

C/EBP delta antibody

Cat. No. GTX17179

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, IHC-P, ELISA
Reactivity	Human, Mouse, Rat

Package
100 µg

Applications

Application Note

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

Suggested dilution	Recommended dilution
WB	0.5 - 1 µg/mL
IHC-P	20 µg/mL
ELISA	Assay dependent

Not tested in other applications.

Calculated MW	28 kDa. (Note)
Product Note	Multiple transcript variants encoding different isoforms have been found for this gene.

Properties

Form	Liquid
Buffer	PBS
Preservative	0.02% Sodium azide
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	Rabbit polyclonal C/EBP delta antibody was raised against an 18 amino acid peptide near the carboxy terminus of human C/EBP delta. The immunogen is located within amino acids 170 - 220 of C/EBP delta.
Purification	Purified by antigen-affinity chromatography
Conjugation	Unconjugated



For full product information, images and publications, please visit our [website](#).

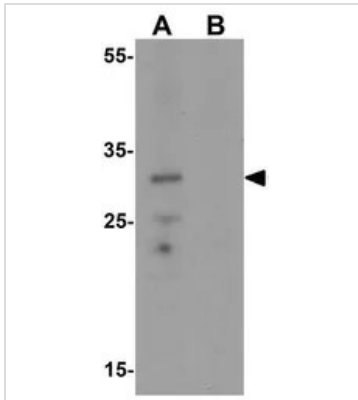
Date 2026 / 01 / 11 Page 1 of 2

For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

Note

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.

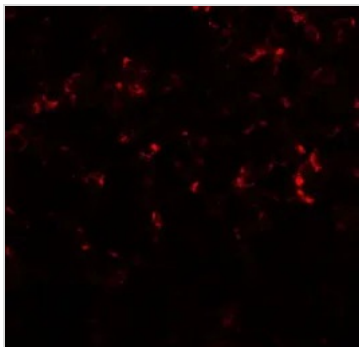
DATA IMAGES



GTX17179 WB Image

WB analysis of rat spleen tissue lysate in (A) the absence and (B) the presence of blocking peptide using GTX17179 C/EBP delta antibody.

Working concentration : 0.5 µg/ml



GTX17179 IHC-P Image

IHC-P analysis of human spleen tissue using GTX17179 C/EBP delta antibody.

Working concentration : 20 µg/ml



For full product information, images and publications, please visit our [website](#).