



# Anti-ACP5 (aa 221-325) polyclonal antibody (DPAB-DC2297)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Antigen Description</b>	This gene encodes an iron containing glycoprotein which catalyzes the conversion of orthophosphoric monoester to alcohol and orthophosphate. It is the most basic of the acid phosphatases and is the only form not inhibited by L(+)-tartrate.
<b>Immunogen</b>	ACP5 (NP_001602, 221 a.a. ~ 325 a.a) partial recombinant protein with GST tag. The sequence is VKQLRPLLATYGV TAYLCGHDHNLQYLQDENG VGYVLSGAGNFMDPSKRHQKVPNGYLR FHYGTEDSLGGFAYVEISSKEMTVTYIEASGKSLFKTRLPRRARP
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB (Recombinant protein), ELISA,
<b>Size</b>	50 µl
<b>Buffer</b>	50 % glycerol
<b>Preservative</b>	None
<b>Storage</b>	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## GENE INFORMATION

**Gene Name** [ACP5 acid phosphatase 5, tartrate resistant \[ Homo sapiens \(human\) \]](#)

<b>Official Symbol</b>	ACP5
<b>Synonyms</b>	ACP5; acid phosphatase 5, tartrate resistant; TRAP; SPENCDI; tartrate-resistant acid phosphatase type 5; TrATPase; tartrate-resistant acid ATPase;
<b>Entrez Gene ID</b>	<a href="#">54</a>
<b>Protein Refseq</b>	<a href="#">NP_001104504</a>
<b>UniProt ID</b>	<a href="#">A0A024R7F8</a>
<b>Chromosome Location</b>	19p13.2
<b>Pathway</b>	Defective AMN causes hereditary megaloblastic anemia 1; Defective CD320 causes methylmalonic aciduria; Defective GIF causes intrinsic factor deficiency; Defective LMBRD1 causes methylmalonic aciduria and homocystinuria type cblF
<b>Function</b>	acid phosphatase activity; ferric iron binding; ferrous iron binding;