In vitro inhibitory effects of imatinib mesylate on stromal cells and hematopoietic progenitors from bone marrow

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Abstract

Imatinib mesylate (IM) is used to treat chronic myeloid leukemia (CML) because it selectively inhibits tyrosine kinase, which is a hallmark of CML oncogenesis. Recent studies have shown that IM inhibits the growth of several non-malignant hematopoietic and fibroblast cells from bone marrow (BM). The aim of the present study was to evaluate the effects of IM on stromal and hematopoietic progenitor cells, specifically in the colony-forming units of granulocyte/macrophage (CFU-GM), using BM cultures from 108 1.5- to 2-month-old healthy Swiss mice. The results showed that low concentrations of IM (1.25 µM) reduced the growth of CFU-GM in clonogenic assays. In culture assays with stromal cells, fibroblast proliferation and α-SMA expression by immunocytochemistry analysis were also reduced in a concentration-dependent manner, with a survival rate of approximately 50% with a dose of 2.5 µM. Cell viability and morphology were analyzed using MTT and staining with acridine orange/ethidium bromide. Most cells were found to be viable after treatment with 5 µM IM, although there was gradual growth inhibition of fibroblastic cells while the number of round cells (macrophage-like cells) increased. At higher concentrations (15 µM), the majority of cells were apoptotic and cell growth ceased completely. Oil red staining revealed the presence of adipocytes only in untreated cells (control). Cell cycle analysis of stromal cells by flow cytometry showed a blockade at the G0/G1 phases in groups treated with 5-15 µM. These results suggest that IM differentially inhibits the survival of different types of BM cells since toxic effects were achieved.

Key words: Bone marrow stromal cell; Hematopoietic progenitors; Imatinib mesylate; Myelosuppression

Introduction

Bone marrow (BM) is composed of functional stroma made up of many cell types and an extracellular matrix, which together form an ideal microenvironment that favors the modulation of quiescence, self-renewal, and commitment of mesenchymal stem cells (MSCs) and the proliferation, differentiation, and apoptosis of hematopoietic cells. These functions correlate with different cell types in the stroma, which consists mainly of macrophages, endothelial cells, fibroblasts, and adipocytes (1). When grown in culture, fibroblasts from different organs are consistently modulated into myofibroblast-like cells with varying degrees of alpha-smooth muscle actin (α-SMA) expression (2). These cells can secrete growth factors, including transforming growth factor, platelet-derived growth factors (PDGF), basic fibroblast growth factor (bFGF), hepatocyte growth factor, keratinocyte growth factor, stem cell factor (SCF), epithelial growth factor (EGF), granulocyte/macrophage colony-stimulating factor (GM-CSF), and other cytokines (3). Bone marrow adipocytes mechanically exert hematopoietic control, which in turn controls the increase in marrow fat inclusions, since the acceleration of hematopoiesis is

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associated with the loss of fat vacuoles, increasing the physical space required for growth (4). Since these cell types are derived from MSCs resident in BM, the rapid rate of cell renewal and differentiation makes the hematopoietic system a target for antineoplastic therapy (5), which can result both from direct drug effects on hematopoietic progenitors (6) and from marrow toxicity (5) due to effects on BM stromal cells.

Imatinib mesylate (IM) is an oral anticancer agent used primarily to treat chronic myeloid leukemia (CML), and has selective activity against a small number of tyrosine kinases, including the BCR-ABL1 oncogene (present in the Philadelphia (Ph) chromosome), c-Kit, c-Ab1, and PDGF receptor. IM acts as a competitive inhibitor by binding with high affinity to the ATP binding site in the kinase domains of these targets. This leads to the inhibition of the phosphorylation of tyrosine residues of the target protein in the signal transduction pathway and, consequently, the excessive cell proliferation occurring in CML is interrupted and apoptosis is induced (7). Blocking the site of action is also the mechanism by which IM interrupts the signaling exacerbated in cells with a c-Kit mutation, which is common in gastrointestinal stromal tumors (GISTs) (8).

At the current standard dose of IM for CML in the chronic phase (400 mg), peak plasma levels at steady state were approximately 4.6 μM and trough levels approximately 2.1 μM, with a half-life of 19.3 h (9). Some studies, however, have reported higher plasma levels (10). The maximal recommended dose is 800 mg/day. However, in these patients the high-dose IM (800 mg/day) may be more effective than standard-dose IM (400 mg/day) (11). Initially approved at doses of 400 and 600 mg daily for advanced GIST patients, the use of high-dose IM (800 mg daily) has been shown to benefit patients with advanced or metastatic GIST that progresses while on the standard dose, and has been recommended under these circumstances by the major management guidelines (12). There are few in vitro studies that use IM at plasma concentrations achieved with doses of 800 mg/day (10), but it is known that in some cases in which a patient has achieved good hematologic control but has failed to achieve or has lost a cytogenetic response, dose escalation may increase the degree of Ph negativity (13). The duration of exposure of CD34+ cells to IM showed a greater effect with prolonged exposure (14).

While it was initially reported that IM minimally affected normal hematopoiesis, recent studies have shown that IM can inhibit the growth of some nonmalignant hematopoietic cells. It was demonstrated that IM may affect the mobilization, proliferation, and differentiation of hematopoietic progenitor cells resulting in myelosuppression in patients receiving treatment. The reversible side effects of IM are dose-dependent and affect all hematopoietic lineages, albeit to varying degrees (15). Data from GIST patients suggest that IM directly suppresses normal hematopoiesis under the assumption that the disease itself has no effect on normal hematopoiesis in these patients. In a phase II study of patients with GIST or other soft tissue sarcomas who received 400 mg IM daily, 92% of patients displayed anemia and 47% displayed granulocytopenia (16).

Several studies have described the use of IM, in addition to CML and GIST, as a possible new therapy for fibrotic diseases (17) and for diseases that require an angiogenic stimulus (18). With the possibility of IM use for many other non-hematological diseases, assessment of its myelosuppressive effects has become important and necessary in order to determine the proper use and prescription of this drug. However, only a few recent studies have investigated the normal BM stroma of animal or human origin (19) after IM treatment, while most have focused mainly on patients with CML in complete cytogenetic response (20).

In this study, we used healthy murine BM as a model of the hematopoietic microenvironment to study the effects of IM on marrow stromal and hematopoietic cells, specifically in the colony-forming units (CFU)-GM.

Material and Methods

Collection of BM cells from mice

Bone marrow cells were harvested from 16 femurs (8 animals/protocol) of 1.5- to 2-month-old male Swiss mice (Mus musculus) killed by cervical dislocation by inserting a 21-gauge needle into the shaft of the bone and flushing it with Iscove’s modified Dulbecco’s medium (IMDM; Sigma-Aldrich, USA) containing 10% fetal bovine serum (FBS; Gibco, USA), 0.5 μM hydrocortisone, 10,000 U/mL penicillin, and 10,000 μg/mL streptomycin. The study was approved by the Ethics Committee of Universidade Federal de Santa Catarina and was conducted on a total number of 108 mice.

Preparation of imatinib mesylate

Stock solutions of 10 mM IM were prepared by dissolving IM (Gleevec®, Novartis, Switzerland) in a solution of dimethyl sulfoxide and Milli-Q water (1:1) and stored at -20°C.

Clonogenic assay of hematopoietic progenitor cells

Bone marrow mononuclear cells were separated by density gradient with Histopaque (1.077 g/mL) according to manufacturer instructions and resuspended in culture medium. The progenitor assay was performed in a semisolid medium on sterile 6-well culture plates. The medium was plated as a base layer composed of 0.3 mL 2X concentrated IMDM, 0.3 mL FBS, and 0.4 mL 1.25% agar in a total volume of 1 mL/well. The cell layer was prepared in 0.2 mL 2X concentrated IMDM supplemented with 1% bovine serum albumin, 0.3 mL FBS, 0.3 mL
1.25% agar, 0.1 mL mononuclear cells in BM (1 × 10^5 mononuclear cells/mL), 0.1 mL GM-CSF (5 ng/mL; Sigma-Aldrich), and various amounts of IM (1.25-25 μM). All experiments were performed in triplicate. Cells cultured without the drug were used as the control group for the experiments. The plates were incubated for 7 days at 37°C in a humid atmosphere containing 5% CO2. After this period, the number of colonies was counted by light microscopy and colony morphology was assessed after staining with May-Grünwald and Giemsa. Cells were considered to have formed a colony when consisting of more than 50 cells, and 10-50 cells were considered to be a cluster.

**Culture of BM stromal cells**

Originally investigated for their role in the formation of hematopoietic cell microenvironment in BM, MSCs and mesenchymal progenitors have the ability to differentiate into cell types phenotypically unrelated to the cells of their tissue of origin. Due to the large number of similar terms and definitions used (21), we chose to use the term “bone marrow stromal cell”, encompassing all types of cells that are present in the microenvironment of BM (except hematopoietic cells) and that grow and are adherent in culture. We cannot refer to MSCs because we did not perform tests to prove this assertion, but these are cells derived from MSCs.

The cells obtained from BM were cultivated in IMDM and maintained in culture on 24-well plates or in 75-cm² culture flasks. The number of cells used was 2 × 10^6 cells·1 mL⁻¹·well⁻¹ and 6 × 10^7 cells·30 mL⁻¹·flask⁻¹. The plates and flasks were incubated for 14 days at 37°C in a humid atmosphere containing 5% CO2, and were permitted to reach 70-80% confluence. Routine maintenance culture was performed every 3-4 days, when half the volume of the supernatant was removed and replaced with an equal volume of IMDM with IM. This technique assured that suspended hematopoietic cells were removed from the culture medium and only the BM stromal cells adherent to plates/flasks were cultivated. Cells cultured without the drug were used as control. The following experiments were carried out on these cultures: 3,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT), cytochemistry to identify adipocytes, and α-SMA. Cultures were evaluated morphologically in parallel after staining with May-Grünwald and Giemsa.

**MTT assay for cell viability**

Cell viability was determined using MTT, which is an indicator of mitochondrial function. Cells were incubated for 14 days as described above, without IM (control group) and with 5-25 μM IM. After this period, the supernatant from culture wells was removed and IMDM with a 10% MTT solution (Sigma-Aldrich; 5 mg/mL in PBS) was added. After 3 h of incubation at 37°C, the supernatant was removed. The precipitates formed in each well were dissolved in an acid solution of isopropanol (0.04 N HCl) and the supernatant was transferred to a 96-well plate. The absorbance spectrum was determined using a Microwell Systems spectrophotometer (Organon Teknika, Belgium) at a wavelength of 540 nm. The absorbance of the control group (cells without treatment) was considered to correspond to 100% viability.

**AO and EB fluorescence staining of viable cells and apoptotic cells**

Like the MTT test, AO and EB fluorescence also reveal viable cells, and allow visualization of the morphology of these cells beyond the specific staining of cells in the apoptotic process. Apoptotic cells were visualized using two fluorescent dyes: AO, which is permeable to the membrane, and EB, which is impermeable. First, the cells were cultured on circular coverslips 13 mm in diameter for 14 days without IM (control group) and with 5-20 μM IM (experimental group). After this period, the circular coverslip was placed on a microscope slide. The reagent was prepared at the time of the test by mixing equal volumes of AO (10 mg/mL) and EB (10 mg/mL). The coverslip with cells and fluorescent dyes was covered with another coverslip and incubated for 10 min at room temperature (RT). The slides were observed under an Olympus BX41 immunofluorescence microscope and photographed.

**Cytochemistry to identify adipocytes**

For the identification of adipocytes in culture, staining was performed using oil red O, a soluble dye. After 14 days of incubation without IM and with 2.5-25 μM IM, whole culture medium was removed. The wells were washed twice with PBS 7.4 and the cells were fixed with 500 μL 10% formalin/well and incubated at RT for 30 min. Formalin was then removed and the cells were washed twice with PBS 7.4. Next, 200 μL 0.3% solution of oil red O (Sigma-Aldrich)/isopropanol diluted with distilled water (3:2) was added to each well and incubated for 10 min at RT. The dye was then removed and the cells were quickly washed with 60% isopropanol followed by washing with distilled water until excess dye was completely removed. After drying, the cells were visualized with the Olympus CX21 light microscope and photographed.

**Cell cycle analysis**

To evaluate the effects of IM on the phases of the cell cycle, we used a propidium iodide (PI)/RNase Solution Kit (Immunostep, Spain) whose action involves coloring staining nucleic acids with PI after RNA digestion. The cells were initially plated in 75-cm² culture flasks and incubated for 14 days without IM (control group) and with 5-20 μM IM (experimental group). After this period, cells were detached with 2 mL 0.25% Trypsin-0.02% EDTA

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Effects of imatinib mesylate (IM) on the cell cycle

The bone marrow cells of mice were incubated for 14 days in the incubator with 1% PBS. Soares et al. (2013) incubated 500 μL cells that were in some stage of the cell cycle was insufficient for flow cytometry analysis. Most cells were already dead and detached from the culture flasks. A negative control was prepared with cells without drug treatment for the characterization of the parameters to be analyzed by flow cytometry. To draw the histograms of the cell cycle, at least 10,000 living cells were necessary. The histograms were analyzed with WinMDI 2.8 (data not shown) and the results are presented in Table 1.

Cell proliferation and α-SMA expression

The proliferation of cultured cells was evaluated by testing the incorporation of BrdU, a thymidine analogue, into the DNA of proliferating cells during the S phase of the cell cycle. The presence of α-SMA was used for the identification of fibroblasts because, when grown in culture, fibroblasts from different organs change to myofibroblasts and show various degrees of expression of α-SMA. After 14 days of culture, according to the protocol for cultivation of BM stromal cells without IM (control group) and with 5–25 μM IM, cells were incubated for 24 h in a 5% CO₂ incubator with 1 μg/mL BrdU (Invitrogen, USA). After incubation, cells were fixed with 4% paraformaldehyde for 30 min at RT. Next, the cells were washed twice with distilled water and incubated with 2 N HCl for 30 min at 37°C. Cells were rinsed 3 times with PBS 7.4 and cell membranes were permeabilized with 0.25% PBS-Triton X-100 for 30 min at RT.

Table 1. Effects of imatinib mesylate (IM) on the cell cycle phases of bone marrow stromal cells of mice.

| IM (μM) | Cells detected |
|--------|----------------|
|        | G₀/G₁ | S | G₂/M |
| Control | 67.21 ± 0.91 | 20.29 ± 4.81 | 12.5 ± 3.9 |
| 5       | 77.72 ± 3.90 | 15.88 ± 8.72 | 6.4 ± 4.8 |
| 10      | 86.73 ± 3.36 | 7.76 ± 0.03 | 5.5 ± 3.4 |
| 15      | 90.21 ± 8.09 | 6.44 ± 2.38 | 3.35 ± 4.03 |
| 20      | #     | #   | #     |

Data are reported as means ± SEM of the percent of live cells. The bone marrow cells of mice were incubated for 14 days in the absence (control) and in the presence of IM. The number of live cells that were in some stage of the cell cycle was insufficient for flow cytometry analysis. Most cells were already dead and detached from the culture flasks. Nonspecific sites were blocked with 5% FBS for 40 min, and the cells were subjected to an immunocytochemical reaction using specific primary and secondary antibodies. Incubation with primary antibodies (anti-BrdU IgG1 1:100, Calbiochem, Germany; anti-α-SMA IgG2a 1:800, Sigma-Aldrich) was performed for 1 h at RT. Cells were then washed 3 times in 0.05% PBS-Tween 20 and incubated for 1 h with secondary antibodies (anti-mouse IgG1, TxRd, 1:75, and anti-mouse IgG2a, FITC, 1:75; Southern Biotech, USA). After 3 further washes with PBS-Tween, cells were incubated with the nuclear stain DAPI (50 ng/mL; Sigma-Aldrich) for 15 s at RT for analysis of total cell nuclei. Fluorescent staining was observed and photographed using the epifluorescence microscope Olympus BX 40. BrdU, DAPI and α-SMA were simultaneously viewed in cells using different microscope filters. All fields were photographed, i.e., the total area of the well culture was photographed. Fields already photographed acquire a change in the fluorescent color; therefore, the fields are not observed more than once. The overlay of images was performed using a picture editor software.

Statistical analysis

Data are reported as means ± SEM. One-way analysis of variance (ANOVA) with the Tukey post hoc test was performed for statistical analysis. The calculations and graphs were performed and generated using the Statistica 6.0 software (StatSoft Inc., USA). P values <0.05 were considered to be statistically significant.

Results

Effects of IM on CFU-GM hematopoietic progenitor cells

Cultured BM mononuclear cells were analyzed using an inverted microscope and the presence of colonies composed of granulocytes and/or macrophages was confirmed. As shown in Figure 1, IM inhibited the formation of CFU-GM (CFU-G/CFU-M) colonies at low concentrations (1.25 μM), with approximately 22% inhibition compared to the control group (100%), maintaining a similar inhibitory profile at concentrations up to 20 μM. Moreover, with respect to morphology, there was a gradual decrease in the size of CFU-GM colonies (Figure 2).

Morphology of BM stromal cells

After culture with IM for 14 days, mouse BM stromal cells showed an apparent change in general morphology (Figure 3). Many cells in the control group showed fibroblastic features, large nuclei, and an abundant and elongated cytoplasm (Figure 3, Panel A). The cultures treated with IM gradually presented round cells with smaller nuclei and less abundant cytoplasm, features suggestive of macrophages (Figure 3, Panels B and C).
This general morphological modification progressed in a concentration-dependent manner.

**MTT assay for cell viability**

The MTT test was performed to assess the viability of BM stromal cells incubated for 14 days with 5-25 μM IM. At doses of 5 and 10 μM IM the number of viable cells increased by 18 and 73%, respectively, compared to the control group (100%). However, after incubation with 15 μM IM, this percentage fell to 89%, and with 25 μM no cells were viable (Figure 1).

**Cell viability and apoptotic cells - AO/EB**

The number of viable BM stromal cells from mice decreased with higher IM concentrations (15-25 μM), as demonstrated by the MTT assay. As seen in Figure 5, the control group (Panel A) almost universally showed a greenish color, which is indicative of viable cells. Panels B and C illustrate that cells incubated with 10 and 20 μM IM, respectively, had some orange color, primarily in the nucleus, which is characteristic of apoptotic cells. In cultures treated with IM, there were more rounded cells (macrophages-like) along with the presence of apoptotic cells. At 10 μM IM (Figure 5, Panel B), there were greater numbers of cells compared to control, but a considerable number of them were entering apoptosis (orange color only in the nucleus). At 20 μM (Figure 5, Panel C) there were few cells in general, and many apoptotic cells, some containing apoptotic bodies. At 25 μM IM there were no cells.

**Cytochemistry for the identification of adipocytes**

As shown in Figure 6, the adipose cells showed reddish lipid vacuoles of varying sizes inside the cytoplasm. At concentrations of 2.5-25 μM, IM had a 100% inhibitory effect on adipose cells and only the control group contained cells with lipid content revealed by oil red O staining (Figure 6, Panels A and B), a result repeatedly observed in experiments performed under the same conditions. Quantitative analysis was not performed since the inhibition was complete.
As shown in Table 1, IM caused a progressive increase in the number of BM stromal cells in the $G_0/G_1$ phase of the cell cycle compared to control and consequently reduced the number of cells in the $S/G_2/M$ phases. Note that even with 5 $\mu$M IM, only 50% of cells (compared to control) were entering mitosis. The greatest blocking effect was observed in the $G_0/G_1$ phase of the cell cycle with an IM concentration of 15 $\mu$M.

Assessment of cell proliferation by BrdU incorporation

The proliferation of BM stromal cells from mice was evaluated by testing BrdU incorporation (nuclei with red color) and the total number of cells was identified by staining with DAPI (nuclei with blue color). The percentage of proliferating cells (Figure 7) was obtained by quantifying cells in the photographed fields (Figure 8) and comparing the count to the control group. Figure 7 shows that IM caused a significant decrease in cell proliferation, a reduction of approximately 50, 80, 95, and 100% at concentrations of 2.5, 5.0, 7.5 to 15, and 20 $\mu$M, respectively.

Evaluation of $\alpha$-SMA expression

$\alpha$-SMA positivity (Figure 9) was determined by counting all $\alpha$-SMA-positive cells (green cells) treated with IM present in each culture well (Figure 8) and compared to the control group. Figure 9 shows that there was a large decrease in the amount of cells expressing $\alpha$-SMA in the treated groups. IM caused a decrease in the expression of $\alpha$-SMA in the stromal cells to 45.74 and 2.51% at concentrations of 2.5 and 7.5 $\mu$M, respectively, and completely inhibited expression at 10 $\mu$M. By overlaying the images, it was discovered that most $\alpha$-SMA-positive cells were also positive for BrdU, confirming the anti-proliferative effect of IM in fibroblasts.

Ratio of fibroblastic cells and round cells

By quantifying cells with nuclei stained with DAPI and the overlap of $\alpha$-SMA-positive cells, a clear difference in the size of the nucleus was observed between cells with characteristics of fibroblastic cells and round cells. Initially, the control exhibited 5.45 times more round cells than fibroblastic cells. As shown in Table 2, this rate
greatly increased in a dose-dependent manner due to an increase in the number of round cells at the expense of decreasing numbers of fibroblast cells, a fact evident in Figures 3 and 8.

**Discussion**

Bone marrow stromal cells appear to regulate the survival, self-renewal, migration, and differentiation of hematopoietic stem cells through several mechanisms, including cell contact interactions or the production of growth factors, chemokines, and extracellular matrix molecules. Four main cell types of the marrow stromal tissue are known to take part in supporting hematopoiesis: one is of hematopoietic origin (macrophages), whereas the others derive from MSCs (reticular/fibroblastic cells, adipocytes, and osteoblasts) (22). Due to the constant renewal of blood cells, the hematopoietic progenitors/precursors exhibit a high proliferation rate, rendering this system very susceptible to the actions of anticancer drugs, which may directly or indirectly affect hematopoiesis (23,24).

The observation of the anti-proliferative effects of IM in BCR-ABL1-negative hematopoietic cells is supported by observations in patients with Ph-negative myeloproliferative disorders who were successfully treated with IM (23,24). Holtz et al. (14) suggested that IM acts mainly by inhibiting the growth of committed progenitor cells (CFCs) and this inhibitory effect increases significantly with prolonged exposure through mechanisms not dependent on BCR-ABL1. In our study, IM reduced the growth of CFU-GM colonies from doses as low as 1.25 μM (Figure 1) and a decrease in colony size was observed, featuring colonies with ever fewer cells in a concentration-dependent manner (Figure 2).

Imatinib mesylate selectively inhibits SCF receptor c-Kit cell signaling. Although SCF itself does not promote colony formation in vitro, it works in synergy with other growth factors such as GM-CSF, G-CSF, IL-3, IL-6, IL-7, and Epo to stimulate formation of both differentiated progenitor cells and more primitive multilineage progenitor cells of myeloid and erythroid lineages (25). The decrease seen in the CFU-GM clonogenic assay can perhaps be explained by IM c-Kit blocking effects.

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**Figure 4.** Effects of imatinib mesylate on bone marrow stromal cells from mice. The absorbance of the control group represents 100% cell viability. Data are reported as the mean ± SEM obtained from at least 3 independent experiments performed in triplicate under the same conditions. *P < 0.05 compared to control group (one-way ANOVA followed by the Tukey test). The same letters indicate no statistically significant difference between groups (P > 0.05).
In order to assess the viability and morphology of stromal cells, we performed both MTT and AO/EB assays. The results were consistent with those observed by light microscopy, showing a change in the total morphology of adherent cells. In untreated cells (Figure 3, Panel A; Figure 5, Panel A) and cells treated with lower concentrations of IM (5 μM; Figure 3, Panel B), the predominant morphology was typically fibroblastic. At 10 μM, an increase was observed in the number of rounded cells with a considerable number of cells entering apoptosis (orange color only in the nucleus) (Figure 3, Panel C; Figure 5, Panel B). The decrease in cell viability was observed only with high concentrations of IM (15-25 μM). At 20 μM (Figure 5, Panel C) there were few cells in general and many apoptotic cells, some containing apoptotic bodies. The evaluation of the results of the MTT and AO/EB assays suggests that cell viability showed a peak (Figure 4) due to the increased number of rounded cells, indicating a possible proliferative effect of IM on these cells, herein referred to as macrophage-like cells (also seen in Figure 8, Panels H and J, with DAPI), and an opposite effect on fibroblast cells confirmed by expression of α-SMA (Figure 8, Panels C, F, I, and K). These results can be related to the data in Table 2, which shows that there was a progressive increase in the ratio of round cells to fibroblastic cells. At the 10 μM dose of IM, the number of round cells was 383 times higher than the number of fibroblast cells, and was reduced only at high concentrations (15-25 μM IM).

In accordance with other studies (26,27), the evaluation of the cell cycle of BM stromal cells with IM showed the blocking of cells in the G0/G1 phases (67.21% in controls and up to 90.21% with 15 μM IM) and a decreased percentage of cells that were in the S (20.29% in controls and up to 6.44% with 15 μM IM) and G2/M phases (12.5% in controls and up to 3.35% with 15 μM IM) (Table 1). The decrease in cells in the S phase...
is consistent with the results of BrdU staining (Figure 7), since BrdU is only incorporated into the nucleus of cells that are undergoing proliferation. In this study, we did not assess the mechanisms involved in blocking the cell cycle; however, the blockade in G0/G1 can lead to apoptosis (seen by the presence of apoptotic bodies stained with EB) (Figure 5, Panel C). In addition, there is a decrease in cells undergoing the mitotic process.

According to Vittal et al. (28), IM induces a concentration-dependent inhibition of fibroblast proliferation with almost complete inhibition at 20 μM. In our study, the proliferation of stromal cells decreased gradually in the presence of IM (Figure 7), reaching 50, 80, 95, and 100% inhibition at concentrations of 2.5, 5.0, 7.5 to 15, and 20 μM, respectively. Confirming the anti-proliferative effects of IM on fibroblasts, the expression of α-SMA, a myofibroblast marker, in the BM stromal cells on day 14 was strongly inhibited (Figure 9). At 2.5 μM, the expression was reduced by approximately half, at 7.5 μM IM, it decreased to 2.51%, and was almost completely inhibited at 10 μM. By overlapping the images (Figure 8, Panels C, F, I, and K), we noted that most cells expressing α-SMA with large nuclei were also in the process of proliferating (BrdU-positive), an event that occurred in the absence, or at very low concentrations, of IM.

It has been demonstrated in the literature that the presence of adipose cells in culture is associated with increased survival of the hematopoietic system, which can serve as a positive indicator of the culture success in maintaining primordial cells (29). Our results demonstrated that adipocytes stained with Oil red O only in the control cultures of BM stromal cells (Figure 6, Panels A and B), suggesting that IM has an effect on the differentiation of stromal cells into adipocytes or, perhaps, induces apoptosis of pre-existing adipocytes.

Cells morphologically characterized as macrophage-like, which show reduced colony formation by CFU-GM clonogenic assay, seem to increase in numbers in stromal cultures. The mechanism by which this happens is not entirely clear, since studies have shown that IM inhibits c-Kit, but does not inhibit c-fms, the receptor for CSF-M (25) and granulocytes/macrophages (CSF-GM) (30), which were used in this experiment. However, studies regarding the origin of resident macrophages in various

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**Figure 7.** Assessment of cell proliferation by BrdU incorporation into bone marrow stromal cells from mice treated with imatinib mesylate. The cells were cultured for 14 days, quantified according to BrdU incorporation into the nucleus and compared to the control group (100%). Data are reported as the mean ± SEM obtained from at least 3 independent experiments performed in triplicate under the same conditions. *P < 0.05 compared to control (one-way ANOVA followed by the Tukey test). The same letters indicate no statistically significant difference between groups (P > 0.05).
tissues have shown that local proliferation plays a significant role in the renewal and maintenance of many types of macrophages, whereas in some cases the recruitment of circulating precursors has little or no role (31). Based on these considerations, there may be a local proliferation of macrophages in tissues, and it is believed that this explains the effects occurring in stroma cultures treated with 10-15 μM IM, as can be seen in Figures 3 (Panel C), 5 (Panel B), and 8 (Panels I and K), where there was an increase of cells morphologically similar to macrophages at the expense of cells with fibroblastic features. If there was proliferation of macrophage-like cells, it would not have occurred on the 14th day of culture since proliferation assessed by BrdU showed a large decrease in a concentration-dependent manner even at lower IM concentrations. This method evaluates the number of cells that proliferate within 24 h after incubation with BrdU.

These results demonstrate that IM is more effective in targeting BM stroma cells, mainly fibroblasts, than hematopoietic cells, since at a dose of 5 μM IM there was growth in only approximately 5% of α-SMA-positive cells while there was 71% growth in CFU-GM cells. These data may be relevant and limiting for IM administration in situations with a large increase in the turnover of cells in the BM stroma due to regeneration after intensive

Figure 8. Analysis of BrdU incorporation and immunofluorescence for alpha-smooth muscle actin (α-SMA) in cells cultured with imatinib mesylate for 14 days. In red, nuclei of cells that incorporated BrdU; in blue, nuclei stained with DAPI; in green, α-SMA-positive cells. Control group cells (A, B, C), cells treated with 5 μM (D, E, F), and 15 μM (G, H, I) imatinib mesylate. No cells treated with 10 μM (J) or 20 μM (L) incorporated BrdU. K, α-SMA-positive cells treated with 10 μM. C, F, I, Representation of the overlapping BrdU, DAPI, and α-SMA images.
chemotherapy/radiotherapy (15), in addition to the direct effect on hematopoietic progenitors involved in myelosuppression. Imatinib mesylate may interfere with the homing of transplanted stem cells by damaging the BM microenvironment of patients previously treated with IM and increase the side effects such as myelosuppression and liver toxicity (32), although there are studies that contradict this hypothesis.

Imatinib mesylate is also being tested for the treatment of several diseases with beneficial results. Studies have shown that IM is a potent inhibitor of PDGF-b signaling and proliferation of fibroblasts and has, therefore, been proposed as a potent antifibrotic therapeutic agent that may not only halt the progression but also induce the regression of tissue fibrosis (17,33-36).

Imatinib has also demonstrated antiangiogenic effects through inhibition of angiogenesis induced by vascular endothelial growth factor (VEGF), bFGF and PDGF in vivo, and by inhibition of tumor growth (18). In previous studies, our group demonstrated that IM acts to decrease the production of reticular fibers produced by stromal cells, suggesting a main activity in fibroblasts. Furthermore, we observed a correlation between fibrosis and vascularization of BM and cytogenetic response in patients with CML (37).

The effects of IM on different cell types, likely through mechanisms of inhibition of the PDGF receptor and a series of growth factors and cytokines (VEGF, bFGF, IL-6, and IL-8) that are also directly or indirectly affected by IM, may have influenced the results obtained here. For

Table 2. Ratio of fibroblastic cells to round cells of bone marrow stromal cells from mice.

| IM (µM) | Round cells/fibroblastic cells |
|---------|-------------------------------|
| Control | 5.45                          |
| 2.5     | 21.51                         |
| 5.0     | 41.15                         |
| 7.5     | 68.76                         |
| 10      | 383.63                        |
| 15      | 80.71                         |
| 20      | 45                            |
| 25      | 0                             |

The ratio of the amount of fibroblastic cells and round cells was determined by the quantification of cells with nuclei stained with DAPI and the overlap of alpha-smooth muscle actin-positive staining cells. IM = Imatinib mesylate.
instance, low levels of growth factors in culture medium alone can induce a decrease in the number of fibroblast cells; however, the low levels of growth factors could be a consequence of the decrease in fibroblasts.

This study demonstrates the inhibitory effect exerted by IM on different cell types present in the BM, represented by hematopoietic cells by inhibiting the growth of CFU-GM, as well as in stromal cells, acting mainly on fibroblasts and adipocytes. These data suggest that IM is more effective on stromal cells than on hematopoietic cells. Thus, in view of the numerous studies that have been performed, the present investigation demonstrates the importance of knowing the effects of IM on non-leukemic cells, since it is a drug used as first-line therapy, for prolonged use in CML and GIST, and is being tested in various other Ph-negative pathologies.

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