



# Mouse anti-Human FPGS polyclonal antibody (DPAB-DC1119)

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

## PRODUCT INFORMATION

<b>Antigen Description</b>	This gene encodes the folylpolyglutamate synthetase enzyme. This enzyme has a central role in establishing and maintaining both cytosolic and mitochondrial folylpolyglutamate concentrations and, therefore, is essential for folate homeostasis and the survival of proliferating cells. This enzyme catalyzes the ATP-dependent addition of glutamate moieties to folate and folate derivatives. Alternative splicing results in transcript variants encoding different isoforms.
<b>Immunogen</b>	FPGS (NP_004948, 135 a.a. ~ 232 a.a) partial recombinant protein with GST tag. The sequence is QVRERIRINGQPISPELFTKYFWRLYHRLEETKDGSCVSMPPYFRFLTMAFHVFLQEKV DLAVVEVGIGGAYDCTNIIRKPVVCGVSSLGIDHTSLL
<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

<b>Gene Name</b>	<a href="#">FPGS folypolyglutamate synthase [ Homo sapiens (human) ]</a>
<b>Official Symbol</b>	FPGS
<b>Synonyms</b>	FPGS; folypolyglutamate synthase; folypolyglutamate synthase, mitochondrial; tetrahydrofolate synthase; folypoly-gamma-glutamate synthetase; tetrahydrofolypolyglutamate synthase;
<b>Entrez Gene ID</b>	<a href="#">2356</a>
<b>Protein Refseq</b>	<a href="#">NP_001018088</a>
<b>UniProt ID</b>	<a href="#">Q05932</a>
<b>Chromosome Location</b>	9q34.1
<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>	ATP binding; tetrahydrofolypolyglutamate synthase activity;