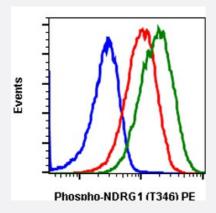


#### RecomAb™

# NDRG1 recombinant monoclonal antibody, clone NDRG1T346-F5 (PE)

Catalog # RAB02933 Size 100 Reactions

# Applications



#### Flow Cytometry

Flow cytometric of THP1 cells unstained and untreated as negative control (blue) or stained and untreated (red) or stained and treated with IFNa plus IL-4 and pervanadate (green) using phospho-NDRG1 (Thr346) (F5) rabbit mAb, NDRG1T346-F5 PE conjugate.

### Specification

Product Description	Rabbit recombinant monoclonal antibody raised against human NDRG1.
Antibody Species	Rabbit
Immunogen	A synthetic phospho-peptide corresponding to residues surrounding Thr346 of human phospho NDR G1
Reactivity	Human
Form	Liquid
Conjugation	PE
Purification	Protein A purification, Protein G purification
lsotype	lgG



#### **Product Information**

Recommend Usage	Flow Cytometry The optimal working dilution should be determined by the end user.
Storage Buffer	1X PBS, 0.09% Sodium azide, 0.2% BSA
Storage Instruction	Store at 4°C. Do not freeze.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

# Applications

Flow Cytometry

Flow cytometric of THP1 cells unstained and untreated as negative control (blue) or stained and untreated (red) or stained and treated with IFNa plus IL-4 and pervanadate (green) using phospho-NDRG1 (Thr346) (F5) rabbit mAb, NDRG1T346-F5 PE conjugate.

# Gene Info — NDRG1

Entrez GenelD	<u>10397</u>
Protein Accession#	<u>Q92597</u>
Gene Name	NDRG1
Gene Alias	CAP43, CMT4D, DRG1, GC4, HMSNL, NDR1, NMSL, PROXY1, RIT42, RTP, TARG1, TDD5
Gene Description	N-myc downstream regulated 1
Omim ID	<u>601455 605262</u>
Orana Oratala ana	
Gene Ontology	<u>Hyperlink</u>
Gene Ontology Gene Summary	Hyperlink This gene is a member of the N-myc downregulated gene family which belongs to the alpha/beta hydrolase superfamily. The protein encoded by this gene is a cytoplasmic protein involved in stres s responses, hormone responses, cell growth, and differentiation. It is necessary for p53-mediate d caspase activation and apoptosis. Mutation in this gene has been reported to be causative for h ereditary motor and sensory neuropathy-Lom. Multiple alternatively spliced variants, encoding the same protein, have been identified. [provided by RefSeq

Disease

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- Alzheimer disease
- <u>Charcot-Marie-Tooth Disease</u>
- <u>Cognition</u>
- <u>Colorectal Neoplasms</u>
- Deafness
- Genetic Predisposition to Disease
- <u>Tobacco Use Disorder</u>