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# Phosphatidylserine-Mediated Phagocytosis of Anticancer Drug-Treated Cells by Macrophages<sup>1</sup>

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Apoptotic cells are rapidly phagocytosed and eliminated from the organism. Although cancer cells apoptose when treated with anticancer drugs, how those cells are recognized by phagocytic cells has remained unclear. The human leukemia cell line Jurkat was cultured with doxorubicin or bufalin and induced to undergo apoptosis accompanied by phosphatidylserine externalization. When apoptotic Jurkat cells were mixed with mouse peritoneal macrophages, efficient phagocytosis was observed. Apoptosis and phagocytosis of Jurkat cells were both inhibited by Z-VAD-FMK, and phagocytosis was significantly reduced in the presence of phosphatidylserine-containing liposomes. These results suggest that anticancer drugs induce apoptosis-dependent and phosphatidylserine-mediated phagocytosis in cancer cells.

**Key words:** anticancer drug, apoptosis, phagocytosis, phosphatidylserine.

Apoptotic cells are engulfed by phagocytic cells, and this event has been presumed to prevent damage to surrounding tissues by the noxious contents of dead cells (reviewed in Refs. 1, 2). Another important role of phagocytosis of apoptotic cells has recently been proposed: that is, phagocytic cells present antigens, which apoptotic cells contain, on their surface and stimulate T lymphocytes (3–7). Furthermore, injection of macrophages that had phagocytosed apoptotic tumor cells led to tumor regression in rats (7). These observations strongly suggest that apoptotic cell phagocytosis occurs to eliminate unwanted cells not only directly but also indirectly by activating the immune response. It is important to elucidate the mechanism by which phagocytic cells recognize and phagocytose apoptotic cells in order to apply this phenomenon to the development of novel therapeutics against cancer.

Phagocytes should distinguish target apoptosing cells from living cells to selectively engulf the former. A variety of molecules, such as proteins, sugars, and phospholipids, have been suggested to act as a “phagocytosis marker,” which is exposed on the surface of apoptotic cells and recognized by phagocytes (reviewed in Refs. 8–10). Among them, the membrane phospholipid PS has been most

extensively characterized (reviewed in Ref. 11). In many apoptosing cells, including lymphocytes (12), vascular smooth muscle cells (13), vascular endothelial cells (14), spermatogenic cells (15), and Fas-overexpressing HeLa cells (16), externalized PS serves as a marker to allow phagocytes to recognize target cells. Many anticancer drugs induce apoptosis in cancer cells (reviewed in Refs. 17–19), and this is often accompanied by PS externalization (20). Apoptotic cells with externalized PS are, however, not necessarily phagocytosed in the PS-mediated manner (21, 22). We investigated here how anticancer drug-treated apoptotic cancer cells are phagocytosed by macrophages.

## MATERIALS AND METHODS

**Cell Culture and Apoptosis Analysis**—Jurkat cells, a human leukemia T-cell line, were grown in RPMI 1640 (Nissui Pharmaceutical, Tokyo) containing 10% fetal bovine serum at 37°C in a humidified atmosphere with 5% CO<sub>2</sub> in air. The cells were treated with either doxorubicin (Sigma, MO) (0.3 µg/ml) or bufalin (provided by K. Nakaya) (10<sup>−7</sup> M). Cell viability and chromatin condensation were analyzed under a microscope after staining cells with trypan blue or Hoechst 33342, respectively. PS externalization was determined by flow cytometry (EPICS-XL; Coulter, FL) as described previously (23, 24). Briefly, test cells were treated with FITC-labeled annexin V (Bender MedSystems, Vienna, Austria) and propidium iodide, and cells that were less intensely stained with propidium iodide were gated and analyzed for the binding of FITC-annexin V. For confirmation of cell surface binding of FITC-annexin V, cells mixed with the fluorescent dyes were further treated with trypan blue (2 mg/ml) at pH 4.5 prior to flow cytometry. Trypan blue, a plasma membrane nonpermeable dye, binds to FITC and diminishes its fluorescence because of energy transfer (quenching) (25). To inhibit apoptosis, the caspase inhibitor Z-VAD-FMK

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Abbreviations: FITC, fluorescein isothiocyanate; PBS, phosphate-buffered saline; PC, phosphatidylcholine; PS, phosphatidylserine; Z-VAD-FMK, benzyloxycarbonyl-Val-Ala-Asp (OMe) fluoromethylketone.

(26) (Peptide Institute, Osaka) was added to cultures 1 h before anticancer drug treatment.

**Macrophage Preparation and Phagocytosis Assay**—Macrophages were prepared from peritoneal fluids of thioglycollate-inoculated BDF<sub>1</sub> mice (female, 8 weeks old) as described previously (16) and cultivated on cover slips (15 mm diameter) in RPMI 1640 containing 10% fetal bovine serum at 37°C until use. The phagocytosis assay was performed essentially as described (15). Briefly, target cells were labeled with biotin (NHS-LS-Biotin; Pierce, Rockford, IL), mixed with macrophages (at a ratio of 10 target cells to one macrophage), and incubated at 37°C for 2–3 h. The mixture was treated with trypsin (0.5 mg/ml) to remove Jurkat cells free from or lightly attaching to macrophages. The remaining cells were further treated with PBS containing paraformaldehyde (2%), glutaraldehyde (0.5%), and Triton X-100 (0.05%), then supplemented with FITC-conjugated avidin (fluorescein-avidin D; Vector Laboratories, Burlingame, CA). To analyze chromatin condensation and nuclear morphology of phagocytosed Jurkat cells, the fixed cells were further treated with propidium iodide, which binds to DNA. The number of macrophages containing engulfed cells was determined under a fluorescence/phase-contrast microscope and expressed relative to that of total macrophages, i.e., as the phagocytic index. The mean and SD of a typical example from at least three independent experiments are presented. For liposome preparations, phospholipids (Avanti Polar Lipids, Alabaster, AL) were dried as films, suspended in PBS, and sonicated. The liposomes were composed of either

PC only (PC-liposomes) or a combination of PC and PS at a molar ratio of 7:3 (PS-liposomes).

## RESULTS

**Apoptosis of Anticancer Drug-Treated Jurkat Cells**—We chose two apoptosis-inducing anticancer drugs, doxorubicin and bufalin. The drugs possess distinct modes of action: doxorubicin intercalates DNA and mainly affects nucleic acid synthesis (27), while bufalin inhibits Na<sup>+</sup>,K<sup>+</sup>-ATPase in the plasma membrane (28). Human leukemic Jurkat cells were cultured in the presence of these drugs, and cell viability and chromatin condensation were examined (Fig. 1A). Both drugs showed similar effects: cell viability decreased (left panel) and the number of cells with condensed chromatin increased (right panel) as the time of drug treatment was prolonged. Chromatin condensation caused by either drug was significantly inhibited in the presence of a caspase inhibitor, Z-VAD-FMK (Fig. 1B). These results indicated that doxorubicin and bufalin induce apoptosis in Jurkat cells.

We next examined whether this apoptosis is accompanied by PS externalization. To do so, drug-treated Jurkat cells were stained with FITC-annexin V and propidium iodide, and analyzed by flow cytometry. When the cells less intensely stained with propidium iodide were examined for the binding of FITC-annexin V, an extra peak with more fluorescence appeared after treatment with either drug, although the size of the peaks was different (left panels in Fig. 2A). Such peaks completely disappeared when the cells

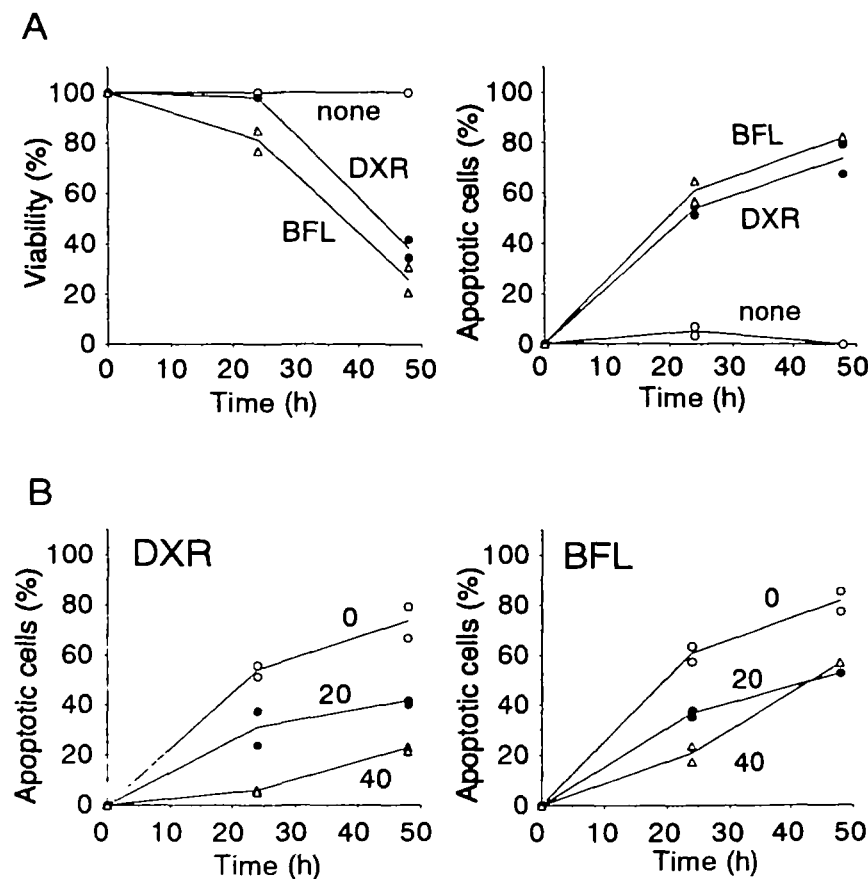
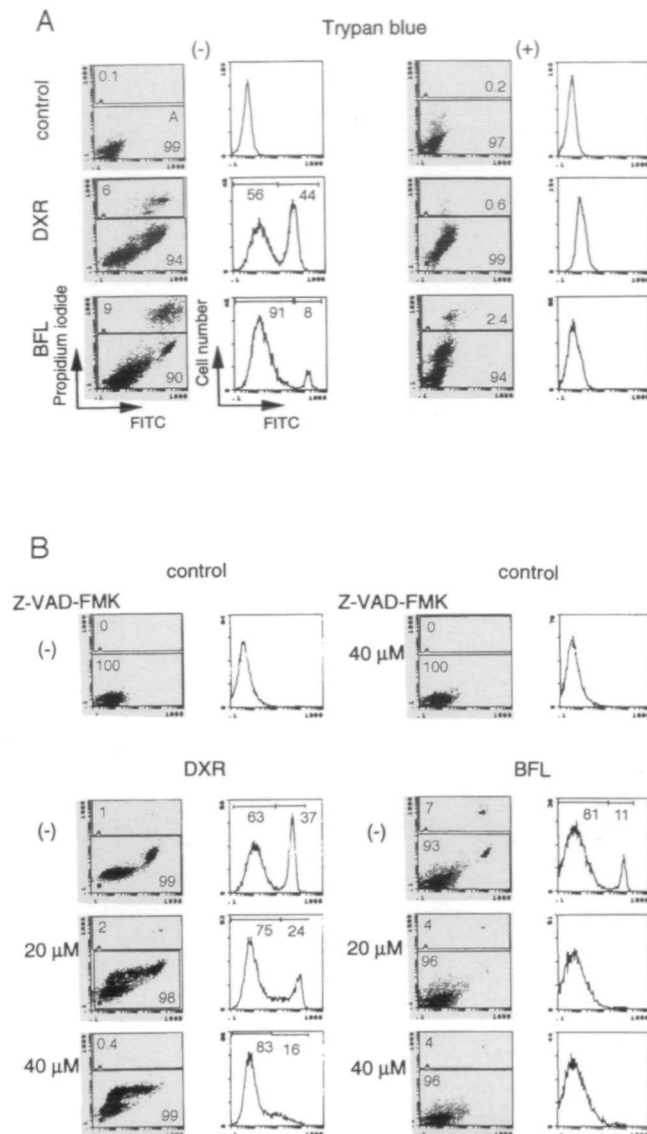


Fig. 1. Anticancer drug-induced apoptosis of Jurkat cells. (A) Jurkat cells treated with doxorubicin (DXR) or bufalin (BFL) for the indicated periods were analyzed for cell viability (trypan blue staining) (left) and occurrence of apoptosis (chromatin condensation) (right). (B) Jurkat cells were incubated with Z-VAD-FMK (20 or 40  $\mu$ M) for 1 h prior to the treatment with doxorubicin (left) or bufalin (right), and the extent of apoptosis was determined.



**Fig. 2. PS externalization in anticancer drug-treated Jurkat cells.** (A) Jurkat cells that had been treated with doxorubicin or bufalin for 24 h were analyzed for PS externalization by flow cytometry in the absence (-) or presence (+) of trypan blue. Cells less intensely stained with propidium iodide (zone A in the left-most panel) were analyzed for the binding of annexin V (histograms at the right). The numbers indicate the percentages of cells in the corresponding areas. (B) The same assay was conducted with cells that were cultured in the presence of Z-VAD-FMK (20 or 40  $\mu$ M).

were treated with trypan blue to quench fluorescence from the cell surface (right panels), indicating that cells corresponding to the extra peaks contained surface-bound FITC-annexin V. Moreover, the addition of Z-VAD-FMK inhibited the appearance of such peaks (Fig. 2B). From these results, we concluded that PS, which is normally restricted to the cytoplasmic side of living cells, was externalized and exposed on the surface of apoptotic Jurkat cells upon treatment with either doxorubicin or bufalin.

**Phagocytosis of Anticancer Drug-Treated Jurkat Cells by Mouse Peritoneal Macrophages**—Jurkat cells that had been cultured with the drugs for 24 h were labeled with biotin and mixed with mouse peritoneal macrophages. The

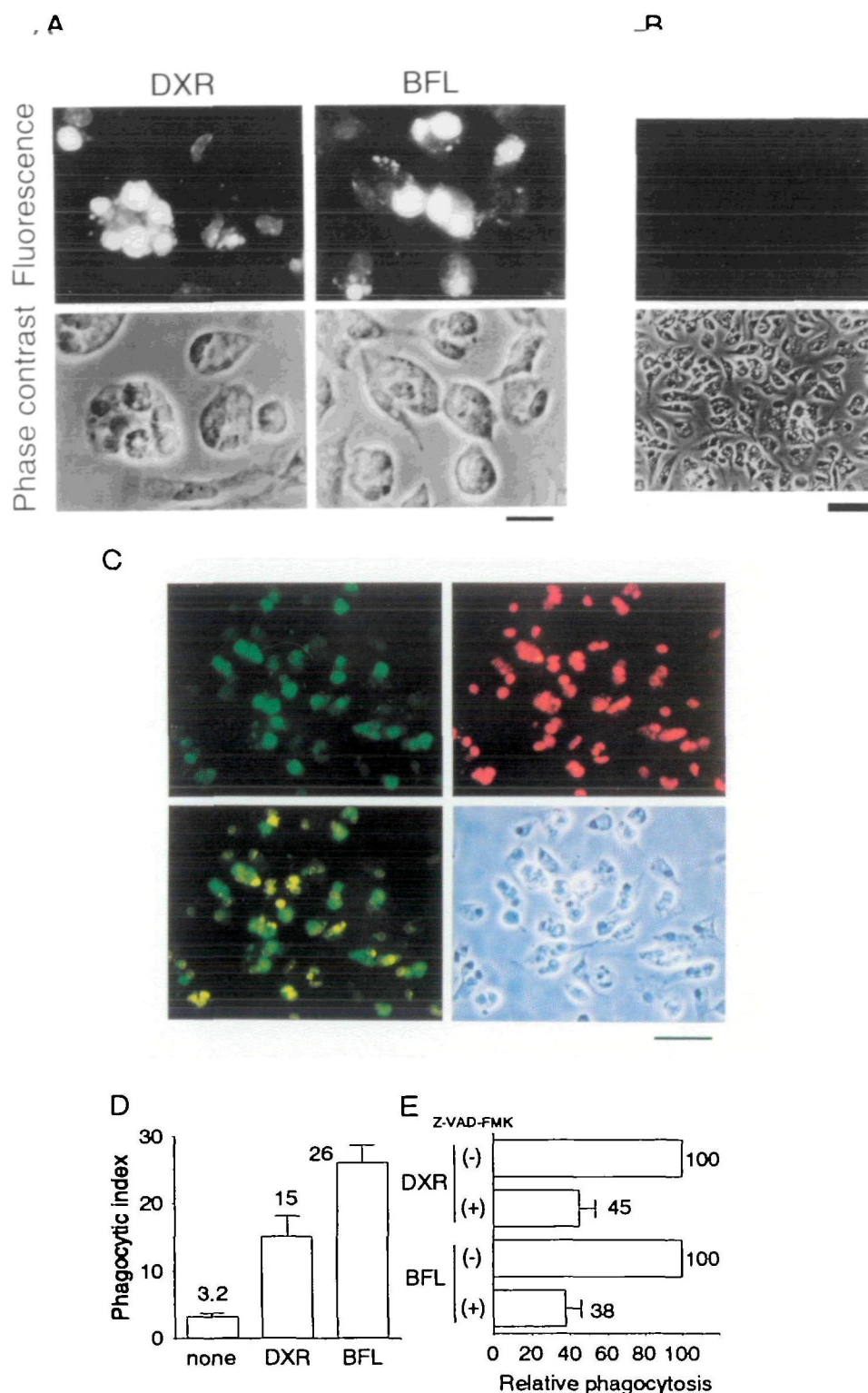
mixture was incubated for 2–3 h, washed, permeabilized, and supplemented with FITC-avidin. When the reacted cells were examined under a fluorescence/phase-contrast microscope, many fluorescent particles were detectable in the cytoplasm of macrophages (Fig. 3A). Jurkat cells left untreated with the drugs were not significantly engulfed by macrophages (Fig. 3B). There was no significant difference between the two drugs. When the macrophages that reacted with doxorubicin-treated Jurkat cells were further stained with propidium iodide, many phagocytosed Jurkat cells whose nuclei were not intensely stained or fragmented were observed (Fig. 3C). A similar result was obtained with bufalin-treated Jurkat cells (data not shown). These results indicated that macrophages phagocytosed drug-treated Jurkat cells before their chromatin condensed and nuclei were fragmented. Quantification of the phagocytosis reaction revealed that Jurkat cells treated with either drug became susceptible to phagocytosis by macrophages (Fig. 3D). Phagocytosis was significantly reduced when Jurkat cells were treated with the anticancer drugs in the presence of Z-VAD-FMK (Fig. 3E), indicating that phagocytosis was dependent on apoptosis. The above results all showed that Jurkat cells treated with either doxorubicin or bufalin were phagocytosed by peritoneal macrophages at early stages of apoptosis.

**Involvement of PS in Phagocytosis of Anticancer Drug-Treated Cells**—Since externalized PS often serves as a marker in phagocytosis of apoptotic cells (11), we next asked if this is the case with anticancer drug-treated Jurkat cells, by conducting the phagocytosis assay in the presence of liposomes. Liposomes containing PS and PC significantly inhibited phagocytosis of Jurkat cells treated with either drug, whereas those composed of only PC showed a minimal effect (Fig. 4). These results suggested that phagocytosis of the drug-treated Jurkat cells by macrophages was mediated, at least partly, by PS exposed on the cell surface.

Significant levels of phagocytosis, however, always occurred in the presence of a maximal amount of PS-liposomes, suggesting the existence of PS-independent phagocytosis. We thus investigated the involvement of other possible phagocytosis markers (Fig. 5). A synthetic peptide containing the integrin-binding sequence RGDS; fucoidan and dextran sulfate, specific inhibitors of class A scavenger receptor types I and II; and *N*-acetyl-D-glucosamine, a ligand for lectin-like receptors, all showed some inhibition, but were much less effective than PS-liposomes. Furthermore, simultaneous addition of PS-liposomes and fucoidan or dextran sulfate gave no additive effect. Similar results were obtained in the phagocytosis assays of Jurkat cells treated with either doxorubicin or bufalin. These results suggest that as-yet unidentified molecules, in addition to PS, are involved in the recognition by macrophages of apoptotic Jurkat cells.

## DISCUSSION

Although anticancer drugs induce apoptosis that is accompanied by externalization of PS in many cancer cell lines (20), it has not yet been shown whether those cells undergo PS-mediated phagocytosis. The present study showed that macrophages phagocytosed Jurkat cells, which were undergoing apoptosis induced by treatment with either doxorubicin or bufalin, in a manner inhibitable by PS-containing



**Fig. 3. Phagocytosis of anticancer drug-treated Jurkat cells by mouse peritoneal macrophages.** (A) The mixture of Jurkat cells and macrophages after the phagocytosis reaction was examined under a fluorescence/phase-contrast microscope. Bar=10  $\mu$ m. (B) Jurkat cells untreated with the drugs were subjected to the phagocytosis assay. Bar=50  $\mu$ m. (C) Nuclear morphology of engulfed Jurkat cells. Jurkat cells treated with doxorubicin were subjected to the phagocytosis assay using FITC-avidin, and the reactive cells were further stained with propidium iodide. Top left, FITC signal from the surface of phagocytosed Jurkat cells; top right, propidium iodide signal from DNA of macrophages and Jurkat cells; bottom left, superimposition of the two signals; bottom right, phase-contrast view. Bar=50  $\mu$ m. (D) Quantification of phagocytosis reactions. Columns and bars are means  $\pm$  SD. (E) The phagocytosis assay was conducted in the presence of Z-VAD-FMK (40  $\mu$ M). The extent of phagocytosis is expressed relative to that of control reactions, taken as 100. The means of the phagocytic index in control reactions were 15 (DXR) and 26 (BFL).

liposomes. Since these two drugs inhibit the growth of cancer cells in different ways, it is likely that anticancer drugs induce PS-mediated phagocytosis of cancer cells irrespective of their mode of action in cell growth inhibition.

PS exists predominantly in the inner leaflet of the plasma membrane bilayer of living cells, and translocates to the

outer leaflet upon induction of apoptosis (reviewed in Refs. 29, 30). Our results showed that PS externalization and phagocytosis were both inhibited in the presence of an apoptosis inhibitor. Induction of apoptosis and phagocytosis are thus attributed to the same actions of anticancer drugs, at least for the drugs used in this study. Cell growth inhibition and apoptosis induction by anticancer drugs



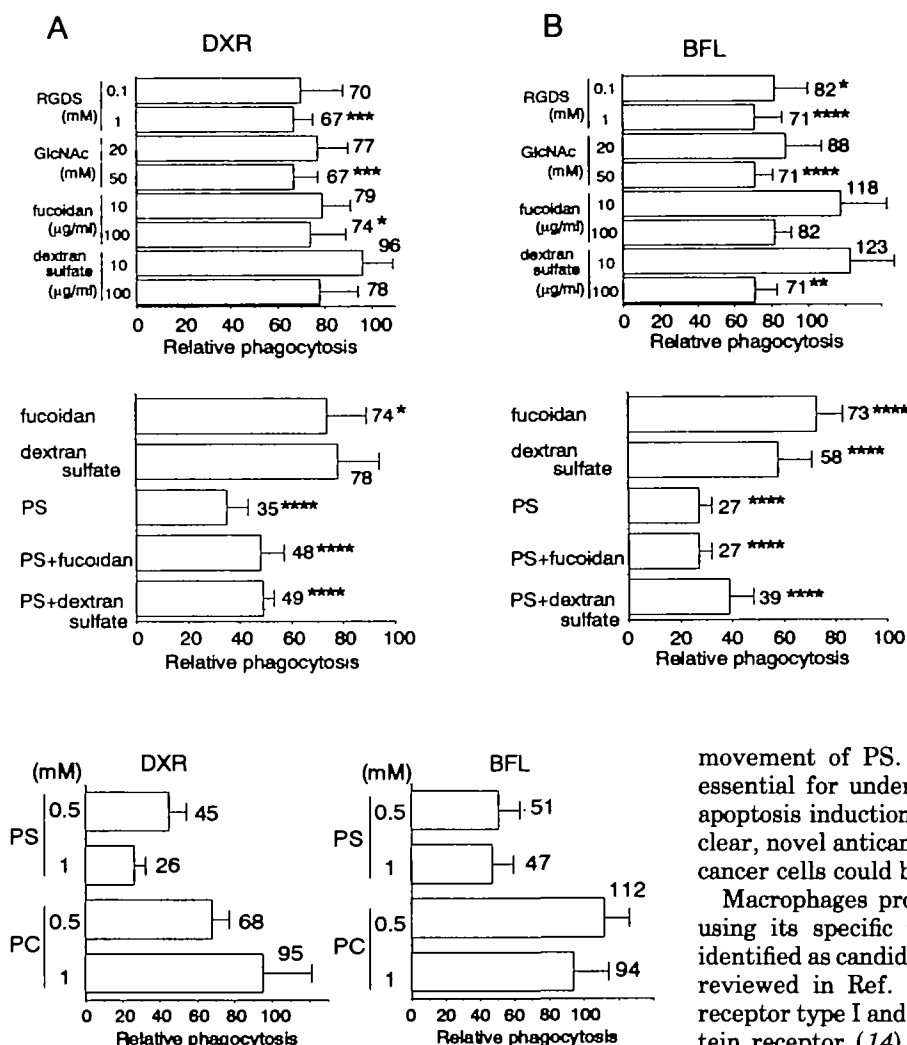


Fig. 4. Effect of liposomes on phagocytosis of anticancer drug-treated Jurkat cells. Phagocytosis reactions were conducted in the presence of PS- or PC-liposomes (0.5 or 1 mM). The extent of phagocytosis is shown relative to that with no added liposomes, which is taken as 100. The means of the phagocytic index in control reactions were 22 (DXR) and 17 (BFL).

might be caused by their distinct activities (31). One could thus modify and improve anticancer drugs simply by removing the former activity, which might cause severe side effects. Moreover, novel drugs may be discovered through screening of chemical and natural products, focusing on apoptosis-inducing activity.

We previously showed that macrophages phagocytose cells that expose PS independently of apoptosis (16). In the present study, drug-treated Jurkat cells were targeted by macrophages before nuclear morphology changed. We speculate that PS externalization by itself, not other apoptotic changes, is sufficient for drug-treated cells to be phagocytosed. It is thus important to elucidate the mechanism by which PS is exposed to the cell surface upon anticancer drug treatment. One suggested mechanism is that changes in the amount and/or activity of enzymes defining localization of the phospholipid lead to its externalization (30). Two enzymes called aminophospholipid translocase (32) and scramblase (33) are believed to control the

Fig. 5. Effect of potent inhibitors on phagocytosis of anticancer drug-treated Jurkat cells. Various substances were added to the phagocytosis assay. The extent of phagocytosis is shown relative to that with no added inhibitor, taken as 100. The means of the phagocytic index in control reactions were 17–30. RGDS, a synthetic peptide containing the integrin-binding sequence RGDS; GlcNAc, *N*-acetyl-D-glucosamine; fucoidan and dextran sulfate, ligands for class A scavenger receptor; PS, PS-liposomes; PC, PC-liposomes. In the bottom two panels, PS-liposomes (0.5 mM), fucoidan (100  $\mu$ g/ml), and dextran sulfate (100  $\mu$ g/ml) were added either alone or in combination. Significance was calculated using Student's *t* test. \**p* < 0.05; \*\**p* < 0.02; \*\*\**p* < 0.002; \*\*\*\**p* < 0.001.

movement of PS. Characterization of these enzymes is essential for understanding how PS is externalized upon apoptosis induction. Once the precise mechanism becomes clear, novel anticancer drugs that induce cell-surface PS in cancer cells could be developed.

Macrophages probably recognize PS on target cells by using its specific receptor. Several proteins have been identified as candidates for the PS receptor (14, 34–41, and reviewed in Ref. 11). Among them, class B scavenger receptor type I and lectin-like oxidized low density lipoprotein receptor (14) were shown to serve as phagocytosis-inducing PS receptors in testicular Sertoli cells (40) and vascular endothelial cells (41), respectively. Although the macrophage PS receptor still remains to be identified, one could modify the quantity and/or quality of such a receptor so that macrophages would phagocytose cancer cells with higher selectivity and efficiency. Further studies are needed to understand how anticancer drug-treated cancer cells are recognized and phagocytosed by phagocytic cells, and such studies should provide us with new approaches for the development of novel therapeutic agents against cancer.

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