

# PYGM Antibody (C-term)

Purified Rabbit Polyclonal Antibody (Pab)  
Catalog # AP1450b

## Product Information

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|--------------------------|------------------------|
| <b>Application</b>       | WB, IHC-P, E           |
| <b>Primary Accession</b> | <a href="#">P11217</a> |
| <b>Other Accession</b>   | <a href="#">Q8HXW4</a> |
| <b>Reactivity</b>        | Human                  |
| <b>Predicted</b>         | Monkey                 |
| <b>Host</b>              | Rabbit                 |
| <b>Clonality</b>         | Polyclonal             |
| <b>Isotype</b>           | Rabbit IgG             |
| <b>Clone Names</b>       | RB11945                |
| <b>Calculated MW</b>     | 97092                  |
| <b>Antigen Region</b>    | 698-727                |

## Additional Information

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|---------------------------|---|
| <b>Gene ID</b>            | 5837  |
| <b>Other Names</b>        | Glycogen phosphorylase, muscle form, Myophosphorylase, PYGM   |
| <b>Target/Specificity</b> | This PYGM antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 698-727 amino acids from the C-terminal region of human PYGM.        |
| <b>Dilution</b>           | WB~~1:1000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.   |
| <b>Format</b>             | Purified polyclonal antibody supplied in PBS with 0.05% (V/V) Proclin 300. This antibody is purified through a protein A column, followed by peptide affinity purification. |
| <b>Storage</b>            | Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.                                     |
| <b>Precautions</b>        | PYGM Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.  |

## Protein Information

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|-----------------|--|
| <b>Name</b>     | PYGM ( <a href="#">HGNC:9726</a> )   |
| <b>Function</b> | Allosteric enzyme that catalyzes the rate-limiting step in glycogen catabolism, the phosphorolytic cleavage of glycogen to produce glucose-1-phosphate, and plays a central role in maintaining cellular and |

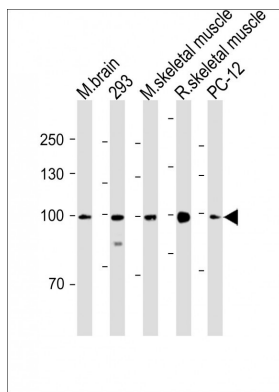
## Background

PYGM catalyzes and regulates the breakdown of glycogen to glucose-1-phosphate. Defects in PYGM are the cause of glycogen storage disease type 5 (GSD5), also known as McArdle disease. GSD5 is a metabolic disorder resulting in myopathy characterized by exercise intolerance, cramps, muscle weakness and recurrent myoglobinuria.

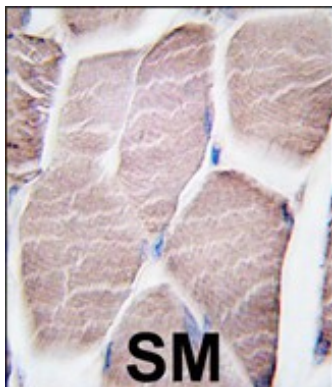
## References

- Tsoi, S.C., et al., *J. Soc. Gynecol. Investig.* 10(8):496-502 (2003).  
Bruno, C., et al., *Neuromuscul. Disord.* 12(5):498-500 (2002).  
Hadjigeorgiou, G.M., et al., *Neuromuscul. Disord.* 12(9):824-827 (2002).  
Deschauer, M., et al., *Mol. Genet. Metab.* 74(4):489-491 (2001).  
Kubisch, C., et al., *Hum. Mutat.* 12(1):27-32 (1998).

## Images



All lanes: Anti-PYGM Antibody (C-term) at 1:2000 dilution  
Lane 1: Mouse brain lysate Lane 2: 293 whole cell lysate  
Lane 3: Mouse skeletal muscle lysate Lane 4: Rat skeletal muscle lysate Lane 5: PC-12 whole cell lysate  
Lysates/proteins at 20 µg per lane. Secondary: Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated (ASP1615) at 1/15000 dilution. Observed band size: 97 KDa  
Blocking/Dilution buffer: 5% NFDN/TBST.



Formalin-fixed and paraffin-embedded human skeletal muscle tissue reacted with PYGM antibody (C-term) (Cat.#AP1450b), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.