Type 3-repeat/C-terminal domain of thrombospondin-1 triggers caspase-independent cell death through CD47/αvβ3 in promyelocytic leukemia NB4 cells

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Running title: TSP-1-induced caspase-independent cell death

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ABSTRACT

By means of its anti-angiogenic activity, thrombospondin-1 (TSP-1) exerts indirect antitumoral action on solid tumors. Here, we investigated potential antitumor action in an in vitro cell model for promyelocytic leukemia (NB4-LR1), resistant to retinoid maturation. Purified soluble TSP-1 added to cultures induced a strong dose-dependent growth inhibition and a slowly developing maturation-independent cell death. Recombinant fragments of TSP-1 allowed to map these activities to its type 3-repeat/C-terminal domain, features that are distinct from those of TSP-1 action on solid tumors, previously ascribed to the type 1-repeat domain. Cell death in leukemia was characterized as a caspase-independent mechanism, without DNA fragmentation, but phosphatidylserine externalization followed by membrane permeabilization. Mitochondria membrane depolarization was inherent to TSP-1 action but did not produce release of death-promoting proteins (e.g. non-caspase apoptosis regulators, AIF, endonuclease G or Omi/HtrA2 or the caspase regulators, cytochrome c or Smac/Diablo). Although detected, ROS production was likely not involved in the death process. Finally, receptor agonist RFYVVM and RGD peptides indicated that TSP-1 death effects are mediated by membrane receptors CD47 and αvβ3. These results demonstrated a new domain specific anti-tumoral activity of TSP-1 on a leukemia cell line, which extends TSP-1 therapeutical potential outside the area of vascularized solid tumors.
INTRODUCTION

Cell interaction with extracellular matrix proteins is essential to regulate proliferation, differentiation and cell death.\(^1\) Thrombospondin-1 (TSP-1) belongs to a family of multidomain and multifunctional glycoproteins that regulate cell proliferation, migration, differentiation and is important for embryonic development, morphogenesis, inflammation, tumor metastasis and multiple other processes.\(^4\) TSP-1 is a secreted component of extracellular matrix that supports attachment and motility of a variety of normal and neoplastic cell types.\(^5\) Since it also functions as soluble protein that antagonizes the pro-adhesive activities of other matrix proteins, TSP-1 has been defined as a matricellular protein, modulating cell-matrix interactions.\(^9\) It can affect matrix structure through its ability to interact with matrix components and to activate TGF-\(\beta\)1.\(^11\) TSP-1 is also a potent physiological inhibitor of angiogenesis by impairing endothelial cell growth and migration and by acting on endothelial cell cycle progression and apoptosis, then down–expression of TSP-1 is important for tumor progression.\(^12\) Many of the TSP-1 functions have been located within specific domains of the molecule which bind to various membrane receptors including integrins of the \(\beta1\) and \(\beta3\) family, CD47/IAP (integrin-associated protein), CD36 and proteoglycans.\(^5,8\) Therefore, cell responses to TSP-1 probably reflect the integration of multiple membrane signals depending on the cell type and its repertoire in membrane receptors.

TSP-1 function in tumor cell differentiation and apoptosis remains poorly explored. TSP-1 has been described to be positively involved in retinoic acid-induced maturation of neuroblastoma cells and the acute myeloblastic leukemic cell line HL-60.\(^16,17\) In the HL60 cell line, it was shown to induce a caspase-dependent cell death mediated by CD36.\(^18\) It is also able to trigger a caspase-independent cell death that is accompanied by selective mitochondrial changes in B-chronic lymphocytic leukemia through binding to CD47.\(^19,20\) TSP-1 also induces CD47-mediated killing of breast cancer cells acting via heterotrimeric G\(_i\)-dependent inhibition of protein kinase A activity.\(^21\)

NB4 cells have been derived from leukemic cells of a patient with acute promyelocytic leukemia (APL)\(^22\) and carry the characteristic t(15;17) translocation giving rise to the expression of the retinoic acid receptor fusion protein (PML-RAR\(_{\alpha}\)).\(^23,24\) Pharmacologic doses of all-trans retinoic acid (ATRA) can convert the fusion protein into an
active receptor and restore granulocytic maturation of these cells, so that its administration to APL patients either alone or combined with chemotherapy is of therapeutic benefit and improve the survival prognosis. The NB4-LR1 cell subline is resistant to maturation by ATRA, and was isolated from the early culture of NB4 cells derived from leukemic cells of the APL patient. It was found that NB4-LR1 cells required cAMP analog and ATRA signaling cross-talk to undertake granulocytic maturation program, suggesting interdependent ATRA and cAMP signaling pathways in acute promyelocytic leukemia maturation. Noteably, a wide spectrum of membrane signals have been shown to modulate or to cooperate with retinoids to induce, in NB4 leukemic cells, maturation or cell death independently of maturation. No investigation has been carried out so far to know whether biological response to TSP-1 signalling could be exploited in APL.

In this work, we investigated the biological outcomes of TSP-1 signaling or ATRA and TSP-1 combined signaling in NB4-LR1 leukemic immature blast cells refractory to ATRA maturation and defective in endogenous TSP-1 synthesis and secretion. The homotrimeric TSP-1 molecule was analysed by means of domain-specific recombinant TSP-1-derived proteins to map the anti-tumoral activities of this protein. The C-terminal domain of TSP-1 comprising the type 3 repeats and the C-terminal globular region is shown herein to signal, by itself or combined with ATRA, a caspase-independent cell death in these leukemic cells through simultaneous ligation of CD47 and the integrin αvβ3.
METHODS

Cell culture and treatment

The APL cell line NB4 and the resistant APL cell line NB4-LR1 were essentially cultured in RPMI 1640 medium (Invitrogen, Carlsbad, CA) as previously described.22,28 Cells (2x10^5/mL) were untreated or treated for 4 days with 1 µM all-trans retinoic acid (ATRA) (Sigma, St Louis, MO) in the absence or presence of soluble human purified platelet TSP-1 or recombinant fragments for specific domains of TSP-1 (see below). In some experiments, cells were cultured with benzoyloxycarbonyl-Val-Ala-Asp-fluoromethylketone (z-VAD-fmk) or RFVVVMWK (4N1-1), GRGDS or VTCG peptides (BACHEM Biochimie, Voisins-le-Bretonneux, France), selenium (Sigma) or Ebselen (Calbiochem, San Diego, CA). Cell morphology was evaluated by May-Grünwald-Giemsa (MGG) staining. Biochemical criterion of granulocytic differentiation was assessed by the nitro blue tetrazolium (NBT, Sigma) dye reduction assay.22,27,28

Purified TSP-1 and TSP-1 recombinant fragments

Human TSP-1 was purified from the supernatant of thrombin-activated platelets as previously described.30 Recombinant proteins including specific sequences for the NH2-terminal (NTSP-1), the three type 1 repeats (3TSR) and the type 3 repeat/C-terminal domain (T3C1) of TSP-1 were produced in Drosophila S2 cells. The preparation of 3TSR has been described previously.31 NTSP-1 was prepared using the forward primer 573htsp1f (GAT GAT CCA TGG AAC CGC ATT CCA GAG TCT GGC) and the reverse primer 574htsp1r (GAT ACC GGT GTT AGT GCG GAT GGC AGG GCT). The PCR product was sequenced and cloned between the \textit{NcoI} and the \textit{AgeI} sites of the vector pMT/BiP/V5-HisA (Invitrogen). The recombinant protein includes the vector-derived sequence RSPW at the N-terminal. T3C1 was prepared using the forward primer 575htsp1f (GAT GAT CCC GGG GAC ACA GAC CTG GAT GGC TGG) and the reverse primer 576htsp1r (GAT ACC GGT GTT AGT GCG GAT GGC AGG GCT). The PCR product was sequenced and cloned between the \textit{NcoI} and the \textit{AgeI} sites of the above vector. The recombinant protein includes the sequence RSPWPG at the N-terminal. Vector transfection, cell selection and protein expression and purification were performed as described previously.31
RNA preparation and Northern blot analysis

Total RNA was extracted from $10^7$ cells using TRIZOL® Reagent (Invitrogen) according to the manufacturer’s instructions. 20 µg of RNA from each sample were denatured in formamide, electrophoresed on a 1% agarose gel containing 2.2 M formaldehyde, and transferred overnight onto nylon membrane (Hybond-N, Amersham Pharmacia Biotech, Orsay, France) using standard protocols. Membrane was probed with a $^{32}$P-labeled fragment of human cDNA specific for TSP-1 encoding amino acids sequence in the NH$_2$-terminus of the molecule (a kind gift of Pr Valerie Castle, University of Michigan Medical School, MI).

Immunofluorescence analysis

Fresh or paraformaldehyde (PFA)-fixed cells in RPMI medium containing 2% bovine serum albumin (BSA) were incubated for 30 min at room temperature with mouse monoclonal antibodies to CD36, the β1 or αv integrin subunit (Immunotech, Marseille, France), the β3 integrin subunit or CD47 (BD Biosciences, Rungis, France), rabbit antiserum to TSP-1 (R1) or αv (Chemicon International, Temecula, CA). Alexa Fluor 488 and 594-conjugated anti-mouse and anti-rabbit IgG were used as secondary antibodies (Molecular Probes, Eugene, OR). Immunofluorescence was analysed using a FACSCalibur flow cytometer (BD Biosciences) or a Bio-Rad MRC 1024 confocal imaging system (Herts, UK), equipped with an inverted Diaphot 300 Nikon microscope and a krypton-argon ion laser.

For analysis of mitochondrion protein localization, cells were permeabilized for 10 min with 0.1% saponin in PBS containing 10% BSA (PBS-BSA-0.1% saponin), then incubated with monoclonal antibody to cytochrome c (BD Biosciences) or rabbit polyclonal antibodies to Smac/Diablo (ProSci, Poway, CA), Omi/HtrA2 (Alexis Biochemicals, Coger, Paris, France), AIF (apoptosis-induced factor) or endonuclease G (Sigma) in PBS-BSA-0.1% saponin. Cells were washed in PBS-BSA-0.01% saponin then incubated with Alexa Fluor-conjugated secondary antibodies.

Subcellular fractionation

Cells in PBS were resuspended in buffer (250 mM sucrose, 10 mM KCl, 1 mM MgCl$_2$, 1.5 mM EDTA, 1.5 mM EGTA, 1 mM DTT, 20 mM Hepes pH 7.4, 0.1 mM PMSF and 10 µg/ml leupeptin, sodium orthovanadate, sodium fluoride and benzamidine) for 20 min on ice, then lysed using a Dounce tissue grinder. Nuclei and unlysed cells were sedimented at 1000 g for 10 min and the supernatant was centrifuged at 100,000 g to recover organelle- and
membrane-free cytosol (S100 fraction). Cytochrome c was immunodetected by Western blotting.

**Immunoprecipitation and Western blot analysis**

Cells \((10^7)\) and culture media corresponding to the collected cells were solubilized for 30 min at 4°C by adding lysis buffer containing 1% Triton X-100, 1 mM MgCl\(_2\), 2 mM CaCl\(_2\), and the following protease inhibitors (Calbiochem): 100 µM leupeptin, 9 KIU (Kallikrein Inhibitory Unit)/mL aprotinin, 2 mM sodium orthovanadate, 10 mM sodium fluoride and 1 mM \(p\)-aminoethylbenzenesulfonyl fluoride (AEBSF) in 20 mM Tris-HCl buffer (pH 7.4). Immunoprecipitation assays were carried out on soluble material essentially as reported\(^{33}\), using P10 mouse monoclonal antibody to TSP-1 (Immunotech) and Protein A Sepharose CL-4B (Amersham Pharmacia Biotech). The immune complexes were analysed by Western blotting, as described\(^{35}\), using P10 antibody (20 µg/mL) and peroxidase-conjugated secondary antibody (1:10,000, Dako, Glostrup, Denmark).

For Western blot analysis of total cell extract, cells were solubilized for 10 min at 100°C in the presence of 2% SDS, 5 mM N-ethylmaleimide acid (NEM), 2 mM EDTA and analysed using mouse monoclonal antibody to PARP (poly ADP-ribose polymerase) (Calbiochem) or rabbit polyclonal antibodies to human caspase-3, caspase-8 or Bcl-2 (Santa-Cruz Biotechnology, Santa-Cruz, CA).

**Assays for cell death**

*Annexin V binding and propidium iodide (PI) staining*

Cell death was assessed by analysing phosphatidylserine (PS) membrane outside exposure and cell membrane permeability\(^{36,37}\) by flow cytometry, using an apoptosis detection kit (Bender MedSystems, Vienna, Austria) for cell double labeling with 1 µg/mL FITC-labeled annexin V and PI.

*Caspase activity*

Caspase-3 activity was measured on cell extract (20-50 µg) by a colorimetric procedure using the specific peptide substrate DEVD-pNa, as described\(^{38}\).

*DNA fragmentation*

DNA was isolated from 2x10\(^6\) cells according to a salting out procedure previously described\(^{39}\), and internucleosomal fragmentation was analysed as already reported\(^{40}\).
Mitochondrial membrane potential determination

Determination of the mitochondrial transmembrane potential ($\Delta \Psi_m$) was assessed using the 3,3’dihexyloxacarbocyanine iodide (DiOC6(3)) lipophilic cationic fluorochrome (20 nM, Uptima, Interchim, Montluçon, France).

Reactive oxygen species (ROS) production

Intracellular ROS was measured through $\text{H}_2\text{O}_2$ production, using the oxidant sensitive nonfluorescent dye 2’-7’-dichlorofluorescein diacetate (DCFH-DA) (2 µM, Molecular Probes) which is oxidized by $\text{H}_2\text{O}_2$ and generate fluorescent dichlorofluorescein (DCF).
RESULTS

TSP-1 and TSP-1 receptor expression in NB4 leukemic cell lines

TSP-1 expression was examined both in ATRA-maturation sensitive NB4 cells and in ATRA-maturation resistant NB4-LR1 cells. In NB4 cells, ATRA dose-dependent TSP-1 mRNA expression was found to correlate with detection of granulocytic maturation, as measured by the NBT dye reduction assay (Figure 1A), and as viewed by MGG staining showing segmented nuclei typical of differentiated phenotype (not shown). TSP-1 protein expression, when examined by quantitative immunoprecipitation during 1 µM ATRA treatment (Figure 1B), showed a modulation in the pattern of TSP-1 protein expression similar to that obtained for TSP-1 transcript. It was noted that upon its induction by ATRA, TSP-1 was mostly secreted into the extracellular culture medium. In NB4-LR1 cells, neither TSP-1 transcript (Figure 1C) nor TSP-1 protein (not shown) expression was detected upon 1 µM ATRA treatment. However, when induced to mature by co-treatment with cAMP analog (8-CPT-cAMP), NB4-LR1 expressed TSP-1 similarly to what was observed in ATRA-treated NB4 cells (not shown).

Potential membrane receptors for TSP-1, such as CD47, the β1 and β3 integrin families were similarly expressed in NB4 and NB4-LR1 cells. Expression of these membrane markers was not modulated by ATRA treatment (Figure 1D), contrasting with the leukocyte β2 integrin, CD11c, for which membrane expression was reported to be upregulated.28,41 Note that the two cell lines were negative for CD36.

We therefore investigated whether TSP-1, upon its synthesis and secretion induced by ATRA in NB4 cells, could play a role in maturation/apoptosis programs triggered by retinoids. To this aim, the ATRA-maturation resistant NB4-LR1 cells which express potential TSP-1 receptors, but do not synthesize TSP-1 in response to ATRA, constitute an exquisite cell model to check whether exogenous TSP-1 signaling could act with ATRA for antitumoral responses.

Cell growth inhibition and cell death induction by TSP-1 in NB4-LR1 cells

NB4-LR1 cells, untreated or treated with ATRA, were cultured in the absence or presence of soluble purified TSP-1. Within 1-2 days of cell treatment with TSP-1, cell growth was inhibited (Figure 2A), and within 3 days, cell death was observed under the microscope, while it was not detected with ATRA alone. Cell morphology showed no cells with chromatin
condensation, typical of nuclear events of apoptosis. Instead, cells with apparent intact nucleus but damaged plasma membrane were observed (Figure 2B), and features of differentiated cells were not seen, as further analysed by NBT dye reduction assay (not shown). However, significant increase in membrane outside exposure of PS was detected at day 2 of both TSP-1 or ATRA treatment (Figure 2C). A further increase in PS exposure was observed upon ATRA/TSP-1 co-treatment essentially due to an additive effect. Cell necrosis was not detected at this time as determined by PI labeling, but was present at day 3 in cells cultured with TSP-1 but not ATRA alone (Figure 2C, up right rectangle, annexin V/PI positive cells).

Based on annexin V labelling (Figure 2D), cell death was found to be TSP-1 dose-dependent. However, morphologically, cell death was only observed with the highest doses of TSP-1 (0.075 and 0.1 µM). Combinatorial effect of 0.1 µM TSP-1 with ATRA caused a greater enhancement of cell death at day 4 (62% annexin V positive cells), when compared with TSP-1 alone (34% positive cells) or ATRA alone (11% positive cells). Quantification of cell growth also indicated a strong dose-dependent inhibition at all TSP-1 concentrations, in untreated (46-70% inhibition) or ATRA-treated cells (56-91% inhibition), when compared with control untreated cells (Figure 2D).

Given the known biological effect of TSP-1 on TGF-β activation, we next examined whether cell death induced in the presence of TSP-1 could be mediated by endogenous TGF-β. Co-treatment of NB4-LR1 cells for 4 days with ATRA and purified recombinant TGF-β (10 µg/mL) showed no increase in the rate of annexin V positive cells, when compared with ATRA alone (not shown).

Together, these results indicated that soluble TSP-1 action on NB4-LR1 leukemic immature blast cells leads to a composite biological response linking growth inhibition and cell death. These responses are increased in the presence of ATRA, independently of any maturation response.

### Involvement of the TSP-1 C-terminal domain in NB4-LR1 cell death induction

To further decipher the molecular mechanisms by which TSP-1 induced NB4-LR1 cell death, three recombinant fragments of TSP-1 (0.5-5 µM) containing the specific sequence for the N-terminal domain (NTSP-1), the type 1 repeat (3TSR) and the type 3 repeat/C-terminal domain (T3C1) of TSP-1, respectively (Figure 3A), were assayed on NB4-LR1 cells.
While only slight growth inhibition was demonstrated either with the NTSP-1 or the 3TSR fragment, a T3C1 dose-dependent strong inhibition of cell growth was measured in untreated cells (15-71% inhibition) and ATRA-treated cells (44-90% inhibition) (Figure 3B). Note that the T3C1/ATRA curve paralleled the T3C1 curve suggesting an additive effect of TSP-1 with ATRA. Thus, ATRA does not appear to make cells more sensitive to T3C1 action. Cell death was observed under the microscope in untreated or ATRA-treated cells within 3 days of culture with 2-5 μM of the T3C1 fragment, similar to what was observed with the entire TSP-1. When quantified, cell death by T3C1 alone was dose-dependent showing 28% of annexin V positive cells for the highest dose (5 μM) (Figure 3C), while no positive cell was detected with the NTSP-1 or the 3TSR fragment. By contrast to the growth inhibition response, the cell death response to T3C1 high dose (5 μM) was amplified by ATRA treatment (10% positive cells with ATRA alone versus 72% positive cells with 5 μM of T3C1). Time-course experiments with T3C1/ATRA co-treatment (not shown) indicated a slight increase of PS exposure at 24 h of culture that clearly rose at 48 h, and necrosis was detected at day 3 with T3C1, similarly to what was found with the entire TSP-1 (see Figure 2C).

Finally, T3C1 treatment also induced maturation-independent cell death, as evaluated through NBT dye reduction and MGG staining assays (not shown). In addition, condensed chromatin or fragmented nuclei were not observed upon Dapi staining of the cells (Figure 3D).

**Caspase-independent cell death induced by TSP-1 in NB4-LR1 cells**

We next investigated the involvement of caspase activity in cell death triggered by TSP-1 by comparison with staurosporine, a well known inducer of cell apoptosis. Experimental conditions for staurosporine treatment were chosen to induce a level of annexin V positive NB4-LR1 cells similar to that obtained with ATRA/T3C1 treatment. Whereas staurosporine-treated cells displayed about 50-fold increase in caspase-3-like activity, compared to untreated cells, ATRA/T3C1-treated cells showed no caspase-3 activity and neither did cells treated with the other TSP-1 recombinant fragments (Figure 4A). In addition, we found that the broad spectrum caspase inhibitor z-VAD-fmk, used at a non-cytotoxic dose (25 μM) which was able to inhibit both caspase activity and cell death in staurosporine-treated cells, did not prevent cell death upon ATRA/T3C1 treatment, as measured by annexin V positive detection and microscopic cell observation (not shown). Furthermore, caspase-3 as well as caspase-8, were found to remain in precursor form, i.e. not activated, when analysed
by Western blotting in ATRA/T3C1- or ATRA/TSP-1-treated NB4-LR1 cell lysates (Figure 4B). Additionally, no proteolysis of caspase substrates such as the DNA repair enzyme PARP or the anti-apoptotic protein Bcl-2 was found. Finally, analysis of low molecular weight DNA extracted from ATRA/T3C1- or ATRA/TSP-1-treated cells showed absence of characteristic ladder pattern resulting from internucleosomal cleavage of genomic DNA (Figure 4C).

**TSP-1 C-terminal domain induction of mitochondrial dysfunction with ROS production in the absence of cytochrome c and other mitochondrial intermembrane protein release**

Many types of cell death are accompanied by the permeabilization of mitochondrial membrane resulting from the opening of mitochondrial pores and disruption of the mitochondrial transmembrane potential ($\Delta\Psi_m$). As shown in Figure 5A, kinetics analysis showed a drop in $\Delta\Psi_m$ of T3C1- or ATRA/T3C1-treated cells, that was detected at day 2 and was more pronounced at day 3. Importantly, ATRA by itself had only very slight effect on $\Delta\Psi_m$ reduction, and staurosporine was found to induce a drop in $\Delta\Psi_m$ similar to that detected after T3C1 treatment (not shown).

Mitochondria are the major source of reactive oxygen species (ROS) which intracellular accumulation, upon $\Delta\Psi_m$ disruption, induces cellular oxidative stress that is implicated both in apoptosis and necrosis. Intracellular ROS production upon T3C1/ATRA treatment was detected at day 2 in close correlation with loss of $\Delta\Psi_m$, and was essentially due to the T3C1 fragment effect (Figure 5B). The lower level of ROS observed at day 3 was very likely resulting from ROS leakage due to membrane alterations. Pretreatment of the cells with selenium, known to reduce oxidative stress or Ebselen, a seleno-organic compound that converts H$_2$O$_2$ to H$_2$O, showed no inhibitory effect, indicating that mitochondria-derived ROS were likely not involved in the cell death process induced by T3C1 (Figure 5C).

We then examined whether cytochrome c was released from mitochondria which release process is independent of caspase activity and needs outer membrane permeabilization without mitochondrial depolarization. Whereas cytosolic accumulation of cytochrome c was clearly detected in staurosporine-treated NB4-LR1 cells through subcellular analysis, it was undetected in T3C1-treated cells (Figure 6A). Supporting this finding, immunofluorescence analysis indicated no change in cytochrome c subcellular localization after T3C1/ATRA treatment, as illustrated in Figure 6B, whereas staurosporine treatment clearly induced change
in subcellular distribution. Mitochondrial release of other soluble intermembrane proteins that occurs upon ΔΨm loss and contributes to both caspase-independent (endonuclease G, AIF, Omi/HtrA2) or caspase-dependent (Smac-Diablo) cell death processes was also examined by immunofluorescence (Figure 6B). Collectively, these data indicated no mitochondrial release upon T3C1 action.

**CD47 and αvβ3 integrin receptors mediate cell death induced by the TSP-1 C-terminal domain**

Given that the T3C1 recombinant fragment contains both TSP-1 binding sequences to CD47 and αvβ3 receptors (see Figure 3A), involvement of these receptors in the death responses of NB4-LR1 cells to T3C1 was next examined.

At first, immunofluorescence experiments were performed that showed colocalization of αvβ3 with CD47 in the membrane of untreated or ATRA-treated NB4-LR1 cells, as illustrated in Figure 7A(a) for ATRA-treated cells. Moreover, when treated for 48 hours with the T3C1 fragment, untreated cells (not shown) or ATRA-treated cells displayed colocalization of T3C1 with CD47 (Figure 7A,b) or αvβ3 (Figure 7A,c). Of note was the absence of TSP-1 signal when cells were not treated with T3C1 (not shown).

Further experiments were carried out using the CD47 agonist peptide RFYVVMWK (4N1-1) from the C-terminal domain of TSP-1, and the GRGDS peptide mimetic for the αvβ3-binding activity of the RGDA sequence identified from type 3-repeat domain of TSP-1. The VTCG peptide corresponding to a binding sequence of TSP-1 for CD36 was used as peptide control. The VTCG sequence is not present in the T3C1 fragment and CD36 is not expressed in NB4 cells (see Figures 1D and 3A). The 4N1-1, GRGDS and VTCG peptides alone were not found to have death induction effect on either untreated or ATRA-treated NB4-LR1 cells, through cell microscopic observation and PS positive detection (not shown). In addition, all the peptides did not display significant cell growth inhibitory effect. When tested for their ability to interfere with the effect of T3C1 by itself or combined with ATRA, the 4N1-1 and GRGDS peptides were found to exert dose-dependent inhibitory action on cell killing, as quantified by annexin V cell staining at day 4. The GRGDS peptide showed a stronger action than 4N1-1, while the VTCG control peptide had no significant effect. When added together, the 4N1-1 and GRGDS peptides showed a greater inhibitory effect that was found to be unchanged by ATRA treatment (Figure 7B). Furthermore, combinatorial effect of these two peptides measured at day 2 indicated substantial inhibition of NB4-LR1 cell death.
induced by T3C1, as defined by PS exposure, inhibition of ΔΨm disruption and ROS production (Figure 7C).
DISCUSSION

In this study, the acute promyelocytic leukemic cell line NB4-LR1, resistant to maturation by ATRA,\textsuperscript{28,29} was used to investigate whether TSP-1 plays a role in the maturation/apoptotic program triggered by ATRA. Indeed, in contrast to the features of the parental NB4 cell line, this subcloned cell line expressed no endogenous TSP-1 in response to ATRA, but constitutively expressed potential membrane receptors for the matrix protein. Therefore, it constituted an appropriate cell model to check whether TSP-1 membrane signaling acts to trigger antitumor responses. We show here that membrane signals by TSP-1 induce strong inhibition of cell growth and caspase-independent cell death in acute promyelocytic leukemia that is not subsequent to cell maturation. Cell death is characterized by an atypical mitochondrial dysfunction, and is mediated by ligation of the TSP-1 membrane receptors CD47 and αvβ3.

Of note, the reduction of tumor cell growth \textit{in vivo} by TSP-1 is a well documented consequence of its inhibitory effect on tumor angiogenesis by acting on endothelial cell proliferation and apoptosis, and/or by activating TGF-β.\textsuperscript{11-13,15,31} Indeed, activated TGF-β has been shown to inhibit tumor cell growth and induce tumor cell apoptosis. These activities are increased by TSP-1 and mediated by specific sequences present in the type 1 repeat domain of TSP-1.\textsuperscript{11,31} Importantly, in this report, we show that TGF-β did not induce NB4-LR1 cell death. Furthermore, we gained evidence that TSP-1 action on NB4-LR1 cells was mapped to its C-terminal domain that does not contain activation sequence for TGF-β.

Anti-tumor effects of TSP-1 were mapped to its type 3-repeat /C-terminal domain, using recombinant fragments of the protein. It was observed that the T3C1 recombinant domain was less active than the intact protein, i.e., about 40 times less at the respective doses of 3 µM for T3C1 and 0.075 µM for TSP-1. Interestingly, this was also observed in a study by Miao \textit{et al.} (2001)\textsuperscript{31} on the tumor growth inhibitory function of the TSP-1 type 1 repeat recombinant protein. This may mean that the two distinct anti-tumor domains identified independently by us and by Miao \textit{et al.} each cooperate with another domain of the native TSP-1 that plays an enhancer role. Alternatively, these data may mean that the effects of these domains are partly dependent upon the trimer structure of the TSP-1 molecule which can not be achieved by the recombinant fragments. In keeping with this notion, note that although substantial growth inhibition was induced by either the entire TSP-1 or the T3C1 recombinant fragment at low concentration, only high doses could induce cell death based on
morphological or biochemical criteria (see Figures 2 and 3). This could indicate that in the leukemic NB4-LR1 cell model, TSP-1 exerts dual effects relevant to differential or composite membrane signals. Further studies are needed to determine whether TSP-1 signals for inhibition of cell proliferation could occur independently of its action leading to cell death.

Cell death, as induced by the soluble form of T3C1 in NB4-LR1 is representative of a new form of caspase-independent death pathway, being similar to that recently described in B-cell chronic lymphocytic leukemia or normal B cells,\textsuperscript{19} cytotoxic T or breast cancer cells\textsuperscript{20,21,48} upon exposure to CD47 agonists. In the latter case, death triggered by TSP-1, RFYVVM peptide or monoclonal antibodies to CD47 is accompanied with PS membrane outside exposure and loss of mitochondrial transmembrane potential ($\Delta\Psi_m$) which are typical of cell apoptosis, but not by cytochrome $c$ or AIF release, or DNA fragmentation. However, at variance with our results in NB4-LR1 cells, cell death rapidly proceeded, within 18 hours, and no plasma membrane breakdown was observed\textsuperscript{19,49}. Importantly, antibodies to CD47 or RFYVVM peptide were either more potent than TSP-1 in inducing cell death\textsuperscript{21,49} or were effective only in immobilized form\textsuperscript{19,20}. We therefore used the antibody to CD47 B6H12 or the RFYVVM 4N1K peptide, as used in these studies, and found them to have no death induction effect on NB4-LR1 cells when used either in soluble or immobilized form (A.S. and V.D., unpublished data, April, 2003). These apparently contradictory findings would mean that other epitopes on CD47 were active in NB4-LR1, as in fact not all the CD47 agonists were able to activate cell death\textsuperscript{48}. Alternatively, this could be relevant to death signals by T3C1 that required CD47/$\alpha\alpha\beta$3 integrin double signaling in NB4-LR1 leukemic cells to take effect. In this regard, NB4-LR1 cell death triggered by this domain of TSP-1 was not inhibited or delayed when cells were treated with a cAMP analog (A.S., M.B.S. and V.D., unpublished data, October, 2003), indicating that TSP-1 action does not involve regulation of cAMP intracellular level and subsequent signal effects, as has been reported for CD47-dependent killing of cytotoxic T or breast cancer cells\textsuperscript{21}.

The mitochondrial dysfunction induced by action of the C-terminal domain of TSP-1 on NB4-LR1 did not lead to cytochrome $c$ or AIF release, as shown in other studies reporting on death effects by CD47 agonists\textsuperscript{20,21}. We extended these studies to other death-promoting factors and found no release of endonuclease G and Omi/HtrA2, or Smac-Diablo. Absence of AIF or endonuclease G leakage is consistent with the absence of chromatin condensation and nuclear fragmentation (see Figure 3D).\textsuperscript{45,46} On the other hand, mitochondrial ROS accumulation was detected in cells, which function as second messengers and regulating molecules for cell death.\textsuperscript{37} However, through use of ROS scavenger or oxidative stress...
inhibitor, we did not find ROS to be involved in cell damage. This result could be related to the absence of cytochrome c release which preserves normal redox state, so that only moderate levels of ROS are produced. It remains to be determined whether a mitochondrial dysfunction featured by a partial (or differential) membrane permeabilization defect could be one rate-limiting step of this TSP-1 dependent cell death.

Finally, many forms of caspase-independent programmed cell death are mediated by other proteases such as cathepsins, calpains, serine proteases that can cleave at least some of the caspase substrates. However, this alternative cell death pathway generally produces mitochondrial intermembrane protein leakage and leads to incomplete chromatin condensation. We investigated the involvement of calpain activity since among numerous calpain substrates are cytoskeletal and membrane-associated proteins suggesting that calpains are important in destruction of cellular architecture. However, no inhibitory effect was exerted by the calpain inhibitors PD150606 and calpeptin on NB4-LR1 cell death induction by the T3C1 (A.S. and V.D., unpublished data, September, 2003).

This is the first report showing the ability of soluble source of the matrix protein TSP-1 to trigger a novel non-caspase cell death pathway in ATRA resistant acute promyelocytic leukemia. This activity is mapped to the type 3-repeat/C-terminal domain of the molecule. Cell death is mediated by the TSP-1 membrane receptors CD47 and the αvβ3 integrin, and is accompanied with ΔΨm collapse that does not result in the release of well-known death-promoting proteins, such as the non-caspase apoptosis regulators AIF, endonuclease G or Omi/HtrA2. This kind of cell death may resemble in some way necrosis-like programmed cell death in which increased PS membrane exposure is detected in the absence of chromatin condensation and occurs before cell damage. However, the level of necrosis is low and not consistent with the apparent substantial cell damage, suggesting it could not be a major event, and cell death might actually be related to an alternative, unidentified, process.

Retinoids have important physiological functions in regulating cell growth, differentiation and death. Upon its induction and secretion following ATRA treatment, TSP-1 could mediate growth inhibition and death responses. Therefore, TSP-1 signalling could well contribute to the therapeutic response of APL cells to retinoids. A pharmacological ligandation of TSP-1 receptors on cells for which ATRA-dependent TSP-1 induction is deficient could be envisioned. Alternatively, identification of the molecular targets of TSP-1 signalling (receptors, signal transduction enzymes) the most appropriate to a pharmacological exploitation will need specific investigation. Mediators of cell death program execution are under current investigation, and their identification would contribute to elaborate new targets.
in the field of cancer, especially to sensitize tumor cells to death and to overcome resistance to drug therapy.

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REFERENCES


42. Defacque H, Piquemal D, Basset A, Marti J, Commes T. Transforming growth factor-
beta1 is an autocrine mediator of U937 cell growth arrest and differentiation induced

43. Lee YC, Tang YC, Chen YH, Wong CM, Tsou AP. Selenite-induced survival of
HuH7 hepatoma cells involves activation of focal adhesion kinase-
phosphatidylinositol 3-kinase-Akt pathway and Rac1. J Biol Chem. 2003;278:39615-
39624.

44. Von Ahsen O, Waterhouse NJ, Kuwana T, Newmeyer DD, Green DR. The 'harmless'

45. van Gurp M, Festjens N, van Loo G, Saelens X, Vandenabeele P. Mitochondrial
intermembrane proteins in cell death. Biochem Biophys Res Commun. 2003;304:487-
497.

46. Lorenzo HK, Susin SA. Mitochondrial effectors in caspase-independent cell death.

47. Gao AG, Frazier WA. Identification of a receptor candidate for the carboxyl-terminal


49. Manna PP, Frazier WA. The mechanism of CD47-dependent killing of T cells:
heterotrimeric Gi-dependent inhibition of protein kinase A. J Immunol.
2003;170:3544-3553.

50. Leist M, Jäättelä M. Four deaths and a funeral: from caspases to alternative

51. Mathiasen IS, Jäättelä M. Triggering caspase-independent cell death to combat cancer.

52. Foghsgaard L, Wissing D, Mauch D, et al. Cathepsin B acts as a dominant execution
2001;153:999-1010.

53. Mathiasen IS, Sergeev IN, Bastholm L, Elling F, Norman AW, Jäättelä M. Calcium
and calpain as key mediators of apoptosis-like death induced by vitamin D compounds

caspase inhibition in vitro with pan-caspase inhibitors or in vivo by XIAP or Survivin
does not affect cell survival or prognosis. Blood. 2003;102:4179-4186.

55. Egger L, Schneider J, Rheme C, Tapernoux M, Hacki J, Borner C. Serine proteases
mediate apoptosis-like cell death and phagocytosis under caspase-inhibiting

FIGURE LEGENDS

**Figure 1. TSP-1 and its membrane receptors expression in NB4 and NB4-LR1 cell lines.** Maturation-sensitive NB4 cells and maturation-resistant NB4-LR1 cells were cultured for 4 days either untreated or treated with 1nM-1µM ATRA or 1µM ATRA, and processed for examination of TSP-1 transcript (A), (C) and protein level expression (B), or TSP-1 receptors membrane expression (D). (A) Cell maturation was monitored by the NBT dye reduction assay. Total RNA (20 µg) were hybridized with a 32P-labeled fragment of human cDNA for TSP-1. Equivalent loading and integrity of each RNA preparation were evaluated by ethidium bromide staining of the 28S rRNA bands. (B) Triton X-100 cell lysates were prepared from maturation-sensitive NB4 cells and processed for immunoprecipitation and Western blotting for analysis of TSP-1 protein expression, as described in the Methods section. Cell secretion of TSP-1 into the extracellular medium was similarly analysed upon Triton X-100 solubilization of the corresponding culture media. Here, Triton X-100 cell extract and Triton X-100-solubilised culture medium coming from 4x10^6 NB4 cells and 1.2x10^5 NB4 cells, respectively, were analysed. (D) Untreated or ATRA(1 µM)-treated NB4 and NB4-LR1 cells (2x10^5) were incubated with monoclonal antibodies to CD36, CD47, the β1 or β3 integrin subunit or isotype-match control antibody (20 µg/mL). Alexa Fluor 488-conjugated anti-mouse IgG were used as secondary antibody (1:700), and cells were analysed by flow cytometry.

**Figure 2. NB4-LR1 cell death induction by TSP-1.** (A) NB4-LR1 cells untreated or treated with ATRA (1 µM) with or without TSP-1 (0.1 µM) were cultured in microwell culture plate for 4 days. Cell growth was quantified by cell counting in a Malassez hemocytometer. (B) Morphology of MGG-stained untreated or ATRA-treated cells after 4 days of culture in the absence or the presence of purified TSP-1 (0.1 µM) showed cells with no chromatin condensation but damaged plasma membrane when cultured with TSP-1 with or without ATRA (arrows). (C) Flow cytometry analysis of FITC-annexin V and PI-stained untreated or ATRA(1 µM)-treated NB4-LR1 cells cultured with or without TSP-1 (0.1 µM) for 3 days. Results, expressed as percent of positive cells for annexin V labeling, are as follows: at day 2: untreated 3%; TSP-1 10%; ATRA 8%; ATRA+TSP-1 22% and at day 3: untreated 6%; TSP-1 19%; ATRA 15%; ATRA+TSP-1 42%. (D) NB4-
LR1 cells untreated or treated with ATRA (1 μM) were cultured for 4 days with or without increasing concentrations of purified TSP-1 (0.025-0.1 μM). Cell growth was expressed as percent of growth compared to untreated cells (●), and cell death, as quantified by annexin V labeling (▲), was expressed as percent of positive cell detection upon flow cytometry analysis: --- untreated cells, — ATRA-treated cells. Results are expressed as mean ± SEM of three experiments.

Figure 3. Induction of NB4-LR1 cell death by the C-terminal domain of TSP-1.
(A), Shematic representation of TSP-1 structural and functional domains. TSP-1 is a disulfide-bonded trimeric protein composed of three identical subunits with a molecular mass of about 140 kDa each. Each subunit comprises multiple domains: a amino-terminal domain, a procollagen homology region (PC), three types of repeated sequence motifs, designated type 1 (◇), type 2 (○) and type 3 (□) repeat and a carboxy-terminal domain. The recombinant fragments used in this study are identified by their amino-terminal and carboxy-terminal amino acids. VTCG, RGDA and RFYVVMWK peptide sequences identified from TSP-1 for its respective binding to CD36, β3 integrin and CD47 membrane receptors are indicated. (B), (C) NB4-LR1 cells untreated (---) or treated (—) with ATRA (1 μM) were cultured for 4 days in the absence or the presence of increasing doses (0.5-5 μM) of TSP-1 recombinant fragments corresponding to the N-terminal (NTSP-1; ▲), the type 1 repeat (3TSR; ■) or the type 3 repeat/C-terminal domain (T3C1; ●) of the molecule, respectively. Cell growth of NB4-LR1 cells was expressed as percent of untreated cells growth (B), and cell death, as quantified by annexin V labeling, was expressed as percent of positive cell detection upon flow cytometry analysis (C). Results are expressed as mean ± SEM of three experiments. (D) Nucleus morphology of NB4-LR1 cells untreated or treated with ATRA with or without the T3C1 TSP-1 recombinant fragment (3 µM) or the entire TSP-1 (0.075 µM), as viewed under the fluorescence microscope after Dapi staining.

Figure 4. Caspase-independent cell death induced by TSP-1 in NB4-LR1 cells
NB4-LR1 cells untreated or treated with ATRA (1 μM) were cultured for 4 days with or without the NTSP-1, 3TSR or T3C1 fragment (3 µM), or the entire purified TSP-1 (0.075 μM). NB4-LR1 cells were also treated with staurosporine (STP, 5 μM) for 3.5 h. (A) Caspase-3-like activity was determined in cell extracts by means of a colorimetric assay of the DEVD-pNa substrate hydrolysis. NB4-LR1 cells treated with STP with or without the broad-
spectrum caspase inhibitor z-VAD-fmk (25 µM) were used as positive control for the detection of caspase-3-like activity. (B) SDS-cell lysates (about 10 µg proteins corresponding to 10^5 cells) prepared from NB4-LR1 cells, treated as described above, were processed for Western blotting for analysis, under reducing conditions, of caspase-3 or caspase-8 activation, through cleavage of their precursor forms, cleavage analysis of the caspase substrate PARP or the anti-apoptotic protein Bcl-2. Polyclonal antibodies specific to caspase-3 or caspase-8 (1:500), monoclonal antibodies specific to PARP (1:200) or Bcl-2 (1:1000) were respectively used. (C) Genomic DNA from NB4-LR1 cells (10^6) treated as described above was extracted, as described in the Methods section, and electrophoresed on 2% agarose gel then analysed upon ethidium bromide visualization.

**Figure 5. Alteration of the mitochondrial transmembrane potential and ROS production induced by the C-terminal domain of TSP-1.**

(A), (B) NB4-LR1 cells untreated or treated with ATRA (1µM) were cultured for 3 days with or without the TSP-1 T3C1 recombinant fragment (3 µM). At each day of the treatment, cells were stained with the lipophilic fluorochrome DiOC_6(3) (A), or the oxidant sensitive dye DCFH-DA (B), as described in the Methods section, then analysed by flow cytometry. (C), left panel, cells were preincubated for 2 h with non-cytotoxic doses of Ebselen (Eb, 10 µM and 50 µM), then treated or not with H_2O_2 (50 µM, 6 h) for detection of ROS production upon cell staining with DCFH-DA. In right panel, untreated or ATRA (1µM)-treated cells were cultured for 3 days with T3C1 with or without selenium (Se, 0.06 µM) or Ebselen (Eb, 50 µM), and cell death, as quantified by annexin V and DiOC_6(3) staining, was analysed at day 3.

**Figure 6. Absence of mitochondrial intermembrane protein release upon NB4-LR1 cell treatment with the C-terminal domain of TSP-1.**

NB4-LR1 cells were either treated with ATRA (1µM) for 3 days with or without the T3C1 recombinant fragment (3 µM), or treated with staurosporine (STP, 5 µM) for 3.5 h. (A) Western blot analysis, under reducing conditions, of total cell extract and cytosolic S100 fraction prepared from 5x10^6 cells, as described in the Methods section, for detection of cytochrome c, using monoclonal antibody (1:100), in the respective fraction each corresponding to 10^5 cells. (B) Immunofluorescence analysis of mitochondria protein in NB4-LR1 cells. Upon fixation by 2% PFA and permeabilization with 0.1% saponin, as described in the Methods section, cells were incubated with antibodies to cytochrome c (1:200), AIF
(1:500), Smac/Diablo (1:150), Omi/HtrA2 (1:150) or endonuclease G (6 µg/mL). Alexa Fluor 594-conjugated anti-rabbit and Alexa Fluor 488-conjugated anti-mouse IgG were used as secondary antibodies. Vectashield containing Dapi for nucleus staining was used as mounting medium for confocal microscopy analysis.

Figure 7. Involvement of CD47 and αvβ3 integrin receptors in mediating cell death induced by the C-terminal domain of TSP-1.

(A) NB4-LR1 cells were treated with ATRA (1µM) for 2 days with or without the T3C1 recombinant fragment (3 µM), then fixed by the addition of 2% PFA. (a) Cells cultured without T3C1 were successively incubated with mouse monoclonal antibody to CD47 (20 µg/mL) and polyclonal antiserum to the αv integrin subunit (1:50). (b), (c) ATRA-treated cells cultured with the T3C1 fragment were successively incubated with rabbit antiserum to TSP-1 (R1, 1:50) and mouse monoclonal antibody to αv or CD47 (20 µg/mL). Alexa Fluor 594-conjugated anti-rabbit and Alexa Fluor 488-conjugated anti-mouse IgG were used as secondary antibodies. Vectashield containing Dapi for nucleus staining was used as mounting medium for confocal microscopy analysis. Yellow colour in the merge confocal images indicate the superimposition of the two fluorochromes. (B), (C) NB4-LR1 cells, untreated or treated with ATRA (1 µM), were incubated for 2 h with 4N1-1 (25-200 µM), GRGDS (0.1-1 mM), 4N1-1 (100 µM) and GRGDS (100 µM), or the VTCG (1 mM) peptide control, then cultured for 4 days (B) or 2 days (C) with the T3C1 recombinant fragment (3 µM). (B) At 4 days of the treatment, cell death was quantified by annexin V labeling and expressed as percent of positive cell detection upon flow cytometry analysis. Results are expressed as mean ± SEM of three experiments. (C) At 2 days of the treatment, cell death induction was monitored by annexin V (upper panel), DiOC<sub>6</sub>(3) (middle panel) or DCFH-DA (lower panel) staining for detection of PS membrane outside exposure, mitochondrial transmembrane potential alteration and ROS production, respectively. Results are expressed as mean ± SEM of two experiments.
A

ATRA dose- and time-course responses (NB4 cells)

TSP-1 mRNA expression (Northern Blot)

NBT positive cells (%) (NBT reduction assay)

B

NB4 cells (ATRA 1 μM)

Culture media

Cell extracts

TSP-1 protein expression

C

NB4-LR1 cells (ATRA 1 μM)

rRNA

hrs 0 24 48 72 96

TSP-1 mRNA expression (Northern Blot)

D

CD36  CD47  β1  β3

NB4

NB4-LR1

isotype control untreated ATRA-treated

Saumet et al., Figure 1
Saumet et al., Figure 2
Saumet et al., Figure 3
Saumet et al., Figure 4
Saumet et al., Figure 5
Saumet et al., Figure 6
Saumet et al., Figure 7
Type 3-repeat/C-terminal domain of thrombospondin-1 triggers caspase-independent cell death through CD47/αvβ3 in promyelocytic leukemia NB4 cells

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