



# Anti-SPG21 (aa 211-306) polyclonal antibody (DPAB-DC2155)

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

## PRODUCT INFORMATION

<b>Antigen Description</b>	The protein encoded by this gene binds to the hydrophobic C-terminal amino acids of CD4 which are involved in repression of T cell activation. The interaction with CD4 is mediated by the noncatalytic alpha/beta hydrolase fold domain of this protein. It is thus proposed that this gene product modulates the stimulatory activity of CD4. Mutations in this gene are associated with autosomal recessive spastic paraplegia 21 (SPG21), also known as mast syndrome. At least three different transcript variants encoding two different isoforms have been found for this gene.
<b>Immunogen</b>	SPG21 (NP_057714, 211 a.a. ~ 306 a.a) partial recombinant protein with GST tag. The sequence is PHKIRDIPVTIMDVFDQSSALSTEAKEEMYKLYPNARRAHLKTGGNFPYLCRSAEVNLYVQ IHLLQFHGTYAAIDPSMVSAEELEVQKGLGISQE
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB (Cell lysate), 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="#">SPG21 spastic paraplegia 21 (autosomal recessive, Mast syndrome) [ Homo sapiens (human) ]</a>
<b>Official Symbol</b>	SPG21
<b>Synonyms</b>	SPG21; spastic paraplegia 21 (autosomal recessive, Mast syndrome); MAST; ACP33; GL010; BM-019; maspardin; acid cluster protein 33;
<b>Entrez Gene ID</b>	<a href="#">51324</a>
<b>Protein Refseq</b>	<a href="#">NP_001121361</a>
<b>UniProt ID</b>	<a href="#">A0A024R5Y1</a>
<b>Chromosome Location</b>	15q21-q22
<b>Function</b>	CD4 receptor binding;