

# CD155 antibody

# Cat. No. GTX129572

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB
Reactivity	Human

Package  $100~\mu l,\,25~\mu l$ 

# **Applications**

## **Application Note**

\*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	1:500-1:3000
Not tested in other applications.	

**Calculated MW** 45 kDa. ( <u>Note</u> )

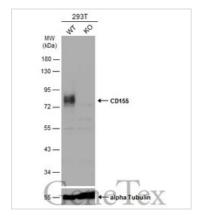
Properties	
Form	Liquid
Buffer	PBS, 20% Glycerol
Preservative	0.025% ProClin 300
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	1 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	Synthetic peptide encompassing a sequence within the Intracellular domain of human CD155. The exact sequence is proprietary.
Purification	Purified by antigen-affinity chromatography.
Conjugation	Unconjugated
Note	For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.
	Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.



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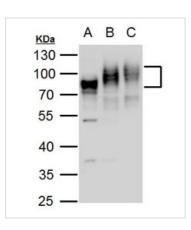
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## DATA IMAGES



#### GTX129572 WB Image

Wild-type (WT) and CD155 knockout (KO) 293T cell extracts (30  $\mu$ g) were separated by 10% SDS-PAGE, and the membrane was blotted with CD155 antibody (GTX129572) diluted at 1:1000. The HRP-conjugated antirabbit IgG antibody (GTX213110-01) was used to detect the primary antibody.



#### GTX129572 WB Image

CD155 antibody detects CD155 protein by western blot analysis.

A. 30  $\mu g$  K562 whole cell lysate/extract

B. 30 µg THP-1 whole cell lysate/extract

C.  $30 \, \mu g$  HL- $60 \, whole \, cell \, lysate/extract$ 

10 % SDS-PAGE

CD155 antibody (GTX129572) dilution: 1:1000



For full product information, images and publications, please visit our <u>website</u>.

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