bs-6592R

[Primary Antibody]

HSD17B6 Rabbit pAb



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– DATASHEET –		400-901-9800	
Host: Rabbit	lsotype: lgG	Applications: Flow-Cyt (1ug/Test)	
Clonality: Polyclonal		Reactivity: Human (predicted: Mouse,	
GeneID: 8630	SWISS: 014756	Rat, Pig, Horse)	
Target: HSD17B6			
Immunogen: KLH conjugated synthetic peptide derived from human HSD17B6: 61-160/317.		Predicted MW.: ^{33 kDa}	
Purification: affinity purified by Protein A			
Concentration: 1mg/ml		Subcellular Location: Cell membrane, Cytoplasm	
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.			
Background: The protein encoded by this gene has both oxidoreductase and epimerase activities and is involved in androgen catabolism. The oxidoreductase activity can convert 3 alpha-adiol to dihydrotestosterone, while the epimerase activity can convert androsterone to epi-androsterone. Both reactions use NAD+ as the		e	

– VALIDATION IMAGES



Molt-4 cells were fixed with 4% PFA for 10min at room temperature,permeabilized with 0.1% PBST for 20 min at room temperature, and incubated in 5% BSA blocking buffer for 30 min at room temperature. Cells were then stained with HSD17B6 Antibody(bs-6592R)at 1:100 dilution in blocking buffer and incubated for 30 min at room temperature, washed twice with 2%BSA in PBS, followed by secondary antibody incubation for 40 min at room temperature. Acquisitions of 20,000 events were performed. Cells stained with primary antibody (green), and isotype control (orange).



preferred cofactor. This gene is a member of the retinol dehydrogenase family. [provided by RefSeq, Aug 2013]

Blank control (Black line): Molt4 (Black). Primary Antibody (green line):Rabbit Anti-HSD17B6 antibody (bs-6592R) Dilution:3µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF647 Dilution: 3µg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with PBST for 20 min at room temperature. 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.