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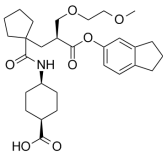
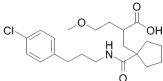
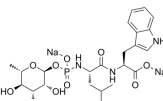
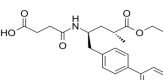
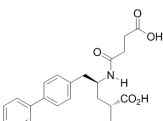
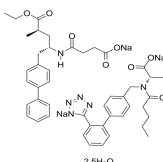
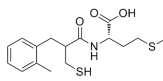
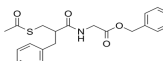
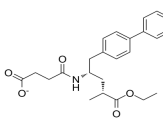
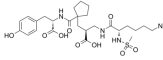
Neprilysin

Neutral endopeptidase; NEP; Cluster of differentiation 10; CD10

Neprilysin (NEP) is a type II membrane metalloendopeptidase composed of 750 residues with an active site containing a zinc-binding motif (HEXXH) at the extracellular carboxyl terminal domain. Neprilysin is capable of degrading the monomeric and the oligomeric forms of A β peptide. Neprilysin is the dominant A β peptide-degrading enzyme in the brain; Neprilysin becomes inactivated and down-regulated during both the early stages of Alzheimer's disease (AD) and aging.

Neprilysin is a neutral endopeptidase and its inhibition increases bioavailability of natriuretic peptides, bradykinin, and substance P, resulting in natriuretic, vasodilatory, and anti-proliferative effects.

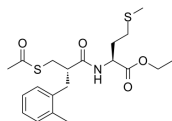
Neprilysin Inhibitors

<p>Candoxatril (UK 79300)</p> <p>Candoxatril is a neutral endopeptidase (NEP) inhibitor.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg</p>	<p>Cat. No.: HY-19649</p> 	<p>NEP-In-1</p> <p>NEP-IN-1 is a neutral endopeptidase (NEP) inhibitor with IC_{50} of 2 nM for dNEP.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg</p>	<p>Cat. No.: HY-U00294</p> 	<p>Phosphoramidon Disodium</p> <p>Phosphoramidon Disodium is a metalloprotease inhibitor. Phosphoramidon inhibits endothelin-converting enzyme (ECE), neutral endopeptidase (NEP), and angiotensin-converting enzyme (ACE) with IC_{50} values of 3.5, 0.034, and 78 μM, respectively.</p> <p>Purity: >98.0% Clinical Data: No Development Reported Size: 5 mg, 10 mg</p>	<p>Cat. No.: HY-N2021A</p> 	<p>Sacubitril (AHU-377)</p> <p>Sacubitril (AHU-377) is a potent NEP inhibitor with an IC_{50} of 5 nM. Sacubitril (AHU-377) is a component of the heart failure medicine LCZ696.</p> <p>Purity: 99.41% Clinical Data: Launched Size: 10 mM \times 1 mL, 5 mg, 10 mg, 50 mg, 100 mg</p>	<p>Cat. No.: HY-15407</p> 	<p>Sacubitrilat (LBQ-657)</p> <p>Sacubitrilat is an active neprilysin (NEP) inhibitor.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 10 mM \times 1 mL, 2 mg, 5 mg, 10 mg, 25 mg, 50 mg</p>	<p>Cat. No.: HY-17620</p> 	<p>LCZ696 (Sacubitril/Valsartan)</p> <p>LCZ696 (Sacubitril/Valsartan), comprised Valsartan (an ARB) and Sacubitril (AHU377) in 1:1 molar ratio, is a first-in-class, orally bioavailable, and dual-acting angiotensin receptor-neprilysin (ARN) inhibitor for hypertension and heart failure.</p> <p>Purity: 99.99% Clinical Data: Launched Size: 10 mM \times 1 mL, 5 mg, 10 mg, 50 mg</p>	<p>Cat. No.: HY-18204A</p> 	<p>NEP-IN-2</p> <p>NEP-IN-2 is an inhibitor of neutral endopeptidase, used in the research of proliferation in atherosclerosis, restenosis.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg</p>	<p>Cat. No.: HY-U00336</p> 	<p>Racecadotril (Acetorphan)</p> <p>Racecadotril (Acetorphan) is a neutral endopeptidase (NEP) inhibitor. Racecadotril and its active metabolite Thiorphane inhibits purified NEP activity from mouse brain with K_is of 4500 and 6.1 nM, respectively. Antidiarrheal agent.</p> <p>Purity: 98.85% Clinical Data: Launched Size: 10 mM \times 1 mL, 200 mg, 1 g</p>	<p>Cat. No.: HY-17399</p> 	<p>Sacubitril hemicalcium salt (AHU-377 hemicalcium salt)</p> <p>Sacubitril hemicalcium salt (AHU-377 hemicalcium salt) is a potent NEP inhibitor with an IC_{50} of 5 nM. Sacubitril hemicalcium salt is a component of the heart failure medicine LCZ696.</p> <p>Purity: 99.69% Clinical Data: Launched Size: 10 mM \times 1 mL, 5 mg, 10 mg, 50 mg, 100 mg, 500 mg, 1 g</p>	<p>Cat. No.: HY-15407A</p> 	<p>Sampatrilat (UK-81252)</p> <p>Sampatrilat (UK-81252) is a potent and orally active vasopeptidase inhibitor of ACE and neutral endopeptidase (NEP). Sampatrilat inhibits C-domain ACE ($K_i=13.8$ nM) 12.4-fold more potent than that for the N-domain ($K_i=171.9$ nM).</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg</p>	<p>Cat. No.: HY-123348</p> 
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SCH 42495

Cat. No.: HY-101682

SCH 42495 is an orally active neutral metalloendopeptidase (NEP) inhibitor with antihypertensive effect. SCH 42495 is the orally active ethylester prodrug of SCH 42354.

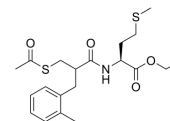


Purity: >98%
Clinical Data: No Development Reported
Size: 1 mg, 5 mg

SCH 42495 racemate

Cat. No.: HY-101682A

SCH 42495 racemate is the racemate of SCH 42495. SCH 42495 is an orally active neutral metalloendopeptidase (NEP) inhibitor with antihypertensive effect. SCH 42495 is the orally active ethylester prodrug of SCH 42354.



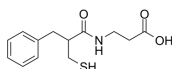
Purity: >98%
Clinical Data: No Development Reported
Size: 1 mg, 5 mg

SQ28603

(SQ28,603; Squibb 28603)

Cat. No.: HY-U00171

SQ28603 is a potent and selective inhibitor of neutral endopeptidase 3.4.24.11 (NEP), an enzyme that degrades atrial natriuretic peptide (ANP).

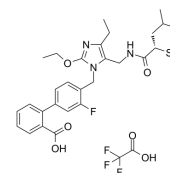


Purity: >98%
Clinical Data: No Development Reported
Size: 1 mg, 5 mg

TD-0212 TFA

Cat. No.: HY-114412A

TD-0212 TFA is an orally active dual pharmacology angiotensin II type 1 receptor (AT₁) antagonist and neprilysin (NEP) inhibitor, with a pK_i of 8.9 for AT₁ and a pIC₅₀ of 9.2 for NEP.



Purity: 98.44%
Clinical Data: No Development Reported
Size: 1 mg, 5 mg, 10 mg