



# HNF-1 $\beta$ Polyclonal Antibody

<b>Catalog No</b>	YP-Ab-02222
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Mouse;Rat
<b>Applications</b>	WB;ELISA
<b>Gene Name</b>	HNF1B
<b>Protein Name</b>	Hepatocyte nuclear factor 1-beta
<b>Immunogen</b>	Synthesized peptide derived from the N-terminal region of human HNF-1 $\beta$ .
<b>Specificity</b>	HNF-1 $\beta$ Polyclonal Antibody detects endogenous levels of HNF-1 $\beta$ protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Dilution</b>	Western Blot: 1/500 - 1/2000. ELISA: 1/10000. Not yet tested in other applications.
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	$\geq 90\%$
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	HNF1B; TCF2; Hepatocyte nuclear factor 1-beta; HNF-1-beta; HNF-1B; Homeoprotein LFB3; Transcription factor 2; TCF-2; Variant hepatic nuclear factor 1; vHNF1
<b>Observed Band</b>	60kD
<b>Cell Pathway</b>	Nucleus.
<b>Tissue Specificity</b>	Colon,Liver,Thalamus,
<b>Function</b>	disease:A genetic variation in HNF1B is associated with susceptibility to hereditary prostate cancer type 11 (HPC11) [MIM:611955].,disease:Defects in HNF1B are a cause of Muellerian aplasia [MIM:158330]. In a Norwegian family with a novel syndrome of mild diabetes and severe non-diabetic renal disease, Muellerian aplasia expressed as vaginal aplasia and rudimentary uterus, were found in 2 females. These findings suggest that a broader spectrum of clinical symptoms may be associated with defects in HNF1B than previously recognized.,disease:Defects in HNF1B are the cause of maturity-onset diabetes of the young type 5 (MODY5) [MIM:604284]. MODY [MIM:606391] is a form of diabetes mellitus characterized by an autosomal dominant mode of inheritance, age of onset of 25 years or younger and a primary defect in insulin secretion.,disease:Defects in HNF1B are the cause of renal cysts and diabetes



## 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.[provided by RefSeq, Sep 2009],

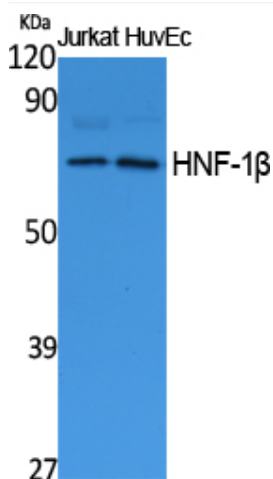
## matters needing attention

Avoid repeated freezing and thawing!

## Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## Products Images



Western Blot analysis of extracts from Jurkat cells, using HNF-1β Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000