

Anti-PCK2/PEPCK-M antibody (Internal) (STJ70781)

STJ70781

GENERAL INFORMATION

Product Type Primary antibodies

Short Goat polyclonal antibody anti-PCK2/PEPCK-M (Internal) is suitable for use in ELISA, Western Blot, Immunofluorescence and Flow

Description Cytometry research applications.

Applications Pep-ELISA/WB/IF/FC

Host/Source Goat

Reactivity Human/Mouse/Rat/Dog/Pig/Cow

PRODUCT PROPERTIES

Clonality Polyclonal

Clone ID

Concentration 0.5 mg/mL **Conjugation** Unconjugated

Purification Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing

peptide

Dilution Peptide ELISA: antibody detection limit dilution 1:4000.

Range WB: Approx 70kDa band observed in lysates of cell lines A431, HEK293 HepG2 and NIH3T3 (calculated MW of 70.7kDa according to

Human NP_004554.3 and 70.5kDa according to Mouse NP_083270.2). R

 $\textbf{Formulation} \quad 0.5 \text{ mg/ml in Tris saline, } 0.02\% \text{ sodium azide, pH7.3 with } 0.5\% \text{ bovine serum albumin. NA}$

Isotype IgG

Storage Store at-20°C on receipt and minimise freeze-thaw cycles.

Instruction

TARGET INFORMATION

Gene ID 5106

Gene Symbol PCK2

Uniprot ID PCKGM_HUMAN

Immunogen

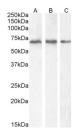
Immunogen Internal

Region

Specificity No cross-reactivity with PCK1.

Immunogen KPWKPGDKEPCAH

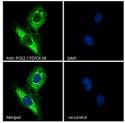
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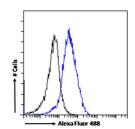
STJ70781 (2µg/ml) staining of A431 (A) , HEK293 (E and (1ug/ml) HepG2 (C) cell lysate (35µg protein in RIPA buffer). Detected by chemiluminescence.



STJ70781 (2ŵg/ml) staining of NIH3T3 cell lysate 35ŵg protein in RIPA buffer). Detected by hemiluminescence.



STJ70781 Immunofluorescence analysis oparaformaldefyde fixed U2OS cells, permeabilized wil 0. 15% Triton. Primary incubation 1hr (10ugm followed by Alexa Fluor 488 secondary antibor (2ug/mi), showing cytoplasmic staining. The nucles stain is DAP (blue). Negative control: Unimmunize goat IgG (10ugm) followed by Alexa Fluor 48 secondary antibody (2ug/mi).



STJ7081 Flow cytometric analysis o paraformaldehyde fixed MCF7 cells (blue line) permeabilized with 0. 5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.