

Product datasheet for AM08430PU-N

OriGene Technologies, Inc.

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GST-Tag Mouse Monoclonal Antibody [Clone ID: 3D4]

Product data:

Product Type: Primary Antibodies

Clone Name: 3D4

Applications: ELISA, IHC, WB

Recommended Dilution: Suitable for most immunological techniques requiring high titer binding and lot-to-lot

consistency.

Recommended Dilutions: ELISA: 1/20,000-1/100,000 Western Blot: 1/1,000-1/10,000

Host: Mouse

Isotype: IgG1

Clonality: Monoclonal

Immunogen: Glutathione-S-Transferase [Schistosoma japonicum]

Specificity: Reacts specifically with GST. Cross reactivity against Glutathione-S-Transferase from other

sources may occur but has not been specifically determined.

Formulation: 0.02M Potassium Phosphate, 0.15M Sodium Chloride, pH 7.2 as buffer and 0.01% (w/v) Sodium

Azide as preservative.

State: Purified

State: Liquid (sterile filtered) purified Ig fraction.

Concentration: lot specific

Purification: Protein A Chromatography.

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C or below for

longer.

Avoid repeated freezing and thawing.

Should this product contain a precipitate we recommend microcentrifugation before use.

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Stability: Shelf life: one year from despatch.

Database Link: P08515

Background: GST (Glutathione-S-Transferase) is a protein expression tag commonly used in molecular

biology. Anti-GST will react with synthetic construct present in most known GST containing cloning or expression vectors. GST is responsible for the conjugation of reduced glutathione to a wide number of exogenous and endogenous hydrophobic electrophiles. The amino acid sequence GST is highly conserved in most organisms including mammals. GST exists as a 26

kDa homodimer.

Synonyms: Glutathione S-transferase Tag, GST26-Tag

Product images:



Western blot using Affinity Purified anti-GST antibody to detect purified GST recombinant protein (Left) and in a lysate of HeLa cells expressing recombinant GST protein (Right). Approximately 0.1 g of protein was separated by SDS-PAGE and transferred onto