

Product datasheet

info@arigobio.com

ARG56053 anti-MITF antibody [D5]

Package: 50 μg Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [D5] recognizes MITF

Tested Reactivity Hu

Species Does Not React With Ms, Rat

Tested Application IHC-P

Host Mouse

Clonality Monoclonal

Clone D5

Isotype IgG1, kappa

Target Name MITF
Species Human

Immunogen An N-terminus fragment of Human MITF protein

Conjugation Un-conjugated

Alternate Names bHLHe32; Class E basic helix-loop-helix protein 32; MI; Microphthalmia-associated transcription factor;

WS2; WS2A; CMM8

Application Instructions

Application table	Application	Dilution
	IHC-P	1 - 2 μg/ml
Application Note	IHC-P: Antigen Retrieval: Boil tissue section in 10 mM Citrate buffer (pH 6.0) for 10-20 min, followed by cooling at RT for 20 min. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Liquid

Purification Purification with Protein G.

Buffer PBS (pH 7.4), 0.05% Sodium azide and 0.1 mg/ml BSA

Preservative 0.05% Sodium azide

Stabilizer 0.1 mg/ml BSA

Concentration 0.2 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

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before use.

Note

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Database links <u>GeneID: 4286 Human</u>

Swiss-port # O75030 Human

Gene Symbol MITF

Gene Full Name microphthalmia-associated transcription factor

Background This gene encodes a transcription factor that contains both basic helix-loop-helix and leucine zipper

structural features. It regulates the differentiation and development of melanocytes retinal pigment epithelium and is also responsible for pigment cell-specific transcription of the melanogenesis enzyme genes. Heterozygous mutations in the this gene cause auditory-pigmentary syndromes, such as Waardenburg syndrome type 2 and Tietz syndrome. Alternatively spliced transcript variants encoding

different isoforms have been identified. [provided by RefSeq, Jul 2008]

Function Transcription factor that regulates the expression of genes with essential roles in cell differentiation,

proliferation and survival. Binds to symmetrical DNA sequences (E-boxes) (5'-CACGTG-3') found in the promoters of target genes, such as BCL2 and tyrosinase (TYR). Plays an important role in melanocyte development by regulating the expression of tyrosinase (TYR) and tyrosinase-related protein 1 (TYRP1). Plays a critical role in the differentiation of various cell types, such as neural crest-derived melanocytes,

mast cells, osteoclasts and optic cup-derived retinal pigment epithelium. [UniProt]

Calculated Mw 59 kDa

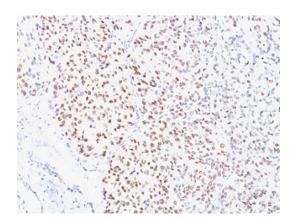
PTM Phosphorylation at Ser-405 significantly enhances the ability to bind the tyrosinase promoter.

Phosphorylated at Ser-180 and Ser-516 following KIT signaling, trigerring a short live activation: Phosphorylation at Ser-180 and Ser-516 by MAPK and RPS6KA1, respectively, activate the transcription factor activity but also promote ubiquitination and subsequent degradation by the proteasome. Ubiquitinated following phosphorylation at Ser-180, leading to subsequent degradation by the

proteasome. Deubiquitinated by USP13, preventing its degradation.

Cellular Localization Nuclear

Images



ARG56053 anti-MITF antibody [D5] IHC-P image

Immunohistochemistry: Formalin-fixed, paraffin-embedded Human melanoma stained with ARG56053 anti-MITF antibody [D5].