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<b>Product name:</b>	Phospho-ERK1/ERK2 (p44/p42 MAPK) (T202/Y204)
<b>Cat number:</b>	MAB-91982
<b>Conjugate:</b>	Unconjugated
<b>Size:</b>	100 ug
<b>Clone:</b>	E10A
<b>Concentration:</b>	1mg/ml
<b>Host:</b>	Rb
<b>Isotype:</b>	IgG
<b>Immunogen:</b>	Synthetic phosphopeptide corresponding to residues surrounding T202/Y204 of human, mouse, and rat extracellular signal- regulated kinase-1.
<b>Reactivity:</b>	Hu, Ms, Rt
<b>Applications:</b>	Western Blot: 1:1000 IHC: 1:100 - 1:200 ICC: 1:100 IF: 1:100 IP: To be tested by the end-user
<b>Form:</b>	liquid
<b>Storage:</b>	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. The antibody can be stored at 2° - 8° C for 1 month without detectable loss of activity. Avoid repeated freeze-thaw cycles.

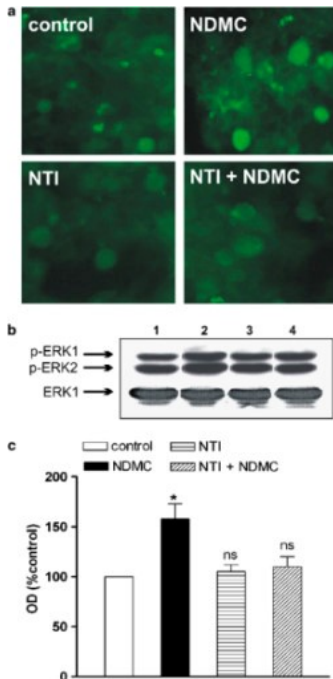


Image: Stimulation of ERK1/2 phosphorylation by NDMC in NG108-15 cells. (a) Immunofluorescence analysis of phospho-ERK1/2 immunoreactivity. Cells were serum-starved for 12 h and then treated with either vehicle (control), 10 M NDMC, 1 M NTI, and NTI+NDMC for 20 min, fixed and immunostained with anti-phospho-ERK1/2 antibody followed by FITC-conjugated secondary antibody. Results are representative of three similar experiments. (b) Western blot analysis of phospho-ERK1/2 immunoreactivity. Serum-starved cells were treated for 20 min with vehicle (lane 1), 10 M NDMC (lane 2), 1 M NTI (lane 3), and NTI+NDMC (lane 4). Thereafter, cell extracts were prepared and equal amounts of proteins (30 g) were loaded in each lane. Samples were subjected to immunoblotting with either anti-phospho-ERK1/2 antibody (top) or anti-ERK1 antibody (bottom). Results are representative of three similar experiments. (c) Densitometric analysis of immunoreactive phospho-ERK1/2 bands. The optical density of the phospho-ERK1/2 bands for each drug treatment was normalized to the density of the corresponding ERK1 band and is reported as percent of control. Values are the mean SEM of three experiments. \* $p < 0.05$  vs control, NS  $p > 0.05$  vs control by one-way ANOVA followed by Dunnett's test.