# abcam

# Product datasheet

# Anti-VCP antibody ab36047

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## 概述

产品名称 Anti-VCP抗体

描述 兔多克隆抗体to VCP

**宿主** Rabbit

经测试应用 适用于: WB, IHC-P

种属反应性 与反应: Mouse, Rat, Human

预测可用于: Cow, Pig 📤

免疫原 Synthetic peptide conjugated to KLH derived from within residues 1 - 100 of Human VCP.参阅

Abcam的专有抗源政策(Peptide available as ab36482.)

**阳性**对照 Ab36047 gave a positive result in the following whole cell lysates: HL60, Ramos, MDA MB 361

(data not shown), SW480 (data not shown), Y79 (data not shown), THP1 (data not shown), and MOLT4 (data not shown). Ab36047 gave a positive signal in the following tissue lysates: Rat

Kidney; Rat Lung; Rat Liver; Mouse Kidney; Mouse Lung; Mouse Liver.

常规说明

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

性能

形式 Liquid

**存放说明** Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

**存储溶液** pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

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纯**度** Immunogen affinity purified

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab36047于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use a concentration of 1 µg/ml. Detects a band of approximately 97 kDa (predicted molecular weight: 89 kDa).
IHC-P		Use a concentration of 1 µg/ml.

#### 靶标

功能

Necessary for the fragmentation of Golgi stacks during mitosis and for their reassembly after mitosis. Involved in the formation of the transitional endoplasmic reticulum (tER). The transfer of membranes from the endoplasmic reticulum to the Golgi apparatus occurs via 50-70 nm transition vesicles which derive from part-rough, part-smooth transitional elements of the endoplasmic reticulum (tER). Vesicle budding from the tER is an ATP-dependent process. The ternary complex containing UFD1L, VCP and NPLOC4 binds ubiquitinated proteins and is necessary for the export of misfolded proteins from the ER to the cytoplasm, where they are degraded by the proteasome. The NPLOC4-UFD1L-VCP complex regulates spindle disassembly at the end of mitosis and is necessary for the formation of a closed nuclear envelope (By similarity). Regulates E3 ubiquitin-protein ligase activity of RNF19A.

疾病相关

Defects in VCP are the cause of inclusion body myopathy with early-onset Paget disease and frontotemporal dementia (IBMPFD) [MIM:167320]; also known as muscular dystrophy, limb-girdle, with Paget disease of bone or pagetoid amyotrophic lateral sclerosis or pagetoid neuroskeletal syndrome or lower motor neuron degeneration with Paget-like bone disease. IBMPFD features adult-onset proximal and distal muscle weakness (clinically resembling limb girdle muscular dystrophy), early-onset Paget disease of bone in most cases and premature frontotemporal dementia.

序列相似性

Belongs to the AAA ATPase family.

翻译后修饰

Phosphorylated by tyrosine kinases in response to T-cell antigen receptor activation (By similarity). Phosphorylated upon DNA damage, probably by ATM or ATR.

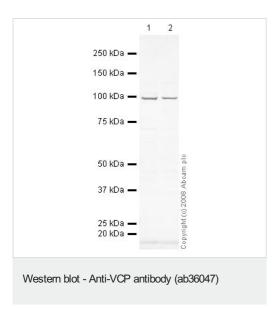
ISGylated.

细胞定位

Cytoplasm > cytosol. Nucleus. Present in the neuronal hyaline inclusion bodies specifically found in motor neurons from amyotrophic lateral sclerosis patients. Present in the Lewy bodies

specifically found in neurons from Parkinson disease patients.

图片



All lanes: Anti-VCP antibody (ab36047) at 1 µg/ml

**Lane 1**: HL60 (Human promyelocytic leukemia cell line) Whole Cell Lysate

Lane 2 : Ramos (Human Burkitt's lymphoma cell line) Whole Cell Lysate

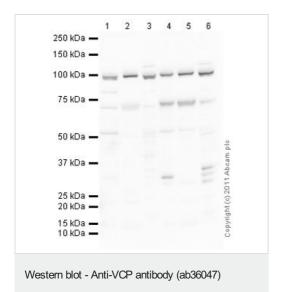
Lysates/proteins at 10 µg per lane.

## Secondary

**All lanes :** IRDye 680 Conjugated Goat Anti-Rabbit lgG (H+L) at 1/10000 dilution

Performed under reducing conditions.

**Predicted band size:** 89 kDa **Observed band size:** 97 kDa



All lanes: Anti-VCP antibody (ab36047) at 1 µg/ml

Lane 1: Kidney (Mouse) Tissue Lysate

Lane 2: Mouse lung normal tissue lysate - total protein (ab29297)

Lane 3 : Liver (Mouse) Tissue Lysate
Lane 4 : Kidney (Rat) Tissue Lysate
Lane 5 : Lung (Rat) Tissue Lysate
Lane 6 : Liver (Rat) Tissue Lysate

Lysates/proteins at 10 µg per lane.

# **Secondary**

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) preadsorbed (ab97080) at 1/5000 dilution

Developed using the ECL technique.

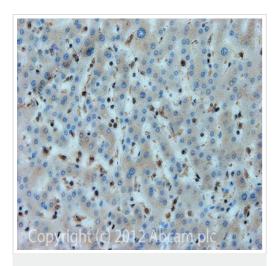
Performed under reducing conditions.

**Predicted band size:** 89 kDa **Observed band size:** 100 kDa

Additional bands at: 52 kDa, 70 kDa. We are unsure as to the

identity of these extra bands.

Exposure time: 10 seconds



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-VCP antibody (ab36047)

IHC image of ab36047 staining in Human normal liver formalin fixed paraffin embedded tissue section, performed on a Leica Bond  $^{TM}$  system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab36047, 1µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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

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