

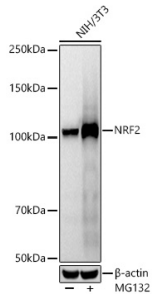
Anti-Nrf2 Antibody [ARC50393] (A308758)

Specifications:

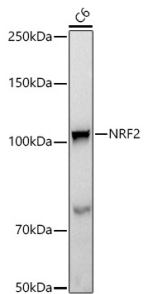
Name:	Anti-Nrf2 Antibody [ARC50393]
Description:	Rabbit monoclonal [ARC50393] antibody to Nrf2.
Applications:	WB
Recommended Dilutions:	WB: 1:500-1:1,000
Reactivity:	Human, Mouse, Rat
Immunogen:	Recombinant fusion protein containing a sequence corresponding to amino acids 243-400 of human NRF2 (NP_006155.2).
Sequence:	SPHFLNAFEDSFSSILSTEDPNQLTVNSLNSDATVNTDFGDEFYSAFIAEPSISNSMP SPATLSHSLSELLNGPIDVSDLKAFNQNHPESTAEFNDSDSGISLNTSPSVASPE HSVESSYGD TLLGLSDSEVEELDSAPGSVKQNGPKTPVHSS
Host:	Rabbit
Clonality:	Monoclonal
Clone ID:	ARC50393
Isotype:	IgG
Conjugate:	Unconjugated
Purification:	Affinity purification.
Molecular Weight:	97 - 100 kDa
Product Form:	Liquid
Formulation:	Supplied in Phosphate Buffered Saline, pH 7.3, with 50% Glycerol, 0.05% BSA, and 0.05% Proclin 300.
Storage:	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
Disclaimer:	This product is for research use only. It is not intended for diagnostic or therapeutic use.

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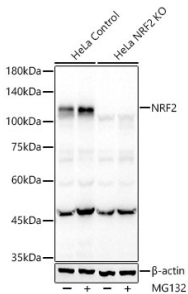
Images:



Western blot analysis of extracts of NIH/3T3 cells, using Anti-Nrf2 Antibody [ARC50393] (A308758) at 1:1,000 dilution. NIH/3T3 cells were treated by MG132(50 μ M) at 37°C for 90 minutes. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25 μ g per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 30s.



Western blot analysis of extracts of C6 cells, using Anti-Nrf2 Antibody [ARC50393] (A308758) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25 μ g per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 30s.



Western blot analysis of extracts from wild type(WT) and NRF2 knockout (KO) HeLa cells, using Anti-Nrf2 Antibody [ARC50393] (A308758) at 1:1,000 dilution. HeLa cells were treated by MG132(50 μ M) at 37°C for 90 minutes. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25 μ g per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 30s.