

# Anti-DCX antibody



Catalog Number: 100526

## Product name

Anti-DCX antibody

## Immunogen

[Human DCX \(aa 45-150, GST Tag\) recombinant protein](#)

## 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°C-8°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

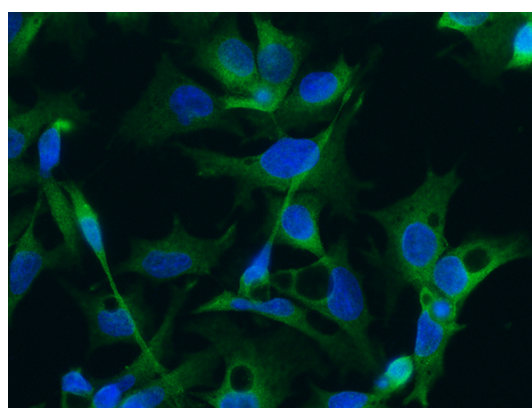
FCM, IF, ICC/IF

## Dilutions

FCM: 0.5-2  $\mu$ g/Test

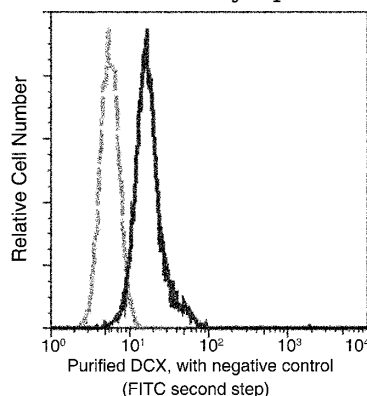
ICC/IF: 10-25  $\mu$ g/mL

## Validations



DCX Antibody, Mouse MAb, Immunofluorescence

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



## 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™ 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.