# Arylsulfatase E (L-13): sc-131421



The Power to Question

#### **BACKGROUND**

Sulfatases hydrolyze sulfate esters from sulfated steroids, carbohydrates, proteoglycans and glycolipids. They are involved in hormone biosynthesis, modulation of cell signaling and degradation of macromolecules. Arylsulfatase E, also known as CDPX, CDPX1 or CDPXR, is a 589 amino acid Golgi apparatus protein expressed in the pancreas, liver and kidney. Belonging to the sulfatase family, Arylsulfatase E may be essential for the correct composition of cartilage and bone matrix during development. Inhibited by warfarin, Arylsulfatase E activity occurs under optimal pH and temperature. Defects in the gene encoding Arylsulfatase E are the cause of chondrodysplasia punctata X-linked recessive type 1 (CDPX1), which is characterized by aberrant bone mineralization, severe underdevelopment of nasal cartilage and distal phalangeal hypoplasia. Arylsulfatase E has no activity toward steroid sulfates.

## **REFERENCES**

- Chen, N.T., et al. 1988. Potentiation of angiostatic steroids by a synthetic inhibitor of arylsulfatase. Lab. Invest. 59: 453-459.
- Parenti, G., et al. 1997. The sulfatase gene family. Curr. Opin. Genet. Dev. 7: 386-391.
- Dahl, H.H., et al. 1999. Late diagnosis of maternal PKU in a family segregating an arylsulfatase [corrected] E mutation causing symmetrical chondrodysplasia punctata. Mol. Genet. Metab. 68: 503-506.
- Seidel, J., et al. 2001. Brachytelephalangic dwarfism due to the loss of ARSE and SHOX genes resulting from an X;Y translocation. Clin. Genet. 59: 115-121.
- Brunetti-Pierri, N., et al. 2003. X-linked recessive chondrodysplasia punctata: spectrum of arylsulfatase E gene mutations and expanded clinical variability. Am. J. Med. Genet. A 117A: 164-168.
- Wolpoe, M.E., et al. 2004. Severe tracheobronchial stenosis in the Xlinked recessive form of chondrodysplasia punctata. Arch. Otolaryngol. Head Neck Surg. 130: 1423-1426.
- Chocholska, S., et al. 2006. Molecular cytogenetic analysis of a familial interstitial deletion Xp22.2-22.3 with a highly variable phenotype in female carriers. Am. J. Med. Genet. A 140: 604-610.
- Nino, M., et al. 2008. Clinical and molecular analysis of arylsulfatase E in patients with brachytelephalangic chondrodysplasia punctata. Am. J. Med. Genet. A 146A: 997-1008.
- Casarin, A., et al. 2009. X-linked brachytelephalangic chondrodysplasia punctata: a simple trait that is not so simple. Am. J. Med. Genet. A 149A: 2464-2468.

# CHROMOSOMAL LOCATION

Genetic locus: Arse (rat) mapping to 2q25.

## **SOURCE**

Arylsulfatase E (L-13) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of Arylsulfatase E of rat origin.

#### **PRODUCT**

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-131421 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## **APPLICATIONS**

Arylsulfatase E (L-13) is recommended for detection of Arylsulfatase E of rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); non cross-reactive with other Arylsulfatase family members.

Molecular Weight of Arylsulfatase E: 66 kDa.

## **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat lgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat lgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat lgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat lgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

# **STORAGE**

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

## **PROTOCOLS**

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

**Santa Cruz Biotechnology, Inc.** 1.800.457.3801 831.457.3801 **Fax** 831.457.3801 **Europe** +00800 4573 8000 49 6221 4503 0 **www.scbt.com**