Mouse Cachexia Induced by Trehalose Dimycolate from Nocardia asteroides

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(Received 21 December 1987)

Trehalose dimycolate (TDM) isolated from Nocardia asteroides induced in mice a severely wasted condition known as cachexia. Intraperitoneal injection of mice with five 10 μg doses of TDM in mineral oil at intervals of 2 d killed 90% of the animals within 26 d. Death followed a precipitous weight loss and an inflammatory process in the peritoneal cavity. When mice were injected intraperitoneally with a single 10 μg dose of TDM, 48 h later, they had begun to lose weight and exhibited extreme hypertriglyceridaemia and hypoglycaemia. Tumour necrosis factor (or cachectin) was detected in the plasma from animals injected with TDM. This cytokine released by mononuclear phagocytes may be involved in the induction of cachexia by TDM.

INTRODUCTION

Trehalose dimycolate (TDM) is one of the products of mycobacteria (see Stewart-Tull, 1983, for a review) that enhances the nonspecific resistance of mice to tumours and to bacterial infections (Bekierkunst et al., 1971; Leclerc et al., 1976; Yarkoni et al., 1978; Parant et al., 1977). Mycobacterial TDM also exerts a toxic effect in experimental animals, leading to cachexia and death (Bloch, 1950). Since the discovery of the toxic TDM from the tubercle bacillus, TDMs have been isolated from strains of Corynebacterium (Thomas et al., 1979; Ioneda & Silva, 1979), Rhodococcus (Silva et al., 1979) and Nocardia (Silva et al., 1979). However, little is known about the biological properties of TDM isolated from Nocardia. Since the chain length of the mycolic acid in TDM from Nocardia is only about 56 carbon atoms (Silva et al., 1979), while it is about 80 carbon atoms in mycolic acids in TDM from mycobacteria, it seemed important to verify whether TDM isolated from N. asteroides would have a toxic effect in mice. The possible role of tumour necrosis factor (TNF) in TDM-induced cachexia is discussed.

METHODS

Bacterial strain and culture conditions. Nocardia asteroides was obtained from a nocardiosis patient at the Clinical Hospital of Ribeirão Preto, University of São Paulo, Brazil, and was maintained on Sabouraud's dextrose agar. It was cultured at 37 °C for 15 d in medium containing 10 g yeast extract, 10 g glucose and 1 g NaCl per litre of distilled water.

Lipid extraction. Bacteria were autoclaved, harvested by centrifugation at 5000 g for 5 min and washed five times with 100 ml distilled water. Lipids were extracted from the killed cells (10 g dry weight) in 300 ml chloroform/methanol (2:1, v/v) by stirring at room temperature for 2 h. The extract was separated by centrifugation at 5000 g for 10 min and the pellet, suspended in the same solvent mixture as described above, was re-extracted three more times as described. The combined extracts were dried on a rotary evaporator and extracted with diethyl ether as described by Silva & Ioneda (1977a) to eliminate non-lipid residues. Briefly, the re-extraction was done by adding 1 vol. 1 M NaCl and 5 vols diethyl ether. The ethyl ether phase was washed with distilled water, dried over anhydrous Na2SO4, filtered and concentrated under reduced pressure.

Abbreviations: TDM, trehalose dimycolate; TNF, tumour necrosis factor.

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Fractionation of lipid extract and purification of TDM. The glycolipid TDM was obtained by a method used previously (Silva et al., 1979). Briefly, the lipid extract, dissolved in a small volume of chloroform, was loaded on a column (2.1 × 28.0 cm) of silicic acid/silica gel H (1:1, w/v) which was successively eluted with 500 ml of each of the following solvents: chloroform; 5% (v/v) ethanol in chloroform; and 60% (v/v) acetone in chloroform. The 60% (v/v) acetone in chloroform fraction was further purified by preparative TLC on plastic sheets coated with silica gel F254 (Merck), using the solvent system chloroform/acetone/methanol/water (50:50:2:5:3, by vol.), yielding a purified glycolipid that had an [α]25 value of +47 (C = 0.5, CHCl3) and a melting point of 38°C. It contained 13% sugar (determined by the phenol/sulphuric acid method; Dubois et al., 1956) and had an infrared spectrum similar to that described for TDM isolated from Mycobacterium tuberculosis (Kato & Asselineau, 1971) and from Nocardia asteroides (Silva et al., 1979). After alkaline hydrolysis (Iioneda et al., 1963), trehalose and mycolic acid were identified in the aqueous and ether phases by TLC and paper chromatography (Silva & Ioneda, 1977b), respectively. Physical and chemical analysis of the isolated mycolic acid showed that the mean carbon chain length was C50, in agreement with previous results (Silva et al., 1979).

Animals. Male Swiss mice 4 to 6 weeks old, and weighing 19.0–21.0 g were used.

Toxicity. The TDM to be tested for toxicity was dissolved in mineral oil (Nujol, from Plough Inc.). Before use, the clear solution was heated to 56°C and subjected to ultrasonic vibration for 5 min at 100 W, using a probe of 9 mm diameter. The toxicity of TDM was evaluated by injecting mice intraperitoneally with different doses in 0.1 ml mineral oil and recording weight loss and deaths. A control group was given 0.1 ml mineral oil alone.

Preparation of plasma. Mice injected with 10 μg TDM were exsanguinated 48 h after TDM injection, and the plasma was obtained. Control plasma was obtained from noninjected mice and from mice injected with mineral oil in a similar manner. All plasma samples were stored at −20°C until used.

Plasma triglyceride and glucose determinations. These were done with triglyceride and glucose kits (Labtest Diagnostica), respectively.

Cytotoxicity assay of TNF. Killing of L929 mouse tumour cells was used to measure the cytotoxicity of soluble factor(s) present in the plasma. This was a standard assay (Ruff & Gifford, 1980). Briefly, L929 cells in RPMI-1640 medium containing 5% (v/v) foetal calf serum were seeded at 3 × 104 cells per well in 96-well microtitre plates (Linbro, Flow Laboratories) and incubated overnight at 37°C in an atmosphere of air/CO2 (95:5, v/v). Serial 1:2 dilutions of plasma were made in the above medium containing 1.0 μg actinomycin D (Sigma) ml−1, and the cell culture medium was replaced with 100 μl volumes of the dilutions in triplicate. The next day, cell survival was assessed by fixing and staining the cells with crystal violet (0.2% in 20% methanol), solubilizing the stained cells with 0.1 ml 1% (w/v) SDS per well and reading the absorbance of each well at 490 nm with a model BT-100 Microelisa Autoreader (Bio-Tek). Percentage cytotoxicity was calculated as [1 − (A490 of sample)/(A490 of control)] × 100.

RESULTS

TDM was injected into mice intraperitoneally at 2 d intervals (five 10 μg doses in 0.1 ml mineral oil), and body weight and mortality were observed for 30 d. Results from a typical experiment are shown in Fig. 1. Mice injected with TDM showed an initial decrease in body weight after the first injection; the animals were reluctant to move, and appeared unwell, with ruffled fur. They had lost 18–26% of their initial body weight by day 12 after the first injection. In addition to inducing cachexia, TDM also caused the death of 90% of the animals between 14 and 28 d after the first injection. Autopsy of the animals injected with TDM showed the presence of peritonitis, with an inflammatory exudate, massive adhesions of the organs in the peritoneal cavity, numerous granulomas and splenomegaly. Control mice given mineral oil alone did not die and showed no toxic effects; they had gained 26% in body weight by day 30 of the experiment. The results obtained were reproducible in three experiments and the pattern of body weight curves and survival rates were very similar.

Cachexia in chronic diseases is associated with a depressed lipoprotein lipase function, resulting in enhanced triglyceride concentration in plasma and enhanced peripheral glucose oxidation (Gallin et al., 1969; Beisel, 1972; Rouzer & Cerami, 1980; Beutler & Cerami, 1986). To examine this effect, mice were injected with 10 μg TDM and killed 48 h later (initial observations showed that 48 h is sufficient to produce cachexia in mice). The triglyceride concentration in the plasma was elevated 2.9-fold, and the plasma glucose concentration was 2.6 times less, in mice injected with TDM compared with control animals (Fig. 2). Killing of tumour cells in vitro is a sensitive bioassay to detect TNF (Beutler & Cerami, 1986). Cytotoxic
Induction of cachexia by trehalose dimycolate

Fig. 1. Mean body weight (± SD) of two groups of ten mice at various times after intraperitoneal injection of five 10 μg doses of TDM in mineral oil (■) or mineral oil alone (○). The mice were injected on the days indicated by arrows. Each cross (±) represents the death of one mouse. It is not possible to examine the results statistically because of the decreasing number of animals in the TDM-injected group. Results are from a typical experiment.

Fig. 2. Plasma triglyceride (open bars) and glucose (hatched bars) concentrations in mice injected intraperitoneally with 10 μg TDM in 0.1 ml mineral oil. Plasma samples were obtained 48 h after injection. Control groups consisted of noninjected mice, and mice injected with mineral oil alone. The data, from a typical experiment, are expressed as mean ± SD for six mice in each group.

Fig. 3. Cytotoxicity for L929 tumour cells of plasma from mice injected intraperitoneally with 10 μg TDM in mineral oil. Plasma samples were obtained 48 h after injection and diluted 1:128 in RPMI-1640 medium. Control groups consisted of noninjected mice, and mice injected with mineral oil alone. The data, from a typical experiment, are expressed as mean ± SD for six mice in each group.

Activity for L929 tumour cells was observed in plasma collected from mice injected 48 h previously with TDM, but not in plasma collected from animals inoculated with mineral oil or from noninjected animals (Fig. 3).
DISCUSSION

Several common physiological and biochemical derangements are seen in the mammalian host responding to a variety of invasive stimuli such as bacterial, viral, and protozoan infections. These responses include a condition known as cachexia in which the animal continuously loses weight, even while consuming an adequate diet. It has been widely supposed that microbial products are directly responsible for the wasting associated with infection (Beutler & Cerami, 1986). The results presented here show that TDM is one of the bacterial products able to induce cachexia in mice: wasting and ultimately death, as well as hypertriglyceridaemia and depressed plasma glucose levels, all occurred after TDM injection; these are all effects associated with cachexia.

In response to a variety of invasive stimuli, reticuloendothelial cells and lymphocytes secrete cytokines that are capable of altering host metabolism. TNF (or cachectin) is one of these cytokines; it has been implicated as being important in inducing shock (Beutler & Cerami, 1986) and cachexia (Rouzer & Cerami, 1980) in animal models, and it causes complete suppression of the enzyme lipoprotein lipase (Torti et al., 1985) thereby preventing the uptake of exogenous triglyceride by fat cells and causing lipaemia (Guy, 1975; Rouzer & Cerami, 1980). We found that plasma from TDM-treated mice had cytotoxic activity for L929 tumour cells, a specific assay for TNF (Beutler & Cerami, 1986).

The predominant physiological role of TNF in TDM-treated animals is still unknown. TNF is released by mononuclear cells (Beutler & Cerami, 1986), and it appears to have many advantages for the host. For instance, it has parasiticidal (Taverne et al., 1981) and tumoricidal activity (Carswell et al., 1975) and it is able to activate neutrophils and increase phagocytosis (Gamble et al., 1986) by stimulating leukocytes to destroy micro-organisms in vitro (Silberstein & David, 1986), and to mobilize substances rich in energy. Thus, the nonspecific immunopotentiating activity of TDM (Stewart-Tull, 1985) may be related, at least in part, to the release of TNF. However, when TNF is liberated in large amounts, the resulting metabolic derangement will result in cachexia, anorexia, fever and diarrhoea, among other phenomena (Beutler & Cerami, 1986), frequently resulting in death.

We wish to thank Rosângela C. P. Mesquita for help and secretarial assistance. This investigation was supported by grants from Fundação de Amparo à Pesquisa do Estado de São Paulo (FAPESP – process no. 87/1853-0) and Conselho Nacional de Pesquisa (CNPq – process nos 401007/87.1 and 30.0351/81).

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**Induction of cachexia by trehalose dimycolate**


