

ACE1 Antibody
Rabbit mAb
Catalog # AP91542**Specification****ACE1 Antibody - Product Information**

Application	WB, IHC, FC
Primary Accession	P12821
Clonality	Monoclonal
Other Names	
Angiotensin-converting enzyme; somatic isoform precursor; CD143 antigen; DCP; DCP1; Dipeptidyl carboxypeptidase I; Kininase II;	
Isotype	Rabbit IgG
Host	Rabbit
Calculated MW	149715 Da

ACE1 Antibody - Additional Information

Dilution	WB~~1:1000 IHC~~1:100~500 FC~~1:10~50
Purification	Affinity-chromatography
Immunogen	A synthesized peptide derived from human ACE1
Description	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.
Storage Condition and Buffer	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.

ACE1 Antibody - Protein Information

Name ACE {ECO:0000303|PubMed:2849100, ECO:0000312|HGNC:HGNC:2707}

Function

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:15615692, PubMed:>20826823, PubMed:>2558109, PubMed:>4322742, PubMed:>7523412, PubMed:>7683654). Composed of two similar catalytic domains, each possessing a functional active site, with different selectivity for substrates (PubMed:>10913258, PubMed:>1320019, PubMed:>1851160, PubMed:>19773553, PubMed:>7683654, PubMed:>7876104). 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:>11432860, PubMed:>1851160, PubMed:>19773553, PubMed:>23056909, PubMed:>4322742). 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:<a

[10336644](http://www.uniprot.org/citations/10336644), PubMed:[2983326](http://www.uniprot.org/citations/2983326), PubMed:[7683654](http://www.uniprot.org/citations/7683654), PubMed:[9371719](http://www.uniprot.org/citations/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](http://www.uniprot.org/citations/10336644), PubMed:[19773553](http://www.uniprot.org/citations/19773553), PubMed:[7876104](http://www.uniprot.org/citations/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

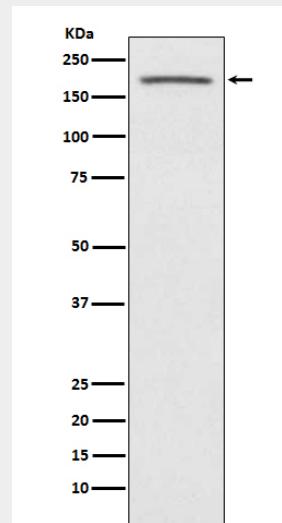
Ubiquitously expressed, with highest levels in lung, kidney, heart, gastrointestinal system and prostate

ACE1 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

ACE1 Antibody - Images



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