

AMBRA1 antibody

Cat. No. GTX17003

Host	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Application	WB, ICC/IF, IHC-P, ELISA	
Reactivity	Human, Mouse, Rat	

Reference (1) Package 100 µg

APPLICATION

Application Note

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

Suggested dilution	Recommended dilution
WB	1 μg/mL
ICC/IF	Assay dependent
IHC-P	5 μg/mL
ELISA	Assay dependent

Not tested in other applications.

Calculated MW 143 kDa. (Note)

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	Ambra1 antibody was raised against a 15 amino acid synthetic peptide from near the carboxy terminus of human Ambra1. The immunogen is located within the last 50 amino acids of Ambra1.	
Purification	Purified by antigen-affinity chromatography	
Conjugation	Unconjugated	



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

Date 2024 / 04 / 25 Page 1 of 2

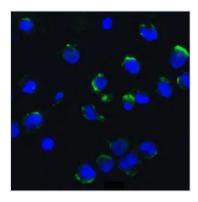


For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

Note

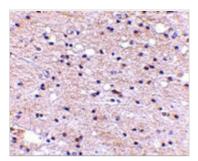
Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.

DATA IMAGES



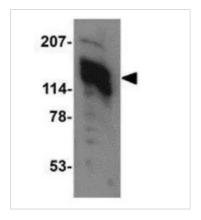
GTX17003 ICC/IF Image

ICC/IF analysis of K562 cells using GTX17003 AMBRA1 antibody. Dilution : 20 $\mu g/ml$



GTX17003 IHC-P Image

IHC-P analysis of human brain tissue using GTX17003 AMBRA1 antibody. Working concentration : 5 μ g/ml



GTX17003 WB Image

WB analysis of 3T3 cell lysate using GTX17003 AMBRA1 antibody. Working concentration : 1 $\mu g/ml$



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

Date 2024 / 04 / 25 Page 2 of 2