HSD17B1 Antibody



PACO55550

Product Information

Size: **Protein Background:**

50ug Favors the reduction of estrogens and androgens. Also has 20-alpha-HSD activity. Uses

preferentially NADH.

Reactivity:

Gene ID: Human

HSD17B1 Source:

Uniprot Rabbit

P14061

Isotype:

ELISA, IHC, IF

Synonyms: lgG

Estradiol 17-beta-dehydrogenase 1 (EC 1.1.1.62) (17-beta-hydroxysteroid **Applications:**

dehydrogenase type 1) (17-beta-HSD 1) (20 alpha-hydroxysteroid dehydrogenase) (20alpha-HSD) (E2DH) (Placental 17-beta-hydroxysteroid dehydrogenase) (Short chain dehydrogenase/reductase family 28C member 1), HSD17B1, E17KSR EDH17B1

EDH17B2 EDHB17 SDR28C1 **Recommended dilutions:**

ELISA:1:2000-1:10000, IHC:1:20-1:200,

IF:1:20-1:200

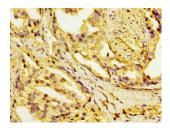
Immunogen:

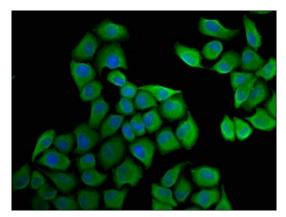
Recombinant Human Estradiol 17-beta-dehydrogenase 1 protein (268-328AA).

Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

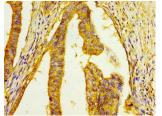
Product Images





IHC image of PACO55550 diluted at 1:100 and staining in paraffinembedded human prostate cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunofluorescence staining of Hela cells with PACO55550 at 1:33, 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).



IHC image of PACO55550 diluted at 1:100 and staining in paraffinembedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.