

Anti-Cellular Apoptosis Susceptibility (CSE1L) Mouse Monoclonal Antibody [Clone ID: OTI4G1]

Catalog Number: M04496

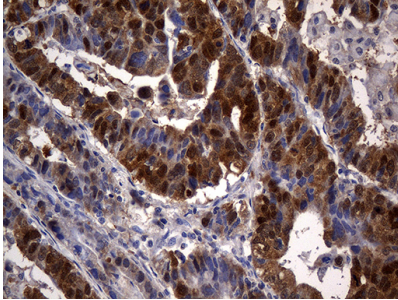
Overview

Product Name	Anti-Cellular Apoptosis Susceptibility (CSE1L) Mouse Monoclonal Antibody [Clone ID: OTI4G1]
Reactive Species	Human, Mouse, Rat
Description	Boster Bio CSE1L mouse monoclonal antibody, clone OTI4G1. Catalog# M04496. Tested in IHC, WB. This antibody reacts with Human, Mouse, Rat.
Conjugate	Unconjugated
Application	IHC, WB
Clonality	Monoclonal OTI4G1
Formulation	PBS (pH 7.3) containing 1% stabilizing protein, 50% glycerol and 0.02% sodium azide. This antibody is supplied in a stabilized formulation. Compatibility with conjugation reactions depends on the chemistry of the conjugation method used. For conjugation methods that are not compatible with the stabilizing components present in this formulation, a carrier-free antibody format is required.
Storage Instructions	Store at -20°C for one year. For short term storage and frequent use, store at 4°C for up to one month. Avoid repeated freeze-thaw cycles.
Host	Mouse
Uniprot ID	P55060

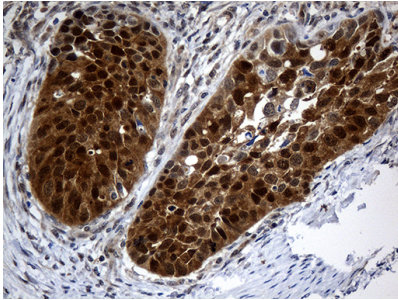
Technical Details

Immunogen	Human recombinant protein fragment corresponding to amino acids 1-300 of human CSE1L (NP_001307) produced in E.coli.
Isotype	IgG1
Concentration	1 mg/ml
Purification	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Suggested Dilutions	WB 1:500 IHC 1:2000

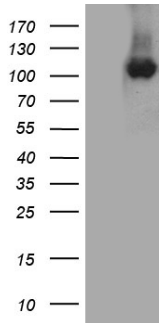
Anti-Cellular Apoptosis Susceptibility (CSE1L) Mouse Monoclonal Antibody [Clone ID: OTI4G1] (M04496) Images



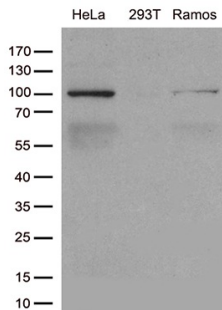
Immunohistochemical staining of paraffin-embedded Human Gastric Carcinoma using anti-CSE1L mouse monoclonal antibody. (Heat-induced epitope retrieval by 1mM EDTA in 10mM Tris buffer (pH8.5) at 120°C for 3min)



Immunohistochemical staining of paraffin-embedded Carcinoma of Human kidney tissue using anti-CSE1L mouse monoclonal antibody. (Heat-induced epitope retrieval by 1mM EDTA in 10mM Tris buffer (pH8.5) at 120°C for 3min)



HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY CSE1L (Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-CSE1L.



Western blot analysis of extracts (35ug) from cell lines by using anti-CSE1L monoclonal antibody (1:500).

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