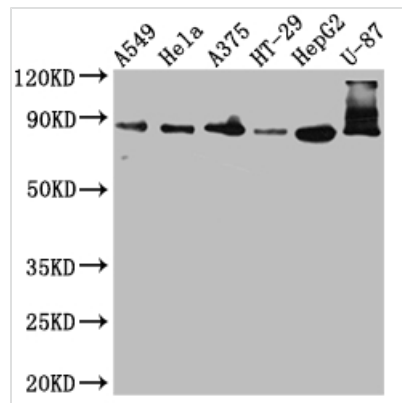




# FURIN Antibody

<b>Product Code</b>	CSB-RA271669A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P09958
<b>Immunogen</b>	A synthesized peptide derived from human Furin
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, WB, IHC; Recommended dilution: WB:1:500-1:5000, IHC:1:50-1:200
<b>Relevance</b>	Furin is likely to represent the ubiquitous endoprotease activity within constitutive secretory pathways and capable of cleavage at the RX(K/R)R consensus motif.
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Rabbit IgG
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Cell biology; Metabolism; Signal transduction
<b>Gene Names</b>	FURIN
<b>Accession NO.</b>	1C3

## Image



### Western Blot

Positive WB detected in: A549 whole cell lysate, HeLa whole cell lysate, A375 whole cell lysate, HT-29 whole cell lysate, HepG2 whole cell lysate, U-87 whole cell lysate

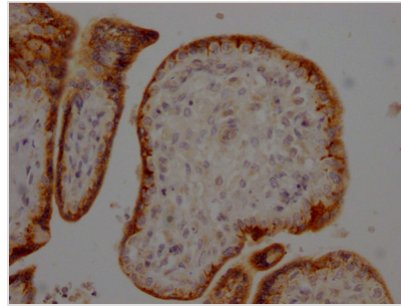
All lanes: Furin antibody at 1:1000

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 87 kDa

Observed band size: 87, 72 kDa



IHC image of CSB-RA271669A0HU diluted at 1:100 and staining in paraffin-embedded human placenta tissue performed on a Leica Bond<sup>TM</sup> system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

## Description

FURIN is a subtilisin-like proprotein convertase with diverse cellular functions. Furin has a well-known role in viral pathogenesis and efficiently cleaves polybasic or multi-basic sites such as those found in influenza virus subtype H5 and H7. Emerging evidence suggests that FURIN plays a critical role in the activation and infectivity of SARS-CoV-2. In SARS-CoV-2, structural studies have suggested that furin cleavage at the S1/S2 boundary primes the spike for an open conformation required for binding to the ACE2 entry receptor.

The generation of this recombinant FURIN antibody occurs in a series of steps: immunization, splenocytes & PBMC, single B cell sorting, mRNA extraction, RT-PCR & insert vector, expression, ELISA validation. And ELISA, WB, IHC was carried out Every step was performed under strict standards to ensure the researchers can have a recombinant FURIN antibody with premium quality.