

Eff. Date: 1 June 2023

Version: 2.1

IFU: Caldesmon ILM8029

Caldesmon (HMW) clone h-CALD

Instructions for Use

Specification:

Recognizes a protein of 150kDa, which is identified as the high molecular weight variant of Caldesmon. Two closely related variants of human caldesmon have been identified which are different in their electrophoretic mobility and cellular distribution. The h-caldesmon variant (120–150kDa) is predominantly expressed in smooth muscle whereas l-caldesmon (70–80kDa) is found in non- muscle tissue and cells. Neither of the two variants has been detected in skeletal muscle. This MAb recognizes only the 150kDa variant (h-caldesmon) in Western blots of human aortic media extracts and is unreactive with fibroblast extracts from cultivated human foreskin. Caldesmon is a developmentally regulated protein involved in smooth muscle and non-muscle contraction.

Availability:

Catalog No.ContentsVolumeILM8029-C01Caldesmon0,1 ml concentrateILM8029-C05Caldesmon0,5 ml concentrateILM8029-C1Caldesmon1,0 ml concentrate

Intended use: For Research Use Only

Reactivity: Human

Clone: h-CALD

Species of origin: Mouse

Isotype: IgG, kappa

Control Tissue: Uterus, Blood vessels in all tissues, smooth muscle or leiomyosarcoma

Staining: Cytoplasmic

Immunogen: Crude human uterus extract

Presentation: Bioreactor Concentrate with 0.05% Azide

Application and suggested dilutions:

Pretreatment: Heat induced epitope retrieval in 10 mM citrate buffer, pH6.0, for 20 minutes is required for IHC staining on formalin-fixed, paraffin embedded tissue sections.

 Immunohistochemical staining of formalin-fixed, paraffin embedded tissue section (dilution up to 1:200 / 1:400)

The optimal dilution for a specific application should be determined by the investigator.

Note: Dilution of the antibody in 10% normal goat serum followed by a goat anti-mouse secondary antibody-based detection is recommended.

Storage & Stability: Store at 2-8 °C. Do not Freeze. Do not use after expiration date printed on the vial.

References:

1) Frid MG, et al. Dev Biol 1992; 153:185





