



# Anti-SLC6A19 (aa 326-414) polyclonal antibody (DPAB-DC1655)

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

## PRODUCT INFORMATION

<b>Antigen Description</b>	This gene encodes a system B(0) transmembrane protein that actively transports most neutral amino acids across the apical membrane of epithelial cells. Mutations in this gene result in Hartnup disorder.
<b>Immunogen</b>	SLC6A19 (NP_001003841, 326 a.a. ~ 414 a.a) partial recombinant protein with GST tag. The sequence is GFRATQRYDDCFSTNILTILINGFDLPEGNVTQENFVDMQQRCNASDPAAYAQLVFQTCDI NAFLSEAVEGTGLAFIVFTEAITKMPLSP
<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="#">SLC6A19 solute carrier family 6 (neutral amino acid transporter), member 19 [ Homo sapiens (human) ]</a>
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<b>Official Symbol</b>	SLC6A19
<b>Synonyms</b>	SLC6A19; solute carrier family 6 (neutral amino acid transporter), member 19; HND; B0AT1; sodium-dependent neutral amino acid transporter B(0)AT1; solute carrier family 6 member 19; system B0 neutral amino acid transporter; system B(0) neutral amino acid transporter AT1; sodium-dependent amino acid transporter system B0; solute carrier family 6 (neurotransmitter transporter), member 19;
<b>Entrez Gene ID</b>	<a href="#">340024</a>
<b>Protein Refseq</b>	<a href="#">NP_001003841</a>
<b>UniProt ID</b>	<a href="#">Q695T7</a>
<b>Chromosome Location</b>	5p15.33
<b>Pathway</b>	Amino acid transport across the plasma membrane; Mineral absorption; Protein digestion and absorption; SLC-mediated transmembrane transport
<b>Function</b>	neurotransmitter:sodium symporter activity; neutral amino acid transmembrane transporter activity;