

Human/Mouse HO-1/HMOX1/HSP32 Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF3776

DESCRIPTION		
Species Reactivity	Human/Mouse	
Specificity	Detects human and mouse HO-1/HMOX1/HSP32 in Western blots. In Western blots, less than 1% cross-reactivity with recombinant huma HO-2 is observed.	
Source	Polyclonal Goat IgG	
Purification	Antigen Affinity-purified	
Immunogen	E. coli-derived recombinant human HO-1/HMOX1/HSP32 Met1-Thr261 Accession # P09601	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.	

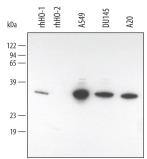
APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample	
Western Blot	0.5 μg/mL	See Below	
Simple Western	25 μg/mL	See Below	
Knockout Validated	HO-1/HMOX1/HSP32 is specifically detected in HeLa human cervical epithelial carcinoma parental cell line but is not detectable in HO-1/HMOX1/HSP32 knockout HeLa cell line.		

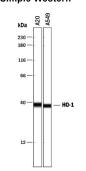
DATA

Western Blot



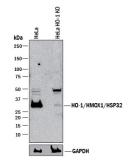
Detection of Human/Mouse HO-1/HMOX1/ HSP32 by Western Blot. Western blot shows lysates of A549 human lung carcinoma cell line, DU145 human prostate carcinoma cell line, and A20 mouse B cell lymphoma cell line. PVDF membrane was probed with 0.5 µg/mL Goat Anti-Human/Mouse HO-1/HMOX1/HSP32 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF3776) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF017). For additional reference, recombinant human HO-1 and HO-2 (5 ng/lane) were included. A specific band for HO-1/HMOX1/HSP32 was detected at approximately 32 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 2.

Simple Western



Detection of Human and Mouse HO-1/HMOX1/HSP32 by Simple WesternTM. Simple Western lane view shows lysates of A20 mouse B cell lymphoma cell line and A549 human lung carcinoma cell line, loaded at 0.2 mg/mL. A specific band was detected for HO-1/HMOX1/HSP32 at approximately 37 kDa (as indicated) using 25 µg/mL of Goat Anti-Human/Mouse HO-1/HMOX1/HSP32 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF3776) followed by 1:50 dilution of HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF109). This experiment was conducted under reducing conditions and using the 12-230 kDa separation system.

Knockout Validated



Western Blot Shows Human HO-1/HMOX1/HSP32 Specificity by Using Knockout Cell Line. Western blot shows lysates of HeLa human cervical epithelial carcinoma parental cell line and HO-1/HMOX1/HSP32 knockout HeLa cell line (KO). PVDF membrane was probed with 0.5 µg/mL of Goat Anti-Human/Mouse HO-1/HMOX1/HSP32 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF3776) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF017). A specific band was detected for HO-. 1/HMOX1/HSP32 at approximately 32 kDa (as indicated) in the parental HeLa cell line, but is not detectable in knockout HeLa cell line. GAPDH (Catalog # AF5718) is shown as a loading control. This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

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Reconstitution
Shipping
Stability & Storage

BACKGROUND

Heme Oxygenase 1 (HO-1), also known as HMOX1 and Heat Shock Protein 32 (HSP32), is a 32 kDa microsomal enzyme required for the metabolism of heme to biliverdin. Heme oxygenase occurs as 2 isozymes, an inducible heme oxygenase-1 (HO-1/HMOX1) and a constitutive heme oxygenase-2 (HO-2/HMOX2). HO-1 expression is induced by heme and other non-heme compounds. Human HO-1 shares 82% amino acid sequence identity with mouse HO-1.

