

# TCF2 Antibody (monoclonal) (M09)

Mouse monoclonal antibody raised against a partial recombinant TCF2. Catalog # AT4185a

### **Product Information**

Application WB
Primary Accession P35680
Other Accession NM\_000458
Reactivity Human
Host mouse
Clonality monoclonal
Isotype IgG2a Kappa

Clone Names 3E5 Calculated MW 61324

### **Additional Information**

Gene ID 6928

Other Names Hepatocyte nuclear factor 1-beta, HNF-1-beta, HNF-1B, Homeoprotein LFB3,

Transcription factor 2, TCF-2, Variant hepatic nuclear factor 1, vHNF1, HNF1B,

TCF2

**Target/Specificity** TCF2 (NP\_000449, 29 a.a. ~ 118 a.a) partial recombinant protein with GST tag.

MW of the GST tag alone is 26 KDa.

**Dilution** WB~~1:500~1000

**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** TCF2 Antibody (monoclonal) (M09) is for research use only and not for use in

diagnostic or therapeutic procedures.

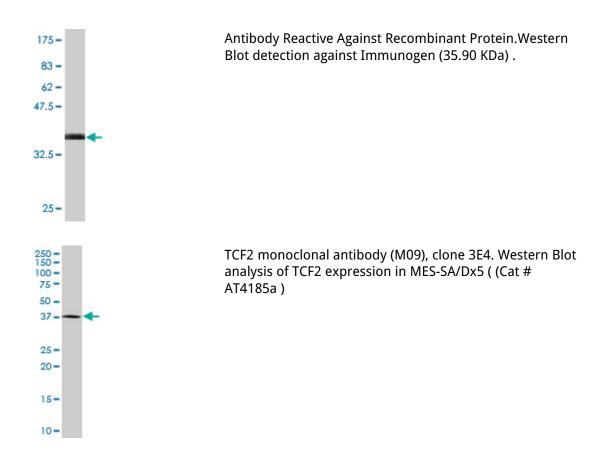
## **Background**

This gene encodes a member of the homeodomain-containing superfamily of transcription factors. The protein binds to DNA as either a homodimer, or a heterodimer with the related protein hepatocyte nuclear factor 1-alpha. The gene has been shown to function in nephron development, and regulates development of the embryonic pancreas. Mutations in this gene result in renal cysts and diabetes syndrome and noninsulin-dependent diabetes mellitus, and expression of this gene is altered in some types of cancer. Multiple transcript variants encoding different isoforms have been found for this gene.

### References

Meta-analysis of genome-wide and replication association studies on prostate cancer. Liu H, et al. Prostate, 2010 Aug 5. PMID 20690139.COMMON VARIANTS IN 40 GENES ASSESSED FOR DIABETES INCIDENCE AND RESPONSE TO METFORMIN AND LIFESTYLE INTERVENTIONS IN THE DIABETES PREVENTION PROGRAM. Jablonski KA, et al. Diabetes, 2010 Aug 3. PMID 20682687.Mutations in the hepatocyte nuclear factor-1beta (HNF1B) gene are common with combined uterine and renal malformations but are not found with isolated uterine malformations. Oram RA, et al. Am J Obstet Gynecol, 2010 Jul 14. PMID 20633866.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.Number of prostate cancer risk alleles may identify possibly 'insignificant' disease. Helfand BT, et al. BJU Int, 2010 Jun 30. PMID 20590552.

### **Images**



Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.