

## **Product datasheet for TA347079**

## OriGene Technologies, Inc.

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## **ASH2L Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

Applications: ELISA, IF, WB

**Recommended Dilution:** ELISA (1:100 ?? 1:500); Western blotting (1:500 - 1:1,000); IF: (1:200)

Reactivity: Mouse
Host: Rabbit
Isotype: IgG

Clonality: Polyclonal

**Immunogen:** The immunogen for anti-ASH2 antibody: mouse Ash2 (absent, small, or homeotic 2), using 3

different KLH-conjugated synthetic peptides, 2 containing an amino acid sequence from the

central and 1 containing an amino acid sequence from the C-terminal part of

**Concentration:** lot specific

**Purification:** Whole antiserum from rabbit containing 0.05% azide.

**Conjugation:** Unconjugated

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

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

Gene Name: ASH2 like histone lysine methyltransferase complex subunit

Database Link: NP 004665

Entrez Gene 23808 Mouse

Q9UBL3

Background: Ash2 (UniProtKB/Swiss-Prot entry Q9UBL3) is a component of the Set1/Ash2 histone

methyltransferase (HMT) complex. This complex specifically methylates K4 of histone H3, thereby activating transcription. Methylation of K4 is blocked by premethylation of the neighboring K9, a repressor of transcription. This indicates that the Set1/Ash2 HMT complex mediates the crosstalk between K9 methylation and K4 methylation. Ash2 plays a role in

hematopoiesis and may be associated with some kinds of leukemia.

Synonyms: ASH2; ASH2L1; ASH2L2; Bre2

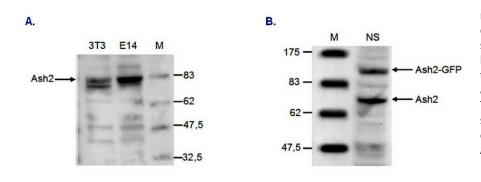
**Protein Families:** Druggable Genome, Transcription Factors



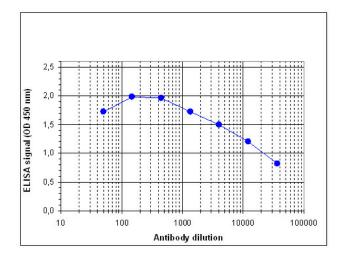


## **Product images:**

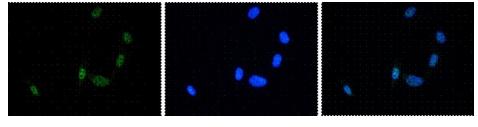
Figure 2



WB was performed on whole cell lysates from mouse fibroblastst (NIH3T3) and embryonic stem cells (E14Tg2a) with the ab at 1:1,000 (predicted size: 68 kDa). B. WB was performed on whole cell lysates from mouse neural stem cells (NS), transfected with GFP tagged Ash2, with the ab against mouse Ash2, diluted 1:500 in BSA/PBS-Tween. The molecular weight marker (in kDa) is shown on the left; the location of the endogenous Ash2 (68 kDa) and of the GFP tagged Ash2 (106 kDa) are indicated on the right.



Determination of the titer To determine the titer, an ELISA was performed using a serial dilution of the antibody against mouse Ash2. The wells were coated with the peptides used for immunisation of the rabbit. By plotting the absorbance against the antibody dilution (Figure 1), the titer of the antibody was estimated to be 1:24,000.



INIH3T3 cells were stained with the ab against Ash2 and with DAPI. Cells were fixed with 4% formaldehyde for 10' and blocked with PBS/TX-100 containing 5% normal goat serum and 1% BSA. The cells were immunofluorescently labelled with the Ash2 antibody (left) diluted 1:200 in blocking solution followed by an anti-rabbit antibody conjugated to Alexa488. The middle panel shows staining of the nuclei with DAPI. A merge of the two stainings is shown on the right.