**Bauhinia purpurea agglutinin**-modified liposomes for human prostate cancer treatment

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**Key words**

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*Lectins are generally defined as proteins that possess the potential to selectively recognize and strongly bind to specific sugar chains of glycoproteins and glycolipids, and the expression of lectins is widely observed in animals, microorganisms and plants. Because interactions between sugar chains and proteins play important roles in not only cell differentiation and proliferation,(1) but also memory formation,(2) the immune system(3) and tumorigenesis,(4) lectins are thought to be one kind of the key molecules that control both physiological and pathological cell signals. Therefore, lectins are often used as detection tools for determining blood type(5) and diagnosing a viral infection(6) or cancer development.(7) In addition, scientific researchers routinely use lectins as biological molecular tools for specific purification, staining and detection of certain proteins.(8,9) As lectins possess the ability to bind to specific sugar chains, they have become available as probes for targeting drugs or nanoparticle drug carriers to diseased organs. In fact, tomato lectin can be used as a probe targeting the lumen of the small intestines,(10) and the application of lectins for use in drug delivery systems (DDS) is ongoing to improve the efficiency of drug delivery to the lungs, brain and other organs for the treatment of various diseases such as cancer.(11,12) Liposomes are lipid-based nanoparticles that are widely used for drug encapsulation and as a DDS, because they have characteristics suitable for clinical use. In particular, liposomal anti-cancer agents are expected to achieve efficient cancer therapy, because liposomes having long blood-circulation potential owing to modification by polyethylene glycol (PEG) effectively accumulate in cancer tissue.(13,14) However, there are considerable problems with the PEG-based modification of liposomes and nanoparticles: for example, decreased interaction of those drug nanocarriers with target cells and impaired intracellular uptake of encapsulated materials into the cells. This PEG dilemma sometimes impairs the therapeutic efficiency of drugs encapsulated in PEGylated nanocarriers. To overcome these disadvantages and to enhance the specific accumulation of nanocarriers, many researchers have applied an active targeting strategy by attaching molecular targeting probes to the top of the PEG chains to improve the efficiency of drug delivery into the target cells for obtaining enhanced therapeutic efficacy.(15) For this reason, identification of targeting molecules that can specifically and strongly recognize the target is desirable to resolve this problem. Boland et al.(16) and Kawa et al.(17) reported that *Bauhinia purpurea agglutinin* (BPA) binds dominantly to cancerous cells in the colon and pancreas, respectively. BPA is a well-known lectin that recognizes galactosyl glycoproteins and glycolipids. In the present study, we firstly found that BPA bound to human prostate cancer specimens but not to normal prostate ones. Therefore, we sought to develop BPA-PEG-modified liposomes (BPA-PEG-LP) encapsulating anticancer drugs for the treatment of prostate cancer. We examined the tumor targetability of BPA-PEG-LP with human prostate cancer DU145 cells, and observed that fluorescently labeled BPA-PEG-LP dominantly associated with the cells via the interaction between liposome-surface BPA and cell-surface galactosyl molecules. We also observed that BPA-PEG-LP accumulated in the prostate cancer tissue after the i.v. injection to DU145 solid cancer-bearing mice, and strongly bound to the cancer cells. In a therapeutic study, DU145 solid cancer-bearing mice were i.v. injected thrice with BPA-PEG-LP encapsulating doxorubicin (BPA-PEG-LPDOX, 2 mg/kg/day as the DOX dosage) or PEG-modified liposomes encapsulating DOX (PEG-LPDOX). As a result, BPA-PEG-LPDOX significantly suppressed the growth of the DU145 cancer cells, whereas PEG-LPDOX at the same dosage as DOX showed little anti-cancer effect. The present study suggested that BPA-PEG-LP could be a useful drug carrier for the treatment of human prostate cancers.

**Materials and Methods**

**Agents.** Dipalmitoylphosphatidylcholine (DPPC), cholesterol and methoxy-polyethylene glycol (2000)-conjugated distearoylphosphatidylethanolamine (DSPE-MPEG) were gifts from..."
Nippon Fine Chemical (Takasago, Hyogo, Japan). DSPE-PEG-NHS (N-hydroxysuccinimide) was purchased from NOF (Tokyo, Japan). 1,1'-Diocadecyl-3,3,3',3'-tetramethylindocarbocyanine perchlorate (DiIC18) was obtained from Life Technologies Japan (Tokyo, Japan). BPA, FITC-conjugated BPA (BPA-FITC) and biotin-conjugated BPA were kindly donated by J-OIL MILLS (Tokyo, Japan).

Animals. BALB/c nu/nu male mice were purchased from Japan SLC (Shizuoka, Japan). All animal experiments were approved by the Animal and Ethics Committee of the University of Shizuoka. The animals were cared for according to the Animal Facility Guidelines of the University of Shizuoka.

Cell culture. Human prostate cancer cell lines DU145 and LNCaP were cultured in RPMI-1640 medium (Wako Pure Chemical Industries, Osaka, Japan) supplemented with 10% heat-inactivated FBS (AusGeneX Pty Ltd, Brisbane, Australia), 100 µg/mL streptomycin (MP Biomedicals, Irvine, CA, USA) and 100 units/mL penicillin (MP Biomedicals) at 37°C in a humidified atmosphere with 5% CO₂.

Histological analysis. A deparaffinized tissue array slide (human normal prostate and cancer tissue; SuperBioChips Laboratories, Seoul, Korea) was blocked with 1% BSA-PBS for 30 min. Then, biotin-conjugated BPA (50 µg/mL) was added to the array slide; and incubation was carried out for 1 h at room temperature for a wash. Then, the slide was next incubated with streptavidin-conjugated HRP (5 µg/mL; Vector Laboratories, Burlingame, CA, USA) for 30 min. To visualize the bound BPA molecules, we incubated the array slide for approximately 5 min in the solution provided in a DAB liquid substrate system (Sigma-Aldrich Co. LLC, St. Louis, MO, USA). The tissues on the array slide were then observed with a light microscope.

Preparation of Bauhinia purpurea agglutinin-PEG-LP. Dipalmitylphosphatidylcholine and cholesterol (2:1 as a molar ratio) dissolved in chloroform were evaporated and dried for at least 1 h under reduced pressure to form a thin lipid film in a round-bottomed flask. The lipids were then hydrated with PBS (pH 7.4), after which the liposome solution was subsequently bottomed flask. The lipids were then hydrated with PBS and lyses with 0.1% SDS buffer (pH 8.4). To determine the binding and uptake of liposomes into the cells, the fluorescence intensity of DiI (Ex. 549 nm; Em. 592 nm) was measured using an Infinite M200 microplate reader (Tecan Japan, Kawasaki, Japan). The detection gains for the fluorimeter were set to obtain the fluorescence intensity automatically by the Tecan i-Control software.

For the binding inhibition assay, DU145 cells were incubated with the liposomes in the presence of glucose or galactose at the concentration of 0.1, 1 or 10 mM; and the fluorescence in the cells was then measured in a similar manner.

Biodistribution analysis. DU145 cells in a Matrigel-containing medium were implanted s.c. (5 × 10⁶ cells/0.2 mL/mouse) into 5-week-old BALB/c nu/nu male mice to prepare solid cancer-bearing mice. [³H]-labeled PEG-LP or BPA-PEG-LP was i.v. injected into the mice via a tail vein; and after distribution of the liposomes for 3 or 24 h, the mice were killed under anesthesia. Blood and organs were then harvested and subjected to further analysis.

For the biodistribution analysis, DU145 cells were cultured in RPMI-1640 medium containing 50% heat-inactivated FBS (AusGeneX Pty Ltd, Brisbane, Australia), 100 µg/mL streptomycin (MP Biomedicals, Irvine, CA, USA) and 100 units/mL penicillin (MP Biomedicals) at 37°C in a humidified atmosphere with 5% CO₂.

Intra-tumor distribution. BPA-PEG-LP or PEG-LP was i.v. injected into DU145-bearing mice via a tail vein; and after 24 h, the mice were killed under anesthesia. Their solid cancers were perfused with an excess amount of PBS in order to remove the circulating liposomes in the blood and then dissected. The cancer tissues were embedded in optimal cutting temperature compound (Sakura, Torrance, CA, USA) and then sectioned in a 10-μm thickness. The sections were stained with DAPI for nuclear staining. Finally, the sections were mounted with PermaFluor Aqueous Mounting Medium (Thermo Fisher Scientific, Cheshire, UK). The localization of liposomes and...
blood vessel cells in the cancer was observed by use of a fluorescence microscope (IX71; Olympus Corporation, Tokyo, Japan).

In vitro anti-proliferative assay. DU145 cells (1.0 × 10⁴ cells/well) were seeded onto a 96-well plate and cultured overnight. DOX-encapsulated PEG-modified liposomes (PEG-LPDOX) or DOX-encapsulated BPA-PEG-LP (BPA-PEG-LPDOX) with the DOX dose at 10 μg/mL was added to the cells; and 3 h later the cells were washed thrice with PBS. Then, they were cultured in fresh medium without liposomes for 48 h. The viable cells were determined by performing a WST-8 assay with a Cell Counting Kit-8 (Dojindo Laboratory, Kumamoto, Japan).

Therapeutic experiment. DU145 cells were implanted s.c. (5 × 10⁵ cells/0.2 mL/mouse) into 5-week-old BALB/c nu/nu male mice, and PEG-LPDOX or BPA-PEG-LPDOX solution (0.2 mL) with a DOX dosage of 2 mg/kg/day was i.v. injected via a tail vein once a week for 3 weeks starting from day 29 after the implantation. The tumor volume and the body weight changes were monitored daily. The tumor volume was calculated according to the following formula: Tumor volume = 0.4 × a × b² (a, largest diameter; b, smallest diameter).

Statistical analysis. One-way ANOVA followed by Tukey honestly significant difference (HSD) post hoc pairwise comparison test was used for multiple group comparisons. In the case of two-group comparisons, Student’s t-test was used.

Results

Selective binding of Bauhinia purpurea agglutinin to human prostate cancer specimens. To demonstrate the potential of BPA to target human prostate cancer, we performed histological analysis by using a tissue array of human prostate cancer specimens (Gleason score = 7–9). As the result, BPA specifically bound to the region of the tissue containing prostate cancer cells, whereas the binding was rarely observed in the normal prostate region (Fig. 1). This result suggests that BPA possessed the potential to bind selectively to human prostate cancer cells.

Characterization of Bauhinia purpurea agglutinin-PEG-LP and Bauhinia purpurea agglutinin-PEG-LPDOX. Bauhinia purpurea agglutinin-PEG-LP and BPA-PEG-LPDOX showed similar particle sizes and ζ-potentials in comparison with those of PEG-LP. PEG-LPDOX and BPA-PEG-LPDOX were 132, 139, 153 and 159 nm, respectively; and the ζ-potentials, −2.1, −4.3, −4.6 and −0.6 mV, respectively. When the stabilities of PEG-LP, PEG-LPDOX and BPA-PEG-LPDOX after 24 h incubation with FBS were examined, the particle sizes of liposomes were not changed and the DOX encapsulation in both PEG-LPDOX and BPA-PEG-LPDOX was kept in over 90% efficiency (data not shown).

Association of Bauhinia purpurea agglutinin-PEG-LP to human prostate cancer cells. To evaluate the specificity of binding of BPA-PEG-LP to prostate cancer cells, we incubated DU145 cells with fluorescently labeled BPA-PEG-LP and then observed the fluorescence in these cells. The results indicated that the association of BPA-PEG-LP to DU145 cells occurred in a time-dependent manner and was significantly higher than that of BPA-free PEG-LP (Fig. 2a). Similar results were obtained in the experiment using LNCaP cells (Fig. 2b).

Biodistribution of Bauhinia purpurea agglutinin-PEG-LP in DU145 cancer-bearing mice. Next, the accumulation of BPA-PEG-LP in the organs after i.v. injection into DU145-bearing mice was examined. As a result, the retention of BPA-PEG-LP in the bloodstream at 3 h after the injection was a little lower than that of PEG-LP; and the other distribution profile was quite similar between them (Fig. 3a). It is notable that BPA-PEG-LP showed a biodistribution similar to that of PEG-LP, suggesting that BPA-PEG-LP had accumulated in the tumor tissue by the
EPR effect, which was a prerequisite for their specific binding to the cancer cells. At 24 h after the injection, whereas the retention of BPA-PEG-LP in the bloodstream was significantly low and the accumulation of BPA-PEG-LP in the liver was higher than that of PEG-LP, the accumulation in the tumor was not much different between BPA-PEG-LP and PEG-LP (Fig. 3b). These results are consistent with the distribution of other cancer-active targeting liposomes: PEG-modification is sufficient to endow liposomes with accumulation in tumors, but intratumoral distribution and the association of liposomes with cancer cells are different between PEG-LP and active targeting probe-conjugated PEG-LP (19).

**Distribution of Bauhinia purpurea agglutinin-PEG-LP in DU145 solid cancers.** In the response to the results of quantitative analysis of liposome accumulation in tumor tissue, we next examined intratumoral distribution of BPA-PEG-LP in the DU145 solid cancers. Fluorescently labeled PEG-LP or BPA-PEG-LP was i.v. injected into the DU145-bearing mice and the distribution of liposomes in the tumor tissues was compared. Immunohistological analysis revealed that higher DiI fluorescence was detected in the tumor sections treated with BPA-PEG-LP than in those treated with PEG-LP (Fig. 4). Moreover, the images which liposomes were associated with the cells were obtained in BPA-PEG-LP-treated mice. These results indicate that the intratumoral distribution of BPA-PEG-LP was obviously different from that of PEG-LP, suggesting that BPA-PEG-LP strongly bound to the prostate cancer cells via the interaction of BPA with target molecules in the tumor tissue.

**Cytotoxicity of Bauhinia purpurea agglutinin-PEG-LPDOX against DU145 cells.** Next, we examined the in vitro cytotoxic effect of BPA-PEG-LPDOX against human prostate cancer cells. As a result, BPA-PEG-LPDOX inhibited the proliferation of DU145 cells at the DOX doses of both 3 and 10 μg/mL, and the inhibition was significantly higher than that obtained with PEG-
To elucidate the cancer therapeutic effect of BPA-PEG-LPDOX, we i.v. injected such liposomes into DU145 solid cancer-bearing mice at the DOX dosage of 2 mg/kg/day via a tail vein once a week three times, and measured the tumor size as an indicator of a side effect of DOX, no remarkable decrease in body weight was observed in any of the mice (data not shown). These results indicate that BPA-PEG-LPDOX possessed a potent therapeutic effect on human prostate cancer cells.

Discussion

Nowadays, the rate of prostate cancer is rising, especially in the developed countries. Corresponding to this situation, the importance of prostate cancer diagnosis has become a matter of focus, because the treatment of prostate cancer in its early stages can significantly improve the survival rate. PSA, a serine proteinase glycoprotein, is a suitable marker for detecting prostate cancer, because the amount of PSA in the bloodstream of the patient begins to increase with the development of prostate cancer.\(^{[20]}\) In addition, it is known that some tumor-associated carbohydrate antigens such as sialyl LewisX (sLeX) can be indicators of tumor malignancy in some types of cancer, including prostate cancer.\(^{[21,22]}\) These findings point to the importance of cell-surface sugar chains in tumor development and the possibility of using them as targets for drug delivery and molecular imaging. In fact, some researchers have used lectins for whole-body imaging of cancer and as an active targeting probe of nanoparticles for drug delivery.\(^{[23]}\)

Our present research aimed at developing an effective prostate cancer therapy by utilizing the interaction between a lectin on the liposomal surface and sugar chains on prostate cancer cells. BPA is a well-known lectin that binds to sugar chains terminating in galactose, such as Gal-GalNac; and of glycoproteins having this terminal sugar, Thomsen–Friedenreich antigen is the most famous.\(^{[24]}\) This antigen is a tumor-associated molecule involved in prostate carcinogenesis; and its expression in prostate cancer cells increases with the development of tumor malignancy.\(^{[25]}\) Pacis et al.\(^{[26]}\) also studied the relationship between the expression of carbohydrates on the surface of prostate cancer cells and tumor malignancy, and reported that the potential of human prostate cancer cells to metastasize to bone increases with an increase in the expression of carbohydrates terminating in galactose. In the present study, we found that BPA strongly bound to human prostate cancer tissue, whereas it did not bind to normal prostate tissue in the same specimen (Fig. 1).

Moreover, we observed that BPA specifically bound to cells of the LNCaP human prostate cancer line and showed that this binding was inhibited in the presence of an excess amount of BPA-PEG-LPDOX at 3 μg/mL of DOX dose (Fig. 5). The result indicates that BPA modification enhanced the anti-cancer effect of PEG-modified liposomal DOX on human prostate cancer cells.

To utilize the binding potential of BPA for the treatment of prostate cancer, we modified the liposomal surface with BPA for delivering anti-cancer drugs to prostate cancer tissue. As the results showed, BPA-PEG-LP of approximately 140 nm in size could be successfully prepared; and the association of BPA-PEG-LP with human prostate cancer cells was significantly higher than that of PEG-LP (Fig. 2a,b). Furthermore, the binding of BPA-PEG-LP to DU145 cells was clearly cancelled by heat-inactivation of BPA or by competitive inhibition.
by an excessive amount of galactose (Fig. 2c,d), suggesting that the binding of BPA-PEG-LP to prostate cancer cells was dependent on the preferential interaction between BPA and the liposomal surface and the cell-surface galactosyl sugar chains.

The biodistribution analysis indicated that BPA-PEG-LP showed a long circulation-time potential, when they were i.v. injected into DU145 solid cancer-bearing mice. Subsequently, BPA-PEG-LP highly accumulated in the solid cancer; and this accumulation was quite similar to that of PEG-LP (Fig. 3); this result was dominantly due to the EPR effect of long-circulating nanoparticles, which was a prerequisite for the specific binding of BPA-PEG-LP to the cancer cells. In fact, our previous study demonstrated that active targeting peptide-conjugated PEG-modified liposomes showed a biodistribution pattern similar to that of PEG-modified liposomes in tumor-bearing mice, although the intratumoral distribution of these liposomes is much different. Thus, to elucidate the difference of liposome distribution in prostate cancer tissue between BPA-PEG-LP and PEG-LP, the liposome localization in DU145 cancer tissues was fluorometrically observed. BPA-PEG-LP was widely distributed in parenchyma of the cancer tissues; and most of the liposomes seemed to bind to or be taken up into the cells (Fig. 4). In contrast, PEG-LP was rarely observed in the cancer tissues. We assumed that this difference in the biodistribution was caused by the ligand-mediated targeting potential of BPA-PEG-LP: BPA-PEG-LP strongly bound to prostate cancer cells via the ligand-receptor interaction and was preferentially incorporated into the cells, whereas PEG-LP was poorly incorporated into the cells because of the shielding effect of PEG on the recognition of these cells by certain proteins and immune cells.

In light of the targetability of BPA-PEG-LP to human prostate cancer cells, we then examined the in vitro cytotoxic effect of BPA-PEG-LPDOX against DU145 cells. BPA-PEG-LPDOX inhibited the proliferation of DU145 cells and this effect of BPA-PEG-LPDOX was higher than that obtained with PEG-LPDOX (Fig. 5). This result suggests that the enhanced cytotoxicity was brought by ligand-mediated targeting of BPA-PEG-LPDOX to prostate cancer cells. There is much evidence that active targeting liposomes such as antibody-modified liposomes are taken up into cells by a receptor-mediated entocytic pathway and that the encapsulated drugs are then released into the cytosol following lysosomal degradation. There are also some reports about lectin-mediated drug delivery into cells. Qaddoumi et al. reported that some lectins were taken up into cells by endocytic pathway after interaction with membrane receptors and Gabor et al. further demonstrated that a kind of lectin, wheat germ agglutinin (WGA), was subject to the proteolytic degradation in lysosome following endocytosis. Thus, we assume that the mechanism by which BPA-PEG-LPDOX was effectively taken up into the cells was endocytosis, which occurred after binding of these modified liposomes to the cell-surface sugar chains. Following endocytosis, the intraliposomal DOX would have been released to exert its cytotoxic action.

Finally, we performed a cancer therapeutic experiment using BPA-PEG-LPDOX. Because liposomal DOX has already been commercialized for the treatment of several cancers all over the world and some candidates of liposomal DOX are now in clinical studies for prostate cancer, we choose DOX as a cytotoxic agent. Three treatments of DU145-bearing mice with BPA-PEG-LPDOX strongly suppressed the tumor growth in these mice; and the tumor volume of the BPA-PEG-LPDOX-treated mice was significantly smaller than that of the PEG-LPDOX-treated mice (Fig. 6). Furthermore, the body-weight decrease as an indicator of side effects was not observed in BPA-PEG-LPDOX-treated mice (data not shown). We strongly believe that this enhanced therapeutic effect was due to the potential of BPA to target prostate cancer and efficient delivery of DOX to the cells.

In the present study, we succeeded in preparing active targeting liposomes bearing the lectin BPA: the liposomes enabled delivery of DOX, an anticancer drug, to prostate cancer cells and suppressed their proliferation. Because BPA could specifically recognize cancer cells in the human prostate but not normal cells in this tissue, BPA could be a potent active targeting probe of nanocarriers for delivering anti-cancer drugs to prostate cancer cells.

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Disclosure Statement

The authors have no conflict of interest to declare.

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cells in the absence or presence of 200 mM galactose on a slide glass, and the cells were then incubated at 37°C for 1 h. The fluorescence was observed under a fluorescence microscope. The green image shows the distribution of the lectin, and blue images show the nuclei stained by DAPI. Scale bars: 100 μm.

Table S1. Binding potential of Bauhinia purpurea agglutinin (BPA) to human cancer tissues. Binding potential of BPA to cancer and normal tissues in each organ are summarized and shown in stages from weak (+) to strong (++++).