

Cleaved-Plasma Kallikrein HC (R390) Polyclonal Antibody

Catalog No: YC0102

Reactivity: Human; Rat; Mouse;

Applications: WB;IHC;IF;ELISA

Target: Plasma Kallikrein HC

Fields: >>Complement and coagulation cascades

P03952

P26262

Gene Name: KLKB1

Protein Name: Plasma kallikrein

Human Gene Id: 3818

Human Swiss Prot

iuman Swiss Frot

No:

Mouse Swiss Prot

No:

Immunogen:

The antiserum was produced against synthesized peptide derived from human

KLKB1. AA range:341-390

Specificity: Cleaved-Plasma Kallikrein HC (R390) Polyclonal Antibody detects endogenous

levels of fragment of activated Plasma Kallikrein HC protein resulting from

cleavage adjacent to R390.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:10000.. IF 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

1/3



Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 45kD

Cell Pathway: Complement and coagulation cascades;

Background : This gene encodes a glycoprotein that participates in the surface-dependent

activation of blood coagulation, fibrinolysis, kinin generation and inflammation. The encoded preproprotein present in plasma as a non-covalent complex with high molecular weight kininogen undergoes proteolytic processing mediated by activated coagulation factor XII to generate a disulfide-linked, heterodimeric serine protease comprised of heavy and light chains. Certain mutations in this gene cause prekallikrein deficiency. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jan 2016],

Function: catalytic activity: Cleaves selectively Arg-|-Xaa and Lys-|-Xaa bonds, including

Lys-|-Arg and Arg-|-Ser bonds in (human) kininogen to release

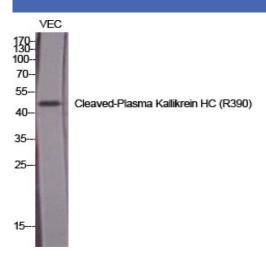
bradykinin., disease: Defects in KLKB1 are the cause of prekallikrein deficiency (PKK deficiency) [MIM:612423]; also called Fletcher factor deficiency. This disorder is a blood coagulation defect., function: The enzyme cleaves Lys-Arg and Arg-Ser bonds. It activates, in a reciprocal reaction, factor XII after its binding to a negatively charged surface. It also releases bradykinin from HMW kininogen and may also play a role in the renin-angiotensin system by converting prorenin into renin., similarity: Belongs to the peptidase S1 family., similarity: Belongs to the peptidase S1 family. Plasma kallikrein subfamily., similarity: Contains 1 peptidase S1 domain., similarity: Contains 4 apple domains., subunit: The zymogen is

activated by factor XIIa, which cleaves t

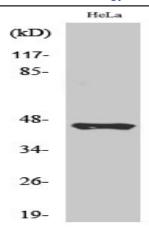
Subcellular Location : Secreted.

Expression : Colon, Liver, Plasma,

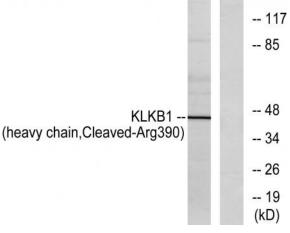
Products Images



Western Blot analysis of various cells using Cleaved-Plasma Kallikrein HC (R390) Polyclonal Antibody diluted at 1:1000



Western Blot analysis of HeLa cells using Cleaved-Plasma Kallikrein HC (R390) Polyclonal Antibody diluted at 1:1000



Western blot analysis of lysates from HeLa cells, using KLKB1 (heavy chain, Cleaved-Arg390) Antibody. The lane on the right is blocked with the synthesized peptide.



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 30min).