

**Table S1: CNS Target Binding Assay**

Receptor Assay	Origin	Ligand	Concentration	Non Specific	Incubation
Adenosine 1 (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]DPCPX	1 nM	DPCPX (1 μM)	60 min./22°C
Adenosine 2A (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]CGS 21680	6 nM	NECA (10 μM)	120 min./22°C
Adenosine 3 (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>125</sup> I]AB-MECA	0.15 nM	IB-MECA (1 μM)	120 min./22°C
Adrenergic α <sub>1</sub> (non-selective) (antagonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]prazosin	0.25 nM	prazosin (0.5 μM)	60 min./22°C
Adrenergic α <sub>2</sub> (non-selective) (antagonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]RX 821002	0.5 nM	(-)epinephrine (100 μM)	60 min./22°C
Adrenergic β <sub>1</sub> (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H](-)CGP 12177	0.15 nM	alprenolol (50 μM)	60 min./22°C
Adrenergic β <sub>2</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H](-)CGP 12177	0.2 nM	alprenolol (50 μM)	120 min./22°C
Angiotensine -1 (antagonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>125</sup> I][Sar <sup>1</sup> ,Ile <sup>8</sup> ]-AT-II	0.05 nM	angiotensin-II (10 μM)	120 min./37°C
Angiotensine-2 (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]CGP 42112A	0.04 nM	angiotensin-II (1 μM)	180 min./37°C
Benzodiazepine (central) (agonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]flunitrazepam	0.4 nM	diazepam (3 μM)	60 min./4°C
Benzodiazepine (peripheral) (antagonist radioligand)	rat heart	[ <sup>3</sup> H]PK 11195	0.2 nM	PK 11195 (10 μM)	15 min./22°C
Bombesin (non-selective) (agonist radioligand)	rat cerebral cortex	[ <sup>125</sup> I][Tyr <sup>4</sup> ]bombesin	0.01 nM	bombesin (1 μM)	60 min./22°C
Bradykinin (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]bradykinin	0.2 nM	bradykinin (1 μM)	60 min./22°C
Calcitonin gene-related peptide (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]hCGRPα	0.03 nM	hCGRPα (1 μM)	90 min./22°C
Cannabinoid (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]CP 55940	0.5 nM	WIN 55212-2 (10 μM)	120 min./37°C
Cholecystokinin-A (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]CCK-8s	0.08 nM	CCK-8s (1 μM)	60 min./22°C
Cholecystokinin-B (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]CCK-8s	0.08 nM	CCK-8s (1 μM)	60 min./22°C
Dopamine-D <sub>1</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]SCH 23390	0.3 nM	SCH 23390 (1 μM)	60 min./22°C
Dopamine-D <sub>2S</sub> (antagonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]spiperone	0.3 nM	(+)-butaclamol (10 μM)	60 min./22°C
Dopamine-D <sub>3</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]spiperone	0.3 nM	(+)-butaclamol (10 μM)	60 min./22°C

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Dopamine-D <sub>4.4</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]spiperone	0.3 nM	(+)-butaclamol (10 μM)	60 min./22°C
Dopamine-D <sub>5</sub> (antagonist radioligand)	human recombinant (GH4 cells)	[ <sup>3</sup> H]SCH 23390	0.3 nM	SCH 23390 (10 μM)	60 min./22°C
Endothelin -ET <sub>A</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]endothelin-1	0.03 nM	endothelin-1 (0.1 μM)	120 min./37°C
Endothelin -ET <sub>B</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]endothelin-1	0.03 nM	endothelin-1 (0.1 μM)	120 min./37°C
GABA (non-selective) (agonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]GABA	10 nM	GABA (100 μM)	60 min./22°C
Galanin <sub>1</sub> (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>125</sup> I]galanin	0.1 nM	galanin (1 μM)	60 min./22°C
Galanin <sub>2</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]galanin	0.05 nM	galanin (1 μM)	120 min./22°C
Growth Factors - PDGF (agonist radioligand)	Balb/c 3T3 cells	[ <sup>125</sup> I]PDGF BB	0.03 nM	PDGF BB (10 nM)	180 min./4°C
Chemokines (IL-8B) (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>125</sup> I]IL-8	0.025 nM	IL-8 (30 nM)	60 min./22°C
TNF-α (agonist radioligand)	U-937 cells	[ <sup>125</sup> I]TNF-α	0.1 nM	TNF-α (10 nM)	120 min./4°C
Chemokines - CCR1 (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>125</sup> I]MIP-1α	0.02 nM	MIP-1α (100 nM)	120 min./22°C
Histamine-H <sub>1</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]pyrilamine	3 nM	pyrilamine (1 μM)	60 min./22°C
Histamine-H <sub>2</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]APT	0.075 nM	tiotidine (100 μM)	120 min./22°C
Melanocortin - MC <sub>4</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]NDP-α-MSH	0.05 nM	NDP-α-MSH (1 μM)	120 min./37°C
Melatonin (ML <sub>1A</sub> ) (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]2- iodomelatonin	0.025 nM	melatonin (1 μM)	60 min./22°C
Muscarinic-M <sub>1</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]pirenzepine	2 nM	atropine (1 μM)	60 min./22°C
Muscarinic-M <sub>2</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]AF-DX 384	2 nM	atropine (1 μM)	60 min./22°C
Muscarinic-M <sub>3</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]4-DAMP	0.2 nM	atropine (1 μM)	60 min./22°C
Muscarinic-M <sub>4</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]4-DAMP	0.2 nM	atropine (1 μM)	60 min./22°C
Muscarinic-M <sub>5</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]4-DAMP	0.3 nM	atropine (1 μM)	60 min./22°C
Neurokinin-NK <sub>1</sub> (agonist radioligand)	U-373MG cells (endogenous)	[ <sup>125</sup> I]BH-SP	0.15 nM	[Sar <sup>9</sup> ,Met(O <sub>2</sub> ) <sup>11</sup> ]- SP (1 μM)	60 min./22°C
Neurokinin-NK <sub>2</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]NKA	0.1 nM	[Nleu <sup>10</sup> ]- NKA (4-10) (10 μM)	60 min./22°C

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Neurokinin-NK <sub>3</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]SR 142801	0.4 nM	SB 222200 (10 μM)	120 min./22°C
Neuropeptide Y-Y <sub>1</sub> (agonist radioligand)	SK-N-MC cells (endogenous)	[ <sup>125</sup> I]peptide YY	0.025 nM	NPY (1 μM)	120 min./37°C
Neuropeptide Y-Y <sub>2</sub> (agonist radioligand)	KAN-TS cells	[ <sup>125</sup> I]peptide YY	0.015 nM	NPY (1 μM)	60 min./37°C
Neurotensin (NT <sub>1</sub> ) (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]Tyr <sup>3</sup> - neurotensin	0.05 nM	neurotensin (1 μM)	60 min./4°C
δ <sub>2</sub> Opioid (DOP) (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]DADLE	0.5 nM	naltrexone (10 μM)	120 min./22°C
κ Opioid (KOP) (agonist radioligand)	rat recombinant (CHO cells)	[ <sup>3</sup> H]U 69593	1 nM	naloxone (10 μM)	60 min./22°C
μ Opioid (MOP) (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]DAMGO	0.5 nM	naloxone (10 μM)	120 min./22°C
Opioid-like (ORL1) (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]nociceptin	0.2 nM	nociceptin (1 μM)	60 min./22°C
Vasoactive intestinal peptide PAC <sub>1</sub> (PACAP) (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]PACAP <sub>1-27</sub>	0.015 nM	PACAP <sub>1-27</sub> (100 nM)	120 min./22°C
Non-steroid nuclear receptor PPAR <sub>γ</sub> (agonist radioligand)	human recombinant ( <i>E. coli</i> )	[ <sup>3</sup> H]rosiglitazone	5 nM	rosiglitazone (10 μM)	120 min./4°C
Glutamate - PCP (antagonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]TCP	10 nM	MK 801 (10 μM)	120 min./37°C
Prostanoid - EP <sub>4</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]PGE <sub>2</sub>	1 nM	PGE <sub>2</sub> (10 μM)	120 min./22°C
Prostanoid - TP (TXA <sub>2</sub> /PGH <sub>2</sub> ) (antagonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]SQ 29548	5 nM	U 44069 (10 μM)	60 min./22°C
Prostanoid - IP (PGI <sub>2</sub> ) (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]iloprost	10 nM	iloprost (10 μM)	60 min./22°C
Purinergic-P2X (agonist radioligand)	rat urinary bladder	[ <sup>3</sup> H]α,β-MeATP	3 nM	α,β-MeATP (10 μM)	120 min./4°C
Purinergic-P2Y (agonist radioligand)	rat cerebral cortex	[ <sup>35</sup> S]dATPαS	10 nM	dATPαS (10 μM)	60 min./22°C
Serotonin 5-HT <sub>1A</sub> (agonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]8-OH-DPAT	0.3 nM	8-OH-DPAT (10 μM)	60 min./22°C
Serotonin 5-HT <sub>1B</sub> (antagonist radioligand)	rat cerebral cortex	[ <sup>125</sup> I]CYP (+ 30 μM (-)propranolol)	0.1 nM	serotonin (10 μM)	120 min./37°C
Serotonin 5-HT <sub>2A</sub> (antagonist radioligand)	human recombinant (HEK-293 cells)	[ <sup>3</sup> H]ketanserin	0.5 nM	ketanserin (1 μM)	60 min./22°C
Serotonin 5-HT <sub>2B</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I](±)DOI	0.2 nM	(±)DOI (1 μM)	60 min./22°C
Serotonin 5-HT <sub>2C</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]mesulergine	1 nM	RS 102221 (10 μM)	60 min./37°C
Serotonin 5-HT <sub>3</sub> (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]BRL 43694	0.5 nM	MDL 72222 (10 μM)	120 min./22°C

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Serotonin 5-HT <sub>5A</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]LSD	1 nM	serotonin (100 µM)	60 min./37°C
Serotonin 5-HT <sub>6</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]LSD	2 nM	serotonin (100 µM)	120 min./37°C
Serotonin 5-HT <sub>7</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]LSD	4 nM	serotonin (10 µM)	120 min./22°C
σ Sigma (non-selective) (agonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]DTG	8 nM	haloperidol (10 µM)	120 min./22°C
Somatostatin (non-selective) (agonist radioligand)	AtT-20 cells	[ <sup>125</sup> I] <sup>11</sup> -Tyr <sup>11</sup> -somatostatin-14	0.05 nM	somatostatin-14 (300 nM)	60 min./37°C
Steroid nuclear receptor GR (agonist radioligand)	IM-9 cells (cytosol)	[ <sup>3</sup> H]dexamethasone	1.5 nM	triamcinolone (10 µM)	6 h./4°C
Vasoactive intestinal peptide (VIP <sub>1</sub> ) (agonist radioligand)	human recombinant (CHO cells)	[ <sup>125</sup> I]VIP	0.04 nM	VIP (1 µM)	60 min./22°C
Vasopressin - V <sub>1a</sub> (agonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]AVP	0.3 nM	AVP (1 µM)	60 min./22°C
Ca <sup>2+</sup> channel (L, verapamil site) (phenylalkylamine) (antagonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H](–)D 888	3 nM	D 600 (10 µM)	120 min./22°C
Potassium- K <sub>V</sub> channel (antagonist radioligand)	rat cerebral cortex	[ <sup>125</sup> I]α-dendrotoxin	0.01 nM	α-dendrotoxin (50 nM)	60 min./22°C
Potassium- SK <sub>Ca</sub> channel (antagonist radioligand)	rat cerebral cortex	[ <sup>125</sup> I]apamin	0.007 nM	apamin (100 nM)	60 min./4°C
Na <sup>+</sup> channel (site 2) (antagonist radioligand)	rat cerebral cortex	[ <sup>3</sup> H]batrachotoxinin	10 nM	veratridine (300 µM)	60 min./22°C
Cl <sup>-</sup> channel (GABA-gated) (antagonist radioligand)	rat cerebral cortex	[ <sup>35</sup> S]TBPS	3 nM	picrotoxinin (20 µM)	120 min./22°C
norepinephrine transporter (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]nisoxetine	1 nM	desipramine (1 µM)	120 min./4°C
dopamine transporter (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]BTCP	4 nM	BTCP (10 µM)	120 min./4°C
Serotonin transporter (antagonist radioligand)	human recombinant (CHO cells)	[ <sup>3</sup> H]imipramine	2 nM	imipramine (10 µM)	60 min./22°C