

## ERK1 (phospho Thr202/Tyr204) + ERK2 (phospho Thr185/Tyr187) antibody

**Cat. No. GTX24819**

<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG
<b>Applications</b>	WB, ICC/IF, IHC-P, IHC, IHC (Free Floating)
<b>Reactivity</b>	Human, Mouse, Rat, Drosophila, Bovine, Chicken, Caenorhabditis elegans, Xenopus

References ( 33 )

Package

50 µl

## Applications

**Application Note**

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

Suggested dilution	Recommended dilution
WB	1:1000
ICC/IF	Assay dependent
IHC-P	1:10-1:100
IHC	Assay dependent
IHC (Free Floating)	Assay dependent

Not tested in other applications.

**Product Note**

The antibody recognizes the same sequence in both proteins (ERK1/2) as long as the T and the Y are phosphorylated. Does not recognize either monophospho- forms of ERK1/2 or the non-phosphorylated proteins.

## Properties

<b>Form</b>	Liquid
<b>Buffer</b>	PBS, 0.1% BSA, 50% Glycerol
<b>Preservative</b>	0.05% Sodium azide
<b>Storage</b>	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.
<b>Concentration</b>	Batch dependent (Please refer to the vial label for the specific concentration.)
<b>Immunogen</b>	The antiserum was produced against a chemically synthesized phosphopeptide derived from the region of human ERK1 and 2 that contains threonine 202/185 and tyrosine 204/187. This region is conserved among many species including rat, mouse, cow, frog, snail, nematode, and fruit fly.
<b>Purification</b>	Purified by antigen-affinity chromatography
<b>Conjugation</b>	Unconjugated



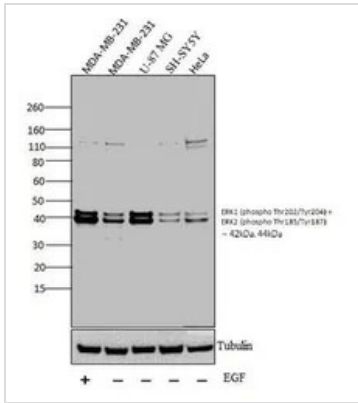
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**Note**  
Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.

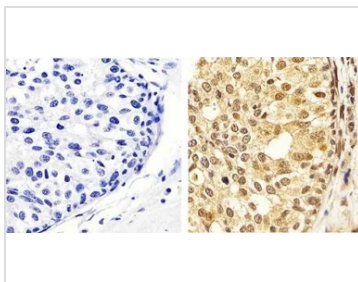
## DATA IMAGES



### GTX24819 WB Image

WB analysis of whole cell extracts (30 µg lysate) of MDA-MB-231 with treatment of EGF(100ng/ml for 15mins) (Lane 1), MDA-MB-231 (Lane 2), U-87 MG (Lane 3), SH-SY5Y (Lane 4) and HeLa (Lane 5) using GTX24819 ERK1 (phospho Thr202/Tyr204) + ERK2 (phospho Thr185/Tyr187) antibody.

Dilution : 1:1000



### GTX24819 IHC-P Image

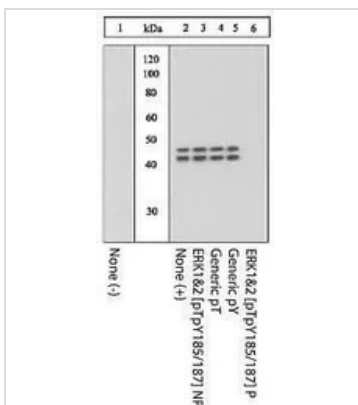
IHC-P analysis of human breast carcinoma tissue using GTX24819 ERK1 (phospho Thr202/Tyr204) + ERK2 (phospho Thr185/Tyr187) antibody.

Right : Primary antibody

Left : Negative control without primary antibody

Antigen retrieval : 10mM sodium citrate (pH 6.0), microwaved for 8-15 min

Dilution : 1:50



### GTX24819 WB Image

WB (peptide competition) analysis of PC12 cells stimulated with 0.5 M sorbitol for 5 minutes (Lane 2-6) using GTX24819 ERK1 (phospho Thr202/Tyr204) + ERK2 (phospho Thr185/Tyr187) antibody prior incubated with the non-phosphopeptide corresponding to the phosphopeptide immunogen (Lane 3), a generic phosphothreonine-containing peptide (Lane 4), a generic phosphotyrosine-containing peptide (Lane 5), or the phosphopeptide immunogen (Lane 6). The data show that only the immunogen phosphopeptide blocks the signal, demonstrating the specificity of the antibody.



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