mixture was stirred at room temperature for 1 h and then N, O-dimethylhydroxyamine hydrochloride (1.3 g, 13.3 mmol) and triethylamine were added. The reaction mixture was stirring overnight and was washed with aqueous Na₂CO₃, water, acetic acid (5%), dried, and concentrated under reduced pressure. The residue was purified by column chromatography to give 8 (2.70 g, 90%). ¹H NMR (CDCl₃): δ 1.46 (s, 9H), 1.64-1.72 (m, 4H), 2.78 (br, 3H), 3.18 (s, 3H), 3.71 (s, 3H), 4.13 (br, 2H).

Procedure F: General Method of Preparing 9a-g

Method 1. tert-butyl

4-(6-methylnicotinoyl)piperidine-1-carboxylate 9a

To a solution of 5-bromo-2-methylpyridine (1.25 g, 7.2 mmol) in anhydrous THF (15 mL) at -40° C. under an argon atmosphere, lithium dibutyl(isopropyl)magnesate (5.2 mL, 0.7 M, 3.6 mmol) was added. The stirring was continued at the same temperature for 1 h. The resulting mixture was cannulated into a solution of 8 (1.5 g, 5.5 mmol) in THF (20 mL) at -78° C. The solution was maintained at -78° C. for 1 h, and then quenched with sat. aqueous NH₄Cl and allowed to warm to room temperature. The organic layer was separated and then the aqueous layer was extracted with ethyl acetate (2×20 mL). The combined organic layer was washed with water, dried over Na₂SO₄, filtered, and evaporated under reduced pressure to give the crude product. The crude compound was purified by chromatography on silica gel to obtain the desired compound 9a (0.7 g, 42%). ¹H NMR (CDCl₃): δ 1.49 (s, 9H), 1.68-1.87 (m, 4H), 2.63 (s, 3H), 3.35 (m, 1H), 4.18 (br, 2H), 7.28 (d, J=8.4 Hz, 1H), 8.11 (dd, J=8.4, 2.1 Hz, 1H), 9.04 (d, J=2.1 Hz, 1H).

Method 2. tert-butyl

4-(6-methoxynicotinoyl)piperidine-1-carboxylate 9b

To a solution of 5-bromo-2-methoxy pyridine (2.6 g, 13.8 mmol) in anhydrous THF at -78° C., n-butyllithium (1.6M, 10 mL) was added. The solution was stirred for 45 minutes at -78° C. The resulting mixture was cannulated into a solution of 8 (2.5 g, 9.2 mmol) in THF (20 mL) at the same temperature and the reaction mixture was stirred at -78° C. for 4 h, then quenched with sat. aqueous NH₄Cl and allowed to warm to room temperature. The mixture was washed with brine and dried over Na₂SO₄, filtered, and evaporated under reduced pressure to give the crude product. The crude compound was purified by chromatography on silica gel to obtain the desired compound 9b (1.8 g, 61%). ¹H NMR (CDCl₃): δ 1.47 (s, 9H), 1.62-1.86 (m, 5H), 2.88 (m, 2H), 3.31 (m, 1H), 4.01 (s, 3H), 4.19 (br, 2H), 6.81 (d, J=8.7 Hz, 1H), 8.13 (dd, J=8.7, 2.4 Hz, 1H), 8.79 (d, J=2.4 Hz, 1H).

tert-Butyl 4-nicotinoylpiperidine-1-carboxylate 9c

Compound 9c was prepared from 3-bromopyridine as describe in procedure B (Method 1). Yield, 31%. ¹H NMR

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(CDCl₃): δ 1.47 (s, 9H), 1.63-1.89 (m, 4H), 2.92 (m, 2H), 3.38 (m, 2H), 4.20 (m, 2H), 7.44 (ddd, J=8.1, 5.1, 0.9 Hz, 1H), 8.22 (dt, J=8.1, 2.1 Hz, 1H), 8.79 (dd, J=5.1, 1.8 Hz, 1H), 9.16 (d, J=1.8 Hz, 1H).

tert-Butyl 4-(1-methyl-1H-pyrrole-2-carbonyl)piperidine-1-carboxylate 9d

Compound 9d was prepared from 1-methyl-1H-pyrrole as describe in procedure B (Method 2). Yield, 34%. ¹H NMR (CDCl₃): δ 1.47 (s, 9H), 1.66-1.80 (m, 4H), 2.83 (s, 2H), 3.17 (m, 1H), 3.94 (s, 3H), 4.18 (br, 2H), 6.15 (dd, J=4.2, 2.4 Hz, 1H), 6.84 (t, J=1.8 Hz, 1H), 6.99 (dd, J=4.2, 1.5 Hz, 1H).

tert-Butyl 4-(1-methyl-1H-pyrrole-4-carbonyl)piperidine-1-carboxylate 9e

Compound 9e was prepared from 3-bromo-1-methyl-1H-pyrrole as describe in procedure B (Method 2). Yield, 40%. ¹H NMR (CDCl₃): δ 1.47 (s, 9H), 1.66-1.77 (m, 4H), 2.83 (s, 2H), 3.15-3.18 (m, 1H), 3.70 (s, 3H), 4.13 (br, 2H), 6.57-6.61 (m, 2H), 7.26-7.29 (m, 1H).

tert-Butyl

4-(3-methoxypicolinoyl)piperidine-1-carboxylate 9f

Compound 9f was prepared from 2-bromo-3-methoxy pyridine as describe in procedure B (Method 2). Yield, 36%. ¹H NMR (CDCl₃): δ 1.47 (s, 9H), 1.66-1.87 (m, 4H), 2.89 (s, 2H), 3.67-3.81 (m, 1H) 3.90 (s, 3H), 4.07 (br, 2H), 7.35-7.45 (m, 2H), 8.23-8.25 (m, 1H).

tert-Butyl

4-(3-fluoronicotinoyl)piperidine-1-carboxylate 9g

Compound 9g was prepared from 3-bromo-5-fluoropyridine as describe in procedure B (Method 2). Yield, 41%. ¹H NMR (CDCl₃): δ 1.44 (s, 9H), 1.56-1.61 (m, 2H), 1.86-1.90 (m, 2H), 2.87 (s, 2H), 3.17-3.24 (m, 1H), 4.06 (br, 2H), 7.54-7.58 (m, H), 8.53-8.60 (m, 2H).

Procedure G: General Method of Preparing 10a-g.

(6-methylpyridin-3-yl)(piperidin-4-yl) methanone 10a

To a solution of 9a (0.7 g, 2.3 mmol) in CH₂Cl₂ (15 mL), TFA (2 mL) was added. The reaction mixture was stirred at room temperature for 4 h. The solvent was removed under reduced pressure and the residue was neutralized with aqueous NaOH solution and extracted with CH₂Cl₂ (2×15 mL). The organic layer was washed with brine, dried and concentrated to give the crude product. The crude product was purified with column chromatography to afforded 10a (0.33 g, 70%). ¹H NMR (CDCl₃): δ 1.62-1.87 (m, 6H), 2.63 (s, 3H), 2.77 (m, 2H), 3.16-3.22 (m, 2H), 3.33 (m, 1H), 7.27 (d, J=8.4 Hz, 1H), 8.12 (dd, J=8.1, 2.4 Hz, 1H), 9.04 (d, J=2.4 Hz, 1H).

(6-Methoxypyridin-3-yl)(piperidin-4-yl)methanone 10b

Compound 10b was prepared from compound 9b as describe in procedure C. Yield, 77%. ¹H NMR (CDCl₃): δ 1.65-1.89 (m, 4H), 2.40 (s, 1H), 2.78 (td, J=12.0, 2.7 Hz, 2H), 3.21 (dt, J=9.0, 3.6 Hz, 2H), 3.31 (m, 1H), 4.00 (s, 3H), 6.79 (d, J=8.4 Hz, 1H), 8.13 (dd, J=8.4, 2.4 Hz, 1H), 8.78 (d, J=2.4 Hz, 1H).

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Piperidin-4-yl(pyridin-3-yl)methanone 10c

Compound 10c was prepared from compound 9c as describe in procedure C. Yield, 85%. ¹H NMR (CDCl₃): δ 1.62-1.77 (m, 2H), 1.86-1.89 (m, 3H), 2.79 (td, J=12.6, 2.7 Hz, 2H), 3.21 (dt, J=12.9, 3.0 Hz, 2H), 3.37 (tt, J=11.4, 3.9 Hz, 1H), 7.43 (ddd, J=8.1, 4.8, 0.9 Hz, 1H), 8.22 (ddd, J=4.8, 2.1, 1.5 Hz, 1H), 8.78 (dd, J=4.5, 1.5 Hz, 1H), 9.16 (dd, J=2.4, 0.9 Hz, 1H).

(1-Methyl-1H-pyrrol-2-yl)(piperidin-4-yl)methanone 10d

Compound 10d was prepared from compound 9d as describe in procedure C. Yield, 72%. ¹H NMR (CDCl₃): δ 1.76-1.87 (m, 4H), 2.81 (m, 2H), 3.17-3.23 (m, 4H), 3.94 (s, 3H), 6.14 (m, 1H), 6.83 (m, 1H), 6.98 (m, 1H).

(1-Methyl-1H-pyrrol-3-yl)(piperidin-4-yl)methanone 10e

Compound 10e was prepared from compound 9e as describe in procedure C. Yield, 80%. ¹H NMR (CDCl₃): δ 1.76-1.80 (m, 4H), 2.65-2.73 (m, 2H), 3.01-3.17 (m, 4H), 3.68 (s, 3H), 6.57-6.59 (m, 2H), 7.25 (s, 1H).

(3-Methoxypyridin-2-yl)(piperidin-4-yl)methanone 10f

Compound 10f was prepared from compound 9f as describe in procedure C. Yield, 78%. ¹H NMR (CDCl₃): δ 1.51-1.81 (m, 4H), 2.69 (s, 2H), 3.08-3.25 (m, 3H), 3.49-3.58 (m, 1H), 3.83 (s, 3H), 7.27-7.36 (m, 2H), 8.16-8.18 (m, 1H).

(5-Fluoropyridin-3-yl)(piperidin-4-yl)methanone 10g

Compound 10g was prepared from compound 9g as describe in procedure C. Yield, 79%. ¹H NMR (CDCl₃): δ 1.56-1.61 (m, 2H); 1.86-1.90 (m, 2H); 2.87 (m, 2H); 3.17-3.21 (m, 4H); 7.56 (s, 1H); 8.53-8.60 (m, 2H).

Procedure H: General Method of Preparing 11a-b, d-g

(OS-5-40, OS-5-88, OS-5-60, Tz-4-1-114, Tz-4-1-92 and TZ-4-1-96) (1-(3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(6-methoxypyridin-3-yl)methanone oxalate 11a OS-5-40

A mixture of 10a (0.32 g, 1.4 mmol), 2 (0.10 g, 0.68 mmol) and EtN₃ (0.3 mL) in ethanol (5 mL) was stirred at 50° C. for 36 h. After cooled to room temperature, the mixture was poured into water, extracted with EtOAc (3×15 mL). The organic layer was dried and concentrated under reduced pressure. The residue was purified by column chromatography to give 11a OS-5-40 (0.11 g, 44%). ¹H NMR (CDCl₃, free base): δ 1.80-2.02 (m, 4H), 2.43 (m, 1H), 2.78-3.06 (m, 7H), 3.20-3.38 (m, 2H), 3.89 (m, 1H), 4.04 (s, 3H), 6.83 (d, J=8.7 Hz, 1H), 7.11-7.18 (m, 4H), 8.17 (dd, J=8.7, 2.4 Hz, 1H), 8.82 (d, J=2.4 Hz, 1H). The free base was converted to the oxalate salt. mp: 223-224° C. Anal. (C₂₂H₂₆N₂O₃·H₂C₂O₄) C, H, N.

(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(6-methylpyridin-3-yl)methanone oxalate 11b OS-5-88

Compound 11b OS-5-88 was prepared from compound 10b as describe in procedure D. Yield, 42%. ¹H NMR

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(CDCl₃, free base): δ 1.74-1.96 (m, 4H), 2.41 (m, 1H), 2.62 (s, 3H), 2.75-3.01 (m, 7H), 3.26-3.33 (m, 2H), 3.86 (m, 1H), 4.17 (br, 1H), 7.06-7.14 (m, 4H), 7.27 (d, J=8.1 Hz, 1H), 8.11 (dd, J=8.1, 2.1 Hz, 1H), 9.03 (d, J=2.1 Hz, 1H). The free base was converted to the oxalate salt. mp: 240-241° C. Anal. (C₂₂H₂₆N₂O₂·2H₂C₂O₄·0.5H₂O) C, H, N.

(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(1-methyl-1H-pyrrol-2-yl)methanone oxalate 11d OS-5-60

Compound 11 d OS-5-60 was prepared from compound 10d as describe in procedure D. Yield, 47%. ¹H NMR (CDCl₃, free base): δ 1.81-2.03 (m, 4H), 2.37 (m, 1H), 2.76-3.13 (m, 8H), 3.30 (m, 1H), 3.88 (m, 1H), 3.95 (s, 3H), 4.45 (br, 1H), 6.14 (m, 1H), 6.83 (m, 1H), 6.98 (m, 1H), 7.09-7.16 (m, 4H). The free base was converted to the oxalate salt. mp: 212-213° C. Anal. (C₂₁H₂₆N₂O₂·H₂C₂O₄) C, H, N.

(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(1-methyl-1H-pyrrol-3-yl)methanone oxalate 11e Tz-4-1-114

Compound 11e Tz-4-1-114 was prepared from compound 10e as describe in procedure D. Yield, 50%. ¹H NMR (CDCl₃, free base): δ 1.83-2.00 (m, 4H), 2.38 (m, 1H), 2.76-3.13 (m, 8H), 3.29-3.35 (m, 1H), 3.71 (s, 3H), 4.32 (br, 1H), 6.61 (m, 2H), 7.12-7.16 (m, 4H), 7.30-7.31 (m, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₁H₂₆N₂O₂·H₂C₂O₄) C, H, N.

(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(3-methoxypyridin-2-yl)methanone oxalate 11f Tz-4-1-92

Compound 11f Tz-4-1-92 was prepared from compound 10f as describe in procedure D. Yield, 36%. ¹H NMR (CDCl₃, free base): δ 1.72-1.96 (m, 4H), 2.33-2.41 (m, 1H), 2.72-2.96 (m, 7H), 3.26-3.33 (m, 1H), 3.55-3.63 (m, 1H), 3.81-3.90 (m, 1H), 4.30 (br, 1H), 7.06-7.14 (m, 4H), 7.26-7.39 (m, 2H), 8.23-8.25 (m, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₂H₂₆N₂O₃·H₂C₂O₄) C, H, N.

(5-Fluoropyridin-3-yl)(1-(3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)methanone oxalate 11g TZ-4-1-96

Compound 11g TZ-4-1-96 was prepared from compound 10g as describe in procedure D. Yield, 44%. ¹H NMR (CDCl₃, free base): δ 1.70-2.04 (m, 4H), 2.34-2.43 (m, 1H), 2.75-3.34 (m, 8H), 3.82-3.90 (m, 1H), 4.13 (br, 1H), 7.06-7.11 (m, 4H), 7.57-7.61 (m, 1H), 8.56-8.62 (m, 2H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₁H₂₃FN₂O₂·H₂C₂O₄) C, H, N.

Procedure I: General Method of Preparing 12a-d

(1-(8-amino-3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(6-methylpyridin-3-yl)methanone oxalate 12a (OS-5-70)

A mixture of 6 (0.20 g, 0.78 mmol), 10a (0.47 g, 2.3 mmol) and EtN₃ (0.5 mL) in ethanol (7 mL) was stirred at 60° C. for 2 d. To the mixture aqueous NaOH (1M, 3 mL) was added and the stiffing was continued overnight. The solvent was removed, the residue was extracted with EtOAc (3×10 mL) and organic layer was washed with aqueous Na₂CO₃ solution,

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dried and concentrated. The crude product was purified by column chromatography to give 12a (OS-5-70). (0.13 g, 44%). ¹H NMR (CDCl₃, free base): δ 1.79-1.99 (m, 5H), 2.35-2.53 (m, 2H), 2.64 (s, 3H), 2.67-3.28 (m, 8H), 3.62 (br, 2H), 3.87 (m, 1H), 6.57 (m, 2H), 6.99 (m, 1H), 7.29 (d, J=8.1 Hz, 1H), 8.13 (dd, J=8.1, 2.1 Hz, 1H), 9.04 (d, J=2.1 Hz, 1H). The oxalate salt was prepared using oxalic acid in ethyl acetate, mp: 98-103° C. Anal. (C₂₂H₂₇N₃O₂·2H₂C₂O₄·H₂O) C, H, N.

(1-(8-Amino-3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(6-methoxypyridin-3-yl) methanone oxalate 12b (OS-5-82)

Compound 12b (OS-5-82) was prepared from compound 10b as describe in procedure E. Yield, 43%. ¹H NMR (CDCl₃, free base): δ 1.65 (br, 1H), 1.80-2.02 (m, 4H), 2.42-2.53 (m, 2H), 2.67-3.03 (m, 6H), 3.21-3.28 (m, 2H), 3.61 (br s, 2H), 3.86 (m, 1H), 4.02 (s, 3H), 6.57 (m, 2H), 6.82 (d, J=8.7 Hz, 1H), 6.99 (t, J=7.8 Hz, 1H), 8.15 (dd, J=8.7, 2.4 Hz, 1H), 8.80 (d, J=2.4 Hz, 1H). The free base was converted to the oxalate salt. mp: 81.4-83.2° C. Anal. (C₂₂H₂₇N₃O₃·2H₂C₂O₄·1.5H₂O) C, H, N.

(1-(8-Amino-3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(pyridin-3-yl)methanone oxalate 12c (OS-5-110)

Compound 12c (OS-5-110) was prepared from compound 10c as describe in procedure E. Yield, 61%. ¹H NMR (CDCl₃, free base): δ 1.80-2.05 (m, 4H), 2.36-2.53 (m, 2H), 2.68-2.92 (m, 5H), 3.01-3.35 (m, 4H), 3.61 (br, 2H), 3.87 (m, 1H), 6.54-6.60 (m, 2H), 6.99 (td, J=7.5, 3.3 Hz, 1H), 7.45 (dd, J=8.1, 4.8 Hz, 1H), 8.24 (dt, J=8.1, 1.8 Hz, 1H), 8.80 (dd, J=4.8, 1.8 Hz, 1H), 9.17 (d, J=2.1 Hz, 1H). The free base was converted to the oxalate salt. mp: 109.0-111.2° C. Anal. (C₂₁H₂₅N₃O₂·2H₂C₂O₄·0.5H₂O) C, H, N.

(1-(8-Amino-3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(1-methyl-1H-pyrrol-2-yl) methanone oxalate 12d (OS-5-72-2)

Compound 12d (OS-5-72-2) was prepared from compound 10d as describe in procedure E. Yield, 37%. ¹H NMR (CDCl₃, free base): δ 1.85-1.96 (m, 4H), 2.43-2.53 (m, 2H), 2.67-2.89 (m, 5H), 2.96-3.28 (m, 3H), 3.60 (br, 2H), 3.85 (m, 1H), 3.95 (s, 3H), 6.14 (m, 1H), 6.57 (m, 2H), 6.83 (br, 1H), 6.96-7.02 (m, 2H). The free base was converted to the oxalate salt. mp: 131-133° C. Anal. (C₂₁H₂₇N₃O₂·2H₂C₂O₄·1.5H₂O) C, H, N.

(1-(5-Amino-3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(1-methyl-1H-pyrrol-2-yl) methanone 13d (OS-5-2-72-1)

Compound 13d (OS-5-2-72-1) was prepared from compound 10d as describe in procedure E. Yield, 20%. ¹H NMR (CDCl₃, free base): δ 1.85-1.96 (m, 4H), 2.36-2.45 (m, 2H), 2.76-2.89 (m, 5H), 2.96-3.16 (m, 3H), 3.62 (br, 2H), 3.92 (m, 1H), 3.96 (s, 3H), 6.15 (m, 1H), 6.56 (m, 2H), 6.83 (br, 1H), 6.96-7.02 (m, 2H). The free base was converted to the oxalate salt. mp: 114-115° C. Anal. (C₂₁H₂₇N₃O₂·2H₂C₂O₄·2H₂O) C, H, N.

Procedure J: General Method for the Synthesis of Compound 17a-b

4-bromo-1-methylpyridin-2(1H)-one (17a)

A solution of 4-bromopyridin-2-amine (600 mg, 3.5 mmol) in a mixture of 2M H₂SO₄ (20 mL) and 2M Na₂NO₂

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(10 mL) was stirred at 0-5° C. for 2 h. The reaction mixture was extracted with CH₂Cl₂, and the organic layer was washed with brine, dried over anhydrous Na₂SO₄ and concentrated. The crude product was used in next step without further purification. The crude product 15a, potassium carbonate (3.6 mmol) and methyl iodide (3.7 mmol) were refluxed in acetone (100 mL) in a sealed tube for 4 h. The reaction mixture was cooled, and potassium carbonate was filtered off. The acetone was evaporated off and a small amount of water was added to the residue. This solution was extracted with CH₂Cl₂. The organic layer was dried, concentrated and purified with column chromatography to afforded 17a (355 mg, 57%). ¹H NMR (CDCl₃): δ 3.49 (s, 3H), 6.31 (d, J=6.0 Hz, 1H), 6.82 (s, 1H), 7.14 (d, J=6.0 Hz, 1H).

6-Bromo-1-methylpyridin-2(1H)-one 17b

Compound 17b was prepared from compound 14b as describe in procedure F. Yield, 54%. ¹H NMR (CDCl₃): δ 3.73 (s, 3H), 6.46-6.52 (m, 2H), 7.10-7.16 (m, 1H).

tert-Butyl 4-(1-methyl-2-oxo-1,2-dihydropyridine-4-carbonyl)piperidine-1-carboxylate 18a

Compound 18a was prepared from compound 17a as describe in procedure B (Method 2). Yield, 38%. ¹H NMR (CDCl₃): δ 1.47 (s, 9H), 1.56-1.87 (m, 4H), 2.82 (s, 2H), 3.05-3.07 (m, 1H), 3.50 (s, 3H), 4.17 (br, 2H), 6.49 (d, J=6.0 Hz, 1H), 6.73 (d, J=6.0 Hz, 1H), 7.36 (s, 1H).

tert-Butyl 4-(1-methyl-6-oxo-1,6-dihydropyridine-2-carbonyl)piperidine-1-carboxylate 18b

Compound 18b was prepared from compound 17b as describe in procedure B (Method 2). Yield, 35%. ¹H NMR (CDCl₃): δ 1.45 (s, 9H), 1.58-1.87 (m, 4H), 2.81 (s, 2H), 3.05-3.07 (m, 1H), 3.47 (s, 3H), 4.14 (br, 2H), 6.44-6.47 (m, 1H), 6.69-6.72 (m, 1H), 7.30-7.36 (m, 1H).

1-Methyl-5-(piperidine-4-carbonyl)pyridin-2(1H)-one (19a)

Compound 19a was prepared from compound 18a as describe in procedure C. Yield, 79%. ¹H NMR (CDCl₃): δ 1.57-1.88 (m, 4H), 2.83 (s, 2H), 3.03-3.09 (m, 4H), 3.50 (s, 3H), 6.50 (d, J=6.0 Hz, 1H), 6.72 (d, J=6.0 Hz, 1H), 7.38 (s, 1H).

1-Methyl-6-(piperidine-4-carbonyl)pyridin-2(1H)-one (19b)

Compound 19b was prepared from compound 18b as describe in procedure C. Yield, 75%. ¹H NMR (CDCl₃): δ 1.58-1.87 (m, 4H), 2.82 (s, 2H), 3.02-3.10 (m, 4H), 3.47 (s, 3H), 6.42-6.48 (m, 1H), 6.67-6.72 (m, 1H), 7.31-7.36 (m, 1H).

4-(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidine-4-carbonyl)-1-methylpyridin-2(1H)-one oxalate (20a TZ-4-1-150)

Compound 20a TZ-4-1-150 was prepared from compound 19a as describe in procedure E. Yield, 37%. ¹H NMR (CDCl₃, free base): δ 1.72-1.99 (m, 4H), 2.33-2.39 (m, 1H), 2.75-3.01 (m, 8H), 3.27-3.34 (m, 2H), 3.50 (s, 3H), 3.82-3.91 (m, 1H),

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6.45 (d, J=6.0 Hz, 1H), 6.72 (d, J=6.0 Hz, 1H), 7.35 (s, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₂H₂₆N₂O₃) C, H, N.

6-(1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidine-4-carbonyl)-1-methylpyridin-2(1H)-one
(20b TZ-4-1-148)

Compound 20b TZ-4-1-148 was prepared from compound 19b as describe in procedure E. Yield, 39%. ¹H NMR (CDCl₃, free base): δ 1.72-2.04 (m, 4H), 2.32-2.39 (m, 1H), 2.81-3.01 (m, 8H), 3.27-3.34 (m, 2H), 3.50 (s, 3H), 3.82-3.89 (m, 1H), 4.08 (br, 1H), 6.43-6.46 (m, 1H), 6.70-6.73 (m, 1H), 7.06-7.14 (m, 4H), 7.31-7.37 (m, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₂H₂₆N₂O₃) C, H, N.

1-(3-Hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)-N-methoxy-N-methylpiperidine-4-carboxamide 22

Compound 21 was prepared from compound 8 as describe in procedure C. The crude product was used in next step without further purification. Compound 22 was prepared from compound 21 as describe in procedure D. Yield, 40%. ¹H NMR (CDCl₃): δ 1.81-1.93 (m, 4H), 2.28-2.31 (m, 1H), 2.75-2.97 (m, 8H), 3.27-3.20 (s, 3H), 3.27-3.34 (m, 1H), 3.72 (s, 3H), 3.85-3.90 (m, 1H), 4.27 (br, 1H), 7.09-7.13 (m, 4H).

1-(3-((tert-Butyldimethylsilyloxy)-1,2,3,4-tetrahydronaphthalen-2-yl)-N-methoxy-N-methylpiperidine-4-carboxamide 23

To a solution of 22 (400 mg, 1.25 mmol) and imidazole (400 mg) in CH₂Cl₂ (50 mL), TBDMSCl (2.5 mmol) was added. The reaction mixture was stirred overnight at room temperature until TLC indicated that the reaction was complete. The reaction mixture was washed by brine and 1M NaH₂PO₄ (30 mL×3). The organic layer was dried and concentrated to a residue. The crude product was purified with column chromatography to afforded 23 (400 mg, 70%). ¹H NMR (CDCl₃): δ 0.12 (s, 6H), 0.91 (s, 9H), 1.65-1.77 (m, 4H), 2.46-3.05 (m, 9H), 3.17 (s, 3H), 3.69 (s, 3H), 4.13 (br, 2H), 7.04-7.12 (m, 4H).

(1-(3-(tert-Butyldimethylsilyloxy)-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(6-fluoropyridin-3-yl) methanone 24a

Compound 24a was prepared from compound 23 as describe in procedure E. Yield, 36%. ¹H NMR (CDCl₃): δ 0.12 (s, 6H), 0.91 (s, 9H), 1.65-1.77 (m, 4H), 2.46-3.05 (m, 9H), 3.17 (s, 3H), 3.69 (s, 3H), 4.13 (br, 2H), 7.04-7.12 (m, 4H), 8.23-8.37 (m, 2H), 8.78 (s, 1H).

(1-(3-(tert-Butyldimethylsilyloxy)-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)(2-fluoropyridin-3-yl) methanone 24b

Compound 24b was prepared from compound 23 as describe in procedure E. Yield, 39%. ¹H NMR (CDCl₃): δ 0.12 (s, 6H), 0.91 (s, 9H), 1.65-1.77 (m, 4H), 2.46-3.05 (m, 9H), 3.17 (s, 3H), 3.69 (s, 3H), 4.13 (br, 2H), 7.04-7.12 (m, 4H), 7.04-7.11 (m, 4H), 7.31-7.397 (m, H), 8.23-8.30 (m, 1H), 8.38-8.40 (m, 1H).

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Procedure K: General Method for the Synthesis of Compound 25a-b

(6-fluoropyridin-3-yl)(1-(3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)methanone oxalate 25a Tz-4-1-154

A mixture of compound 24a and HCl (12N, 5 mL) in THF were stirred for 4 h at room temperature until TLC indicated that the reaction was complete, and then carefully neutralized with NaOH (1N) and extracted with CH₂Cl₂. The organic layer was dried and concentrated under reduce pressure. The crude product was purified with column chromatography to afforded 25a (yield, 79%). ¹H NMR (CDCl₃, free base): δ 1.84-2.03 (m, 4H), 2.38-2.46 (m, 1H), 2.75-3.02 (m, 7H), 3.21-3.34 (m, 2H), 3.55-3.63 (m, 1H), 3.82-3.89 (m, 1H), 4.30 (br, 1H), 7.03-7.11 (m, 4H), 8.33-8.39 (m, 2H), 8.79 (s, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₁H₂₃FN₂O₂·H₂C₂O₄) C, H, N.

(2-fluoropyridin-3-yl)(1-(3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)methanone 25b Tz-4-1-122

Compound 25b Tz-4-1-122 was prepared from compound 24b as describe in procedure G. Yield, 75%. ¹H NMR (CDCl₃, free base): δ 1.84-2.03 (m, 4H), 2.38-2.46 (m, 1H), 2.75-2.99 (m, 7H), 3.25-3.33 (m, 2H), 3.80-3.90 (m, 1H), 4.31 (br, 1H), 7.04-7.11 (m, 4H), 7.31-7.397 (m, H), 8.23-8.30 (m, 1H), 8.38-8.40 (m, 1H). The free base was converted to the oxalate salt. mp: ° C. Anal. (C₂₁H₂₃FN₂O₂·H₂C₂O₄) C, H, N.

2-bromo-3-(2-fluoroethoxy)pyridine 28

A mixture of 2-bromopyridin-3-ol 27 (0.1 mmol), 1-bromo-2-fluoroethane (0.2 mmol) and K₂CO₃ (0.4 mmol) in acetonitrile were stirred at refluxing for one day until TLC indicated that the reaction was complete, and then filter, dried and concentrated under reduce pressure. The crude product was purified with column chromatography to afforded 28 (yield, 68%). ¹H NMR (CDCl₃): δ 4.25-4.34 (m, 2H), 4.73-4.89 (m, 2H), 7.20-7.22 (m, 2H), 8.01 (s, 1H).

tert-butyl 4-(3-(2-fluoroethoxy)picolinoyl)piperidine-1-carboxylate 29

To a solution of 28 (7.3 mmol) in THF was added 1.6 N butyllithium/hexane solution (7 ml) at -78° C. under N₂ atmosphere. The solution was stirred at -78° C. for 1 hour and a solution of tert-butyl 4-(methoxy(methyl)carbamoyl)piperidine-1-carboxylate (1 g, 3.6 mmol) in THF was added. The solution was maintained at -78° C. for 4 hour, and then quenched with sat. aqueous NH₄Cl and allowed to warm to rt with stirring. The organic layer was separated and then the aqueous layer was extracted with ethyl acetate. The combine extract was washed with water, dried over Na₂SO₄, filtered, concentrated, and the residue chromatographed on silica eluted with an ethyl acetate-hexane (1:4, v/v) and the aimed product was obtained. The yield was 40%. NMR (CDCl₃): δ 1.47 (s, 9H), 1.62-1.70 (m, 4H), 2.83 (s, 2H), 3.64-3.73 (m, 1H), 4.17 (br, 2H), 4.25-4.36 (m, 2H), 4.68-4.87 (m, 2H), 7.23-7.26 (m, 1H), 7.38-7.40 (m, 1H), 8.25-8.30 (m, 1H).

(3-(2-fluoroethoxy)pyridin-2-yl)(1-(3-hydroxy-1,2,3,4-tetrahydronaphthalen-2-yl)piperidin-4-yl)methanone oxalate 31 (TZ-4-2-12)

Compound 31 (TZ-4-2-12) was prepared from compound 29 as describe in procedure C and E. The yield was 43%. ¹H

NMR (CDCl₃, free base): δ 1.73-1.98 (m, 4H), 2.32-2.40 (m, 1H), 2.71-2.96 (m, 8H), 3.31-3.33 (m, 1H), 3.49-3.56 (m, 1H), 3.80-3.89 (m, 1H), 4.24-4.36 (m, 2H), 4.69-4.87 (m, 2H), 7.09-7.25 (m, 4H), 7.34-7.38 (m, 2H), 8.27-8.29 (m, 1H). The free base was converted to the oxalate salt. mp: ° C.

Anal. (C₂₃H₂₇FN₂O₃·H₂C₂O₄) C, H, N.

In Vitro Biological Evaluations

Sigma Receptor Binding Assays.

The compounds were dissolved in DMF, DMSO, or ethanol, and then diluted in 50 mM Tris-HCl buffer containing 150 mM NaCl and 100 mM EDTA at pH 7.4, prior to performing the σ₁ and σ₂ receptor binding assays. The procedures for isolating the membrane homogenates and performing the σ₁ and σ₂ receptor binding assays have been previously described in detail.^{40,51}

Briefly, the σ₁ receptor binding assays were conducted in 96-well plates using guinea pig brain membrane homogenates (~300 μg protein) and ~5 nM (+)-[³H]pentazocine (34.9 Ci/mmol, Perkin-Elmer, Boston, Mass.). The total incubation time was 90 min at room temperature. Nonspecific binding was determined from samples that contained 10 μM of cold haloperidol. After 90 min, the reaction was quenched by adding 150 μL of ice-cold wash buffer (10 mM Tris-HCl, 150 mM NaCl, pH 7.4) using a 96 channel transfer pipet (Fisher Scientific, Pittsburgh, Pa.). The samples were harvested and filtered rapidly through a 96-well fiberglass filter plate (Millipore, Billerica, Mass.) that had been presoaked with 100 μL of 50 mM Tris-HCl buffer at pH 8.0 for 1 h. Each filter was washed 3 times with 200 μL of ice-cold wash buffer, and the filter counted in a Wallac 1450 MicroBeta liquid scintillation counter (Perkin-Elmer, Boston, Mass.).

The σ₂ receptor binding assays were conducted using rat liver membrane homogenates (~300 μg protein) and ~5 nM [³H]DTG (58.1 Ci/mmol, Perkin-Elmer, Boston, Mass.) in the presence of 1 μM of (+)-pentazocine to block σ₁ sites. The incubation time was 2 h at room temperature. Nonspecific binding was determined from samples that contained 10 μM of cold haloperidol. All other procedures were identical to those described for the σ₁ receptor binding assay above.

Data from the competitive inhibition experiments were modeled using nonlinear regression analysis to determine the concentration that inhibits 50% of the specific binding of the radioligand (IC₅₀ value). Competitive curves were best fit to a one-site fit and gave pseudo-Hill coefficients of 0.6-1.0. K_i values were calculated using the method of Cheng and Prusoff and are presented as the mean (±1 SEM). For these calculations, we used a K_d value of 7.89 nM for (+)-[³H]pentazocine and guinea pig brain and a K_d value of 30.73 nM for [³H]DTG and rat liver.

Vesicular Acetylcholine Transporter Binding Assays

In vitro binding assays of these novel compounds on VACHT were conducted with human vesicular acetylcholine transporter (VACHT) permanently expressed in PC12^{4123.7} cells at about 50 pmol/mg of crude extract. The radioligand used was 5 nM (-)-[³H]vesamicol, and the assay was conducted as described at final concentrations of 10⁻¹¹ M to 10⁻⁵ M novel compound (53). Unlabeled (-)-vesamicol was used as an external standard, for which K_i=15 nM, and the mixture was allowed to equilibrate for 20 hrs. Duplicate data were averaged and fitted by regression with a rectangular hyperbola to estimate the K_i value of the novel compound. All compounds were independently assayed at least two times.

Ligands for σ₁ receptors that are potent and selective are disclosed. The structures of prezmocol and trozamicol scaffolds of carbonyl-containing vesicular acetylcholine transporter (VACHT) inhibitors are explored. Of the 23 analogues synthesized and tested, 5 display very high affinity for σ₁

(K_i=0.48-4.05 nM) and high selectivity for σ₁ relative to σ₂ receptors (σ₁/σ₂ selectivity>749-fold). Four of the five compounds (14a, 14b, 14c and 14e) showed very low affinity for VACHT (K_i>290 nM) and the fifth compound (14g) show moderate affinity for VACHT (K_i=44.2 nM). The compound [1'-(4-fluorobenzyl)-3'-hydroxy[1,4']bipiperidiny-4-yl]-(4-fluorophenyl)-methanone (14a) displays very high affinity and selectivity for σ₁ receptor (K_i=0.48 nM, σ₁/σ₂>3600). All four of these compounds (14a, 14b, 14c and 14e) can be radiosynthesized with fluorine-18 or carbon-11, and can be used as PET probes for imaging σ₁ receptor in vivo.

Biological Binding Studies.

The σ₁ and σ₂ binding affinities (K_i, nM) of the new compounds were determined by using the competitive inhibition method with tritiated a ligands according to reported procedures.^{6,40,51} The σ₁ binding sites were assayed by using guinea pig brain membranes with the selective radioligand (+)-[³H]pentazocine. The σ₂ binding sites were assayed in rat liver membranes, a rich source of these sites, with [³H]DTG in the presence of (+)-pentazocine (100 nM) to mask σ₁ sites. VACHT binding was assayed using highly expressed human VACHT assayed with homogenized and partially clarified PC12^{123.7} cells by displacement of bound 5 nM (-)-[³H]vesamicol. Apparent dissociation constants for binding of the novel compounds in vitro are shown in Table 1.

Binding data identified a number of structure-activity trends. First, these new analogues generally have very low affinity for σ₂ receptors (K_i>1000 nM). For VACHT, only compounds 14g, 16e, and 16g display moderate affinity of 44.2±3.03, 48.6±8.37, and 35.5±11.1 nM. For the other ligands, the K_i values are greater than 100 nM. On the other hand, compounds 14a, 14b, 14c, 14e and 14g displayed very high affinity for a receptor (K_i<5 nM) and very good selectivity for σ₁ versus σ₂ receptors (>1000 fold). Among these latter compounds, 14a displayed the highest affinity for σ₁ receptors (K_i=0.48±0.14 nM), very good selectivity for σ₁ versus σ₂ receptors (>3000 fold), and good selectivity for σ₁ receptors vs. VACHT (2800 fold), indicating that 14a is selective for σ₁ receptor.

14a has two fragments that contain a fluorine atom. One is in the para-position of the carbonyl group in 4-fluorobenzoylpiperidine fragment, which is easy to label by displacement of a nitro group with ¹⁸F. The other fluorine atom is in the para-position of the 1-(4-fluorobenzyl)piperidin-3-ol fragment. This other site allows incorporation of fluorine-18 in two steps. The first step is to make 4-[¹⁸F]fluorobenzyl iodide,⁵² and the second step is to make the final labeled radiotracer by N-alkylation of 13a with 4-[¹⁸F]fluorobenzyl iodide. Labeling 14a with fluorine-18 at different positions also provides a unique way to investigate the metabolic stability of 14a in vivo by determining which site releases ¹⁸F—. In addition, the moderate lipophilicity of 14a (Log P value=2.83) indicates that compound 14a has good ability to enter into the brain and can be a suitable PET imaging probe or therapeutic agent for CNS disorders.

Comparing compounds 14a-h with 15a-g, N-benzyl substituted ligands display significantly higher binding affinities for a receptors, particularly for σ₁ receptor sites, whereas, N-benzoyl substituted ligands display low binding affinities for a receptors. For example, when the 4-fluorobenzoyl group in 15a is replaced by the 4-fluorobenzyl group in 14a, the σ₁ receptor affinity is increased dramatically, changing the K_i value from 3144 nM for 15a to 0.48 nM for 14a. The same trend was observed in all other corresponding compounds. In general, compounds containing substituted benzyl groups preferred to bind to a receptor sites; 14a-h, displayed very high σ₁ binding affinities in which the K_i values ranged from

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0.48 nM for 14a to 59.6 nM for 14h. However, compound 14g showed moderate VACHT binding affinity (44.2 ± 3.03 nM), whereas the other compounds 14a-f and 14h displayed lower VACHT binding affinities. The observation that compounds containing the substituted benzoyl groups 15a-g led to dramatic decrease in σ_1 receptor binding affinities ($K_i > 1000$ nM) is consistent with results reported in the literature.⁴⁰ This confirms that a carbonyl group between the substituted aromatic ring and the piperidine moiety plays an important role by affecting σ_1 receptor binding affinities. Without being limited by theory, the inventors hypothesize that incorporating the carbonyl group leads to more rigidity in the structure and might prevent the interaction of ligands with the σ_1 receptor binding site.

When the electron-withdrawing group fluoride (14a-d) was replaced with the electron-donating group methoxy (14e-h) at the para-position of the 4-substituted benzoylpiperidine fragment, the σ_1 binding affinities (nM) were decreased. For compounds 14a-d, a, binding affinities (nM) were 0.48 ± 0.14 , 4.03 ± 0.47 , 1.36 ± 0.28 and 22.8 ± 2.32 respectively; for compounds 14e-h, the σ_1 binding affinities (nM) were 2.51 ± 0.34 , 25.9 ± 0.96 , 4.05 ± 0.88 and 59.64 ± 2.22 respectively. The decrease in affinity ranged from 2.6 fold for 14h vs. 14d to 6.4 fold for 14b vs. 14f.

Comparing the structures of 14a, 14b, and 14c, the only difference among them is the substitution at the para-position of the benzyl moiety on the terminal piperidine ring of (3'-hydroxy-1,4'-bipiperidin-4-yl)(4-fluorophenyl)methanone. The σ_1 K_i -values (nM) are displayed in the order of -F>-H>-OCH₃ with the dissociation constant values being 0.48 ± 0.14 for 14a, 1.36 ± 0.28 for 14c and 4.03 ± 0.47 for 14b. Compared with 14b, incorporating an electron-withdrawing fluorine atom at the 4-position of the benzyl group (14a) diminished the basicity of the nitrogen atom in the terminal piperidine ring and the ligand affinity for σ_1 receptor was increased by 2.8 fold; on the contrary, incorporating an electron-donating methoxyl group (14b) enhanced the basicity of the nitrogen atom in the terminal piperidine ring and the ligand affinity for σ_1 receptor was decreased by approximately 3 fold. A similar trend is observed in 4-methoxybenzoyl containing compounds 14e, 14f and 14g. To further test the effect of the electron density on the benzyl ring, compounds 14d and 14h containing the relatively high electron-density pyridine ring, with the N-atom in the meta-position, were synthesized and screened. However, decrease in σ_1 binding affinities was observed. For 14d, the σ_1 binding affinity was decreased more than 17 fold as expressed by K_i value changing from 1.36 ± 0.28 nM for 14c to 22.8 ± 2.32 nM for 14d. For 14h, the σ_1 binding affinity was decreased 15 fold as expressed by K_i value changing from 4.05 ± 0.88 nM for 14g to 59.64 ± 2.22 nM for 14h.

The presamical scaffolds (14a-h) display substantially higher affinities for σ_1 receptors than the trozamicol scaffolds (16a-h) do. The only structural difference between presamical and trozamicol is the position of the nitrogen atom in the terminal piperidine ring. Some trozamicol scaffold analogues display moderate σ_1 receptor binding affinities, for example 16a (50.0 ± 7.9 nM) and 16b (91.1 ± 19.9 nM). The difference in binding affinities for presamical and trozamicol analogues is 11 fold for 14f vs. 16f (25.9 ± 0.96 nM vs. 297 ± 27 nM) and 104 fold for the most potent 14a vs. 16a (0.48 ± 0.14 nM vs. 50.0 ± 7.9 nM).

Overall, (1) N-alkylation with benzyl groups of (3'-hydroxy-1,4'-bipiperidin-4-yl)(4-substituted phenyl)methanone to form tertiary amines generally gives very high σ_1 receptor binding affinities and high selectivity for σ_1 receptor vs. σ_2 receptor and VACHT sites, (2) N-acylation with benzoyl

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groups of (3'-hydroxy-1,4'-bipiperidin-4-yl)(4-substituted phenyl)methanone to form tertiary amides diminishes the σ_1 receptor binding affinity, and (3) prezamicol analogues display high σ_1 receptor binding affinities, whereas, trozamicol analogues display moderate to low σ_1 receptor binding affinities. Although it is well known that σ_1 receptor distinguishes between enantiomers,^{30,31} only racemic mixtures are used in this study.

Compounds were tested for VACHT binding using highly purified Torpedo synaptic vesicles. The σ_1 and σ_2 binding affinities were tested in rat brain and in guinea pig membranes, respectively. Structures and Equilibrium binding constants (K_i) are reported in Table 5 and Table 6, respectively. As in previous studies, we define ligand selectivity in terms of a selectivity index which is calculated as K_i (σ_1 or σ_2)/ K_i (VACHT).

In the previous study, ((±)-trcms-2-Hydroxy-3-(4-(4-[¹⁸F] fluorobenzoyl)piperidino)tetralin (FBBV) as a potent VACHT tracer is discovery. In the present study, fluorine group (FBBV) was replaced with MeNH and OCH₂CH₂F. Inspiring, TZ-3-156 ($K_i = 3.03$ nm) displayed comparable affinity with FBBV ($K_i = 2.70$ nm), suggesting that this structure with the amino tetralin structure in the region of ring C would produce a potent yet more hydrophilic drug that will improve the specific to nonspecific ratio for binding.

To extend our structure modification studies, the phenyl ring in fragment C was replaced with the heterocyclic ring and the results of the binding assays for compounds 11a-b, d-g, 12a-d, 13d, 20a-b, 25a-b, 31, TZ-5-3, TZ-3-156 and TZ-3-148 are shown in Table 5. Substituting the 4-position of pyridine group with methyl group, methoxy group, fluorine group generated 11a-b, f-g, 12a-c, 25a-b, 31 and TZ-5-3 (OS-5-40, OS-5-88, TZ-4-1-92, TZ-4-1-96, OS-5-70, OS-5-82, OS-5-110, TZ-4-1-154, TZ-4-1-122, TZ-4-2-12, TZ-5-3). The affinities for VACHT are 11b (OS-5-88) > 11g (TZ-4-1-96) > 25b (TZ-4-1-122) > 25a (TZ-4-1-154) > 11a (OS-5-40) > TZ-5-3 > 12a (OS-5-70) = 12b (OS-5-82) > 31 > TZ-4-2-12 > 12c (OS-5-110) > 11f (TZ-4-1-92). In contrast to 4'-benzoylpiperidine ($K_i = 4.30 \pm 1.00$ nm),⁶⁶ 4-pyridine-carbonylpropionylvesamicol 12c (OS-5-110) ($K_i = 118 \pm 23.6$ nm) displayed 27-fold lower affinity for VACHT than 4'-benzoylpiperidine, while methyl group substituting on pyridine ring 11a (OS-5-40) ($K_i = 27.9 \pm 7.99$ nm) was 4-fold more potent than the no substituting 12c (OS-5-110). After replaced by methoxy group, compound 11b (OS-5-88) has the highest affinity for the VACHT ($K_i = 8.36 \pm 0.68$ nm). Surprisingly, When carbonylpropionylvesamicol group close to nitrogen atoms of the pyridine, methoxy group compound 11f (TZ-4-1-92) ($K_i = 121 \pm 29.2$ nm) displayed lower activity than other methoxy group compound 11b (OS-5-88). To pyridine ring including, fluorine group, the introduction of a fluorine atom in a different location on the pyridine ring 11g (TZ-4-1-96), 25b (TZ-4-1-122 and 25a (TZ-4-1-154) (K_i values: 10.1 ± 1.54 , 12.7 ± 2.41 nm) displayed higher affinity for VACHT than ethylfluorine compounds TZ-5-3 and 31 > TZ-4-2-12 ($K_i = 38 \pm 3.84$ and 76.4 ± 8.33 nm). To further probe suitable six-membered heterocyclic including nitrogen in fragment C vesamicol analogues, 1-methylpyridin-2(1H)-one derivatives 20a (TZ-4-1-148) and 20b (TZ-4-1-154) were also synthesized. It was observed that two compounds displayed lower affinity, suggesting that substitute group directly coupling nitrogen was unfavorable than no substitute nitrogen at enhancing affinity for VACHT. As observed in the pyridine ring in fragment C, the ortho-carbonylpropionylvesamicol group displayed lower affinity for VACHT than meta- and para-position (TZ-5-3 vs 31 > TZ-4-2-12, 11b OS-5-88 vs 11f TZ-4-1-92, 20a TZ-4-1-148 vs 20b TZ-4-1-154). Taken together, we there-

fore conclude that, with suitable substitution, the pyridine can replace the phenyl group in fragment C.

Replacement of the six-member heterocyclic ring in fragment C with the five-member heterocyclic ring 11e and 11d (TZ-4-1-114 and OS-5-60) displayed high affinity (K_i values: 18.4±2.54 and 26.6±3.83 nM), suggesting that methyl group directly coupling nitrogen in five-member heterocyclic ring is more effective in altering affinity for VACHT than the six-member heterocyclic ring in fragment C.

The introduction of an amino group into 2 (benzovesamicol) increases affinity for VACHT by an order of magnitude, making 3 (K_i=0.0065±0.0005 nM) the most potent VACHT inhibitor reported to date. We therefore sought to investigate the effect of the 5-amino group on the affinity of 4-benzoylpiperidines. In the previous study, no improvement was observed for the 4-benzoylpiperidines or the 4-(thien-2-yl) carbonylpiperidines. In the present study, 4-benzoyl group was replaced by heteroaromatic ring on fragment-C. It is consistent with previous results that 4-(heteroaromatic)-5-amino-carbonylpiperidines (12b, 12a and 12d OS-5-82, OS-5-70 and OS-5-72-2) still displayed lower affinity for VACHT than no substitute amino compounds 11b, 11a and 11d OS-5-88, OS-5-40 and OS-5-60. There was, nevertheless, a separation in affinity between the regioisomeric pairs (13d OS-5-72-1 vs 12d OS-5-72-2) similar to that observed with 2 and its 8-amino isomer. Thus, it would appear that the structure-activity relationships of the 4-heteroaromaticpiperidines largely reduced, but are not identical to, those of the 4-phenylpiperidines.

The moderate to high affinity of vesamicol and many of its analogues for σ₁ and σ₂ receptors reduces the selectivity of these compounds for the VACHT and potentially compromises their utility as VACHT ligands. Consequently, we also screened the target compounds for sigma binding to identify selective VACHT ligands from this new structural class of agents. Among the compounds tested, there was wide variability in affinity for σ₁ or σ₂ receptors; only one compound (TZ-3-156) displayed high affinity for σ₁ receptors (K_i=13.23±0.48 nM), while all other compounds exhibited poor affinity for a receptors. As in previous studies of vesamicol analogues, we have derived estimates of ligand selectivity, expressed as a selectivity index, by comparing K_i values for VACHT obtained from highly purified Torpedo synaptic vesicles, with corresponding values for σ₁ and σ₂ receptors determined from rat or guinea pig brain preparations. In the present study, among the six-member heterocyclic ring analogues, 3 compounds (31 TZ-4-2-12, 11 f TZ-4-1-92 and 20b TZ-4-1-148) showed poor selectivity while 10 others (11a-b, g, 12a-c, 20a, 25a-b and TZ-5-3) (OS-5-40, OS-5-88, TZ-4-1-96, OS-5-70, OS-5-82, OS-5-110, TZ-4-1-150, TZ-4-1-154, TZ-4-1-122, TZ-5-3)) displayed moderate or high VACHT/σ selectivity (Table 5). Among the five-member heterocyclic ring analogues 11d OS-5-60, 11e TZ-4-1-114, 12d OS-5-72-2 and 13d OS-5-72-1, all compounds except 12d OS-5-72-2 displayed moderate or high VACHT/σ selectivity. In contrast, among phenyl analogues TZ-3-156 and TZ-3-148, TZ-3-156 exhibited high selectivity while TZ-3-148 showed poor selectivity for VACHT.

TABLE 5

Compounds tested for VACHT binding		Ar ₁	R ₅	R ₆
	OS-5-40	11a	H	H
	OS-5-88	11b	H	H

TABLE 5-continued

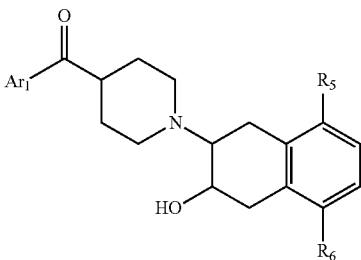
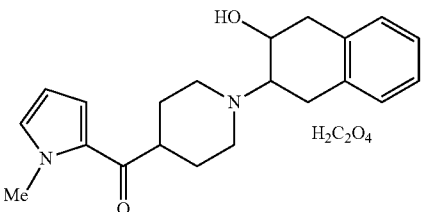
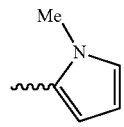
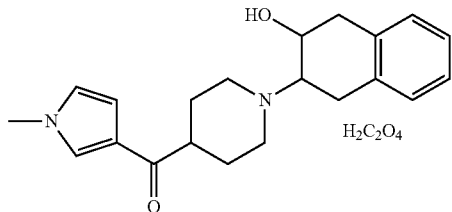
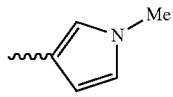
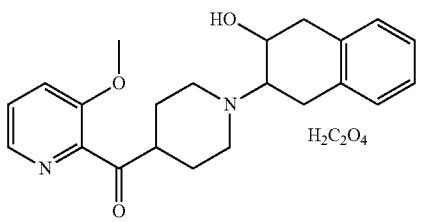
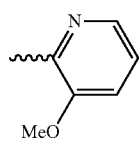
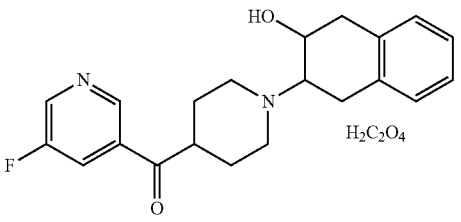
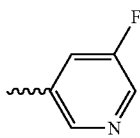
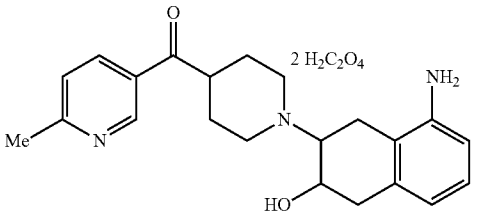
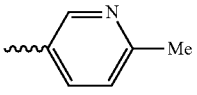
Compounds tested for VAcHT binding		Ar ₁	R ₅	R ₆
				
	OS-5-60 11d		H	H
	TZ-4-1-114 11e		H	H
	TZ-4-1-92 11f		H	H
	TZ-4-1-96 11g		H	H
	OS-5-70 12a		NH ₂	H

TABLE 5-continued

Compounds tested for VAcHT binding

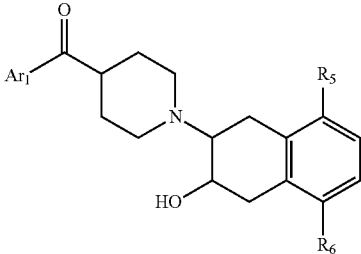
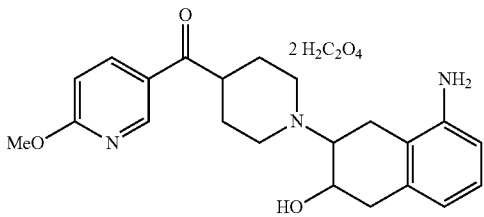
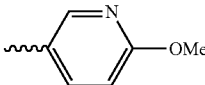
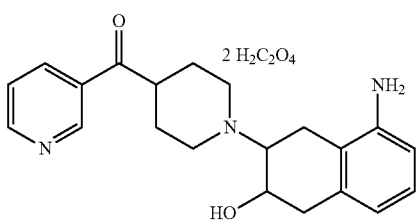
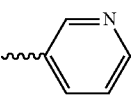
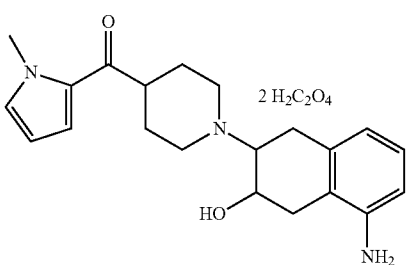
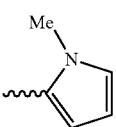
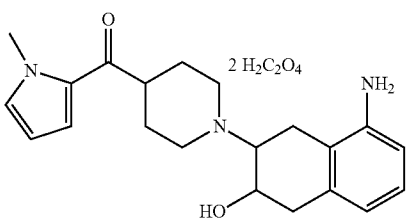
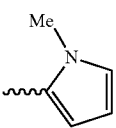
		Ar ₁	R ₅	R ₆
				
 OS-5-82 2 H ₂ C ₂ O ₄	OS-5-82	12b		NH ₂ H
 OS-5-110 2 H ₂ C ₂ O ₄	OS-5-110	12c		NH ₂ H
 OS-5-72-2 C ₂₅ H ₃₁ N ₃ O ₁₀ MW: 533.53 2 H ₂ C ₂ O ₄	OS-5-72-2	12d		NH ₂ H
 OS-5-72-1 2 H ₂ C ₂ O ₄	OS-5-2-72-1	13d		H NH ₂

TABLE 5-continued

Compounds tested for VAcHT binding		Ar ₁	R ₅	R ₆
	TZ-4-1-150 20a		H	H
	TZ-4-1-148 20b		H	H
	TZ-4-1-154 25a		H	H
	TZ-4-1-122 25b		H	H
	TZ-4-2-12 31		H	H

TABLE 5-continued

Compounds tested for VAcHT binding							
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Ar_1	R_5	R_6					
	H	H					
	TZ-3-156 <table border="0" style="margin-left: 20px;"> <tr> <td style="text-align: center;">Ar_1</td> <td style="text-align: center;">R_5</td> <td style="text-align: center;">R_6</td> </tr> <tr> <td style="text-align: center;"> </td> <td style="text-align: center;">H</td> <td style="text-align: center;">H</td> </tr> </table>	Ar_1	R_5	R_6		H	H
Ar_1	R_5	R_6					
	H	H					
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Ar_1	R_5	R_6					
	H	H					

TABLE 6

Binding constants of structures in Table 5							
Compd.	LogP	VAcHT	σ_1 Ki (nM) \pm SEM	σ_2	$\sigma_1/VAcHT$	$\sigma_2/VAcHT$	
OS-5-40	11a	2.29	27.9 \pm 7.99	2115 \pm 289	3771 \pm 886	75.8	135.2
OS-5-88	11b	2.64	8.36 \pm 0.68	986 \pm 160	2572 \pm 139	117.9	307.6
OS-5-60	11d	1.42	18.4 \pm 2.54	1302 \pm 93	2553 \pm 339	70.7	138.8
TZ-4-1-114	11e	1.84	26.6 \pm 3.83	3040 \pm 187	7708 \pm 1059	114.2	289.7
TZ-4-1-92	11f	2.96	121 \pm 29.2	20975 \pm 3809	1564 \pm 63	173.3	12.9
TZ-4-1-96	11g	1.99	10.1 \pm 1.54	709 \pm 79	1690 \pm 56	70.2	167.3
OS-5-70	12a	1.33	48.0 \pm 11.4	11362 \pm 2040	27525 \pm 6943	236.7	573.4
OS-5-82	12b	1.66	48.0 \pm 11.4	5682 \pm 26	10813 \pm 1282	118.4	225.3
OS-5-110	12c	0.87	118 \pm 23.6	20346 \pm 8133	5078 \pm 221	172.4	43.0
OS-5-72-2	12d	0.53	2310 \pm 393	9237 \pm 793	11396 \pm 2500	4.0	4.9
OS-5-2-72-1	13d	0.53	38.7 \pm 6.95	7664 \pm 233	6987 \pm 1482	198.0	180.5
TZ-4-1-150	20a	0.60	66.2 \pm 10.2	3345 \pm 46	2124 \pm 251	50.5	32.1
TZ-4-1-148	20b	0.64	182 \pm 64.8	2155 \pm 182	3339 \pm 127	11.8	18.3
TZ-4-1-154	25a	2.02	26.1 \pm 2.41	682 \pm 27	1750 \pm 93	26.2	67.0
TZ-4-1-122	25b	1.65	12.7 \pm 1.06	1299 \pm 49	3529 \pm 331	102.2	277.9
TZ-4-2-12	31	3.18	76.4 \pm 8.33	20016 \pm 6798	303 \pm 130	262.0	4.0
TZ-5-3		2.87	38 \pm 3.84	2122 \pm 49	1859 \pm 144	55.8	48.9
TZ-3-156		2.61	3.03 \pm 0.48	304.63 \pm 21.94	6181.73 \pm 537.07	100.5	2040
TZ-3-148		3.19		13.23 \pm 0.20	1885.84 \pm 239.40		

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HPLC Conditions to Confirm the Purity of Final Compounds

The analytical HPLC system consisted of an Alltech Econosil reversed phase C18 column (250×4.6 mm, 10 uA) (Alltech Associates, Deerfield, Ill.). The mobile phase was 35% acetonitrile and 65% 0.1 M ammonium formate buffer (PH=4.5) at 1.0 mL/min flow rate. The UV wavelength is 254 uM. Under these conditions the benzamide analogs was eluted with a retention time varying from 10-50 min. The purity of the final compounds was >95%.

Chemistry

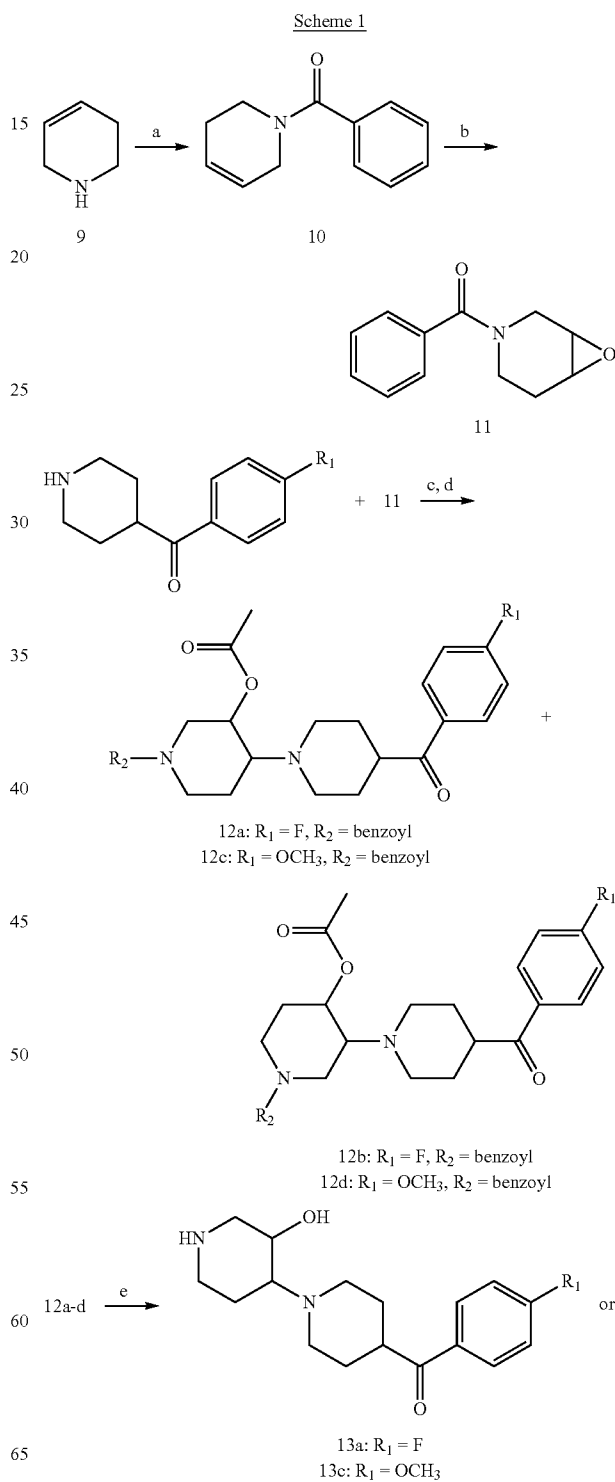
The target compounds were synthesized as depicted in Schemes 1-3. The synthesis started with 1,2,3,6-tetrahydropyridine, in which the secondary amine was first protected by reacting with benzoyl chloride to form benzamide 10. The alkenyl double bond of compound 10 was oxidized by meta-chloroperoxybenzoic acid (m-CPBA) to form epoxide 11. The epoxide 11 was refluxed with substituted 4-benzoylpiperidine hydrochloride salts, namely (4-(4'-fluorobenzoyl)piperidine hydrochloride or 4-(4'-methoxybenzoyl)piperidine hydrochloride), in ethanol with triethylamine as the base to afford tertiary amino alcohol intermediates. To improve the yield, commercially available 4'-substituted 4-benzoylpiperidine hydrochloride salts were used in excess and the reaction temperature was kept below 75° C. This reaction afforded (3'-hydroxy-1,4'-bipiperidin-4-yl)(4-substituted phenyl) methanone intermediates and their regioisomers, (4'-hydroxy-1,3'-bipiperidin-4-yl)(4-substituted phenyl)methanone intermediates. The separation of these regioisomers by silica gel chromatography was difficult. The mixture of corresponding regioisomers was reacted with acetic anhydride to convert the free hydroxyl group to the corresponding acetates 12a-d. Subsequently, 12a and 12b were separated easily by silica gel chromatography. To confirm the structures of the regioisomers (12a(c) and 12b(d)), in addition to 1D NMR, 2D NMR was performed for regioisomers, 12c and 12d. The ¹H, ¹³C NMR and HMQC spectra showed $\delta_H=5.31/2.68, 3.79/3.12, 2.06/1.47, 3.70, 2.48$ and $\delta_H=4.89/2.66, 4.09/2.82, 1.95/1.41, 3.43$ and 2.41 belong to the CH₂-a, CH₂-c, CH-d and CH-e (FIG. 2). The exact regioisomer (12a(c) or 12b(d)) were further verified by ¹H-¹H COSY. For regioisomer 12c, the correlations were observed between $\delta_H=5.31$ (CH-a)/ $\delta_H=2.68$ (CH-a') and $\delta_H=2.48$ (CH-e); $\delta_H=3.79$ (CH-b)/ $\delta_H=3.12$ (CH-b') and $\delta_H=2.06$ (CH-c)/ $\delta_H=1.47$ (CH-c'). In contrast, for regioisomer 12d, $\delta_H=4.89$ (CH-a)/ $\delta_H=2.66$ (CH-a') has no correlation with $\delta_H=2.41$ (CH-e), and $\delta_H=4.09$ (CH-b)/ $\delta_H=2.82$ (CH-b') and has no correlation with $\delta_H=1.95$ (CH-c)/ $\delta_H=1.41$ (CH-c'), but it has correlation with $\delta_H=1.95$ (CH-c)/ $\delta_H=1.41$ (CH-c') and $\delta_H=1.41$ (CH-d).

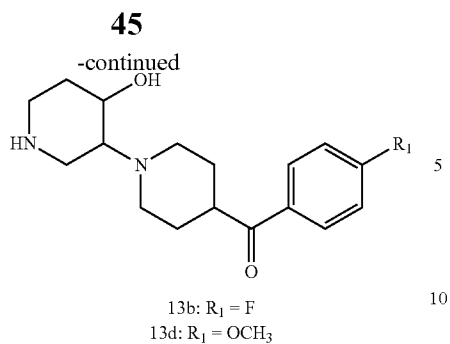
12a and 12b were hydrolyzed under strongly acidic conditions to remove both the acetyl and benzoyl groups, which afforded key intermediates 13a and 13b. A similar procedure was used to make corresponding regioisomers 13c and 13d. Intermediates 13a and 13c are similar in structure to presamicol, and 13b and 13d are similar in structure to trozamicol (Scheme 1). Starting with key intermediates 13a and 13c, the target compounds 14a-h were obtained via N-alkylation with various substituted benzyl halides in yields ranging from 34% to 84%. All the substituted benzyl halides are commercially available, and they were used in slight excess (Scheme 2). The target compounds 15a-g were obtained by coupling the various substituted benzoic acids with the piperidine intermedi-

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ates 13a and c using bis(2-oxo-3-oxazolidinyl)phosphonic chloride (BOP-Cl) in yields ranging from 22% to 90%. Compounds 16a-h were synthesized by following similar procedure to synthesize compounds 14a-h, except that the trozamicol analogues 13b and 13d were utilized (Scheme 3). All the products were converted into oxalates for determining the bioactivity affinities.

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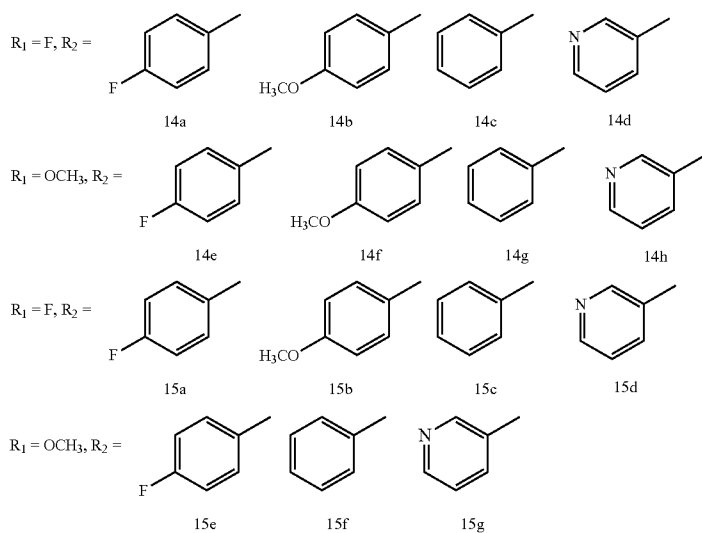
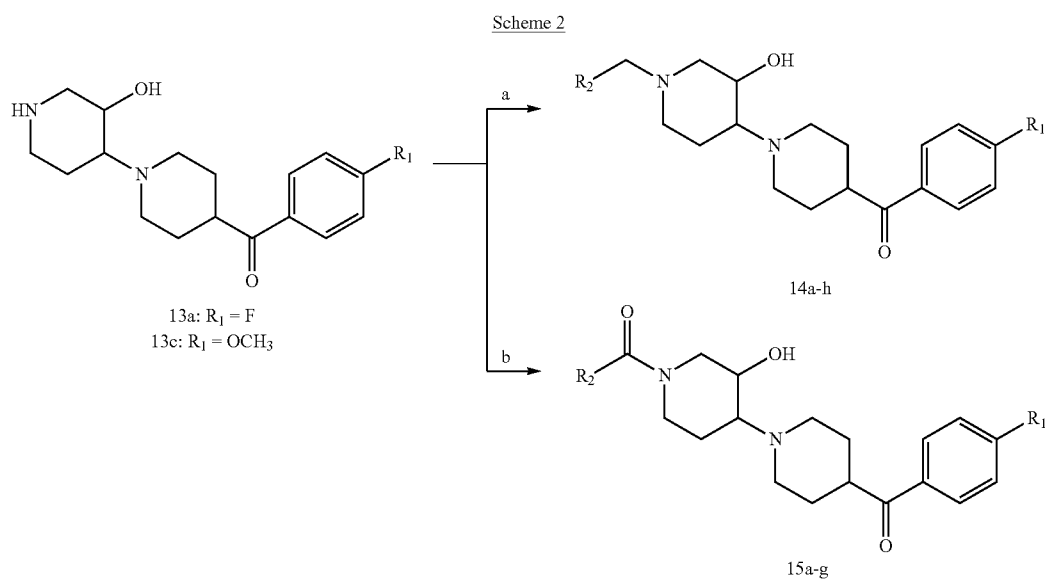




Reactions and reagents:

(a) benzoyl chloride, (C₂H₅)₃N; (b) m-CPBA, CH₂Cl₂; (c) 4-substituted benzoyl piperidine hydrochloride, (C₂H₅)₃N, ethanol, 0° C.; (d) acetic anhydride, CH₂Cl₂; (3) 6N HCl, reflux.

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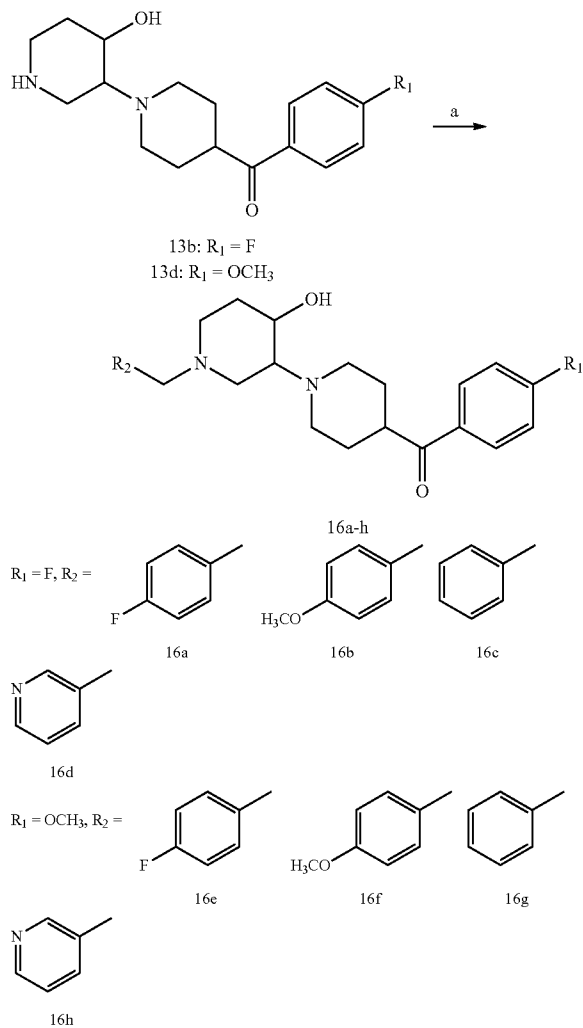


Reactions and reagents:

(a) substituted benzyl halide (C₂H₅)₃N, CH₂Cl₂;
(b) substituted benzoic acid, BOP-Cl, (C₂H₅)₃N, CH₂Cl₂

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Scheme 3



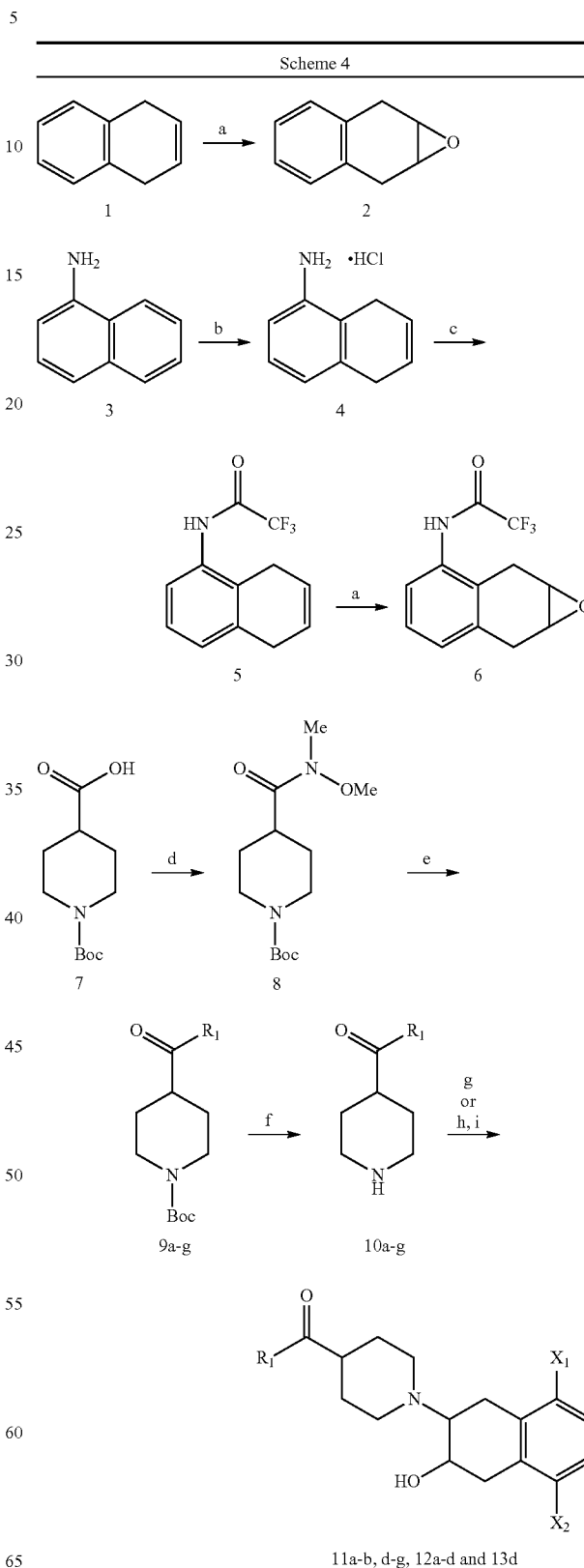
Reactions and reagents:

(a) substituted benzyl halide, ((C₂H₅)₃N, CH₂Cl₂)

The target compounds 11a-b, d-g, 12a-d and 13d (OS-5-40, OS-5-88, OS-5-60, TZ-4-1-114, TZ-4-1-92, TZ-4-1-96, OS-5-70, OS-5-82, OS-5-110, OS-5-72-2, OS-5-2-72-1) were synthesized as depicted in Scheme 4. Piperidine-4-carboxylic acid was protected by a BOC group and then condensed with N,O-dimethylhydroxylamine after activation of the acid by CDI to give 8 that provided a versatile intermediate by halogen-metal exchange with tert-butyllithium except for 9d, to generate 9a-c, 9e-g. To 9d, the first step of the reaction consists on the formation of lithium intermediate by reacting N-methylpyrrole with tert-butyl lithium. Side reactions were avoided by cooling the reaction down to -78° C. during the addition of tert-butyl lithium, and subsequent warming up to room temperature and followed by a nucleophilic addition of the lithiated intermediate to give 9d. These key intermediates 9a-g were deprotected with TFA to give piperidine 10a-b, d-g which were reacted with epoxide 2 under S_N2 conditions to give target compounds 11a-b, d-g (OS-5-40, OS-5-88, OS-5-60, TZ-4-1-114, TZ-4-1-92, TZ-4-1-96) in moderate yields. Similarly, secondary amines 10a-d were reacted with 6 and then hydrolyzed by sodium hydrox-

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ide aqueous solution to provide the regioisomeric pairs 12a-d and 13d. (OS-5-70, OS-5-82, OS-5-110, OS-5-72-2, OS-5-2-72-1)



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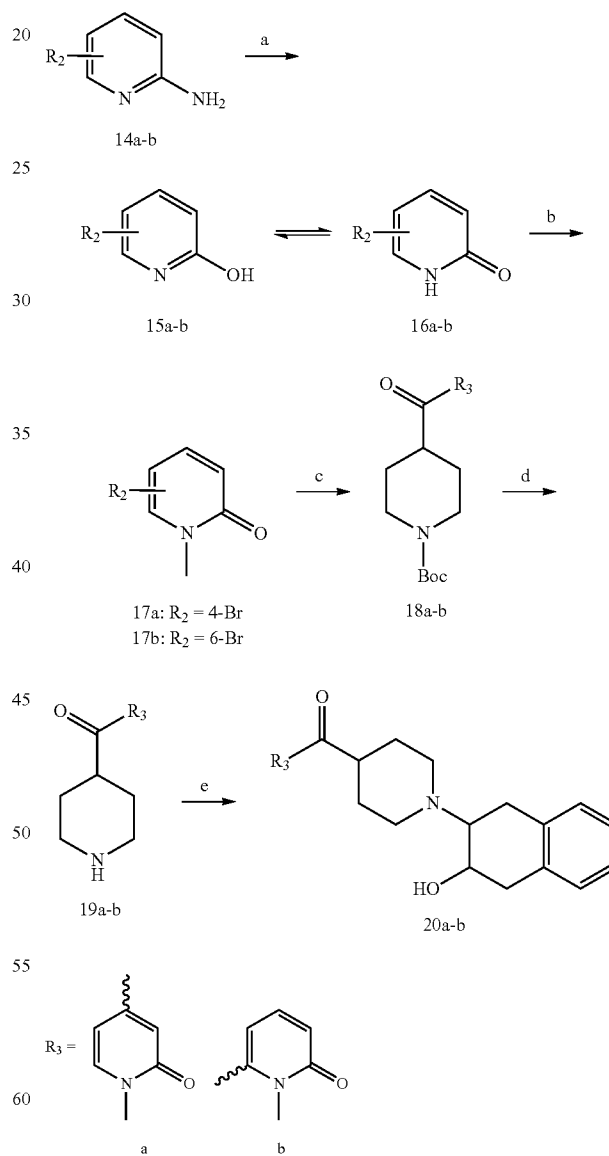
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Cpd	R ₁	X ₁	X ₂
11a		H	H
11b		H	H
11d		H	H
11e		H	H
11f		H	H
11g		H	H
12a		NH ₂	H
12b		NH ₂	H
12c		NH ₂	H
12d		NH ₂	H
13d		H	NH ₂

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Synthesis of 20a-b (TZ-4-1-150 and TZ-4-1-148) began with pyridin-2-amine 14a-b as shown in Scheme 5. Diazotization of pyridin-2-amine 14a-b in aqueous sulphuric acid gave tautomer 15a-b and 16a-b. According to the literature, the more polar lactam form 16a-b, which is in tautomeric equilibrium with the lactim form 15a-b, is favored in polar solvents. Therefore, N-Arylation of this substrate with iodomethane was found to strongly predominate over O-arylation, in acetone at refluxing and N-arylation produced 17a-b which could be converted into the corresponding ketone 18a-b by halogen-metal exchange in aqueous THF. Boc deprotection of ketone 18a-b with TFA followed by S_N2 conditions with epoxide 2 afforded target compounds 20a-b. (T1-4-1-150 and TZ-4-1-148)

Scheme 5

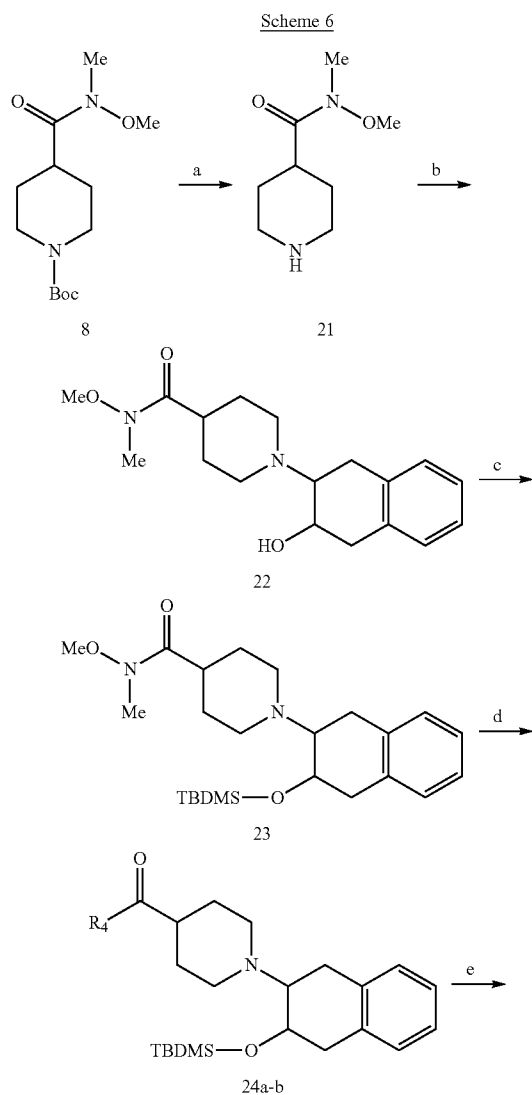


^aReagents and conditions: (a) mCPBA, CH₂Cl₂; (b) Na, EtOH, xylene; (c) (CF₃CO)₂O, NEt₃, CH₂Cl₂; (d) CDI, N,O-Dimethylhydroxylamine hydrochloride, NEt₃, CH₂Cl₂; (e) ArLi, THF; (f) TFA, CH₂Cl₂; (g) 1,4-dihydronaphthalene oxide, NEt₃, EtOH; (h) N-(trifluoroacetyl)-l-amino-5,8-dihydronaphthalene oxide, NEt₃, EtOH; (i) NaOH, H₂O, EtOH

^aReagents and conditions: (a) H₂SO₄/NaNO₂; (b) CH₃I, K₂CO₃; (c) BuLi, THF; (d) TFA, CH₂Cl₂; (e) 1,4-dihydronaphthalene oxide, NEt₃, EtOH.

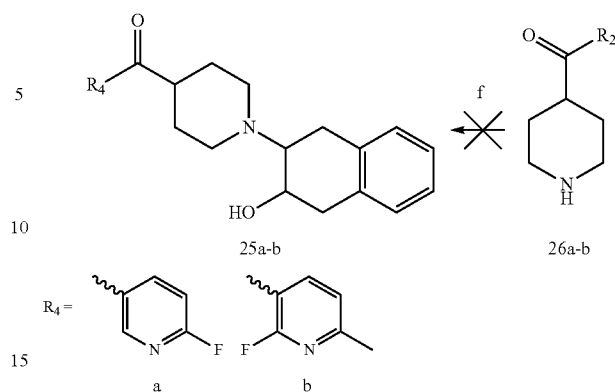
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Direct condensation of secondary amines and epoxide is a feasible method for obtaining vesamicol analogues. Although the preparation of the vesamicol 25a-b (TZ-4-1-154 and TZ-4-



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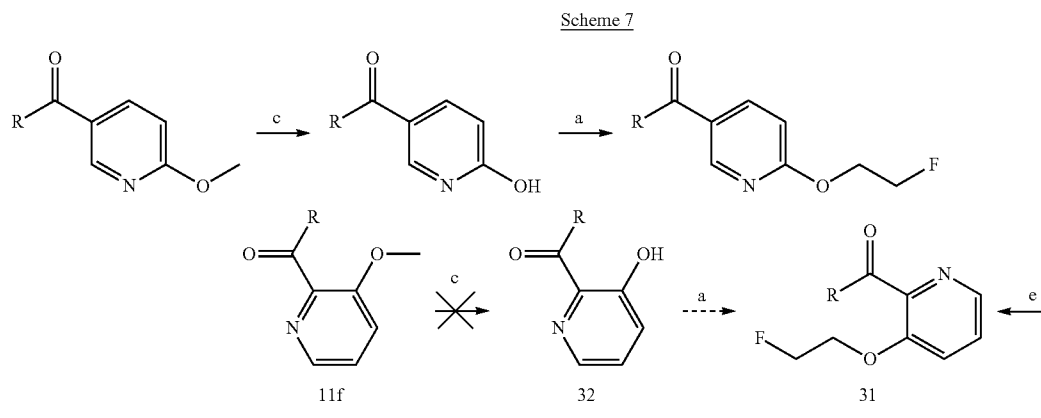
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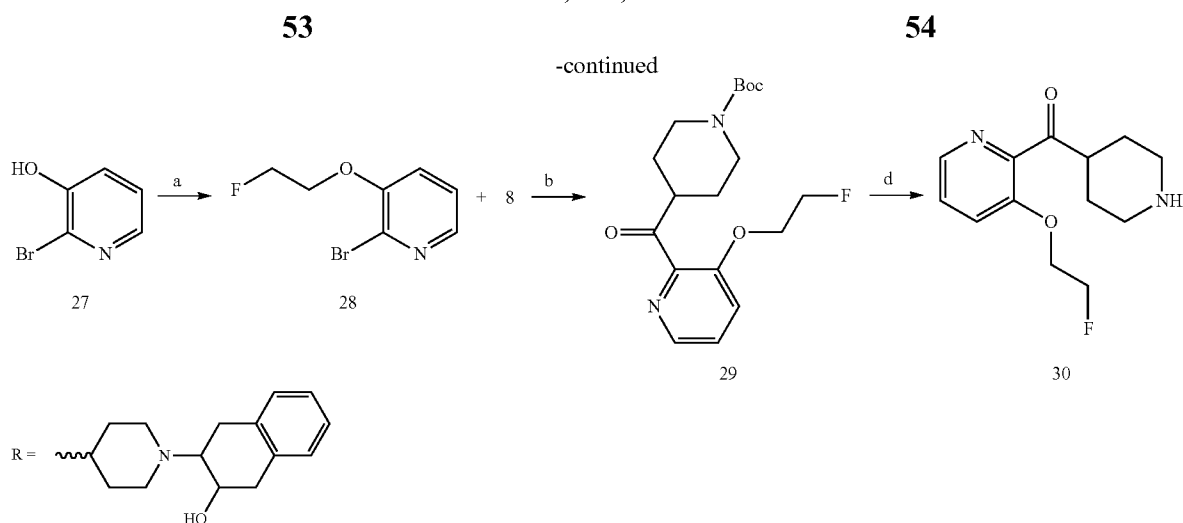


1-122) seemed trivial, it proved more problematic in practice, which probably arose from low solubility of secondary amines 26a-b (methanol, ethanol, dichloromethane, DMF, DMSO), than we had hoped. In an alternative route, the key intermediate 23 was utilized for producing vesamicol 25a-b (TZ-4-1-154 and TZ-4-1-122) which began with Boc deprotection of compound 8 with TFA followed by condensation with epoxide 2 afforded intermediate 22. Hydroxyl group of intermediate 22 was protected by a TBDMS group and then halogen-metal exchange with tert-butyllithium to provide compound 24a-b followed by an acidic workup, to liberate the hydroxy group, furnished the desired fluorine vesamicol 25a-b (TZ-4-1-154 and TZ-4-1-122) in Scheme 6.

Reagents and conditions in Scheme 6: (a) TFA, CH₂Cl₂; (b) 1,4-dihydronaphthalene oxide, NEt₃, EtOH; (c) TBDM-SCI, imidazole, CH₂Cl₂; (d) n-BuLi, THF; (e) 12N HCl, THF; (f) 1,4-dihydronaphthalene oxide, NEt₃, EtOH.

As for the preparation of vesamicol 31 (TZ-4-2-12), although direct nucleophilic fluorination is generally a better method from 11f, we found that methyl group deprotection of compound 11f could not be obtained by BBr₃, TMSI, etc. We chose another way of synthesis which introduction of the ethyl fluorine moiety was realized starting from commercially available 1-bromo-2-fluoroethane, using K₂CO₃, to afford compound 28. Subsequently, ethyl fluorine vesamicol 31 (TZ-4-2-12) was prepared according to similar procedure of making 11a-b, d-g from compound 28 (Scheme 7).





*Reagents and conditions: (a) Br(CH₂)₂F, K₂CO₃, rt; (b) BuLi, THF; (c) B₃Br or TMSI; (d) TFA, CH₂Cl₂, rt; (e) 1,4-dihydronaphthalene oxide, NEt₃, EtOH.

Example 1

Biodistribution Studies in Rats of [¹¹C]11e

The biodistribution and regional brain uptake of [¹¹C]11e was evaluated in 150-200 gram adult male Sprague-Dawley rats. Anesthetized rats were injected with 150-200 μCi of [¹¹C]11e and euthanized at 5, 30 and 60 min post-i.v. injection (p.i.). Target and non-target tissues were harvested and weighed; brain regions of interest were identified using gross dissection techniques. Radioactivity was measured in a well counter along with a standard dilution of the injectate. The results (Table 7 and FIG. 3) indicate high initial uptake and good retention of [¹¹C]11e in the brain with low radioactivity in the blood. Blocking was evaluated in rats pretreated with 1 mg/kg 11a 5 min prior to injection of [¹¹C]11e. The uptake at 30 min was significantly reduced for brain stem, cerebellum, cortex, striatum, hippocampus and total brain; the percentage reduction in uptake (% I.D./gram) was 82, 62, 87, 87, 68 and 63 respectively. By 30 minutes post-i.v., most of the radioactivity (31% of the total injected dose) was in the liver. Due to page limitations, distribution of radioactivity in the peripheral organs is not shown.

TABLE 7

The biodistribution of [¹¹ C]11e in male S.D. rats.			
organs	5 min	30 min	60 min
blood	0.12 ± 0.01	0.09 ± 0.01	0.13 ± 0.01
brain stem	2.73 ± 0.27	1.47 ± 0.16	0.88 ± 0.11
cerebellum	2.60 ± 0.13	1.29 ± 0.10	0.75 ± 0.09
cortex	2.14 ± 0.24	1.69 ± 0.15	0.86 ± 0.02
striatum	2.33 ± 0.10	1.58 ± 0.31	0.77 ± 0.07
hippocampus	2.36 ± 0.08	1.50 ± 0.05	0.79 ± 0.07
total brain	2.37 ± 0.08	1.46 ± 0.17	0.80 ± 0.07

*selected % ID/g values (mean ± SD) with n = 4 rats per group.

Example 2

We report the radiosynthesis and in vivo evaluation of [¹¹C]14e as a PET tracer to quantify the σ₁ density in human brain. Radiosynthesis of [¹¹C]14e was accomplished by alkylation of the corresponding desmethyl precursor with [¹¹C]CH₃I. Mice were injected with 80-320 μCi of [¹¹C]14e and euthanized at 5, 30 and 60 min. post-injection. Tissue

samples were taken and converted to % ID/g for each organ. Ex vivo autoradiography studies were also conducted in mice at 30 min post-injection of the radiotracer. A microPET imaging study was also performed on a male cynomolgus monkey.

[¹¹C]14e was synthesized in high yield (~45%) and high specific activity (>20 Ci/μmol, EOB). High regional brain uptake (% ID/g) was observed in cerebellum, striatum, cortex and hippocampus respectively as 6.34±0.05, 6.63±0.56, 6.69±0.72 and 5.41±0.30 at 5 min; tracer levels remained high between 5 and 30 min and washed out between 30 and 60 min, indicating [¹¹C]14e has a suitable clearance from brain. Autoradiography studies revealed the distribution of [¹¹C]14e is consistent with the distribution of σ₁ receptors in brain. MicroPET studies matched the σ₁ distribution in brain; the tissue time-activity curves indicated [¹¹C]14e has a suitable clearance rate from monkey brain (FIG. 4).

These results indicate that [¹¹C]14e is useful as a brain imaging radiotracer. The results show that [¹¹C]14e has high specific binding with receptor and suitable clearance rates from the brain.

All references cited are incorporated by reference, each in its entirety.

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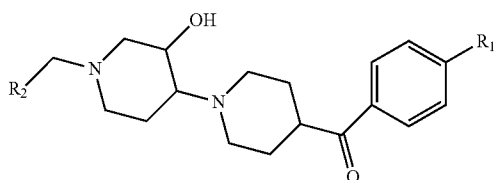
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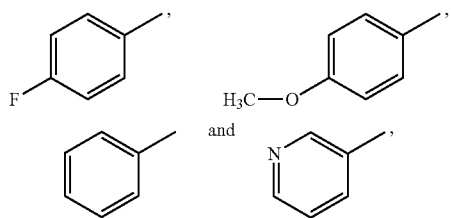
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What is claimed is:

1. A compound of structure



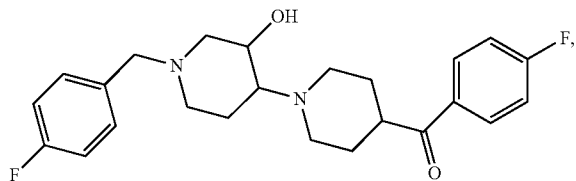
or a salt thereof, wherein R_1 is selected from the group consisting of F and OCH_3 , and R_2 is selected from the group consisting of



and wherein the compound exhibits selectivity for σ_1 in comparison to vesicular acetylcholine transporter (VAChT), and wherein the compound comprises a positron-emitting nuclide.

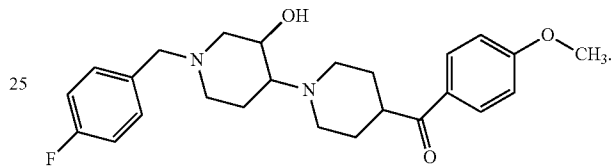
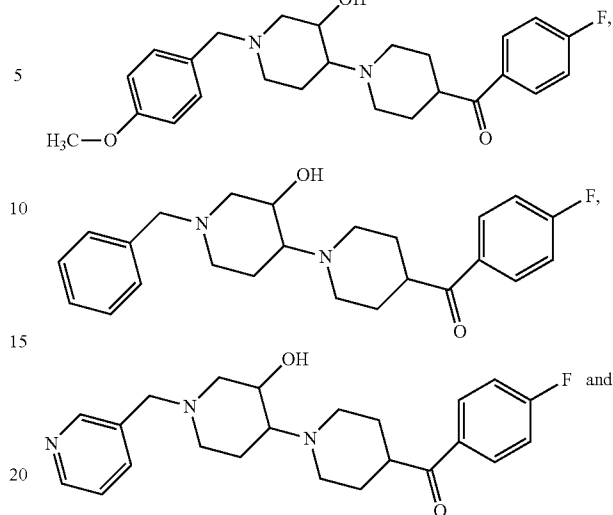
2. A compound or salt thereof in accordance with claim 1, wherein R_1 is F.

3. A compound or salt thereof in accordance with claim 1, wherein the compound is selected from the group consisting of



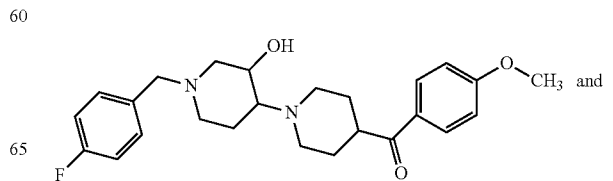
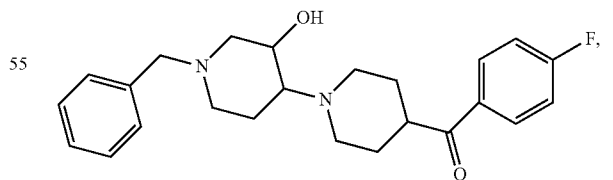
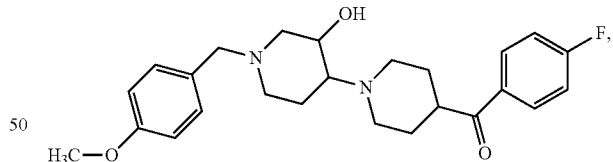
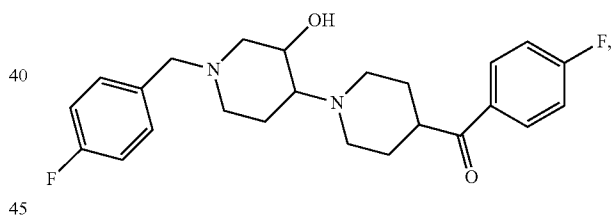
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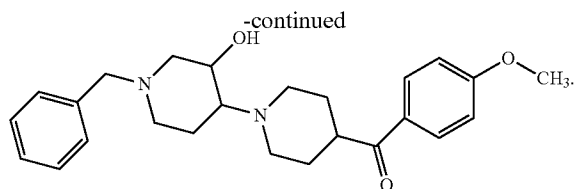


4. A compound or salt thereof in accordance with claim 3, wherein an F is an ^{18}F .

5. A compound or salt thereof in accordance with claim 1, wherein the compound is selected from the group consisting of

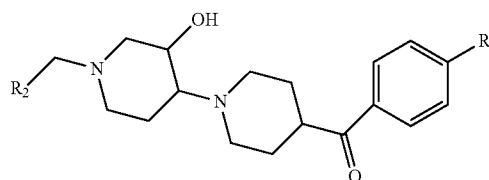


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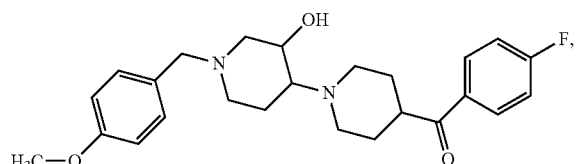


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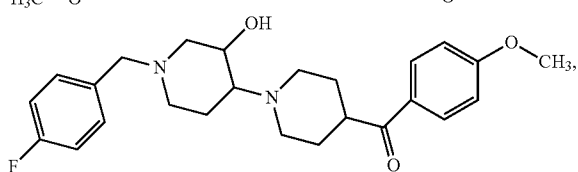
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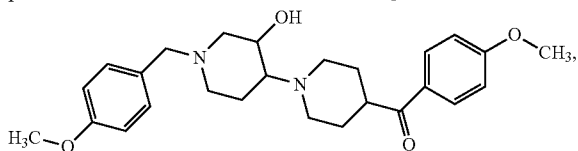
6. A compound or salt thereof in accordance with claim 1, wherein the compound is selected from the group consisting of



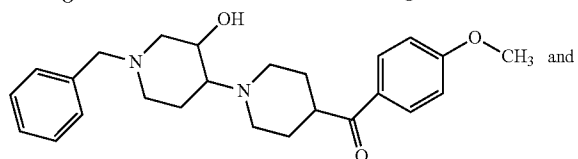
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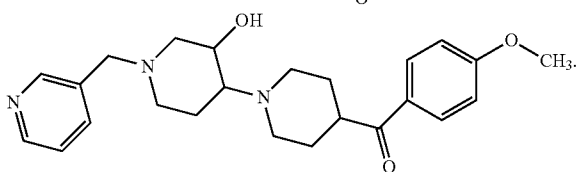
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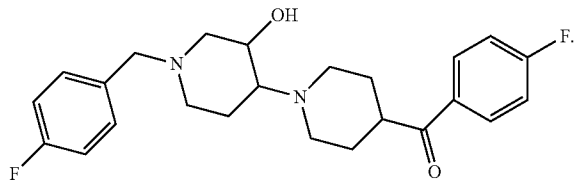
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7. A compound or salt thereof in accordance with claim 6, wherein a methoxy is an ^{11}C methoxy.

8. A compound or salt thereof in accordance with claim 1, wherein the compound is



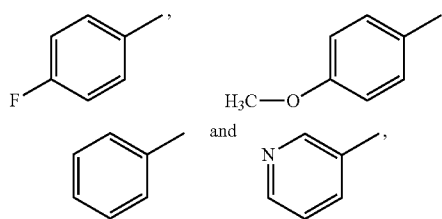
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9. A compound or salt thereof in accordance with claim 1, wherein at least 1 carbon is a ^{11}C .

10. A method of imaging distribution of sigma-1 receptors in a subject, comprising:

administering to a subject a radiolabeled compound of structure

or a salt thereof, wherein R_1 is selected from the group consisting of F and OCH_3 , and R_2 is selected from the group consisting of



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and wherein the compound exhibits selectivity for σ_1 in comparison to vesicular acetylcholine transporter (VACHT), and wherein the compound comprises a positron-emitting nuclide; and

subjecting the subject to positron emission tomography (PET) scanning.

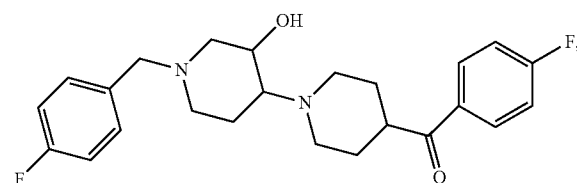
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11. A method in accordance with claim 10, wherein R_1 is an ^{18}F .

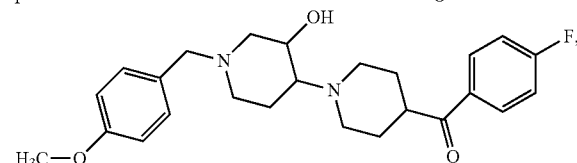
12. A method in accordance with claim 10, wherein R_1 is an ^{11}C .

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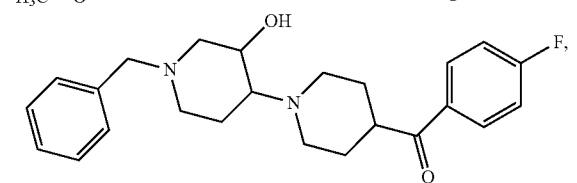
13. A method in accordance with claim 10, wherein the compound is selected from the group consisting of



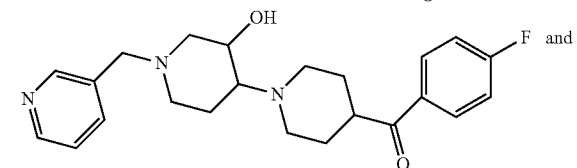
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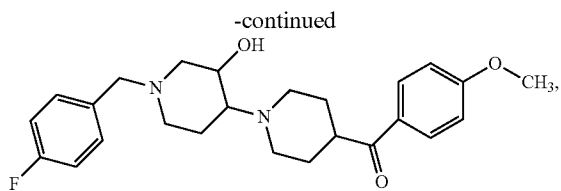
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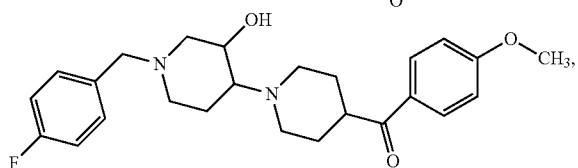
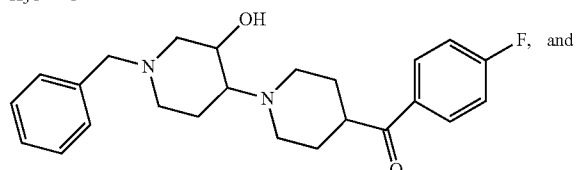
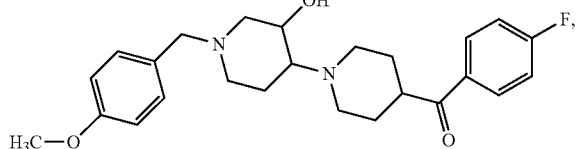
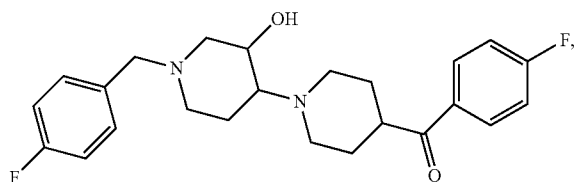
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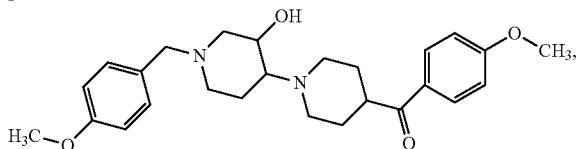
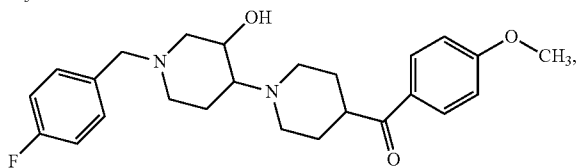
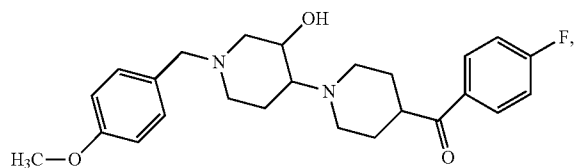
wherein an F is an ^{18}F .

14. A method in accordance with claim 10, wherein the compound is selected from the group consisting of

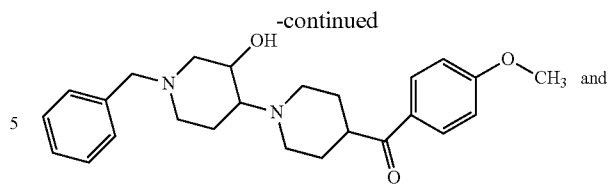


wherein an F is an ^{18}F .

15. A method in accordance with claim 10, wherein the compound is selected from the group consisting of

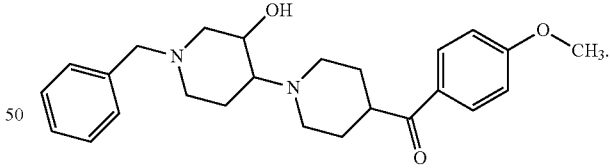
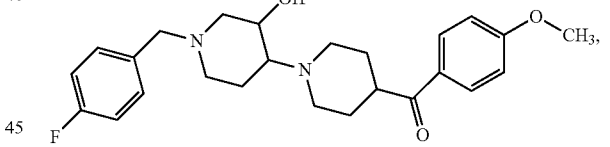
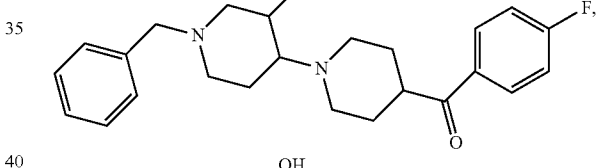
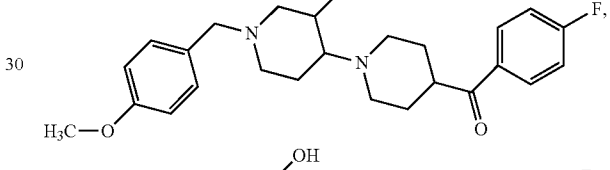
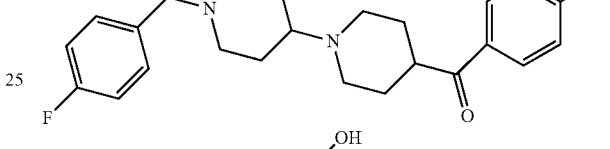
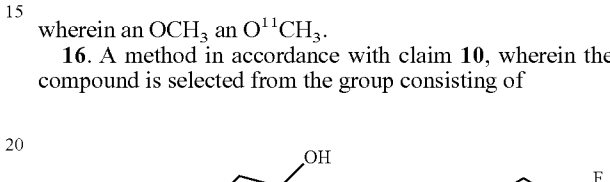


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wherein an OCH_3 an O^{11}CH_3 .

16. A method in accordance with claim 10, wherein the compound is selected from the group consisting of



17. A method in accordance with claim 10, wherein the compound is

