VHL monoclonal antibody (AT82B10)

Von Hippel-Lindau disease (VHL) is a dominant inherited syndrome characterized by the predisposition to develop various kinds of benign and malignant tumors. VHL syndrome is caused by germline mutation in the VHL tumor suppressor, and VHL tumors are associated with loss or mutation of the remaining wild-type allele. VHL protein is also involved in the degradation of hypoxia-inducible factor (HIF).

This antibody is covered by our Worry-Free Guarantee.

Ordering Information

Order Online »

ENZ-ABS746-0100	100µl

Manuals, SDS & CofA

View Online »

Handling & Storage

Handling	Avoid freeze/thaw cycles.
Long Term Storage	-20°C
Shipping	Blue Ice

Regulatory Status

RUO - Research Use Only

Product Details

Alternative Name	Von Hippel-Lindau tumor suppressor isoform 1
Application	ELISA, ICC, IF, WB
Clone	AT82B10
Formulation	Liquid. In PBS, pH 7.4, containing 0.02% sodium azide and 10% glycerol.
Host	Mouse
Immunogen	Recombinant human VHL (aa 1-154) purified from E. coli.
Isotype	IgG2bκ
Purity Detail	Protein A affinity purified
Source	Purified from ascites.
Species Reactivity	Human
UniProt ID	P40337
Worry-free Guarantee	**This antibody is covered by our Worry-Free Guarantee.**

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