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MAGE-A3 Antibody



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Applications: Reactivity: Sensitivity: MW (kDa): Source: **UniProt ID:** Entrez-Gene Id: WB. IP Н Endogenous 45 Rabbit #P43357 4102 **Product Usage Application** Dilution Information Western Blotting 1:1000 Immunoprecipitation 1:50 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at -**Storage** 20°C. Do not aliquot the antibody. Specificity / Sensitivity MAGE-A3 Antibody recognizes endogenous levels of total MAGE-A3 protein. This antibody cross-reacts with MAGE-A6. Source / Purification Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human MAGE-A3 protein. Antibodies are purified by protein A and peptide affinity chromatography.

Background

Cancer/testis antigens (CTAs) are a family of more than 100 proteins whose normal expression is largely restricted to immune privileged germ cells of the testis, ovary, and trophoblast cells of the placenta. Although most normal somatic tissues are void of CTA expression, due to epigenetic silencing of gene expression, their expression is upregulated in a wide variety of human solid and liquid tumors (1,2). As such, CTAs have garnered much attention as attractive targets for a variety of immunotherapy-based approaches to selectively attack tumors (3).

Melanoma antigen-A3 (MAGE-A3) is a cancer testis antigen and belongs to the type I MAGE family of proteins. The expression of MAGE-A3 is normally restricted to the human testis but is aberrantly upregulated in a number of human cancers, such as lung cancer, colorectal cancer, and multiple myeloma (4-6). Research studies have recently demonstrated that MAGE-A3 drives tumorigenesis as part of the MAGE-A3-TRIM28 ubiquitin ligase complex that promotes proteasomal degradation of the tumor suppressor kinase AMPK (7). Due to its upregulated and selective expression in human tumors and high degree of immunogenicity, MAGE-A3 has received significant attention as a novel immunotherapy target through the use of vaccines and adoptive cell therapy (8,9).

Background References

- 1. Caballero, O.L. and Chen, Y.T. (2009) Cancer Sci 100, 2014-21.
- 2. De Smet, C. et al. (1999) Mol Cell Biol 19, 7327-35.
- 3. Gjerstorff, M.F. et al. (2015) ${\it Oncotarget}$ 6, 15772-87.
- 4. Jang, S.J. et al. (2001) Cancer Res 61, 7959-63.
- 5. Shantha Kumara, H.M. et al. (2012) Cancer Immun 12, 16.
- 6. Atanackovic, D. et al. (2007) *Blood* 109, 1103-12.
- 7. Pineda, C.T. et al. (2015) Cell 160, 715-28.
- 8. Straetemans, T. et al. (2012) Clin Dev Immunol 2012, 586314.
- 9. Esfandiary, A. and Ghafouri-Fard, S. (2015) Immunotherapy 7, 683-704.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

WB: Western Blotting IP: Immunoprecipitation

3/23/24. 10:52 AM

Cross-Reactivity Key

Trademarks and Patents

Limited Uses

MAGE-A3 Antibody (#25800) Datasheet Without Images Cell Signaling Technology

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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