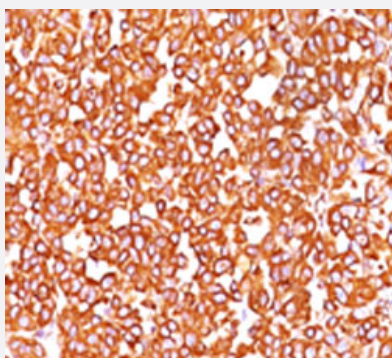


VIM monoclonal antibody, clone SPM576

Catalog # MAB12064 Size 100 ug

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human melanoma with VIM monoclonal antibody, clone SPM576 (Cat# MAB12064).

Specification

Product Description	Mouse monoclonal antibody raised against recombinant VIM.
Immunogen	Recombinant protein corresponding to human VIM.
Host	Mouse
Theoretical MW (kDa)	57-60
Reactivity	Bovine, Dog, Human, Pig
Form	Liquid
Purification	Protein A/G purification
Isotype	IgG1
Recommend Usage	Immunohistochemistry (0.5-1 ug/mL) Western Blot (0.5-1 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.05% BSA, 0.05% sodium azide).

Storage Instruction

Store at 4°C.
Aliquot to avoid repeated freezing and thawing.

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot
- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human melanoma with VIM monoclonal antibody, clone SPM576 (Cat# MAB12064).

Gene Info — VIM

Entrez GeneID[7431](#)**Gene Name**

VIM

Gene Alias

FLJ36605

Gene Description

vimentin

Omim ID[193060](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

This gene encodes a member of the intermediate filament family. Intermediate filaments, along with microtubules and actin microfilaments, make up the cytoskeleton. The protein encoded by this gene is responsible for maintaining cell shape, integrity of the cytoplasm, and stabilizing cytoskeletal interactions. It is also involved in the immune response, and controls the transport of low-density lipoprotein (LDL)-derived cholesterol from a lysosome to the site of esterification. It functions as an organizer of a number of critical proteins involved in attachment, migration, and cell signaling. Mutations in this gene causes a dominant, pulverulent cataract

Other Designations

OTTHUMP00000019224

Disease

- [Alzheimer disease](#)
- [Anorexia Nervosa](#)

- [Bulimia](#)
- [Cognition](#)
- [Genetic Predisposition to Disease](#)