abcam

Product datasheet

Anti-beta Actin antibody [EPR21242] ab241153





4 图像

概述

产品名称 Anti-beta Actin抗体[EPR21242]

描述 兔单克隆抗体[EPR21242] to beta Actin

宿主 Rabbit

特异性 In-house testing showed ab241153 to react with gamma actin in western blot.

经测试应用 适用于: WB, ICC/IF

种属反应性 与反应: Mouse, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

阳性对照 ICC/IF: Methanol-fixed HeLa and NIH3T3 cells. WB: HAP1, MEF1 and NIH3T3 whole cell lysates

常规说明 This product was made using synthetic libraries and phage display technology.

This antibody is a recombinant chimeric antibody. Rabbit chimeric monoclonal antibody (Human

Fab/ Rabbit Fc).

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

存储溶液 Preservative: 0.02% Sodium azide

Constituents: PBS, 1% BSA

克隆 单克隆

克隆编号 EPR21242

同种型 lgG1

应用

The Abpromise guarantee Abpromise™承诺保证使用ab241153于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度:实际最佳的稀释度/浓度应由使用者检定。

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应 用	Ab评论	说 明
WB		Use a concentration of 1 µg/ml. Predicted molecular weight: 42 kDa.
ICC/IF		Use at an assay dependent concentration.

靶标

功能 Actins are highly conserved proteins that are involved in various types of cell motility and are

ubiquitously expressed in all eukaryotic cells.

疾病相关 Defects in ACTB are a cause of dystonia juvenile-onset (DYTJ) [MIM:607371]. DYTJ is a form of

dystonia with juvenile onset. Dystonia is defined by the presence of sustained involuntary muscle contraction, often leading to abnormal postures. DYTJ patients manifest progressive, generalized,

dopa-unresponsive dystonia, developmental malformations and sensory hearing loss.

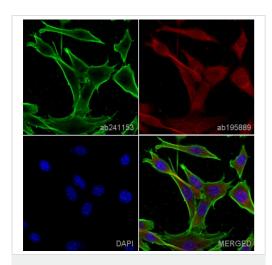
序列相似性 Belongs to the actin family.

翻译后修饰 ISGylated.

细胞定位 Cytoplasm > cytoskeleton. Localized in cytoplasmic mRNP granules containing untranslated

mRNAs.

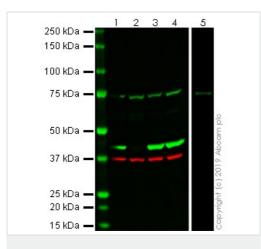
图片



Immunocytochemistry/ Immunofluorescence - Antibeta Actin antibody [EPR21242] (ab241153)

ab213243 staining beta-Actin in NIH3T3 cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab241153 at 0.5ugml then detected with an Alexa Fluor® 488 goat anti-rabbit secondary antibody (ab150081) at a 1/1000 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue), and ab195889, Mouse monoclonal to alpha Tubulin (Alexa Fluor® 594), at a 1/250 dilution (shown in red).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



Western blot - Anti-beta Actin antibody [EPR21242] (ab241153)

All lanes:

Lane 1: HAP1 whole cell lysate at 20 µg

Lane 2: HAP1 ACTNB whole cell lysate at 20 µg

Lane 3: NIH3T3 whole cell lysate at 20 µg

Lane 4: MEF whole cell lysate at 20 µg

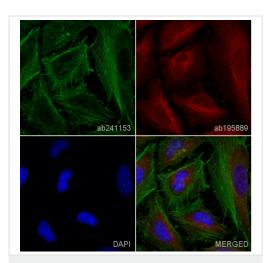
Lane 5: Recombinant Human gamma Actin protein (ab157841)

Performed under reducing conditions.

Predicted band size: 42 kDa

ab241153 was shown to react with ACTNB (beta actin) when tested against HAP1 wild-type and HAP1 ACTNB (beta actin) knockout cells. ACTNB wild-type and knockout samples were subjected to SDS-PAGE and incubated with ab231153. Incomplete reduction of signal was seen in the ACTNB (beta actin) knockout samples at beta actin's expected molecular weight. SDS-PAGE of gamma actin recombinant protein and incubation with ab241153 showed cross-reactivity to gamma actin.

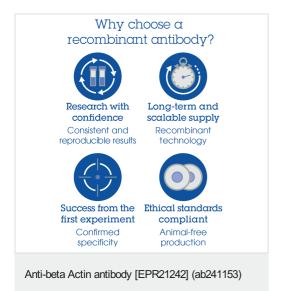
ab241153 and <u>ab8245</u> (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at a 1ug/ml and 1/10000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Antibeta Actin antibody [EPR21242] (ab241153)

ab213243 staining beta-Actin in HeLa cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab241153 at 0.01ugml then detected with an Alexa Fluor® 488 goat anti-rabbit secondary antibody (ab150081) at a 1/1000 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue), and ab195889, Mouse monoclonal to alpha Tubulin (Alexa Fluor® 594), at a 1/250 dilution (shown in red).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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