# **UBE3A Polyclonal Antibody**

Catalog No: #30079

Package Size: #30079-1 50ul #30079-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

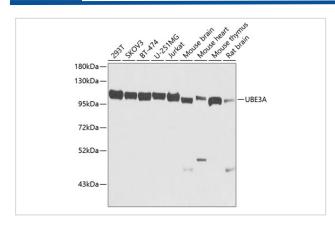
## Description

| Product Name          | UBE3A Polyclonal Antibody   |
|-----------------------|---|
| Host Species          | Rabbit  |
| Clonality             | Polyclonal  |
| Isotype               | IgG   |
| Purification          | Affinity purification   |
| Applications          | WB,IHC  |
| Species Reactivity    | Human,Mouse,Rat   |
| Immunogen Description | Recombinant fusion protein of human UBE3A (NP_570853.1).              |
| Other Names           | UBE3A; ANCR; AS; E6-AP; EPVE6AP; HPVE6A; ubiquitin-protein ligase E3A |
| Accession No.         | Swiss-Prot#:Q05086NCBI Gene ID:7337                                   |
| Uniprot               | Q05086  |
| GeneID                | 7337;   |
| Calculated MW         | 110kDa  |
| Formulation           | Avoid freeze / thaw cycles. Buffer: PBS with 50% glycerol, pH7.4.     |
| Storage               | Store at -20°C  |

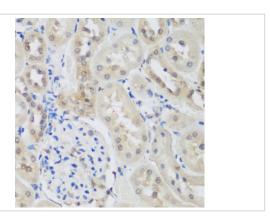
## **Application Details**

WB 1:500 - 1:2000IHC 1:50 - 1:200

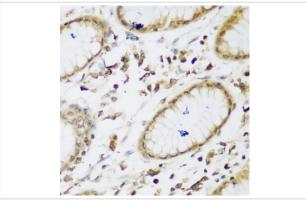
## **Images**



Western blot analysis of extracts of various cell lines, using UBE3A at 1:1000 dilution.



Immunohistochemistry of paraffin-embedded rat kidney using UBE3A at dilution of 1:200 (40x lens).



Immunohistochemistry of paraffin-embedded human gastric using UBE3A at dilution of 1:200 (40x lens).

### Background

This gene encodes an E3 ubiquitin-protein ligase, part of the ubiquitin protein degradation system. This imprinted gene is maternally expressed in brain and biallelically expressed in other tissues. Maternally inherited deletion of this gene causes Angelman Syndrome, characterized by severe motor and intellectual retardation, ataxia, hypotonia, epilepsy, absence of speech, and characteristic facies. The protein also interacts with the E6 protein of human papillomavirus types 16 and 18, resulting in ubiquitination and proteolysis of tumor protein p53. Alternative splicing of this gene results in three transcript variants encoding three isoforms with different N-termini. Additional transcript variants have been described, but their full length nature has not been determined.

Note: This product is for in vitro research use only