

For Research Use Only

# VPRBP Monoclonal antibody

Catalog Number: 66392-1-Ig

Featured Product

1 Publications



## Basic Information

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| <b>Catalog Number:</b><br>66392-1-Ig   | <b>GenBank Accession Number:</b><br>BC022792     | <b>Purification Method:</b><br>Protein A purification  |
| <b>Size:</b><br>150ul , Concentration: 2500 ug/ml by Nanodrop and 2000 ug/ml by Bradford method using BSA as the standard; | <b>GeneID (NCBI):</b><br>9730                    | <b>CloneNo.:</b><br>1A7A8  |
| <b>Source:</b><br>Mouse  | <b>UNIPROT ID:</b><br>Q9Y4B6                     | <b>Recommended Dilutions:</b><br>WB 1:1000-1:4000<br>IHC 1:50-1:500<br>IF-P 1:200-1:800<br>IF/ICC 1:400-1:1600 |
| <b>Isotype:</b><br>IgG2a   | <b>Full Name:</b><br>Vpr (HIV-1) binding protein |  |
| <b>Immunogen Catalog Number:</b><br>AG2184   | <b>Calculated MW:</b><br>1506 aa, 169 kDa        |  |
|  | <b>Observed MW:</b><br>169 kDa                   |  |

## Applications

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| <b>Tested Applications:</b><br>WB, IHC, IF/ICC, IF-P, FC (Intra), ELISA  | <b>Positive Controls:</b><br>WB : HepG2 cells, HeLa cells, HEK-293 cells, K-562 cells, PC-3 cells, DU 145 cells<br>IHC : human breast cancer tissue,<br>IF-P : human breast cancer tissue,<br>IF/ICC : HepG2 cells, human breast cancer tissue |
| <b>Cited Applications:</b><br>WB   |  |
| <b>Species Specificity:</b><br>human   |  |
| <b>Cited Species:</b><br>human, mouse  |  |
| <b>Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0</b> |  |

## Background Information

VprBP was first identified as a protein that can interact with HIV-1 viral protein R (PMID: 11223251). It is a component of the CUL4A-RBX1-DDB1-VprBP/DCAF1 E3 ubiquitin-protein ligase complex that could interact with HIV-1 virus Vpr protein and HIV-2 virus Vpx protein (PMID: 18332868; 17314515; 18606781). VprBP is a 1,507-amino acid protein that contains conserved domains, including YXXY repeats, the Lis homology motif, and WD40 repeats. Through binding to Vpr, VprBP allows Vpr to modulate the catalytic activity of the CUL4-DDB1 complex, which in turn leads to the induction of G2 phase arrest in the virus-infected cells (PMID: 17630831). Recently it has been reported that VprBP is able to regulate the p53-induced transcription and apoptotic pathway (PMID: 22184063).

## Notable Publications

| Author          | Pubmed ID | Journal    | Application |
|-----------------|-----------|------------|-------------|
| Nikhil B Ghatge | 37069142  | Nat Commun | WB          |

## Storage

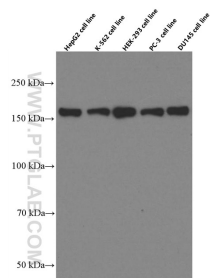
**Storage:**  
Store at -20°C. Stable for one year after shipment.  
**Storage Buffer:**  
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.  
**Aliquoting is unnecessary for -20°C storage**

\*\*\* 20ul sizes contain 0.1% BSA

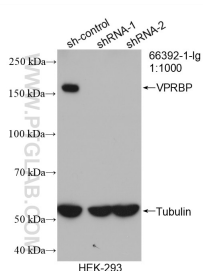
For technical support and original validation data for this product please contact:  
T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)  
E: [proteintech@ptglab.com](mailto:proteintech@ptglab.com)  
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This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

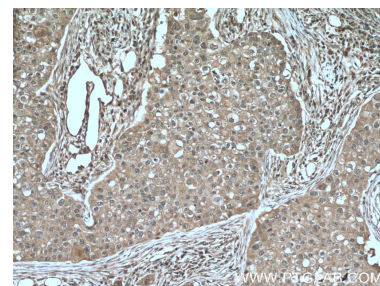
## Selected Validation Data



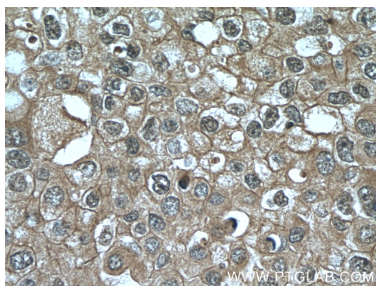
Various lysates were subjected to SDS PAGE followed by western blot with 66392-1-Ig (VPRBP antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours.



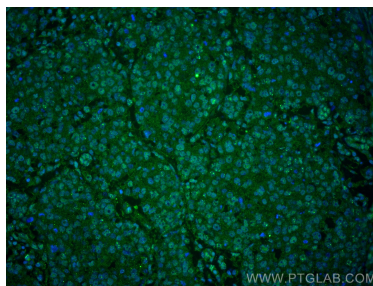
WB result of VPRBP antibody (66392-1-Ig; 1:1000; incubated at room temperature for 1.5 hours) with sh-Control and sh-VPRBP transfected HEK-293 cells.



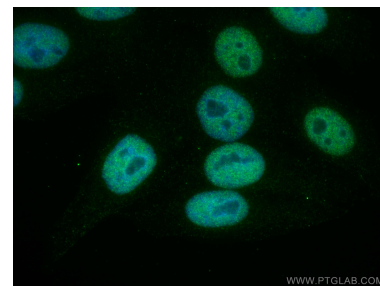
Immunohistochemical analysis of paraffin-embedded human breast cancer tissue slide using 66392-1-Ig (VPRBP antibody) at dilution of 1:200 (under 10x lens. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



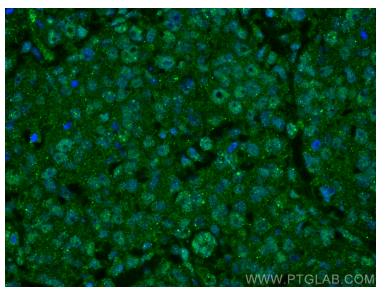
Immunohistochemical analysis of paraffin-embedded human breast cancer tissue slide using 66392-1-Ig (VPRBP antibody) at dilution of 1:200 (under 40x lens. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



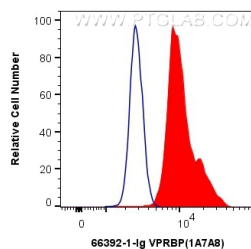
Immunofluorescent analysis of (4% PFA) fixed human breast cancer tissue using VPRBP antibody (66392-1-Ig, Clone: 1A7A8) at dilution of 1:400 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



Immunofluorescent analysis of (4% PFA) fixed HepG2 cells using VPRBP antibody (66392-1-Ig, Clone: 1A7A8) at dilution of 1:800 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



Immunofluorescent analysis of (4% PFA) fixed human breast cancer tissue using VPRBP antibody (66392-1-Ig, Clone: 1A7A8) at dilution of 1:400 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



1X10<sup>6</sup> K-562 cells were intracellularly stained with 0.4 ug Anti-Human VPRBP (66392-1-Ig, Clone:1A7A8) and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000 (red), or 0.4 ug Mouse IgG2a Isotype Control (66360-2-Ig, Clone: K11A1B2A2) and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000 (blue). Cells were fixed with 4% PFA and permeabilized with Flow Cytometry Perm Buffer (PF00011-C).