

## PAM16 Polyclonal Antibody

catalog number: E-AB-53027

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

### Description

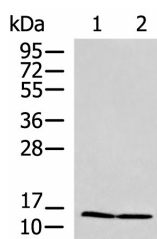
|              |  |
|--------------|--|
| Reactivity   | Human;Mouse  |
| Immunogen    | Fusion protein of human PAM16  |
| Host         | Rabbit   |
| Isotype      | IgG  |
| Purification | Antigen affinity purification  |
| Conjugation  | Unconjugated   |
| Buffer       | Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol. |

### Applications

### Recommended Dilution

|     |              |
|-----|--------------|
| WB  | 1:500-1:2000 |
| IHC | 1:50-1:200   |

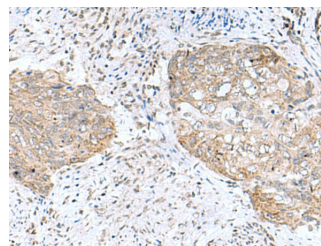
### Data



Western blot analysis of HepG2 and Jurkat cell lysates using PAM16 Polyclonal Antibody at dilution of 1:700

**Observed-MW:Refer to figures**

**Calculated-MW:14 kDa**



Immunohistochemistry of paraffin-embedded Human cervical cancer tissue using PAM16 Polyclonal Antibody at dilution of 1:50(×200)

### Preparation & Storage

|          |  |
|----------|--|
| Storage  | Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.  |
| Shipping | The product is shipped with ice pack, upon receipt, store it immediately at the temperature recommended. |

### Background

This gene encodes a mitochondrial protein involved in granulocyte-macrophage colony-stimulating factor (GM-CSF) signaling. This protein also plays a role in the import of nuclear-encoded mitochondrial proteins into the mitochondrial matrix and may be important in reactive oxygen species (ROS) homeostasis. Mutations in this gene cause Megarbane-Dagher-Melike type spondylometaphyseal dysplasia, an early lethal skeletal dysplasia characterized by short stature, developmental delay and other skeletal abnormalities.

### For Research Use Only