## bs-6528R

DATACHEET

## [ Primary Antibody ]

## **EZH1 Rabbit pAb**

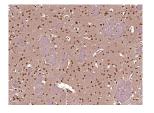


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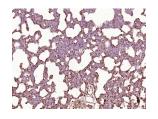
- DATASHEE	: I ————		
Host:	Rabbit	Isotype: IgG	Applications: IHC-P
Clonality:	Polyclonal		IHC-F IF (1:1)
GenelD:	2145	SWISS: Q92800	Flow-
Target:	EZH1		ICC/IF
Immunogen:	<b>Dgen:</b> KLH conjugated synthetic peptide derived from human EZH1: 471-510/747.		Reactivity: Human (predic
Purification:	affinity purified by Prote	Cow, C Guinea	
Concentration:	1mg/ml		Predicted
<b>Storage:</b> 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.			<b>MW.:</b> <sup>85 KDa</sup>
			Subcellular Location: Nucleu
Background:	EZH1 complex, which m transcriptional repression mono-, di- and trimethy H3K27me1, H3K27me2 a embryonic stem cell der is involved in safeguardi Compared to EZH1-conter embryonic stem cells, h plays a less critical role i embryonic stem cell ide Sequence similarities	protein. Catalytic subunit of the PRC2/EEE nethylates 'Lys-27' of histone H3, leading t on of the affected target gene. Able to vlate 'Lys-27' of histone H3 to form and H3K27me3, respectively. Required fo rivation and self-renewal, suggesting that ing embryonic stem cell identity. taining complexes, it is less abundant in as weak methyltransferase activity and in forming H3K27me3, which is required f entity and proper differentiation.	o r it

Belongs to the histone-lysine methyltransferase family. EZ subfamily. Contains 1 SET domain.

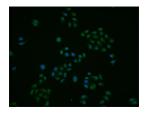
## - VALIDATION IMAGES



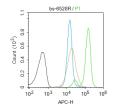
Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (EZH1) Polyclonal Antibody, Unconjugated (bs-6528R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Rat lung); 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; Antibody incubation with (EZH1) Polyclonal Antibody, Unconjugated (bs-6528R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (EZH1) polyclonal Antibody, Unconjugated (bs-6528R) 1:25, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control:Molt4. Primary Antibody (green

400-901-9800

**P** (1:100-500) **F** (1:100-500) 100-500) -Cyt (2ug/Test) **F** (1:25)

an, Mouse, Rat licted: Rabbit, Pig, Chicken, Dog, eaPig)

eus

line): Rabbit Anti-EZH1 antibody (bs-6528R) Dilution: 2μg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-APC Dilution: 1μg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at-20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.