Production of oxygen free radicals by Ehrlich ascites tumour cells: effect of lipids

Gopal K. Marathe and Cletus J. M. D'Souza

Department of Studies in Biochemistry, Manasagangotri, University of Mysore, Mysore-570 006, India

Corresponding Author

Introduction

Phagocytic cells kill invading microorganisms in a metabolic event characterized by a marked increase in oxygen consumption termed 'respiratory burst'.1 The membrane-bound enzyme NADPH-oxidase catalyzes this reaction.2 This enzyme is dormant in resting cells, but can be activated by a wide variety of stimulants.2,3 During the activation of NADPH-oxidase, oxygen free radicals (like the superoxide anion \( \text{O}_2^- \)) are generated.1,2 Naturally occurring antioxidants like vitamin E, vitamin C and vitamin A can inhibit free radical generation.4,5 In addition to these vitamins, membrane lipids are also known to modulate free radical generation.6 As vitamin A has been known to affect membrane integrity7 and to impair immunity,8 there may be a relationship between membrane lipids, vitamin A and free radicals.

In order to test this relationship, the free radical generation in Ehrlich ascites tumour cells (EAT cells) using stimulants like phorbol-12-myristate-13-acetate (PMA) calcium ionophore A23187, and PAF in cells grown in vitamin A deficient and vitamin A sufficient animals has been studied. The effect of various lipids and stimulants on free radical generation in this cell line is reported in this paper.

Materials and Methods

Chemicals: Platelet activating factor (1-O-hexadecyl-2-acetyl-Sn-glycero-3-phosphocholine (PAF), calcium ionophore A23187, phorbol-12-myristate-13-acetate (PMA), phosphatidylserine (PS), phosphatidylethanolamine (PE), phosphatidylinositol (PI), phosphatidylcholine (PC), phosphatidylglycerol (PG), sphingomyelin (SM), sphingosine (SS), phosphoinositides (PIP), and superoxide dismutase (SOD) of bovine erythrocytes were obtained from Sigma Chemical Co., USA. BN 52021 was a generous gift from Dr P. Braquet (Institute Henri Beaufour, France). BN 52021, PMA, and calcium ionophore were dissolved in DMSO. PAF and Lyso-PAF were dissolved in 0.9% NaCl containing 2.5 mg/ml BSA and diluted with Tyrode-Ringer's buffer (pH 7.5) just before use. Nitro-blue tetrazolium (NBT) was from SR Laboratories, India. All other chemicals were of analytical grade and solvents were redistilled before use. The
concentration of DMSO never exceeded 0.1% v/v in these experiments. Such a concentration of DMSO did not interfere with the experiments.

Development of vitamin A deficiency in mice and culture of EAT cells: Weanling Swiss albino mice were made vitamin A deficient as reported earlier. The animals were divided into three groups: group 1, vitamin A deficient; group 2, pair-fed controls; and group 3, controls receiving a normal commercial diet (supplied by Lipton, India Ltd). The pair-fed controls received the same diet as the vitamin A deficient mice, except 120 I.U. of retinyl palmitate was given orally three times a week. EAT cells were cultured in the peritoneal cavity of such mice (12–14 weeks old) by serial transplantation. Viability of the cells was determined by trypan blue dye exclusion and was greater than 95%.

Determination of oxygen free radicals by NBT reduction The release of oxygen free radicals was determined spectrophotometrically by measuring the reduction of NBT at 540 nm. Freshly harvested EAT cells (4 × 10⁶ cells) were suspended in Tyrode–Ringer's buffer (pH 7.5) in a total volume of 1 ml, containing 0.25% BSA. The cells, untreated or stimulated by various stimulants, were incubated with NBT (60 nmol) for 20 min. The cells were washed with the same buffer and lysed by the addition of 2 ml of 1,4-dioxane and maintained in a boiling water bath for 8 min. The lysate was centrifuged at 250 × g for 5 min and the extracted blue colour was read at 540 nm. A calibration curve of absorbance at 540 nm was obtained using PAF (4 μM), PMA (2 μM) and calcium ionophore (2 μM) as stimulants, and different concentrations of NBT (0-80 nmol) on EAT cells (4 × 10⁶).

Addition of lipids: The various lipids in chloroform dried under nitrogen were prepared at two concentrations, 12.5 μg/ml and 50 μg/ml, and sonicated. These lipids were added to cells (4 × 10⁶ per ml) in Tyrode–Ringer buffer (pH 7.5) containing 0.25% BSA. The EAT cells suspended in this buffer were pre-incubated with these lipids (PS, PE, PI, PC, PG, SM, SS and PIP) prior to the addition of PAF (4 μM). The neutral lipid (NL), glycolipid (GL) and phospholipid fractions (PL) from total lipid extract of EAT cells were separated as described in an earlier paper and the effect of these lipids on free radical generation was also studied. The free radicals were also measured in the presence of vitamin A and SOD.

Statistical analysis: The results are expressed as means ± S.D. and significance was assessed using Student's t test.

Results

Production of vitamin A deficiency in mice: Although the literature with respect to vitamin A deficiency in rat is extensive, reports on vitamin A deficiency in mice are scanty. In fact, it was difficult to develop vitamin A deficient mice. Even though we were able to develop severe vitamin A deficiency in mice, the animals could not survive when EAT cells were injected intraperitoneally. Hence, a marginal vitamin A deficient condition was chosen in this study. It took 10–12 weeks to develop such a deficiency.

Stimulation of free radicals in EAT cells by PAF and its inhibition by BN 52021: A dose response curve showing the effect of increasing PAF concentrations on free radical generation in EAT cells grown in vitamin A deficient, pair-fed controls and in the control mice receiving the commercial diet is shown in Fig. 1. Although the optimal level of PAF required to stimulate the free radical generation is 4 μM in all three groups, the basal level of free radicals is more in vitamin A deficient mice (10.5 ± 10.8 per 10⁶ cells) compared with control cells (5.8 ± 1.50 per 10⁶ cells). The production of free radicals was a maximum at 20 min and then gradually declined (Fig. 2). The specific PAF antagonist BN 52021 progressively inhibited the free radical generation over the concentration range of 5–50 μM (Fig. 3).

Effect of lipids on free radical generation: The effect of various lipids on the generation of free radicals in EAT cells is shown in Fig. 4. When PE, PS, PI, PG and PIP were used in the incubation medium prior to the addition of PAF (4 μM), there was a...
stimulation of free radical generation. PS was the most effective of the phospholipids used \((p < 0.001)\). However, PC at the lower concentration \((12.5 \mu g/ml)\) was inhibitory \((p < 0.001)\), while at a higher concentration \((50 \mu g/ml)\) the stimulatory effect was not significant \((p > 0.09)\). Sphingolipids were inhibitory at both the concentrations used \((p < 0.001)\). A more pronounced effect was seen when the phospholipid fraction (PL) derived from a total lipid extract of EAT cells was used. The free radical generation was increased 2.8 fold at a concentration of \(12.5 \mu g/ml\) \((p < 0.001)\). On the other hand, glycolipid (GL) and neutral lipid (NL) fractions derived from EAT cells were ineffective in free radical generation in PAF stimulated cells \((p > 0.05)\). The effect of these lipids on un-stimulated cells was examined, but they did not affect the basal level of free radicals (data not shown).

Comparison of the free radical generation by different stimulants: The effect of various stimulants on free radical generation is shown in Table 1. PMA appears to be the most effective activator of the respiratory burst, while PAF was the least. Exogenous addition of vitamin A and SOD suppress the enhanced respiratory burst, and also affect the basal levels of free radicals. However, lyso-PAF (the biologically inactive metabolite of PAF) totally failed to cause respiratory burst at both the concentrations used.

Discussion

PAF has been reported to activate the respiratory burst in various cells such as macrophages, neutrophils and eosinophils.\(^{14}\) In contrast, certain investigators could not detect the effect of PAF on free radical generation in human monocytes and rat alveolar macrophages.\(^{14}\) Thus, some uncertainty still persists as to the role of PAF in free radical
Table 1. Effect of different stimulants on the production of free radicals in EAT cells grown in mice receiving a commercial diet under various conditions

| Stimulus                  | Treatment | nmol of NBT reduced/10^6 cells/20 min |
|---------------------------|-----------|-------------------------------------|
| None                      |           | 6.25 ± 1.20                         |
| PAF (4 μM)                |           | 22.0 ± 2.60                         |
| Calcium ionophore A23187  | (2 μM)    | 42.3 ± 3.20                         |
|                           | (6 μM)    | 51.0 ± 7.45                         |
| PMA                       | (1 μM)    | 36.4 ± 3.20                         |
|                           | (2 μM)    | 74.1 ± 2.58                         |
|                           | (4 μM)    | 85.0 ± 3.40                         |
| Lyso-PAF                  | (5 μM)    | 6.45 ± 1.35                         |
|                           | (10 μM)   | 6.95 ± 1.20                         |
| None                      | Vitamin A (10 μg/10^6 cells) | 4.20 ± 0.50                         |
|                           | Vitamin A (20 μg/10^6 cells) | 3.31 ± 1.12                         |
| PAF (4 μM)                | Vitamin A (20 μg/10^6 cells) | 9.82 ± 5.20                         |
| None                      | SOD (5 units) | 3.6 ± 1.05                          |
| PAF (4 μM)                | SOD (5 units) | 10.4 ± 1.18                          |

Values are mean ± S.D. (n = 3). (Whenever vitamin A and SOD are used, cells are pretreated with these and then stimulated.)

Generation. It may not be a universal activator, but may act as an activator in a few cells. A new role for PAF has also been assigned in amplifying the respiratory burst induced by other stimulants. In a previous report the authors showed that EAT cells produce PAF (95 pmol per 10^6 cells) on stimulation with calcium ionophore A23187 (10 μM). Here it is demonstrated that PAF produced by EAT cells can act on those cells and generate free radicals. Although the concentration of PAF used in the in vitro assay was far greater than PAF generated in vivo by an equivalent number of cells, it is possible that the local concentration of PAF may be even greater than the one employed in this study. Such an observation has been reported in rabbit leukocytes. The production of free radicals is the main function of the phagocytic cells; its generation in non-phagocytic cells such as human fibroblasts, and transformed cells such as human breast carcinoma and EAT cells (present study), is interesting.

Lipids, especially the phospholipids and their metabolites, seem to play a crucial role in many cell functions, particularly in intracellular signalling. Phospholipids also stimulate a variety of enzyme catalyzed oxidative reactions. In the present study, most of the lipids used activate the respiratory burst oxidase except for sphingolipids which are inhibitory. Such an activation of the respiratory burst oxidase by PS was also observed by Tamura et al.

Protein kinase C has been implicated as essential in activation of NADPH oxidase. However, Tamura et al. have questioned the involvement of protein kinase C, since activation observed during PS addition could not be inhibited by EGTA, which is known to inhibit protein kinase C. Hence, it appears that although direct stimulation of NADPH oxidase by PS and its inhibition by sphingolipids is possible, involvement of protein kinase C cannot be ruled out.

Augmented production of free radicals during vitamin A deficiency is interesting. Vitamin A deficiency is known to alter the membrane integrity and to bring about associated changes including changes in membrane lipid composition. These effects may activate the respiratory burst oxidase. In fact, the basal level of free radicals is more during vitamin A deficiency and could be suppressed by exogenous addition of vitamin A, suggesting a role for vitamin A in free radical generation.

Besides PAF, other stimulants like PMA and calcium ionophore were also capable of eliciting the respiratory burst (Table 1). Although PAF appears to be a weak stimulant, it is a physiological one.

Activation/inhibition of the respiratory burst oxidase displayed by various phospholipids and dietary factors such as vitamin A probably play a regulatory role in free radical generation. As the respiratory burst is lethal for both the invader and the host, it should not be turned on unnecessarily. Membrane lipids and vitamin A probably regulate this event.

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