

**For Research Use Only.**  
**Not for use in diagnostic procedures.**



## Anti-Vinculin pAb

**CODE No.** PM088

**CLONALITY** Polyclonal  
**ISOTYPE** Rabbit Ig, affinity purified  
**QUANTITY** 100 µL

**SOURCE** Purified Ig from rabbit serum  
**FORMURATION** PBS containing 50% Glycerol (pH 7.2). No preservative is contained.  
**STORAGE** This antibody solution is stable for one year from the date of purchase when stored at -20°C.

### APPLICATION-CONFIRMED

Western blotting 1:1,000 for chemiluminescence detection system

### SPECIES CROSS REACTIVITY on WB

Species	Human	Mouse	Rat	Hamster
Sample	HeLa	NIH/3T3	PC12	CHO
Reactivity	+	+	+	+

**Entrez Gene ID** 7414 (Human), 22330 (Mouse), 305679 (Rat), 100759958 (Hamster)

For more information, please visit our web site <http://ruo.mbl.co.jp/>



**RELATED PRODUCTS**

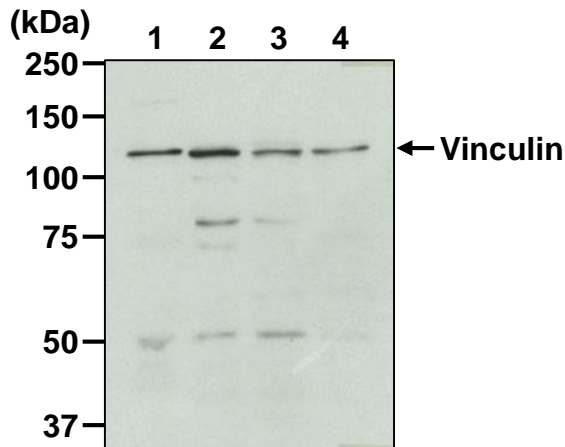
Antibodies

PM088	Anti-Vinculin pAb
PM054	Anti- $\alpha$ -Tubulin pAb
PM054-7	Anti- $\alpha$ -Tubulin pAb-HRP-DirecT
M175-3	Anti- $\alpha$ -Tubulin mAb
PM053	Anti- $\beta$ -Actin pAb
PM053-7	Anti- $\beta$ -Actin pAb-HRP-DirecT
M177-3	Anti- $\beta$ -Actin mAb
M171-3	Anti-GAPDH mAb
M171-7	Anti-GAPDH mAb-HRP-DirecT
PM064	Anti-Lamin B1 pAb

### **SDS-PAGE & Western blotting**

- 1) Wash cells 3 times with PBS and suspend them in Extraction buffer [50 mM Tris-HCl (pH7.5), 150 mM NaCl, 0.05% NP-40], then sonicate briefly (up to 10 sec.).
- 2) Centrifuge the tube at 12,000 x g for 10 min. at 4°C and transfer the supernatant to another tube. Measure the protein concentration of the supernatant.
- 3) Add equal volume of 2 x Laemmli's sample buffer and mix well.
- 4) Boil the sample for 3 min. and centrifuge. Load 10 µg of the sample per lane in a 1-mm-thick SDS-polyacrylamide gel (7.5% acrylamide) for electrophoresis.
- 5) Blot the protein to a polyvinylidene difluoride (PVDF) membrane at 1 mA/cm<sup>2</sup> for 1 hr. in a semi-dry transfer system (Transfer Buffer: 25 mM Tris, 190 mM glycine, 20% MeOH). See the manufacturer's manual for precise transfer procedure.
- 6) To reduce nonspecific binding, soak the membrane in 10% skimmed milk (in PBS, pH 7.2) overnight at 4°C.
- 7) Incubate the membrane with primary antibody diluted with 1% skimmed milk (in PBS, pH 7.2) as suggested in the **APPLICATION** for 1 hr. at room temperature. (The concentration of antibody will depend on the conditions.)
- 8) Wash the membrane with PBS-T [0.05% Tween-20 in PBS] (5 min. x 3 times).
- 9) Incubate the membrane with the 1:10,000 Anti-IgG (Rabbit) pAb-HRP (MBL; code no. 458) diluted with 1% skimmed milk (in PBS, pH 7.2) for 1 hr. at room temperature.
- 10) Wash the membrane with PBS-T (5 min. x 3 times).
- 11) Wipe excess buffer on the membrane, and then incubate it with appropriate chemiluminescence reagent for 1 min. Remove extra reagent from the membrane by dabbing with paper towel, and seal it in plastic wrap.
- 12) Expose to an X-ray film in a dark room for 3 min. Develop the film as usual. The condition for exposure and development may vary.

(Positive controls for Western blotting; HeLa, NIH/3T3, PC12 and CHO)



#### ***Western blot analysis of Vinculin***

- Lane 1: HeLa
- Lane 2: NIH/3T3
- Lane 3: PC12
- Lane 4: CHO

Immunoblotted with Anti-Vinculin pAb (PM088)