
Product Name: KD-Validated DAXX Recombinant Rabbit Monoclonal Antibody**Catalog #: KVA00214**

For research use only.

Summary

Description	KO&KD-Validated antibody
Host	Rabbit
Application	WB,FCM,ICC
Reactivity	Human,Mouse,Rat
Conjugation	Unconjugated
Modification	Unmodified
Isotype	Rabbit IgG
Clonality	Rabbit mAb
Form	Liquid
Storage	Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.
Shipping	Ice bags
Buffer	Supplied in PBS (pH 7.4) containing 50% glycerol, and 0.02% sodium azide.
Purification	Affinity purification

Application

Dilution Ratio	WB 1:1,000-1:5,000; FC 1:200-1:2,000; ICC 1:100-1:1,000
Molecular Weight	Calculated MW: 81.4kDa

Antigen Information

Gene Name	DAXX DAXX; Death Domain Associated Protein; DAP6; Fas Death Domain-Associated Protein;
Alternative Names	Death Domain-Associated Protein 6; Death-Associated Protein 6; ETS1-Associated Protein 1; BING2; EAP1; CENP-C Binding Protein; Fas-Binding Protein; SMIM40; HDaxx; Daxx
Gene ID	1616.0
SwissProt ID	Q9UER7
Immunogen	A synthesized peptide derived from human DAXX

Background

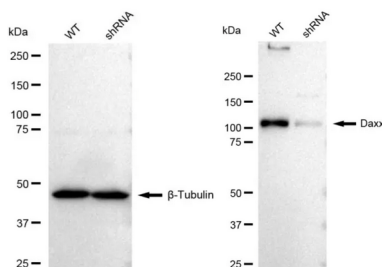
This gene encodes a multifunctional protein that resides in multiple locations in the nucleus and in the cytoplasm. It interacts with a wide variety of proteins, such as apoptosis antigen Fas, centromere protein C, and transcription factor erythroblastosis

virus E26 oncogene homolog 1. In the nucleus, the encoded protein functions as a potent transcription repressor that binds to sumoylated transcription factors. Its repression can be relieved by the sequestration of this protein into promyelocytic leukemia nuclear bodies or nucleoli. This protein also associates with centromeres in G2 phase. In the cytoplasm, the encoded protein may function to regulate apoptosis. The subcellular localization and function of this protein are modulated by post-translational modifications, including sumoylation, phosphorylation and polyubiquitination. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov 2008]

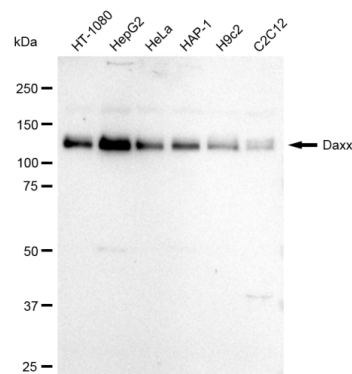
Research Area

Cell Biology, Microbiology, Epigenetics and Nuclear Signaling

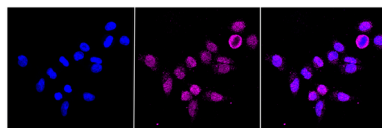
Image Data



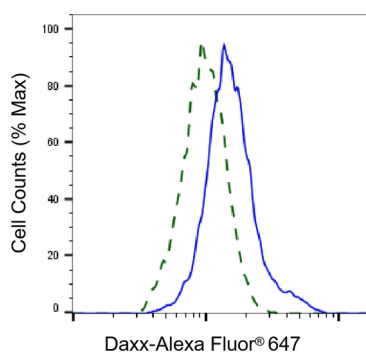
Western blotting analysis using Daxx antibody (KVA00214). Daxx expression in wild type (WT) and Daxx shRNA knockdown (KD) HeLa cells with 30 µg of total cell lysates. β-Tubulin serves as a loading control. The blot was incubated with Daxx antibody (KVA00214, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (APS0635, 1:10,000) respectively.



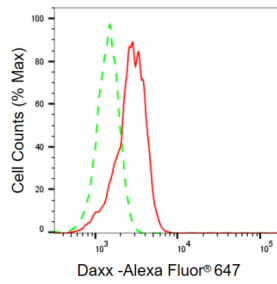
Western blotting analysis using Daxx antibody (KVA00214). Total cell lysates (30 µg) from various cell lines were loaded and separated by SDS-PAGE. The blot was incubated with Daxx antibody (KVA00214, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (APS0635, 1:10,000) respectively.



Immunocytochemical staining of HepG2 cells with Daxx antibody (KVA00214, 1:1,000). Nuclei were stained blue with DAPI; Daxx was stained magenta with Alexa Fluor® 647. Images were taken using Leica stellaris 5. Protein abundance based on laser intensity and smart gain: Medium. Scale bar: 20 µm.



Validation of Daxx knockdown using flow cytometry. Wild-type (WT, Blue) and knockdown (KD, Green) HeLa cells were stained with Daxx antibody (KVA00214, 1:2,000) and analyzed using BD flow cytometer.



Flow cytometric analysis of Daxx expression in HepG2 cells using Daxx antibody (KVA00214, 1:2,000). Green, isotype control; red, Daxx.