

## LRP5 Conjugated Antibody

Catalog No: #C36957



Package Size: #C36957-AF350 100ul #C36957-AF405 100ul #C36957-AF488 100ul  
 #C36957-AF555 100ul #C36957-AF594 100ul #C36957-AF647 100ul  
 #C36957-AF680 100ul #C36957-AF750 100ul #C36957-Biotin 100ul

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## Description

Product Name	LRP5 Conjugated Antibody
Host Species	Rabbit
Clonality	Polyclonal
Species Reactivity	Hu Ms
Specificity	The antibody detects endogenous levels of total LRP5 protein.
Immunogen Description	Synthetic peptide corresponding to a region derived from internal residues of human low density lipoprotein receptor-related protein 5
Conjugates	Biotin AF350 AF405 AF488 AF555 AF594 AF647 AF680 AF750
Other Names	HBM, LR3, OPS, EVR1, EVR4, LRP7, OPPG, BMND1, OPTA1, VBCH2
Accession No.	Swiss-Prot#:O75197NCBI Gene ID:4041NCBI Protein#:NP_002326
Uniprot	O75197
GeneID	4041;
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 transmembrane low-density lipoprotein receptor that binds and internalizes ligands in the process of receptor-mediated endocytosis. This protein also acts as a co-receptor with Frizzled protein family members for transducing signals by Wnt proteins and was originally cloned on the basis of its association with type 1 diabetes mellitus in humans. This protein plays a key role in skeletal homeostasis and many bone density related diseases are caused by mutations in this gene. Mutations in this gene also cause familial exudative vitreoretinopathy.

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Note: This product is for in vitro research use only