

VCP Monoclonal Antibody (5)

Product Details		
Size	100 μg	
Species Reactivity	Human, Mouse, Rat	
Published Species	Yeast, Mouse, Human	
Host/Isotype	Mouse / IgG2a	
Class	Monoclonal	
Туре	Antibody	
Clone	5	
Conjugate	Unconjugated	
Immunogen	Synthetic peptide corresponding to residues C G(792) G S V Y T E D N D D D L Y G(806) of mouse VCP.	
Form	Liquid	
Concentration	1.0 mg/mL	
Purification	Protein G	
Storage buffer	PBS, pH 7.4, with 1mg/mL BSA	
Contains	0.05% sodium azide	
Storage conditions	-20° C, Avoid Freeze/Thaw Cycles	
RRID	AB_2214638	

Applications	Tested Dilution	Publications
Western Blot (WB)	1:1,000	16 Publications
Immunohistochemistry (IHC)	-	7 Publications
Immunohistochemistry (Paraffin) (IHC (P))	1:500	1 Publication
Immunohistochemistry (Frozen) (IHC (F))	1:500	-
Immunocytochemistry (ICC/IF)	1:20-1:200	5 Publications
Flow Cytometry (Flow)	1/400	-
Immunoprecipitation (IP)	Assay-dependent	2 Publications

Product Specific Information

MA3-004 detects VCP protein in human, mouse, and rat samples.

MA3-004 has successfully been used in Western blot immunoprecipitation, and immunohistochemical procedures. By Western blot, this antibody detects an ~97 kDa protein representing VCP from total lysate of cultured human B cells.

The MA3-004 immunogen is a synthetic peptide corresponding to residues C G(792) G S V Y T E D N D D D L Y G(806) of mouse VCP. This peptide (Cat. # PEP-239) is available for use in neutralization and control experiments.

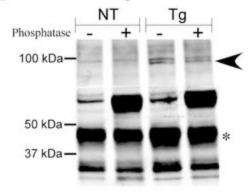
Advanced Verification Data

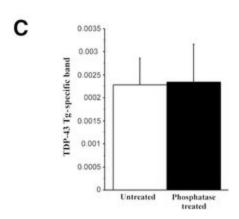
A IP: VCP IB: TDP-43 NT Tg NT Tg NT Tg 100 kDa— * 50 kDa— 37 kDa— IB: VCP — — — —

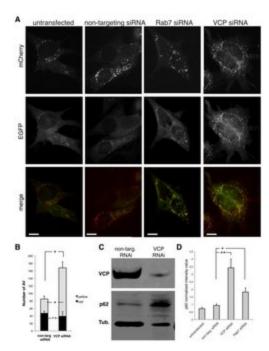
VCP Antibody (MA3-004)

Figure 7 High-molecular-weight TDP-43 interacts with VCP. A: Immunoprecipitation (IP) with a VCP antibody and immunoblot (IB) for TDP-43 uncovered an interaction between VCP and the Tg-specific TDP-43 high-molecular-weight isoform (arrowhead). B: High-molecular-weight TDP-43 is not affected by alkaline phosphatase treatment (arrowhead). C: Quantification of B (n = 4 per group). Asterisks indicate full-length TDP-43 bands. Data are given as means +- SEM. Cell treatment validation info.

B Alkaline Phosphatase treatment

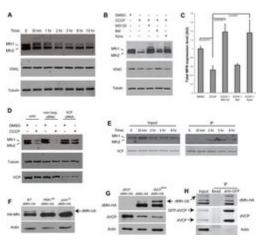






VCP Antibody (MA3-004)

RNAi knock down of VCP impairs autophagosome maturation (A) mCherry-EGFP-LC3b stable MEFs transfected with non-targeting, Rab7 or VCP siRNA. Scale bars equal 10 m. (B) Quantification of autophagosomes and autophagolysosomes under basal conditions in cells transfected with non-targeting or VCP-targeting siRNA. * indicates p<0.01; n.s. indicates non-significant difference, error bars indicate standard errors. (C). Immunoblot against VCP, p62 and tubulin in non-targeting and VCP-targeting siRNA in MEFs. p62 and tubulin were observed simultaneously using a Li-Cor Odyssey system. (D) Quantification of p62 normalized against tubulin in cells transfected with non-targeting, VCP-targeting or Rab7-targeting siRNA. * indicates p<0.01; ** p<0.001; error bars indicate standard deviation from triplicates. Knockdown validation info.



VCP Antibody (MA3-004)

Mitofusin degradation by the proteasome is dependent on VCP A. Western blots in YFP-Parkin stable HeLa cells against MFN1 and 2, VDAC and tubulin at different time points after CCCP treatment. Ubiquitinated forms of MFNs 1/2 can be observed migrating more slowly. B. Western blots in YFP-Parkin stable HeLa cells against MFN1/2, VDAC, and tubulin, Cells were treated for 12 h with CCCP and either proteasome inhibitors (MG132 or epoxomicin) or the autophagy inhibitor bafilomycin. Ubiquitinated forms of MFN1/2 can be again observed migrating more slowly, particularly with proteasome inhibition. C. Quantification of total MFN expression levels normalized against tubulin in YFP-Parkin stable HeLa cells treated for 12 h with CCCP and either proteasome inhibitors (MG132 or epoxomicin), or the autophagy inhibitor bafilomycin. Error bars indicate standard deviation from triplicates. D. Knockdown of VCP stabilizes MFNs 1 and 2. Western blots in YFP-Parkin stable HeLa cells against MFN1/2, VCP and tubulin. Cells were transfected with non-targeting or VCP-targeting siRNA and treated for 12 h with DMSO or CCCP. E. FLAG IP in HeLa cells cotransfected with YFP-Parkin and VCP-FLAG and treated with CCCP for the indicated times. Immunoprecipitation samples were immunoblotted against MFN1/2 and VCP. Following mitochondrial depolarization VCP interacts with MFN2. F. Total dMfn-HA accumulates in PINK1B9 (lane 2) and Park25 (lane 3) null mutants. Notably, ubiquitinated dMfn is decreased in PINK1B9 n Knockdown validation info.

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□ 31 References

Western Blot (16)

International journal of molecular sciences

Parkin Coordinates Platelet Stress Response in Diabetes Mellitus: A Big Role in a Small Cell.

"MA3-004 was used in Western Blotting to investigate the expression of parkin within healthy and diabetic platelets, hearts, muscles and brains to find the significance of parkin in diabetes meticullus and its possible therapeutic use.

Authors: Lee SH.Du J.Hwa J.Kim WH

Species Mouse

Dilution 1:1000

Year 2020

Cell chemical biology

Site-Specific Photo-Crosslinking Proteomics Reveal Regulation of IFITM3 Trafficking and Turnover by VCP/p97 ATPase.

"MA3-004 was used in Western Blotting to reveal the interaction of IFITM3 with VCP and uncovered key functional roles of the interaction on modulating IFITM3 trafficking.

Authors: Wu X,Spence JS,Das T,Yuan X,Chen C,Zhang Y,Li Y,Sun Y,Chandran K,Hang HC,Peng T

Species Human

Dilution Not Cited

Year 2020

View more WB references on thermofisher.com

Immunohistochemistry (7)

Neurology. Genetics

PFKM gene defect and glycogen storage disease GSDVII with misleading enzyme histochemistry.

"Published figure using VCP monoclonal antibody (Product # MA3-004) in Immunohistochemistry"

Authors: Auranen M, Palmio J, Ylikallio E, Huovinen S, Paetau A, Sandell S, Haapasalo H, Viitaniemi K, Piirilä P, Tyynismaa H,Udd B

Species

Human

Dilution Not Cited

Year 2015

Neuropathology: official journal of the Japanese Society of Neuropathology

Ubiquitin-negative, eosinophilic neuronal cytoplasmic inclusions associated with stress granules and autophagy: an immunohistochemical investigation of two cases.

"MA3-004 was used in immunohistochemistry to investigate the association of eosinophilic neuronal cytoplasmic inclusions with stress granules and autophagy in two clinical cases"

Authors: Mori F, Watanabe Y, Miki Y, Tanji K, Odagiri S, Eto K, Wakabayashi K

Species Human

Dilution 1:5000

Year 2014

View more IHC references on thermofisher.com

More applications with references on thermofisher.com

IHC (P) (1) ICC/IF (5) IP (2)

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