



## Review

# The role of phosphatidylserine in recognition of apoptotic cells by phagocytes

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## Abstract

Exposure of phosphatidylserine on the outer leaflet of the plasma membrane is a surface change common to many apoptotic cells. Normally restricted to the inner leaflet, phosphatidylserine appears as a result of decreased aminophospholipid translocase activity and activation of a calcium-dependent scramblase. Phosphatidylserine exposure has several potential biological consequences, one of which is recognition and removal of the apoptotic cell by phagocytes. It is still not clear which receptors mediate PS recognition on apoptotic cells; however, several interesting candidates have been proposed. These include the Class B scavenger and thrombospondin receptor CD36, an oxLDL receptor (CD68), CD14, annexins,  $\beta$ 2 glycoprotein I, gas-6 and a novel activity expressed on macrophages stimulated with digestible particles such as  $\beta$ -glucan. Whether PS is the sole ligand recognized by phagocytes or whether it associated with other molecules to form a complex ligand remains to be determined.

**Keywords:** phosphatidylserine; apoptosis; phagocytosis; receptors; membranes

**Abbreviations:**  $\beta$ 2-GPI: b2-glycoprotein IBMDM: bone marrow-derived macrophages; CL: cardiolipin; GPS: glycerophosphorylserine; HMDM: human monocyte-derived macrophages; PC: phosphatidylcholine; PE: phosphatidylethanolamine; PI: phosphatidylinositol; PS: phosphatidylserine; SR: scavenger receptors; TCR: T cell receptor; TSP: thrombospondin; UV: ultraviolet light; VnR: vitronectin receptor.

Recognition and removal by phagocytes is the final common event in the lives of many apoptotic cells. Apoptotic cells are removed prior to their lysis, suggesting that they express specific surface changes that signal phagocytes to bind and engulf them. These phagocytes can be professional (the macrophages) or amateur (including such cells as fibroblasts,

epithelial cells and vascular smooth muscle cells). To date, there have been a number of receptors described for macrophages and other cells which bind to apoptotic cells and mediate their uptake. These include lectin-like receptors (Duvall *et al*, 1985; Dini *et al*, 1992; 1995; Hall *et al*, 1994; Morris *et al*, 1994; Falasca *et al*, 1996), the vitronectin receptor  $\alpha$ v $\beta$ 3 (Savill *et al*, 1990; Hall *et al*, 1994; Hughes *et al*, 1997), CD36 (Savill *et al*, 1992), an uncharacterized phosphatidylserine-recognizing receptor (Fadok *et al*, 1992a,b, 1993; Pradhan *et al*, 1997), CD14 (Flora and Gregory 1994; Devitt *et al*, 1998), and scavenger receptors (Sambrano and Steinberg, 1995; Fukasawa *et al*, 1996; Platt *et al*, 1996; Murao *et al*, 1997). The ABC1 transporter, also involved in uptake of mammalian apoptotic cells, has recently been shown to mediate anion transport (Luciani and Chimini, 1996; Becq *et al*, 1997). It is homologous to the ced-7 gene associated with phagocytosis of apoptotic bodies in *C. elegans* (Ellis *et al*, 1991). How any of these receptors mediate phagocytosis of apoptotic cells is not known. In addition, the ligands for these receptors are not well characterized; however, exposure of phosphatidylserine on the external leaflet of the plasma cell membrane appears to be common to many apoptotic cells (Fadok *et al*, 1992a,b; Schlegel *et al*, 1993; Mower *et al*, 1994; Koopman *et al*, 1994; Bennet *et al*, 1995; Homburg *et al*, 1995; Martin *et al*, 1995a; Shiratsuchi *et al*, 1997), and this phospholipid appears to be recognized in a stereospecific fashion by subsets of macrophages (Fadok *et al*, 1992a,b, 1993; Pradhan *et al*, 1997), by melanoma cells (Fadok, 1995), by smooth muscle vascular cells (Bennett *et al*, 1995), and by Sertoli cells (Shiratsuchi *et al*, 1997).

Normally, the phospholipids of the plasma membrane are distributed asymmetrically. Choline-containing lipids, such as phosphatidylcholine (PC) and sphingomyelin, are concentrated on the outer leaflet, whereas the aminophospholipids, including phosphatidylethanolamine (PE) and phosphatidylserine (PS), are present in highest abundance on the inner leaflet. Phosphatidylserine, in fact, is restricted exclusively to the inner leaflet in most cells. Loss of phospholipid asymmetry and exposure of phosphatidylserine was first demonstrated for apoptotic lymphocytes (Fadok *et al*, 1992a; Schlegel *et al*, 1993; Mower *et al*, 1994); however, these findings have been confirmed for a number of cell types, including neutrophils, tumor cell lines, smooth muscle vascular cells, spermatogonia, and Jurkat T cells and their cytoplasts (Fadok *et al*, 1992b; Bennet *et al*, 1995; Homburg *et al*, 1995; Martin *et al*, 1995a, 1996; Shiratsuchi *et al*, 1997). Because exposure of phosphatidylserine appears to be a universal feature of apoptotic cells, it is now used as a marker in flow cytometry assays utilizing fluorochrome-labelled annexin V, which binds to PS in a calcium-dependent manner (Vermees *et al*, 1995; Zhang *et al*, 1997).

## Mechanisms for phosphatidylserine exposure in apoptotic cells

The mechanisms for PS exposure on the surface of apoptotic cells appear complex and will remain an active area of investigation for some time. To determine these, an understanding of the maintenance of normal membrane phospholipid asymmetry is required. Two mechanisms have been proposed. A membrane protein termed the aminophospholipid translocase transfers PS and to some extent, PE, from the outer leaflet to the inner leaflet (Devaux, 1991; Zachowski, 1993; Williamson and Schlegel, 1994). This protein appears to be a Mg<sup>2+</sup>-ATPase which is inhibited by Ca<sup>2+</sup> at concentrations achieved in many apoptotic cells (Daleke and Huestis, 1985; Zachowski *et al*, 1986; Bitbol *et al*, 1987; Bevers *et al*, 1989; Morrot *et al*, 1990; Comfurius *et al*, 1990; Auland *et al*, 1994). Tang and coworkers recently cloned a novel P-type ATPase with aminophospholipid translocase activity (Tang *et al*, 1996). The second mechanism involves the tethering of PS in the inner leaflet by its association with membrane skeletal proteins such as fodrin (non-erythrocyte spectrin), based on the known ability of PS to bind to spectrin (Momers *et al*, 1980; Williamson *et al*, 1987; Maksymiw *et al*, 1987; Middlekoop *et al*, 1988). The tethering mechanism fell out of favor for a period of time (see Calvez *et al*, 1988; Devaux, 1991; Williamson and Schlegel, 1994); for example, Pradhan and coworkers labelled cytoskeletal proteins with a photoactivable analogue of phosphatidylethanolamine (2-(2-azido-4-nitrobenzoyl)-1-acyl-sn-glycero-3-phospho [<sup>14</sup>C]ethanolamine) and found no differences in spectrin labelling between normal red cells and PS-expressing red cells from patients with sickle cell anemia (Pradhan *et al*, 1991). However, the tethering hypothesis appears to be enjoying a resurgence of interest, given the observation that fodrin is cleaved during apoptosis. Other potential intracellular tethering molecules for phosphatidylserine could include annexins or polyamines (Chung *et al*, 1985; Meers *et al*, 1986; Bratton, 1994).

Loss of phospholipid asymmetry can be achieved in red cells and platelets by treatment with calcium ionophores. Externalization of PS is rapid, suggesting facilitated transport rather than passive diffusion of lipids across the bilayer. This activity is believed to be mediated by a protein termed a scramblase, which is activated by calcium, is ATP-independent, and is nonspecific with regard to the phospholipids transported (Verhoven *et al*, 1995; Williamson *et al*, 1992, 1995). It is believed to be a protein-mediated flip site for the rapid transbilayer movement of phospholipids. The protein has been purified from red cell and platelet membranes, and the gene has recently been cloned (Basse *et al*, 1996; Comfurius *et al*, 1996; Zhou *et al*, 1997). Abnormalities in the function of this protein *in vivo* mediate the bleeding disorder known as Scott syndrome, in which failure of coagulation results from lack of PS expression on the surface of activated platelets (e.g. see Sims *et al*, 1989; Kojima *et al*, 1994; Stout *et al*, 1997).

Verhoven and coworkers (Verhoven *et al*, 1995) studied PS exposure in the T cell hybridoma DO11.10 which undergoes activation-induced cell death following anti-CD3 treatment. They utilized NBD-labelled phospholipids to

study PS and PC transport during development of apoptosis. They found that prior to the development of nuclear condensation, PS translocation to the inner leaflet was impaired. At the same time, the transport of PC was increased. These data showed for the first time that during apoptosis, aminophospholipid translocase activity was impaired as the scramblase was activated. The net result was PS exposure on the surface of the cell.

We have recently compared the roles of the aminophospholipid translocase and scramblase in three additional cell types known to express PS when undergoing apoptosis: ultraviolet light (UV)-treated HL-60 cells, anti-Fas treated Jurkat T cells, and UV-treated human peripheral blood neutrophils (Bratton *et al*, 1997). Utilizing NBD-PS uptake as a measure of aminophospholipid translocase activity in both bulk uptake assays and flow cytometric assays, we found that translocase activity was impaired in all three cell types. The increase in NBD-PC uptake by all three cell types supported the notion that scramblase was activated. Changes in these activities preceded the expression of PS, as determined by annexin V binding. Because both the aminophospholipid translocase activity and the scramblase are reported to be affected by alterations in calcium levels, we studied the effects of Ca<sup>2+</sup> depletion on PS exposure. Chelation of intracellular Ca<sup>2+</sup> with BAPTA had no effect on PS expression, loss of aminophospholipid translocase activity, or activation of nonspecific phospholipid flip-flop. In contrast, depletion of extracellular Ca<sup>2+</sup> was associated with a marked decrease in annexin V binding (the annexin assay however was done in the presence of Ca<sup>2+</sup>, as annexin binding is calcium-dependent) and loss of nonspecific phospholipid transport by apoptotic cells. There was no alteration in the percentage of apoptotic cells nor was there restoration of aminophospholipid translocase activity, suggesting that loss of aminophospholipid translocase activity was not sufficient for PS expression on apoptotic cells. Restoration of extracellular Ca<sup>2+</sup> levels restored annexin binding and nonspecific phospholipid transport, but the actual levels of PS expressed appeared to be determined by the degree of aminophospholipid translocase impairment. We therefore concluded that nonspecific phospholipid transport (as mediated by the proposed scramblase) is dependent on extracellular Ca<sup>2+</sup>, which is in agreement with the findings of Hampton *et al* (1996) in U937 cells. Activation of the scramblase activity appears to be the driving force for PS expression during apoptosis; however, the activity of the aminophospholipid translocase ultimately modulates the degree of PS expression.

The contribution of membrane skeletal changes to exposure of PS remains to be determined. Fodrin is cleaved early in the development of apoptosis (Martin *et al*, 1995b, 1996; Vanags *et al*, 1996) by calpain or a similar protease (Squier *et al*, 1994; Squier and Cohen, 1997). The relationship of this finding to PS exposure is not known at this time, as the inhibitors used in these studies appear to inhibit the entire apoptotic process rather than just fodrin cleavage. One of the central mechanisms believed to mediate many of the changes of apoptosis is the activation of ICE-like proteases termed caspases. The relationship of

these proteins to apoptosis was first determined in the nematode *C. elegans*, in which the *ced-3* gene, which encodes a caspase, was shown to be essential for cell death (Yuan and Horvitz, 1990; Yuan *et al*, 1993). The role of caspases in the eventual exposure of PS on the surface of the apoptotic cell was first suggested by the findings of Martin *et al* (1996). They found that the caspase inhibitors ZVAD-FK, YVAD-CK and DEVD-FK prevented CPP 32 activation, fodrin cleavage, and PS expression in anti-Fas treated Jurkat T cells and their cytoplasts. In contrast, Schlegel recently reported that caspase inhibitors had no effect on PS expression induced by glucocorticoids in D010.11 T cell hybridoma cells, although they did inhibit PS expression induced by anti-TCR antibodies (Schlegel *et al*, 1997). Vanags *et al* (1996) examined the kinetics of PS exposure, fodrin cleavage, plasma membrane blebbing, and nuclear fragmentation in TNF $\alpha$ -treated U937 cells and anti-Fas-treated Jurkat T cells. They could not determine which event occurred first and they were not able to demonstrate uncoupling of the nuclear events from the extranuclear events. They did observe, however, that all events in U937 cells appeared more sensitive to calpain inhibitors than to caspase inhibitors, whereas Jurkat cells appeared more sensitive to caspase inhibitors.

Our own work with apoptotic human neutrophils (Frasch *et al*, 1998) also supports the notion that regulation of apoptosis and PS expression may be cell and signal specific. Stressful stimuli such as exposure to UV or hyperosmolarity induced DNA fragmentation and PS exposure in neutrophils that was associated with activation of p38 MAPkinase; a p38 MAPkinase inhibitor was found to inhibit nuclear changes of apoptosis (Frasch *et al*, 1998) PS expression (Frasch, unpublished data). Neither anti-Fas nor spontaneous apoptosis in human neutrophils was inhibited by the p38 MAPK inhibitor. In contrast, caspase inhibitors inhibited nuclear apoptosis and PS expression in stress-induced, anti-Fas, and spontaneous neutrophil apoptosis (Frasch *et al*, 1998, and unpublished data). If and how the caspases affect the aminophospholipid translocase or the scramblase remains to be determined; however, Schlegel has shown preliminary evidence that the translocase is not cleaved in apoptotic T cells (Schlegel *et al*, 1997). To summarize then, it appears that caspases are involved in many but not all cells in which PS exposure has been demonstrated; clarification of caspase involvement in PS exposure will require the step-by-step dissection of the pathway between caspase activation and loss of phospholipid asymmetry.

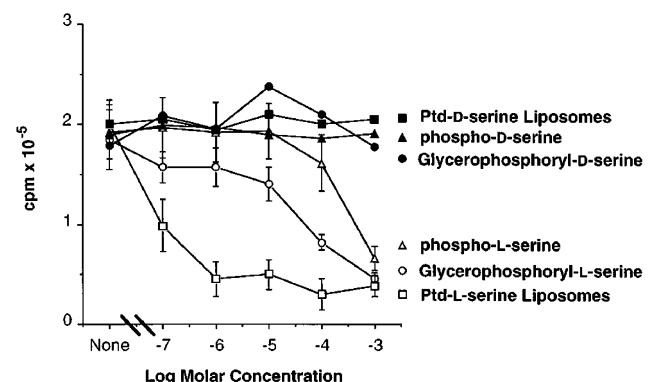
BCL-2 protein, as well as other family members such as BCL-X<sub>L</sub>, are known to inhibit apoptosis in many systems. Their role in regulating membrane changes is not clear. In T and B lymphocytic tumor cell lines, overexpression of BCL-2 appears to inhibit PS expression and prevent recognition by macrophages, as well as inhibit nuclear apoptosis (Martin *et al*, 1995a; Flora *et al*, 1996). Aged neutrophils overexpressing BCL-2 do not show evidence of nuclear apoptosis; however they are still phagocytosed by macrophages (Lagasse and Weissman, 1994). Whether they express PS remains to be determined, but the data

reinforce the notion that membrane changes can be divorced from nuclear changes of apoptosis in some cells.

## Recognition of phosphatidylserine by phagocytes

It has been known for some time that macrophages appear to recognize cells which have lost membrane asymmetry (Fidler *et al*, 1980; McEvoy *et al*, 1986; Schlegel and Williamson, 1987; Pradhan *et al*, 1994); however, a large body of data suggest that macrophages can recognize PS specifically. Several groups of investigators have found that human and rodent macrophages, as well as insect phagocytes, preferentially take up negatively charged liposomes, particularly those containing PS (e.g. Fidler *et al*, 1980; Schroit and Fidler, 1982; Mehta *et al*, 1982; Ratner *et al*, 1986; Allen *et al*, 1988; Mietto *et al*, 1989). In addition, human and rodent macrophages (including freshly isolated human alveolar and splenic macrophages, 10–14 day cultured human bone-marrow-derived macrophages, cultured human monocytes, resident and thioglycollate-elicited mouse peritoneal macrophages) can bind to and engulf symmetric red cell ghosts, red cells with PS inserted externally, oxidized red cells, or sickled red cells, all of which express PS externally (Tanaka and Schroit, 1983; Hebbel and Miller, 1984; Schwartz *et al*, 1985; Connor *et al*, 1994; Pradhan *et al*, 1994; Sambrano *et al*, 1994; Sambrano and Steinberg, 1995). Binding and/or phagocytosis of these PS-expressing red blood cells could be inhibited by preincubation of the macrophages with liposomes containing PS.

This work inspired our search for exposure of PS during apoptosis and for the mechanism by which this phospholipid was recognized. We initially studied the phagocytosis of apoptotic mouse thymocytes by thioglycollate-elicited mouse peritoneal exudate macrophages, which had been



**Figure 1** The binding of <sup>3</sup>H-glycerophosphorylserine to macrophages is inhibited by L, but not D, isomers of phosphatidylserine, glycerophosphorylserine, and phosphoserine. <sup>3</sup>H-glycerophosphorylserine (10<sup>-1</sup> M in HBSS +2mg/ml bovine serum albumin) was added to thioglycollate-elicited mouse macrophages adhered to tissue culture plastic in the presence or absence of unlabelled ligands for 30 min at 37°C, then washed and lysed. The lysates were incubated overnight in scintillation fluid, then analyzed for radioactivity in a  $\beta$ -counter. Ptd-D-serine=phosphatidyl-D-serine; Ptd-L-serine=phosphatidyl-L-serine

demonstrated previously to take up PS and to bind to PS-expressing red cells (Tanaka and Schroit, 1983). Recognition and uptake of apoptotic cells appeared specific for PS: liposomes containing PS inhibited phagocytosis in a dose-dependent manner, whereas liposomes containing other anionic phospholipids, such as phosphatidic acid, phosphatidylinositol, or phosphatidylglycerol, had very little effect (Fadok *et al*, 1992a). More importantly, by utilizing L and D isomers of PS and its structural analogues, we found that inhibition was stereospecific, which supported the notion that macrophages had receptors specific for PS. We performed ligand binding assays on whole macrophages using tritiated glycerophosphorylserine (GPS), a water soluble structural analogue containing the glycerol backbone and phosphoserine head group without the fatty acid side chains. This compound bound to macrophage membranes and its binding was competed successfully by L but not D isomers of PS, GPS and phosphoserine (Figure 1). These data lend further support to the hypothesis that macrophages had a receptor specific for PS.

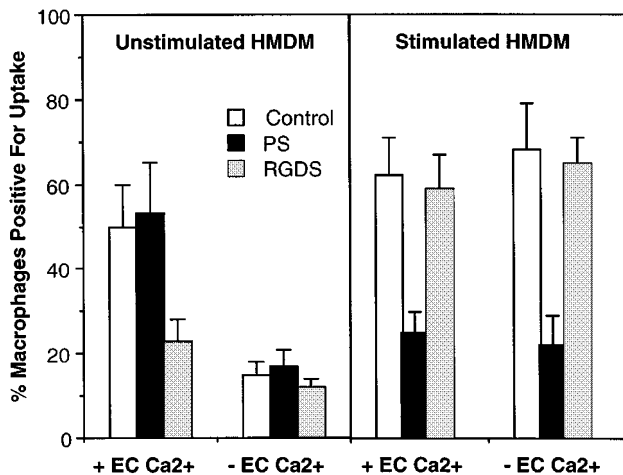
Savill and coworkers had established that human monocyte-derived macrophages (HMDM) could recognize apoptotic human neutrophils and lymphocytes by virtue of their vitronectin ( $\alpha v \beta 3$ ) receptors (VnR) and their Class B scavenger receptors (CD36, also a receptor for thrombospondin) (Savill *et al*, 1990, 1992). These receptors bound to thrombospondin (TSP), which served as a bridge between the macrophage and the apoptotic cell. We wondered if the differences in our systems related to the species we studied, the type of macrophage, or the type of apoptotic cell. We did not know the identity of the ligand on the apoptotic cell to which the thrombospondin bound; however, the data were most compatible with a negatively charged ligand (Savill *et al*, 1989). Because PS has a negative charge, we collaborated to determine if PS could be recognized by both sets of macrophages (Fadok *et al*, 1992b). We examined the uptake of apoptotic human neutrophils and mouse thymocytes by mouse bone marrow-derived macrophages (BMDM) (similar in many respects to HMDM) and thioglycollate-elicited macrophages. We found that regardless of apoptotic cell type, bone marrow cells were inhibited by RGDS and anti-VnR antibody, but NOT by PS liposomes. In contrast, the exudate cells were inhibited by PS liposomes but not RGDS or anti-VnR antibody. This led us to several conclusions: (1) that PS was probably not the ligand recognized by thrombospondin, CD36, or VnR; (2) that the differences in recognition mechanisms were not determined by differences between mice and humans or between types of apoptotic cells; (3) that recognition appeared to be determined by macrophage phenotype not the type of apoptotic cell. Since that time, however, evidence is accumulating that the picture is, in fact, more complex, and that the type of apoptotic cell may also contribute to which mechanisms are used for its removal. For example, Flora and Devitt with Christopher Gregory, have shown that CD14 is involved in the recognition of apoptotic cells by human monocyte-derived macrophages, but that this mechanism appears to be more efficient for lymphocytes compared to neutrophils (Flora and Gregory, 1994; Devitt *et al*, 1998). They also have suggested that ICAM-3 plays a

role in the recognition of apoptotic lymphocytes by macrophages (Gregory *et al*, 1995). Lastly, Hart *et al* (1997) have demonstrated that ligation of CD44 increases uptake of apoptotic neutrophils but not apoptotic lymphocytes.

Because thioglycollate is a proinflammatory stimulus, we wished to test the hypothesis that inflammatory macrophages recognized PS. We also wondered if a macrophage using the VnR-CD36-thrombospondin mechanism could be stimulated to recognize PS. Laszlo *et al* (1993) had demonstrated that stimulation of mouse BMDM with digestible particulate stimuli such as  $\beta$ -glucan could induce an increase in lysosomal hydrolases. Noble and coworkers went on to show that these macrophages secreted TGF $\beta$ , which was essential to the maturation of the inflammatory phenotype (Noble *et al*, 1993). We therefore stimulated mouse BMDM with  $\beta$ -glucan and found that over the next 48 h, inhibition of phagocytosis by RGDS was lost whereas inhibition by PS liposomes was acquired (Fadok *et al*, 1993). We confirmed these results using PS-expressing red cells in a rosetting assay. PS recognition was induced by digestible particulate stimuli and required the autocrine/paracrine production of either TGF $\beta$  or PAF (Fadok *et al*, 1993; Rose *et al*, 1995). We concluded that the PS receptor was one component of the inflammatory phenotype. Interestingly, although these macrophages no longer used the VnR receptor to bind and engulf apoptotic cells, there was no decrease in VnR expression on the cell surface (Fadok *et al*, 1993).

These findings raised several questions. Investigators studying a variety of human macrophages have shown that they could bind and phagocytose PS-expressing red cells, yet Savill and coworkers were never able to inhibit phagocytosis of apoptotic cells using phospho-L-serine by HMDM or by macrophages isolated from inflammatory sites (Ren and Savill, 1995, Savill, personal communication). We found that human macrophages, like mouse macrophages, required induction to recognize PS on apoptotic cells (Figure 2). Treating HMDM with digestible particulate stimuli such as  $\beta$ -glucan, zymosan, or even apoptotic cells (Warner and Fadok, unpublished data) resulted in phagocytosis of apoptotic cells that was inhibited stereospecifically by PS and its analogues, but not by other anionic phospholipids and not by anti-VnR antibodies or RGDS. As we found for the mouse bone marrow-derived macrophages, the addition of exogenous TGF $\beta$  plus latex could also induce PS recognition by these macrophages. The presence of a phagocytic stimulus was essential as TGF $\beta$  alone did not induce PS recognition (Ren and Savill, 1995). These data suggested to us that the PS receptor we proposed for mouse macrophages could also be induced on human macrophages, and led to the development of the model shown in Figure 3.

It is difficult to reconcile the disparities between the reports of human macrophage-mediated uptake of PS-expressing red cells, which is inhibited by PS liposomes, and uptake of apoptotic cells, which is not inhibited by PS liposomes unless the macrophages are stimulated first. One possible explanation is that macrophages do not see PS on the surface of the red cell in the same way that they

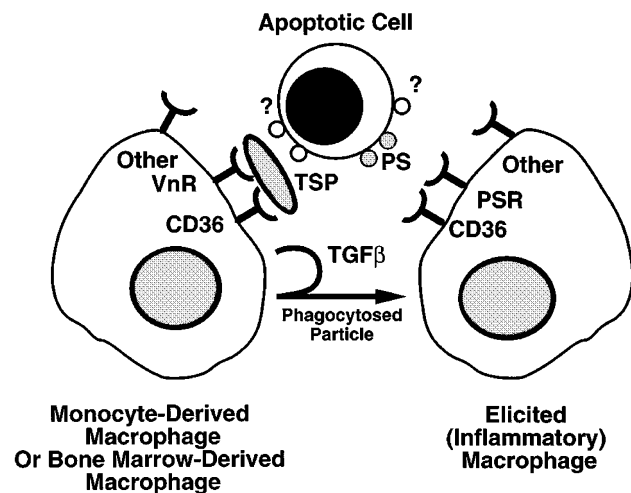


**Figure 2** PS-mediated phagocytosis of apoptotic neutrophils by human monocyte-derived macrophages (HMDM) does not depend on extracellular calcium. HMDM were stimulated or not with TGF $\beta$  (10 ng/ml) and  $\beta$ -glucan (10  $\mu$ g/ml) for 48 h prior to the phagocytosis assay, which was performed in RPMI (+EC Ca<sup>2+</sup>) or Ca<sup>2+</sup>, Mg<sup>2+</sup> free RPMI with 5 mM EDTA (-EC Ca<sup>2+</sup>). The preparation of PS-containing liposomes is described in Fadok *et al* (1992a); total lipid concentration was 0.1 mM; RGDS was used at 1 mM. Control indicates macrophage uptake in the absence of either inhibitor

see it on an apoptotic cell. This hypothesis is supported by the work of Terpstra *et al* (1997); they found that macrophage binding of PS-expressing oxidized red cells was inhibited by the presence of EDTA, whereas the binding of apoptotic cells was not. Also, we have found that PS-inhibitable uptake by stimulated human macrophages (Figure 2) and thioglycollate-elicited macrophages (Fadok, unpublished data) is not dependent on extracellular Ca<sup>2+</sup> or Mg<sup>2+</sup>; this is in contrast to the Ca<sup>2+</sup> and Mg<sup>2+</sup> dependence observed for the VnR-CD36-thrombospondin mechanism described by Savill *et al* (1990). Another possibility is suggested by the work of Pradhan *et al* (1997); they have attempted to synthesize all these findings in a series of experiments comparing uptake of apoptotic cells by the mouse macrophage cell line J774 with that by thioglycollate-elicited peritoneal exudate macrophages. They found that uptake of apoptotic cells by J774 macrophages was inhibited by RGDS and by symmetric red cell ghosts, but not by PS liposomes. In contrast, uptake by the exudate macrophages was inhibited by PS liposomes, but not by symmetric red cell ghosts or RGDS. They proposed that each macrophage type can recognize PS; the macrophages just see it in a different context or in association with a different additional ligand.

### The nature of the phosphatidylserine receptor

It is clear that macrophages can recognize phosphatidylserine in a dose-dependent and stereospecific manner (Mietto *et al*, 1989; Fadok *et al*, 1992a) which suggests that a PS receptor exists. As mentioned above, we have identified macrophages



**Figure 3** Uptake of particulate stimuli induce the autocrine/paracrine release of TGF $\beta$  and promotes the switch to PS recognition. This simplified model suggests that macrophages which utilize the VnR-CD36-TSP mechanism can be induced to recognise PS by treatment with digestible particulate stimuli, which can include zymosan,  $\beta$ -glucan, or even apoptotic cells themselves. CD36 appears to be required for uptake by either mechanism. The role of other receptors in this model has not yet been clarified. TSP=thrombospondin

which phagocytose apoptotic cells in a manner inhibited stereospecifically by PS and its structural analogues. Other anionic phospholipids, including phosphatidylinositol, phosphatidylglycerol and phosphatidic acid were inactive (Fadok *et al*, 1992a). Is this recognition mediated by a unique receptor, or by receptors which have been identified previously? Several receptors have been suggested to play a role in mediating PS recognition, including scavenger receptors, CD68 (an oxLDL receptor), CD14, annexins,  $\beta$ 2 glycoprotein I and gas-6.

### Scavenger receptors

Some (but not all) of the scavenger receptors (SR) are known to mediate the uptake of PS liposomes (for a review, see Pearson, 1996) and a variety of SR have been implicated in the uptake of apoptotic cells (Savill *et al*, 1992; Sambrano and Steinberg, 1995; Fukasawa *et al*, 1996; Platt *et al*, 1996).

### Class A scavenger receptors

Platt and coworkers showed that the class A macrophage scavenger receptor (SR-A) mediates the uptake of apoptotic thymocytes by mouse thymic macrophages and thioglycollate-elicited peritoneal macrophages. In fact, thymic macrophages from SR-A deficient mice showed a 50% decrease in uptake of apoptotic thymocytes (Platt *et al*, 1996). Terpstra and coworkers recently examined the ability of resident peritoneal macrophages from SR-A deficient mice to bind apoptotic thymocytes; they found a 21% decrease in binding (Terpstra *et al*, 1997). The ligand recognized by this receptor, however, remains to be

**Table 1** Potential candidates for recognition of phosphatidylserine on apoptotic cells

PS-binding protein	Bind PS	Ca <sup>2+</sup> dependence of PS binding	Bind other anionic phospholipids
Class A scavenger receptors	No	Not applicable	No
Class B scavenger receptors	Yes	? <sup>b</sup>	Yes
CD36	Yes	?	Yes
CLA-1	Yes	?	Yes
CD68	Yes	No <sup>c</sup>	Yes
CD14	?	?	Yes <sup>d</sup>
Annexins	Yes	Yes	Yes
$\beta$ 2-GPI	Yes	No	Yes
Gas-6	Yes	Yes	Yes
Stimulated macrophage PSR <sup>a</sup>	Yes	No	No

<sup>a</sup>PSR (phosphatidylserine receptor) indicates the PS-inhibitable uptake mechanism described by Fadok *et al* (1992a) in stimulated macrophages (see text). The actual receptor has not been identified. <sup>b</sup>Ca<sup>2+</sup> dependence was not assessed for the Class B scavenger receptors (Rigotti *et al*, 1995; Murao *et al*, 1997). <sup>c</sup>Binding of PS determined by ligand blotting in the presence of EDTA (see Sambrano and Steinberg, 1995). <sup>d</sup>Other anionic phospholipids not assessed (Sambrano and Steinberg, 1995).

identified, as the class A SR do not bind phosphatidylserine (Lee *et al*, 1992).

### Class B scavenger receptors

The Class B scavenger receptors include SRB1 (and its human homologue CLA-1 [CD36 and LIMP-II analogous-1]) and CD36. SRB1 can bind to PS liposomes (Rigotti *et al*, 1995; Fukasawa *et al*, 1996) and when transfected into CHO cells can phagocytose apoptotic cells (Fukasawa *et al*, 1996); however, SRB1 is expressed primarily in liver and steroidogenic tissues (Acton *et al*, 1996), making it less likely as a candidate for a general macrophage PS receptor. CLA-1 is found in the liver and testis; however, it also has been found on circulating monocytes (Murao *et al*, 1997). When transfected into HEK 293 cells, it facilitated the binding of apoptotic cells, which was inhibited by liposomes containing either PS or PI (Murao *et al*, 1997). The most interesting potential candidate for PS recognition is the other Class B scavenger receptor CD36, which is also a thrombospondin receptor. CD36 has been implicated in uptake of apoptotic neutrophils by human macrophages (Savill *et al*, 1992) and has been suggested to mediate the uptake of PS-expressing photoreceptor outer segments (ROS) in retinal pigment epithelium (Ryeom *et al*, 1996). Rigotti *et al* (1995) showed that the binding of acLDL to cells transfected with human CD36 was blocked by PS liposomes; however, phosphatidylinositol (PI) was more potent in this assay. The uptake of ROS by retinal pigment epithelium was also inhibited by both PS and PI liposomes, as would be expected for a CD36-mediated mechanism (Ryeom *et al*, 1996). In contrast, we have never observed significant inhibition of phagocytosis of apoptotic cells by PI liposomes in either human or mouse macrophages (Fadok *et al*, 1992a; and unpublished data). Ren, Navazo and coworkers transfected human CD36 into melanoma cells and COS cells, and showed that phagocytosis of apoptotic cells was upregulated. However, phagocytosis was not inhibited by phospho-L-serine (Ren *et al*, 1995; Navazo *et al*, 1996), a ligand we have found effective in both human and mouse macrophages stimulated to

recognize PS on apoptotic cells (Fadok *et al*, 1992a; unpublished data). Unfortunately, the effects of PS liposomes in the CD36-transfected melanoma and COS systems were not examined.

We have attempted to determine the role of CD36 in PS recognition by studying phagocytosis of apoptotic cells by human macrophages stimulated to recognize PS with  $\beta$ -glucan (Warner *et al*, unpublished data). Stimulation resulted in a macrophage population whose phagocytosis of apoptotic cells was no longer inhibited by RGDS or anti-VnR antibodies, but was inhibited by PS liposomes. Antibodies against CD36 inhibited the phagocytosis of either stimulated or unstimulated human macrophages. Flow cytometric analysis showed that particulate stimuli caused the downregulation of VnR on macrophages from 10 of the 12 donors we have examined; however, CD36 expression remained the same. If macrophages were pretreated with PS liposomes in an attempt to downregulate the PS receptor, the binding of anti-CD36 monoclonal antibodies was not decreased. PS liposomes also did not compete with the antibody for binding to CD36. These data suggested to us that CD36 was not the PS-recognizing receptor in our system. If, however, CD36 was downregulated by incubating the cells on antibody-coated plates, phagocytosis was reduced and PS inhibition was no longer statistically significant. These data, together with our observations that phosphatidylinositol-containing liposomes did not inhibit phagocytosis, suggested to us that CD36 did not recognize PS directly, but that it did cooperate with our proposed PS receptor (see Figure 3), perhaps in a fashion similar to that proposed for its cooperation with VnR (Savill *et al*, 1992).

### CD68 (oxidized LDL receptor)

Sambrano and Steinberg (1995) have shown that the 94–97 kd oxLDL receptor found on resident peritoneal macrophages may mediate uptake of apoptotic cells. This oxLDL receptor is identical to mouse macrosialin or its human homologue CD68 (Ramprasad *et al*, 1995). The oxLDL

receptor appeared to mediate the recognition of apoptotic cells and phagocytosis was inhibited by PS liposomes (Sambrano and Steinberg, 1995). Direct binding of PS liposomes to a 94–97 kd band from macrophage membrane preparations was also demonstrated (Sambrano and Steinberg, 1995). In our studies of human macrophages, surface expression of CD68 was not detectable by flow cytometry, and anti-CD68 antibodies did not inhibit phagocytosis of apoptotic cells (Warner *et al*, unpublished data). These data suggest that, although CD68 may mediate binding to PS on apoptotic cells, it is not the only receptor to do so, as CD68-negative macrophages can also phagocytose apoptotic cells in a PS-inhibitable manner.

### CD14

The antigen identified by monoclonal antibody 61D3 has been shown to be involved in phagocytosis of apoptotic lymphocytes by human macrophages, and has recently been identified as CD14 (Flora *et al*, 1994; Devitt *et al*, 1998). Soluble CD14 has been shown to bind directly to phosphatidylinositol and to a fluorescent derivative of phosphatidylethanolamine, and it has been suggested to be a soluble shuttle for lipids (Yu *et al*, 1997). Two pieces of preliminary evidence from the laboratories of Williamson and Schlegel have suggested that CD14 might recognize PS on apoptotic cells (Callahan *et al*, 1997). First, removal of CD14 from the surface of J774 cells (a mouse macrophage line) with phospholipase C resulted in decreased phagocytosis of apoptotic cells and lipid-symmetric red cell ghosts. Second, an LPS-resistant variant of J774, which lacks functional CD14, showed reduced capacity to phagocytose apoptotic cells or symmetric red cell ghosts. Direct demonstration of PS binding to CD14 is still required, particularly demonstration of the ability of CD14 to mediate binding to PS on a cell surface.

### Annexins

Annexins comprise a group of proteins which bind to phospholipids in the presence of  $\text{Ca}^{2+}$  (Reutelingsperger and van Heerde, 1997; Swairjo and Seaton, 1994). In fact, this property has been exploited to develop a rapid and simple assay for the assessment of PS exposure on apoptotic cells (Vermes *et al*, 1995; Zhang *et al*, 1997). Prydzial and coworkers have some intriguing data to suggest that annexin II can mediate the binding of herpes viruses to host cells. They found that cytomegalovirus and herpes simplex types 1 and 2 expressed phosphatidylserine on their surfaces, which facilitated the association of human annexin II to virus particles within human fibroblasts (Sutherland *et al*, 1997; Wright *et al*, 1995; Prydzial and Wright, 1994). They were able to inhibit viral infection of human fibroblasts with a rabbit antiserum raised against human annexin II, suggesting that annexin II might facilitate the uptake of these viruses. Is it possible that annexins could mediate the binding of apoptotic cells to phagocytes? Although primarily intracellular in location, some annexins can be found in the blood and extracellular spaces (Reutelingsperger and van Heerde, 1997). Our studies with human and mouse macrophages suggest that annexins are unlikely candidates for mediating

the binding of apoptotic cells, as PS-inhibitable uptake is not lost in the absence of  $\text{Ca}^{2+}$  (Figure 2). Furthermore, preliminary experiments showed that anti-annexin V did not inhibit uptake of apoptotic neutrophils by human monocyte-derived macrophages, whether they were stimulated to recognize PS or not (Fadok, unpublished data). Reutelingsperger and van Heerde (1997) have made the intriguing suggestion that annexins, in fact, may help control coagulation in inflammatory sites *in vivo* by binding to the procoagulant surfaces of apoptotic cells.

### $\beta$ 2-glycoprotein I

$\beta$ 2-glycoprotein I ( $\beta$ 2-GPI) is a serum protein believed to regulate coagulation. It binds to anionic phospholipids such as cardiolipin, phosphatidylserine, phosphatidylglycerol, phosphatidylinositol, and phosphatidic acid (Balasubramanian *et al*, 1997), and forms an important epitope for many of the 'anti-cardiolipin' and 'anti-phospholipid' antibodies identified in patients with the antiphospholipid antibody syndrome (aPL) (Roubey, 1996). Price *et al* (1996) showed that anti-phospholipid antibodies from patients with primary aPL or aPL associated with systemic lupus erythematosus could opsonize apoptotic cells in the presence of  $\beta$ 2-GPI. Subsequently, Balasubramanian *et al* (1997) showed that the binding of  $\beta$ 2-GPI to apoptotic cells or lipid-symmetric red cells, followed by the binding of anti- $\beta$ 2-GPI, facilitated uptake into macrophages.  $\beta$ 2GPI alone was able to increase the uptake of symmetric red cells by 50%; however, uptake of apoptotic thymocytes was only slightly enhanced in the absence of the opsonizing antibody. These data suggest that PS on apoptotic cells could contribute to FcR-mediated uptake in patients with the appropriate autoantibodies; this would be expected to be a proinflammatory pathway which would contribute to the disease process, as uptake of opsonized apoptotic cells results in the release of proinflammatory cytokines and lipid mediators (Meagher *et al*, 1992; Stern *et al*, 1996; Fadok *et al*, 1998).

### Gas-6

Gas-6, the product of growth arrest-specific gene 6 (Manfioletti *et al*, 1993), is the newest potential candidate for mediating the binding of macrophages to PS on apoptotic cells. It is a soluble protein related to protein S, a negative regulator of coagulation, and has been shown to mediate the binding of U937 cells (a human histiocytic line) to PS-coated ELISA plates in the presence of  $\text{Ca}^{2+}$  (Nakano *et al*, 1997). Nakano and coworkers have made the intriguing suggestion that gas 6 might mediate the binding of PS-expressing cells to phagocytes. Gas-6 binds to a set of unique receptor tyrosine kinases (axl, sky and mer) which contain extracellular domains similar to those found in cell adhesion molecules (Varnum *et al*, 1995; Nagata *et al*, 1996). Axl mRNA expression has been demonstrated by PCR in normal human bone marrow and at low levels in peripheral blood monocytes; treatment of human monocytes with  $\text{IFN}\alpha$  was associated with upregulation of message (Neubauer *et al*, 1994). Whether gas-6 is involved in the uptake of apoptotic cells by macrophages remains to be determined. If so, gas-6 may

serve as a bridging molecule between the apoptotic cell and the macrophage, in a manner similar to that observed for thrombospondin.

To summarize our thoughts, then, on the nature of PS receptor in macrophages, we believe that the phagocytic activity we have described, the uptake of apoptotic cells which is inhibited stereospecifically by PS and its structural analogues, GPS and phospho-L-serine; which is not inhibited by other negatively charged phospholipids, polyanions, or acLDL; and which is expressed on macrophages stimulated with digestible particulate stimuli, is distinct from the currently described scavenger receptors and CD14, and has not been identified to date. This activity can be removed from the surface of the macrophages with proteases. Incubation with PS liposomes inhibits its function with maximal inhibition reached by 20 min after addition of the liposomes, followed by recovery of phagocytic activity 24 h later. We also believe that the activity we have described is not likely to be mediated by annexins or gas-6, as the binding of those proteins to PS requires  $\text{Ca}^{2+}$  and phagocytosis by PS recognizing macrophages (which we define strictly as inhibition by PS liposomes) is not inhibited in the absence of calcium. We think that  $\beta 2$ -GPI would most likely facilitate phagocytosis in diseased states where autoantibodies are produced, rather than in the normal state. What seems likely is that multiple PS-recognizing receptors exist; determining how they may interact, what exactly they see and when they operate, is a critical objective.

Lastly, there is also some evidence to suggest that amateur phagocytes can recognize PS on apoptotic cells. Vascular smooth muscle cells phagocytose their apoptotic counterparts in a manner inhibited by preincubation with PS liposomes or with annexin V (Bennett *et al*, 1995). Neither PI liposomes nor anti-annexin V antibodies inhibited phagocytosis in this system, suggesting that neither the known scavenger receptors nor annexins were mediating phagocytosis by vascular smooth muscle cells. In contrast, the recognition of spermatogenic cells by Sertoli cells was inhibited by PS-, PI- and CL-containing liposomes (Shiratsuchi *et al*, 1997), suggesting the use of a different receptor from that used by stimulated macrophages or vascular smooth muscle cells. However, phagocytosis by Sertoli cells was also inhibited by glycerophosphorylserine and weakly by phospho-L-serine, whereas phospho-D-serine had no effect, suggesting some stereospecificity.

## The biological consequences of PS expression

What does exposure of PS on the outer leaflet of an apoptotic cell mean, from a biological point of view? We favor it as a recognition ligand for rapid removal; however, it is not clear at this point what the role of PS as a recognition ligand is in vivo and whether all macrophages, in fact, see it. There must be other ligands on the apoptotic cell in addition to PS. The early work of Duvall and coworkers suggested changes in surface carbohydrates (Duvall *et al*, 1985); most recently Falasca and coworkers have shown increased expression of mannose, N-acetylgalactose, and possibly galactose on the surface of

apoptotic cells (Falasca *et al*, 1996). However, the ligands which mediate the binding of thrombospondin to apoptotic cells remain to be determined. It is intriguing to speculate that loss of membrane phospholipid asymmetry may facilitate exposure of these ligands as well.

Exposure of PS on cell surfaces has other potential biological consequences. It has been known for some time that PS serves as a cofactor for the coagulation cascade (Zwaal and Schroit, 1997), and there is a small but growing body of evidence to suggest that PS can have a variety of anti-inflammatory or immunosuppressive effects. In his recent review, Kornbluth (1994) cited a number of instances in which PS could inhibit macrophage parasitocidal activity and cytokine production, as well as lymphocyte proliferation and cytokine production. More recently, two different groups of investigators (Aramaki *et al*, 1996; DiNapoli *et al*, 1997) have shown that phosphatidylserine-containing liposomes inhibited transcription of inducible nitric oxide synthase and nitric oxide production in mouse macrophages stimulated with LPS. Phagocytosis of apoptotic cells by non-FcR mediated uptake mechanisms not only fails to stimulate inflammatory mediator production in normal human macrophages (Meagher *et al*, 1992; Stern *et al*, 1996) but also inhibits inflammatory mediator production stimulated by LPS or zymosan (Voll *et al*, 1997; Fadok *et al*, 1998). Perhaps PS is one of the inhibitory signals found on apoptotic cells. Lastly, exposure of PS on the surface of apoptotic cells may contribute to the inflammation and coagulation defects seen in patients with systemic lupus erythematosus and primary aPL by binding autoantibodies in the presence of proteins such as  $\beta 2$ GPI (see section on  $\beta 2$ GPI above).

## Conclusions

It seems clear that exposure of phosphatidylserine is a common feature on apoptotic cells and that it serves as a recognition signal for phagocytosis by at least a subset of macrophages. Several questions remain. How is this phospholipid recognized? In particular, how does the macrophage get past the glycocalyx to access this lipid? There are two potential clues. First, there appears to be an alteration of surface carbohydrates on apoptotic cells (Falasca *et al*, 1996). Second, Casciola-Rosen *et al* (1996) have shown by confocal microscopy that annexin V binding is strongest on the surface of small membrane blebs, suggesting the appearance of domains. Is PS recognized on its own merits or is it seen in conjunction with another ligand, such as a protein or carbohydrate? If PS expression is a critical recognition signal, why do not all macrophages appear to recognize it? Is inhibition by PS liposomes too strict a definition, as has been suggested by the work of Pradhan *et al* (1997)? Perhaps in addition to signalling phagocytosis on its own, loss of phospholipid asymmetry promotes the exposure of other apoptotic ligands. Is there a receptor specific for PS or are multiple receptors (including the scavenger receptors) capable of recognizing this lipid on apoptotic cells? It seems most likely that multiple receptors could bind PS, but whether they can recognize it on a cell surface remains to be determined. How does PS exposure



contribute to the biological effects of phagocytosis of apoptotic cells, which include suppression of proinflammatory mediator release? These questions will provide fruitful research for us and other 'phagocytophilic' investigators in the coming years.

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