Bcl10 (H-2): sc-55511



The Power to Question

BACKGROUND

BcI10, also designated CIPER, c-CARMEN and mE10, was first identified as a gene truncated or mutated in MALT B cell lymphomas and other tumor types. BcI10 is homologous to the equine herpes virus-2 E10 gene, and like E10 it contains an amino-terminal caspase recruitment domain (CARD). Expression of BcI10 was shown to induce NF κ B activation in a NIK-dependent pathway, and the CARD domain was shown to be essential for this activation. In a separate study, BcI10 by itself did not induce JNK or NF κ B activation. Overexpression of BcI10 was shown to induce apoptosis, in a manner that was dependent on CARD-mediated oligomerization. BcI10 was also shown to play a role in processing of caspase-9 to its active dimer. Other studies have shown that BcI10 is not mutated in many human tumors and lymphomas.

REFERENCES

- Ye, H., et al. 2000. Bcl10 expression in normal and neoplastic lymphoid tissue. Nuclear localization in MALT lymphoma. Am. J. Pathol. 157: 1147-1154.
- 2. Ruland, J., et al. 2001. Bcl10 is a positive regulator of antigen receptorinduced activation of NFκB and neural tube closure. Cell 104: 33-42.
- 3. Lucas, P.C., et al. 2001. Bcl10 and MALT1, independent targets of chromosomal translocation in malt lymphoma, cooperate in a novel NFκB signaling pathway. J. Biol. Chem. 276: 19012-19019.
- 4. Yui, D., et al. 2001. Interchangeable binding of Bcl10 to TRAF2 and cLAPs regulates apoptosis signaling. Oncogene 20: 4317-4323.
- 5. Thome, M., et al. 2002. Bcl10. Curr. Biol. 12: R45.
- Zhou, H., et al. 2004. Bcl10 activates the NFκB pathway through ubiquitination of NEMO. Nature 427: 167-171.
- 7. Fischer, K.D., et al. 2004. New roles for Bcl10 in B-cell development and LPS response. Trends Immunol. 25: 113-116.

CHROMOSOMAL LOCATION

Genetic locus: BCL10 (human) mapping to 1p22.3; Bcl10 (mouse) mapping to 3 H2.

SOURCE

Bcl10 (H-2) is a mouse monoclonal antibody raised against amino acids 1-197 mapping at the N-terminus of Bcl10 of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_{2b}$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

APPLICATIONS

Bcl10 (H-2) is recommended for detection of Bcl10 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000)

Suitable for use as control antibody for Bcl10 siRNA (h): sc-29793, Bcl10 siRNA (m): sc-29794, Bcl10 shRNA Plasmid (h): sc-29793-SH, Bcl10 shRNA Plasmid (m): sc-29794-SH, Bcl10 shRNA (h) Lentiviral Particles: sc-29793-V and Bcl10 shRNA (m) Lentiviral Particles: sc-29794-V.

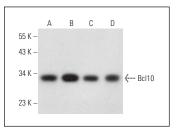
Molecular Weight of Bcl10: 33 kDa.

Positive Controls: Bcl10 (h): 293T Lysate: sc-116437, Hep G2 cell lysate: sc-2227 or Daudi cell lysate: sc-2415.

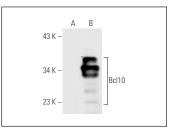
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA







BcI10 (H-2): sc-55511. Western blot analysis of BcI10 expression in non-transfected: sc-117752 (**A**) and human BcI10 transfected: sc-116437 (**B**) 293T whole call lysates

SELECT PRODUT CITITATIONS

 Hong, T., et al. 2024. PARP9 knockdown confers protection against chemoresistance and immune escape of breast cancer cells by blocking the PI3K/AKT pathway. Arch. Med. Sci. 20: 1228-1248.



See **BcI10 (331.3): sc-5273** for BcI10 antibody conjugates, including AC, HRP, FITC, PE, Alexa Fluor® 488 and Alexa Fluor® 647.