

Expression of CA125 and cisplatin susceptibility of pleural effusion-derived human lung cancer cells from a Thai patient

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Abstract. Advances in understanding lung cancer biology and tumor markers aid clinicians in managing the disease. Cancer-associated antigen (CA)125 has garnered increasing attention in lung cancer research and may benefit the treatment and follow-up of this type of cancer. In Thai lung cancer patients, knowledge regarding ethnic differences in cancer cell biology is largely absent. We generated lung cancer cells from the pleural effusion fluids of a Thai patient and designated these as P1 cells. P1 cells were assessed for growth rate, response to chemotherapy, and the presence of tumor markers, in particular CA125 expression. Results of immunofluorescence indicated that P1 cells exhibited strong expression levels of CA125, comparable to that of established H460 lung cancer cells. Furthermore, P1 cells were analyzed for the expression of additional markers. Results revealed that H460 cells exhibited strong immunofluorescent signals from cytokeratin-19 fragments (CYFRA 21-1) and squamous cell carcinoma antigen (SCCA) while P1 presented only CYFRA 21-1 signals. We also found evidence of relative cisplatin resistance in P1 compared to the susceptibility level of established lung cancer cells. Thus, the results and methodology described in this study may aid the development of lung cancer diagnostic and therapeutic approaches and, in particular, advance understanding of ethnic differences.

Introduction

The identification of tumor markers leads to a significant improvement in cancer therapy and aids investigators in understanding cancer biology (1). Tumor markers are useful

in cancer detection and classification (2). Moreover, in treating patients, serum levels of such tumor markers are helpful in the determination of cancer prognosis and likelihood of recurrence (3). Among several well-known tumor markers, cancer-associated antigen (CA)125 has garnered increasing attention in lung cancer research. CA125 is a glycoprotein found on the cell membrane which has been used as a standard tumor marker for the diagnosis and follow-up of ovarian cancers (4,5). Moreover, a significant increase of CA125 has been reported in other cancers including breast and lung cancers has been noted (6,7).

Among the various anti-cancer agents used for lung cancer treatment, cisplatin [cis-diamminedichloroplatinum (II)] is commonly prescribed (8). Cisplatin mediates cancer cell apoptosis through reactive oxygen species (ROS)-dependent and DNA adduct pathways (9,10). Cisplatin-induced apoptosis is mainly mediated through the caspase signaling pathway (11). Although available lung cancer cell lines have been widely used for the investigation of cancer cell biology and chemotherapeutic susceptibility, none of the cell lines were from Thai patients. Since ethnic diversity is known to be a factor affecting tumor marker expression and chemotherapeutic response (12,13) and knowledge regarding such cancer signatures of Thai-originated cancer cells remains elusive, the present study aimed to generate knowledge about the tumor marker expression and chemotherapeutic response of lung cancer cells obtained from a Thai patient. Consequently, we generated primary cancer cells collected from the pleural effusion fluid of a Thai patient and characterized the lung cancer signatures of the cells compared to the established lung cancer H460 cells. The obtained cells were evaluated for CA125 expression and response to cisplatin, one of the most widely used chemotherapeutic agents. The results and methodology described in this study may aid the development of lung cancer diagnostic and therapeutic approaches and, in particular, advance understanding of ethnic differences.

Materials and methods

Clinical specimens and reagents. Pleural effusions were collected from a 76-year-old male Thai patient with suspected

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lung adenocarcinoma. Informed consent was obtained from the patient and the study was approved by the ethics committee of the Faculty of Medicine and the ethics committee of the faculty of Pharmaceutical Sciences, Chulalongkorn University. Human proximal tubular epithelial renal cells (HK-2) and human lung cancer epithelial (H460) cells were obtained from the American Type Culture Collection (Manassas, VA, USA). H460 cells were cultured in RPMI-1640 while HK-2 cells were cultured in DMEM, supplemented with 10% fetal bovine serum (FBS), 2 mM L-glutamine and 100 U/ml penicillin/streptomycin in a 5% CO₂ environment at 37°C. Cisplatin, propidium iodide (PI) and Hoechst 33342 were obtained from Sigma Chemical, Inc. (St. Louis, MO, USA). Resazurin was purchased from Invitrogen (Carlsbad, CA, USA). Specific antibodies for CA125, CYFRA 21-1, and SCCA were obtained from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Resazurin and Alexa Fluor 488 goat anti-rabbit IgG (H+L) and Alexa Fluor 590 goat anti-mouse IgG (H+L) conjugated secondary antibody were purchased from Invitrogen.

Specimen preparation. Pleural effusion was centrifuged at 1600 x g for 10 min at room temperature. The pellet was resuspended with 4 ml sterile balanced salt solution and then centrifuged on a Ficoll gradient (Ficoll-Paque™, GE Healthcare, Piscataway, NJ, USA) at 400 x g for 40 min at 20°C to separate the tumor cells from erythrocytes. The layer of mononuclear cells were collected and washed twice with 3 volumes of RPMI-1640 by centrifugation at 400 x g for 10 min at 20°C. The pellet was then resuspended and the cells were cultured in ACL-4 medium supplemented with 5% FBS at 37°C and 5% CO₂.

Growth properties and cell morphology. Cells were initially seeded at a density of 1x10⁴ cells in a 24-well plate and the population doubling time was determined. At various time points, cells were trypsinized by 0.25% trypsin-EDTA treatment and the number of cells was calculated by the trypan blue exclusion method. Population doubling times were determined from an exponential regression of viable cell counts over nine days.

Immunofluorescence. Cells (5x10⁴/well) were seeded in 6-well plates for 24 h to allow the cells to completely adhere to the surface. The cells were then fixed in 3.7% formaldehyde for 10 min at room temperature, permeabilized and blocked in a solution containing 0.5% saponin, 1% FBS and 1.5% goat serum for 30 min. Following primary antibody incubation with CA125, CYFRA 21-1 or SCCA antibody at 1:100 dilution for 1 h, cells were washed and incubated together with Alexa Fluor 488 goat anti-rabbit IgG (H+L)-conjugated secondary antibody (Invitrogen) or Alexa Fluor 590 goat anti-mouse IgG (H+L)-conjugated secondary antibody (Invitrogen) for 30 min. Nuclei were stained with Hoechst dye (Invitrogen). Images were visualized by fluorescence microscopy (Olympus IX51 with DP70).

Cytotoxicity and apoptosis assay. Cytotoxicity was determined by a Presto Blue fluorescence assay. Following specific treatments, cells in a 96-well plate were incubated with 1:50 resazurin for 1 h at 37°C. Fluorescence intensity of resazurin product (resorufin) was measured at 530 nm (excitation wavelength) and 590 nm (emission wavelength) using a microplate

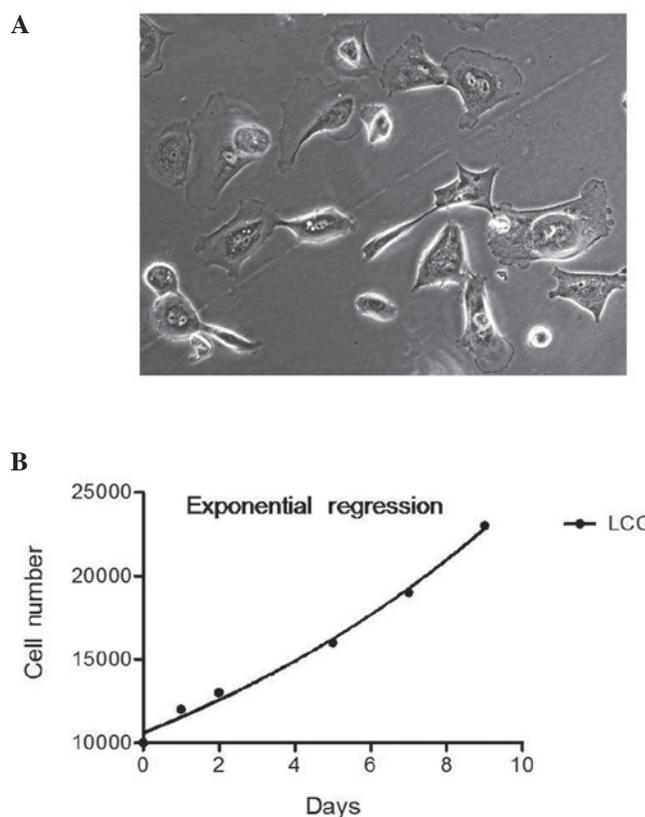


Figure 1. Phase-contrast micrographs showing morphology of a pleural effusion-derived lung carcinoma cells. (A) P1 cells exhibited epitheloid morphology with a bipolar, spindle-shaped appearance. (B) Population doubling time of P1 cells were calculated based on the exponential regression of cell growth.

reader. Cell viability was calculated as a percentage relative to non-treated cells. Analyses were performed independently in triplicate. Apoptosis was determined by a Hoechst 33342 DNA fragmentation assay. Briefly, cells were incubated with 10 µg/ml of Hoechst 33342 for 30 min and analyzed for apoptosis by scoring the percentage of cells having intensely condensed chromatin and/or fragmented nuclei on fluorescence microscopy (Olympus IX51 with DP70). The apoptotic index was calculated as the percentage of cells with apoptotic nuclei over the total number of cells.

Statistical analysis. The data are presented as the means ± SD from three or more independent experiments. Statistical analysis was performed using the Student's t-test at a significance level of P<0.05.

Results

Growth properties and morphology of lung carcinoma cells. Pleural effusion fluids of an untreated Thai patient were first centrifuged and the suspended cells were collected. The cells obtained were cultured under adherent conditions in lung cancer cell-selective ACL-4 medium as described by the National Cancer Institute (NCI) for the selective growth of non-small cell lung cancer, while rarely sparing the growth of normal cells, such as fibroblasts and macrophages. In the present study, those cells continuously propagating for at least

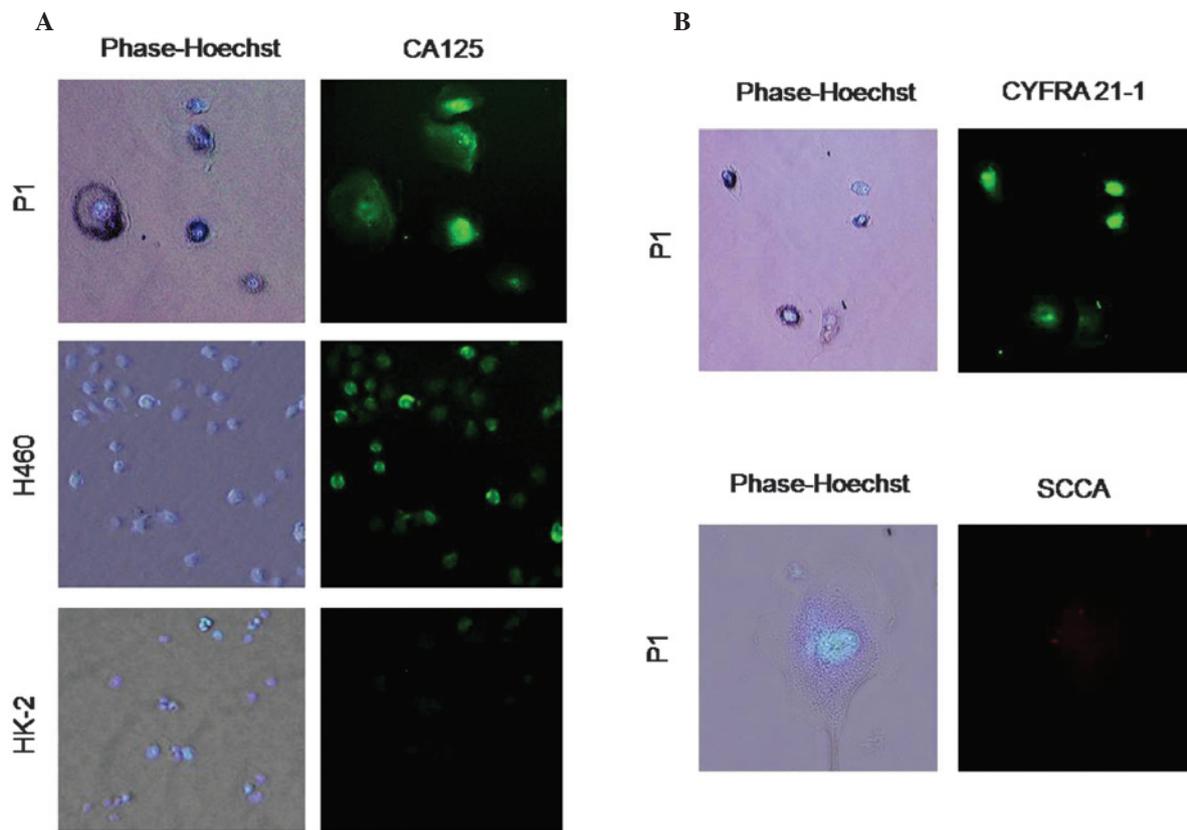


Figure 2. Immunocytochemistry evaluation for CA125, CYFRA 21-1 and SCCA expression. Primary lung carcinoma P1, lung cancer H460, and proximal epithelial HK-2 cells were evaluated for CA125 expression. The expression of CYFRA 21-1 and SCCA in P1 cells was evaluated by immunofluorescence as described in Materials and methods. The fluorescence signal was detected under a fluorescence microscope.

six months were designated as P1 cells. The clinical and pathological details of the patient from whom P1 cells were obtained are described in Materials and methods. Six months later, cells were routinely cultured in ACL-4 medium supplemented with 10% FBS. P1 cells exhibited epitheloid morphology with a bipolar, spindle-shaped appearance (Fig. 1A). Cells grew individually with little cell-cell contact. This morphology of P1 cells was markedly similar to the previously established non-small cell lung cancer cell lines from solid tumor specimens (14). Population doubling time of P1 cells was calculated based on the exponential regression of cell growth (Fig. 1B). The best-fit population doubling value was 8.127 days. Although the cells were slow-growing, they could be maintained in ACL-4 medium and continue to be cultured for a long period of time.

Expression of CA125 in P1 cells. Recent evidence suggests that CA125 may be an important tumor marker for lung cancer. Data obtained from the present study revealed that the expression profile of primary lung cancer cells originating from a Thai patient confirms this hypothesis and may facilitate the development of a lung cancer marker; we therefore evaluated the expression of CA125 in P1 cells. Immunohistochemical investigation using anti-CA125 antibody indicated that CA125 was highly expressed in P1 cells as in the positive control lung carcinoma H460 cells. Notably, CA125 signals originated from the nuclear region of the cells as well as the cell surface (Fig. 2A). Moreover, we evaluated

the specificity of CA125 expression on tumor cells by staining the non-cancerous proximal tubule epithelial HK-2 cells. Results indicated that even though HK-2 cells were stained by the CA125 antibody, the signal was only slightly increased (Fig. 2A). To provide supporting evidence for the histological sub-type of P1 cells, we performed an additional immunofluorescence study using CYFRA 21-1 (a marker for non-small cell carcinoma) and anti-SCCA (a marker for squamous cell carcinoma) antibodies. P1 cells were found to express CYFRA 21-1 but not SCCA protein (Fig. 2B). The results indicated that P1 may be either adenocarcinoma or large cell carcinoma, but not squamous carcinoma.

Cisplatin response of P1 cells. Disruption of apoptosis contributes to malignant cell growth and chemotherapeutic resistance. To determine the apoptosis phenotype of P1 cells, the cells were treated with various concentrations of cisplatin (0-200 μ M) and analyzed for cell toxicity by resazurin assay and apoptosis by Hoechst 33342 assay. H460 cells were used as a standard phenotype for non-small cell lung cancer. Cells having intensely condensed and/or fragmented nuclei were considered apoptotic. Treatment with cisplatin induced a dose-dependent decrease in cell viability and a concomitant increase of apoptotic cells in P1 and H460 cells (Fig. 3). P1 cells acquired higher apoptosis resistance to cisplatin as compared to H460 cells (Fig. 3B). We summarized the IC_{50} values of P1, H460, and the additional A549 lung cancer cell line in Table I. Since apoptotic resistance is a key

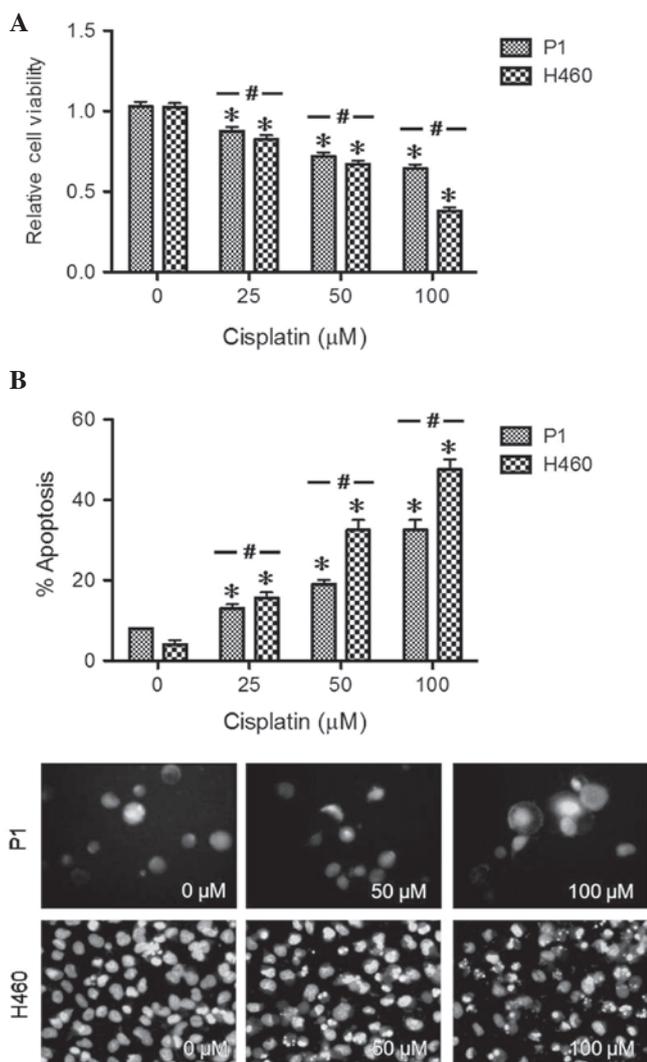


Figure 3. Susceptibility of cells in response to cisplatin-mediated apoptosis. Primary lung carcinoma P1 and lung cancer H460 cells were evaluated for their apoptotic response to various concentrations of cisplatin (0-100 μM). Cell viability and apoptosis were evaluated by resazurin and Hoechst 33342 assays, respectively. Values are the means ± SD (n=3).

characteristic of all cancer cells, this finding suggested the lung cancer-like signature of P1 cells.

Discussion

Improved understanding of cancer cell biology is likely to benefit the overall improvement of cancer therapy. In Thailand, the number of diagnosed lung cancer patients is on the increase and has become a primary cause of cancer-related mortality (15). Satisfactory clinical outcomes for lung cancer treatment are currently limited by two main obstacles which include late detection and the presence of chemotherapeutic resistance (16). Overcoming these barriers is vital, with improvement of basic knowledge of lung cancer biology, tumor markers and cellular response to chemotherapeutic agents being crucial. The present study generated P1 cells from the pleural effusion of a Thai male who was diagnosed as a lung cancer patient. The cells demonstrated typical cancerous characteristics including CYFRA 21-1 expression.

Table I. Comparison of cisplatin susceptibility of cells.

Cells	IC ₅₀ (μM)
H460	80±8.61
P1	151±10.51
A549	62±5.32

P1, H460, and A549 lung cancer cells were incubated with various concentrations of cisplatin for 24 h. The reduction of cell viability was determined by a resazurin viability assay and the IC₅₀ values were then calculated. Values are the means ± SD (n=3).

CA125 expression in lung cancer cells has previously been reported and the accumulated data suggested that this tumor marker may be of significance for lung cancer therapy (6). Furthermore, CA125 expression in lung cancer patients may be a good predictive tool for patient outcome (17). CA125 is an important tumor marker recognized by a monoclonal antibody OC125. CA125 is elevated in the serum of ovarian cancer patients and has been identified as a useful tool for the screening of ovarian cancer (4,5,18). In lung cancer, investigators have found a significant expression of CA125 in a number of lung cancer cell lines (7). In accordance with these findings, we found a strong expression of such tumor markers in lung carcinoma H460 (obtained from ATCC) and Thai-originated lung cancer P1 cells.

Accumulating evidence indicates that ethnic difference is a notable factor in the determination of the chemotherapeutic response. In the case of anti-cancer agents, regimens including doses and drug manipulating protocols are often used for different ethnic populations. In response to chemotherapeutic agents, a number of cell mechanisms are activated, supporting the idea that genetic variation by ethnicity may affect the drug responsiveness. Cisplatin is widely used for the treatment of numerous solid tumors including lung cancers (19). The mechanisms of action of the drug on tumor cells is through ROS production and DNA adduct formation (9,10). To induce cell death, several signaling pathways are activated in the regulation of survival and apoptosis and the variation in genetics may alter the cell response. Variation in response to cisplatin among cells was revealed in the present study. Two lung cancer cell models obtained from ATCC, H460 and A549, cells were used as standard cells in the evaluation of cisplatin susceptibility. Results of viability in response to cisplatin revealed that P1 exhibited slight resistance to cisplatin-mediated death compared to H460 (Fig. 3) and A549 cells (data not shown). For the apoptotic evaluation, the number of apoptotic cells in response to cisplatin in the P1 population significantly decreased as compared to those of H460 cells (Fig. 3B), suggesting that P1 cells exhibited relative cisplatin resistance. Consistent results obtained from further investigation revealed that the cisplatin IC₅₀ of P1, H460, and A549 cells was 151±10.51, 80±8.61, and 62±5.32 μM, respectively.

Based on these data, it is possible that the variation in ethnicity plays a significant role in drug susceptibility. However, investigations as to this role are required. The present study

provides a basis for the better understanding of ethnic difference in cancer cell biology.

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