## **NYX Rabbit Polyclonal Antibody**



## **CAB7830**

**Product Information** 

Size:

20uL, 50uL, 100uL, 200uL

**Observed MW:** 

52kDa

**Calculated MW:** 

52kDa

**Applications:** 

**WB IHC** 

Reactivity:

Human, Mouse

Antibody Information

Recommended dilutions:

WB 1:500 - 1:2000 IHC 1:50

- 1:100

Source:

Rabbit

Isotype:

IgG

**Purification:** 

Affinity purification

**Protein Background** 

The product of this gene belongs to the small leucine-rich proteoglycan (SLRP) family of proteins. Defects in this gene are the cause of congenital stationary night blindness type 1 (CSNB1), also called X-linked congenital stationary night blindness (XLCSNB). CSNB1 is a rare inherited retinal disorder characterized by impaired scotopic vision, myopia, hyperopia, nystagmus and reduced visual acuity. The role of other SLRP proteins suggests that mutations in this gene disrupt developing retinal interconnections involving the ON-bipolar cells, leading to the visual losses seen in patients with complete CSNB.

Immunogen information

Gene ID:

60506

Uniprot

Q9GZU5

**Synonyms:** 

NYX; CLRP; CSNB1; CSNB1A; CSNB4; NBM1

Immunogen:

Recombinant fusion protein containing a sequence corresponding

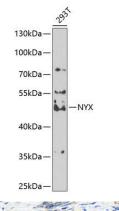
to amino acids 282-481 of human NYX (NP\_072089.1).

Storage:

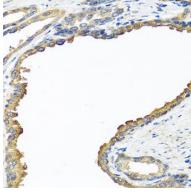
Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%

sodium azide, 50% glycerol, pH7.3.

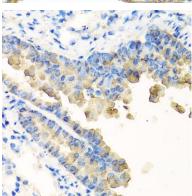
## **Product Images**



Western blot analysis of extracts of 293T cells, using NYX antibody (CAB7830) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Enhanced Kit (CABM00021). Exposure time: 90s.



Immunohistochemistry of paraffin-embedded human prostate using NYX antibody (CAB7830) at dilution of 1:100 (40x lens).



Immunohistochemistry of paraffin-embedded mouse lung using NYX antibody (CAB7830) at dilution of 1:100 (40x lens).