

재조합 대장암 항원 단백질 GA733-2의 발현과 쥐에서의 면역반응 효과

¹경희대학교 생명공학원, ²원광대학교 생명과학부, ³전남대학교 생명과학기술학부,
⁴한국생명공학원 단백질치료연구센터
정하영¹, 김경일¹, 이현호¹, 석연주¹, 고기성², 정영희³, 강형식³, 오두병⁴, 정인식^{1*}

**Expression of Recombinant Colorectal Cancer Antigens GA733-2 and
Its Effect on Immune Responses in Mice**

¹Graduate School of Biotechnology, Kyung Hee University, Yongin 446-701, Korea,
²Department of Life Science, Won Kwang University, Iksan 156-756, Korea, ³School of
Biological and Science and Technology, Chonnam National University, Gwangju 500-757,
Korea, ⁴Protein Therapeutics Research Center, Korea Research Institute of Bioscience and
Biotechnology, Daejeon 305-333, Korea
Ha-Young Chung¹, Kyung-Il Kim¹, Hyun-Ho Lee¹, Yeon-Ju Seok¹, Ki-Sung Ko², Young-Hee
Joung³, Hyung-Sik Kang³, Doo-Byoung Oh⁴ and In-Sik Chung^{1*}

Objectives

The GA733-2 antigen is a 40 kDa human cell surface glycoprotein defined by the murine mAb3 GA733. This antigen has been found to be associated with a variety of human carcinoma such as colorectal, pancreatic, and breast carcinoma GA733-2 can be an immuno therapeutic target of the most frequent human cancer.

Materials and Methods

- Materials - pGEM-T vector, Drosophila melanogaster Schneider (S2) cell, Ni-NTA resin
- Methods - Cell culture, transfection, western blot analysis, ELISA

Results

In this study, we examined the expression of recombinant GA733-2 using DES insect cell and plant systems. The gene insertion was confirmed in stably tranfected insect cells and plants by Southern hybridization analysis. Recombinant GA733-2 was expressed with a molecular size of approximately 35~37 kDa and 40 kDa insect cells and plants, respectively. We are currently investigating immunization of recombinant GA733-2 from insect cells and plants in mice. Our presentation will also include the potential merits of both systems expressing recombinant cancer antigens necessary for the research of future vaccine development.

* 시험성적

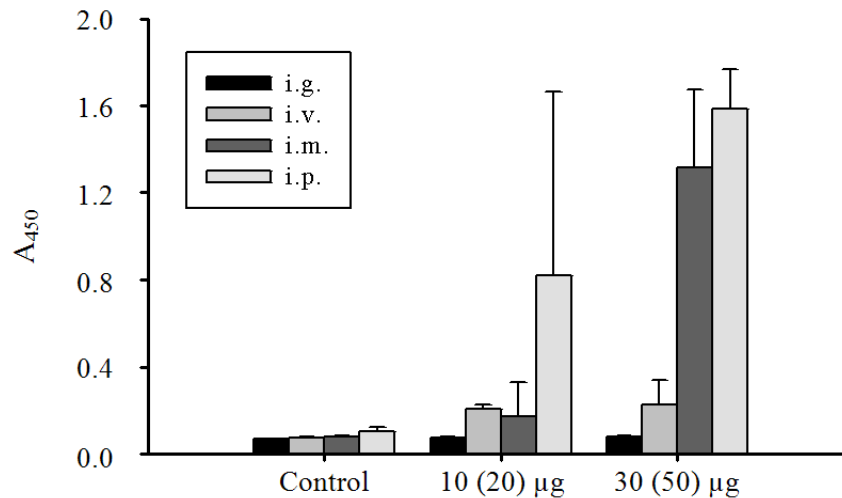


Figure 1. Production of anti-GA733-2 IgG antibodies in serum induced by immunization of GA733-2. BALB/c mice were immunized by intragastric (i.g.), intravenous (i.v.), intramuscular (i.m.) and intraperitoneal (i.p.) delivery on days 0, 7 and 21 with GA733-2. Serum samples were collected at day 28 and diluted 1:1000. ELISA showed serum IgG levels against GA733-2. The samples were tested in triplicate runs. The bars represent the means of OD₄₅₀ plus standard deviations.

References

1. Birebent, B., Somasundaram, R., Purev, E., Li, W., Mitchell, E., Hoey, D., Bloom, E., Mastrangelo, M., Maguire, H., Harris, D. T., Staib, L., Braumuller, H., Leeser, C., Kuttner, N., Beger, H. G. and Herlyn, D. (2001) Anti-idiotypic antibody and recombinant antigen vaccines in colorectal cancer patients. *Crit Rev Oncol Hematol.* 39, 107-113.
2. Brodzik, R., Spitsin, S., Golovkin, M., Bandurska, K., Portocarrero, C., Okulicz, M., Steplewski, Z. and Koprowski, H. (2008) Plant-derived EpCAM antigen induces protective anti-cancer response. *Cancer Immunol Immunother.* 57,317-323.
3. Cirulli, V., Crisa, L., Beattie, G. M., Mally, M. I., Lopez, A. D., Fannon, A., Ptasznik, A., Inverardi, L., Ricordi, C., Deerinck, T., Ellisman, M., Reisfeld, R. A. and Hayek, A. (1998) KSA antigen Ep-CAM mediates cell-cell adhesion of pancreatic epithelial cells: morphoregulatory roles in pancreatic islet development. *J. Cell Biol.* 140, 1519-1534.
4. Gottlinger, H. G., Funke, I., Johnson, J. P., Gokel, J. M. and Riethmuller, G. (1986) The epithelial cell surface antigen 17-1A, a target for antibody-mediated tumor therapy: its biochemical nature, tissue distribution and recognition by different monoclonal antibodies. *Int. J. Cancer.* 38, 47-53.