

## **Product Datasheet**

## Normal Mouse Spleen Whole Cell Lysate (orb348720)



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**Description**nts. Normal Mouse Spleen Whole Cell

**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** Tissues were washed exhaustively with

PBS to remove blood and other debris.

A lysate was prepared by

homogenizing the tissue and washing the cells in cold PBS. Washed cells were incubated at 4° C in modified RIPA buffer containing 150 mM sodium chloride, 50 mM Tris Cl, pH 7.4, 1 mM EDTA, 1.0% NP-40, 0.5% sodium deoxycholic acid and 0.1% SDS to lyse the cells. Protein integrity is ensured using a cocktail of protease inhibitors with broad specificity for the inhibition of aspartic, cysteine, and serine proteases as well as aminopeptidases

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