

## **Product datasheet for TA302286S**

## **PARG Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** IHC, WB

Recommended Dilution: WB: 1:1000, IHC: 1:10~50

Reactivity: Human (Predicted: Mouse)

**Host:** Rabbit

**Isotype:** lg

Clonality: Polyclonal

**Immunogen:** This Parg antibody is generated from rabbits immunized with a KLH conjugated synthetic

peptide between 390-421 amino acids from the C-terminal region of human Parg.

**Formulation:** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

**Purification:** This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by

dialysis against PBS.

**Conjugation:** Unconjugated

**Storage:** Store at -20°C as received.

**Stability:** Stable for 12 months from date of receipt.

Predicted Protein Size: 111110 Da

**Gene Name:** poly(ADP-ribose) glycohydrolase

Database Link: NP 003622

Entrez Gene 26430 MouseEntrez Gene 8505 Human

Q86W56

**Background:** Poly(ADP-ribose) glycohydrolase (PARG) is the major enzyme responsible for the catabolism

of poly(ADP-ribose), a reversible covalent-modifier of chromosomal proteins. The protein is found in many tissues and may be subject to proteolysis generating smaller, active products.

Synonyms: PARG99



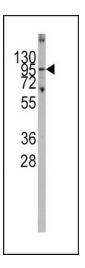
**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

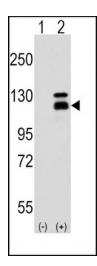
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## **Product images:**

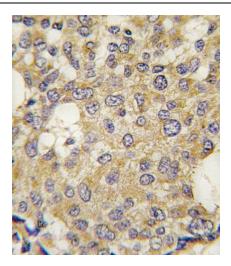


Western blot analysis of Parg (arrow) using rabbit polyclonal Parg Antibody (C-term) (Cat.# [TA302286]).293 cell lysates (2 ug/lane) either nontransfected (Lane 1) or transiently transfected with the Parg gene (Lane 2) (Origene Technologies).



Western blot analysis of Parg Antibody (C-term) in 293 cell line lysates (35ug/lane). Parg (arrow) was detected using the purified Pab.





Formalin-fixed and paraffin-embedded human breast carcinoma tissue reacted with Parg antibody (C-term) (Cat.#[TA302286]), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.