[Primary Antibody]

phospho-DAXX (Ser671) Rabbit pAb



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– DATASHEET ––––––		400-901-9800
Host: Rabbit	lsotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)
GenelD: 1616	SWISS: Q9UER7	IF (1:100-500)
Target: phospho-DAXX (Ser671)		Reactivity: Human (predicted: Cow, Horse)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human DAXX around the phosphorylation site of Ser671: PP(p-S)PL.		
Purification: affinity purified by	Protein A	
Concentration: 1mg/ml		Predicted MW.: ^{81 kDa}
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Location: ^{Cytoplasm} ,Nucleus
regulating the RING Under non-stress of deubiquitinating U enhances the intri- thereby promoting proteasomal degra MDM2 and USP7 is autoubiquitination leads to TP53 stab JNK pathway and a from TNFRSF6 and prevent interaction mediated apoptos and TNFRSF6-med regulate transcript with PML and may thereby. Down-reg to act as a transcri through direct pro activity. Its transcr	protein in a MDM2-DAXX-USP7 complex by G-finger E3 ligase MDM2 ubiquitination activity. condition, in association with the ISP7, prevents MDM2 self-ubiquitination and nsic E3 ligase activity of MDM2 towards TP53, g TP53 ubiquitination and subsequent adation. Upon DNA damage, its association with disrupted, resulting in increased MDM2 n and consequently, MDM2 degradation, which ilization. Proposed to mediate activation of the apoptosis via MAP3K5 in response to signaling TGFBR2. Interaction with HSPB1/HSP27 may n with TNFRSF6 and MAP3K5 and block DAXX- is. In contrast, in lymphoid cells JNC activation iated apoptosis may not involve DAXX. Seems to ion in PML/POD/ND10 nuclear bodies together influence TNFRSF6-dependent apoptosis ulates basal and activated transcription. Seems ptional corepressor and inhibits PAX3 and ETS1 tein-protein interaction. Modulates PAX5 iption repressor activity is modulated by nuclear compartments like the nucleolus or uclear bodies through interactions with MCSR1 rely.	

- VALIDATION IMAGES -

kDa	5H-262 12 40 5 4 10 10 10 10
$^{310}_{245} =$	
180	
140 —	phospho-DAXX
100	
60 —	
45	
35 —	the set off as the set

Sample: Lane 1: Human SH-SY5Y cell lysates Lane 2: Human K562 cell lysates Lane 3: Human U-2 OS cell lysates Lane 4: Human MOLT4 cell lysates Lane 5: Human Jurkat cell lysates Lane 6: Human HeLa cell lysates Primary: Anti-phospho-DAXX (Ser671) (bs-5293R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 81 kDa Observed band size: 120 kDa



Paraformaldehyde-fixed, paraffin embedded (human breast carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (phospho-DAXX (Ser671)) Polyclonal Antibody, Unconjugated (bs-5293R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.