

Mouse (monoclonal) anti-Topoisomerase-1

PRODUCT ANALYSIS SHEET

Catalog Number: 44321M Volume: 50 μL Clone Number: 23B11

Isotype: IgG1 (mouse)

Form of Antibody: Mouse monoclonal immunoglobulin in PBS, pH 7.3, with PEG and Sucrose

Preservative: 0.09% sodium azide (Caution: sodium azide is a poisonous and hazardous substance. Handle with care

and dispose of properly.)

Purification: The antibody was purified from serum-free cell culture supernatant by subsequent thiophilic

adsorption and size exclusion chromatography.

Immunogen: Peptide containing amino acid residues 699-725, conjugated to KLH.

Target Summary: DNA Topoisomerases catalyze strand breakage of DNA molecules. During DNA cleavage, a tyrosyl

residue of the enzyme breaks the DNA strand by forming a covalent phosphotyrosine residue.

Rejoining of the DNA strands occurs by a second transesterification reaction.

Specificity: This Mab specifically recognizes Topoisomerase 1 (90 kDa).

Species Reactivity: Human

Applications: The antibody is suitable for Western blotting. Other applications may be possible but have not been

testea

Suggested Working

Dilutions:

determined empirically for each specific application.

Recommended Positive

Control:

Cell lysate from untreated HepG2 cells.

Storage: Upon arrival, we recommend a brief centrifugation before opening to settle vial contents. Then,

apportion the antibody into working aliquots and store at -20°C. Avoid repeated freeze / thaw cycles.

Immunoblotting: 0.5 µg/mL for HRP/ECL detection. The optimal antibody concentration should be

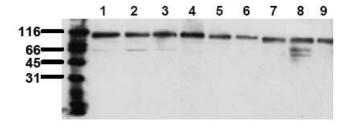
Expiration Date: Expires one year from date of receipt when stored as instructed.

PI44321M

MAN0007942 Rev 1.00 Effective Date: 27 MAR 2013

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Manufacturing site: 7335 Executive Way | Frederick, MD 21704 | Toll Free in USA 800.955.6288 www.lifetechnologies.com



Western Blotting

Whole cell lysates of serum starved HeLa (1), HepG2 (2), HEK293 (3), SH-SY5Y (4), MDCK (5) PC12 (6), CMT 93 (7), Neuro 2A (8) and 3T3 (9) tumor cells (approximately 20,000 cells per lane) were resolved by SDS-PAGE and transferred to PVDF. The membrane was blocked with a casein/Tween 20 buffer, then incubated with the Mab at 0.5 μ g/mL for 1 hour at room temperature. After washing, the membrane was incubated with an anti-mouse HRP-conjugated secondary antibody and signals were detected using an ECL detection method (exposure time: 30 seconds).

The data show that the Mab recognizes various levels of endogenous Topoisomerase 1 in these cell systems.

Explanation of symbols

Symbol	Description	Symbol	Description	Symbol	Description
***	Manufacturer	REF	Catalog number	LOT	Batch code
Ξ	Use by	1	Temperature limitation		
$\bigcap i$	Consult instructions for use	<u>^</u>	Caution, consult accompanying documents		

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