### **Product datasheet**

### Anti-EIF4A1 Antibody (Clone#3F11)

Catalog Number: M03922-2



Building C21, 3rd and 4th floors, Optics Valley Biomedical Accelerator, Wuhan East Lake High-tech Development Zone

Web: www.boster.com Phone: 027-67845390 Email: boster@boster.com

<b>Basic Inform</b>	nation	
Product Name	Anti-EIF4A1 Antibody (Clone#3F11)	
Gene Name	EIF4A1	
Source	Mouse	
Clonality	Monoclonal	
Isotype	IgG2a	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, FCM	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence of human EIF4A1, identical to the related mouse and rat sequences.	
Concentration	500 ug/ml	
Purification	protein G purified.	
Observed MW	46 kDa	
Dilution Ratios	Western blot (WB): Immunohistochemistry (IHC): Immunocytochemistry/Immunofluorescence (ICC/IF): Flow Cytometry (Fixed): (Boiling the paraffin sections in 10mM citrate buffer,pH6. for 20 mins is required for the staining of formalin/paraffidilutions must be determined by end user.	· · · · · · · · · · · · · · · · · · ·

## **Storage**

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

# **Background Information**

Eukaryotic initiation factor 4A-I is a protein that in humans is encoded by the EIF4A1 gene. It is mapped to 17p13.1. EIF4A1 has been shown to interact with EIF4E and eukaryotic translation initiation factor 4 gamma.

# **Selected Validation Data**

### **Anti-EIF4A1 Antibody (Clone#3F11)**

Catalog Number: M03922-2

**BOSTER BIOLOGICAL TECHNOLOGY**Building C21, 3rd and 4th floors, Optics Valley Biomedical Accelerator,

Wuhan East Lake High-tech Development Zone

Web: www.boster.com Phone: 027-67845390 Email: boster@boster.com

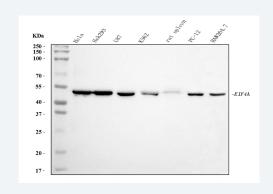


Figure 1. Western blot analysis of EIF4A1 using anti-EIF4A1 antibody (M03922-2). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human HELA whole cell lysates,

Lane 2: human HEK293 whole cell lysates,

Lane 3: human U87 whole cell lysates,

Lane 4: human K562 whole cell lysates,

Lane 5: rat spleen tissue lysates,

Lane 6: rat PC-12 whole cell lysates,

Lane 7: mouse RAW264.7 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with mouse anti-EIF4A1 antigen affinity purified monoclonal antibody (M03922-2) at a dilution of 1:1000 and probed with a goat antimouse IgG-HRP secondary antibody (Catalog # BA1050). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for EIF4A1 at approximately 46 kDa. The expected band size for EIF4A1 is at 46 kDa.

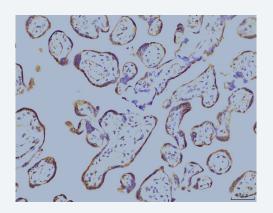


Figure 2. IHC analysis of EIF4A1 using anti-EIF4A1 antibody (M03922-2).

EIF4A1 was detected in a paraffin-embedded section of human placenta tissue. Biotinylated goat anti-mouse IgG was used as secondary antibody. The tissue section was incubated with mouse anti-EIF4A1 Antibody (M03922-2) at a dilution of 1:200 and developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1021) with DAB (Catalog # AR1022) as the chromogen.

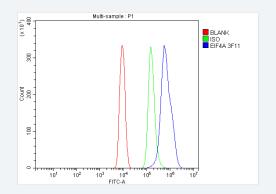


Figure 7. Flow Cytometry analysis of ANA-1 cells using anti-EIF4A1 antibody (M03922-2).

Overlay histogram showing ANA-1 cells stained with M03922-2 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with mouse anti-EIF4A1 Antibody (M03922-2) at 1:100 dilution for 30 min at

### **Product datasheet**

## **Anti-EIF4A1 Antibody (Clone#3F11)**

Catalog Number: M03922-2



**BOSTER BIOLOGICAL TECHNOLOGY** 

Building C21, 3rd and 4th floors, Optics Valley Biomedical Accelerator, Wuhan East Lake High-tech Development Zone

Web: www.boster.com Phone: 027-67845390 Email: boster@boster.com

20°C. DyLight®488 conjugated goat anti-mouse IgG (BA1126) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was mouse IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.