



# **AGO2 Polyclonal Antibody**

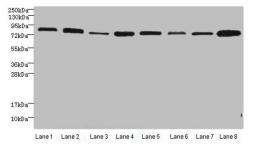
<b>Product Details</b>	
Size	100 μg
Species Reactivity	Human, Mouse
Host/Isotype	Rabbit / IgG
Class	Polyclonal
Type	Antibody
Conjugate	Unconjugated
Immunogen	Recombinant Human Protein argonaute-2 protein (517-818aa)
Form	Liquid
Concentration	1 mg/mL
Purification	Protein G
Storage buffer	10mM PBS, pH 7.4, with 50% glycerol
Contains	0.03% ProClin 300
Storage conditions	-20°C or -80°C if preferred
RRID	AB_2902354

Applications	Tested Dilution	Publications
Western Blot (WB)	1:1,000-1:5,000	-
Immunohistochemistry (Paraffin) (IHC (P))	1:100-1:1,500	-
ChIP assay (ChIP)	Assay-Dependent	-

## **Product Specific Information**

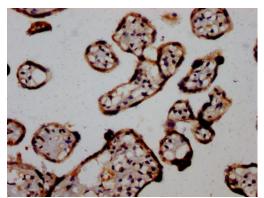
This antibody has been tested in direct-ELISA

## **Product Images For AGO2 Polyclonal Antibody**



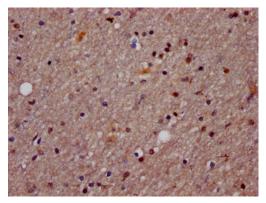
#### AGO2 Antibody (PA5-117725) in WB

Western Blot analysis of AGO2 using AGO2 Polyclonal Antibody (Product # PA5-117725) at 2 µg/mL. Lane 1: A549 whole cell lysate. Lane 2: Jurkats whole cell lysate. Lane 3: MCF-7 whole cell lysate. Lane 4: HepG2 whole cell lysate. Lane 5: Raw264.7 whole cell lysate. Lane 6: K562 whole cell lysate. Lane 7: Mouse liver tissue. Lane 8: Mouse kidney tissue. Secondary: Goat polyclonal to rabbit lgG at 1:10,000 dilution. Predicted band size: 98, 94 kDa. Observed band size: 94 kDa



## AGO2 Antibody (PA5-117725) in IHC (P)

Immnohistochemistry (Paraffin) analysis of AGO2 in paraffin-embedded human placenta tissue using AGO2 Polyclonal Antibody (Product # PA5-117725) diluted at 1:1200. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



## AGO2 Antibody (PA5-117725) in IHC (P)

Immnohistochemistry (Paraffin) analysis of AGO2 in paraffin-embedded human brain tissue using AGO2 Polyclonal Antibody (Product # PA5-117725) diluted at 1:1200. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

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