

# ACE1 Antibody

Rabbit mAb

Catalog # AP91542

## Product Information

<b>Application</b>	WB, IHC, FC
<b>Primary Accession</b>	<a href="#">P12821</a>
<b>Reactivity</b>	Human, Mouse
<b>Clonality</b>	Monoclonal
<b>Other Names</b>	Angiotensin-converting enzyme; somatic isoform precursor; CD143 antigen; DCP; DCP1; Dipeptidyl carboxypeptidase I; Kininase II;
<b>Isotype</b>	Rabbit IgG
<b>Host</b>	Rabbit
<b>Calculated MW</b>	149715

## Additional Information

<b>Dilution</b>	WB 1:500~1:1000 IHC 1:50~1:200 FC 1:30
<b>Purification</b>	Affinity-chromatography
<b>Immunogen</b>	A synthesized peptide derived from human ACE1
<b>Description</b>	Converts angiotensin I to angiotensin II by release of the terminal His-Leu, this results in an increase of the vasoconstrictor activity of angiotensin. Also able to inactivate bradykinin, a potent vasodilator. Has also a glycosidase activity which releases GPI-anchored proteins from the membrane by cleaving the mannose linkage in the GPI moiety.
<b>Storage Condition and Buffer</b>	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at +4°C short term. Store at -20°C long term. Avoid freeze / thaw cycle.

## Protein Information

<b>Name</b>	ACE {ECO:0000303   PubMed:2849100, ECO:0000312   HGNC:HGNC:2707}
<b>Function</b>	Dipeptidyl carboxypeptidase that removes dipeptides from the C-terminus of a variety of circulating hormones, such as angiotensin I, bradykinin or enkephalins, thereby playing a key role in the regulation of blood pressure, electrolyte homeostasis or synaptic plasticity (PubMed: <a href="#">15615692</a> , PubMed: <a href="#">20826823</a> , PubMed: <a href="#">2558109</a> , PubMed: <a href="#">4322742</a> , PubMed: <a href="#">7523412</a> , PubMed: <a href="#">7683654</a> ). Composed of two similar catalytic domains, each possessing a functional active site, with different selectivity for substrates (PubMed: <a href="#">10913258</a> , PubMed: <a href="#">1320019</a> , PubMed: <a href="#">1851160</a> , PubMed: <a href="#">19773553</a> , PubMed: <a href="#">7683654</a> , PubMed: <a href="#">7876104</a> ). Plays a major role in the angiotensin-renin system that regulates blood pressure and sodium retention by the kidney by converting angiotensin I to angiotensin II, resulting in an increase of the vasoconstrictor activity of angiotensin (PubMed: <a href="#">11432860</a> , PubMed: <a href="#">1851160</a> , PubMed: <a href="#">19773553</a> , PubMed: <a href="#">23056909</a> , PubMed: <a href="#">4322742</a> ).

Also able to inactivate bradykinin, a potent vasodilator, and therefore enhance the blood pressure response (PubMed:[15615692](#), PubMed:[2558109](#), PubMed:[4322742](#), PubMed:[6055465](#), PubMed:[6270633](#), PubMed:[7683654](#)). Acts as a regulator of synaptic transmission by mediating cleavage of neuropeptide hormones, such as substance P, neurotensin or enkephalins (PubMed:[15615692](#), PubMed:[6208535](#), PubMed:[6270633](#), PubMed:[656131](#)). Catalyzes degradation of different enkephalin neuropeptides (Met-enkephalin, Leu-enkephalin, Met-enkephalin-Arg-Phe and possibly Met-enkephalin-Arg-Gly-Leu) (PubMed:[2982830](#), PubMed:[6270633](#), PubMed:[656131](#)). Acts as a regulator of synaptic plasticity in the nucleus accumbens of the brain by mediating cleavage of Met-enkephalin-Arg-Phe, a strong ligand of Mu-type opioid receptor OPRM1, into Met-enkephalin (By similarity). Met-enkephalin-Arg-Phe cleavage by ACE decreases activation of OPRM1, leading to long-term synaptic potentiation of glutamate release (By similarity). Also acts as a regulator of hematopoietic stem cell differentiation by mediating degradation of hemoregulatory peptide N-acetyl-SDKP (AcSDKP) (PubMed:[26403559](#), PubMed:[7876104](#), PubMed:[8257427](#), PubMed:[8609242](#)). Acts as a regulator of cannabinoid signaling pathway by mediating degradation of hemopressin, an antagonist peptide of the cannabinoid receptor CNR1 (PubMed:[18077343](#)). Involved in amyloid-beta metabolism by catalyzing degradation of Amyloid-beta protein 40 and Amyloid-beta protein 42 peptides, thereby preventing plaque formation (PubMed:[11604391](#), PubMed:[16154999](#), PubMed:[19773553](#)). Catalyzes cleavage of cholecystokinin (maturation of Cholecystokinin-8 and Cholecystokinin-5) and Gonadoliberin-1 (both maturation and degradation) hormones (PubMed:[10336644](#), PubMed:[2983326](#), PubMed:[7683654](#), PubMed:[9371719](#)). Degradation of hemoregulatory peptide N-acetyl-SDKP (AcSDKP) and amyloid-beta proteins is mediated by the N-terminal catalytic domain, while angiotensin I and cholecystokinin cleavage is mediated by the C-terminal catalytic region (PubMed:[10336644](#), PubMed:[19773553](#), PubMed:[7876104](#)).

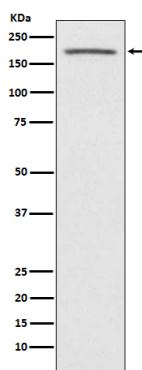
## Cellular Location

Cell membrane; Single-pass type I membrane protein. Cytoplasm {ECO:0000250|UniProtKB:P09470}. Note=Detected in both cell membrane and cytoplasm in neurons. {ECO:0000250|UniProtKB:P09470} [Isoform Testis-specific]: Cell membrane; Single-pass type I membrane protein. Secreted. Note=The testis-specific isoform can be cleaved before the transmembrane region, releasing a soluble form

## Tissue Location

Ubiquitously expressed, with highest levels in lung, kidney, heart, gastrointestinal system and prostate

## Images



Western blot analysis of ACE1 expression in human fetal kidney lysate.