

## **Product Name: HoxD10 Rabbit Polyclonal Antibody**

Catalog #: APRab12177

For research use only.

#### **Summary**

**Description** Rabbit polyclonal Antibody

HostRabbitApplicationWB,ELISA

Reactivity Human, Mouse
Conjugation Unconjugated
Modification Unmodified

**Isotype** IgG

ClonalityPolyclonalFormLiquidConcentration1mg/ml

**Storage** Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.

Shipping Ice bags

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type **Buffer** 

preservative N.

**Purification** Affinity purification

### **Application**

**Dilution Ratio** WB 1:500-1:2000,ELISA 1:20000-1:40000

Molecular Weight 32kDa

# **Antigen Information**

**Alternative Names** 

Gene Name HOXD10

HOXD10; HOX4D; HOX4E; Homeobox protein Hox-D10; Homeobox protein Hox-4D;

Homeobox protein Hox-4E

 Gene ID
 3236.0

 SwissProt ID
 P28358

The antiserum was produced against synthesized peptide derived from human HOXD10. AA Immunogen

range:291-340

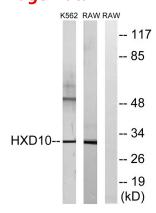
# **Background**



This gene is a member of the Abd-B homeobox family and encodes a protein with a homeobox DNA-binding domain. It is included in a cluster of homeobox D genes located on chromosome 2. The encoded nuclear protein functions as a sequence-specific transcription factor that is expressed in the developing limb buds and is involved in differentiation and limb development. Mutations in this gene have been associated with Wilm's tumor and congenital vertical talus (also known as "rocker-bottom foot" deformity or congenital convex pes valgus) and/or a foot deformity resembling that seen in Charcot-Marie-Tooth disease. [provided by RefSeq, Jul 2008],developmental stage:Expressed in the developing limb buds.,disease:Defects in HOXD10 are a cause of congenital vertical talus (CVT) [MIM:192950]; also known as "rocker-bottom foot" deformity or congenital convex pes valgus. CVT is a dislocation of the talonavicular joint, with rigid dorsal dislocation of the navicular over the neck of the talus. This condition is usually associated with multiple other congenital deformities and only rarely is an isolated deformity.,function:Sequence-specific transcription factor which is part of a developmental regulatory system that provides cells with specific positional identities on the anterior-posterior axis,,similarity:Belongs to the Abd-B homeobox family,,similarity:Contains 1 homeobox DNA-binding domain.,tissue specificity:Strongly expressed in the adult male and female urogenital tracts.,

#### **Research Area**

#### **Image Data**



Western blot analysis of lysates from K562 and RAW264.7 cells, using HOXD10 Antibody. The lane on the right is blocked with the synthesized peptide.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838