Advanced malignant melanoma responds to *Prunus mume* Sieb. Et Zucc (Ume) extract: Case report and *in vitro* study

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**Abstract.** Malignant melanoma (MM) is an aggressive chemoresistant skin cancer characterized by rapid metastasis and a poor prognosis. Therefore, the development of innovative effective therapies is critical. MK615 is an extract from the Japanese apricot *Prunus mume* Sieb. Et Zucc. At a neutral pH, it contains natural chemical substances such as triterpenoids that exert anti-neoplastic effects in several types of cancers. We found that in patients with advanced MM, MK615 dramatically suppressed the cutaneous in-transit metastasis of the disease. Pre- and post-treatment comparison of tumors showed that the apoptotic index was significantly increased by MK615. *In vitro* studies, MTT assay, flow cytometric cell cycle analysis and immunofluorescence microscopy revealed that MK615 inhibited the growth of SK-MEL28 cells in a dose-dependent manner, increased the proportion of cells in sub-G1 phase and induced apoptosis. We further examined the expression of the receptor for advanced glycation end products (RAGE). RAGE is a multi-ligand receptor that binds to a novel cytokine, high mobility group box protein 1 (HMGB1), as well as advanced glycation end products. There is evidence that RAGE/HMGB1 interactions enhance cell invasion in MM. Here, we present Western blotting and immunofluorescence microscopy data indicating that MK615 inhibited the expression of RAGE in SK-MEL28 cells, and suppressed the release of HMGB1 by SK-MEL28 cells. Our findings suggest that MK615 may be a valuable tool for treating MM and other malignant tumors.

**Introduction**

Malignant melanoma (MM) is a highly invasive fatal skin cancer arising from the abnormal proliferation of epidermal melanocytes (1). Its resistance to conventional cancer therapies including irradiation and chemotherapy (2) renders its treatment difficult. MK615, an extract mixture of the Japanese apricot *Prunus mume* Sieb., contains several triterpenoids (3), compounds derived from 30-carbon precursors (4). These compounds have been demonstrated to exert anti-neoplastic effects (5,6). The extract of Ume has been reported to exert anticancer effects against gastric (3), breast (7), hepatocellular (8), colon (9) and pancreatic cancers (10). The mechanisms underlying these effects include the induction of apoptosis (3,7) and autophagy (9) and the suppression of Aurora A kinase (8,10) in cancer cells.

In the present study, after obtaining marked results by treating a patient with advanced MM with MK615, we investigated the mechanisms underlying the effect of MK615 using the human MM cell line SK-MEL28.

**Patients and methods**

**Patients.** The patient, a 67-year-old Japanese woman diagnosed with MM, provided prior written informed consent for use of her data in this report.

**Chemicals.** MK615 and Misatol® GL, containing an additive to render it drinkable, were provided by AdaBio Co. Ltd. (Takasaki, Japan).

**TUNEL staining and evaluation of apoptosis in vivo.** Apoptosis in skin cells from the patient was evaluated by...
the TdT-mediated dUTP-biotin nick end-labeling (TUNEL) method using the ApoTag® Plus Peroxidase in situ Apoptosis Detection kit (Chemicon, Temecula, CA) according to the manufacturer’s instructions. Cell nuclei were stripped of proteins by incubation with 20 µg/ml Proteinase K (Invitrogen, Carlsbad, CA). The immune complexes were visualized by incubating skin sections with DAB-nickel (Vector, Burlingame, CA). They were then counterstained with methyl-green. The apoptotic index was estimated by determining the percentage of apoptotic cells under a light microscope at x400 magnification. A minimum of 2,000 cells was counted in tumor sections. Positively stained tumor cells exhibiting the morphological characteristics of apoptosis were identified using standard criteria (11).

**Cell lines.** Human SK-MEL28 melanoma cells were maintained in DMEM medium containing 10% fetal calf serum. The cells were confirmed to be mycoplasma-free.

**Cell growth analysis.** The growth of SK-MEL28 cells in the presence or absence of MK615 was determined by 3-(4,5-di methylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) colorimetric assay.

**Cell cycle analysis.** The cell cycle was analyzed by the propidium iodide (PI) method of Biswas et al (12). SK-MEL28 cells (1x10^5/ml) were grown for 24 h in 60-mm culture dishes containing 0, 10 or 20 µl/ml MK615, scraped off, and centrifuged at 1,500 rpm for 10 min. The cell pellets were fixed in 70% ethanol at −20°C for 1 h, incubated in 0.25 mg/ml ribonuclease A in PBS at room temperature, stained with PI (0.125 mg/ml in PBS) for 30 min, and subjected to flow cytometric analysis. Cells in the sub-G1 phase were considered to be apoptotic.

**Determination of apoptosis in vitro by fluorescence microscopy.** We determined apoptosis in vitro by immunofluorescence microscopy (13). Briefly, SK-MEL28 cells (5x10^4/well) in 4-well culture dishes were treated with 0, 10 or 20 µl/ml MK615 for 2, 4 and 8 h and incubated with Annexin V-fluorescein isothiocyanate (FITC) (Immunotech, France) according to the manufacturer’s instructions. Cell nuclei were stained with 4,6-diamidino-2-phenylindole (DAPI; Dojindo, Kumamoto, Japan) and examined under an Axioskop microscope (Carl Zeiss, Oberkochen, Germany).

**Western blotting.** Cell lysate samples were subjected to 12% SDS-polyacrylamide gel electrophoresis (PAGE), and the separated proteins were then transferred to a nitrocellulose membrane (GE Healthcare Bio-sciences KK, Piscataway, NJ) as described previously (14). The membrane was blocked with 5% non-fat dry milk in Tris-buffered saline (TBS; pH 7.4) containing 0.02% Tween-20 (TBST) for 1 h at room temperature and then incubated with the first antibody in TBST containing 1% non-fat dry milk (overnight at 4°C). After washing, the membrane was incubated with horseradish peroxidase (HRP)-conjugated anti-rabbit IgG polyclonal antibody (Invitrogen) diluted 1:3000 in TBST containing 2.5% non-fat dry milk (1 h at room temperature). The membrane was washed again, and the immunoreactive bands were visualized using the ECL detection system (GE Healthcare Bio-sciences KK).

**Immunofluorescence microscopy.** SK-MEL28 cells on culture slides were fixed in 4% paraformaldehyde and blocked with 1% bovine serum albumin (Sigma, St. Louis, MO) in PBS. After washing, the cells were incubated with RAGE antibody (Santa Cruz, CA) and washed. The slides were then incubated with Alexa-fluor 488 goat anti-rabbit IgG (Invitrogen) diluted 1:200, washed, and stained with DAPI. Cells were visualized under an Axioskop microscope.

**Statistical analysis.** Statistical analysis was carried out using the Student’s t-test (15); p-values <0.05 were regarded as significant.

**Results**

**Improvement of clinical manifestations in the MK615-treated MM patient.** This 67-year-old Japanese woman presented with a black nodule and a freckle on her left sole. Based on clinical examinations a diagnosis of MM was made (Fig. 1a). We resected the primary cutaneous lesions and dissected the popliteal, inguinal and iliac lymph nodes. Multiple nodules appeared on her left thigh 15 months later (Fig. 1b). Histological examination of biopsy specimens disclosed intra-dermal tumor nests consisting of highly atypical cells positive for S100 protein, HMB45 and Melan-A (Fig. 1c). Positron emission tomography (PET) showed highly increased glucose uptake by multiple lesions on the left thigh (Fig. 1d). The lesions were diagnosed as cutaneous in-transit metastasis of MM, and although we delivered 3x10^8 units of IFN-β intravenously daily, the number of in-transit metastatic lesions gradually increased. After 4 months, we administered 13 g Misatol® GL perorally daily. After 5 months, the number of in-transit metastatic lesions was significantly decreased (Fig. 2a), and de-pigmentation was observed on some of the regressed tumors. On PET scans, the lesions on her left thigh were strikingly improved after MK615 administration (Fig. 2b). The apoptotic index determined by the TUNEL technique was clearly higher after MK615 treatment (0.462±0.203 vs. 0.165±0.039%; p=0.00568).

**Inhibition of SK-MEL28 cell growth by MK615.** The growth of SK-MEL28 cells in the presence or absence of MK615 was determined by MTT assay. The viability of MK615-treated SK-MEL28 cells was significantly decreased in a dose-dependent manner after a 24- and 48-h incubation (Fig. 3a).

**Cell cycle retardation of SK-MEL28 cells by MK615.** To examine the effects of MK615 on apoptosis we assessed the cell cycle of SK-MEL28 cells by flow cytometry using PI staining. In the presence of 10 and 20 µl/ml MK615, the proportion of sub-G1 cells was increased in a dose-dependent manner (Fig. 3b); 19.97±3.90 and 23.07±2.73% at 10 and 20 µl/ml MK615, respectively, and significantly higher than in cells cultured in the absence of MK615 (p=0.00284 and 0.00041, Table I).

**MK615-induced SK-MEL28 apoptosis.** Fluorescence staining with Annexin V showed that at 10 and 20 µl/ml, MK615
induced apoptosis as reflected by phosphatidylserine externalization. The number of apoptotic cells increased in a time-dependent manner (Fig. 3c).

Attenuation of RAGE expression in SK-MEL28 cells by MK615. Western blotting showed that MK615 inhibited RAGE expression by SK-MEL28 cells in a dose-dependent manner (Fig. 4a). Immunofluorescence analysis revealed that at 10 µl/ml, MK615 abolished the expression of RAGE on the cell membrane (Fig. 4b).

MK615 inhibits HMGB1 release by SK-MEL28 cells. The concentration of HMGB1 in SK-MEL28 cell supernatants is induced apoptosis as reflected by phosphatidylserine externalization. The number of apoptotic cells increased in a time-dependent manner (Fig. 3c).

Attenuation of RAGE expression in SK-MEL28 cells by MK615. Western blotting showed that MK615 inhibited RAGE expression by SK-MEL28 cells in a dose-dependent manner (Fig. 4a). Immunofluorescence analysis revealed that at 10 µl/ml, MK615 abolished the expression of RAGE on the cell membrane (Fig. 4b).

MK615 inhibits HMGB1 release by SK-MEL28 cells. The concentration of HMGB1 in SK-MEL28 cell supernatants is

<table>
<thead>
<tr>
<th>MK615 (µl/ml)</th>
<th>sub-G1 (%)</th>
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<tr>
<td>0</td>
<td>1.80±0.50a</td>
</tr>
<tr>
<td>10</td>
<td>19.97±3.90b</td>
</tr>
<tr>
<td>20</td>
<td>23.07±2.73c</td>
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Each value represents the mean ± SD of 3 independent experiments. Statistical significance was determined with the Student's t-test. a vs b, P=0.00284; a vs c, P=0.00041; b vs c, P=0.409.
shown in Fig. 5. The SK-MEL28 cells spontaneously released 4.2±0.6 pg/ml HMGB1. MK615 significantly suppressed HMGB1 release in a dose-dependent manner.

Discussion

The Japanese apricot Ume has been accepted for centuries as a traditional medicine in Japan. It contains several chemical substances, e.g. citric and malic acid, cyanogenic glycosides and triterpenoids (3). Triterpenoids are natural or synthetic compounds derived from 30-carbon precursors (4). They have been shown to exert anti-neoplastic effects (5,7). MK615 is a neutral pH extract mixture of the Ume fruit that contains several triterpenoids. It has been reported to exert anti-cancer effects against gastric (3), breast (7), hepatocellular (8), colon (9) and pancreatic cancer (10).

Here, we demonstrated that the Ume extract MK615 at a neutral pH suppressed cutaneous and in-transit metastatic lesions in a patient with MM. Pre- and post-treatment comparison showed that the apoptotic index of the tumor
cells was significantly increased by MK615. We attributed the de-pigmentation of the regressed tumor portions to the activation of T-cell immunity against MM and melanocytes (16). Triterpenoids have been reported to regulate the cell cycle. Sun et al (17) showed that, in HeLa cells, triterpenoids induced apoptosis by cell cycle deviation resulting in S-phase accumulation and a decrease in G0-G1 phase cells. Other researchers (6) have documented triterpenoid-induced G2-M phase arrest, suggesting that cell cycle deviation plays a key role in the induction of apoptosis. Gutterman et al (5) demonstrated that, in pancreatic cancer cell lines, Ume extract increased the proportion of cells in the G2-M phase and inhibited Aurora A and B kinases. Aurora kinases are key mediators of cell deviation by controlling chromatid segregation (18-20). In our study, the exposure of SK-MEL28 melanoma cells to MK615 resulted in an increase in cells in the sub-G1 phase. Moreover, we detected Annexin V-positive apoptotic cells 2 h after the start of MK615 exposure, and their number continued to increase up to 8 h. These data suggest that MK615 induces apoptosis in MM cells by another pathway.

RAGE, a multi-ligand receptor, is highly expressed and is associated with cell invasion and migration (21). In a primary tumor model, anti-RAGE and soluble RAGE partially inhibited local tumor growth (21). Therefore, RAGE blockade may contribute to reducing tumor growth and invasion.

HMGB1, the ligand of RAGE, has two distinct functions in cellular systems. In the nucleus, HMGB1 acts as an intracellular regulator of the transcription process and plays a crucial role in the maintenance of DNA functions (22). In the extracellular space, HMGB1 is released by all eukaryotic cells upon necrosis or by macrophages in response to inflammatory stimuli such as endotoxin. Extracellular HMGB1 plays a major role in several diseases such as sepsis (22), rheumatoid arthritis, disseminated intravascular coagulation, periodontitis, xenotransplantation and atherosclerosis (13,22-25).

Extracellular matrix associated HMGB1 promotes neurite outgrowth and provides a surface for the assembly of protease complexes in the fibrinolytic system, which can contribute to cell mobility. In view of the increased expression of HMGB1 and RAGE in tumors, the tumor bed may be an ideal locus for the effects of RAGE/HMGB1 interactions on cell migration and invasion.

Our study demonstrated that MK615 decreased the expression of RAGE in SK-MEL28 cells. Moreover, it significantly suppressed the release of HMGB1 by SK-MEL28 cells. Our results suggest that MK615 inhibits cell migration and invasion.

In summary, MK615 was effective in a patient with advanced MM. Our in vitro studies showed that it inhibited the growth of MM cells and increased the proportion of apoptotic cancer cells. MK615 may be a valuable tool for treating MM and other malignant tumors and further studies to evaluate its clinical effectiveness and to elucidate the precise mechanisms of its actions are warranted.

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References


