

Product Data Sheet

Vector

RelA

45 kDa

HEK293 cells were transfected with RelA or empty vector and 24hrs later cell extracts harvested using a 1%

CHAPS lysis buffer. Extracts were resolved by non-denaturing, non-reducing electrophoresis,

transferred to nitrocellulose, and probed with a 1:500 dilution purified W6/32 . Proteins were visualized

using a goat anti-mouse secondary antibody conjugated to HRP and a

system. These data document that

was provided by Dr. Ezra Burstein, University of Michigan Medical School, Ann Arbor, MI).

MHC class I was upregulated in cells constitutively expressing RelA. (Data

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Log Fluoresence Intensity Human peripheral blood lymphocytes stained with purified W6/32, followed

by anti-mouse IgG FITC

chemiluminescence detection

100

Purified anti-human HLA-A,B,C

Catalog # / Size: 311402 / 100 µg

Clone: W6/32

Isotype: Mouse IgG2a, κ

Reactivity: Human, Cross-Reactivity: Chimpanzee, Baboon, Cynomolgus, Rhesus,

Cattle (Bovine, Cow), Cat (Feline)

Preparation: The antibody was purified by affinity chromatography.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

Concentration: 0.5 mg/ml

Storage: The antibody solution should be stored undiluted at 4°C.

Applications:

Applications: FC - Quality tested IP, WB, IHC - Reported in the literature

Recommended Usage: Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For immunofluorescent staining, the

suggested use of this reagent is ≤ 2.0 μg per 106 cells in 100 μl volume or 100 µl of whole blood. It is recommended that the reagent be titrated for

optimal performance for each application.

Application Notes: Clone W6/32 recognizes a monomorphic epitope on the 45 kD polypeptide products of HLA-A, B, C¹⁸.

(Endotoxin < 0.01 EU/µg).

Additional reported applications (for the relevant formats) include: immunoprecipitaton², Western blotting (non-reducing)³, immunohistochemical staining of acetone-fixed frozen tissue sections^{4,5}, blocking^{6,7}, inhibition of NK cell-mediated lysis¹⁰, and activation^{8,9}. Clone W6/32 has been reported not to be suitable for immunohistochemistry on paraffin sections¹⁷. The LEAF The Leaf purified antibody (Endotoxin <0.1 EU/μg, Azide-Free, 0.2 μm filtered) is recommended for functional assays (Cat. No. 311412). For highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 311428) with a lower endotoxin limit than standard LEAF™ purified antibodies

- Application References: 1. Darrow TL, et al. 1989. J. Immunol. 142:3329.

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 - 6. DeFelice M, et al. 1990. Cell. Immunol. 126:420. 7. Fayen J, et al. 1998. Int. Immunol. 10:1347.

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 14. Lunemann A, et al. 2008. J. Immunol. 181:6170. PubMed
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 17. Vambutas A, et al. 2000. Clin. Diagn. Lab. Immun. 7:79.

 - 18. Coppieters KT, et al. 2012. J. Exp. Med. 209:51. (epitope) 19. Crivello P, et al. 2013. Hum Immunol. 22:100. PubMed.

Description: MHC class I antigens associated with β2-microglobulin are expressed by all human nucleated cells. MHC class I molecules are involved in presentation of antigens to CD8+ T cells. They play an important role in cell-mediated

immune responses and tumor surveillance.

Antigen References: 1. Barclay AN, et al. Eds. 1993. The Leukocyte Antigen FactsBook. Academic Press Inc. San Diego.

Related Products: Product Clone Application FC, IHC, IP, WB, CyTOF® Purified anti-human HLA-DR L243



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APC Goat anti-mouse IgG (minimal x-reactivity) Biotin Goat anti-mouse IgG (minimal x-reactivity) FITC Goat anti-mouse IgG (minimal x-reactivity) Purified Mouse IgG2a, κ Isotype Ctrl PE Goat anti-mouse IgG (minimal x-reactivity) Cell Staining Buffer RBC Lysis Buffer (10X)

Poly4053 Poly4053 Poly4053 MOPC-173 Poly4053 FC FC, ELISA, IHC, IF, WB FC FC, ICC, IF, IHC, IP, WB FC FC, ICC, ICFC FC, ICFC



