Suramin Rapidly Alters Cellular Tyrosine Phosphorylation in Prostate Cancer Cell Lines

Oliver Sartor, Catherine A. McLellan, Charles E. Myers, and Markus M. Borner
Clinical Pharmacology Branch, Division of Cancer Treatment, National Cancer Institute, Bethesda, Maryland 20892

Abstract

Suramin, a synthetic polysulfonated anionic compound, is known to abrogate the activity of a variety of growth factors that serve as ligands for receptor-class protein-tyrosine kinases. Based on this information, we initially hypothesized that suramin treatment would be associated with decreased tyrosine phosphorylation. Upon testing this hypothesis in prostate cancer cell lines, we found that the most conspicuous effect of suramin was to increase the tyrosine phosphorylation of several distinct proteins. Further analyses indicate that suramin-induced increases in tyrosine phosphorylation represent a generalized, but not universal, phenomenon found in cell lines derived from a variety of human tissues. These rapid and specific suramin-induced alterations represent a novel finding for a non-polypeptide pharmaceutical agent and question the hypothesis that suramin exerts its antitumor action simply by abrogation of growth factor action. (J. Clin. Invest. 1992. 90:2166–2174.)

Key words: epithelial cell lines • polyanions • prostate • suramin • tyrosine phosphorylation

Introduction

Suramin is the hexasodium salt of carbonyl bis[8-(3-aminobenzamido) 4-methylbenzamido]-naphthalene-4,6,8 trisulfonate (Fig. 1). A synthetic compound, suramin was originally synthesized in 1916 by Bayer AG as part of an ongoing search for antitrypanosomal chemotherapeutic agents. After demonstration of activity against trypanosomes in model systems, and the subsequent discovery of its activity against onchocerciasis, suramin has been used to treat parasite-infected patients throughout the world since the 1920s (1).

In 1984 suramin was reported to dissociate platelet-derived growth factor (PDGF)1 from its receptor and block the binding of PDGF to cell membranes (2). The mechanism whereby suramin inhibits PDGF binding to its receptor is secondary to a direct attachment of suramin to the PDGF molecule (3). In v-sis transformed fibroblasts, where activation of the PDGF receptor is a requisite for transformation, suramin induces phenotypic reversion (4) and decreases the extent of cellular tyrosine phosphorylation (5). Subsequent studies indicate that suramin can inhibit the activity of a number of peptides known to stimulate cellular tyrosine phosphorylation including basic fibroblast growth factor (bFGF) (6, 7), the product of the hst/K-sgf gene (8), interleukin 2 (9), and scatter factor (also known as hepatocyte growth factor) (10).

Beginning in 1987, prompted by reports that suramin reversed a variety of growth factor-induced actions, suramin was administered in clinical trials and found to have antitumor activity (11). More recently, in both model systems (12–14) and in clinical trials (15, 16), suramin has been demonstrated to have antitumor activity against prostatic cancer.

Given the fact that suramin antagonizes the activity of a variety of growth factors known to increase cellular tyrosine phosphorylation (see reference 17), and given the fact that suramin has antitumor activity against cell lines derived from prostatic cancer patients (12, 13), we initially hypothesized that suramin treatment would decrease tyrosine phosphorylation in prostate cancer cells. Upon testing this hypothesis, we were surprised to find that the most conspicuous effect of suramin was a rapid (within seconds) and marked increase in the tyrosine phosphorylation of several cellular proteins.

Methods

Cell culture. The androgen-insensitive PC3 and DU145 and the androgen-sensitive LNCaP prostate cancer cell lines were obtained from the American Type Culture Collection (ATCC). PC3M, a highly metastatic variant of PC3 cells, was obtained from Dr. J. M. Kozlowski, Northwestern Medical School, Evanston, IL. The colon cell lines HT29, COLO 205, and SW620, and the leukemia cell lines (CEM and HL60) were obtained from ATCC. The colon cell lines NCI-H630 (designated 630) and NCI-H716 (designated 716), originally described by Dr. A. Gazdar and colleagues (18), were obtained from the laboratory of Dr. C. Allegra. The myeloma cell lines H929 and U266 were obtained from Dr. R. Nordan, National Cancer Institute (NCI). The Burkitt’s lymphoma cell lines, Dufaw and Wilson, were obtained from Dr. I. Magrath, NCI. The lung cancer cell lines NCI-H82, NCI-N417, and NCI-H226, originally derived from the laboratories of Drs. D. Carney and A. Gazdar (19, 20), were obtained from Dr. E. Sausville, NCI. The breast cancer cell lines H5578, T47-D, MDA-231, and ZZ-75 as well as the HeLa cell line were obtained from Dr. K. Cowan, NCI. The gastric cell lines TMK-1, A2521, NUGC4, MKN74, and MKN28 were obtained from Dr. T. Toko, NCI. The medulloblastoma D283Med, the neuroepithelioma cell lines CHP-100, SKNMC, and TC32, and the melanoma cell lines RPMI-7951 and HS69ST were obtained from Dr. L. Whitescnt. NCI. The rhabdomyosarcoma cell line was obtained from Dr. L. Helman, NCI. All colon, prostate, breast, gastric, myeloma, lung, leukemia, and rhabdomyosarcoma cell lines were routinely maintained in RPMI 1640 supplemented with 10–20% FCS, 2 mM glutamine, 100 U/ml penicillin, and 100 μg/ml streptomycin. Melanoma cells were maintained in ATCC-advised media. NIH 3T3 cells were obtained from the laboratory of Dr. S. Aaronson, NCI, and cultured in DME supplemented with 10% calf serum.

1. Abbreviations used in this paper: FGF, fibroblast growth factor; GAP, ras GTPase activating protein; LSB, Laemmli sample buffer; NCI, National Cancer Institute; PDGF, platelet-derived growth factor; PLC-γ, phospholipase C-γ; TGF-α, transforming growth factor; TNESV, Tris 50 mM, pH 7.6, NP-40 1%, EDTA 2 mM, NaCl, 100 mM, and vanadate 1 mM.

Address reprint requests to Dr. Sartor, Clinical Pharmacology Branch, Building 10, Room 12N226, National Cancer Institute, Bethesda, MD 20892.

Received for publication 16 January 1992 and in revised form 24 July 1992.

The Journal of Clinical Investigation, Inc.
Volume 90, December 1992, 2166–2174
Media and all media supplements (except calf serum and human-derived blood-products) were obtained from Life Technologies Inc. (Gaithersburg, MD). Calf serum was obtained from ABI, Inc. (Columbia, MD). Fresh normal human serum was donated by one of the authors. Human platelet-poor plasma was obtained from the Division of Transfusion Medicine, NIH Clinical Center.

**Immunoblotting.** Cells were routinely lysed on ice with a buffer containing 50 mM Tris, pH 7.6, 1% NP-40, 2 mM EDTA, 100 mM NaCl, and 1 mM vanadate (TNEV) freshly supplemented with 1 mM phenylmethylsulfonyl fluoride (PMSF), 20 µg/ml aprotinin, and 20 µg/ml leupeptin. In other cases (where designated), cells were lysed with a buffer containing 1% Triton X-100, 0.5% sodium deoxycholate, 100 mM NaCl, 42 mM Na2HPO4, and 8 mM NaH2PO4 supplemented with protease inhibitors as described above. Protease inhibitors were obtained from Boeringer-Mannheim (Indianapolis, IN), detergents from Sigma Chemical Co. (St. Louis, MO). After clearing cellular debris by centrifugation at 15,000 g for 20 min at 4°C, protein concentrations were determined by the Bradford method using reagents supplied by Bio-Rad Laboratories (Richmond, CA). After adding Laemmlli sample buffer (LSB), samples were heated to 100°C and 40–60 µg of protein were loaded per lane on 8% SDS-PAGE gels prepared according to the method of Laemmlli (21). Bethesda Research Laboratories (Gaithersburg, MD) size markers were used for determination of protein size. After transfer to nitrocellulose, filters were blotted as previously described (22).

**Preparation of membrane-enriched and cytosolic cell fractions.** Cells were lysed in ice-cold hypotonic lysing buffer (20 mM pH 7.6 Hepes with 1 mM EDTA, 1 mM vanadate, 1 mM PMSF, 20 µg/ml leupeptin, and 20 µg/ml aprotinin) and all subsequent steps were performed at 4°C. Cells were scraped from the tissue culture plate, repetitively passed through a 23-gauge needle, and nuclei were pelleted for 10 minutes at 1,000 g. The supernatant was then removed and centrifuged at 176,000 g for 1 h. The remaining supernatant (cytosolic fraction) and the TNEV extractable portions of the pellet (membrane fraction) were size-fractionated by SDS-PAGE after boiling in LSB.

**Antibodies.** The anti-phosphotyrosine monoclonal antibody 4G10, the anti-human fibroblast growth factor (FGF) receptor (f/g) monoclonal antibody, the anti-ras GTPase-activating protein (anti-GAP) antisera, and the monoclonal anti-phospholipase C-γ (PLC-γ) were obtained from Upstate Biotechnology Incorporated (Lake Placid, NY). The anti-phosphotyrosine monoclonal antibody PY20 was obtained from ICN Biomedicals Inc. (Costa Mesa, CA). The monoclonal antibodies 3B12, 2B12, 2A7, and 4F4, recognizing (respectively) p80/85 kD, 120 kD, 125 kD, and 130 kD proteins that are tyrosine-phosphorylated in chicken embryo fibroblasts expressing activated versions of the c-src kinase were kindly provided by Dr. T. Parsons, University of Virginia (23). The monoclonal antibodies ASC9 and MT-18, recognizing the 80-kD component of the IL-6 receptor were provided by Dr. R. Nordan, NCI. The anti-met antisera was provided by Dr. D. Botaro, NCI.

**Immunoprecipitations.** Cellular lysates were prepared with TNEV supplemented with protease-inhibitors as described above. After equalization of protein between samples (300–600 µg of protein per sample), antibodies were added for 1–4 h at 4°C. Rabbit antibodies were immunoprecipitated with the aid of protein A-Sepharose beads (Pharmacia, Inc., Piscataway, NJ). Mouse monoclonal antibodies were immunoprecipitated with the aid of protein A-Sepharose beads coated with affinity-purified rabbit anti-mouse antisera (Cappel Laboratories, West Chester, PA). Beads were then washed twice with lysing buffer before boiling in LSB and fractionation by SDS-PAGE.

Immune-complex kinase assays and V8 digests. Immune-complex kinase assays and proteolytic digests with V8 protease (Worthington Biochemical Corp., Freehold, NJ) were performed exactly as previously described (22).

**Metabolic labeling.** Metabolic labeling with [32P]orthophosphate (New England Nuclear, Boston, MA) was performed by 2-h labeling (0.5 mCi/ml) at 37°C in phosphate-free RPMI (Gibco, Gaithersburg, MD).

**Radioimmunoassay of transforming growth factor-α (TGF-α).** Conditioned medium was collected from PC3 cells, cleared of debris by centrifugation, and assayed for TGF-α with an RIA kit obtained from Biomedical Technologies, Inc. (Stoughton, MA).

**Determination of suramin concentration.** Suramin levels were determined by an HPLC based assay as previously described (24).

**Chemicals.** Suramin and pentosan polysulfate were obtained from the Division of Cancer Treatment, NCI. Heparin from beef lung was obtained from The Upjohn Co. (Kalamazoo, MI), and heparan sulfate from the Sigma Chemical Co. Protamine sulfate was obtained from Elkins-Sinn, Inc. (Cherry Hill, NJ).

**Results.** To determine the effects of suramin on tyrosine phosphorylation in an androgen-insensitive prostate cancer cell line, lysates derived from PC3 cells were examined by anti-phosphotyrosine immunoblotting (see Fig. 2 A). PC3 cells express several prominent tyrosine phosphorylated proteins of ~60 and 120 kD as detected by the 4G10 monoclonal antibody. Within seconds after treatment of cells with 300 µg/ml suramin, a suramin concentration utilized in the clinical setting (15, 16, 25), several new tyrosine-phosphorylated bands were readily detected in the cellular lysates and (in some cases) the p120 band became less prominent. The most conspicuous of the newly tyrosine-phosphorylated proteins had molecular masses of ~75 and 135 kD. To verify the specificity of the 4G10 antibody, phenylphosphate (10 mM) was preincubated with 4G10 before immunoblotting. Under these conditions, no bands were detected in suramin-treated or untreated PC3 cells (data not shown). To determine whether these bands were detected by a second and distinct anti-phosphotyrosine antibody, we immunoblotted these same lysates using monoclonal antibody PY20 (see Fig. 2 B). In untreated PC3 cells, the 120-kD band was readily detectable after immunoblotting with PY20. In contrast, the 60-kD band was much less prominent in blots with PY20 as compared to 4G10. After suramin addition, the prominent tyrosine phosphorylated bands at ~75 and 135 kD

---

**Figure 1.** Structure of suramin. Note the six sulfonic acid groups which make this compound a highly charged polyanion.
To ascertain whether the suramin-induced changes in tyrosine phosphorylation were indicative of broader changes in cellular phosphorylation, detergent lysates derived from metabolically labeled ([32P]orthophosphate) PC3 cells were examined by SDS-PAGE after suramin treatment. No differences were noted in the phosphorylated protein pattern detected in these assays, indicating that the suramin-induced changes in phosphotyrosine were not the result of global changes in phosphorylation (data not shown).

The rapidity of the changes in tyrosine phosphorylation after suramin treatment suggested that suramin interacts with a component of the culture media (such as serum) and/or a cellular component in direct contact with the extracellular environment. Because the initial experiments were performed in the presence of 10% fetal calf serum, we questioned whether or not serum would affect the suramin-induced changes in tyrosine phosphorylation. Thus, PC3 cells were washed twice with PBS and placed in RPMI without serum supplementation ("0"), RPMI plus 10% human platelet-poor plasma ("HP"), or RPMI plus 10% human serum ("HS"). Regardless of the media employed, 300 µg/ml of suramin induced a rapid increase in the phosphotyrosine content of the 75- and 135-kD bands (Fig. 3). In other experiments, similar suramin-induced changes were noted for cells cultured in serum-free media for 72 h (data not shown). Thus, suramin-induced changes are not directly dependent upon interactions between suramin and serum components. Because of the rapidity of the changes, and because the entry of suramin into cells is limited by size and charge, these studies suggest that rapid suramin-induced increases in tyrosine phosphorylation are the result of an interaction between suramin and a component of the cell (such as the plasma membrane) readily accessible to the extracellular environment.

To determine if the tyrosine phosphorylated proteins detected after suramin treatment would segregate with membrane-enriched or cytosolic cell fractions, we performed immu-

---

**Figure 2.** Suramin rapidly stimulates tyrosine phosphorylation in the prostate cancer cell line PC3. Suramin at a concentration of 0 (−) or 300 (+) µg/ml was added to the culture medium (RPMI 1640 plus 10% FCS) and cells were lysed 0.5, 1, 2, 4, or 8 min later. Samples were size-fractionated by SDS-PAGE, transferred to nitrocellulose, and immunoblotted with (A) the anti-phosphotyrosine monoclonal antibody 4G10 or (B) the anti-phosphotyrosine monoclonal antibody PY20. The electrophoretic mobility of molecular size markers is shown in kilodaltons.

---

**Figure 3.** Suramin stimulation of tyrosine phosphorylation is not dependent upon the presence of serum or a specific culture media. Suramin at a concentration of 0 (−) or 300 (+) µg/ml was added to various culture medias and PC3 cells were lysed 10 min later. After fractionation by SDS-PAGE and transfer to nitrocellulose, samples were blotted with monoclonal antibody 4G10. Suramin was added to cells bathed in RPMI without serum (0), RPMI plus 10% human platelet-poor plasma (HP), or RPMI plus 10% human serum (HS). The electrophoretic mobility of molecular size markers is shown in kilodaltons.
noblots on proteins derived from each of these fractions after hypotonic lysis and differential centrifugation. From these data (not shown), it was clear that both the 75-kD band (prominent) and the 135-kD band (far less prominent) were present within the membrane fraction. Neither of these bands were present in the cytosolic fraction.

To determine the duration of the suramin response, PC3 cells were treated with 300 µg/ml suramin for varying periods of time. As shown in Fig. 4 A, the tyrosine phosphorylated bands of 75 and 135 kD were readily detected between 10 and 40 min after suramin addition. However, 160 min later, the tyrosine phosphorylation pattern was not substantially different from baseline. Also of note, the prominence of the p120 band varied in an inverse manner with that of the p75 and p135 bands. After 72 h of continuous exposure to 50, 150, or 300 µg/ml suramin (Fig. 4 B), no differences were noted in PC3 proteins as assessed by the anti-phosphotyrosine immunoblots. These data indicate that the suramin-induced changes in tyrosine phosphorylation are both rapidly induced and relatively short-lived.

The relatively short-lived drug-induced changes could be the result of alterations in the drug itself, or to a cellular "desensitization" to the drug's action. To distinguish between these possibilities, we applied suramin to PC3 cells and measured the suramin concentration in the culture media 1 min and 3 h later and found no differences (279 vs. 294 µg/ml, respectively). Furthermore, by this HPLC-based assay, we found no evidence for suramin degradation or metabolism. To determine whether the suramin-induced tyrosine phosphorylation desensitizes, the response to suramin was studied at various times after a 2-h suramin pretreatment. As shown in Fig. 5, 2 h of suramin pretreatment clearly diminishes the subsequent suramin-induced response; however, 20 h after pretreatment, the suramin-induced tyrosine phosphorylation pattern was unchanged, indicating reversibility of the desensitization process. The minor differences between the untreated lanes in this figure can be attributed to sample variation. These studies demonstrate that the relatively brief duration of the suramin-induced responses is not the result of suramin degradation, but rather the result of desensitization to the action of the drug.

To determine the dose-response curve for suramin-induced tyrosine phosphorylation in PC3 cells, cells were treated with 0, 30, 100, 300, 1,000, and 3,000 µg/ml of suramin (Fig. 6 A). From these studies it is clear that the suramin-induced alterations in tyrosine phosphorylation of the 75- and 135-kD proteins occurred at dosages of ≥ 100 µg/ml in the PC3 cells. These experiments also emphasize the reciprocal relationship between tyrosine phosphorylation of the p120 and p135 bands. Because suramin is clinically used at a serum level of ~ 300 µg/ml (15, 16, 24), it is clear that suramin-induced changes in tyrosine phosphorylation can occur at clinically relevant drug concentrations within this cell line. To determine whether or not these suramin-induced alterations in tyrosine phosphorylation were detectable in other prostate cancer cell lines, lysates derived from suramin treated PC3M, DU145, and LNCaP cells were immunoblotted with the 4G10 monoclonal antibody (Fig. 6 B-D). The untreated PC3M and LNCaP cells displayed a tyrosine phosphorylation pattern very similar to that of the PC3 cells, with prominent proteins detected at ~ 60 and 120 kD. In contrast, the pattern of tyrosine phosphorylated proteins in the DU145 cells was distinct. In addition to the tyrosine phosphorylated 60- and 120-kD proteins, prominent bands were also detected at 54 and 57 kD. After 10 min of suramin treatment, PC3M, DU145, and LNCaP cells responded by substantially increasing tyrosine phosphorylation of proteins in the 75- and 135-kD size range.

**Figure 4.** Time course of the suramin-induced tyrosine phosphorylation in PC3 cells. Suramin was added to the media and cells were lysed at various time points thereafter. After SDS-PAGE and transfer to nitrocellulose, filters were immunoblotted with the monoclonal antibody 4G10. (A) Suramin (300 µg/ml) was added 10, 20, 40, 55, or 160 min before cell lysis. (B) Suramin (0, 50, 150, or 300 µg/ml) was added to the culture media 72 h before cellular lysis. The electrophoretic mobility of molecular size markers is shown in kilodaltons.
Table 1

<table>
<thead>
<tr>
<th>Time</th>
<th>0</th>
<th>-2h</th>
<th>-4h</th>
<th>-20h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre Rx</td>
<td>-</td>
<td>+</td>
<td>-</td>
<td>+</td>
</tr>
</tbody>
</table>

The PC3M and DU145 cells, but not the LNCaP cells, responded by decreasing the tyrosine phosphorylation of the p120 band. Of interest, the dose response of the various cell lines was distinct, with the androgen-sensitive cell line (LNCaP) being the least sensitive to the suramin-induced changes in these assays.

We next questioned whether or not the suramin-induced alterations in tyrosine phosphorylations were relegated to cells derived from prostatic cancer. To answer this question, a variety of cell lines were treated with suramin and assayed for changes in tyrosine phosphorylation. In Table 1, we summarized data from a series of suramin treated cell lines (Fig. 7, A-D). From these data we conclude that cell lines derived from a number of prostate, breast, gastric, and colon cancers respond to suramin by increasing the tyrosine phosphorylation of 75- and 135-kD proteins, whereas cell lines derived from a number of neuroepitheliomas, lymphomas, and leukemias do not respond in this characteristic fashion. Of interest, some of the cancer cell lines (originally derived from myelomas) did not exhibit suramin-induced alterations of 75- or 135-kD bands, but did show alterations in bands of ~94 and 170 kD (Fig. 7 D). In several cell lines, for instance the gastric-derived cell lines NUGC4 and MKN74 (Fig. 7 C), suramin-induced alterations were detectable within a 75-kD protein but not within a 135-kD protein. No cell lines had alterations in the 135-kD band alone. Several cell lines (HT29, 630, WIDR) also exhibited obvious suramin-induced tyrosine phosphorylations in proteins of ~58 kD (Fig. 7 A). Suramin also induced a modest decrease in tyrosine phosphorylation of a 120-kD band, however, these effects were limited to PC3 (Fig. 7 C) and HT29 cells (Fig. 7 A). Thus, although acute suramin-induced decreases in tyrosine phosphorylation are detectable, they are relatively limited in comparison to suramin-induced increases in tyrosine phosphorylation. From these data we conclude that suramin-induced alterations in tyrosine phosphorylations are a common feature of many, but not all, cell lines. As a generalization, cells with the suramin-induced 75- and 135-kD bands were derived from epithelial adenocarcinomas.

Figure 5. Desensitization, and subsequent recovery, of the suramin-induced tyrosine phosphorylation. PC3 cells were pretreated with suramin at a concentration of 0 (-) or 300 (+) µg/ml for 2 h. After various periods of delay (2, 4, or 20 h), all cells were then treated with suramin (300 µg/ml) and lysed 10 min later. Samples were size-fractionated by SDS-PAGE, transferred to nitrocellulose, and immunoblotted with the 4G10 monoclonal antibody. The electrophoretic mobility of molecular size markers is shown in kilodaltons.

The PC3M and DU145 cells, but not the LNCaP cells, responded by decreasing the tyrosine phosphorylation of the p120 band. Of interest, the dose response of the various cell lines was distinct, with the androgen-sensitive cell line (LNCaP) being the least sensitive to the suramin-induced changes in these assays.

We next questioned whether or not the suramin-induced alterations in tyrosine phosphorylations were relegated to cells derived from prostatic cancer. To answer this question, a variety of cell lines were treated with suramin and assayed for changes in tyrosine phosphorylation. In Table 1, we summarized data from a series of suramin treated cell lines (Fig. 7, A-D). From these data we conclude that cell lines derived from a number of prostate, breast, gastric, and colon cancers respond to suramin by increasing the tyrosine phosphorylation of 75- and 135-kD proteins, whereas cell lines derived from a number of neuroepitheliomas, lymphomas, and leukemias do not respond in this characteristic fashion. Of interest, some of the cancer cell lines (originally derived from myelomas) did not exhibit suramin-induced alterations of 75- or 135-kD bands, but did show alterations in bands of ~94 and 170 kD (Fig. 7 D). In several cell lines, for instance the gastric-derived cell lines NUGC4 and MKN74 (Fig. 7 C), suramin-induced alterations were detectable within a 75-kD protein but not within a 135-kD protein. No cell lines had alterations in the 135-kD band alone. Several cell lines (HT29, 630, WIDR) also exhibited obvious suramin-induced tyrosine phosphorylations in proteins of ~58 kD (Fig. 7 A). Suramin also induced a modest decrease in tyrosine phosphorylation of a 120-kD band, however, these effects were limited to PC3 (Fig. 7 C) and HT29 cells (Fig. 7 A). Thus, although acute suramin-induced decreases in tyrosine phosphorylation are detectable, they are relatively limited in comparison to suramin-induced increases in tyrosine phosphorylation. From these data we conclude that suramin-induced alterations in tyrosine phosphorylations are a common feature of many, but not all, cell lines. As a generalization, cells with the suramin-induced 75- and 135-kD bands were derived from epithelial adenocarcinomas.

Figure 6. Dose-response curve for suramin-induced tyrosine phosphorylation in four human prostate cancer cell lines. Suramin at various concentrations (0, 30, 100, 300, 1,000, or 3,000 µg/ml) was added to the media of (A) PC3, (B) DU145, (C) PC3M, or (D) LNCaP. Cells were lysed 10 min after suramin addition and immunoblotted with the 4G10 monoclonal antibody after size fractionation by SDS-PAGE and transfer to nitrocellulose. The electrophoretic mobility of molecular size markers is shown in kilodaltons.
Suramin is a highly sulfated polyanionic molecule at physiologic pH. To determine whether or not other polyanionic compounds could induce acute alterations in tyrosine phosphorylation, heparin (1–1,820 U/ml), heparan sulfate (10–50 µg/ml), glycosaminoglycans purified from urine of suramin-treated patients (10–50 µg/ml, see reference 26) and pentosan polysulfate (1–1,200 µg/ml) were assayed in the 10-min PC3 tyrosine phosphorylation assay. None of these compounds caused any alterations in tyrosine phosphorylation (data not shown). Protamine sulfate (3–100 µg/ml), a compound (like suramin) that reverses bFGF and PDGF action (27, 28), also had no effect on tyrosine phosphorylation in PC3 cells. In addition, simultaneous additions of heparin and suramin did not demonstrate interactions between these two molecules (data not shown). These data demonstrate that the suramin-induced alterations in tyrosine phosphorylation are specific for suramin, and not simply dependent upon the presence of a highly charged polyanion.

Alterations in protein-tyrosine phosphorylation may be due to either activation of protein-tyrosine kinases or inhibition of protein-tyrosine phosphatases. To assess the possibility of kinase activation, we performed in vitro immune-complex kinase assays utilizing 32P-labeled ATP after anti–phosphotyrosine immunoprecipitations. Under such circumstances, auto-phosphorylating kinases, and in vitro kinase substrates that are immunoprecipitated by anti–phosphotyrosine antibodies will be detected as 32P-labeled proteins. After size-fractionation of the reaction products by SDS-PAGE (Fig. 8), a prominent 75-kD band was detectable in the samples derived from suramin-treated PC3 cells as well as the colon-derived cell lines (HT 29, SW 620, and COLO 205). Samples derived from the WiDR cell line (also colon-derived) had a less prominent, but detectable, band at 75 kD as well. Of note, the 135-kD band detected in immunoblots after suramin treatment was not detected in the immune-complex kinase assays, suggesting that the mechanism(s) of suramin-induced phosphorylation may differ for the 75- and 135-kD proteins. Similar results were obtained in the PC3 and SW620 cells (other cells were not tested) when using antibody PY20 or 4G10 (not shown). These results indicate that anti–phosphotyrosine immunoprecipitates derived from suramin-treated PC3 cells contain increased kinase activity relative to untreated PC3 cells and suggest that suramin-induced increases in cellular phosphorylation may be the result of enhanced kinase activity. To determine if the 75-kD protein detected after suramin treatment is the same in prostate and colon cells, digestions with V8 protease were performed (using p75 labeled in immune-complex kinase reactions). Analysis of the 32P-labeled fragments by SDS-PAGE revealed peptides of identical size (Fig. 9), suggesting that suramin-induced tyrosine phosphorylation of p75 represents the same protein in cells derived from prostatic and colonic neoplasms. V8 proteolysis also confirms that the 75-kD protein recognized by the PY20 monoclonal antibody is highly similar, if not identical, to that recognized by the 4G10 antibody (Fig. 9).

**Discussion**

The exact mechanism of suramin action has not been determined from these studies. The simplest hypothesis is that suramin directly alters the enzymatic activity of a protein-tyrosine kinase. However, our results do not rule out the possibility that a tyrosine phosphatase(s) is directly or indirectly involved, particularly in the phosphorylation changes detected in the 120-kD band. It is also possible that suramin acts via an “intermediate” compound which in turn modulates the activity of a protein-tyrosine kinase or phosphatase. Such an intermediate mechanism has recently been postulated for suramin-induced epidermal growth factor (EGF) receptor activation in the A431 cell line (29). In A431 cells, suramin treatment increases TGF-α (as detected by radioimmunounassay) in the culture media in a time-dependent fashion and increases cellular tyrosine phosphorylation. In that study (29), neither TGF-α nor alterations in tyrosine phosphorylation were noted within the first several

---

**Table I. Suramin Increases Tyrosine Phosphorylation of a 135-and/or 75-kD Proteins in a Multiplicity of Cell Lines**

<table>
<thead>
<tr>
<th>Cell Line</th>
<th>75 kD</th>
<th>135 kD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prostate (PC3)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Prostate (PC3M)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Prostate (DU145)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Prostate (LNCaP)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (HT 29)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (630)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (WiDR)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (SW 620)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (COLO 205)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Colon (716)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Gastric (TMK-1)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Gastric (A2521)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Gastric (NUGC4)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Gastric (MKN74)</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>Breast (ZR-75)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Breast (H5578)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Breast (T47D)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Breast (MDA-231)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Medulloblastoma</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Neuroepithelioma</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Neuroepithelioma</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lung (small cell)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lung (small cell)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lung (squamous)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Leukemia (HL60)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Leukemia (CEM)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Myeloma (H929)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Myeloma (U266)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lymphoma (Wilson)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Lymphoma (Defauw)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Melanoma (RPMI 7951)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Melanoma (H5695T)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Cervical (HeLa)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Rhabdomyosarcoma</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Fibroblast (NIH 3T3)</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Cells were lysed 10 min after 1,000 µg/ml suramin addition to the media. After size-fractionation (SDS-PAGE) and transfer to nitrocellulose, filters were blotted with an anti–phosphotyrosine antibody (4G10). Cell lines were categorized as having enhanced (+) or not enhanced (−) bands at 75 or 135 kD.
Plausible candidates for the tyrosine-phosphorylated proteins detected in these studies include a variety of tyrosine kinases, and their substrates, with molecular masses of ~75 and 135 kD. Tyrosine kinases of ~135 kD include members of the FGF receptor family and the met proto-oncogene (hepatocyte growth factor receptor) (see references 17 and 30). Members of the trk family (31, 32), ret (33), and the recently described 72-kD syk kinase (34) are also within the expected size ranges; however, these proteins are expressed in a tissue-specific manner that is not compatible with that found for the suramin-induced tyrosine phosphorylations. Potential tyrosine-phosphorylated "substrate" molecules include PLC-γ (35, 36), the ras GTPase-activating protein (GAP) (37, 38), and/or the 80-, 120-, 125-, and 130-kD tyrosine-phosphorylated proteins initially identified in fibroblasts expressing activated src genes (23). In our preliminary experiments (utilizing antibodies to specific proteins in combination with anti-phosphotyrosine immunoblotting), we have failed to detect suramin-induced alterations in tyrosine phosphorylation for each of the "substrate" molecules mentioned above. Of interest, however, we did detect prominent expression and tyrosine phosphorylation of a p120 protein when utilizing the anti-p125 (2A7) antibody on
PC3 cell lysates. In addition, we failed to detect alterations in
 tyrosine phosphorylation when utilizing antibodies directed
 against the product of the /tg gene (an FGF receptor), the met
 proto-oncogene, or the 80-kD subunit of the IL-6 receptor
 (data not shown). Thus the identities of the 75- and 135-kD
 suramin-induced bands are not known at this time and more
 experiments will be necessary to identify these molecules.

 These studies clearly indicate that suramin effects are more
 complex than a simple abrogation of heparin-binding growth
 factors, a finding underscored by the recent demonstrations
 that suramin can block ATP binding to certain purinergic re-
 ceptors (39) and can induce production of antiproliferative
 heparan sulfates (26). The rapid effects of suramin on tyrosine
 phosphorylation, at first glance, appear to be antithetical to
 suramin’s role as an antitumor agent. However, these results
 suggest an alternative hypothesis for suramin’s antitumor
 mechanism of action. Based upon our data, we hypothesize
 that a portion of the antitumor effect of suramin may be due to
 the activation, and subsequent down-regulation, of a signal
 transduction pathway essential for cancer cell growth. It is also
 plausible that alterations in signal transduction may play a role
 in suramin’s antiparasitic action.

 In summary, these studies clearly demonstrate an acute and
 marked alteration in the tyrosine phosphorylation of several
 cellular proteins in response to suramin treatment. In some cell
 lines, this response occurs at levels of the drug used in the
 clinical setting. To our knowledge, these studies represent the
 first demonstration of enhanced tyrosine phosphorylation by a
 nonpolypeptide pharmaceutical agent. Though our initial stud-
 ies were restricted to prostate cancer cell lines, it is readily ap-
 parent that suramin can elicit similar changes in an assortment
 of cell lines derived from a variety of human tissues.

 Acknowledgments

 We would like to thank D. Hill, P. Gernt, M. Khan, and Dr. M. Cooper
 for their laboratory assistance; Dr. M. Ranson for providing glycosami-
 noglycans purified from the urine of suramin treated cancer patients;
 Dr. T. Parsons for providing antibodies to tyrosine-phosphorylated
 proteins; Dr. R. Nordan for providing IL-6 receptor antibodies; Dr. D.
 Bottaro for providing antibodies to the product of the met gene; and J.
 Amber for editorial assistance.

 References


