

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



# RiboCluster Profiler™

**RBP** Antibody

# Anti-TARDBP (TDP-43) pAb

CODE No.	RN107PW
CLONALITY	Polyclonal
ISOTYPE	Rabbit Ig, affinity purified
QUANTITY	100 μL, 1 mg/mL
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.

# APPLICATIONS

Western blotting1:500 for chemiluminescence detection systemImmunoprecipitation $5 \ \mu L/500 \ \mu L$  of nuclear extract from  $1 \ x \ 10^7$  cells/sample

# SPECIES CROSS REACTIVITY on WB

Species	Human	Mouse	Rat	Hamster
Cell	HeLa, HEK293T, Jurkat, K562	NIH/3T3, WR19L	Rat1	СНО
Reactivity	+	+	-	+

**Entrez Gene ID** 23435 (Human), 230908 (Mouse)

For more information, please visit our web site http://ruo.mbl.co.jp/je/rip-assay/

**LICENSING OPPORTUNITY:** The RIP-Assay uses patented technology (US patent No. 6,635,422, US patent No. 7,504,210, JP patent No. 5,002,105) of Ribonomics, Inc. MBL manufactures and distributes this product under license from Ribonomics, Inc. Researchers may use this product for their own research. Researchers are not allowed to use this product or RIP-Assay technology for commercial purpose without a license. For commercial use, please contact us for licensing opportunities at RIP@mbl.co.jp



MEDICAL & BIOLOGICAL LABORATORIES CO., LTD. URL <u>http://ruo.mbl.co.jp/je/rip-assay/</u> e-mail <u>support@mbl.co.jp</u>, TEL 052-238-1904

#### **RELATED PRODUCTS**

<b>RIP-Assay</b>	<u>    Kit                                </u>
RN1001	RIP-Assay Kit
RN1005	RIP-Assay Kit for microRNA

**RIP-Certified Antibody** 

in continee	<u> </u>
RN001P	Anti-EIF4E pAb
RN002P	Anti-EIF4G1 pAb
RN003P	Anti-EIF4G2 pAb
RN004P	Anti-ELAVL1 (HuR) pAb
RN005P	Anti-ELAVL2 (HuB) pAb
RN006P	Anti-ELAVL3 (HuC) pAb
RN007P	Anti-IGF2BP1 (IMP1) pAb
RN008P	Anti-IGF2BP2 (IMP2) pAb
RN009P	Anti-IGF2BP3 (IMP3) pAb
RN010P	Anti-MSI1 (Musashi1) pAb
RN011P	Anti-PTBP1 pAb
RN012P	Anti-STAU1 pAb
RN013P	Anti-STAU2 pAb
RN014P	Anti-TIA1 pAb
RN015P	Anti-YBX1 pAb
RN016P	Anti-FMR1 pAb
RN017P	Anti-FXR1 pAb
RN018P	Anti-FXR2 pAb
RN019P	Anti-HNRNPK pAb
RN020P	Anti-ILF3 pAb
RN021P	Anti-KHDRBS1 pAb
RN022P	Anti-PABPC4 pAb
RN024P	Anti-PCBP1 pAb
RN025P	Anti-PCBP2 pAb
RN026P	Anti-PUM1 pAb
RN027P	Anti-PUM2 pAb
RN028P	Anti-EIF2C1 (AGO1) pAb
RN032P	Anti-CIRBP pAb
RN033P	Anti-TNRC6A (GW182) pAb
RN037P	Anti-AUH pAb
RN038P	Anti-CPEB1 pAb
RN041P	Anti-KHDRBS2 (SLM1) pAb
RN045P	Anti-SLBP pAb
RN001M	Anti-IGF2BP1 (IMP1) mAb (6H6)
RN003M	Anti-EIF2C2 (AGO2) mAb (1B1-E2H5)
RN004M	Anti-Ribosomal P0/P1/P2 mAb (9D5)
RN005M	Anti-EIF2C2 (AGO2) mAb (2A8)
RN006M	Anti-EIF4E mAb (C107-3-5)
RN007M	Anti-ELAVL1 (HuR) mAb (C67-1)
RN009M	Anti-PABPC1 mAb (10E10)
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#### RBP Antibody

RBP Antibody works on WB and /or IP, but not certified for working on RIP-Assay.

RN008MWAnti-ELAVL1 (HuR) mAb (C54-6)RN023PWAnti-PABPN1 pAbRN031PWAnti-ZFP36 (TTP) pAbRN046PWAnti-SYNCRIP (HNRNPQ) pAbRN047PWAnti-PTBP2 pAbRN048PWAnti-G3BP1 pAbRN049PWAnti-G3BP2 pAbRN050PWAnti-GRSF1 pAbRN051PWAnti-HDLBP (Vigilin) pAb

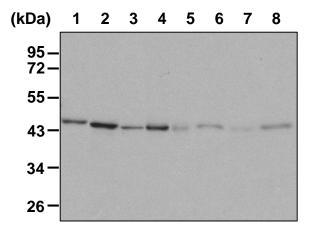
RN052PW Anti-HNRNPC pAb RN053PW Anti-PAIP1 pAb RN054PW Anti-PCBP3 pAb RN056PW Anti-SERBP1 pAb RN057PW Anti-TARBP1 pAb RN058PW Anti-TARBP2 pAb RN059PW Anti-TIAL1 pAb RN060PW Anti-HNRNPD (AUF1) pAb RN061PW Anti-HNRNPA0 pAb RN062PW Anti-DGCR8 pAb RN063PW Anti-DHX9 pAb RN064PW Anti-FUSIP1 (SRSF10) pAb RN065PW Anti-KHSRP pAb RN066PW Anti-KIAA0020 pAb RN067PW Anti-PPP1R10 pAb RN068PW Anti-PPP1R8 pAb RN069PW Anti-RBM14 pAb RN077PW Anti-SMN1 pAb RN078PW Anti-SMNDC1 pAb RN079PW Anti-SRSF7 (9G8) pAb RN080PW Anti-SRSF3 (SRp20) pAb RN081PW Anti-SRSF9 (SRp30c) pAb RN082PW Anti-SRSF5 (SRP40) pAb RN084PW Anti-SRRM1 (SRM160) pAb RN085PW Anti-U2AF1 pAb RN086PW Anti-U2AF2 pAb RN087PW Anti-ALYREF (THOC4) pAb RN088PW Anti-NXF1 (TAP) pAb RN089PW Anti-MAGOH pAb RN090PW Anti-DDX21 pAb RN091PW Anti-DDX23 pAb RN091PW Anti-DDA25 pA6 RN092PW Anti-NONO (P54NRB) pAb RN093PW Anti-PRPF4 pAb RN094PW Anti-PRPF8 pAb RN095PW Anti-SNRNP200 pAb RN096PW Anti-SNRNP40 pAb RN097PW Anti-SNRNP70 pAb RN098PW Anti-EDC4 pAb RN099PW Anti-EIF4A1 pAb RN100PW Anti-EXOSC5 (RRP46) (Human) pAb RN101PW Anti-FBL (Fibrillarin) pAb RN102PW Anti-GEMIN2 (Human) pAb RN103PW Anti-NCBP1 (CBP80) pAb RN104PW Anti-PAN2 (USP52) (Human) pAb RN105PW Anti-PARN pAb RN106PW Anti-SFPQ (PSF) pAb RN107PW Anti-TARDBP (TDP-43) pAb RN108PW Anti-UPF1 pAb RN109PW Anti-XRN1 (Human) pAb Anti-hnRNP-A2/B1 mAb (C20308) D216-3 M162-3 Anti-p62 (SQSTM1) (Human) mAb (5F2) PM045 Anti-p62 (SQSTM1) pAb PM066 Anti-p62 C-terminal pAb

For the latest information of RiboCluster Profiler<sup>™</sup>, please visit our website at <u>http://ruo.mbl.co.jp/je/rip-assay/</u>

# **SDS-PAGE & Western blotting**

- 1) Wash 1 x  $10^7$  cells 3 times with PBS and suspend them in 1 mL of Laemmli's sample buffer, then sonicate briefly (up to 20 sec.).
- 2) Boil the samples for 3 min. and centrifuge. Load 10 µL of the sample per lane in a 1-mm-thick SDS-polyacrylamide gel (12.5% acrylamide) for electrophoresis.
- 3) 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.
- 4) To reduce nonspecific binding, soak the membrane in 5% skimmed milk (in PBS, pH 7.2) for overnight at 4°C
- 5) Wash the membrane with PBS-T (0.05% Tween-20 in PBS) [5 min. x 3 times].
- 6) Incubate the membrane with primary antibody diluted with 1% skimmed milk (in PBS, pH 7.2) as suggested in the **APPLICATIONS** for 1 hr. at room temperature. (The concentration of antibody will depend on the conditions.)
- 7) Wash the membrane with PBS-T (10 min. x 3 times).
- 8) Incubate the membrane with the 1:5,000 of 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.
- 9) Wash the membrane with PBS-T (10 min. x 3 times).
- 10) Wipe excess buffer on the membrane, 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.
- 11) Expose to an X-ray film in a dark room for 3 min. Develop the film as usual settings. The condition for exposure and development may vary.

(Positive controls for Western blotting; HeLa, HEK93T, Jurkat, K562, NIH/3T3, WR19L and CHO)



# Western blot analysis of TARDBP (TDP-43)

Lane 1:	HeLa
Lane 2:	HEK293T
Lane 3:	Jurkat
Lane 4:	K562
Lane 5:	NIH/3T3
Lane 6:	WR19L
Lane 7:	Rat1
Lane 8:	СНО

Immunoblotted with Anti-TARDBP (TDP-43) pAb (RN107PW)

# **Immunoprecipitation**

- 1) Wash 2 x  $10^7$  cells 4 times with PBS and resuspend them with 1 mL of ice-cold Lysis Buffer (+) (MBL; code no. RN1001) containing appropriate protease inhibitors and DTT. Vortex thoroughly, then incubate on ice for 5 min.
- 2) Centrifuge the tube at 12,000 x g for 5 min. at  $4^{\circ}$ C and discard the supernatant.
- 3) Wash the pellet 3 times with PBS and resuspend them with 500  $\mu$ L of RIPA buffer, then sonicate briefly.
- 4) Centrifuge the tube at 12,000 x g for 5 min. at 4°C. and transfer the supernatant to another tube.
- 5) Add 500 µL of ice-cold Lysis buffer (+) into the supernatant. Mix well by pipetting up and down.
- 6) Add 20  $\mu$ L of 50% protein G agarose beads slurry resuspended in ice-cold Lysis Buffer (+) into the supernatant. Incubate it at 4°C with rotating for 1 hr.
- 7) Centrifuge the tube at 2,000 x g for 2 min. at 4°C and transfer the supernatant to another tube (precleared sample).
- 8) Mix 20 μL of 50% protein G agarose beads slurry resuspended in 1 mL of ice-cold Wash Buffer (+) (MBL; code no. RN1001) containing DTT at the appropriate concentration with normal rabbit IgG (RIP-Assay Kit) or anti-TARDBP (TDP-43) pAb (MBL; code no. RN107PW) as suggested in the **APPLICATIONS**. Incubate at 4°C with rotating for 1 hr.
- 9) Wash the beads 1 time with ice-cold Lysis Buffer (+). Carefully discard the supernatant.
- 10) Add 500  $\mu$ L of the precleared sample (prepared in step 7)) to the tube containing antibody conjugated beads, then incubate with gentle agitation for 2 hr. at 4°C.
- 11) Wash the beads 4 times with 1 mL of ice-cold Wash Buffer (+).
- 12) Resuspend the beads in 20 µL of Laemmli's sample buffer, boil for 3 min. and centrifuge.
- 13) Load 10 µL of the sample per lane in a 1-mm-thick SDS-polyacrylamide gel (12.5% acrylamide) for electrophoresis.
- 14) 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.
- 15) To reduce nonspecific binding, soak the membrane in 5% skimmed milk (in PBS, pH 7.2) for overnight at 4°C.
- 16) Wash the membrane with PBS-T (0.05% Tween-20 in PBS) [5 min. x 3 times].
- 17) Incubate the membrane with primary antibody diluted with 1% skimmed milk (in PBS, pH 7.2) as suggested in the **APPLICATIONS** for 1 hr. at room temperature. (The concentration of antibody will depend on the conditions.)
- 18) Wash the membrane with PBS-T (10 min. x 3 times).
- 19) Incubate the membrane with 1: 1,000 of Rabbit TrueBlot<sup>®</sup> anti-Rabbit IgG-HRP (eBioscience; code no. 18-8816-33) diluted with 1% skimmed milk (in PBS, pH 7.2) for 1 hr. at room temperature.
- 20) Wash the membrane with PBS-T (10 min. x 3 times).
- 21) Wipe excess buffer on the membrane, then incubate it with appropriate chemiluminescence reagent for 1 min.
- 22) Remove extra reagent from the membrane by dabbing with paper towel, and seal it in plastic wrap.
- 23) 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 control for Immunoprecipitation; HeLa nuclear extract)

