BACTERIAL PATHOGENESIS

Regulation by protein kinase of phagocytosis of Mycobacterium leprae by macrophages

K. PRABHAKARAN, E. B. HARRIS and B. RANDHAWA

US Public Health Service, GWL Hansen’s Disease Center @ Louisiana State University, PO Box 25072, Baton Rouge, LA 70894-5072, USA

Mycobacterium leprae multiplies within host macrophages. The mechanism of internalisation of the bacteria by the phagocytic cells is unknown. In this study, M. leprae was purified from the foot pads of experimentally infected nu/nu mice. Peritoneal macrophages were harvested from BALB/c mice or C57 beige (bg/bg) mice. The effect of protein kinase inhibitors (erbstatin, genistein or staurosporine for BALB/c and bg/bg mice, plus herbimycin for bg/bg mice) on phagocytosis of the mycobacteria by the macrophage monolayers was tested. The untreated (control) macrophages phagocytosed M. leprae. Phagocytosis by BALB/c macrophages was inhibited by erbstatin and staurosporine but not by genistein; all the protein kinase inhibitors prevented uptake of M. leprae by bg/bg cells. The results demonstrate that protein kinase regulates phagocytosis of M. leprae by macrophages. The mechanism might prove to be a rational drug target for mycobacteria that multiply intracellularly.

Introduction

Protein phosphorylase (EC 2.7.1.27) regulates a multitude of cell functions such as cell proliferation and differentiation, signal transduction, transcription and synaptic transmission. Phosphorylation involves the introduction of a phosphoryl group into a substrate through the formation of an ester bond between the compound and phosphoric acid. In the protein kinase superfamily there are two subdivisions: protein tyrosine kinases and protein serine/threonine kinases. The terminal phosphoryl group of ATP is transferred to specific tyrosine residues by one class of protein kinases and to serine/threonine residues by the other.

Protein tyrosine phosphorylation was first discovered in 1979 [1]. At one time it was believed that only eukaryotic cells possess protein kinases; now the activity has been detected in eubacteria and archaea-bacteria as well [2, 3]. Protein kinase has been recognised as an evolutionarily conserved enzyme activity. Uncontrolled activation of tyrosine kinase is associated with many kinds of cancers and decrease in insulin receptor tyrosine kinase is associated with diabetes [1, 4].

Pathogenic bacteria have adopted various strategies to invade their hosts. Many intracellular micro-organisms interact with host cell receptor molecules to induce their own internalisation; bacteria exploit eukaryotic protein kinases to enter mammalian cells. Protein kinase inhibitors prevent cell infection by blocking stimulation of receptor sites on the phagocytic cells of the host. Protein kinase is activated during phagocytosis [5]. Mycobacterium leprae phagocytosed by macrophages suppresses phagosome–lysosome fusion and the bacteria multiply within the cells. The present study examined the effect of kinase inhibitors on uptake of M. leprae by macrophages and the possible regulatory role of protein phosphorylase in phagocytosis.

Materials and methods

Inhibitors

Staurosporine, erbstatin, genistein and herbimycin were purchased from Calbiochem, La Jolla, CA, USA.

Bacteria

M. leprae was purified from the foot pads of experimentally infected nu/nu BALB/c mice by homogenisation of the tissues and differential centrifugation of the homogenate. The bacterial pellet was washed thoroughly to remove host tissue elements.
Macrophages

Peritoneal macrophages were harvested with heparinised Hank’s balanced salts solution from BALB/c (Simonsen Lab, Gilroy, CA, USA) or C57 beige (bg/bg) mice (Jackson Laboratory, Bar Harbour, ME, USA). The cells were sedimented by centrifugation at 200 g for 10 min at 4°C. The macrophages were suspended in a small volume of RPMI 1640 medium containing heated fetal bovine serum 15%, 20 mM HEPES (N-2-hydroxyethylpiperazine-N-2-ethane sulfonic acid), NaHCO3 2 mg/ml, 2 mM glutamine and gentamycin 50 μg/ml. A cell count was made with acetic acid-methylene blue stain, and the number of cells in the medium was adjusted to 4 × 10⁶/ml (BALB/c) or 2 × 10⁶/ml (bg/bg).

Phagocytosis assay

BALB/c macrophages. A round, 13-mm sterile cover-slip (Thermanox Plastic, Nalge Nunc International, Naperville, IL, USA) was introduced into each well of a sterile, 24-well tissue culture plate; 0.5 ml of the cell suspension was pipetted into each well. After 24 h, non-adherent cells were washed off twice with phosphate-buffered saline (PBS). The protein kinase inhibitors were dissolved in dimethyl sulphoxide (DMSO) 0.25% at concentrations as follows: 516 μM erbsatin, 306 μM genistein, 25 μM staurosorpine. Each inhibitor solution (0.5 ml) was added to three wells; three wells received the culture medium containing DMSO 0.25%. After 60 min, the coverslips were washed twice with PBS and 0.5 ml of the medium was added to the wells. In another group, three of the wells were replenished with medium containing each inhibitor. M. leprae (2 × 10⁷/20 μl) was added to all the wells. After incubation at 37°C for 2 h, the coverslips were washed with PBS and stained by the Ziehl-Neelsen method for acid-fast bacteria. The coverslips were mounted on slides and photographed with a Zeiss Photomicroscope. The experiments were repeated three times, with similar results.

bg/bg macrophages. The procedures followed were the same as for the BALB/c macrophages, except for the following changes. The number of macrophages in the medium was 2 × 10⁶/ml; concentrations of inhibitors used were: 250 μM and 500 μM erbsatin, 100 nM and 200 nM herbimycin, 100 μM and 500 μM genistein, 500 nM and 5 μM staurosorpine. The IC₅₀ values of the inhibitors vary widely and their effects are not always concentration-dependent [6]. There was no pre-incubation of the macrophages with the inhibitors, which were added to the reaction mixture. M. leprae (1.8 × 10⁷/0.5 ml) was added to the wells and incubated for 4 h at 37°C. The coverslips were washed twice with PBS and fixed in formaldehyde 10% in ethanol. After staining by the Ziehl-Neelsen method, the number of acid-fast bacilli in 100 macrophages on each slide was counted. The results are expressed as the mean (SD) of three counts, from representative experiments.

Results

BALB/c macrophages

The control macrophages showed good internalisation of M. leprae (Fig. 1). Phagocytosis of the bacteria was suppressed in the macrophages exposed to staurosorpine and erbstatin (Figs. 2 and 3). Genistein showed no effect, the bacteria being internalised as in the controls (Fig. 4). The macrophages exposed to staurosorpine or erbstatin remained attached to the coverslips, indicating that they were still viable. Even after the inhibitors had been removed after pre-incubation with the cells, those cells treated with erbstatin or staurosorpine did not phagocytose the bacteria. The same results were obtained whether the cells remained exposed to the inhibitory compounds or the inhibitors were washed off after pre-incubation.

bg/bg macrophages

Beige (bg/bg) mice are mutants derived from the C57 parent strain and are considered to be deficient in natural killer (NK) cells. In the control macrophages, not exposed to the protein kinase inhibitors, 83±3SD%)% of the cells phagocytosed M. leprae (Table 1). Phagocytosis of the bacteria was suppressed by all the inhibitors at both the concentrations tested. Genistein, which had no effect on macrophages of BALB/c mice, produced the greatest inhibition in the bg/bg macrophages. The extreme susceptibility of these phagocytic cells from the genetically deficient bg/bg mice to protein kinase inhibitors requires further study.

Discussion

Protein phosphorylase regulates a variety of cellular functions. The kinases phosphorylate specific amino acids in proteins; the enzyme phosphatase that dephosphorylates the protein returns the system to its original state. Phosphorylase and phosphatase are involved in

---

Fig. 1. Phagocytosis of M. leprae by BALB/c macrophages; magnification ×1500.

Fig. 2. Staurosorpine inhibition of phagocytosis of M. leprae; magnification ×1500.

Fig. 3. Erbsatin inhibition of phagocytosis of M. leprae; magnification ×1500.

Fig. 4. Phagocytosis of M. leprae by macrophages in the presence of genistein; magnification ×1500.
the most crucial reactions carried out by organisms. Protein kinases and phosphatases have been found not only in eukaryotes but also in a wide range of bacteria [3]. In *Listeria monocytogenes* which invades non-phagocytic cells, cell penetration by the bacteria is associated with host cell tyrosine phosphorylation [7]. Protein kinase has been shown to mediate phagocytosis of *M. bovis* BCG [8].

The mechanism of action may not be the same for the different protein kinase inhibitors. Many tyrosine kinase inhibitors are not very selective, acting on several tyrosine kinases. Staurosporine is a potent broad-range inhibitor of protein kinases [9]. *In vitro*, genistein is a competitive inhibitor of ATP in the kinase reaction, but it has no efficacy *in vivo* [10] and did not prevent uptake of *M. leprae* by BALB/c macrophages in the present study. Erbstatin was reported to be competitive with both protein and ATP [11]. Herbimycin inhibits tyrosine kinase irreversibly [12].

Tyrosine phosphorylation was shown to be one of the signalling events mediating the uptake of *M. tuberculosis* by macrophages [13, 14]. The results of the present study demonstrate that protein kinase is a factor that regulates phagocytosis of *M. leprae* by macrophages. Further investigation of the various types of protein kinases has not been pursued in this study. As mycobacteria such as *M. tuberculosis* and *M. leprae* multiply within the phagocytic cells of the host, blocking protein phosphorylase could be a rational drug target against infections caused by the bacilli [11].

We thank Mr J. P. Pasqua for harvesting the macrophages and Mr Gregory McCormick for preparing the figures.

**References**


**Table 1. Effects of protein kinase inhibitors on the phagocytosis of *M. leprae* by macrophages of beige (bg/bg) mice**

<table>
<thead>
<tr>
<th>Inhibitors</th>
<th>Concentration of inhibitors</th>
<th>Mean (SD) percent of cells taking up <em>M. leprae</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>–</td>
<td>83 (3)</td>
</tr>
<tr>
<td>Erbstatin</td>
<td>250 μM</td>
<td>8 (2)</td>
</tr>
<tr>
<td></td>
<td>500 μM</td>
<td>5 (2)</td>
</tr>
<tr>
<td>Herbimycin</td>
<td>100 nM</td>
<td>7 (3)</td>
</tr>
<tr>
<td></td>
<td>200 nM</td>
<td>13 (2)</td>
</tr>
<tr>
<td>Genistein</td>
<td>100 μM</td>
<td>5 (1)</td>
</tr>
<tr>
<td></td>
<td>500 μM</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Staurosporine</td>
<td>500 nM</td>
<td>20 (10)</td>
</tr>
<tr>
<td></td>
<td>5 μM</td>
<td>10 (4)</td>
</tr>
</tbody>
</table>