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Product datasheet

Jurkat whole cell lysate ab7899

3 References

概述

产品名称 Jurkat全细胞裂解物

常规说明 Cell line: Jurkat, clone E6-1 (Human acute T cell leukemia).

Growth media: RPMI and 10% NCS (Newborn calf serum).

Jurkat cell lysate was prepared by homogenization in modified RIPA buffer (150 mM sodiumchloride, 50 mM Tris-HCI, pH 7.4, 1 mM ethylenediaminetetraacetic acid, 1 mM phenylmethylsulfonyl flouride, 1% Triton X-100, 1% sodium deoxycholic acid, 0.1% sodium dodecylsulfate, 5 μ g/ml of aprotinin, 5 μ g/ml of leupeptin). Cell debris was removed by

centrifugation. Protein concentration was determined with Bio-Rad protein assay. The cell lysate was boiled for 5 min in 1 x SDS sample buffer (0.045 M Tris-HCl pH 6.8, 10% glycerol, 1%

sodium dodecylsulfate, 0.01% bromophenol blue), containing 0.05 M DTT.

经测试应用 适用于: WB

性能

Mycoplasma free Yes

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot. Store at -80°C. Avoid freeze / thaw cycle.

存储溶液 pH: 7.20

Constituent: 100% SDS Sample Buffer

裂解物说明 Jurkat cell lysate was prepared by homogenization in modified RIPA buffer (150 mM

sodiumchloride, 50 mM Tris-HCl, pH 7.4, 1 mM ethylenediaminetetraacetic acid, 1 mM phenylmethylsulfonyl flouride, 1% Triton X-100, 1% sodium deoxycholic acid, 0.1% sodium dodecylsulfate, 5 µg/ml of aprotinin, 5 µg/ml of leupeptin). Cell debris was removed by

centrifugation. Protein concentration was determined with Bio-Rad protein assay. The cell lysate was boiled for 5 min in 1 x SDS sample buffer (50 mM Tris-HCl pH 6.8, 12.5% glycerol, 1%

sodium dodecylsulfate, 0.01% bromophenol blue) containing 5% b-mercaptoethanol.

背景 Often this cell line is called 'JM' (Jurkat and JM are derived from the same patient and are sister

clones), occasionally JM may be a subclone with somewhat divergent features. Jurkat cells can be transfected and are therefore useful for studies of blood proto-oncogenes expression, apoptosis and cell survival (e.g HIV). They produce IL-2 and can also be used for studies of

differentiation.

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应用	Ab评论	说明
WB		Use at an assay dependent dilution. Jurkat cell lysate is ready to load on SDS-PAGE for Western blotting, 20 µg per lane is recommended for mini gel.

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