Coud-Clone Corp.

APB159Hu61 100µg Active Integrin Alpha X (CD11c) Organism Species: *Homo sapiens* (Human) *Instruction manual* 

#### FOR RESEARCH USE ONLY NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

13th Edition (Revised in Aug, 2023)

#### [PROPERTIES]

Source: Eukaryotic expression.

Host: 293F cell

Residues: Gln143~Ser344

Tags: N-terminal His-tag

**Purity: >95%** 

Endotoxin Level: <1.0EU per 1µg (determined by the LAL method).

Buffer Formulation: PBS, pH7.4, containing 5% Trehalose .

Original Concentration: 200µg/mL

Applications: Cell culture; Activity Assays.

(May be suitable for use in other assays to be determined by the end user.)

Predicted isoelectric point: 7.1

Predicted Molecular Mass: 24.4kDa

Accurate Molecular Mass: 27&29kDa as determined by SDS-PAGE reducing conditions.

Phenomenon explanation:

The possible reasons that the actual band size differs from the predicted are as follows:

1. Splice variants: Alternative splicing may create different sized proteins from the same gene.

- 2. Relative charge: The composition of amino acids may affects the charge of the protein.
- 3. Post-translational modification: Phosphorylation, glycosylation, methylation etc.

4. Post-translation cleavage: Many proteins are synthesized as pro-proteins, and then cleaved to give the active form.

5. Polymerization of the target protein: Dimerization, multimerization etc.

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## [ <u>USAGE</u> ]

Reconstitute in 10mM PBS (pH7.4) to a concentration of 0.1-1.0 mg/mL. Do not vortex.

### [ STORAGE AND STABILITY ]

Storage: Avoid repeated freeze/thaw cycles.

Store at 2-8°C for one month.

Aliquot and store at -80°C for 12 months.

**Stability Test:** The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration date under appropriate storage condition.

#### [SEQUENCE]

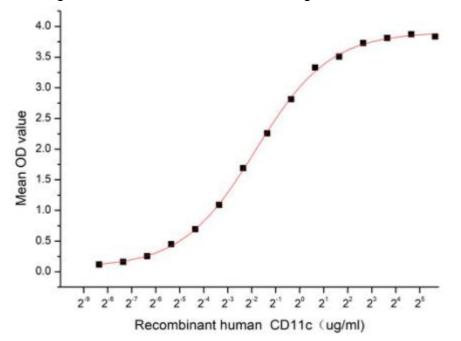
QECPRQEQ DIVFLIDGSG SISSRNFATM MNFVRAVISQ FQRPSTQFSL MQFSNKFQTH FTFEEFRRSS NPLSLLASVH QLQGFTYTAT AIQNVVHRLF HASYGARRDA AKILIVITDG KKEGDSLDYK DVIPMADAAG IIRYAIGVGL AFQNRNSWKE LNDIASKPSQ EHIFKVEDFD ALKDIQNQLK EKIFAIEGTE TTSS

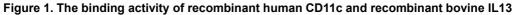
## [ACTIVITY]

Integrin Alpha X (CD11c) is a member of the leukointegrin family, it is an approximately 150 kDa type I transmembrane glycoprotein that heterodimerizes with Integrin beta 2. One of the earliest described properties of CD11c is its ability to mediate phagocytosis of iC3b-opsonized particles. Accordingly, CD11c is known as complement receptor 4. CD11c has also been implicated in phagocytosis of latex beads and bacteria in the absence of complement. CD11c binds a diverse array of ligands such as cell adhesion molecules and Interleukin 13 (IL13). Thus a functional binding ELISA assay was conducted to detect the interaction of recombinant human CD11c and recombinant bovine IL13. Briefly, CD11c was

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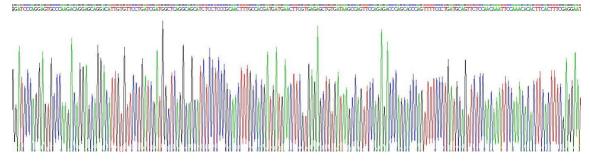
diluted serially in PBS with 0.01% BSA (pH 7.4). Duplicate samples of 100  $\mu$  I were then transferred to IL13-coated microtiter wells and incubated for 1h at 37 °C. Wells were washed with PBST and incubated for 1h with anti-CD11c pAb, then aspirated and washed 3 times. After incubation with HRP labelled secondary antibody for 1h at 37 °C, wells were aspirated and washed 5 times. With the addition of substrate solution, wells were incubated 15-25 minutes at 37 °C. Finally, add 50  $\mu$ I stop solution to the wells and read at 450/630 nm immediately. The binding activity of recombinant human CD11c and recombinant bovine IL13 was shown in Figure 1, the EC50 for this effect is 0.28 ug/mL.





#### [IDENTIFICATION]

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#### Figure 2. Gene Sequencing (extract)

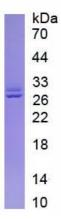


Figure 3. SDS-PAGE

Sample: Active recombinant CD11c, Human

### [<u>IMPORTANT NOTE</u>]

The kit is designed for research use only, we will not be responsible for any issue if the kit was used in clinical diagnostic or any other procedures.