Anti-DCX antibody

Catalog Number: 100526



Product name

Anti-DCX antibody

Immunogen

<u>Human DCX (aa 45-150, GST Tag) recombinant</u> <u>protein</u>

Specificity

Human DCX

Antibody description

Mouse monoclonal to DCX

Preparation

This antibody was produced from a hybridoma resulting from the fusion of a mouse myeloma with B cells obtained from a mouse immunized with purified, recombinant Human DCX (rh DCX; O43602-2; Ala45-Val150). The IgG fraction of the cell culture supernatant was purified by Protein A affinity chromatography.

Formulation

 $0.2 \mu m$ filtered solution in PBS

Storage

This antibody can be stored at $2^{\circ}\text{C-8}^{\circ}\text{C}$ for one month without detectable loss of activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C .

Preservative-Free.

Sodium azide is recommended to avoid contamination (final concentration 0.05%-0.1%). It is toxic to cells and should be disposed of properly. Avoid repeated freeze-thaw cycles.

Clonality

Monoclonal

Ig Type

Mouse IgG2a

Applications

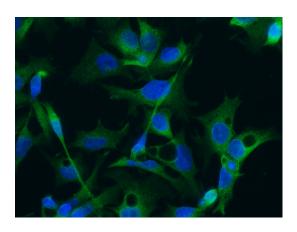
FCM, IF, ICC/IF

Dilutions

FCM: 0.5-2 µg/Test

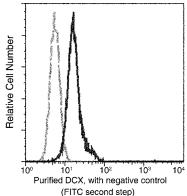
ICC/IF: $10-25 \mu g/mL$

Validations



DCX Antibody, Mouse MAb, Immunofluorescence

Immunofluorescence staining of Human DCX in HepG2 cells. Cells were fixed with 4% PFA, permeabilzed with 0.3% Triton X-100 in PBS, blocked with 10% serum, and incubated with Mouse anti-Human DCX monoclonal antibody (15 µg/ml) at 37°C 1 hour. Then cells were stained with the Alexa Fluor® 488-conjugated Goat Antimouse IgG secondary antibody (green) and counterstained with DAPI (blue). Positive staining was localized to cytoplasm.



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DCX Antibody, Mouse MAb

Flow cytometric analysis of Human DCX expression on HepG2 cells. The cells were treated according to manufacturer's manual (BD Pharmingen $^{\text{TM}}$ Cat. No. 554714), stained with

purified anti-Human DCX, then a FITC-conjugated second step antibody. The fluorescence histograms were derived from gated events with the forward and side light-scatter characteristics of intact cells.