



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Lieferung & Zahlungsart

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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# MAGE-H1 (h): 293T Lysate: sc-159163

## BACKGROUND

The melanoma-associated antigen (MAGE) family consists of a number of antigens recognized by cytotoxic T lymphocytes. The MAGE genes were initially isolated from different kinds of tumors and, based on their virtually exclusive tumor-specific expression in adult tissues, they have been used as targets for cancer immunotherapy. MAGE genes encode for tumor-rejection antigens that are expressed in tumors of different histologic types and in normal testis and placenta. MAGE-H1 (melanoma-associated antigen H1), also known as restin or APR1 (apoptosis-related protein 1), is a 219 amino acid protein that contains a type II MAGE homology domain (MHD). Enhanced ligand stimulation promotes MAGE-H1 interaction with the type II death domain of NGFR p75. It is suggested that MAGE-H1 accelerates differentiation in response to nerve growth factor in cells.

## REFERENCES

1. De Plaen, E., et al. 1994. Structure, chromosomal localization, and expression of 12 genes of the MAGE family. *Immunogenetics* 40: 360-369.
2. Lucas, S., et al. 1999. A new MAGE gene with ubiquitous expression does not code for known MAGE antigens recognized by T cells. *Cancer Res.* 59: 4100-4103.
3. Serrano, A., et al. 1999. Quantitative evaluation of the expression of MAGE genes in tumors by limiting dilution of cDNA libraries. *Int. J. Cancer* 83: 664-669.
4. Kobayashi, Y., et al. 2000. Expression of MAGE, GAGE and BAGE genes in human liver diseases: utility as molecular markers for hepatocellular carcinoma. *J. Hepatol.* 32: 612-617.
5. Tcherpakov, M., et al. 2002. The p75 neurotrophin receptor interacts with multiple MAGE proteins. *J. Biol. Chem.* 277: 49101-49104.
6. Barker, P.A. and Salehi, A. 2002. The MAGE proteins: emerging roles in cell cycle progression, apoptosis, and neurogenetic disease. *J. Neurosci. Res.* 67: 705-712.
7. Shao, J.B. and Chen, Z. 2003. Expression of MAGE, GAGE, and BAGE genes in human hepatocellular carcinoma. *Zhonghua Gan Zang Bing Za Zhi* 11: 142-144.
8. Albrecht, D.E. and Froehner, S.C. 2004. DAMAGE, a novel  $\alpha$ -dystrobrevin-associated MAGE protein in dystrophin complexes. *J. Biol. Chem.* 279: 7014-7023.

## CHROMOSOMAL LOCATION

Genetic locus: MAGEH1 (human) mapping to Xp11.21.

## PRODUCT

MAGE-H1 (h): 293T Lysate represents a lysate of human MAGE-H1 transfected 293T cells and is provided as 100  $\mu$ g protein in 200  $\mu$ l SDS-PAGE buffer.

## STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

## APPLICATIONS

MAGE-H1 (h): 293T Lysate is suitable as a Western Blotting positive control for human reactive MAGE-H1 antibodies. Recommended use: 10-20  $\mu$ l per lane.

Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.