

**Development of new immunoradiometric assay for CA 125
antigen using two monoclonal antibodies produced
by immunizing lung cancer cells**

Mihoko KUNIMATSU, Keigo ENDO, Tetsuo NAKASHIMA,** Toshikazu AWAJI, Tsuneo SAGA,
Yuji WATANABE, Yasutaka KAWAMURA, Hitoya OHTA, Mitsuru KOIZUMI,
Harumi SAKAHARA, Junji KONISHI, Shingo FUJII,* Takahide MORI,* Kanji TORIZUKA,**
Yoichiro MATSUOKA,*** Tsuyoshi NAKAGAWA*** and Nobuo YAMAGUCHI***

Departments of Nuclear Medicine and Gynecology, Kyoto University Hospital, Kyoto
Fukui Medical College**, Fukui
Department of Radiology, Mie University Hospital***, Mie*

CA 125 is an antigen associated with non-mucinous epithelial ovarian cancer, which is defined by OC 125 antibody developed by immunizing ovarian cancer cells. We have produced two monoclonal antibodies, 130-22 and 145-9, by using the human lung adenocarcinoma cell line PC-9. Both 130-22 and 145-9 antibodies recognized CA 125 antigen. However, the binding sites seemed to be separate from those of OC 125. Testing by 9 immunoradiometric assays (IRMA), using different combinations of the 3 monoclonal antibodies 130-22, 145-9 and OC 125 demonstrated that the best standard curve for detecting CA 125 could be obtained by a "simultaneous sandwich" assay based on a mixture of ¹²⁵I-labeled OC 125 and 130-22 or 145-9 coated beads. One-step IRMA, using 130-22 as a tracer and 145-9 as an immunoadsorbent, also showed good reproducibility and sensitivity for measuring CA 125. Antigens were detectable in the culture supernatants of PC-9 cells and 5 of 6 ovarian cancer and endometrial adenocarcinoma cells. These results indicate that one-step IRMA using 130-22 and 145-9 is useful for detecting CA 125 antigen.

Key words: Monoclonal antibody, CA 125, Immunoradiometric assay, Ovarian cancer, Lung cancer