

## PYGM rabbit pAb

| Catalog No         | YP-Ab-09040  |
|--------------------|--|
| Isotype            | IgG  |
| Reactivity         | Human; Mouse;Rat   |
| Applications       | WB   |
| Gene Name          | PYGM   |
| Protein Name       | PYGM   |
| Immunogen          | Synthesized peptide derived from human PYGM AA range: 426-476  |
| Specificity        | This antibody detects endogenous levels of PYGM at Human/Mouse/Rat   |
| Formulation        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.  |
| Source             | Polyclonal, Rabbit,IgG   |
| Purification       | The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.  |
| Dilution           | WB 1: 500-2000   |
| Concentration      | 1 mg/ml  |
| Purity             | ≥90%   |
| Storage Stability  | -20°C/1 year   |
| Synonyms           |  |
| Observed Band      |  |
| Cell Pathway       | cytoplasm,cytosol,extracellular exosome,   |
| Tissue Specificity |  |
| Function           | catalytic activity:(1,4-alpha-D-glucosyl)(n) + phosphate = (1,4-alpha-D-glucosyl)(n-1) + alpha-D-glucose 1-phosphate.,cofactor:Pyridoxal phosphate.,disease:Defects in PYGM are the cause of glycogen storage disease type 5 (GSD5) [MIM:232600]; also known as McArdle disease. GSD5 is a metabolic disorder resulting in myopathy characterized by exercise intolerance, cramps, muscle weakness and recurrent myoglobinuria.,enzyme regulation:Activity of phosphorylase is controlled both by allosteric means (through the noncovalent binding of metabolites) and by covalent modification. Thus AMP allosterically activates, whereas ATP, ADP, and glucose-6-phosphate allosterically inhibit, phosphorylase B.,function:Phosphorylase is an important allosteric enzyme in carbohydrate metabolism. Enzymes from different sources differ in their regulatory mechanisms and in their natural substrates. However, all know |
| Background         | This gene encodes a muscle enzyme involved in glycogenolysis. Highly similar enzymes encoded by different genes are found in liver and brain. Mutations in this  |
|                    |  |



## UpingBio technology Co.,Ltd

C Tel: 400-999-8863 🛎 Email:UpingBio@163.com



gene are associated with McArdle disease (myophosphorylase deficiency), a glycogen storage disease of muscle. Alternative splicing results in multiple transcript variants.[provided by RefSeq, Sep 2009],

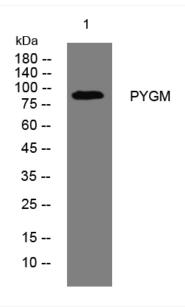
matters needing attention

Avoid repeated freezing and thawing!

**Usage suggestions** 

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## **Products Images**



Western blot analysis of lysates from MCF-7 cells, primary antibody was diluted at 1:1000, 4° over night