

Product Data Sheet: Phospho-ERK1/ERK2 (p44/p42 MAPK) (T202/Y204)

Cat. No: AB-10762-A
Conjugate: Unconjugated

Size: 100 ug
Clone: Poly

Concentration: 1mg/ml

Host: Rb
Isotype: IgG

Immunogen: Peptide derived from the protein area including conserved pT-E-pY motif of activated

Erk 1,2

Reactivity: Hu, Ms, Rt

Western blotting: 1:2000 Immunohistochemistry: 1:100-500 Immunofluorescence: 1:100-500 Immunoprecipitation: to be determined Immunocytochemistry: 1:100

Applications: Dilutions listed only as a recommendation. Optimal dilution should be determined by

investigator. Recognizes ERK1 and ERK2 dually phosphorylated at T202/Y204 and

T185/Y187, respectively.

Form: liquid

Buffer:

Storage:

Western immunoblotting solutions: - Wash buffer: 1x Tris Buffered Saline (TBS); 0.1%

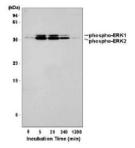
Triton X 100 - Blocking buffer: 1xTBS; 0.1% Triton X-100; 5% BSA (used with the

primary antibody)

The antibody can be stored at 2° - 8° C for 1 month without detectable loss of activity. Antibody can also be aliquotted and stored frozen at -20° C to -70° C in a

manual defrost freezer for six months without detectable loss of activity. Avoid

repeated freeze-thaw cycles.



Detection of ERK1 and ERK2 phosphorylated at T202/Y204 and T185/Y187, respectively. Human HeLa cells were incubated with 200 nM PMA for the indicated times. Total cell lysates in gel sample buffer were resolved by SDS-PAGE, transferred to Immobilon membrane and Immunoblotted with 0.5µg/Ml antiphosphERK1/2.



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WESTERN BLOT (WB) PROTOCOL

Western immunoblotting solutions:

Wash buffer: 1x Tris Buffered Saline (TBS); 0.1% Triton X 100 -

Blocking buffer: 1xTBS; 0.1% Triton X-100; 5% BSA (used with the primary antibody)

For western blots , incubate the membrane with antibody diluted in blocking buffer 2 hours at room temperature

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