

## HMGB3 rabbit pAb

<b>Catalog No :</b>	YT7408
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB
<b>Target :</b>	HMGB3
<b>Gene Name :</b>	HMGB3 HMG2A HMG4
<b>Protein Name :</b>	HMGB3
<b>Human Gene Id :</b>	3149
<b>Human Swiss Prot No :</b>	O15347
<b>Mouse Gene Id :</b>	15354
<b>Mouse Swiss Prot No :</b>	O54879
<b>Immunogen :</b>	Synthesized peptide derived from human HMGB3 AA range: 123-173
<b>Specificity :</b>	This antibody detects endogenous levels of HMGB3 at Human/Mouse
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1:500-2000
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)

**Molecularweight :** 22kD

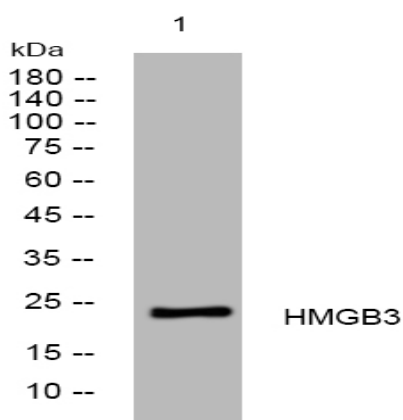
**Background :** This gene encodes a member of a family of proteins containing one or more high mobility group DNA-binding motifs. The encoded protein plays an important role in maintaining stem cell populations, and may be aberrantly expressed in tumor cells. A mutation in this gene was associated with microphthalmia, syndromic 13. There are numerous pseudogenes of this gene on multiple chromosomes. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2014],

**Function :** function: Binds preferentially single-stranded DNA and unwinds double stranded DNA., similarity: Belongs to the HMGB family., similarity: Contains 2 HMG box DNA-binding domains., tissue specificity: Expressed predominantly in placenta.,

**Subcellular Location :** Nucleus . Chromosome . Cytoplasm .

**Expression :** Expressed predominantly in placenta.

## Products Images



Western blot analysis of lysates from U2OS cells, primary antibody was diluted at 1:1000, 4° over night