

Product datasheet

Anti-DUSP22 antibody ab70124

画像数 2

製品の概要

製品名	Anti-DUSP22 antibody
製品の詳細	Rabbit polyclonal to DUSP22
特異性	ab70124 detects endogenous levels of total DUSP22 protein.
アプリケーション	適用あり: ICC/IF, WB, ELISA
種交差性	交差種: Mouse, Human
免疫原	Synthetic peptide derived from an internal sequence of human DUSP22.
ポジティブ・コントロール	Jurkat and RAW264.7 cell extracts.

製品の特性

製品の状態	Liquid
保存方法	Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.
バッファー	Preservative: 0.02% Sodium Azide Constituents: 50% Glycerol, PBS (without Mg ²⁺ and Ca ²⁺), 150mM Sodium chloride, pH 7.4
精製度	Immunogen affinity purified
特記事項 (精製)	ab70124 was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
ポリ/モノ	ポリクローナル
アイソタイプ	IgG

アプリケーション

Our [Abpromise guarantee](#) covers the use of **ab70124** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

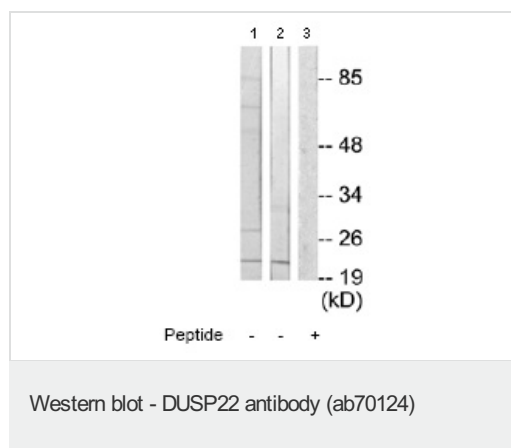
アプリケーション	Abreviews	特記事項
ICC/IF		Use a concentration of 1 µg/ml.
WB		1/500 - 1/1000. Detects a band of approximately 20 kDa (predicted molecular weight: 20 kDa).

アプリケーション	Abreviews	特記事項
ELISA		1/5000.

ターゲット情報

機能	Activates the Jnk signaling pathway. Dephosphorylates and deactivates p38 and stress-activated protein kinase/c-Jun N-terminal kinase (SAPK/JNK).
組織特異性	Ubiquitous. Highest expression seen in heart, placenta, lung, liver, kidney and pancreas.
配列類似性	Belongs to the protein-tyrosine phosphatase family. Non-receptor class dual specificity subfamily.
	Contains 1 tyrosine-protein phosphatase domain.
細胞内局在	Cytoplasm. Nucleus.

画像



All lanes : Anti-DUSP22 antibody (ab70124)
at 1/500 dilution

Lane 1 : Jurkat cell extracts

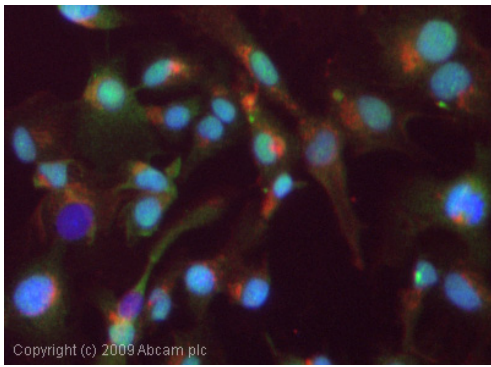
Lane 2 : RAW264.7 cell extracts

Lane 3 : RAW264.7 cell extracts with
immunising peptide at 10 µg

Lysates/proteins at 30 µg per lane.

Predicted band size : 20 kDa

Observed band size : 20 kDa



Immunocytochemistry/ Immunofluorescence-
DUSP22 antibody(ab70124)

ICC/IF image of ab70124 stained HepG2 cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab70124, 1 µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

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