

## Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

#### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com



## Data Sheet

### Anti-H3K4me1 monoclonal antibody

Catalog #: 25252

Lot #: 150227	Host Species: Mouse
<b>Conc.:</b> 50 μg/50 μl	Species Reactivity: Human
<b>Size:</b> 50 μg	Immunogen: Synthetic peptide
Clonality: Monoclonal	Purification: Protein A purified

**Description:** Monoclonal antibody raised in Mouse against histone H3, monomethylated at lysine 4 (H3K4me1), using a KLH-conjugated syntheticpeptide

#### Background:

Formulation: PBS containing 0.05% azide and 0.05% ProClin 300

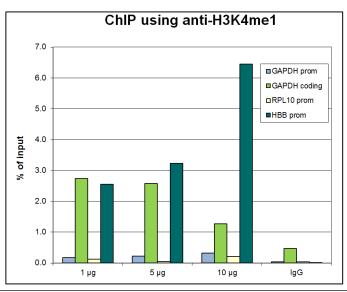
**Applications:** ChIP (1 µg/IP), ELISA (1:5000), WB (1:1000), IF (1:500)

**Storage/Stability:** Store at -80 °C for up to 2 years. Centrifuge after first thaw to maximize product recovery. Aliquot to avoid repeated freeze/thaw cycles. Aliquots may be stored at -20 °C for at least one month.

Warnings: Avoid freeze/thaw cycles

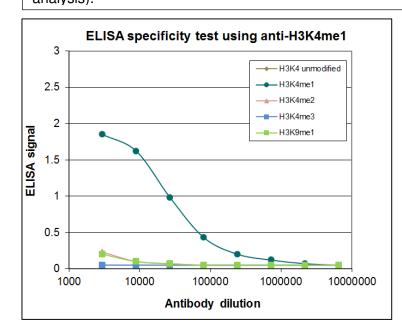
**Notes:** The optimal antibody amount per IP should be determined by the end-user. We recommend testing 1-5  $\mu$ g per IP





#### **Quality Assurance:**

ChIP results obtained with the monoclonal antibody directed against H3K4me1. ChIP assays were performed using HeLa cells, the monoclonal antibody against H3K4me1 (Cat. No. 25252) and optimized PCR primer sets for qPCR. ChIP was performed using sheared chromatin from 1.6 million cells. A titration of the antibody consisting of 1, 5 and 10  $\mu$ g per ChIP experiment was analysed. IgG (5  $\mu$ g/IP) was used as negative IP control. QPCR was performed with primers for the promoter and the coding region of the GAPDH gene, and for the RPL10 and HBB promoters. Figure 1 shows the recovery, expressed as a % of input (the relative amount of immunoprecipitated DNA compared to input DNA after qPCR analysis).



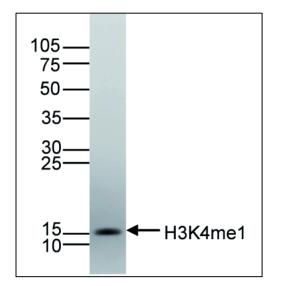
# Cross reactivity of the monoclonal antibody directed against H3K4me1.

To test the specificity an ELISA was performed using a serial dilution of the monoclonal antibody against H3K4me1 (Cat. No. 25252). The wells were coated with peptides containing the unmodified H3K4 as well as the mono-, di- and trimethylated H3K4 and the monomethylated H3K9. Figure 2 shows a high specificity of the antibody for the modification of interest.

For R&D use only. Not for human, drug, diagnostic, household or other uses. MSDS available. Product of USA. 6044 Cornerstone Court West, Suite E, San Diego, CA 92121 Tel: (858) 829 - 3082 • Fax: (858) 481 - 8694

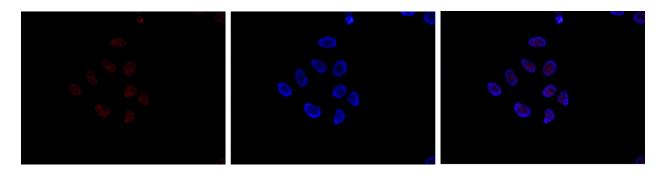
Email: info@bpsbioscience.com • Visit: www.bpsbioscience.com • © 2012 BPS Bioscience Inc.





## Western blot analysis using the monoclonal directed antibody against H3K4me1.

Histone extracts (15  $\mu$ g) from HeLa cells were analysed by Western blot using the monoclonal antibody against H3K4me1 (Cat. No. 25252) diluted 1:1,000 in TBS-Tween containing 5% skimmed milk. The position of the protein of interest is indicated on the right; the marker (in kDa) is shown on the left.



#### Immunofluorescence using the monoclonal antibody directed against H3K4me1.

HeLa cells were stained with the antibody against H3K4me1 (Cat. No. 25252) and with DAPI. Cells were fixed with 4% formaldehyde for 10 minutes and blocked with PBS/TX-100 containing 5% normal goat serum and 1% BSA. The cells were immunofluorescently labelled with the H3K4me1 antibody (left) diluted 1:500 in blocking solution followed by an anti-mouse antibody conjugated to Alexa594. The middle panel shows staining of the nuclei with DAPI. A merge of the two stainings is shown on the right.