Effect of Nicotine- and Tar-Removed Cigarette Smoke Extract on Cancer Metastasis

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Abstract
Cigarette smoking is known to impact the promotion of carcinogenesis and tumor metastasis. On the other hand, some components in smoke were found to have health-promoting effects, and cancer suppressor effects of components in tobacco smoke have attracted attention. Although some studies showed the cancer suppressive effect of cigarette smoke extract (CSE) in vitro study, the effect of CSE administration on cancer is controversial. In this study, we investigated the effect of CSE-administration on tumor metastasis in a spontaneous tumor metastasis model using B16-BL6 cells, which is more clinical conditions. C57BL/6NCr mice were subcutaneously inoculated B16-BL6 cells into the footpad of the right rear leg. CSE was intraperitoneally administrated for 28 days from the day of inoculation. At 2 weeks after inoculation, the primary focus was excised. Subsequently, survival days of the mice were recorded to determine the effect of CSE-administration on spontaneous metastasis. The effect of CSE, α, β-unsaturated ketones, and aldehydes on B16-BL6 cell invasiveness were confirmed by matrigel invasion assay. Survival days of mice injected with 100% CSE was significantly shortened than that of control. B16-BL6 cell invasiveness was accelerated by the treatment with 0.1% CSE and 3 µM of crotonaldehyde. Intraperitoneal CSE-administration may progress spontaneous metastasis of B16-BL6 cells via enhancement of B16-BL6 cell invasiveness. As the cause, we found that crotonaldehyde contained in CSE may enhance the invasion ability of cancer cells. To clarify the cancer-suppressing effect of tobacco components, the effect of crotonaldehyde-removed CSE on tumor should be assessed in detail.

Keywords: cigarette smoke extract (CSE), metastasis, crotonaldehyde (CA), B16-BL6 mouse melanoma cells, invasion

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1. Introduction

Cigarette smoking is widely known to impact the promotion of carcinogenesis and tumor metastasis in several cancer types [1-4]. Especially in lung cancer, it is reported that smoking rate and carcinogenic risk are significantly correlated, and about 70% of lung cancer were attributed to smoking in a study of Japanese men with a relatively low increase in cancer risk due to smoking [1]. Similarly, for the esophagus, liver, pancreas, head and neck, cervix, and bladder cancers, smoking has been reported to be one of the significant carcinogenic risks [5-10].

The Mainstream of cigarette smoke is composed of about 4,300 kinds of particulate components and about 1,000 kinds of gas components [11]. Of these, approximately 70 components have been reported to be carcinogenic or may have some adverse health effects [12]. In a study of cigarette smoke, cigarette smoke exposure was reported to increase lung metastasis and tumor volume in colon and pancreatic cancer cell lines in mice [13, 14]. Nicotine, known as one of the causes of tobacco dependence, is also known to impact tumor growth and progression [15, 16], and Nguyen et al. reported that nicotine promotes the proliferation and migration of melanoma cell line by regulating PD-L1 expression via α9 nicotinic acetylcholine receptors [17].

On the other hand, Sayed et al. found health-promoting components in cigarette smoke [18], and reported that some components, such as cembratriene-4, 6-diol and methyl vinyl ketone (MVK), have a tumor-suppressing activity [19-22]. Thus, cancer suppressor effects of components in tobacco smoke have attracted attention, and several studies about cancer suppression using tobacco smoke have been conducted. Saito et al. showed that the tumor-promoting activity of 12-O-tetradecanoylphorbol 13-acetate, one of the phorbol esters, was suppressed by cembratriene-4, 6-diol, isolated from cembratriene acetate contained in tobacco smoke concentrate, in vitro study [20], and Sayed et al. reported that cembratriene-4, 6-diol could inhibit tumor cell invasion [19]. In addition, our group has also reported that the pretreatment of highly metastatic B16-BL6 mouse melanoma (B16-BL6) cells with nicotine- and tar-removed cigarette smoke extract (CSE) could reduce the number of lung nodules of B16-BL6 cells in hematogenous lung metastasis model mice injected with B16-BL6 cells through the tail vein [21]. Furthermore, as a study to evaluate the effect of CSE on tumor metastasis in mice, Hatai et al. conducted an in vivo study on the tumor-suppressive activity of intraperitoneal (i.p.) administration of CSE in mice and showed the possibility of i.p. administration of CSE to suppress liver metastasis in a model of transsplenic liver metastasis using colon-26 [22]. They further searched for components involved in cancer metastasis suppression and found MVK, an α, β-unsaturated ketone contained in tobacco smoke, can suppress metastasis through suppression of invasion of colon-26 cells [22]. However, in the study by Hatai et al., the survival time of mice was not found to be prolonged by CSE administration [22], and the cancer metastasis-suppressing effect of CSE administration on living organisms has
not been clarified yet. Furthermore, since CSE was administered intraperitoneally in their study, it cannot be ruled out that liver metastasis may have been suppressed by direct exposure of CSE to the primary organ of tumor cell engraftment, spleen. Thus, the cancer metastasis-suppressing effect of CSE administration on living organisms remains controversial.

In this study, we investigated the effect of CSE on tumor metastasis a spontaneous tumor metastasis model in which B16-BL6 cells were seeded subcutaneously in the footpad and subsequently developed lung metastatic nodules, which is more clinical conditions and can rule out the effects of direct exposure of CSE to the primary organ of tumor cell engraftment.

2. Materials and Methods

2.1. Materials

Frontier Lights brand cigarettes containing 1 mg of tar and 0.1 mg of nicotine per cigarette, were purchased from Japan Tobacco, Inc. (Tokyo, Japan). Cambridge filters, used to remove almost all particles and nicotine from cigarette smoke, were obtained from Heinr. Borgwaldt GmbH (Hamburg, Germany). Fetal bovine serum (FBS) was from BioWest Co. (Nuaille, France). EDTA trypsin solution (EDTA: 2.2 mM, trypsin: 0.25%) was from Mediatech, Inc. (Manassas, VA, USA). Penicillin/streptomycin solution (penicillin: 50,000 U/mL, streptomycin: 50 mg/mL) was from Cosmo Bio Co., Ltd. (Tokyo, Japan). Dulbecco’s phosphate-buffered saline without calcium and magnesium [DPBS (-)] was from Nissui Pharmaceutical Co., Ltd. (Tokyo, Japan). Growth factor-reduced Matrigel matrix and FALCON cell culture inserts were from Becton Dickinson Labware (Bedford, MA, USA).

2.2. Preparation of CSE

The CSE was prepared according to the method described in a previous report [21]. Briefly, CSE was obtained by bubbling the filtered mainstream of smoke (gas phase) into DPBS (-) (1 mL per 3 cigarettes). As a filter, Cambridge filter was used to remove the particle phase containing tar and nicotine. The suction speed was kept constant (1 L/min) using a suction pump (Nippon Rikagaku Kikai Co., Ltd., Tokyo, Japan), and smoke was bubbled for 1 min. The CSE contained DPBS (-) solution was immediately filtrated with a 0.22 μm filter. The filtered solution, 100% CSE, was stored at -80°C until use and diluted to various concentrations with DPBS (-) at the time of use.

2.3. Animals

Specific pathogen-free male C57BL/6NCr mice (7 weeks old) purchased from Japan SLC, Inc. (Hamamatsu, Japan) were used as metastatic melanoma syngeneic animals. Mice were maintained in an air-conditioned room (23±2°C and 60±10% humidity) under an artificial 12-hour light/dark cycle (7:00 a.m. - 7:00 p.m.). Food and water were given ad libitum during the experimental period. This study was approved by the Animal Experiment
Committee of Mukogawa Women's University (Approval No. P-11-2012-06-A), and all procedures followed the Guidelines for the Care and Use of Laboratory Animals at the University.

2.4 Cells
A highly metastatic B16-BL6 mouse melanoma cell line was kindly provided by Dr. Futoshi Okada of Tottori University (Yonago, Japan). Cells less than 50 passages were used in all experiments. B16-BL6 cells were cultured in DMEM containing 10% FBS and 0.1% penicillin/streptomycin solution in a humidified incubator at 37°C in the presence of 5% CO₂.

2.5. Evaluation of spontaneous metastasis of tumor cells
Sub-confluent B16-BL6 cells were harvested with EDTA trypsin solution and resuspended in DPBS (+) to the appropriate concentrations. Fifty microliters of cell suspension (2 × 10⁷ cells/mL) were subcutaneously injected into the footpad of the right hind leg of syngeneic C57BL/6NCr mice. Two weeks after the inoculation, the mice were anesthetized with diethyl ether and the enlarged primary tumor was excised. After 4 weeks from tumor inoculation, 10, 30, and 100% CSE were administrated with i.p. to the mice at the dose of 16 mL/kg daily. As a control, DPBS (-) was administered. The survival of the mice in each group was followed up for 100 days and the date of death was recorded. The survival duration was determined as the number of days after tumor cell inoculation. Dead mice were dissected for confirmation of tumor metastases. Each group contained 7 animals at the start of the experiment.

2.6. Proliferation assay
B16-BL6 cells were seeded on a 12 well plate at 1 × 10⁵ cells/well. Then, cells were treated with CSE (0.01, 0.03 and 0.1% as final concentration), MVK (1, 3, 10 µM), CA (1, 3, 10 µM), ACR (1, 3, 10 µM) or DPBS(-). After 72 h of exposure to CES, cells were collected and the number of cells in each well was determined using a coulter counter.

2.7. Matrigel invasion assay
B16-BL6 cells were resuspended in FBS-free DMEM to obtain a concentration of 4.0 × 10⁵ cells/mL, and 500 µL of cell suspension was added to the upper layer within the cell culture insert coated with Matrigel on the filter. To the lower layer, DMEM containing fibronectin as a chemoattractant was added. After 24 h of incubation with CSE (0.01, 0.03 and 0.1% as final concentration), 3 µM of MVK, 3 µM of CA, 3 µM of ACR or 0.1% of DPBS(-). Uninfiltrated cells, which remain on the top of the filter, were wiped with a swab. The infiltrated cells on the bottom surface of the filter were Giemsa stained and counted under a microscope.

2.8. Statistical Analyses
Data are expressed as the mean ± S.E. Survival data were analyzed by the log-rank test. Data from in vitro experiments were analyzed by Dunnett's test. Statistical analyses were performed using the Graphpad Prism 4 software package (Graphpad
Software, Inc., San Diego, CA, USA). A difference was considered significant when \( p < 0.05 \).

3. Results

3.1. Effect of intraperitoneal CSE administration on survival time of spontaneous metastasis model mice

Although administration of CSE with a concentration of under 30% did not affect the survival time of mice, the survival time was significantly shortened by administration of 100% CSE compared to controls (Fig. 1). There was no significant change in the weight and thickness of the primary tumor collected 14 days after inoculation of B16-BL6 cells in each group (data not shown).

Figure 1. Effect of intraperitoneal CSE administration on survival time of spontaneous metastasis model mice

\*\( p < 0.05 \) vs control (\( n = 5-7 \)).

Figure 2. Effect of CSE exposure on the (a) invasion and (b) proliferation of B16-BL6 cells.

\*\( p < 0.05 \) vs control ((a) \( n = 3 \) and (b) \( n = 6 \), respectively).
3.2. Effect of CSE on invasion and proliferation of B16-BL6 cells

Invasiveness was significantly increased in the 0.1% CSE exposure group compared to the control (Fig. 2a). On the other hand, the cell proliferation rate was not affected by CSE exposure (Fig. 2b).

3.3. Effect of MVK, ACR, and CA on B16-BL6 cell invasion

CA increased invasiveness of B16-BL6 cells about 4-fold compared to the control, though ACR and MVK did not show a significant effect on the invasiveness of B16-BL6 cells (Fig. 3).

3.4. Effect of MVK, ACR, and CA on B16-BL6 cell proliferation

The exposure of 10 μM of MVK or ACR significantly suppressed the proliferation of B16-BL6 cells, whereas 3 μM or under of MVK and ACR did not affect the proliferation rate of B16-BL6 cells, and CA did not inhibit proliferation regardless of exposure concentration (Fig. 4).

4. Discussions

In this study, we assessed the effect of intraperitoneal administration of CSE on tumor metastasis using a spontaneous cancer
metastasis model in which B16-BL6 cells are disseminated subcutaneously in the footpad as primary cancer. As a result, as Hatai et al. showed in a study on intraperitoneal administration of CSE to transsplenic liver metastasis model mice using colon-26 cells, CSE administration could not prolong the survival time of mice [22], but rather significantly shortened the survival of C57BL/6NCr mice when 100% CSE was used. Since Hatai et al. have reported that intraperitoneal administration of 100% CSE was not toxic to non-cancer-planted mice [22], the shortening of survival time by 100% CSE administration confirmed in this study was considered to be the result of CSE affecting the proliferation and metastasis of B16-BL6 cells. In addition, because the primary lesion was resected 14 days after cancer cell dissemination and the thickness or weight of the primary lesion did not change at that point, the shortened survival time was considered to be caused by the effect of CSE on cancer metastasis rather than the primary lesion. So, we assessed the effect of CSE and its major components on the invasion and proliferation of cancer cells to find the cause of the shortened survival time. First, we analyzed the effect of CSE on cancer cell proliferation and infiltration. As a result, although CSE exposure did not affect cancer cell proliferation as Hatai reported, CSE promoted cancer cell invasion, which was different from the report by Hatai et al. These results suggested that the decrease in survival time by CSE may be due to an increase in invasion ability of B16-BL6 cells.

Next, to clarify the causative agent of this invasiveness promotion, we assessed the effects of MVK, ACR, and CA, which are α, β-unsaturated ketones and aldehydes and considered as main components of CSE, on the invasion of B16-BL6 cells. As a result, while those substances did not affect the proliferation of cancer cells, the invasion ability of B16-BL6 cells was significantly increased by CA. From these results, it was considered that CA was involved in the decrease in survival time of CSE-administered mice through the promotion of B16-BL6 cell invasion. Although CA has been reported to have carcinogenicity and lung cell injury [23, 24], its impact on the invasion ability of tumors has not been reported. To the best of our knowledge, this is the first report that CA has significantly improved the infiltration capacity of cancer cells.

Despite preparing CSE by a method shown in Hatai’s report, CSE enhanced cancer cell invasion in our study, while CSE showed inhibitory efficacy on cancer invasion in Hatai’s report [22]. As the reason for this difference, it was considered to be a large difference in CA concentration in CSE in addition to the difference in cell lines. Slight changes in conditions when extracting CSE or a difference in the lot of tobacco to use may cause a difference in the components contained in tobacco smoke. We need to examine in detail the factor affecting CA content in smoke and differences in CA content in smoke between lots of tobacco. In any case, in order to assess the tumor-suppressive efficacy of CSE in detail, it was considered important to remove CA, which
has an enhancing effect on cancer invasion ability, not only nicotine and tar.

Although CSE suppressed metastasis of cancer cells in the previous study, CSE could not suppress metastasis in our study, and the cause of the discrepancy in these results is thought to be due to the difference in the level of CSE exposure to cancer cells. **In vitro** study about the effect of tobacco smoke, CSE was exposed directly to cancer cells, and cancer cells are exposed to CSE with high level. Similarly, in the study of Hatai et al., intraperitoneal administration of CSE could expose the spleen, where cancer cells first engrafted in the model, and cancer cells to high levels of CSE. And this high level of CSE exposure can have a high metastasis-suppressing effect. Bourgeois et al. reported that cellular response to CSE exposure is dependent not only on the nominal concentration of CSE, but also on specific experimental variables, including the total cell number, and the volume of CSE solution used [25]. They also reported that the effective dose of CSE is more accurately related to the amount of bioavailable chemicals per cell. Similarly, Lee et al. reported that CSE-induced cytotoxicity was reduced at high cell densities [26]. Based on these reports, it is considered that the metastasis of cancer cells was not suppressed in this study because the transfer of CSE to the subcutaneous footpad where cancer cells engrafted was insufficient. In this study, we planned to weigh the lung, which is the site of metastasis, to evaluate the effect of CSE on metastasis. However, because of severe damage to the lung due to cancer metastasis, lung tissue could not be collected, and the cancer metastasis suppressing effect of CSE could not be evaluated in detail. In addition, since CA could not be removed from CSE, the effect of CA-removed CSE on tumor could not be assessed. We need to investigate in detail whether exposure to CA-removed CSE can suppress cancer.

**5. Conclusion**

In conclusion, intraperitoneal administration of CSE significantly shortened the survival time of spontaneous lung metastasis model C57BL/6NCr mice seeded with B16-BL6 cells. As the cause, we found that CA contained in CSE may enhance the invasion ability of cancer cells. To the best of our knowledge, this is the first report that CA has significantly improved the infiltration capacity of cancer cells. In the future, to clarify the cancer-suppressing effect of tobacco components, it was considered important to evaluate the cancer suppression effect of CA-removed CSE in detail and to establish a delivery system, which can transfer the components in CSE to cancer tissues efficiently.

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