

## **Product Datasheet**

Beta galactosidase antibody (Biotin) (orb344930)



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Western

Blot of Rabbit anti-

Beta

Galacto...

**Description**nts. Beta galactosidase antibody (Biotin)

Species/Host Rabbit

**Conjugation** Biotin

**Tested** ELISA, IHC, WB **Applications** 

Immunogen Beta Galactosidase (E.coli)

**Preservatives** 0.01% (w/v) Sodium Azide

Form/Appearance Lyophilized

**Concentration** 1.0 mg/mL

**Storage** Store vial at 4° C prior to restoration. For extended

storage aliquot contents and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid.

Dilute only prior to immediate use.

**Note** For research use only

**Application notes** Anti-Beta Galactosidase Biotin Conjugated Antibody

has been tested by Western blot and is suitable for ELISA, immunohistochemistry, immunomicroscopy as

well as other antibody based assays using

streptavidin or avidin conjugates requiring lot-to-lot consistency. The antibody recognizes both frozen tissue sections, paraffin embedded tissue and 4%

paraformaldehyde fixed tissue for most

immunohistochemical analysis. A 1:5,000 dilution has

been reported to be successful for staining by immunoblot of beta-galactosidase fusion proteins after transfer using a semi-dry transfer apparatus. A 1:1,500 dilution has been reported to detect beta-galactosidase in adult rat spinal cord tissue after infection with helper-dependent adenovirus expressing lacZ. In this particular experiment, tissue was perfused with 4% paraformaldehyde and

cryostat-cut (35 µm) to produce free-floating sections. A 1:5,000 dilution has been reported to be successful for staining of brain sections from

transgenic mice expressing nuclear betagalactosidase when assayed by immunofluorescence

microscopy. A 1:5,000 dilution has been reported for immunofluorescent staining of methanol fixed, devitellinized Drosophila embryos. Although a wide range of conditions was reported to be effective, a 1:10,000 dilution was noted to show no background

1:10,000 dilution was noted to show no background and to be suitable for double labeling experiments.

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