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Additive apoptotic effect of STI571 with the cytoprotective agent amifostine in K-562 cell line

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Abstract Purpose: To study the apoptotic effect of the 2-phenylaminopyrimidine derivative STI571 in combination with antioxidant agents on K-562 cell line derived from a Philadelphia chromosome-positive chronic myeloid leukemia patient. **Materials and methods:** K-562 (BCR/ABL+), U-937, and HL60 (BCR/ABL–) leukemic cell lines were incubated with STI571 and the antioxidant agents catalase, glutathione, superoxide dismutase, and amifostine (AMI). Apoptotic effect was analyzed by morphological and flow cytometric criteria. **Results:** STI571 at concentrations higher than $0.25 \mu\text{mol L}^{-1}$ produced apoptosis ($P < 0.05$) in K-562 cells only after treatment for 72 h. At the mentioned concentrations, STI571 also induced an increase in the loss of mitochondrial transmembrane potential from 24.6 to 40%. Combination of STI571 ($0.5 \mu\text{mol L}^{-1}$) with antioxidant agents showed that the cytoprotective agent AMI (0.75 mg mL^{-1}) produced an additive effect in the proapoptotic activity of STI571 in K-562 cells at nuclear ($58.8\% \pm 2.0$ vs. $28.9\% \pm 3.3$) and mitochondrial ($53.3\% \pm 3.6$ vs. $29.5\% \pm 1.2$) levels. **Conclusions:** Our results show that only AMI in combination with STI571, at submicromolar concentration, has an additive effect in K-562 cell line, and it does not have severe toxic effects on Philadelphia chromosome negative cells.

Keywords STI571 · Apoptosis · K-562 cell line · Antioxidant agents · Amifostine · Cell proliferation

Abbreviations AlloBMT: Allogeneic bone-marrow transplantation · AMI: Amifostine · ATP: Adenosine triphosphate · BC: Fatal blastic phase · BCR/ABL–:

BCR/ABL negative cells · BCR/ABL+ : BCR/ABL positive cells · CA: Continuous administration · CAT: Catalase · CML: Chronic myeloid leukemia · CP: Chronic phase · $\Delta\Psi_m$: Mitochondrial transmembrane potential · DMSO: Dimethyl sulfoxide · EDTA: Ethylenediaminetetraacetic acid · GSH: Glutathione · HDC: Highly damaged cells · IFN: Interferon · LMA: Low-melting agarose · MDC: Middle damaged cells · PBS: Phosphate-buffer solution · PI-3k: Phosphatidylinositol-3 kinase · PMN: Polymorphonuclear cells · Rh123: Rhodamine 123 · RNase A: Ribonuclease A · ROS: Reactive oxygen species · SOD: Superoxide dismutase · STI571: 2-Phenylaminopyrimidine derivative · UC: Undamaged cells

Introduction

Chronic myeloid leukemia (CML) is a clonal myeloproliferative disorder resulting from neoplastic transformation of the primitive hematopoietic stem cell [1]. The cytogenetic hallmark of CML is a reciprocal translocation between chromosomes 9 and 22 t(9;22)(q34;q11) [2]. This translocation results in a BCR/ABL hybrid gene and the fusion protein, p210^{BCR/ABL}. This protein has abnormal tyrosine kinase activity compared with normal c-ABL protein [3] and there is evidence that p210^{BCR/ABL} itself can act as an antiapoptotic factor [4, 5].

The course of CML involves progression from a chronic phase (CML-CP) to an accelerated phase with duration of 1–1.5 years followed by a fatal blastic phase (CML-BC) [6, 7].

Current therapies for CML include allogeneic bone-marrow transplantation (AlloBMT) and drug regimens including interferon (IFN)-alpha. AlloBMT is associated with substantial morbidity and mortality and is limited to patients for whom a suitable donor is available. IFN-alpha prolongs overall survival but has

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serious adverse effects. Thus, the overall cure for CML is less than 20% [8].

STI571, a 2-phenylaminopyrimidine derivative ($C_{30}H_{35}N_7SO_4$), is a potent ABL tyrosine kinase inhibitor with which encouraging results have been obtained in phase I and II clinical trials [9–11]. However, 60% of patients with myeloid CML-BC and most of patients with lymphoid CML-BC eventually relapse [9], indicating there are leukemic cells that overcome the proapoptotic effect of STI571 and survive in the presence of pharmacological concentrations of the drug [12].

To increase the cytotoxic effects of STI571, combinations of this drug with commonly employed anti-leukemic agents were used and shown to exert increased effects against BCR/ABL positive cells as compared with STI571 alone [13, 14]. On the other hand, exogenous expression of p210^{BCR/ABL} in BA/F3, 32Dcl3, and M07e hematopoietic cell lines results in an increase of reactive oxygen species (ROS) in comparison with their untransformed counterparts; this would enhance BCR/ABL-mediated transformation events by inhibiting cell phosphatases [15].

Taking these data into account, the aim of this work was to study the proapoptotic effect of the combination of STI571 with antioxidant agents on the erythroblastic K562 cell line, which is derived from a Philadelphia chromosome-positive CML patient [16].

Materials and methods

Cell cultures

K-562, BCR/ABL+ (ATCC, USA), U-937, and HL60, BCR/ABL– (kindly provided by Dr Mirta Giordano from the Instituto de Investigaciones Hematológicas, Argentina) cells were maintained in a 5% CO₂ atmosphere at 37°C in RPMI 1640 medium supplemented with 10% heat-inactivated (30 min, 65°C) fetal bovine serum. Cells were used for the studies in passages 3–10 after thawing. Polymorphonuclear cells from peripheral blood from six normal donors (PMN) were isolated by centrifugation in Ficoll–Histopaque and further sedimentation in a dextran gradient.

Drug treatments

K-562, U-937, and HL60 cells were incubated for 24, 48, and 72 h in the presence of STI571 (Imatinib, Gleevec, Novartis, Basel, Switzerland), at concentrations ranging between 0 and 2.5 $\mu\text{mol L}^{-1}$. Antioxidant agents were used in the dose range reported in previous papers: [17–19]. Catalase, CAT 200 U mL^{-1} (Sigma, St Louis, USA), glutathione, GSH, 100 $\mu\text{mol L}^{-1}$ and 1 mmol L^{-1} (Sigma), superoxide dismutase, SOD, 200 $\mu\text{g mL}^{-1}$ (Sigma), and amifostine, AMI, 0.75 mg mL^{-1} (Actifos, Laboratorios Filaxis, Argentina) were added 15–30 min before treatment with STI571 (0.5 $\mu\text{mol L}^{-1}$) and were

present throughout the culture time. In addition, AMI, 0.75 mg mL^{-1} , was added for 15 min and removed from the medium before STI571 treatment. PMN cells were incubated with antioxidant agents and STI571 for 24 h.

Apoptosis assessment

Apoptosis was assessed by fluorescence microscopy, after staining of the cells with a mixture of acridine orange (100 $\mu\text{g mL}^{-1}$) and ethidium bromide (100 $\mu\text{g mL}^{-1}$) in PBS. Loss of nuclear structure or fragmented nuclei were taken as indicative of apoptotic cells. Apoptosis were also detected by flow cytometry, using loss of mitochondrial transmembrane potential ($\Delta\Psi_m$) as a marker of apoptosis. Loss of $\Delta\Psi_m$ was assessed by incubating the treated cells with 0.3 $\mu\text{g mL}^{-1}$ rhodamine 123 (Rh123) for 15 min. Cytometric analysis was performed with a FacScan flow cytometer (Becton Dickinson, San Diego, USA). Cell Quest software was used for data acquisition and analysis.

Single-cell gel electrophoresis assay (comet assay)

For detection of DNA damage and apoptosis, the comet assay was carried out under alkaline conditions. After treatment, cells (10 μL) were mixed with 75 μL 0.75% low-melting agarose (LMA), spread on a slide precoated with normal melting agarose (0.75%), and a third layer of LMA was then applied. The microscope slides were placed in cold lysis solution (2.5 mol L^{-1} NaCl, 100 mmol L^{-1} EDTA, 10 mmol L^{-1} Tris pH 10, 1% Triton X-100, 10% DMSO) and kept at 4°C for at least 1 h. Slides were then transferred to an electrophoresis box containing an alkaline solution at pH 13 (300 mmol L^{-1} NaOH, 1 mmol L^{-1} EDTA) and kept for 20 min unwinding time at 4°C. A current of 30 V (250 mA) was applied for 25 min. Slides were removed, neutralized with 0.4 mol L^{-1} Tris pH 7.5 for 5 min, and stained with 20 μL ethidium bromide (20 $\mu\text{g mL}^{-1}$).

Comets were evaluated with a fluorescence microscope. A total of 100 comets on each slide were scored visually as belonging to one of three classes according to tail fluorescence intensity, tail length, and nucleoid integrity, and classified as undamaged cells (UC), cells with an intermediate grade of DNA damage (MDC), and highly damaged cells (HDC).

Cell proliferation: tritiated thymidine (³H-thymidine) incorporation and cell-cycle studies

K-562 cells (200×10^3 per well) were seeded in 200 μL culture medium and treated with STI571 at concentrations ranging from 0 to 2.5 $\mu\text{mol L}^{-1}$ for 72 h. Eighteen hours before finishing the incubation 20 μL per well of a working solution of ³H-thymidine (0.05 $\mu\text{Ci mL}^{-1}$) was added and activity was detected with a β counter.

To assess DNA ploidy levels STI571-treated K-562 cells were fixed in ethanol 70%, treated with RNase A (0.2 mg mL^{-1} for 15–30 min), and resuspended in propidium iodide ($20 \text{ } \mu\text{g mL}^{-1}$ in PBS). Cytometric analysis was performed with a FacScan flow cytometer (Becton Dickinson). Modfit software was run for data acquisition and analysis.

Statistical analysis

Differences between mean results from the different treatments were evaluated by applying the paired Student's *t*-test.

Results

STI571-induced apoptosis

K-562, U-937, and HL60 cells were treated with STI571 at the stated concentrations (0, 0.01, 0.05, 0.25,

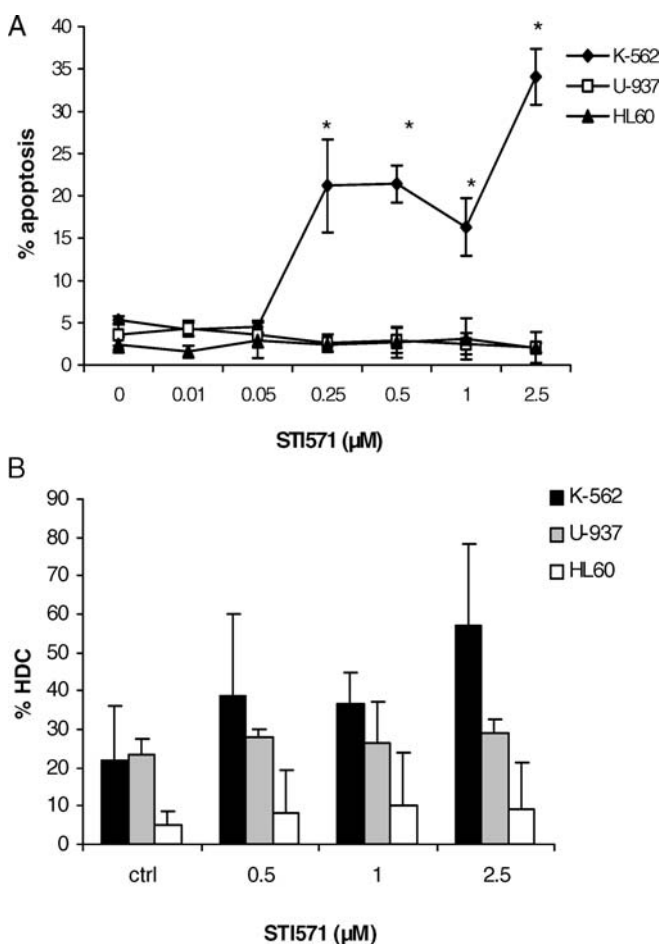


Fig. 1 A, STI571 ($0\text{--}2.5 \text{ } \mu\text{mol L}^{-1}$)-induced apoptosis in K-562, U-937, and HL60 cells after incubation for 72 h (* $P < 0.05$). Representative of four independent experiments. B, Percentage of HDC (mean $\pm$ SD) in K-562, U-937, and HL60 cell lines after incubation for 72 h with the indicated concentrations of STI571. Representative of three independent experiments

0.5, 1.0 and $2.5 \text{ } \mu\text{mol L}^{-1}$) for 24, 48, and 72 h. Apoptosis was assessed by morphological analysis and only after 72 h of treatment; STI571 induced significant levels of apoptosis ($P < 0.05$) starting from $0.25 \text{ } \mu\text{mol L}^{-1}$ in K-562 cells in comparison with untreated controls. No increase of apoptosis percentage was observed in STI571-treated U-937 and HL60 cell lines (Fig. 1A). The response of the leukemic cell lines to STI571 was also studied by the comet assay. A trend toward increased percentages of HDC in K-562 cells only after incubation with the drug for 72 h was observed, but no significant differences were found (Fig. 1B). STI571 at the mentioned concentrations also induced an increase in the loss of $\Delta\Psi_m$ from 24.6 to 40% in K-562 cells (Fig. 2).

Cell-proliferation studies

^3H -Thymidine incorporation studies were performed with K-562 cells treated with STI571. It was found that after incubation with the drug for 72 h incorporation of ^3H -thymidine decreased ($P < 0.05$) if concentrations of STI571 were higher than $0.25 \text{ } \mu\text{mol L}^{-1}$ (Fig. 3).

DNA content analysis by flow cytometry of K-562 cells in the presence of STI571 (0, 0.5, 1, and $2.5 \text{ } \mu\text{mol L}^{-1}$) showed a lower percentage of cell subpopulations in S and G₂/M phases of cell cycle in comparison with untreated cells (Fig. 4)

STI571 in combination with antioxidant agents

To enhance the proapoptotic effect of STI571 in K-562 cells combined treatment with STI571 and antioxidant agents was studied. Cells were incubated for 30 min with

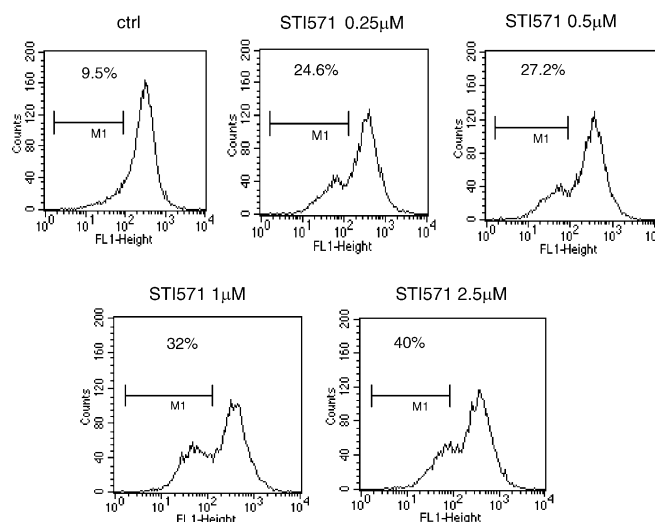


Fig. 2 Loss of $\Delta\Psi_m$ in K-562 measured by Rh123 after incubation for 72 h with STI571 (representative of three independent experiments). M1: fraction of cells considered to have a decrease in their mitochondrial transmembrane potential, compared with untreated controls

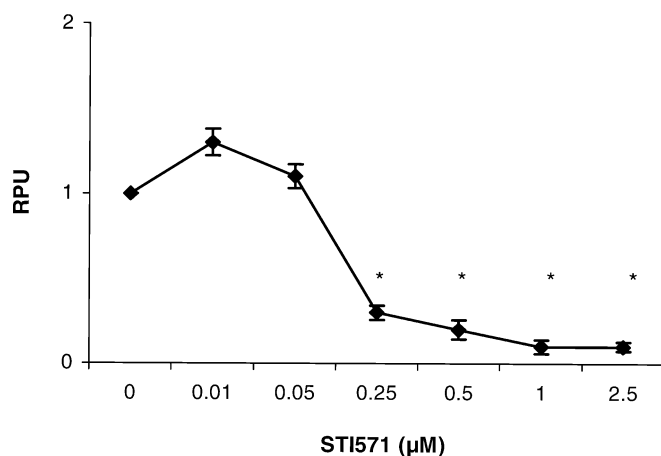


Fig. 3 Incorporation of ^3H -thymidine by K-562 cells treated with STI571 for 72 h (* $P < 0.05$ in comparison with untreated control). (RPU relative proliferation units; referred to incorporation of ^3H -thymidine in untreated cells as 1)

SOD, CAT, and GSH then the ABL tyrosine kinase inhibitor was added. Addition of these antioxidant agents did not modify the apoptotic value induced by STI571 alone (Fig. 5). Incubation with the cytoprotective agent AMI was carried out in two ways—continuous administration (CA) for 72 h and for 15 min, after which it was removed from the culture medium before addition of STI571. Apoptosis was assessed by morphological and cytometric analysis, to study cell death at the nuclear and mitochondrial levels. By morphological detection the percentage of apoptosis induced by treatment with a combination of AMI for 15 min and STI571 (mean $\pm$ SE, $28.7 \pm 3.3\%$) was no different from that

resulting from treatment with STI571 alone (mean $\pm$ SE, $28.9 \pm 4.0\%$). CA of AMI had an additive effect in combination with STI571 (mean $\pm$ SE, $58.8\% \pm 2.0$). This additive effect of STI571 and AMI (CA) was also observed at the mitochondrial level (mean $\pm$ SE, $53.3\% \pm 3.6$ vs. $29.5\% \pm 1.2$ in STI571-treated cultures) (Fig. 6).

To determine the specificity of the additive effect of STI571 and AMI in BCR/ABL+ cells, U-937 cells (BCR/ABL– cells) were incubated with both drugs for 72 h. By morphological detection of apoptosis it was observed that AMI was a strong apoptotic inducer in U-937 cells; however, combination with STI571 did not increase levels of apoptosis (Fig. 7).

PMN cells from six normal donors were also treated with both drugs, taking into consideration that severe neutropenia is one of the most frequent adverse effects associated with treatment with STI571. This drug at $0.5 \mu\text{mol L}^{-1}$ induced an increase of apoptosis and the effect was reversed by pretreatment with AMI after incubation for 24 h (Fig. 8).

These data indicate that AMI increased the apoptotic effect of STI571 and is specific for BCR/ABL+ cells.

Discussion

Because the structure of several protein kinases has been determined, it is possible to design compounds based on the structure of the ATP binding site or their active site. One of these compounds, the 2-phenylaminopyrimidine derivative STI571, is a specific inhibitor of the ABL family of tyrosine kinases (p210^{BCR/ABL}, p185^{BCR/ABL}, v-ABL, and c-ABL), although it also inhibits the kinase activity of platelet-derived growth factor receptor (PDGFR) and c-kit [20].

To study the antileukemic effect of STI571, K-562, U-937, and HL60 leukemic cell lines were treated with concentrations of the drug up to $2.5 \mu\text{mol L}^{-1}$.

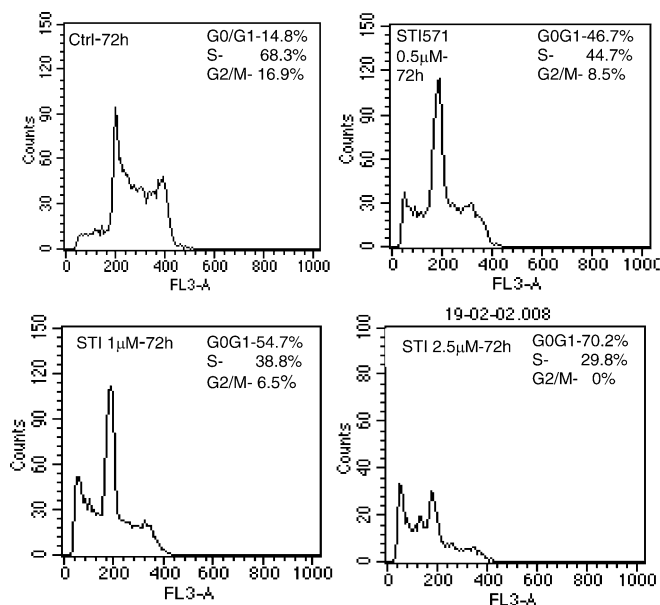


Fig. 4 DNA content histograms of K-562 cell line treated for 72 h with the indicated concentrations of STI571 (representative of three independent experiments). The apoptotic fraction was not considered in the cell cycle analysis

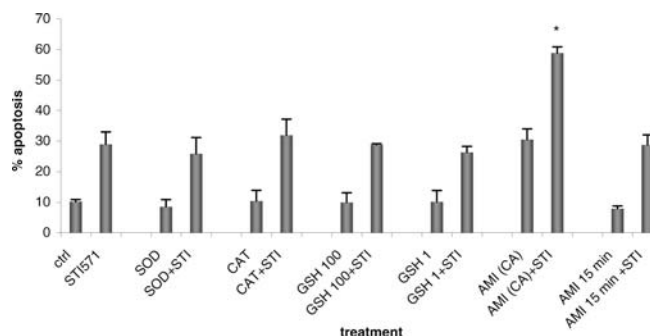


Fig. 5 Apoptosis, detected by morphological analysis (mean $\pm$ SE), induced in K-562 cells by STI571 ($0.5 \mu\text{mol L}^{-1}$) in combination with antioxidant agents after incubation for 72 h (CAT catalase; SOD superoxide dismutase; GSH 100 glutathione $100 \mu\text{mol L}^{-1}$; GSH 1 glutathione 1 mmol L^{-1} ; AMI amifostine) (* $P < 0.05$ in comparison with cells treated with STI571). Representative of five independent experiments

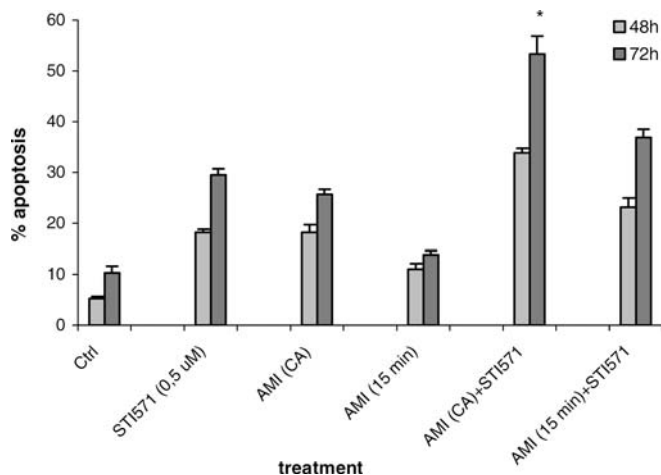


Fig. 6 Apoptosis (mean $\pm$ SE) induced by STI571 ($0.5 \mu\text{mol L}^{-1}$) in combination with AMI in K-562 cell line after incubation for 48 and 72 h. Percentages represent loss of $\Delta\Psi_m$ detected as a decrease in Rh123 fluorescence (* $P < 0.05$ in comparison with cells treated with STI571). Representative of three independent experiments

STI571 induced significant levels of apoptosis in K-562 cells after incubation for 72 h with the drug at concentrations higher than $0.25 \mu\text{mol L}^{-1}$. U-937 and HL60 cells did not trigger apoptosis in response to STI571 after treatment for 72 h at the same concentrations. These data confirm previous reports showing induction of apoptosis in BCR/ABL transfected cell lines; while STI571 did not affect the viability of their wild-type counterparts [21].

Concentrations of STI571 employed were equivalent to those found in plasma from CML patients 24 h after administration of 400 mg of the drug ($1.46 \mu\text{mol L}^{-1}$) [9].

When apoptosis was measured at the mitochondrial level, Tabrizi et al. [14] reported that STI571 at $1 \mu\text{mol L}^{-1}$ during 18 h of incubation was able to induce 10 to 40% apoptosis in four BCR/ABL-expressing cell lines—AR230, LAMA84, K562, and KCL22. This percentage was comparable with that obtained by us in K-562 cells.

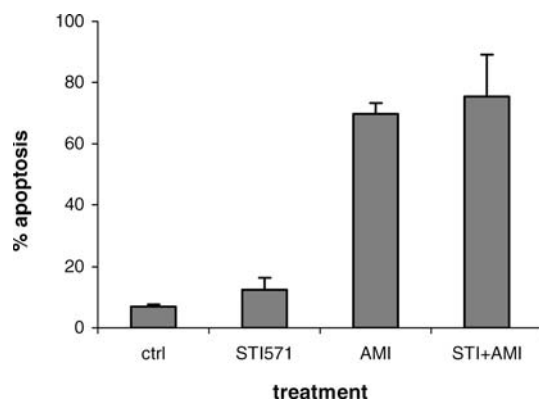


Fig. 7 Apoptosis (mean $\pm$ SE) induced by STI571 ($0.5 \mu\text{mol L}^{-1}$) plus AMI in U-937 cells. Representative of three independent experiments

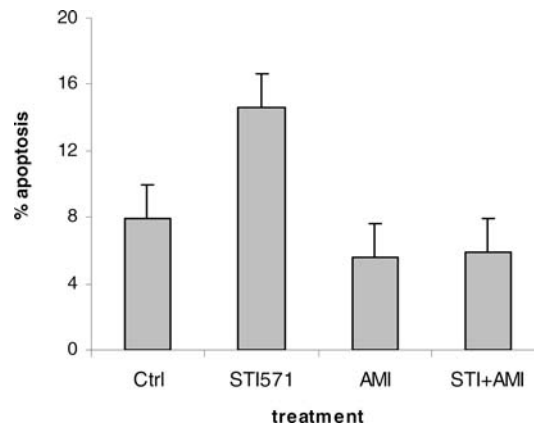


Fig. 8 Effect of STI571 ($0.5 \mu\text{mol L}^{-1}$) plus AMI on PMN cells from the peripheral blood of six healthy donors. Apoptosis (mean $\pm$ SE) was determined by morphological observation after 24 h

Apoptotic response of the leukemic cell lines to STI571 was also analyzed by the comet assay; an increase in the percentage of highly damaged K-562 cells was observed after incubation with the drug for 72 h. Such an increase was not observed for U-937 and HL60 cells.

Moreover, STI571 markedly decreased incorporation of ^3H -thymidine in K-562 cells after incubation for 72 h with up to $2.5 \mu\text{mol L}^{-1}$ of the drug. DNA content analysis by flow cytometry showed a decrease in S and G₂/M subpopulations in STI571-treated cells. Consistent with our results, Gambacorti-Passerini et al. [22] have shown that STI571-treatment induced a decrease in cycling cell lines derived from BCR-ABL + leukemias.

STI571 has given encouraging results in clinical trials on patients with CP-CML [9], with CP-CML refractory to IFN- α [10], and in those with the accelerated phase of CML. However, 60% of CML myeloid blast crisis and most lymphoid blast crisis eventually relapse [9] indicating there are cells which are able to grow continuously in the presence of STI571. Mechanisms of resistance to this drug vary between the different BCR/ABL + cell lines. The most frequent of these mechanisms, detected in BaF3 murine leukemic cells transformed with the BCR/ABL hybrid gene and in the human CML cell lines LAMA84 and AR230 is overexpression of the BCR/ABL protein. Resistance to STI571 without an increase of p210^{BCR/ABL} levels in subpopulations of K-562 and KCL22 cells has, however, been reported [12]. It has been shown in studies with cells from CML patients that resistance to STI571 was because of mutation (8/11 patients) or amplification of the BCR/ABL gene (3/11 patients) [23].

To improve the efficacy of treatment of Philadelphia chromosome-positive leukemias with STI571 there have been many studies of combination of STI571 with commonly used antileukemic agents, for example γ -irradiation, busulfan, treosulfan [13], and daunorubicin [14]. Combination of STI571 with the phosphatidylinositol-3 kinase (PI-3k) inhibitors, wortmannin and

LY294002 has also been studied, because PI-3k activity is essential for the growth of Philadelphia chromosome-positive cells [8].

It has, on the other hand, been reported that transformation of BA/F3, 32Dcl3 and MO7e cell lines with BCR/ABL gene results in an increase of ROS compared with their untransformed counterparts. This would enhance p210^{BCR/ABL}-mediated phosphorylation events, by ROS-mediated inhibition of cell phosphatases. This increase in ROS was because of p210^{BCR/ABL} activity, because it was blocked by STI571 [15].

In this work the effect of combining STI571 with antioxidant agents on the leukemic cell lines K-562 and U-937 and on PMN from peripheral blood of normal donors has been evaluated. The concentration of STI571 used for the combination treatment was 0.5 $\mu\text{mol L}^{-1}$, almost one-third of the average concentration (1.46 $\mu\text{mol L}^{-1}$) reached in the plasma of CML patients 24 h after administration of 400 mg of the ABL kinase inhibitor [9]. The antioxidant agents used were SOD, CAT, GSH, and the cytoprotective agent AMI. Among these, AMI was the only one with an additive effect on the proapoptotic activity of STI571 in K-562 cell line at both nuclear and mitochondrial levels. This effect was not observed in U-937 cells or in PMN. AMI, a phosphorylated aminothioliol prodrug, is dephosphorylated by membrane-bound alkaline phosphatases, releasing the active metabolite WR-1065, which quickly accumulates in normal tissues [24].

Continuous administration of AMI to p53-deficient cells K-562 and U937 induced an increase in the percentage of apoptotic cells, but this increase was not observed for the p53-proficient cells PMN. These data are in agreement with Lee et al. [25] who identify p53 protein as a mechanism of resistance to AMI-induced apoptosis.

AMI protects bone marrow progenitor cells from the cytotoxic effect of chemotherapeutic agents [26]. Clinical trials have shown that AMI reduces myelosuppression resulting from irradiation, alkylating agents, and platinum analogs, while keeping their antitumor activity [27]. It has also been shown that this aminothioliol exerts a protective effect on early and late hematological progenitor cells from the cytotoxic effects of cyclophosphamide derivatives [28]. All these data obtained in vitro support the use of AMI in association with chemotherapeutic agents to reduce toxicity in normal cells.

Our results show that AMI in combination with STI571 at submicromolar concentrations has an additive effect on the K-562 cell line whereas it does not have severe toxic effects on Philadelphia chromosome negative cells or normal PMN. The cytoprotective effect of AMI observed in PMN is very important, taking into account that severe neutropenia is one of the most frequent adverse effects associated with STI571 treatment. [10, 29]. Similarly, Blasiak et al. [30] reported that AMI reduced the extent of DNA damage induced by the chemotherapeutic agent idarubicin in normal human lymphocytes and murine BaF3 cells, but when BaF3

cells were transformed with TEL/ABL gene, AMI potentiated the DNA-damaging effect of idarubicin.

Future studies should be performed to clarify whether the additive effect of using AMI in combination with STI571 in BCR/ABL+ and p53- cells is related to the scavenger ROS properties and/or the proapoptotic activity of AMI on them. It should also be determined whether this combination of agents could be clinically relevant.

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