

We have assessed the effect of platelet-activating factor (PAF), a biologically active phospholipid present in the human marrow, on the growth of human marrow and blood CD34⁺ progenitors. While the metabolization rate of PAF by CD34⁺ cells is low (weak acetylhydrolase and acylation processes) it is readily catabolized by the acetylhydrolase activity present in the growth medium (10% fetal calf serum + 10% 5637-conditioned medium). Treatment of marrow CD34⁺ cells with the non-metabolizable PAF agonist C-PAF (1 nM to 100 nM) immediately before semi-solid culture significantly ($P < 0.01$) decreased the number of BFU-E but not of CFU-GM colonies. Treatment of marrow or blood CD34⁺ cells with C-PAF (10–100 nM) for 3 days in liquid medium before semi-solid culture significantly ($P < 0.01$) decreased the number of BFU-E and CFU-GM colonies. Treatment of blood CD34⁺ cells with the two PAF receptor antagonists CV 3988 and BN 52021 (1 μ M) had no significant effect on the number of BFU-E and CFU-GM colonies suggesting no role of endogenous PAF in these processes. These results show that exogenous PAF downregulates human erythropoiesis and myelopoiesis, a result that might be of importance during inflammatory states.

Key words: PAF, CD34⁺ cells, Myeloid progenitor, Erythroid progenitor

Effect of platelet-activating factor on the growth of human erythroid and myeloid CD34⁺ progenitors

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Introduction

Studies highlight the role of lipidic compounds on human haematopoiesis.^{1–6} They demonstrate that leukotrienes and prostaglandin E upregulate and downregulate human myelopoiesis, respectively. By contrast leukotrienes and prostaglandin E downregulate and upregulate human erythropoiesis, respectively. Platelet-activating factor (PAF) is a phospholipid molecule produced by inflammatory stimulated cells.⁷ PAF levels are regulated by an acetylhydrolase activity (AHA) which converts PAF into lyso PAF which is then reacylated into membrane phospholipids by an acyltransferase activity (ATA).⁷ Numerous studies report the immunoregulatory properties of PAF. Its inhibiting or activating effects are shown in various cell types such as monocytes/macrophages, polymorphonuclear neutrophils, eosinophils and T- and B-lymphocytes.^{7–9} Studies report that complex interrelations exist between PAF and cytokines. Thus, PAF stimulates the production of several cytokines such as interleukin (IL)-1, IL-6 and tumour necrosis factor-alpha (TNF- α) which, in turn, may enhance PAF synthesis.^{7–9}

While PAF is present in the human bone marrow,^{10,11} its effects on the growth of myeloid and erythroid progenitors is not documented. CD34, a surface antigen expressed on haemopoietic stem/

progenitor cells that disappears at later stages of differentiation can be used to select this population.^{12,13} In this study we have examined the effect of PAF on human haematopoiesis by investigating its effects on the growth of granulocyte-macrophage (CFU-GM) and erythroid progenitor cells (BFU-E) from purified human marrow and blood CD34⁺ cells. We also examined PAF metabolism by CD34⁺ cells.

Materials and Methods

Collection and preparation of mononuclear cells

Sternum bone marrow cells were harvested by aspiration into heparinized tubes (Vacutainer system, Becton Dickinson, Meylan, France) from patients referred for diagnosis according to the Helsinki recommendations. Mononuclear cells were isolated by separation on a Ficoll gradient (400 \times g, 20 min) and washed twice with Hanks's balanced salts solution (Gibco, Cergy Pontoise, France). None of these patients had an haematological malignancy and their myelograms were normal. Peripheral blood stem cells were collected from lymphoma patients in complete remission undergoing apheresis for autologous trans-

plantation. Several aliquots used for bacteriological and CFU-GM controls were harvested in freezing tubes from the final apheresis product just before freezing. They were then stored in liquid nitrogen with apheresis autograft bags until used.

Selection of CD34⁺ cells

Mononuclear bone marrow cells were used immediately after Ficoll and washing procedures. Peripheral blood stem cells were rapidly defrosted in a 37°C water bath, then diluted and washed in saline with 4% human serum albumin (HSA). CD34⁺ cells were obtained by magnetic cell sorting (MiniMACS, Tebu, Le Perray en Yvelines, France) as previously described.¹⁴ The entire procedure was made according to the manufacturer's instructions. Briefly, cells were resuspended at a concentration of $1 \times 10^8/0.3$ ml. The blocking reagent and the antibody reagent (0.1 ml for 1×10^8 cells) were added to the cell suspension and incubated at 4°C for 15 min. Cells were then washed with 4 ml of washing buffer (PBS/0.5% human serum albumin (HSA)/5 mM EDTA) and centrifuged. Cells were recovered in cold washing buffer (0.4 ml for 1×10^8 cells) and incubated with a suspension of submicroscopic magnetic beads (0.1 ml for 1×10^8 cells) for 15 min at 4°C. After washing, cells were recovered in 0.5 ml of cold washing buffer, filtered, applied to the separation column previously placed in a magnetic field, and washed with 2 ml of buffer. Cells which did not bind CD34⁺ antibody passed through the column while CD34⁺ cells were retained. The CD34⁺ cells were eluted by removing the column from the magnetic field and washing the column with 1 ml buffer. After a new step of washing CD34⁺ cells were recovered in IMDM and counted.

Flow cytometric analysis of selected CD34⁺ cells

Samples were analysed on a Profile Coulter using Epics Profile Software to determine the efficiency of the selection. One-hundred µl of cell suspension was incubated at room temperature for 30 min with 5 µl phycoerythrin (PE)-labelled anti-CD34 (HPCA-2) monoclonal antibody or PE-labelled monoclonal control (mouse immunoglobulin) and washed twice with PBS supplemented with 2% HSA. Fixation with paraformaldehyde (PFA 1%) was performed at the end of staining. The purified population contained $84 \pm 2\%$ of CD34⁺ cells (mean \pm SEM of 17 experiments, range 72–98%).

Semisolid cultures

Two experimental conditions were used for CD34⁺ cells. Firstly, freshly isolated marrow CD34⁺ cells were incubated 30 min at 37°C in 1 ml IMDM with

various concentrations of PAF (Sigma, St Quentin-Fallavier, France) and of the non-metabolizable PAF agonist 1-O-hexadecyl-2-N-methylcarbamyl-sn-glycerol-3-phosphocholine (C-PAF) (Sigma) before methylcellulose culture. Secondly, freshly isolated marrow CD34⁺ cells and frozen blood mononuclear CD34⁺ cells were grown for 3 days in liquid medium with various concentrations of PAF, C-PAF, CV 3988 and BN 52021 (two potent PAF receptor antagonists) (Tebu) or the appropriate vehicle (10 µl of 2% HSA) before colony formation in methylcellulose cultures. Marrow or blood CD34⁺ cells (4×10^3) were grown in 96-round-bottomed well microtitre plates in 100 µl IMDM with 10% fetal calf serum (FCS) (Gibco), 10% 5637-conditioned medium and antibiotics (100 U/ml penicillin and 100 µg/ml streptomycin) at 37°C in 5% CO₂ in air.

Colony assays were performed in both experiments with CD34⁺ cells (5×10^2) cultured in 35 mm Petri dishes in sixplicates for 14 days at 37°C in 5% CO₂ in air in 100% humidity. The culture medium (1 ml/dish) was a mixture of IMDM with 0.8% methylcellulose, 30% FCS, 1% detoxified bovine serum albumin, 10 ng/ml Stem Cell Factor (Tebu), 40 ng/ml interleukin-3 (Sandoz, Rueil-Malmaison, France), 3 U/ml erythropoietin (R&D systems, Oxfordshire, UK), α -thioglycerol (0.1 mM) and 10% 5637-conditioned medium. CFU-GM and BFU-E derived colonies were counted under an inverted microscope in the same dish. The differences of colony number between controls and test dishes were analysed using Wilcoxon's test.

[³H]PAF metabolism and AHA assay

CD34⁺ cells (1×10^5) were washed twice with HBSS and were incubated at 37°C in 1 ml of IMDM for various periods of time in the presence of [³H]alkyl-PAF (final concentration 0.05 nmol/tube; 0.5 µCi) (Amersham) complexed to HSA (final concentration 2 mg/ml). Experiments were performed in duplicate. The PAF metabolism was assessed after lipid extraction.¹⁵ Recovery of added radioactivity after lipid extraction was 85%. The labelled compounds derived from [³H]alkyl-PAF were separated using TLC plates (Silica Gel 60 (20 × 20 cm, 0.25 mm)) eluted with chloroform/methanol/acetic acid/water (50:25:8:4, v/v).¹¹ Each lane was divided in areas of 0.5 cm length which were scraped into vials and radioactivity was measured on a Packard liquid scintillation counter. PAF, lyso PAF, phosphatidylcholine (PC) and neutral lipid (NL) were used as standards. AHA was assessed in culture medium according to the method of Miwa *et al.*¹⁶ Results were expressed as picomoles PAF degraded per min per ml as means of duplicate determinations. The variation between duplicates was less than 5%.

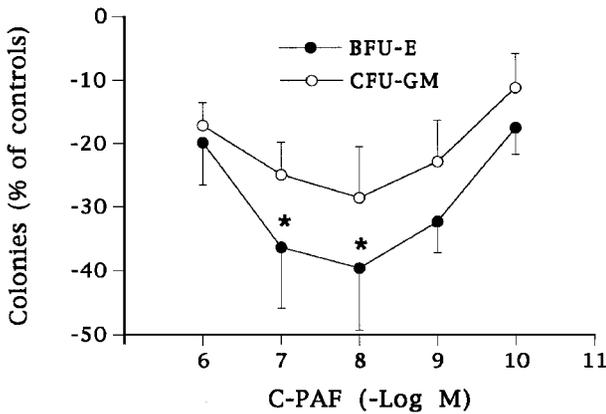


FIG. 1. Direct effect of C-PAF on the number of BFU-E and CFU-GM colonies derived from freshly isolated marrow CD34⁺ cells. Percentage of variation of colony numbers compared with control values (seven independent experiments). * $P < 0.01$ (Wilcoxon's test). Colony numbers in controls were 37 ± 7 and 85 ± 11 for BFU-E and CFU-GM, respectively.

Results

We first investigated PAF catabolism by human CD34⁺ cells. Cells were incubated with [³H]alkyl PAF in IMDM alone. After 24 h of incubation 73%, 18%, 5% and 4% of the label migrated with PAF, lyso PAF, 1-alkyl analogue of PC and NL, respectively (mean of three experiments). When cells were incubated in IMDM + 10% FCS + 10% 5637-conditioned medium (growth medium), 0%, 90%, 6% and 4% of the label migrated with PAF, lyso PAF, 1-alkyl analogue of PC and NL, respectively. After 1 h of incubation in growth medium, 100% of the [³H]PAF was converted into [³H]lyso PAF demonstrating a high AHA. The mean AHA levels detected in IMDM with 10% 5637-conditioned medium, 10% FCS, and 10% 5637-conditioned medium + 10% FCS were 204 ± 16 , 1676 ± 190 and 2160 ± 115 pmol/min/ml, respectively (mean \pm SEM of three experiments). No significant AHA was detected in IMDM alone.

We next assessed the effect of PAF and C-PAF on the growth of human CFU-GM and BFU-E. Due to its rapid metabolism, no significant effect was found with PAF. By contrast treatment of marrow CD34⁺ cells with the non-metabolized PAF agonist C-PAF (1–100 nM) significantly ($P < 0.01$, seven independent experiments) decreased the number of BFU-E but not of CFU-GM colonies (Fig. 1). Treatment of marrow CD34⁺ cells with C-PAF (10–100 nM) for 3 days in liquid medium before semi solid cultures, significantly ($P < 0.01$, three independent experiments) decreased the number of BFU-E and CFU-GM colonies (Fig. 2A). In similar experimental conditions, C-PAF significantly ($P < 0.01$, three independent experiments) decreased the growth of BFU-E and CFU-GM colonies from blood CD34⁺ cells (Fig. 2B). In these experimental conditions the PAF receptor antagonist CV 3988 (1 μ M)

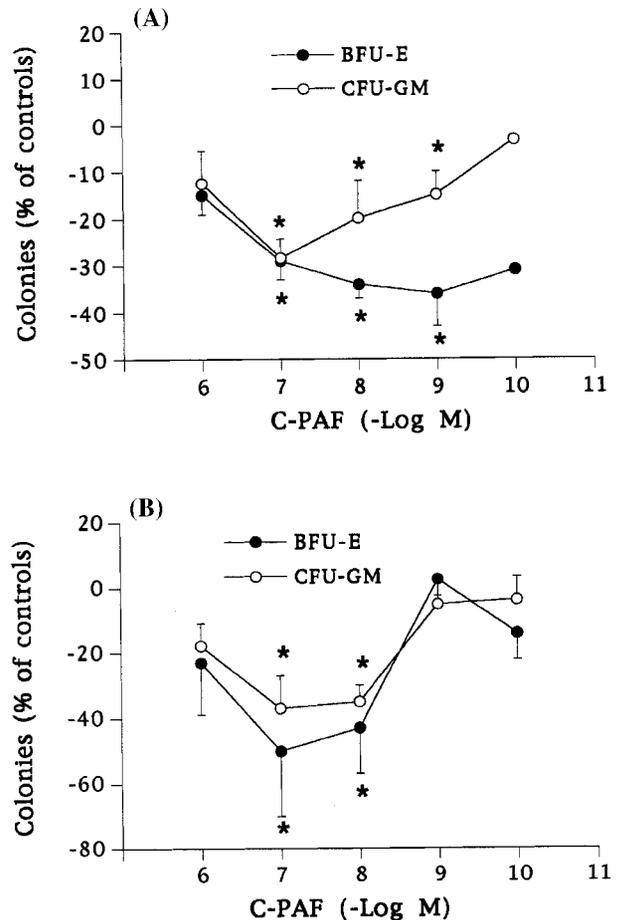


FIG. 2. Effect of C-PAF on the growth of BFU-E and CFU-GM colonies from marrow (A) and blood (B) CD34⁺ cells cultured 3 days in liquid medium before colony assay. (A) Percentage of variation of colony numbers compared with control values (three independent experiments). * $P < 0.01$ (Wilcoxon's test). Colony numbers in controls were 28 ± 4 and 51 ± 3 for BFU-E and CFU-GM, respectively. (B) Percentage of variation of colony number compared with control values (three independent experiments). * $P < 0.01$ (Wilcoxon's test). Colony number in controls were 16 ± 2 and 37 ± 3 for BFU-E and CFU-GM, respectively.

slightly ($P < 0.05$, four independent experiments) enhanced the growth of BFU-E colonies compared with control dishes (23 ± 2 colonies vs. 19 ± 2 colonies for CV 3988-treated cells and controls, respectively) while BN 52021 (1 μ M) had no effect (20 ± 2 colonies). CV 3988 and BN 52021 had no effect on CFU-GM growth (54 ± 5 colonies and 56 ± 5 colonies vs. 52 ± 5 colonies for CV 3988-treated cells, BN 52021-treated cells and controls, respectively).

Discussion

Lipidic mediators (such as prostaglandins and leukotrienes) affect the growth of human CFU-GM and BFU-E progenitors.^{1–6} The rationale for analysing the role of PAF on the growth of human myeloid and erythroid progenitors is based on several points. First, PAF is

present in the human bone marrow.^{10,11} Second, PAF modulates the growth of various human cell types including adherent bone marrow cells.¹⁷ Third, PAF stimulates the eosinophilic and basophilic differentiation from human blood progenitors.¹⁸ Finally, PAF increases the erythroid colony formation from cultured CD34⁺ haematopoietic progenitors by enhancing the prolactin synthesis by human bone marrow stromal cell feeders.¹⁹

The use of the CD34 antigen is a useful way to obtain haematopoietic stem/progenitor cells.^{12,13} It is expressed on 1–3% of adult bone marrow cells,²⁰ and about 0.1% of normal blood mononuclear cells.²¹ As the population of bone marrow CD34⁺ progenitors differ from that in blood by a higher percentage of more committed progenitors,²² we have investigated the effect of PAF on both marrow and blood CD34⁺ cells. Our percentage of CD34⁺ cells after magnetic cell sorting is similar to recent studies using this method.^{14,23} All CD34⁺ cells are not haematopoietic progenitors. Thus, cells that give rise to fibroblast-like stromal elements of the bone marrow may be recovered in the CD34⁺ cell fraction but represent less than 5% of total CD34⁺ cells.²⁴ Moreover colonies of fibroblastic cells were not documented in our CD34⁺ cell cultures.

Our studies are done with a non-metabolizable PAF agonist since PAF is readily catabolized by an AHA present in the culture medium. By contrast the metabolization rate of PAF by human CD34⁺ cells is low (weak intracellular AHA and ATA). The nanomolar concentrations of C-PAF active in this study are physiologically significant since PAF is detected in the human marrow at a mean level of about 1 nM.¹¹ The use of 1 nM of C-PAF may be compared with a continuous stimulation with this physiologic dose of PAF. In the present study treatment of marrow CD34⁺ cells with C-PAF immediately before semi solid culture decreased the growth of BFU-E but not of CFU-GM colonies suggesting a direct effect of C-PAF on human erythroid progenitor cells. An effect of C-PAF on marrow or blood CFU-GM colonies was found after 3 days of treatment in liquid medium before semi solid culture suggesting an indirect effect through an elevated production of inhibitory factor(s) or a decreased production of stimulatory factor(s). *In vitro* studies report that PAF modulates the production of several molecules such as IL-1, IL-3, IL-4, IL-6, TNF- α , prostaglandins and leukotrienes that regulate human CFU-GM growth.^{7–9,25} No role of endogenous PAF may be documented on the growth of CD34⁺ cells. Thus, the two PAF receptor antagonists CV 3988 and BN 52021 have no effect on the number of CFU-GM colonies. In contrast to BN 52021, CV 3988 has a weak effect on BFU-E colonies. A similar difference between these two drugs has been found concerning their ability to inhibit superoxide anion generation in a human B cell line.²⁶ In this previous study the weak

effect of CV 3988 was attributed to its incorporation in cell membrane leading to a non-specific effect. Such an effect may also be suggested in our study.

Bellone *et al.*¹⁹ have recently reported that PAF stimulates the erythroid colony formation from CD34⁺ haematopoietic progenitors by enhancing the prolactin synthesis by human bone marrow stromal cells. Our present results indicate that, in absence of a marrow stromal cell feeder, PAF decreases the growth of erythroid and myeloid CD34⁺ progenitors highlighting not only the putative role of PAF during early steps of human haematopoiesis but also the complexity of the regulation of the bone marrow progenitor growth.

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