abcam

Product datasheet

Anti-HUWE1/Mule antibody ab65153

4 Images

Overview

Product name Anti-HUWE1/Mule antibody

Description Rabbit polyclonal to HUWE1/Mule

Host species Rabbit

Specificity This antibody reacts with HUWE1/Mule.

Tested applications Suitable for: ICC/IF, WB, IHC-P

Species reactivity Reacts with: Rat, Human

Immunogen Synthetic peptide corresponding to HUWE1/Mule (C terminal). Peptide corresponding to 19

amino acids near the C-terminus of human HUWE1.

Database link: Q7Z6Z7

Positive control Brain, cortex; heart.

General notes

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

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C or -80°C. Avoid freeze /

thaw cycle.

Storage buffer pH: 7.4

Preservative: 0.02% Sodium azide

Constituent: PBS

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

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The Abpromise guarantee

Our Abpromise guarantee covers the use of ab65153 in the following tested applications.

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

Application	Abreviews	Notes
ICC/IF		Use a concentration of 1 µg/ml.
WB		Use a concentration of 1 - 2 μg/ml. Predicted molecular weight: 481 kDa.
IHC-P		Use a concentration of 5 µg/ml.

Target

Function

E3 ubiquitin-protein ligase which mediates ubiquitination and subsequent proteasomal degradation of target proteins. Regulates apoptosis by catalyzing the polyubiquitination and degradation of MCL1. Mediates monoubiquitination of DNA polymerase beta (POLB) at 'Lys-41', 'Lys-61' and 'Lys-81', thereby playing a role in base-excision repair. Also ubiquitinates the p53/TP53 tumor suppressor and core histones including H1, H2A, H2B, H3 and H4. Binds to an upstream initiator-like sequence in the preprodynorphin gene. Regulates neural differentiation and proliferation by catalyzing the polyubiquitination and degradation of MYCN. May regulate abundance of CDC6 after DNA damage by polyubiquitinating and targeting CDC6 to degradation.

Tissue specificity

Weakly expressed in heart, brain and placenta but not in other tissues. Expressed in a number of cell lines, predominantly in those from colorectal carcinomas.

Pathway

Protein modification; protein ubiquitination.

Involvement in disease

Defects in HUWE1 are the cause of mental retardation syndromic X-linked Turner type (MRXST) [MIM:300706]; also known as mental retardation and macrocephaly syndrome. MRXST shows clinical variability. Associated phenotypes include macrocephaly and variable contractures. A chromosomal microduplication involving HUWE1 and HSD17B10 is the cause of mental retardation X-linked type 17 (MRX17) [MIM:300705]; also known as mental retardation X-linked type 31 (MRX31). Mental retardation is characterized by significantly sub-average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. In contrast to syndromic or specific X-linked mental retardation which also present with associated physical, neurological and/or psychiatric manifestations, intellectual deficiency is the only primary symptom of non-syndromic X-linked mental retardation.

Sequence similarities

Belongs to the TOM1/PTR1 family.

Contains 1 HECT (E6AP-type E3 ubiquitin-protein ligase) domain.

Contains 1 UBA domain.

Contains 1 UIM (ubiquitin-interacting motif) repeat.

Contains 1 WWE domain.

Domain

The HECT domain mediates inhibition of the transcriptional activity of p53.

Post-translational modifications

Phosphorylated on tyrosine; phosphorylation is probably required for its ability to inhibit TP53

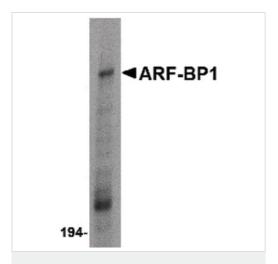
transactivation.

Phosphorylated upon DNA damage, probably by ATM or ATR.

Cellular localization

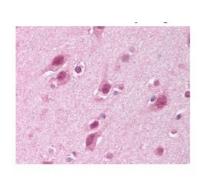
Cytoplasm. Nucleus. Mainly expressed in the cytoplasm of most tissues, except in the nucleus of

Images



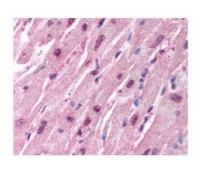
Western blot - Anti-HUWE1/Mule antibody (ab65153)

Western blot analysis using ab65153 in Daudi cell lysate at 1 ug/ml



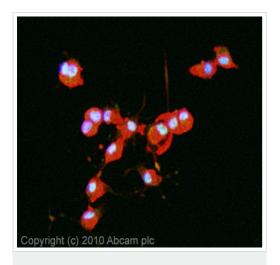
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HUWE1/Mule antibody (ab65153)

ab65153 at $5\mu g/ml$ staining HUWE1/Mule in human Brain, cortex tissue.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HUWE1/Mule antibody (ab65153)

Immunohistochemistry analysis of paraffin embedded formalin fixed human heart tissue using 5 μ g/ml ab65153.



Immunocytochemistry/ Immunofluorescence - Anti-HUWE1/Mule antibody (ab65153) ICC/IF image of ab65153 stained PC12 cells. The cells were 4% formaldehyde fixed (10 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 (ab65153, 1 μ g/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit lgG (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 1.43 μ M.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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