# ABclonal www.abclonal.com

# **UFD1L Rabbit pAb**

Catalog No.: A3255

# **Basic Information**

#### **Observed MW**

37kDa

# **Calculated MW**

35kDa

#### Category

Primary antibody

#### **Applications**

ELISA, WB, IF/ICC

#### **Cross-Reactivity**

Human, Mouse

# **Background**

The protein encoded by this gene forms a complex with two other proteins, nuclear protein localization-4 and valosin-containing protein, and this complex is necessary for the degradation of ubiquitinated proteins. In addition, this complex controls the disassembly of the mitotic spindle and the formation of a closed nuclear envelope after mitosis. Mutations in this gene have been associated with Catch 22 syndrome as well as cardiac and craniofacial defects. Alternative splicing results in multiple transcript variants encoding different isoforms. A related pseudogene has been identified on chromosome 18.

# **Recommended Dilutions**

**WB** 1:500 - 1:2000

**IF/ICC** 1:50 - 1:100

# **Immunogen Information**

**Gene ID**7353
Swiss Prot
Q92890

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-307 of human UFD1L (NP\_005650.2).

#### **Synonyms**

UFD1L

### **Contact**

www.abclonal.com

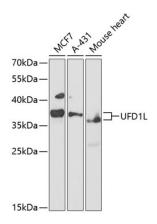
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

# Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.



Western blot analysis of extracts of various cell lines, using UFD1L antibody (A3255) at 1:1000

dilution

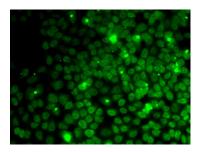
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 90s.



Immunofluorescence analysis of MCF7 cells using UFD1L antibody (A3255).