MONOCLONAL ANTIBODY



# **Anti-Peroxiredoxin V (3F11)**

**Background**: Peroxiredoxin (Prx) is a growing peroxidase family, whose mammalian members have been known to connect with cell proliferation, differentiation, and apoptosis.

Many isoforms (about 50 proteins), collected in accordance to the amino acid sequence homology, particularly amino-terminal region containing active site cysteine residue, and the thiol-specific antioxidant activity, distribute throughout all the kingdoms. Among them, mammalian Prx consists of 6 different members grouped into typical 2-Cys, atypical 2-Cys Prx, and 1-Cys Prx. Except Prx VI belonging to 1-Cys Prx subgroup, the other five 2-Cys Prx isotypes have the thioredoxin-dependent peroxidase utilizing activity thioredoxin, thioredoxin reductase, and NADPH as a reducing system. Mammalian Prxs are 20 -30 kilodalton in molecular size and vary in subcellular localization: Prx I, II, and VI in cytosol, Prx III in mitochondria, Prx IV in ER and secretion, Prx V showing complicated distribution including peroxisome, mitochondria and cytosol.

Species cross reactivity		
Human	Mouse	Rat

**Immunogen :** Recombinant human Prx V fragment protein (without mitochondrial leader sequence) purified from *E.coli* 

# **Applications:**

ELISA

Western Blotting (1:1000) Immunoprecipitation (1-2ul/400ml lysates) Immunocytochemistry (20ul/ml)

**Host:** Mouse

**Isotype :** IgG1, k

Clone number: 3F11

**Concentration :** PBS containing 50% glycerol

**Positive control :** HeLa cell lysates

**Size**: 100ul

# **Background Reference:**

(1) Wood, Z. A. et al. (2003) *Trends Biochem Sci.* **28**(1):32-40

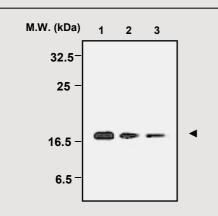
(2) Rhee Sue Goo et al. (2001) *IUBMB Life*. **52**:35-41

**Storage :** Store for 1 year at -20°C from date

of shipment

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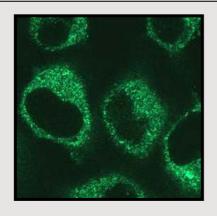


IMMUNOBLOT ANALYSIS of cell lysates using Prx V (LF-MA0002) Antibody

Lane 1: HeLa cell lysates

Lane 2: 293T cell lysates

Lane 3: Jurkat cell lysates



IMMUNOCYTOCHEMICAL STAINING of Human HeLa cells using Prx V(LF-MA0002) Antibody

# **Immunocytochemistry Protocol**

#### **Fixation**

- Cells were cultured on coverslips.
- Immerse coverslips in 4% paraformaldehyde for 40 minutes at room temperature.
- Wash coverslips with 1xPBS 2 times.
- Quench cells in 50mM NH<sub>4</sub>Cl for 10 minutes at room temperature.
- Aspirate off completely.
- Wash the coverslips with 1xPBS 2 times.
- Aspirate off completely.
- Permeabilize cells on coverslips with 0.2% triton X-100 for 10 minutes at room temperature.

## **Blocking**

- Block all coverslips with blocking solution for 30 minutes at room temperature.

## **Staining**

- Incubate with primary antibody in blocking solution(20ug/ml) for 1 hour at room temperature.
- Wash all coverslips with blocking solution for 10 minutes 3 times.
- Incubate all coverslips with a dilution of the Alexa green conjugated secondary antibody in blocking solution.
- Wash all coverslips with 1xPBS for 10 minutes 3 times.

# Mounting and Observation

- Mount coverslips on slides.
- Store slides at room temperature in the dark.
- Evaluate with fluorescence microscope.
- \* Blocking Solution : 2% BSA in PBS

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