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# **ProductInformation**

## Anti-GRP94 (C-terminal)

Produced in Rabbit, Affinity Isolated antibody

Product Number G 4420

### **Product Description**

Anti-GRP94 (C-terminal) is developed in rabbit using as immunogen a synthetic peptide corresponding to amino acids 733-750 located near the C-terminus of human GRP94, conjugated to KLH. This sequence is identical in mouse, rat, dog, porcine, and bovine GRP94 and highly conserved (1 amino acid substitution) in chicken GRP94. The antibody is affinity-purified using the immunizing peptide immobilized on agarose.

Anti-GRP94 (C-terminal) specifically recognizes GRP94 by various applications including immunoprecipitation, immunoblotting (94 kDa), and immunocytochemistry. Staining of the GRP94 band in immunoblotting is specifically inhibited with the GRP94 immunizing peptide (human, amino acids 733-750).

Heat shock proteins (HSP) are a class of stress proteins, which includes HSP20, HSP60, HSP70, and HSP90. These proteins are considered to function as molecular chaperones by transiently binding to newly synthesized proteins to facilitate their correct folding and assembly. The GRP94 (glucose-regulated protein 94, also known as gp96, endoplasmin precursor, tumor rejection antigen 1), is a 94 kDa Ca<sup>2+</sup>-binding glycoprotein that belongs to a subfamily of the heat shock proteins Hsp90. 1-4 Other members of the GRPs family include GRP75 and GRP78/BiP. GRPs are unresponsive to heat stress and are induced by stress related to glucose starvation or defects in glycoprotein processing. GRP94 is a chaperone protein constitutively localized to the endoplasmic reticulum (ER) of mammalian cells. 1,2,5 GRP94 displays diverse functions in the ER including protein folding, sorting and secretion, and binding of immunogenic peptides. The N-terminal region of GRP94 is highly conserved in Hsp90 and

serves as the peptide binding site. GRP94 is involved in peptide antigen presentation to major histocompatibility complex class I molecules of antigen presenting cells (APCs). GRP94/gp96 binds specifically to CD91 ( $\alpha$ 2-macroglobulin receptor), and possibly to the Toll-like receptors (TLRs) on the surface of APCs. GRPs are induced as a survival response to various stress conditions including glucose starvation, depletion of Ca $^{2+}$  stores, acidosis, and hypoxia conditions that are also common in poorly vascularized tumor tissues. Overexpression of GRP94 has been associated with cellular transformation and tumorigenesis. In a variety of cancer cell lines (rodent tumor models and human biopsies), the level of GRP94 is elevated correlating with increased tumor proliferation and increased resistance to cancer treatments.

# Reagent

The antibody is supplied as a solution in 0.01 M phosphate buffered saline, pH 7.4, containing 15 mM sodium azide.

Antibody concentration: ~1.0 mg/mL

## **Precautions and Disclaimer**

Due to the sodium azide content, a material safety data sheet (MSDS) for this product has been sent to the attention of the safety officer of your institution. Consult the MSDS for information regarding hazardous and safe handling practices.

### Storage/Stability

For continuous use, store at 2-8 °C for up to one month. For extended storage, freeze in working aliquots. Repeated freezing and thawing is not recommended. Storage in "frost-free" freezers is also not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use. Working dilutions should be discarded if not used within 12 hours.

#### **Product Profile**

By immunoblotting, a working antibody concentration of 0.5-1  $\mu$ g/mL is recommended using whole cell extracts of the human epitheloid carcinoma HeLa cells, Madin-Darby canine kidney (MDCK) cells, and mouse fibroblast NIH3T3 cells.

By immunoprecipitation, 10-20  $\mu g$  of the antibody can immunoprecipitate GRP94 protein from a MDCK whole cell lysate.

Note: In order to obtain the best results using various techniques and preparations, we recommend determining the optimal working dilutions by titration.

#### References

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