TXN Antibody



PACO31564

Reactivity:

Human

Product Information

Size: Protein Background:

50ug Participates in various redox reactions through the reversible oxidation of its active center dithiol to a disulfide and catalyzes dithiol-disulfide exchange reactions. Plays a

role in the reversible S-nitrosylation of cysteine residues in target proteins, and thereby contributes to the response to intracellular nitric oxide. Nitrosylates the active site Cys

of CASP3 in response to nitric oxide (NO), and thereby inhibits caspase-3 activity.

Source: Induces the FOS/JUN AP-1 DNA binding activity in ionizing radiation (IR) cells through

Rabbit its oxidation/reduction status and stimulates AP-1 transcriptional activity.

Gene ID: Isotype:

TXN IgG

Applications:
P10599

ELISA, WB, IHC, IF

Synonyms: Recommended dilutions:

Thioredoxin (Trx) (ATL-derived factor) (ADF) (Surface-associated sulphydryl protein) ELISA:1:2000-1:10000, WB:1:500-1:5000, IF:1:50-1:200 (SASP), TXN, TRDX TRX TRX1

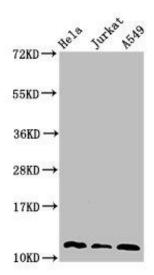
Immunogen:

Recombinant Human Thioredoxin protein (2-105AA).

Storage:

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

Product Images



Western Blot

Positive WB detected in: Hela whole cell lysate, Jurkat whole cell lysate,

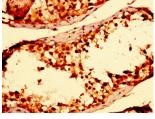
A549 whole cell lysate

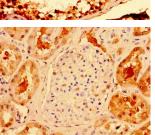
All lanes: TXN antibody at 3µg/ml

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 12, 10 kDa Observed band size: 12 kDa





IHC image of PACO31564 diluted at 1:300 and staining in paraffinembedded human testis tissue 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.

IHC image of PACO31564 diluted at 1:300 and staining in paraffinembedded human kidney tissue 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.