





TXN Antibody

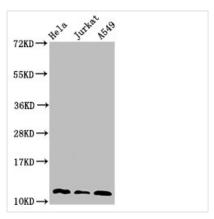
Product Code	CSB-PA02814A0Rb
Abbreviation	Thioredoxin` protein
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P10599
Immunogen	Recombinant Human Thioredoxin protein (2-105AA)
Raised In	Rabbit
Species Reactivity	Human
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:200-1:500, IF:1:50-1:200
Relevance	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. Induces the FOS/JUN AP-1 DNA binding activity in ionizing radiation (IR) cells through its oxidation/reduction status and
	stimulates AP-1 transcriptional activity.
Form	
Form Conjugate	stimulates AP-1 transcriptional activity.
	stimulates AP-1 transcriptional activity. Liquid
Conjugate	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300
Conjugate Storage Buffer	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Conjugate Storage Buffer Purification Method	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified
Conjugate Storage Buffer Purification Method Isotype	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG
Conjugate Storage Buffer Purification Method Isotype Clonality	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG Polyclonal Thioredoxin (Trx) (ATL-derived factor) (ADF) (Surface-associated sulphydryl
Conjugate Storage Buffer Purification Method Isotype Clonality Alias	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG Polyclonal Thioredoxin (Trx) (ATL-derived factor) (ADF) (Surface-associated sulphydryl protein) (SASP), TXN, TRDX TRX TRX1
Conjugate Storage Buffer Purification Method Isotype Clonality Alias Species	stimulates AP-1 transcriptional activity. Liquid Non-conjugated Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG Polyclonal Thioredoxin (Trx) (ATL-derived factor) (ADF) (Surface-associated sulphydryl protein) (SASP), TXN, TRDX TRX TRX1 Human

CUSABIO TECHNOLOGY LLC









Western Blot

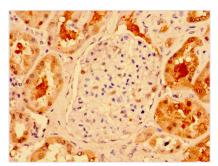
Positive WB detected in: Hela whole cell lysate, Jurkat whole cell lysate, A549 whole cell lysate

All lanes: TXN antibody at 3µg/ml

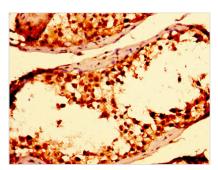
Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

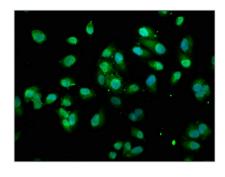
Predicted band size: 12, 10 kDa Observed band size: 12 kDa



IHC image of CSB-PA02814A0Rb diluted at 1:300 and staining in paraffin-embedded 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.



IHC image of CSB-PA02814A0Rb diluted at 1:300 and staining in paraffin-embedded 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.



Immunofluorescence staining of Hela cells with CSB-PA02814A0Rb at 1:100, 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).