ELF3 Antibody



PACO63295

Isotype:

lgG

Product Information

Size: Protein Background:

50ul Transcriptional activator that binds and transactivates ETS sequences containing the

Reactivity:consensus nucleotide core sequence GGA[AT]. Acts synergistically with POU2F3 to transactivate the SPRR2A promoter and with RUNX1 to transactivate the ANGPT1

Human promoter. Also transactivates collagenase, CCL20, CLND7, FLG, KRT8, NOS2, PTGS2,

SPRR2B, TGFBR2 and TGM3 promoters. Represses KRT4 promoter activity. Involved in **Source:** mediating vascular inflammation. May play an important role in epithelial cell

differentiation and tumorigenesis. May be a critical downstream effector of the ERBB2

Rabbit signaling pathway. May be associated with mammary gland development and

involution. Plays an important role in the regulation of transcription with TATA-less

promoters in preimplantation embryos, which is essential in preimplantation

development.

Applications: Gene ID:

ELISA, IHC, IF ELF3

Recommended dilutions: Uniprot

ELISA:1:2000-1:10000, IHC:1:500-1:1000, P78545

IF:1:50-1:200

Synonyms:

ETS-related transcription factor Elf-3 (E74-like factor 3) (Epithelial-restricted with serine box) (Epithelium-restricted Ets protein ESX) (Epithelium-specific Ets transcription factor 1) (ESE-1), ELF3, ERT ESX JEN

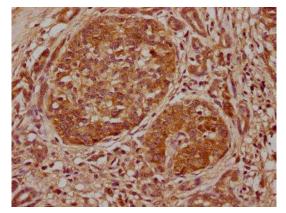
Immunogen:

Recombinant Human ETS-related transcription factor Elf-3 protein (147-371AA).

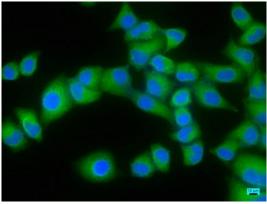
Storage:

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

Product Images



IHC image of PACO63295 diluted at 1:500 and staining in paraffinembedded human pancreatic 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 PACO63295 at 1:166, 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).