

Product Datasheet

NIH/3T3 Whole Cell Lysate PDGF Stimulated (orb348723)



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Descriptionnts. NIH/3T3 Whole Cell Lysate PDGF

Conjugation Unconjugated

Tested Applications

SDS-PAGE, WB

Preservatives Preservative: None. Stabilizer: 10%

(v/v) Glycerol. 1X SDS-PAGE Sample Buffer (62.5 mM Tris HCl, 2% SDS,

10% Glycerol and 0.005% bromophenol blue, pH 6.8)

Form/Appearance Liquid (sterile filtered)

Concentration 1.0 mg/ml

Storage Store vial at -70° C or COLDER. For

extended storage, aliquot contents to

minimize freeze/thaw cycles.

Note For research use only

Application notes ready-to-use lysates are especially

prepared as positive controls for separation by SDS-PAGE and subsequent western blot analysis. Lysates are prepared in denaturing buffer WITHOUT dissociating agents

(i.e. no 2-mercaptoethanol or dithiothreitol has been added). Heat lysate to 95°C for 5 minutes and rapidly cool. If dissociating conditions are desired, add reducing agent prior to heating. The recommended loading volume per lane is 10-20 µl depending

on the size format of your gel.

Purity The cells were grown in Dulbecco's

medium supplemented with 10% fetal bovine serum. Cells were washed with PBS and then incubated on ice in modified RIPA buffer, containing 150 mM sodium chloride, 50 mM Tris HCl, pH 7.4, 1 mM EDTA, 1.0% NP-40, 0.5% sodium deoxycholic acid , 0.1% SDS and 0.01% (w/v) sodium azide to lyse the cells. Protein integrity was

ensured using a cocktail of protease inhibitors with broad specificity for the inhibition of aspartic, cysteine, and

serine proteases as well as

aminopeptidases (0.1 mM AEBSF HCl,

0.00 vM Assotisis F vM Bostotis 1 F

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