

## **MVK Polyclonal Antibody**

YT2924 Catalog No:

Reactivity: Human; Monkey

**Applications:** WB;IHC;IF;ELISA

MVK **Target:** 

Fields: >>Terpenoid backbone biosynthesis;>>Metabolic pathways;>>Peroxisome

Gene Name: MVK

**Protein Name:** Mevalonate kinase

Q03426

Q9R008

**Human Gene Id:** 4598

**Human Swiss Prot** 

No:

**Mouse Swiss Prot** 

No:

Immunogen: The antiserum was produced against synthesized peptide derived from human

Mevalonate Kinase. AA range:151-200

MVK Polyclonal Antibody detects endogenous levels of MVK protein. **Specificity:** 

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution:** WB 1:500 - 1:2000, IHC 1:100 - 1:300, IF 1:200 - 1:1000, ELISA: 1:10000, Not

yet tested in other applications.

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

**Concentration:** 1 mg/ml

**Storage Stability:** -15°C to -25°C/1 year(Do not lower than -25°C)

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**Observed Band**: 42kD

**Cell Pathway:** Terpenoid backbone biosynthesis;

**Background:** This gene encodes the peroxisomal enzyme mevalonate kinase. Mevalonate is a

key intermediate, and mevalonate kinase a key early enzyme, in isoprenoid and sterol synthesis. Mevalonate kinase deficiency caused by mutation of this gene results in mevalonic aciduria, a disease characterized psychomotor retardation, failure to thrive, hepatosplenomegaly, anemia and recurrent febrile crises. Defects in this gene also cause hyperimmunoglobulinaemia D and periodic fever syndrome, a disorder characterized by recurrent episodes of fever associated with lymphadenopathy, arthralgia, gastrointestinal dismay and skin rash.

Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul

2014],

**Function :** catalytic activity:ATP + (R)-mevalonate = ADP +

(R)-5-phosphomevalonate., disease: Defects in MVK are the cause of

hyperimmunoglobulinemia D and periodic fever syndrome (HIDS) [MIM:260920]. HIDS is an autosomal recessive disease characterized by recurrent episodes of unexplained high fever associated with skin rash, diarrhea, adenopathy (swollen, tender lymph nodes), athralgias and/or arthritis. Concentration of IgD, and often IgA, are above normal., disease: Defects in MVK are the cause of mevalonic aciduria [MIM:610377]. It is an accumulation of mevalonic acid which causes a variety of symptoms such as psychomotor retardation, dysmorphic features, cataracts, hepatosplenomegaly, lymphadenopathy, anemia, hypotonia, myopathy,

and ataxia.,enzyme regulation:Farnesyl- and geranyl-pyrophosphates are

competitive inhibitors.,function:May be a regulatory site in cholesterol biosynthetic

pathway.,onl

Subcellular Location:

Cytoplasm . Peroxisome .

**Expression :** Brain, Hepatoma, Skin, Testis,

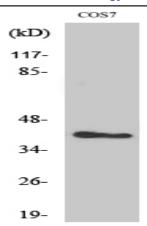
**Sort**: 10417

No4:

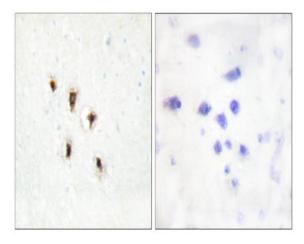
Host: Rabbit

Modifications: Unmodified

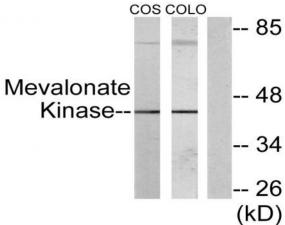
## **Products Images**



Western Blot analysis of various cells using MVK Polyclonal Antibody



Immunohistochemistry analysis of paraffin-embedded human brain tissue, using Mevalonate Kinase Antibody. The picture on the right is blocked with the synthesized peptide.



Western blot analysis of lysates from COS7 and COLO205 cells, using Mevalonate Kinase Antibody. The lane on the right is blocked with the synthesized peptide.