

## DRAK1 antibody

Cat. No. GTX31646

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, ICC/IF, ELISA
Reactivity	Human

Package  
100 µg

## Applications

## Application Note

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

Suggested dilution	Recommended dilution
WB	1 µg/mL
ICC/IF	2 µg/mL
ELISA	Assay dependent

Not tested in other applications.

Calculated MW	47 kDa. ( <a href="#">Note</a> )
Product Note	No cross responses to DRAK2, DAP or ZIP kinases.

## 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	Batch dependent (Please refer to the vial label for the specific concentration.)
Immunogen	DRAK1 antibody was raised against a peptide corresponding to amino acids near the amino terminus of human DRAK1. The immunogen is located within the first 50 amino acids of DRAK1.
Purification	Purified by antigen-affinity chromatography
Conjugation	Unconjugated



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

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

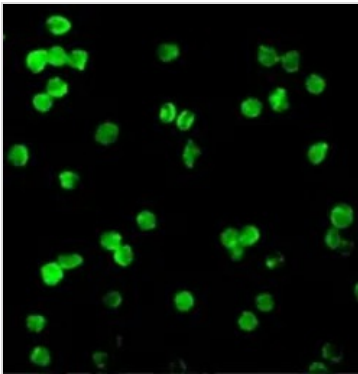
**Note**

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.

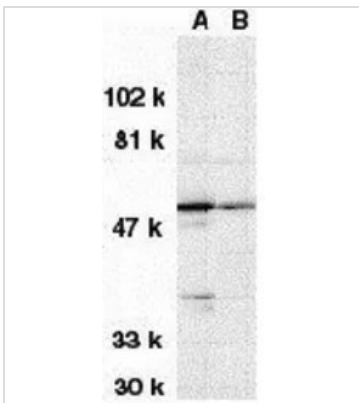
**DATA IMAGES**

**GTX31646 ICC/IF Image**

ICC/IF analysis of MOLT4 cells using GTX31646 DRAK1 antibody.  
Working concentration : 2 µg/ml


**GTX31646 ICC/IF Image**

ICC/IF analysis of MOLT4 cells using GTX31646 DRAK1 antibody.  
Working concentration : 20 µg/ml


**GTX31646 WB Image**

WB analysis of (A) MOLT4 and (B) A431 whole cell lysates using GTX31646 DRAK1 antibody.  
Working concentration : 1 µg/ml



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