### bsm-33118M

## [ Primary Antibody ]

# Bioss ANTIBODIES

www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

### Peroxiredoxin 1 Mouse mAb

- DATASHEET -

Host: Mouse Isotype: IgG
Clonality: Monoclonal CloneNo.: 10F9
GeneID: 5052 SWISS: Q06830

**Target:** Peroxiredoxin 1

Purification: affinity purified by Protein G

Concentration: 1mg/ml

Storage: Size: 50ul/100ul/200ul

0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Size: 200ug (PBS only)

0.01M PBS

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

**Background:** This gene encodes a member of the peroxiredoxin family of

antioxidant enzymes, which reduce hydrogen peroxide and alkyl hydroperoxides. The encoded protein may play an antioxidant protective role in cells, and may contribute to the antiviral activity of CD8(+) T-cells. This protein may have a proliferative effect and play a role in cancer development or progression. Four transcript variants encoding the same protein have been identified for this

gene. [provided by RefSeq, Jan 2011].

Applications: WB (1:500-1000)

IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500)

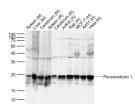
Flow-Cyt (0.5ug/Test)

Reactivity: Human, Mouse, Rat

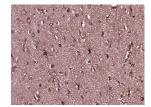
Predicted MW.: 22 kDa

**Subcellular Location:** Cell membrane ,Cytoplasm

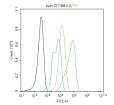
#### VALIDATION IMAGES



Sample: Lane 1: Mouse Spleen tissue lysates
Lane 2: Mouse Liver tissue lysates Lane 3: Mouse
Cerebrum tissue lysates Lane 4: Rat Spleen
tissue lysates Lane 5: Rat Cerebrum tissue
lysates Lane 6: Human Jurkat cell lysates Lane 7:
Human Raji cell lysates Lane 8: Human MCF-7
cell lysates Lane 9: Human A431 cell lysates Lane
10: Human Hela cell lysates Primary: AntiPeroxiredoxin 1 (bsm-33118M) at 1/1000 dilution
Secondary: IRDye800CW Goat Anti-Mouse IgG at
1/20000 dilution Predicted band size: 22 kD
Observed band size: 23 kD



Paraformaldehyde-fixed, paraffin embedded (Human brain glioma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Peroxiredoxin 1) Monoclonal Antibody, Unconjugated (ascites of bsm-33118M 10F9) at 1:2000 overnight at 4°C, followed by a conjugated secondary (sp-0024) for 20 minutes and DAB staining.



Blank control (black line) :Hela. Primary Antibody (green line): Mouse Anti-Peroxiredoxin 1 antibody (bsm-33118M) Dilution:0.5ug/Test; Secondary Antibody (white blue line): Goat anti-mouse IgG-FITC Dilution: 0.5ug/Test. Isotype control (orange line): Normal Mouse IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.