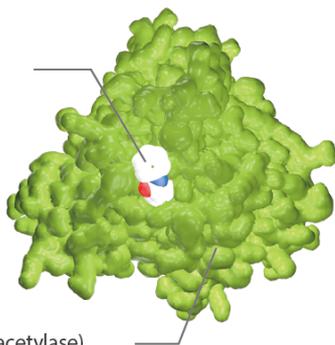


# Amino peptidase

HDAC Inhibitor:  
Vorinostat (SAHA)



HDAC (Histone deacetylase)

Amino peptidases catalyze the cleavage of amino acids from the amino terminus of protein (N-terminus) or peptide substrates. They are widely distributed throughout the animal and plant kingdoms and are found in many subcellular organelles, in cytoplasm, and as membrane components. Amino peptidases are used in essential cellular functions. Many, but not all, of these peptidases are zinc Metalloenzymes. Some amino peptidases are monomeric, and others are assemblies of relatively high mass (50 kDa) subunits. CDNA sequences are available for several amino peptidases and a crystal structure of the open state of human endoplasmic reticulum Amino peptidase 1 ERAP1 is presented here. Amino acid sequences

determined directly or deduced from cDNAs indicate some amino acid sequence homologies in organisms as diverse as Escherichia coli and mammals, particularly in catalytically important residues or in residues involved in metal ion binding. One important amino peptidase is a zinc-dependent enzyme produced and secreted by glands of the small intestine. It helps the enzymatic digestion of proteins. Additional digestive enzymes produced by these glands include dipeptidases, maltase, sucrase, lactase, and enterokinase.

## Aminopeptidase Inhibitors & Modulators

<p><b>Bestatin</b> (Ubenimex) <span style="float: right;">Cat. No.: HY-B0134</span></p>	<p><b>Bestatin hydrochloride</b> (Ubenimex hydrochloride) <span style="float: right;">Cat. No.: HY-B0134A</span></p>
<p><b>Bioactivity:</b> Bestatin is a natural, broad-spectrum, and competitive <b>aminopeptidase</b> inhibitor.</p> <p><b>Purity:</b> 99.96%</p> <p><b>Clinical Data:</b> Launched</p> <p><b>Size:</b> 10mM x 1mL in DMSO, 10 mg, 50 mg, 100 mg</p> 	<p><b>Bioactivity:</b> Bestatin hydrochloride is an inhibitor of <b>CD13 (Aminopeptidase N)/APN</b> and <b>leukotriene A4 hydrolase</b>, used for cancer treatment.</p> <p><b>Purity:</b> &gt;98%</p> <p><b>Clinical Data:</b> Launched</p> <p><b>Size:</b> 10 mg, 50 mg, 100 mg</p> 
<p><b>Bestatin trifluoroacetate</b> (Ubenimex trifluoroacetate) <span style="float: right;">Cat. No.: HY-B0134B</span></p>	<p><b>DG051</b> <span style="float: right;">Cat. No.: HY-10825</span></p>
<p><b>Bioactivity:</b> Bestatin trifluoroacetate is an inhibitor of <b>CD13 (Aminopeptidase N)/APN</b> and <b>leukotriene A4 hydrolase</b>, used for cancer treatment.</p> <p><b>Purity:</b> &gt;98%</p> <p><b>Clinical Data:</b> Launched</p> <p><b>Size:</b> 10 mg, 50 mg, 100 mg</p> 	<p><b>Bioactivity:</b> DG051 is a potent <b>leukotriene A4 hydrolase</b> inhibitor of leukotriene B4 biosynthesis in the enzyme assay with an <b>IC<sub>50</sub></b>=47 nM.</p> <p><b>Purity:</b> 99.65%</p> <p><b>Clinical Data:</b> No Development Reported</p> <p><b>Size:</b> 10mM x 1mL in DMSO, 1 mg, 5 mg</p> 
<p><b>HFI-142</b> <span style="float: right;">Cat. No.: HY-110259</span></p>	<p><b>NGR peptide Trifluoroacetate</b> <span style="float: right;">Cat. No.: HY-P1043A</span></p>
<p><b>Bioactivity:</b> HFI-142 is an insulin-regulated aminopeptidase (IRAP) inhibitor with a <b>K<sub>i</sub></b> of 2.01 μM <sup>[1]</sup>.</p> <p><b>Purity:</b> 99.0%</p> <p><b>Clinical Data:</b> No Development Reported</p> <p><b>Size:</b> 10mM x 1mL in DMSO, 5 mg</p> 	<p><b>Bioactivity:</b> NGR peptide Trifluoroacetate containing the asparagine-glycine-arginine (NGR) motif is recognized by <b>CD13/ aminopeptidase N ( APN) receptor</b> isoforms that are selectively overexpressed in tumor neovasculature.</p> <p><b>Purity:</b> 98.55%</p> <p><b>Clinical Data:</b> No Development Reported</p> <p><b>Size:</b> 10mM x 1mL in Water, 1 mg, 5 mg</p> 
<p><b>Tosedostat</b> (CHR-2797) <span style="float: right;">Cat. No.: HY-14807</span></p>	
<p><b>Bioactivity:</b> Tosedostat is an <b>aminopeptidase</b> inhibitor.</p> <p><b>Purity:</b> 99.65%</p> <p><b>Clinical Data:</b> Launched</p> <p><b>Size:</b> 10mM x 1mL in DMSO, 1 mg, 5 mg, 10 mg, 25 mg, 50 mg, 100 mg</p> 	