

## ALG11 Conjugated Antibody

Catalog No: #C37327



Package Size: #C37327-AF350 100ul #C37327-AF405 100ul #C37327-AF488 100ul  
 #C37327-AF555 100ul #C37327-AF594 100ul #C37327-AF647 100ul  
 #C37327-AF680 100ul #C37327-AF750 100ul #C37327-Biotin 100ul

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## Description

Product Name	ALG11 Conjugated Antibody
Host Species	Rabbit
Clonality	Polyclonal
Species Reactivity	Hu Ms
Specificity	The antibody detects endogenous levels of total ALG11 protein.
Immunogen Description	Synthetic peptide corresponding to a region derived from internal residues of human ALG11, alpha-1,2-mannosyltransferase
Conjugates	Biotin AF350 AF405 AF488 AF555 AF594 AF647 AF680 AF750
Other Names	GT8; CDG1P
Accession No.	Swiss-Prot#:Q2TAA5NCBI Gene ID:440138NCBI Protein#:NP_037471
Uniprot	Q2TAA5
GeneID	440138;
Excitation Emission	AF350: 346nm/442nm AF405: 401nm/421nm AF488: 493nm/519nm AF555: 555nm/565nm AF594: 591nm/614nm AF647: 651nm/667nm AF680: 679nm/702nm AF750: 749nm/775nm
Formulation	0.01M Sodium Phosphate, 0.25M NaCl, pH 7.6, 5mg/ml Bovine Serum Albumin, 0.02% Sodium Azide
Storage	Store at 4°C in dark for 6 months

## Application Details

Suggested Dilution:

AF350 conjugated: most applications: 1: 50 - 1: 250

AF405 conjugated: most applications: 1: 50 - 1: 250

AF488 conjugated: most applications: 1: 50 - 1: 250

AF555 conjugated: most applications: 1: 50 - 1: 250

AF594 conjugated: most applications: 1: 50 - 1: 250

AF647 conjugated: most applications: 1: 50 - 1: 250

AF680 conjugated: most applications: 1: 50 - 1: 250

AF750 conjugated: most applications: 1: 50 - 1: 250

Biotin conjugated: working with enzyme-conjugated streptavidin, most applications: 1: 50 - 1: 1,000

## Background

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This gene encodes a GDP-Man:Man3GlcNAc2-PP-dolichol-alpha 1,2-mannosyltransferase which is localized to the cytosolic side of the endoplasmic reticulum (ER) and catalyzes the transfer of the fourth and fifth mannose residue from GDP-mannose (GDP-Man) to Man3GlcNAc2-PP-dolichol and Man4GlcNAc2-PP-dolichol resulting in the production of Man5GlcNAc2-PP-dolichol. Mutations in this gene are associated with congenital disorder of glycosylation type I<sub>p</sub> (CDGIP).?

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Note: This product is for in vitro research use only