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Product datasheet

HRP Anti-Cardiac Troponin I antibody [16A11] ab24460

1 References

製品の概要

製品名 HRP Anti-Cardiac Troponin I antibody [16A11]

製品の詳細 HRP Mouse monoclonal [16A11] to Cardiac Troponin I

由来種 Mouse 標識 HRP

特異性 Reacts with free cardiac troponin I (cTnI) and cTnI forming complexes with other troponin

components (in the presence of 5 mM EDTA). Not affected by heparin, phosphorylation, oxidation

and troponin complex formation. Does not cross-react with skeletal muscle troponin I.

アプリケーション 適用あり: WB, ELISA, Sandwich ELISA

種交差性 交差種: Rat, Rabbit, Goat, Cow, Cat, Dog, Human, Pig

非交差種: Fish

免疫原 Full length native protein (purified) corresponding to Human Cardiac Troponin I.

エピトープ 87-91aa

特記事項The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C.

バッファー pH: 7.40

Preservative: 0.05% Proclin 300

Constituents: 0.8% Sodium chloride, 0.02% Potassium chloride, 0.18% Dibasic monohydrogen

sodium phosphate, 0.024% Monobasic dihydrogen potassium phosphate

精製度 Protein A purified

ポリ/モノ モノクローナル

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クローン名16A11ミエローマSp2/0アイソタイプIgG1

アプリケーション

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アプリケーション	Abreviews	特記事項
WB		Use at an assay dependent concentration.
ELISA		Use at an assay dependent concentration.
Sandwich ELISA		Use at an assay dependent concentration.
AP		Use at an assay dependent concentration.

ターゲット情報

機能

関連疾患

Troponin I is the inhibitory subunit of troponin, the thin filament regulatory complex which confers calcium-sensitivity to striated muscle actomyosin ATPase activity.

Defects in TNNI3 are the cause of cardiomyopathy familial hypertrophic type 7 (CMH7)

[MIM:613690]. Familial hypertrophic cardiomyopathy is a hereditary heart disorder characterized by ventricular hypertrophy, which is usually asymmetric and often involves the interventricular septum. The symptoms include dyspnea, syncope, collapse, palpitations, and chest pain. They can be readily provoked by exercise. The disorder has inter- and intrafamilial variability ranging from benign to malignant forms with high risk of cardiac failure and sudden cardiac death. Defects in TNNI3 are the cause of cardiomyopathy familial restrictive type 1 (RCM1) [MIM:115210]. RCM1 is an heart muscle disorder characterized by impaired filling of the ventricles with reduced diastolic volume, in the presence of normal or near normal wall thickness

and systolic function.

Defects in TNNI3 are the cause of cardiomyopathy dilated type 2A (CMD2A) [MIM:611880]. Dilated cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature death.

Defects in TNNI3 are the cause of cardiomyopathy dilated type 1FF (CMD1FF) [MIM:613286]. Dilated cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature death.

配列類似性

Belongs to the troponin I family.

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