

NR2F2 Antibody (monoclonal) (M02)

Mouse monoclonal antibody raised against a partial recombinant NR2F2. Catalog # AT3104a

Specification

NR2F2 Antibody (monoclonal) (M02) - Product Information

Application E
Primary Accession P24468
Other Accession NM_021005
Reactivity Human
Host mouse
Clonality Monoclonal
Isotype IgG2b Kappa

Calculated MW 45571

NR2F2 Antibody (monoclonal) (M02) - Additional Information

Gene ID 7026

Other Names

COUP transcription factor 2, COUP-TF2, Apolipoprotein A-I regulatory protein 1, ARP-1, COUP transcription factor II, COUP-TF II, Nuclear receptor subfamily 2 group F member 2, NR2F2, ARP1, TFCOUP2

Target/Specificity

NR2F2 (NP_066285, 153 a.a. \sim 240 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Dilution

E~~N/A

Format

Clear, colorless solution in phosphate buffered saline, pH 7.2.

Storage

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Precautions

NR2F2 Antibody (monoclonal) (M02) is for research use only and not for use in diagnostic or therapeutic procedures.

NR2F2 Antibody (monoclonal) (M02) - Protocols

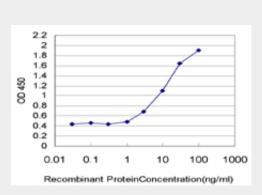
Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides



- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

NR2F2 Antibody (monoclonal) (M02) - Images



Detection limit for recombinant GST tagged NR2F2 is approximately 1ng/ml as a capture antibody.

NR2F2 Antibody (monoclonal) (M02) - Background

This gene encodes a member of the steroid thyroid hormone superfamily of nuclear receptors. The encoded protein is a ligand inducible transcription factor that is involved in the regulation of many different genes. Alternate splicing results in multiple transcript variants.

NR2F2 Antibody (monoclonal) (M02) - References

Variation at the NFATC2 Locus Increases the Risk of Thiazolinedinedione-Induced Edema in the Diabetes REduction Assessment with ramipril and rosiglitazone Medication (DREAM) Study. Bailey SD, et al. Diabetes Care, 2010 Jul 13. PMID 20628086.Web-based, participant-driven studies yield novel genetic associations for common traits. Eriksson N, et al. PLoS Genet, 2010 Jun 24. PMID 20585627.Genome-wide searching of rare genetic variants in WTCCC data. Feng T, et al. Hum Genet, 2010 Sep. PMID 20549515.An exquisite cross-control mechanism among endothelial cell fate regulators directs the plasticity and heterogeneity of lymphatic endothelial cells. Kang J, et al. Blood, 2010 Jul 8. PMID 20351309.COUP-TFs regulate eye development by controlling factors essential for optic vesicle morphogenesis. Tang K, et al. Development, 2010 Mar. PMID 20147377.