abcam

Product datasheet

Human CXCL7 ELISA Kit ab100613

4 References 3 Images

Overview

Product name Human CXCL7 ELISA Kit

Detection methodColorimetric

Sample type Cell culture supernatant, Serum, Plasma, Cell Lysate

Assay type Sandwich (quantitative)

Sensitivity < 8.5 pg/ml

Range 4.1 pg/ml - 1000 pg/ml

Recovery 93 %

Sample specific recovery

Sample type	Average %	Range
Cell culture supernatant	95.28	84% - 103%
Serum	92.76	82% - 102%
Plasma	93.45	83% - 103%

Assay duration Multiple steps standard assay

Species reactivity Reacts with: Human

Product overview Abcam's CXCL7 Human ELISA (Enzyme-Linked Immunosorbent Assay) kit is an in vitro enzyme-

linked immunosorbent assay for the quantitative measurement of Human CXCL7 in serum,

plasma, and cell culture supernatants.

This assay employs an antibody specific for Human CXCL7 coated on a 96-well plate. Standards and samples are pipetted into the wells and CXCL7 present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-Human CXCL7 antibody is added. After washing away unbound biotinylated antibody, HRP-conjugated streptavidin is pipetted to the wells. The wells are again washed, a TMB substrate solution is added to the wells and color develops in proportion to the amount of CXCL7 bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is measured at 450 nm.

Notes Optimization may be required with urine samples.

Platform Microplate

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Properties

Storage instructions

Store at -20°C. Please refer to protocols.

Components	1 x 96 tests
200X HRP-Streptavidin Concentrate	1 x 200µl
20X Wash Buffer	1 x 25ml
5X Assay Diluent B	1 x 15ml
Assay Diluent A	1 x 30ml
Biotinylated anti-Human CXCL7	2 vials
CXCL7 Microplate (12 x 8 wells)	1 unit
Recombinant Human CXCL7 Standard (lyophilized)	2 vials
Stop Solution	1 x 8ml
TMB One-Step Substrate Reagent	1 x 12ml

Function

LA-PF4 stimulates DNA synthesis, mitosis, glycolysis, intracellular cAMP accumulation, prostaglandin E2 secretion, and synthesis of hyaluronic acid and sulfated glycosaminoglycan. It also stimulates the formation and secretion of plasminogen activator by human synovial cells. NAP-2 is a ligand for CXCR1 and CXCR2, and NAP-2, NAP-2(73), NAP-2(74), NAP-2(1-66), and most potent NAP-2(1-63) are chemoattractants and activators for neutrophils. TC-1 and TC-2 are antibacterial proteins, in vitro released from activated platelet alpha-granules. CTAP-III(1-81) is more potent than CTAP-III desensitize chemokine-induced neutrophil activation.

Sequence similarities

Post-translational modifications

Belongs to the intercrine alpha (chemokine CxC) family.

Proteolytic removal of residues 1-9 produces the active peptide connective tissue-activating peptide III (CTAP-III) (low-affinity platelet factor IV (LA-PF4)).

 $Proteolytic\ removal\ of\ residues\ 1-13\ produces\ the\ active\ peptide\ beta-thromboglobulin,\ which\ is$

NAP-2(1-66) is produced by proteolytical processing, probably after secretion by leukocytes other

released from platelets along with platelet factor 4 and platelet-derived growth factor.

than neutrophils.

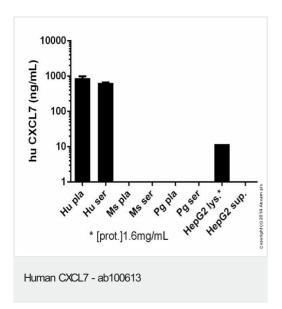
NAP-2(73) and NAP-2(74) seem not be produced by proteolytical processing of secreted

precursors but are released in an active form from platelets.

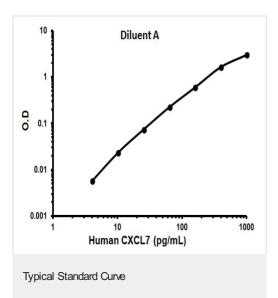
Cellular localization

Secreted.

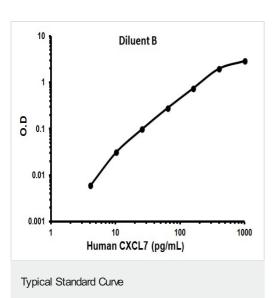
Images



CXCL7 measured in biological fluids showing quantity (ng) per mL of tested sample



Representative Standard Curve using ab100613.



Representative Standard Curve using ab100613.

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