# abcam

## Product datasheet

## Human IL-17A Receptor ELISA Kit ab100558

## 2 Images

#### Overview

Product name Human IL-17A Receptor ELISA Kit

**Detection method**Colorimetric

Sample type Cell culture supernatant, Serum, Plasma

**Assay type** Sandwich (quantitative)

Sensitivity < 10 pg/ml

**Range** 12.35 pg/ml - 3000 pg/ml

Recovery 81 %

Sample specific recovery

Sample type	Average %	Range
Cell culture supernatant	81.54	70% - 92%
Serum	87.64	83% - 104%
Plasma	73.27	67% - 87%

**Assay duration** Multiple steps standard assay

Species reactivity Reacts with: Human

Product overview Abcam's IL-17A Receptor Human ELISA (Enzyme Linked Immunosorbent Assay) kit is an in vitro

enzyme-linked immunosorbent assay for the quantitative measurement of Human IL-17A in serum,

plasma, and cell culture supernatants.

This assay employs an antibody specific for Human IL-17A coated on a 96-well plate. Standards and samples are pipetted into the wells and IL-17A present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-Human IL-17A 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 IL-17A 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 C	1 x 30ml
Biotinylated anti-Human IL-17A	2 vials
IL-17A Microplate (12 x 8 wells)	1 x 200µl
Recombinant Human IL-17A Standard (lyophilized)	2 vials
Stop Solution	1 x 8ml
TMB One-Step Substrate Reagent	1 x 12ml

Function Receptor for IL17A and IL17F. Binds its IL17A ligand with low affinity, suggesting that additional

components are involved in IL17A-induced signaling.

**Tissue specificity** Widely expressed.

**Involvement in disease** Defects in IL17RA are the cause of familial candidiasis type 5 (CANDF5) [MIM:613953].

CANDF5 is a rare disorder with altered immune responses and impaired clearance of fungal infections, selective against Candida. It is characterized by persistent and/or recurrent infections of the skin, nails and mucous membranes caused by organisms of the genus Candida, mainly

Candida albicans.

Sequence similarities Contains 1 SEFIR domain.

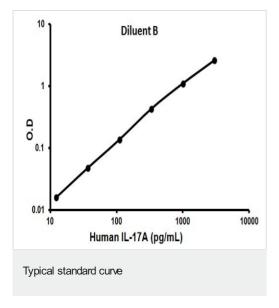
Post-translational

modifications

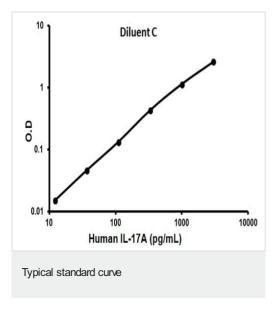
Glycosylated.

Cellular localization Membrane.

## **Images**



Representative standard curve using ab100558



Representative standard curve using ab100558

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