

**Datasheet for 200-301-V02****5-mC Antibody****Overview**

<b>Description:</b>	Anti-5-mC (MOUSE) Monoclonal Antibody - 200-301-V02
<b>Item No.:</b>	200-301-V02
<b>Size:</b>	100 µg
<b>Applications:</b>	ChIP, Dot Blot, ELISA
<b>Reactivity:</b>	Plants, Vertebrates
<b>Host Species:</b>	Mouse

**Product Details**

<b>Background:</b>	The 5-methylcytosine antibody (clone 33D3) is the most published and widely used antibody for DNA methylation analysis. Anti-5mC is ideal for research in Epigenetics due to its applicability to DNA methylation analysis.
<b>Synonyms:</b>	5-methylcytosine, 5-methyl cytosine
<b>Host Species:</b>	Mouse
<b>Clonality:</b>	Monoclonal
<b>Clone ID:</b>	33D3
<b>Format:</b>	IgG1

**Target Details**

<b>Reactivity:</b>	Plants, Vertebrates
<b>Immunogen Type:</b>	Other
<b>Immunogen:</b>	Anti-5-mC Antibody was produced in mice by repeated immunization with 5-methylcytosine.
<b>Purity/Specificity:</b>	Anti-5-mC Antibody was purified by gel filtration. Cross-reactivity with 5-mC from other sources has not been determined.

**Application Details**

<b>Tested Applications:</b>	ChIP, Dot Blot, ELISA
<b>Application Note:</b>	Anti-5-mC Antibody is tested for Dot Blotting, ELISA, Immunofluorescence, Methylated DNA Immunoprecipitation (MeDIP), and Sequencing (MeDIP-seq). Specific conditions for reactivity should be optimized by the end user.
<b>Assay Dilutions:</b>	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.
<b>ChIP:</b>	1 µg/ChIP
<b>ELISA:</b>	1:3,000
<b>WB:</b>	1:1,000

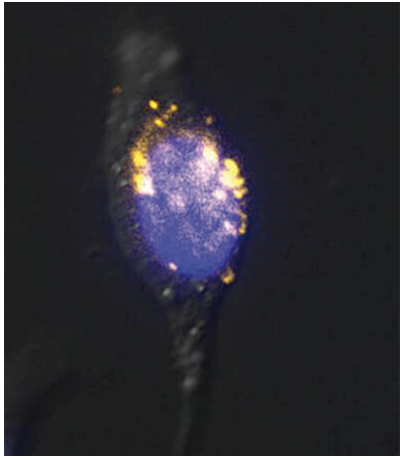
## Formulation

<b>Physical State:</b>	Liquid (sterile filtered)
<b>Concentration:</b>	1.24 mg/ml by UV absorbance at 280 nm
<b>Buffer:</b>	0.01 M Sodium Phosphate, 0.25 M Sodium Chloride, pH 7.2
<b>Preservative:</b>	0.05% (w/v) Sodium Azide
<b>Stabilizer:</b>	None

## Shipping & Handling

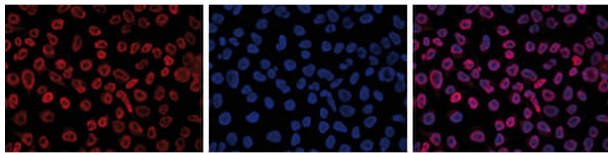
<b>Shipping Condition:</b>	Dry Ice
<b>Storage Condition:</b>	Store vial at -20° C prior to opening. Aliquot contents and freeze at -20° C or below for extended storage. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.
<b>Expiration:</b>	Expiration date is one (1) year from date of receipt.

## Images



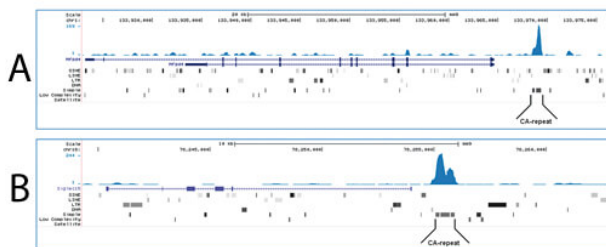
### Immunofluorescence Microscopy

Immunofluorescence Microscopy results of Mouse anti-5-mC antibody. Tissue: Interphase HeLa cell. Fixation: 4% formaldehyde in PBS for 10 min at RT. Retrieval: 0.5% Triton X-100 for 1 hour and treated with 2N HCl for 1 hour. Block: PBS/0.1% TritonX-100/1% BSA. Primary antibody: 5-mC antibody at 1:500 for 1 hr at RT. Secondary antibody: Goat anti-Mouse FITC secondary antibody at 1:10,000 for 45 min at RT. Staining: : 5-mC antibody as yellow fluorescent signal with Hoescht staining (blue).



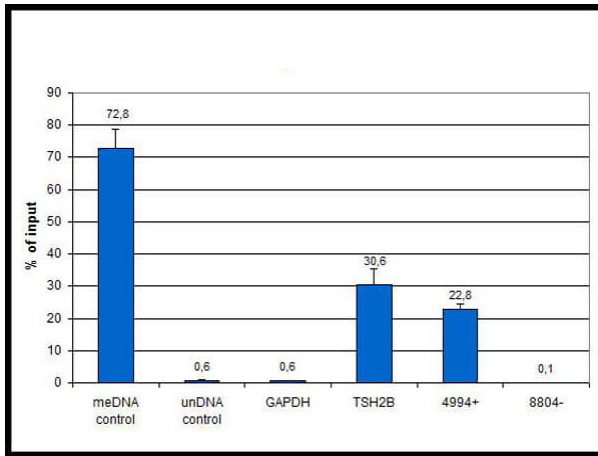
### Immunofluorescence Microscopy

Immunofluorescence Microscopy results of Mouse anti-5-mC antibody. Tissue: HeLa cells. Fixation: 4% formaldehyde in PBS for 10 min at RT. Retrieval: 0.5% Triton X-100 for 1 hr, treated with 2N HCl for 1 hr. Block: PBS/0.1% Triton X-100/1% BSA. Primary antibody: 5-mC antibody at 1:500 for 1 hr at RT (left). Secondary antibody: Goat anti-Mouse Alexa594 secondary antibody at 1:10,000 for 45 min at RT. Staining: 5-mC antibody as red fluorescent signal (left), DAPI blue (middle), merge of the two stainings (right).



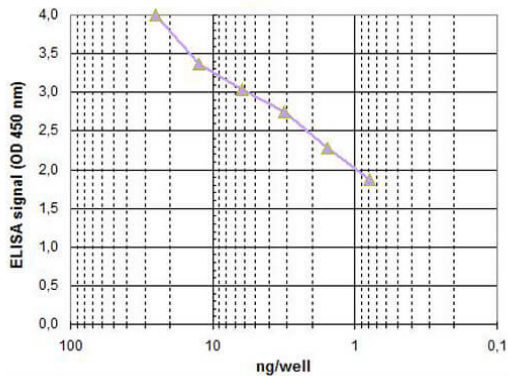
### ChIP

Methylated DNA immunoprecipitation sequencing with anti-5-mC Antibody. Genomic DNA from E14 ES cells were sheared to generate random fragments (size range 300 to 700 bp). One  $\mu\text{g}$  of the fragmented DNA was ligated to Illumina adapters and the resulting DNA was used for a standard MeDIP assay, using 2  $\mu\text{g}$  of the 5-mC. Figure A and B show Genome browser views of CA simple repeat elements with read distributions specific for 5-mC at 2 gene locations (Siglec15 and Mfsd4). Visual inspection of the peak profiles in a genome browser reveals high enrichment of CA simple repeats in affinity-enriched methylated fragments after MeDIP with the Mouse Anti-5-mC Antibody.



### Immunoprecipitation

Methylated DNA Immunoprecipitation (MeDIP) was performed on fragmented human genomic DNA using the monoclonal antibody against 5-mC. The fragmented DNA was spiked with controls methylated DNA (meDNA) as a positive and unmethylated DNA (unDNA) as a negative control) prior to performing the IP. QPCR was performed with primer sets and for a known methylated (TSH2B) and unmethylated (GAPDH) genomic region. An additional internal positive and negative control locus (4994+ and 8804 -, respectively) were also tested (4994+: forward primer 5'-GGGAATATAAGGAGCGCACA-3' and reverse primer 5'-TCGGTTAAAACGGTCAGGTC-3'; 8804-: forward primer 5'-CGAGGCGTGAGTTATTCCTG-3' and reverse primer 5'-CTCTGTGGCTGAGCTCCTT-3'). This figure shows the recovery (mean of 3 experiments), expressed as a % of input (the relative amount of immunoprecipitated DNA compared to input DNA after qPCR analysis).



### ELISA

ELISA results of mouse anti-5-mC antibody. ELISA was performed using monoclonal antibody against 5-mC diluted 1:100. The wells were coated with a serial dilution of the methylated DNA control.



### Dot Blot

Dot Blot results of Mouse anti-5-mC monoclonal antibody. Antigens: hmC, mC, and, C control forms of the immunizing peptide. Load: 100 ng, 20 ng, and 4 ng as indicated. Primary antibody: 5-mC antibody at 1:250 for 45 min at 4°C. Secondary antibody: Dylight™488 anti-mouse secondary antibody at 1:10,000 for 45 min at RT. Block: 5% BLOTTO overnight at 4°C.

## Disclaimer

This product is for research use only and is not intended for therapeutic or diagnostic applications. Please contact a technical service representative for more information. All products of animal origin manufactured by Rockland Immunochemicals are derived from starting materials of North American origin. Collection was performed in United States Department of Agriculture (USDA) inspected facilities and all materials have been inspected and certified to be free of disease and suitable for exportation. All properties listed are typical characteristics and are not specifications. All suggestions and data are offered in good faith but without guarantee as conditions and methods of use of our products are beyond our control. All claims must be made within 30 days following the date of delivery. The prospective user must determine the suitability of our materials before adopting them on a commercial scale. Suggested uses of our products are not recommendations to use our products in violation of any patent or as a license under any patent of Rockland Immunochemicals, Inc. If you require a commercial license to use this material and do not have one, then return this material, unopened to: Rockland Inc., P.O. BOX 5199, Limerick, Pennsylvania, USA.