



href="http://www.uniprot.org/citations/7523412" target="\_blank">7523412</a>, PubMed:<a href="http://www.uniprot.org/citations/7683654" target="\_blank">7683654</a>). Composed of two similar catalytic domains, each possessing a functional active site, with different selectivity for substrates (PubMed:<a href="http://www.uniprot.org/citations/10913258" target="\_blank">10913258</a>, PubMed:<a href="http://www.uniprot.org/citations/1320019" target="\_blank">1320019</a>, PubMed:<a href="http://www.uniprot.org/citations/1851160" target="\_blank">1851160</a>, PubMed:<a href="http://www.uniprot.org/citations/19773553" target="\_blank">19773553</a>, PubMed:<a href="http://www.uniprot.org/citations/7683654" target="\_blank">7683654</a>, PubMed:<a href="http://www.uniprot.org/citations/7876104" target="\_blank">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="http://www.uniprot.org/citations/11432860" target="\_blank">11432860</a>, PubMed:<a href="http://www.uniprot.org/citations/1851160" target="\_blank">1851160</a>, PubMed:<a href="http://www.uniprot.org/citations/19773553" target="\_blank">19773553</a>, PubMed:<a href="http://www.uniprot.org/citations/23056909" target="\_blank">23056909</a>, PubMed:<a href="http://www.uniprot.org/citations/4322742" target="\_blank">4322742</a>). Also able to inactivate bradykinin, a potent vasodilator, and therefore enhance the blood pressure response (PubMed:<a href="http://www.uniprot.org/citations/15615692" target="\_blank">15615692</a>, PubMed:<a href="http://www.uniprot.org/citations/2558109" target="\_blank">2558109</a>, PubMed:<a href="http://www.uniprot.org/citations/4322742" target="\_blank">4322742</a>, PubMed:<a href="http://www.uniprot.org/citations/6055465" target="\_blank">6055465</a>, PubMed:<a href="http://www.uniprot.org/citations/6270633" target="\_blank">6270633</a>, PubMed:<a href="http://www.uniprot.org/citations/7683654" target="\_blank">7683654</a>). Acts as a regulator of synaptic transmission by mediating cleavage of neuropeptide hormones, such as substance P, neurotensin or enkephalins (PubMed:<a href="http://www.uniprot.org/citations/15615692" target="\_blank">15615692</a>, PubMed:<a href="http://www.uniprot.org/citations/6208535" target="\_blank">6208535</a>, PubMed:<a href="http://www.uniprot.org/citations/6270633" target="\_blank">6270633</a>, PubMed:<a href="http://www.uniprot.org/citations/656131" target="\_blank">656131</a>). Catalyzes degradation of different enkephalin neuropeptides (Met- enkephalin, Leu-enkephalin, Met-enkephalin-Arg-Phe and possibly Met- enkephalin-Arg-Gly-Leu) (PubMed:<a href="http://www.uniprot.org/citations/2982830" target="\_blank">2982830</a>, PubMed:<a href="http://www.uniprot.org/citations/6270633" target="\_blank">6270633</a>, PubMed:<a href="http://www.uniprot.org/citations/656131" target="\_blank">656131</a>). 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:<a href="http://www.uniprot.org/citations/26403559" target="\_blank">26403559</a>, PubMed:<a href="http://www.uniprot.org/citations/7876104" target="\_blank">7876104</a>, PubMed:<a href="http://www.uniprot.org/citations/8257427" target="\_blank">8257427</a>, PubMed:<a href="http://www.uniprot.org/citations/8609242" target="\_blank">8609242</a>). Acts as a regulator of cannabinoid signaling pathway by mediating degradation of hemopressin, an antagonist peptide of the cannabinoid receptor CNR1 (PubMed:<a href="http://www.uniprot.org/citations/18077343" target="\_blank">18077343</a>). Involved in amyloid-beta metabolism by catalyzing degradation of Amyloid-beta protein 40 and Amyloid-beta protein 42 peptides, thereby preventing plaque formation (PubMed:<a href="http://www.uniprot.org/citations/11604391" target="\_blank">11604391</a>, PubMed:<a href="http://www.uniprot.org/citations/16154999" target="\_blank">16154999</a>, PubMed:<a href="http://www.uniprot.org/citations/19773553" target="\_blank">19773553</a>). Catalyzes cleavage of cholecystokinin (maturation of Cholecystokinin-8 and Cholecystokinin-5) and Gonadoliberin-1 (both maturation and degradation) hormones (PubMed:<a href="http://www.uniprot.org/citations/10336644" target="\_blank">10336644</a>, PubMed:<a href="http://www.uniprot.org/citations/2983326" target="\_blank">2983326</a>, PubMed:<a href="http://www.uniprot.org/citations/7683654" target="\_blank">7683654</a>, PubMed:<a

<http://www.uniprot.org/citations/9371719> target="\_blank">9371719</a>). 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:<a href="http://www.uniprot.org/citations/10336644" target="\_blank">10336644</a>, PubMed:<a href="http://www.uniprot.org/citations/19773553" target="\_blank">19773553</a>, PubMed:<a href="http://www.uniprot.org/citations/7876104" target="\_blank">7876104</a>).

#### 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

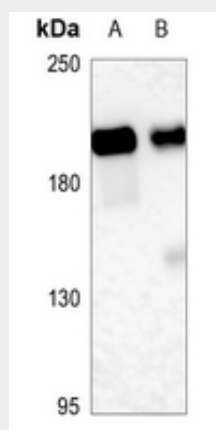
Ubiquitously expressed, with highest levels in lung, kidney, heart, gastrointestinal system and prostate

### Anti-CD143 Antibody - Protocols

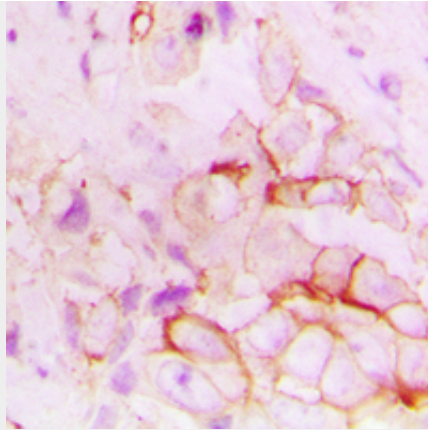
Provided below are standard protocols that you may find useful for product applications.

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

### Anti-CD143 Antibody - Images



Western blot analysis of CD143 expression in mouse kidney (A), mouse testis (B) whole cell lysates.



Immunohistochemical analysis of CD143 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

#### **Anti-CD143 Antibody - Background**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human CD143. The exact sequence is proprietary.