abcam

Product datasheet

HRP Anti-MEK2 antibody [Y78] ab200607





3 Images

Overview

Product name HRP Anti-MEK2 antibody [Y78]

Description HRP Rabbit monoclonal [Y78] to MEK2

Host species Rabbit HRP Conjugation

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Predicted to work with: Mouse

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

Positive control WB: Jurkat whole cell lysate.

General notes Our RabMAb® technology is a patented hybridoma-based technology for making rabbit

monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

Properties

Form Liquid

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Storage instructions

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer

Preservative: 0.1% Proclin 300 Solution

Constituents: PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

Purity Protein A purified

Clonality Monoclonal

Y78 Clone number Isotype ΙgG

Applications

Our **Abpromise guarantee** covers the use of ab200607 in the following tested applications. The Abpromise guarantee

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

Application	Abreviews	Notes
WB		1/7500. Detects a band of approximately 45 kDa (predicted molecular weight: 44 kDa).

Target

Function

Catalyzes the concomitant phosphorylation of a threonine and a tyrosine residue in a Thr-Glu-Tyr sequence located in MAP kinases. Activates the ERK1 and ERK2 MAP kinases.

Involvement in disease

Defects in MAP2K2 are a cause of cardiofaciocutaneous syndrome (CFC syndrome) [MIM:115150]; also known as cardio-facio-cutaneous syndrome. CFC syndrome is characterized by a distinctive facial appearance, heart defects and mental retardation. Heart defects include pulmonic stenosis, atrial septal defects and hypertrophic cardiomyopathy. Some affected individuals present with ectodermal abnormalities such as sparse, friable hair, hyperkeratotic skin lesions and a generalized ichthyosis-like condition. Typical facial features are similar to Noonan syndrome. They include high forehead with bitemporal constriction, hypoplastic supraorbital ridges, downslanting palpebral fissures, a depressed nasal bridge, and posteriorly angulated ears with prominent helices. The inheritance of CFC syndrome is autosomal dominant.

Sequence similarities

Belongs to the protein kinase superfamily. STE Ser/Thr protein kinase family. MAP kinase kinase

subfamily.

Contains 1 protein kinase domain.

Post-translational modifications

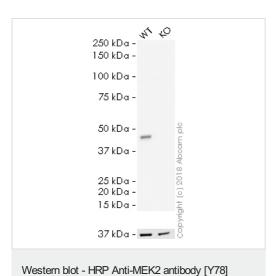
MAPKK is itself dependent on Ser/Thr phosphorylation for activity catalyzed by MAP kinase kinases (RAF or MEKK1).

Acetylation of Ser-222 and Ser-226 by Yersinia yopJ prevents phosphorylation and activation,

thus blocking the MAPK signaling pathway.

Images

(ab200607)



All lanes : HRP Anti-MEK2 antibody [Y78] (ab200607) at 1/7500 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: MAP2K2 (MEK2) knockout HAP1 whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 44 kDa **Observed band size:** 44 kDa

Exposure time: 30 seconds

ab200607 was shown to specifically react with MEK2 in wild-type HAP1 cells as signal was lost in MAP2K2 (MEK2) knockout cells. Wild-type and MAP2K2 (MEK2) knockout samples were subjected to SDS-PAGE. Ab200607 and ab184095 (Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control (Alexa Fluor 680) loading control) were incubated overnight at 4°C at 1/7500 dilution and 1/1000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.



Western blot - HRP Anti-MEK2 antibody [Y78] (ab200607)

HRP Anti-MEK2 antibody [Y78] (ab200607) at 1/7500 dilution + Jurkat (Human T cell lymphoblast-like cell line) Whole Cell Lysate at 10 μg

Developed using the ECL technique.

Performed under reducing conditions.

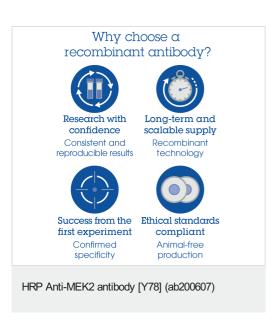
Predicted band size: 44 kDa **Observed band size:** 45 kDa

Additional bands at: 100 kDa. We are unsure as to the identity of

these extra bands.

Exposure time: 10 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab200607 overnight at 4°C. Antibody binding was visualised using ECL development solution ab133406.



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