

# Phospho-p44/42 MAPK (Erk1/2) (Thr202/Tyr204) (A11) rabbit mAb SureLight 488 conjugate

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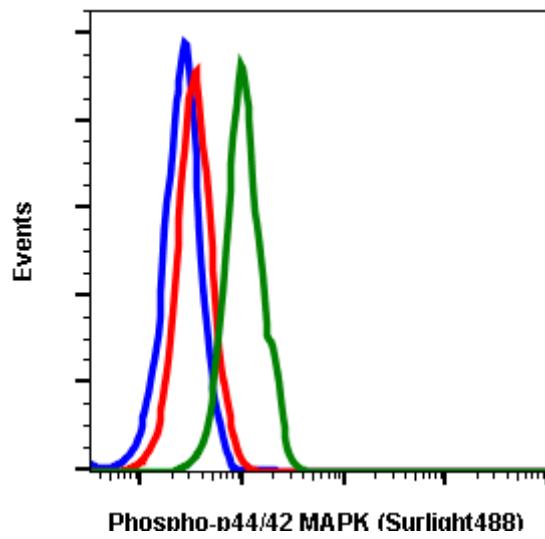
Catalog: #1115

Store at: 2-8°C

*For Research Use Only. Not For Use In Diagnostic Procedures.*

Applications	Detection	Clonality	Isotype
Flow Cytometry	Anti-Rabbit IgG	Monoclonal	Rabbit IgGκ

<b>Format:</b>	SureLight 488
<b>Cross Reactivity:</b>	Predicted to work with mouse, rat, and other homologues.
<b>Formulation:</b>	PBS, 0.09% NaN <sub>3</sub> , 0.2% BSA
<b>Preparation:</b>	Protein A
<b>Reactivity:</b>	Human, Mouse
<b>Recommended Usage:</b>	For flow cytometric staining, the suggested use of this reagent is 5 µL per million cells or 5 µL per 100 µL of staining volume. It is recommended that the reagent be titrated for optimal performance for each application.
<b>Immunogen:</b>	A synthetic phospho-peptide corresponding to residues surrounding Thr202/Tyr204 of human phospho Erk1/2.
<b>Description:</b>	Human Erk1 and Erk2 Ser/Thr kinases share 84% sequence identity and nearly all functions. These MAP kinases are activated in response to mitogens and growth factors as part of the Ras-Raf-MEK-ERK signal transduction cascade(1-3). This pathway regulates cell survival, differentiation, adhesion, cell cycle progression, and many other cellular processes. Upon phosphorylation, Erk1/2 translocate to the nucleus to activate transcription factors including c-Fos, Elk1, Ets1, and SP-1 (4,5). There are more than 175 known cytoplasmic and nuclear substrates of Erk1/2. The Erk1/2 cascade is upregulated in many human cancers, even when oncogenic mutations are not found. Multiple small-molecule inhibitors of Erk1/2 have been developed, including ones targeting the ATP-binding site either competitively or irreversibly (6).
<b>References:</b>	<ol style="list-style-type: none"><li>1. Blagoev B, et al., 2003, Nat Biotechnol, 21:315-318.</li><li>2. Thelemann A. et al, 2005, Mol Cell Proteomics 4:356-376.</li><li>3. Morandell S., et al., 2008, Proteomics 8:4383-4401.</li><li>4. Ramos J.W., 2008, Biochem Cell Biol 40:2707-2719.</li><li>5. Nakano H., et al., 1998, Proc Natl Acad Sci U S A. 104:19837-19842.</li><li>6. Roskoski Jr R. 2012, Pharmacol Res 66:105-143.</li></ol>



Flow cytometric analysis of Jurkat cells secondary antibody only negative control (blue) or treated with U0126 (red) or treated with TPA (green) using Phospho-ERK1/2 (Thr202/Tyr204) antibody ERK12T202Y204-A11 SureLight®488. Cat. #1115.