Product datasheet Anti-HO-1/HMOX1 Antibody Catalog Number: A00253-2

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BOSTER BIOLOGICAL TECHNOLOGY

Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

Web: www.boster.com Phone: 027-67845390/1/2 Email: boster@boster.com

Product Name	Anti-HO-1/HMOX1 Antibody	
Gene Name	HMOX1	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E.coli-derived human Heme Oxygenase 1/HMOX1 recombinant protein (Position: E19-R269).	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	33 kDa	
Dilution Ratios	Western blot (WB): Flow Cytometry (Fixed):	1:500-2000 1:50-200

Storage

12 months from date of receipt, -20° C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Enzyme linked immunosorbent assay (ELISA):1:100-1000

Background Information

HMOX1 (heme oxygenase (decycling) 1), also known as HO-1, is a human gene that encodes for the enzyme heme oxygenase 1. It is an essential enzyme in heme catabolism, it cleaves heme to form biliverdin. HMOX1 belongs to the heme oxygenase family. The HMOX1 gene is located on the long (q) arm of chromosome 22 at position 12.3, from base pair 34,101,636 to base pair 34,114,748. HMOX1, an essential enzyme in heme catabolism, cleaves heme to form biliverdin, which is subsequently converted to bilirubin by biliverdin reductase, and carbon monoxide, a putative neurotransmitter. HMOX1 activity is induced by its substrate heme and by various nonheme substances.

Reference

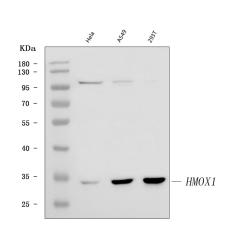
Anti-HO-1/HMOX1 Antibody被引用在2文献中。



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Selected Validation Data



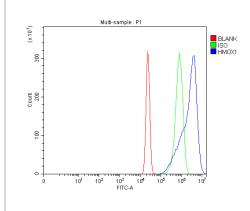
Western blot analysis of HO-1/HMOX1 using anti-HO-1/HMOX1 antibody (A00253-2). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Hela whole cell lysates,

Lane 2: human A549 whole cell lysates,

Lane 3: human 293T whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-HO-1/HMOX1 antigen affinity purified polyclonal antibody (A00253-2) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for HO-1/HMOX1 at approximately 33 kDa. The expected band size for HO-1/HMOX1 is at 33 kDa.



Flow Cytometry analysis of U2OS cells using anti-HO-1/HMOX1 antibody (A00253-2).

Overlay histogram showing U2OS cells stained with A00253-2 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-HO-1/HMOX1 Antibody (A00253-2) at 1:100 dilution for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.