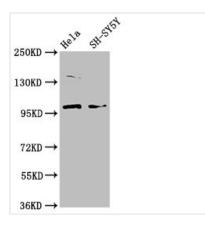
**Image** 



## PDE6C Antibody

<b>Product Code</b>	CSB-PA23139A0Rb
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P51160
Immunogen	Recombinant Human Cone cGMP-specific 3',5'-cyclic phosphodiesterase subunit alpha' protein (285-451AA)
Raised In	Rabbit
Species Reactivity	Human
<b>Tested Applications</b>	ELISA, WB, IF; Recommended dilution: WB:1:500-1:5000, IF:1:50-1:200
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300
213.4 <b>3</b> 0 <b>2</b> 41101	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	
	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified
Purification Method Isotype	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified  IgG
Purification Method Isotype Clonality	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4  >95%, Protein G purified  IgG  Polyclonal  Cone cGMP-specific 3',5'-cyclic phosphodiesterase subunit alpha' (EC 3.1.4.35)
Purification Method Isotype Clonality Alias	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4  >95%, Protein G purified  IgG  Polyclonal  Cone cGMP-specific 3',5'-cyclic phosphodiesterase subunit alpha' (EC 3.1.4.35) (cGMP phosphodiesterase 6C), PDE6C, PDEA2
Purification Method Isotype Clonality Alias Immunogen Species	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4  >95%, Protein G purified  IgG  Polyclonal  Cone cGMP-specific 3',5'-cyclic phosphodiesterase subunit alpha' (EC 3.1.4.35) (cGMP phosphodiesterase 6C), PDE6C, PDEA2  Homo sapiens (Human)



Western Blot

Positive WB detected in: Hela whole cell lysate,

SH-SY5Y whole cell lysate

All lanes: PDE6C antibody at 3.7µg/ml

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 100 kDa Observed band size: 100 kDa



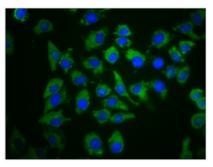
## **CUSABIO TECHNOLOGY LLC**











Immunofluorescence staining of A549 cells with CSB-PA23139A0Rb at 1:66, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).