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

RiboCluster Profiler™ RIP-Certified Antibody

Anti-STAU1 (Human) pAb

Code No.	Quantity	Concentration	Form
RN012P	200 µL	1 mg/mL	Affinity Purified

BACKGROUND: Mammalian Staufen1 (Stau1) is a ubiquitous double-stranded RNA binding protein associated with polysomes. It has two protein isoforms, molecular weights of 55 and 63 kDa. The 55 kDa isoform associates with 40S and 60S ribosomal subunit and colocalizes with the rough endoplasmic reticulum. Stau1 is thought to function in transporting mRNA, controlling and eliciting mRNA decay. Previous study showed that the Stau1 recruits upf1 to 3' UTR region of Arf1 mRNA and induces the Stau1-mediated mRNA decay (SMD). Recent study suggests that Stau1 regulates diverse class of mammalian transcripts. In order to identify mRNA targets of Staufen proteins, RIP-Assay technology was employed and identified distinct but overlapping subset of cellular mRNAs associated with Stau1 and Stau2-containing ribonucleoprotein complexes.

RIP-CERTIFIED ANTIBODY:
Posttranscriptional regulation of gene expression is a ribonucleoprotein-driven process, which involves RNA binding proteins (RBPs) and non-coding RNAs that affect splicing, nuclear export, subcellular localization, mRNA decay and translation. The RNP Immunoprecipitation-Chip (RIP-Chip), RIP-Seq and RIP-RTPCR allow the identification of multiple RNA targets of RBPs globally and within the context of a cell extract. Antibodies specific to the RNA binding protein of interest are used to co-immunoprecipitate the RNA binding protein and the associated subset of mRNAs. The mRNA content is interrogated using standard microarray or sequencing technology. RIP-Certified Antibody is validated for use in RNP Immunoprecipitation (RIP) in conjunction with the RIP-Assay Kit distributed from MBL. Its ability to immunoprecipitate mRNAs and RBPs complex was confirmed by quantitative and qualitative analysis on NanoDrop, Bioanalyzer and RT-PCR or microarray.

SOURCE: This antibody was purified from rabbit serum by affinity column chromatography. The rabbit was immunized with KLH conjugated synthetic peptide, MKLGKKPMYKPVDPYSRMQ corresponding to 82-100 aa.

FORMULATION: 200 µL volume of 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.

INTENDED USE:
For Research Use Only. Not for use in diagnostic procedures.

REACTIVITY: This antibody reacts with human STAU1 (~63 kDa) on Western blotting, Immunoprecipitation and RNP Immunoprecipitation.

APPLICATIONS:
RNP Immunoprecipitation; 15 µg/500 µL of cell extract from 1.5 x 10^7 cells
Western blotting; 1:1,000 for chemiluminescence detection system
Note: This antibody is suitable for IP-WB. Non-specific bands are detected when this antibody is used on WB for whole cell lysate samples.
Immunoprecipitation; 1.5 µg/50 µL of cell extract from 1.25 x 10^6 cells
Immunohistochemistry; Not tested
Immunocytochemistry; Not tested
Flow cytometry; Not tested

Detailed procedure is provided in the following **PROTOCOLS**.

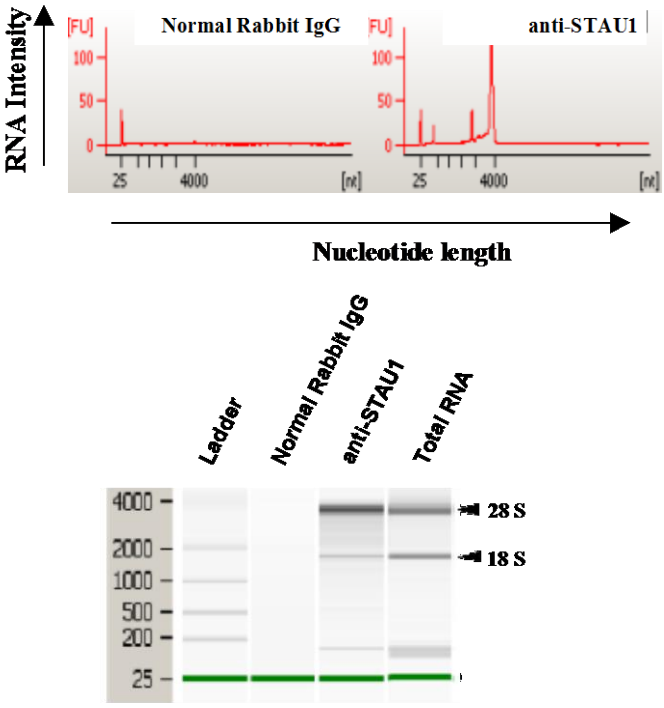
REFERENCES:
1) Ramen, A. V., *et al., Brain Behav.* **3**, 562-574 (2013) [RIP]
2) Yu, Z., *et al., J Biol Chem.* **287**, 22560-22572 (2012) [RIP]
3) Furic, L., *et al., RNA* **14**, 324-335 (2008)
4) Kim, Y. K., *et al., EMBO J.* **26**, 2670-2681 (2007)
5) Kim, Y. K., *et al., Cell* **120**, 195-208 (2005)
6) Kiebler, M. A., *et al., J. Neurosci.* **19**, 288-297 (1999)

SPECIES CROSS REACTIVITY:

Species	Human	Mouse	Rat	Hamster
Cells	Transfectant	NIH/3T3, WR19L	Not tested	Not tested
Reactivity on WB	+	-		

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

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



Analysis of isolated RNA with Bioanalyzer.

Average of the RNA Quantity (n=2)	
Antibody	RNA (ng)
Normal Rabbit IgG	79.0
anti-STAU1	1215.0
Total RNA	254710.0

PROTOCOLS:

RNP Immunoprecipitation

Some buffer and reagents are included in the RIP-Assay Kit (code. RN1001). Please also refer to the protocol packaged in the RIP-Assay Kit.

[Material Preparation]

1. Lysis Buffer (+)
   Before using the Lysis Buffer, protease inhibitors, RNase inhibitors, and DTT are added to the Lysis Buffer at the appropriate concentration.
2. Wash Buffer (+)
   Before using the Wash Buffer, DTT is added to the Wash Buffer at the appropriate concentration.

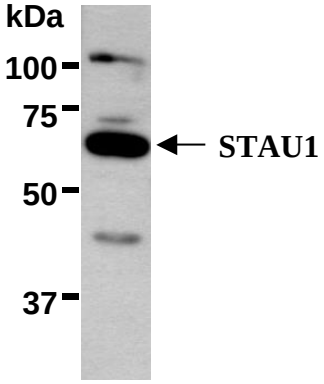
[Precaution]

RNP Immunoprecipitation using this antibody requires the addition of 30 µL of High-Salt Solution (RIP-Assay Kit) to each mL of Lysis Buffer (+) and Wash Buffer (+) just before use.

Protocol

1) Wash 1.5 x 10^7 cells 2 times with PBS and resuspend them with 500 µL of ice-cold Lysis Buffer (+) containing appropriate protease inhibitors, RNase inhibitors, and DTT. Vortex for 10 seconds. Leave on ice for 10 minutes.
2) Centrifuge the tube at 12,000 x g for 5 minutes at 4°C and transfer the supernatant to another tube.
3) Add 25 µL of 50% protein A agarose beads slurry resuspended in Lysis Buffer (+) into the supernatant. Incubate it at 4°C with rotating for 1 hour.
4) Centrifuge the tube at 2,000 x g for 1 minute at 4°C and transfer the supernatant to another tube (precleared sample).
5) Mix both 25 µL of 50% protein A agarose beads slurry resuspended in nuclease-free PBS and Normal Rabbit IgG (RIP-Assay Kit) or Anti-STAU1 (Human) pAb at the amount of suggested in the **APPLICATIONS,** and then add 1 mL of Wash buffer (+) into each tube. Incubate with gentle agitation for 1 hour at 4°C.
6) Wash the beads once with ice-cold Lysis Buffer (+) (centrifuge the tube at 2,000 x g for 1 minute). Carefully discard the supernatant using a pipettor without disturbing the beads.
7) Add 500 µL of cell lysate (precleared sample of step 4), then incubate with gentle agitation for 3 hours at 4°C.
8) Wash the beads 4 times with Wash Buffer (+) (centrifuge the tube at 2,000 x g for 1 minute).
9) Add 400 µL of Master mix solution (Solution I: Solution II = 10 µL: 390 µL). Vortex for 10 seconds.
10) Add 250 µL of Solution III. Vortex for 10 seconds.
11) Centrifuge the tube at 2,000 x g for 2 minutes.
12) Transfer the supernatant to the tube containing 2 µL of Solution IV.
13) Add 600 µL of ice-cold 2-propanol and place at -20°C for 20 minutes. Centrifuge the tube at 12,000 x g for 10 minutes.
14) Wash the pellet 2 times with 0.5 mL of ice-cold 70% Ethanol and dry up the pellet for 5-15 minutes.
15) Dissolve the pellets in nuclease-free water.
16) RNA was quantified with NanoDrop (Thermo Fisher Scientific Inc.) and the RNA quality was analyzed with Bioanalyzer (Agilent Technologies, Inc.).

(Positive control for RNP Immunoprecipitation; 293T)



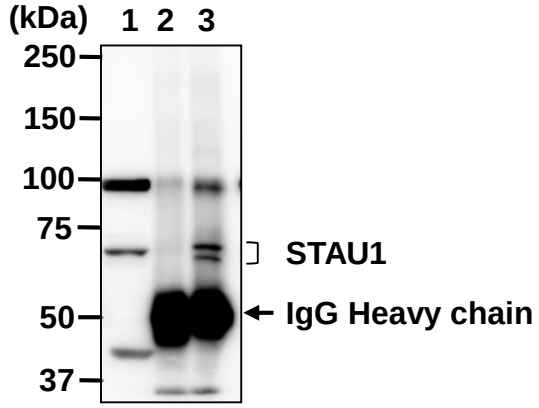
Western blot analysis of STAU1 expression in transfectant using RN012P.

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.
2) Boil the samples for 2 minutes and centrifuge. Load 10 µL of the sample per lane in a 1 mm thick SDS-polyacrylamide gel (5% acrylamide) for electrophoresis.

3) Blot the protein to a polyvinylidene difluoride (PVDF) membrane at 1 mA/cm^2 for 1 hour in a semi-dry transfer system (Transfer Buffer: 25 mM Tris, 190 mM glycine, 20% MeOH). See the manufacture's manual for precise transfer procedure.
4) To reduce nonspecific binding, soak the membrane in 10% skimmed milk (in PBS, pH 7.2) for 1 hour at room temperature, or overnight at 4°C.
5) Incubate the membrane with primary antibody diluted with PBS, pH 7.2 containing 1% skimmed milk as suggested in the **APPLICATIONS** for 1 hour at room temperature. (The concentration of antibody will depend on condition.)
6) Wash the membrane with PBS-T [0.05% Tween-20 in PBS] (5 minutes x 3 times).
7) 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 hour at room temperature.
8) Wash the membrane with PBS-T (5 minutes x 3 times).
9) Wipe excess buffer on the membrane, then incubate it with appropriate chemiluminescence reagent for 1 minute. Remove extra reagent from the membrane by dabbing with paper towel, and seal it in plastic wrap.
10) Expose to an X-ray film in a dark room for 1 minute. Develop the film as usual. The condition for exposure and development may vary.

(Positive control for Western blot; 293T transfectant)



Immunoprecipitation of STAU1 from 293T cells with Normal Rabbit IgG (2) or RN012P (3). *After immunoprecipitated with the antibody, immunocomplex was resolved on SDS-PAGE and immunoblotted with RN012P. Lane 1 is the input sample (2% of total cell lysate). ImageQuant LAS4000 imaging system (Fujifilm) was used for detection (exposure time: 20 seconds).*

Immunoprecipitation

1) Wash 1.25 x 10^7 cells 2 times with PBS and resuspend them with 1 mL of ice-cold Lysis buffer (RIP-Assay Kit) containing appropriate protease inhibitors, then sonicate briefly (up to 10 seconds).
2) Centrifuge the tube at 12,000 x g for 5 minutes at 4°C and transfer the supernatant to another tube.
3) Add 25 µL of 50% protein A agarose beads slurry resuspended in Lysis Buffer into the supernatant. Incubate it at 4°C with rotating for 1 hour.
4) Centrifuge the tube at 2,000 x g for 1 minute at 4°C and transfer the supernatant to another tube (precleared sample).
5) Mix both 25 µL of 50% protein A agarose beads slurry resuspended in nuclease-free PBS and Normal Rabbit IgG (RIP-Assay Kit) or Anti-STAU1 (Human) pAb at the amount of suggested in the APPLICATIONS, and then add 1 mL of Wash buffer into each tube. Incubate with gentle agitation for 1 hour at 4°C.
6) Wash the beads once with ice-cold Lysis Buffer (centrifuge the tube at 2,000 x g for 1 minute). Carefully discard the supernatant using a pipettor without disturbing the beads.
7) Add 50 µL of cell lysate (precleared sample of step 4)), then incubate with gentle agitation for 1 hour at 4°C.
8) Wash the beads 4 times with Wash Buffer (+) (centrifuge the tube at 2,000 x g for 1 minute).
9) Resuspend the beads in 20 µL of Laemmli's sample buffer, boil for 3-5 minutes, and centrifuge for 5 minutes. Use 20 µL/lane for the SDS-PAGE analysis.
(See SDS-PAGE & Western blotting.)

(Positive control for Immunoprecipitation; 293T)

RELATED PRODUCTS:

RIP Certified Antibody

RN012P	Anti-STAU1 (Human) pAb
RN013P	Anti-STAU2 (Human) pAb
RN014P	Anti-TIA1 pAb
RN016P	Anti-FMR1 pAb
RN026P	Anti-PUM1 pAb
RN027P	Anti-PUM2 pAb
RN045P	Anti-SLBP pAb
RN009M	Anti-PABPC1 mAb (10E10)

RBP Antibody

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

RN031PW	Anti-ZFP36 (TTP) pAb
RN034PW	Anti-CUGBP1 pAb
RN036PW	Anti-ACO1 (IRP1) pAb
RN051PW	Anti-HDLBP (Vigilin) pAb
RN060PW	Anti-HNRNPD (AUF1) pAb
RN065PW	Anti-KHSRP pAb
RN002MW	Anti-CUGBP1 mAb (3B1)

Kit

RN1001	RIP-Assay Kit
RN1005	RIP-Assay Kit *for microRNA*
RN1011	RiboTrap Kit

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

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