

## Basic Information

<b>Product Name</b>	Anti-KRAS/HRAS/NRAS Antibody (Clone#EBE-14)		
<b>Gene Name</b>	KRAS/HRAS/NRAS		
<b>Source</b>	Rabbit		
<b>Clonality</b>	Monoclonal		
<b>Isotype</b>	IgG		
<b>Species Reactivity</b>	human, mouse, rat		
<b>Tested Application</b>	WB, ICC/IF, IP, FCM		
<b>Contents</b>	500 ug/ml; Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide, 0.4-0.5 mg/ml BSA and 50% glycerol.		
<b>Immunogen</b>	A synthesized peptide derived from human KRAS+HRAS+NRAS		
<b>Concentration</b>	500 ug/ml		
<b>Purification</b>	Affinity-chromatography		
<b>Observed MW</b>	21 kDa		
<b>Dilution Ratios</b>	Western blot (WB):	1:500-2000	
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-200	
	ImmunoPrecipitation (IP):	1:20	
	Flow Cytometry (FCM):	1:20	

## Storage

12 months from date of receipt, -20°C as supplied.

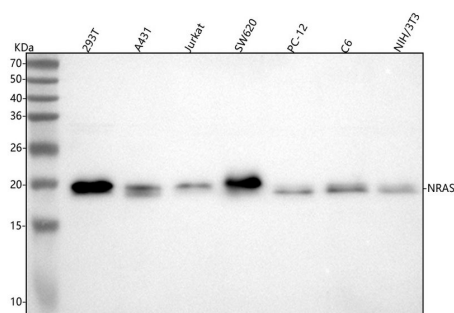
## Background Information

This gene, a Kirsten ras oncogene homolog from the mammalian ras gene family, encodes a protein that is a member of the small GTPase superfamily. A single amino acid substitution is responsible for an activating mutation. The transforming protein that results is implicated in various malignancies, including lung adenocarcinoma, mucinous adenoma, ductal carcinoma of the pancreas and colorectal carcinoma. Alternative splicing leads to variants encoding two isoforms that differ in the C-terminal region. [provided by RefSeq, Jul 2008]

## Reference

Anti-KRAS/HRAS/NRAS Antibody (Clone#EBE-14)被引用在6文献中。

## Selected Validation Data



Western blot analysis of anti-KRAS/HRAS/NRAS antibody (BM4388).

The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human 293T whole cell lysates,

Lane 2: human A431 whole cell lysates,

Lane 3: human Jurkat whole cell lysates,

Lane 4: human SW620 whole cell lysates,

Lane 5: rat PC-12 whole cell lysates,

Lane 6: rat C6 whole cell lysates,

Lane 7: mouse NIH/3T3 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-KRAS/HRAS/NRAS antigen affinity purified monoclonal antibody (BM4388) 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 KRAS/HRAS/NRAS at approximately 21 kDa. The expected band size for KRAS/HRAS/NRAS is at 21 kDa.