

EGFR (phospho Tyr1086) antibody

Cat. No. GTX25650

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

Package 50 μl

Applications

Application Note

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

Suggested dilution		Recommended dilution	
WB		1:1000	
ICC/IF		1:100	
IHC		Assay dependent	
Not tested in other applications.			
Calculated MW	134 kDa. (<u>Note</u>)		

Product Note	The full-length EGFR includes the 24 aa signal peptide that is cleaved during EGFR maturation. The Tyr1086 of mature EGFR
	is equivalent to Tyr1110 of uncleaved EGFR.

FormLiquidBufferPBS, 0.1% BSA, 50% GlycerolPreservative0.05% Sodium azideStorageStore 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.	Properties	
Preservative 0.05% Sodium azide Storage Storage Storage 1.2 weeks), store at 4°C. For short-term storage (1-2 weeks), store at 4°C.	Form	Liquid
Storage Storag	Buffer	PBS, 0.1% BSA, 50% Glycerol
Storage	Preservative	0.05% Sodium azide
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Concentration Batch dependent (Please refer to the vial label for the specific concentration.)	Concentration	Batch dependent (Please refer to the vial label for the specific concentration.)
Immunogen The antiserum was produced against a chemically synthesized phosphopeptide derived from the region of human EGFR that contains tyrosine 1086. The sequence is conserved in mouse and rat.	Immunogen	
Purification Purified by antigen-affinity chromatography	Purification	Purified by antigen-affinity chromatography
Conjugation Unconjugated	Conjugation	Unconjugated



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

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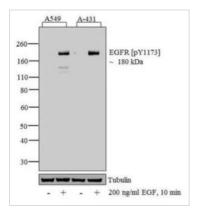


For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

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.

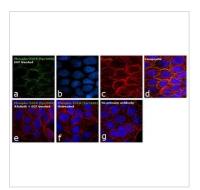
DATA IMAGES



GTX25650 WB Image

WB analysis of 30 μ g of A549 (Lane 1) treated for 10 minutes with 200 ng/ml of EGF (Lane 2), A431 (Lane 3) treated for 10 minutes with 200 ng/ml of EGF (Lane 4) cell lysate using GTX25650 EGFR (phospho Tyr1086) antibody.

Dilution: 1:1000



GTX25650 ICC/IF Image

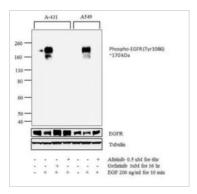
ICC/IF analysis of A431 cells treated with Afatinib (1uM for 6hrs) followed by EGF (200ng/ml for 10 minutes) using GTX25650 EGFR (phospho Tyr1086) antibody.

Green: Primary antibody

Blue : Nuclei Red : Actin

Fixation: 4% paraformaldehyde

Permeabilization: 0.1% Triton X-100 for 15 minutes Dilution: 1:100 in 0.1% BSA incubated at 4°C



GTX25650 WB Image

WB analysis of membrane enriched extracts (30 ug lysate) of A-431 (Lane 1) treated with EGF (200ng/ml for 10 minutes) (Lane 2), A-431 treated with Gefitinib followed by EGF (1uM for 16 hours, 200ng/ml for 10 minutes) (Lane 3), A-431 treated with Afatinib followed by EGF (0.5 uM for 6 hours, 200ng/ml for 10 minutes) (Lane 4), A549 (Lane 5) treated with EGF (200ng/ml for 10 minutes) (Lane 6), and A549 treated with Afatinib followed by EGF (0.5 uM for 6 hours, 200ng/ml for 10 minutes) (Lane 7) using GTX25650 EGFR (phospho Tyr1086) antibody.

Dilution: 1:1000



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