

## GAA Polyclonal Antibody

<b>Catalog No.</b>	A7674	<b>Category</b>	Polyclonal Antibodies
<b>Applications</b>	WB, IF, IP	<b>Observed MW</b>	105kDa
<b>Cross-reactivity</b>	Human, Mouse, Rat	<b>Calculated MW</b>	105kDa

### Immunogen Information

<b>Immunogen</b>	A synthetic peptide corresponding to a sequence within amino acids 350-450 of human GAA (NP_000143.2).
<b>Gene ID</b>	2548
<b>Swiss prot</b>	P10253
<b>Synonyms</b>	GAA; LYAG; glucosidase alpha, acid

### Product information

<b>Source</b>	Rabbit
<b>Isotype</b>	IgG
<b>Purification method</b>	Affinity purification
<b>Storage</b>	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

### Background

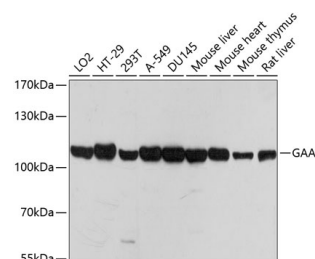
This gene encodes lysosomal alpha-glucosidase, which is essential for the degradation of glycogen to glucose in lysosomes. The encoded preproprotein is proteolytically processed to generate multiple intermediate forms and the mature form of the enzyme. Defects in this gene are the cause of glycogen storage disease II, also known as Pompe's disease, which is an autosomal recessive disorder with a broad clinical spectrum. Alternative splicing results in multiple transcript variants.

### Recommended Dilutions

WB 1:500 -  
1:2000

IF 1:50 -  
1:200

IP 1:50 -  
1:200



Western blot - GAA Polyclonal Antibody (A7674)