

Serotonin Transporter

5-HTT; SERT; SLC6A4

Serotonin Transporters (SERTs) are integral membrane proteins that transport serotonin from synaptic spaces into presynaptic neurons. SERTs function by reuptaking serotonin in the synaptic cleft, effectively terminating the function of serotonin and halting neuronal transmission. Serotonin reuptake is a critical process to prevent overstimulation of nerves.

Serotonin transporter (SERT) regulates extracellular levels of serotonin (5-hydroxytryptamine, 5HT) in the brain by transporting 5HT into neurons and glial cells. The human SERT (hSERT) is the primary target for drugs used in the treatment of emotional disorders, including depression. hSERT belongs to the solute carrier 6 family that includes a bacterial leucine transporter (LeuT), for which a high resolution crystal structure has become available.

Serotonin Transporter Inhibitors

Amitifadine hydrochloride

(DOV-21947 hydrochloride; EB-1010 hydrochloride) Cat. No.: HY-18332A

Amitifadine hydrochloride is a serotonin-norepinephrine-dopamine reuptake inhibitor (SNDRI), with IC $_{50}$ s of 12, 23, 96 nM for serotonin, norepinephrine and dopamine in HEK 293 cells , respectively.

CI

H-CI

Purity: 99.92% Clinical Data: Phase 3

Size: 10 mM × 1 mL, 5 mg, 10 mg, 50 mg

Azaphen (Azafen; Pipofezin hydrochloride; Pipofezine hydrochloride)

Pipofezine(Azafen or Azaphen) is a potent inhibitor of the reuptake of serotonin.

N H-CI

Cat. No.: HY-A0022

Purity: >98%
Clinical Data: Launched
Size: 100 mg, 500 mg

Azaphen dihydrochloride monohydrate (Azafen dihydrochloride

monohydrate; Pipofezin dihydrochloride monohydrate; ...) Cat. No.: HY-A0022A

Pipofezine(Azafen or Azaphen) is a potent inhibitor of the reuptake of serotonin.

$$\begin{array}{c|c} O & N_{c} N \\ \hline N & N_{c} N \\ H-CI & H_{2}O \end{array}$$

Purity: 99.60% Clinical Data: Launched Size: 100 mg, 500 mg

Centanafadine

(EB-1020) Cat. No.: HY-16736

Centanafadine is dual **norepinephrine** (NE)/dopamine (DA) transporter inhibitor, also inhibits serotonin transporter, with IC_{50} s of 6 nM, 38 nM and 83 nM for human NE, DA and serotonin transporter , respectively.

NH

Purity: >98%

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Centanafadine hydrochloride

(EB-1020 hydrochloride) Cat. No.: HY-16736A

Centanafadine (hydrochloride) is dual norepinephrine (NE)/dopamine (DA) transporter inhibitor, also inhibits serotonin transporter, with IC_{so} s of 6 nM, 38 nM and 83 nM for human NE, DA and serotonin transporter , respectively.

NH

HCI

Purity: 99.93%

Clinical Data: No Development Reported

Size: $10 \text{ mM} \times 1 \text{ mL}$, 5 mg, 10 mg, 50 mg, 100 mg

Cinchonidine

(α-Quinidine) Cat. No.: HY-N0173

Cinchonidine (α -Quinidine) is a cinchona alkaloid found in Cinchona officinalis and Gongronema latifolium. A building block used in asymmetric synthesis in organic chemistry.



Purity: >98.0%

Clinical Data: No Development Reported
Size: 10 mM × 1 mL, 100 mg

Citalopram hydrobromide

((±)-Citalopram hydrobromide; Lu 10-171) Cat. No.: HY-B1287

Citalopram hydrobromide is an antidepressant drug of the selective serotonin reuptake inhibitor (SSRI) class. It has US FDA approval to treat major depression.

N H-Br

Cat. No.: HY-B0457S

Purity: >99.0%
Clinical Data: Launched

Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg

Clomipramine D3

Cat. No.: HY-B0457AS

Clomipramine D3 is the deuterium labeled

Clomipramine. Clomipramine is a serotonin transporter (SERT), norepinephrine transporter (NET) dopamine transporter (DAT) blocker with $\rm K_i$ of 0.14, 54 and 3 nM, respectively.



Purity: >98%

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Clomipramine hydrochloride

Cat. No.: HY-B0457

Clomipramine D3 hydrochloride is a deuterium
labeled Clomipramine hydrochloride. Clomipramine
hydrochloride is a serotonin transporter (SERT),
norepinephrine transporter (NET) dopamine
transporter (DAT) blocker with K, of 0.14, 54 and 3

nM, respectively.

Purity: >98%

Clinical Data: No Development Reported

Clomipramine D3 hydrochloride

Size: 1 mg, 5 mg

Clomipramine hydrochloride is a **serotonin** transporter (SERT), norepinephrine transporter (NET) and **dopamine** transporter (DAT) blocker with \mathbf{K}_i of 0.14, 54 and 3 nM, respectively.



Purity: 99.72% Clinical Data: Launched

Size: 10 mM × 1 mL, 100 mg, 500 mg

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Dapoxetine hydrochloride

(LY-210448 hydrochloride)

Dapoxetine hydrochloride is a short-acting novel selective serotonin reuptake inhibitor(SSRI). Target: SSRI Dapoxetine hydrochloride is a short-acting novel selective serotonin reuptake inhibitor marketed for the treatment of premature ejaculation in men.

HCI

Cat. No.: HY-B0304A

Purity: 99 93% Clinical Data: Launched

10 mM × 1 mL, 50 mg, 100 mg Size:

Dasotraline

(SEP 225289)

Dasotraline is a triple reuptake inhibitor that blocks dopamine, norepinephrine, and serotonin transporters with IC₅₀ values of 4, 6, and 11 nM, respectively.

Cat. No.: HY-12850

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Dasotraline hydrochloride

(SEP-225289 hydrochloride)

Dasotraline hydrochloride (SEP-225289 hydrochloride) is a triple reuptake inhibitor that blocks dopamine, norepinephrine, and serotonin transporters with IC₅₀ values of 4, 6, and 11 nM, respectively.

Cat. No.: HY-12850A

Purity: 99 55% Clinical Data: Phase 3

10 mM × 1 mL, 1 mg, 5 mg, 10 mg, 25 mg, 50 mg

Desipramine hydrochloride

Desipramine hydrochloride is an inhibitor of norepinephrine transporter (NET), 5-HT

transporter (SERT) and dopamine transporter (DAT) with K_.s of 4, 61 and 78,720 nM, respectively.

Purity: 99.68% Clinical Data: Launched

10 mM × 1 mL, 100 mg, 500 mg

Cat. No.: HY-B1272

Desvenlafaxine

(O-Desmethylvenlafaxine)

Desvenlafaxine is a serotonin (5-HT) and norepinephrine (NE) reuptake inhibitor with Ki of 40.2 nM and 558.4 nM, respectively.

Cat. No.: HY-B0602

Purity: > 98.0% Clinical Data: Launched

Size: 10 mM × 1 mL, 100 mg

Desvenlafaxine succinate hydrate

(O-Desmethylvenlafaxine succinate hydrate)

Desvenlafaxine succinate hydrate is an antidepressant of the serotonin-norepinephrine reuptake inhibitor (SNRI).

Cat. No.: HY-B0602A

99.76% Purity: Clinical Data: Launched

Size 10 mM × 1 mL, 100 mg

DOV-216,303 Free Base

Cat. No.: HY-18332C

DOV-216,303 (Free Base) is a potent triple serotonin, norepinephrine, and dopamine reuptake inhibitor, with IC_{so} values of 14 nM, 20 nM and 78 nM for hSERT, hNET and hDAT, respectively.

98.47% Purity:

Clinical Data: No Development Reported 10 mM × 1 mL, 5 mg Size:

DSP-1053

Cat. No.: HY-111419

DSP-1053 is a potent Serotonin Transporter (SERT) (K = 1.02 nM) inhibitor with partial 5-HT1A receptor (K_i=5.05 nM) agonistic activity.

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

LY248686 D3 hydrochloride)

Duloxetine

Purity:

((S)-Duloxetine; LY248686)

Duloxetine is a serotonin-norepinephrine reuptake inhibitor with a K, of 4.6 nM, used for treatment of major depressive disorder and generalized anxiety disorder (GAD).



Cat. No.: HY-B0161

Duloxetine D3 hydrochloride ((S)-Duloxetine D3 hydrochloride) is a deuterium labeled Duloxetine hydrochloride.

Duloxetine D3 hydrochloride ((S)-Duloxetine D3 hydrochloride;

Cat. No.: HY-B0161AS

Purity: >98%

Clinical Data: No Development Reported

1 mg, 5 mg

Clinical Data: Launched Size: 1 mg, 5 mg

>98%

Duloxetine hydrochloride

((S)-Duloxetine hydrochloride; LY-248686 hydrochloride) Cat. No.: HY-B0161A

Duloxetine hydrochloride ((S)-Duloxetine hydrochloride) is a serotonin-norepinephrine reuptake inhibitor (SNRI) with a K, of 4.6 nM, used for treatment of major depressive disorder and generalized anxiety disorder (GAD).

Purity: 99 92% Clinical Data: Launched

Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg

Purity: 99 95%

Eplivanserin mixture

(SR-46349 (mixture))

Clinical Data: No Development Reported 5 mg, 10 mg, 50 mg, 100 mg

Eplivanserin mixture is a selective serotonin

antagonist, extracted from patent WO 2005/002578

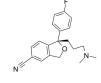
reuptake inhibitor and a 5-HT₂₄ receptor

Cat. No.: HY-10792A

Escitalopram

((S)-Citalopram; S-(+)-Citalopram) Cat. No.: HY-14258

Escitalopram is a selective serotonin reuptake inhibitor (SSRI) with Ki of 0.89 nM. Target: SSRIs Escitalopram, the S-enantiomer of citalopram, belongs to a class of antidepressant agents known as selective serotonin-reuptake inhibitors (SSRIs).



Purity: >98% Clinical Data: Launched Size: 1 mg, 5 mg

Escitalopram oxalate ((S)-(+)Citalopram oxalate)

Escitalopram oxalate ((S)-(+)Citalopram oxalate) is a selective serotonin reuptake inhibitor (SSRI) with Ki of 0.89 nM.

Cat. No.: HY-14258A

Purity: 99.98% Clinical Data: Launched

10 mM × 1 mL, 10 mg, 50 mg, 100 mg

Fluoxetine

(LY-110140 (free base)) Cat. No.: HY-B0102

Fluoxetine (LY-110140 free base) is a selective serotonin reuptake inhibitor (SSRI) class used for antidepressant research.

Purity: >98% Clinical Data: Launched Size: 1 mg, 5 mg

Fluoxetine hydrochloride

(LY 110140) Cat. No.: HY-B0102A

Fluoxetine hydrochloride (LY 110140) is an antidepressant and a selective serotonin reuptake inhibitor.



Purity: 99.84% Clinical Data: Launched

 $10~\text{mM}\times1~\text{mL},\,50~\text{mg},\,100~\text{mg},\,500~\text{mg}$

Fluvoxamine

(DU-23000) Cat. No.: HY-B0103

Fluvoxamine (DU-23000) is an antidepressant which functions pharmacologically as a selective serotonin reuptake inhibitor.

>98% Purity: Clinical Data: Launched Size: 1 mg, 5 mg

Fluvoxamine maleate

(DU-23000 maleate) Cat. No.: HY-B0103A

Fluvoxamine maleate (DU-23000 maleate) is an antidepressant which functions pharmacologically as a selective serotonin reuptake inhibitor.



Purity: Clinical Data: Launched

Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg

99.85%

Imipramine hydrochloride

Cat. No.: HY-B1490

Imipramine hydrochloride inhibits serotonin transporter with an IC_{50} value of 32 nM in vitro.

Purity: 99.93% Clinical Data: Launched

Size: 10 mM × 1 mL, 100 mg

Milnacipran

Milnacipran is a serotonin-norepinephrine reuptake inhibitor (SNRI) used in the clinical treatment of fibromyalgia.



Cat. No.: HY-B0168

Purity: >98% Clinical Data: Launched 1 mg, 5 mg

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Milnacipran (1S-cis) hydrochloride

(Levomilnacipran hydrochloride; F-2695 hydrochloride)

Milnacipran (1S-cis) hydrochloride is a serotonin-norepinephrine reuptake inhibitor (SNRI), used in the clinical treatment of fibromyalgia.

Cat. No.: HY-B0168B

Purity: 99 94% Clinical Data: Launched

Size: 10 mM × 1 mL, 5 mg, 10 mg, 50 mg

Milnacipran hydrochloride

Cat. No.: HY-B0168A

Milnacipran hydrochloride is a serotonin-norepinephrine reuptake inhibitor (SNRI) used in the clinical treatment of fibromyalgia.



Purity: 99 87% Clinical Data: Launched

Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg, 200 mg

Nitroxazepine

(CIBA 2330Go) Cat. No.: HY-101684

Nitroxazepine is a tricyclic antidepressant (TCA) for the treatment of depression. Nitroxazepine acts as a serotonin-norepinephrine reuptake inhibitor.

Purity: >98%

Clinical Data: No Development Reported

1 mg, 5 mg

Paroxetine hydrochloride

(BRL29060 hydrochloride; BRL29060A)

Paroxetine hydrochloride is a potent selective serotonin-reuptake inhibitor, commonly prescribed as an antidepressant and has GRK2 inhibitory ability with IC_{50} of 14 μ M.

Cat. No.: HY-B0492

Purity: >99.0% Clinical Data: Launched

10 mM × 1 mL, 100 mg, 500 mg

Paroxetine hydrochloride hemihydrate (BRL29060 hydrochloride

hemihydrate; BRL29060A hemihydrate) Cat. No.: HY-B0492A

Paroxetine hydrochloride hemihydrate is a potent selective serotonin-reuptake inhibitor, commonly prescribed as an antidepressant and has GRK2 inhibitory ability with IC_{50} of $14\mu M$.

0.5H₂O

>98% Purity: Clinical Data: Launched Size: 1 mg, 5 mg

Sertraline hydrochloride

Cat. No.: HY-B0176A

Sertraline hydrochloride is an antidepressant of the selective serotonin reuptake inhibitor (SSRI) class. Target: SSRI Sertraline is an antidepressant of the selective serotonin reuptake inhibitor (SSRI) class. It was introduced to the market by Pfizer in 1991.

99.90% **Purity:** Clinical Data: Launched

 $10 \text{ mM} \times 1 \text{ mL}$, 5 mg, 10 mg, 50 mg



Sibutramine hydrochloride monohydrate

(BTS 54-524 hydrochloride monohydrate)

Sibutramine hydrochloride monohydrate is a novel 5-HT (serotonin) and noradrenaline reuptake inhibitor (SNRI). The IC_{so} for Sibutramine block of voltage-gated K+ channel (K_v)4.3 is 17.3 μM.

Cat. No.: HY-14470

HCI H₂O

Purity: Clinical Data: No Development Reported

>98%

Size: 1 mg, 5 mg

SPD-473 citrate

Cat. No.: HY-101612

SPD-473 citrate is a

serotonin/dopamine/norepinephrine reuptake inhibition

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Venlafaxine

(Wy 45030) Cat. No.: HY-B0196

Venlafaxine is an antidepressant of the serotonin-norepinephrine reuptake inhibitor (SNRI) class. Target: SNRI Venlafaxine is an antidepressant of the serotonin-norepinephrine reuptake inhibitor (SNRI) class.



Purity: >98% Clinical Data: Launched Size: 1 mg, 5 mg

Venlafaxine hydrochloride

(Wy 45030 hydrochloride)

Venlafaxine hydrochloride is an antidepressant of the serotonin-norepinephrine reuptake inhibitor (SNRI) class. Target: SNRI Venlafaxine is an antidepressant of the serotonin-norepinephrine reuptake inhibitor (SNRI) class.

99.87% Clinical Data: Launched

10 mM × 1 mL, 10 mg, 50 mg

Cat. No.: HY-B0196A

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Vilazodone

(EMD 68843; SB659746A) Cat. No.: HY-14262

Vilazodone (EMD 68843; SB 659746A) is a combined serotonin specific reuptake inhibitor (SSRI) and 5-HT1A receptor partial agonist currently under clinical evaluation for the treatment of major depression.

Purity: 99 91% Clinical Data: Launched

10 mM × 1 mL, 10 mg, 50 mg Size:

Cat. No.: HY-14261

Vilazodone Hydrochloride (EMD 68843 Hydrochloride)

Purity: 99 95% Clinical Data: Launched

10 mM × 1 mL, 10 mg, 50 mg

Vortioxetine is a inhibitor of 5-HT_{1A}, 5-HT_{1B}, 5-HT₃₄, 5-HT₇ receptor and SERT, with K₁ values of 15 nM, 33 nM, 3.7 nM, 19 nM and 1.6 nM, respectively.

Purity: 98.81% Clinical Data: Launched

Vilazodone Hydrochloride

(EMD 68843 Hydrochloride; SB659746A Hydrochloride)

is a serotonin transporter (SER) inhibitor and 5-HT₁₄ receptor partial agonist.

Vortioxetine D8

(Lu AA 21004 D8) Cat. No.: HY-15414S

Vortioxetine D8 is a deuterium labeled Vortioxetine. Vortioxetine is an inhibitor of $\mathbf{5\text{-}HT}_{\mathbf{1A'}}, \mathbf{5\text{-}HT}_{\mathbf{1B'}}, \mathbf{5\text{-}HT}_{\mathbf{3A'}}, \mathbf{5\text{-}HT}_{\mathbf{7}}$ receptor and **SERT**, with **K**₁ values of 15 nM, 33 nM, 3.7 nM, 19 nM and 1.6 nM, respectively.

Purity: >98%

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Wf-516

Wf-516 is an inhibitor of 5-HT reuptake, and an antagonist of 5-HT1A and 5-HT2A receptors, with K, of 5 nM and 40 nM for 5-HT1A receptor and 5-HT2A receptor in humans, respectively, and has potent antidepressant activity.

Cat. No.: HY-19417A

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Vilazodone D8

Vilazodone D8 is the a deuterium labeled vilazodone, which is a combined serotonin specific reuptake inhibitor (SSRI) and 5-HT1A receptor partial agonist.

Cat. No.: HY-14261S

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Vortioxetine

(Lu AA 21004) Cat. No.: HY-15414

10 mM × 1 mL, 5 mg, 10 mg, 50 mg

Vortioxetine hydrobromide

(Lu AA21004 hydrobromide)

Vortioxetine hydrobromide is a multimodal serotonergic agent, inhibits 5-HT_{1A}, 5-HT_{1B}, 5-HT_{3A}, 5-HT₇ receptor and SERT with K₁ values of 15 nM, 33 nM, 3.7 nM, 19 nM and 1.6 nM, respectively.



Cat. No.: HY-15414A

99.94% Purity: Clinical Data: Launched

Size: 10 mM \times 1 mL, 5 mg, 10 mg, 50 mg