# PDE6β (N-17): sc-30717



The Power to Question

#### **BACKGROUND**

Cyclic guanosine monophosphate (cGMP)-specific phosphodiesterase (PDE6) plays a crucial role in the phototransduction cascade in the vertebrate retina. The enzyme consists of an  $\alpha$  and a  $\beta$  subunit, with catalytic and cGMP binding activity, respectively, as well as two inhibitory  $\gamma$  subunits and a  $\delta$  subunit. PDE6 reduces intracellular cytoplasmic cGMP levels, specifically in photoreceptor cells. Mutations in the human PDE6A gene, which encodes the  $\alpha$  subunit, account for roughly 3-4% of the cases of recessive retinitis pigmentosa (RP) in North America.

# REFERENCES

- Mohamed, M.K., Taylor, R.E., Feinstein, D.S., Huang, X. and Pittler, S.J. 1998. Structure and upstream region characterization of the human gene encoding rod photoreceptor cGMP phosphodiesterase α-subunit. J. Mol. Neurosci. 10: 235-250.
- 2. Dryja, T.P., Rucinski, D.E., Chen, S.H. and Berson, E.L. 1999. Frequency of mutations in the gene encoding the  $\alpha$  subunit of rod cGMP-phosphodiesterase in autosomal recessive retinitis pigmentosa. Invest. Ophthalmol. Vis. Sci. 40: 1859-1865.
- Dekomien, G. and Epplen, J.T. 2000. Exclusion of the PDE6A gene for generalised progressive retinal atrophy in 11 breeds of dog. Anim. Genet. 31: 135-139.
- 4. Pittler, S.J., Zhang, Y., Chen, S., Mears, A.J., Zack, D.J., Ren, Z., Swain, P.K., Yao, S., Swaroop, A. and White, J.B. 2004. Functional analysis of the rod photoreceptor cGMP phosphodiesterase  $\alpha$ -subunit gene promoter: Nrl and Crx are required for full transcriptional activity. J. Biol. Chem. 279: 19800-19807.

## **CHROMOSOMAL LOCATION**

Genetic locus: PDE6B (human) mapping to 4p16.3; Pde6b (mouse) mapping to 5 F.

# **SOURCE**

PDE6 $\beta$  (N-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of PDE6 $\beta$  of human 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-515648 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

# **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.

#### **APPLICATIONS**

PDE6β (N-17) is recommended for detection of precursor and mature PDE6β of mouse, rat and human 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).

PDE6 $\beta$  (N-17) is also recommended for detection of precursor and mature PDE6 $\beta$  in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for PDE6 $\beta$  siRNA (h): sc-106850, PDE6 $\beta$  siRNA (m): sc-152131, PDE6 $\beta$  shRNA Plasmid (h): sc-106850-SH, PDE6 $\beta$  shRNA Plasmid (m): sc-152131-SH, PDE6 $\beta$  shRNA (h) Lentiviral Particles: sc-106850-V and PDE6 $\beta$  shRNA (m) Lentiviral Particles: sc-152131-V.

Molecular Weight of PDE6β: 98 kDa.

## **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-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 IgG-FITC: sc-2024 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

## **SELECT PRODUCT CITATIONS**

1. Ying, M., Chen, G., Qiu, Y., Shi, X., Zhang, C., Wang, Q., Yang, S., Lu, L., Yuan, Q., Xu, G., Jin, Z., Wu, Q. and Liu, X. 2015. Drug-inducible synergistic gene silencing with multiple small hairpin RNA molecules for gene function study in animal model. Transgenic Res. 24: 309-317.

# **PROTOCOLS**

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

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