

RNF20 Polyclonal Antibody

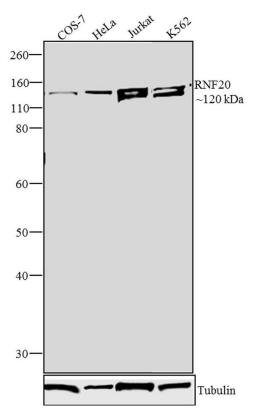
Product Details		
Size	100 μg	
Species Reactivity	Human, Non-human primate	
Host/Isotype	Rabbit / IgG	
Class	Polyclonal	
Туре	Antibody	
Conjugate	Unconjugated	
Immunogen	Recombinant protein corresponding to Human RNF20 (aa 96-433)	
Form	Liquid	
Concentration	0.5 mg/mL	
Purification	Affinity chromatography	
Storage buffer	PBS, pH 7.2, with 30% glycerol, 0.1% BSA	
Contains	0.09% sodium azide	
Storage conditions	Store at 4°C short term. For long term storage, store at -20°C, avoiding freeze/thaw cycles.	
RRID	AB_2607444	

Applications	Tested Dilution	Publications
Western Blot (WB)	0.5-1 μg/mL	-
Immunohistochemistry (Paraffin) (IHC (P))	1:20	-
Immunocytochemistry (ICC/IF)	1 μg/mL	-

Product Specific Information

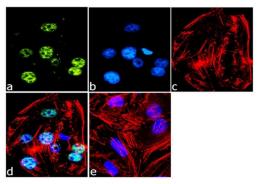
These Polyclonal antibodies are of rabbit origin developed by immunizing animals with proteins or peptides. The polyclonal antibody is purified by affinity purification from the rabbit sera generated after immunizing the rabbits with a specific type of protein or peptide. The purified antibody is tested for its functionality in various relevant research applications. The antibody is developed for Research Use Only and is non-hazardous or non-infectious in nature.

Product Images For RNF20 Polyclonal Antibody



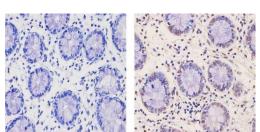
RNF20 Antibody (720146) in WB

Western blot analysis was performed on whole cell extracts (30 µg lysate) of COS-7 (Lane 1), HeLa (Lane 2), Jurkat (Lane3) and K-562 (Lane 4). The blots were probed with Anti- RNF20 Rabbit Polyclonal Antibody (Product # 720146, 0.5-1 µg/mL) and detected by chemiluminescence using Goat anti-Rabbit IgG (Heavy Chain) Superclonal Secondary Antibody, HRP conjugate (Product # A27036, 0.4 µg/mL, 1:2500 dilution). A 120 kDa band and an additional isoform at 72 kDa corresponding to RNF-20 was observed in cell lines tested. Known quantity of protein samples were electrophoresed using Novex® NuPAGE® 4-12% Bis-Tris gel (Product # NP0321BOX), XCell SureLock Electrophoresis System (Product # El0002) and Novex® Sharp Pre-Stained Protein Standard (Product # LC5800). Resolved proteins were then transferred onto a nitrocellulose membrane with iBlot® Dry Blotting System (Product # IB21001). The membrane was probed with the relevant primary and secondary Antibody following blocking with 5% skimmed milk. Chemiluminescent detection was performed using Pierce™ ECL Western Blotting Substrate (Product # 32106).



RNF20 Antibody (720146) in ICC/IF

Immunofluorescence was performed on fixed and permeabilized HeLa cells for detection of RNF20 using Anti-RNF20 Rabbit Polyclonal Antibody (Product #720146, 1 µg/mL), alpha-Tubulin was detected using Anti-alpha Tubulin Monoclonal Antibody (Product #32-2500, 1 µg/mL) and labeled with Goat anti-Rabbit IgG (Heavy Chain) Superclonal Secondary Antibody, Alexa Fluor® 488 conjugate (Product #A27034, 1:2000), Goat anti-Mouse IgG Secondary Antibody, Alexa Fluor®594conjugate (Product #A-11032, 1:400) respectively. Panel a) shows representative cells that were stained for detection and localization of RNF20 protein (green), Panel b) is stained for nuclei (blue) using SlowFade® Gold Antifade Mountant with DAPI (Product #S36938,). Panel c) represents cytoskeletal alpha-tubulin staining (red). Panel d) is a composite image of Panels a, b and c clearly demonstrating Nuclear localization of RNF-20. Panel e) represents control cells with no primary Antibody to assess background.



RNF20 Antibody (720146) in IHC (P)

Immunohistochemistry analysis of RNF20 showing staining in nucleus of paraffinembedded human colon tissue (right) compared to a negative control without primary antibody (left). To expose target proteins, antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature, washed with ddH2O and PBS, and then probed with a RNF20 Rabbit Polyclonal Antibody (Product # 720146) diluted in 3% BSA-PBS at a dilution of 1:20 for 1 hour at 37°C in a humidified chamber. Tissues were washed extensively in PBST and detection was performed using an HRP-conjugated secondary antibody followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.

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