INTRODUCTION

Several studies have shown that micro-organisms have developed the ability to interact with host-cell receptor molecules to induce their own internalization (Moulder, 1985; Falkow, 1991; Isberg, 1991; Pace et al., 1993; Káposzta et al., 1999). Entry of the pathogen into the host cell is initiated by binding to its surface, which is followed by internalization. These events allow a pathogen to sequester itself within a particular organ site or to gain access to deeper tissues. Both host and pathogen encode factors that determine which type of host cells will be infected. Factors that determine whether the micro-organism will remain surface-bound or be internalized include: (i) the nature of the host-cell receptors, (ii) the strength of the physical interaction that occurs between the host and microbial ligands and (iii) whether the micro-organisms trigger signals that either stimulate or antagonize internalization (Isberg & Tran Van Nhung, 1994). Host-cell actin filaments facilitate the internalization step, since agents that disrupt actin filaments, such as cytochalasin D, inhibit internalization of most pathogens (Clerc & Sansonetti, 1987; Finlay & Falkow, 1988; Finlay et al., 1989; Farbiarz et al., 1992; Rosenshine et al., 1992). Given the active involvement of the host cytoskeleton, it appears that invasive pathogens generate an ‘uptake signal’ that induces the host cell to internalize them (Rosenshine et al., 1992). It is clear that binding of the parasite to the host-cell surface is not necessarily followed by internalization, as many parasites bind to the cell surface but do not enter the cell. These observations suggest that, following parasite attachment, this ‘uptake signal’ must be generated in order for internalization to take place (Vieira et al., 1994).

Cell-mediated immunity (T cells) and non-specific cellular immunity (e.g. macrophages, natural killer cells and neutrophils) are generally believed to provide the main defence mechanisms against fungi. The importance of cellular defence mechanisms for protection against fungi is supported by the clinical observation that most invasive fungal infections occur in individuals with defective cellular immunity (Levitz, 1992; Casadevall, 1995).

The human pathogen Fonsecaea pedrosoi is one of the aetiological agents of chromoblastomycosis, a subcutaneous disease that shows no major tendency to disseminate to deeper organs (McGinnis, 1983; Rippon, 1988). All agents of subcutaneous mycosis induce the influx of cells belonging to the mononuclear phagocytic system, producing a granulomatous reaction. In tissues, conidia and sclerotic cells of F. pedrosoi have been found. Immunofluorescence micrographs of the interaction between F. pedrosoi conidia and IC21 macrophages are available as supplementary material in JMM Online (http://jmm.sgmjournals.org).
pedrosoi can be detected within macrophages and neutrophils as well as extracellularly. Previous studies have demonstrated that resident macrophages have no fungidal or fungistatic activity against F. pedrosoi and that, 24 h after the beginning of the interaction, the macrophages are completely destroyed by the fungi (Farbízár et al., 1990; Rozental et al., 1994). Different cell types have been used to study the F. pedrosoi–host cell interaction, showing that this fungus can invade epithelial cells like Chinese hamster ovary (CHO) cells (Limongi et al., 1997), Madine–Darby canine kidney cells and also cytochalasin-treated macrophages (Farbízár et al., 1992). These results suggest that F. pedrosoi might penetrate host non-phagocytic cells through an active process. We demonstrated recently that the interaction of F. pedrosoi conidia with mannose and N-acetylglycosamine surface carbohydrates in CHO epithelial cells is mediated by a 50 kDa lectin-like protein found in the conidium cell wall (Limongi et al., 2001).

In eukaryotes, phosphorylation/dephosphorylation cycles represent a major mechanism for switching cellular pathways in response to changing circumstances, both internal developmental cues and external environmental stimuli (Dickman & Yarden, 1999). Different classes of protein kinases (PKs) play a major role in transducing extracellular signals and regulating essential cellular processes in mammalian cells, such as cell growth, differentiation, transcription, disease and death (Courtneidge, 1994; Jarvis et al., 1994; Van der Geer et al., 1994; Hunter, 1995; Dickman & Yarden, 1999). The phosphorylation process has recently been implicated in defence strategies against a variety of pathogens that have evolved mechanisms to alter host-signalling pathways in order to facilitate their invasion and survival within host cells. We have investigated the involvement of PKs in the attachment to and invasion of CHO epithelial cells and IC21 macrophages by F. pedrosoi. We examined the involvement of these enzymes present in host cells and in the fungal cell in transducing signals needed for conidium internalization by epithelial cells and macrophages. The use of the PK inhibitors staurosporine, genistein and calphostin C prior to infection gave us significant information about the role played by PKs in the F. pedrosoi–host cell interaction. In this paper, we present evidence that PK inhibitors can significantly reduce invasion by F. pedrosoi.

METHODS

Micro-organisms. A pathogenic strain of F. pedrosoi was isolated from a human case of chromoblastomycosis (Oliveira et al., 1990; Rozental et al., 1994). Different cell types have been used to study the F. pedrosoi–host cell interaction, showing that this fungus can invade epithelial cells like Chinese hamster ovary (CHO) cells (Limongi et al., 1997), Madine–Darby canine kidney cells and also cytochalasin-treated macrophages (Farbízár et al., 1992). These results suggest that F. pedrosoi might penetrate host non-phagocytic cells through an active process. We demonstrated recently that the interaction of F. pedrosoi conidia with mannose and N-acetylglycosamine surface carbohydrates in CHO epithelial cells is mediated by a 50 kDa lectin-like protein found in the conidium cell wall (Limongi et al., 2001).

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METHODS

Micro-organisms. A pathogenic strain of F. pedrosoi was isolated from a human case of chromoblastomycosis (Oliveira et al., 1993). Stock cultures were maintained on Sabouraud's glucose agar under mineral bovine serum (FBS; Gibco), at 37 °C with 5 % CO2, until they reached confluence. They were subcultured at least once a week. IC21 macrophage cells were also purchased from the ATCC (TIB-186). The cells were grown in RPMI medium supplemented with sodium pyruvate (0·11 g l−1), glucose (3·7 g l−1), L-glutamine (2·0 mmol l−1), sodium bicarbonate (1·1 g l−1) and 10 % FBS, at 37 °C with 5 % CO2. They were subcultured at least once a week. Prior to the interaction with the fungus, CHO cells and IC21 macrophages were plated onto 24-well multidishes in the presence of FBS and incubated at 37 °C for 24 h. After this time, the cells were washed in MEAM or RPMI and counted in an inverted phase-contrast microscope.

PK inhibitors. Genistein (4',5,7-trihydroxyisoflavone), a protein-tyrosine kinase (PTK) inhibitor; staurosporine (antibiotic AM-2282 from Streptomyces sp.), a broad-range kinase inhibitor, calphostin C (UCN-1082c, from Cladostrum cladostrum), a Ca2+/phospholipid-dependent PK (PKC) inhibitor, and okadaic acid, a phosphatase inhibitor (Tamaoki, 1991), were purchased from Sigma. Stock solutions were made in 100 % DMСO and stored at −20 °C until use. DMСO did not affect cultures at the final concentration used (0·01 %). Prior to use, the inhibitors were diluted in MEAM or RPMI and the mixture was added to the mammalian cells or to the conidia. The final concentrations used in the assays were staurosporine, 0·5, 1·0 and 2·0 μM; genistein, 50, 100 and 200 μM; calphostin C, 25, 50 and 100 mM; and okadaic acid, 5, 10 and 20 nM.

Host cell– F. pedrosoi interaction. CHO epithelial cells and IC21 macrophages were incubated for 30 min at 37 °C in the presence of 0·01 % DMСO, 0·5, 1·0 or 2·0 μM staurosporine, 50, 100 or 200 μM genistein or 25, 50 or 100 nM calphostin C, washed three times and allowed to interact with F. pedrosoi conidia for 1 h at 37 °C, 5 % CO2. The conidium:host cell ratio was adjusted to 10:1 by counting the number of conidia with a Neubauer chamber (Sigma). Alternatively, after PK inhibitor treatment, the cells were re-incubated in the presence of 5, 10 or 20 nM okadaic acid, for 30 min, before the interaction. In other assays, the conidia were incubated with the same PK inhibitors at the same concentrations, washed and allowed to interact with non-treated CHO cells and IC21 macrophages for 1 h, at 37 °C, 5 % CO2 and, alternatively, were also re-incubated with okadaic acid prior to the interaction. After the interaction, the cultures were rinsed three times in MEAM or RPMI to remove free conidia, fixed in Bouin’s solution and stained with Giemsa (Merck). Bouin’s solution was made in-house and contains 375 ml saturated aqueous picric acid (Merck), 125 ml stock formaldehyde (37 %, w/v) (Reagen) and 25 ml glacial acetic acid (Reagen). The percentage of infected cells was determined by randomly counting a minimum of 300 cells on each duplicated cover slip and experiments were repeated at least three times. The adhesion index (AI) was calculated by multiplying the mean number of attached conidia per CHO cell by the percentage of infected cells observed by microscopic examination using the immersion objective of a Zeiss Universal photomicroscope. The endocytic index (EI) was determined in the same way, but was based on the mean number of ingested conidia.

Transmission electron microscopy. After the interaction, performed in 25 cm2 flasks, the cultures were fixed for 1 h at room temperature with a solution containing 1 % glutaraldehyde, 4 % paraformaldehyde and 1 mM CaCl2 in 0·1 M cacodylate buffer, pH 7·2. The cells were then gently scrapped off with a rubber policeman, washed with the same buffer and post-fixed for 1 h in a solution containing 1 % OsO4, 0·8 % potassium ferricyanide and 1 mM CaCl2 in a cacodylate buffer, pH 7·2. Subsequently, the cells were rinsed, dehydrated in acetone and embedded in Epon. Ultrathin sections were stained with uranyl acetate and lead citrate and observed with a Zeiss 900 electron microscope.

Immunofluorescence microscopy. F. pedrosoi conidia and IC21 macrophages were allowed to interact for 30 min as described above,
washed three times with 1 mM sodium vanadate in PHEM buffer (60 mM PIPES-NaOH, 20 mM Hepes-NaOH, 10 mM EGTA, 5 mM MgCl₂, 70 mM KCl, pH 7·2) and fixed with freshly prepared 4 % formaldehyde in 0·1 M phosphate buffer. Cells were permeabilized with 0·1 % Triton X-100 in PHEM buffer for 90 s and then treated with a blocking solution containing 50 mM ammonium chloride and 1·5 % BSA for 120 min at room temperature. The samples were then incubated for 60 min in the presence of phallolidin–rhodamine (diluted 1 : 100), washed and then incubated in the presence of mAbs recognizing phosphotyrosine, phosphoserine or phosphothreonine residues (diluted 1 : 20 in PBS), washed three times in PBS/3 % BSA and incubated for 60 min in the presence of FITC-labelled goat anti-mouse IgG (diluted 1 : 200), washed and then incubated in the presence of mAbs recognizing bound vacuoles in mammalian cells (Limongi et al., 1997, 2001). Previous observations of our group have shown that prolonged incubation of infected mammalian cells allows both ingested fungi, as well as fungi that remain attached to the macrophage surface, to differentiate into hyphae, since resident macrophages have little or no cytotoxic effect in vitro on F. pedrosoi (Farbian et al., 1990). On the other hand, when we tested times shorter than 1 h, fungi were not observed inside epithelial cells, probably because they are non-professional phagocytic cells (data not shown).

RESULTS

Effect on cell adhesion of previous treatment of host cells with PK inhibitors

Staurosporine treatment decreased binding of conidia to IC21 macrophages (Fig. 1a). At 2·0 μM staurosporine, treatment of IC21 macrophages prior to infection reduced the AI by 77 %. However, no significant effect on the AI was observed for Pro-5 (Fig. 1b). The cells were washed extensively before interactions to avoid possible direct effects of PK inhibitors upon conidia. Genistein also inhibited conidium attachment to IC21 macrophages, with a reduction of the AI by 68 % when used at a concentration of 200 μM. Genistein did not interfere with the attachment of the fungus to Pro-5 (Fig. 1b), as seen for staurosporine. Previous treatment of the cells with calphostin C reduced the AI to IC21 macrophages by 41 % (Fig. 1a) and to Pro-5 by 47 % (Fig. 1b). We decided to use a period of 1 h for binding; we have previously used 2 h and found that the values were very similar (Limongi et al., 1997, 2001). Previous observations of our group have shown that prolonged incubation of infected mammalian cells allows both ingested fungi, as well as fungi that remain attached to the macrophage surface, to differentiate into hyphae, since resident macrophages have little or no cytotoxic effect in vitro on F. pedrosoi (Farbian et al., 1990). On the other hand, when we tested times shorter than 1 h, fungi were not observed inside epithelial cells, probably because they are non-professional phagocytic cells (data not shown).

Effect on fungus internalization of previous treatment of host cells with PK inhibitors

In a previous work, we demonstrated that, by light microscopy of Giemsa-stained preparations of interaction assays, we can distinguish between attached or ingested fungi, since the ingested fungi are located within well-stained membrane-bound vacuoles in mammalian cells (Limongi et al., 1997). Our results show that, when used at a concentration of 2 μM, staurosporine decreased conidia internalization by IC21 macrophages (Fig. 2a) and CHO cells (Fig. 2b) by 78 and 73 %, respectively. Genistein also inhibited conidia phagocytosis by IC21 macrophages, by 77 % (Fig. 2a). In addition, previous incubation of the epithelial cells with genistein led to a remarkable decrease of the EI, by 52 % (Fig. 2b). Previous

![Fig. 1](http://www.microbiologyresearch.org:80/jmm.sgmjournals.org/article/Fig1.jpg)  
**Fig. 1.** Effect of pre-treatment of host cells with PK inhibitors for 30 min at 37 °C on the AI for the interaction of F. pedrosoi conidia for 1 h at 37 °C with IC21 cells (a) and parental Pro-5 CHO cells (b). PK concentrations are indicated in μM (genistein and staurosporine) or mM (calphostin C).

![Fig. 2](http://www.microbiologyresearch.org:80/jmm.sgmjournals.org/article/Fig2.jpg)  
**Fig. 2.** Effect of pre-treatment of host cells with PK inhibitors for 30 min at 37 °C on the EI for the interaction of F. pedrosoi conidia for 1 h at 37 °C with IC21 cells (a) and parental Pro-5 CHO cells (b).
treatment of macrophages with calphostin C decreased the EI by 75% (Fig. 2a) and also inhibited internalization of the fungi by epithelial cells (Fig. 2b). These drug effects were not dose-dependent. However, in most of the cases, the higher dose was the one that led to the greater inhibitory effect.

Effect on cell adhesion of previous treatment of the fungus with PK inhibitors

Previous incubation of F. pedrosoi conidia with staurosporine, genistein or calphostin C did not interfere significantly with their adhesion to IC21 macrophages (Fig. 3a). On the other hand, previous incubation with staurosporine decreased the AI to epithelial cells by almost 37%. At 200 μM, genistein treatment caused a decrease of 50% in adhesion of the fungus to these cells (Fig. 3b). Conidium viability after PK inhibitor treatment was greater than 90%, as determined by staining with 0·04 % Trypan blue (J. T. Baker Chemical Co.). As a control, conidia were not incubated with the PK inhibitors prior to the interaction. Conidium incubation with DMSO only prior to the interaction was used as a second control.

Effect on fungus internalization of previous treatment of the fungus with PK inhibitors

Previous incubation of the fungus with staurosporine, genistein or calphostin C did not interfere with its entry into IC21 macrophages (Fig. 4a). On the other hand, all the PK antagonists used markedly reduced the EI for CHO cells. Staurosporine and genistein treatment inhibited the EI by about 50% (Fig. 4b). Once again, 100 nM calphostin C caused the greatest reduction of the EI on epithelial cells, reaching 70% (Fig. 4b).

Antagonism by okadaic acid of the effect of PK inhibitors on the interaction of mammalian cells with conidia

Staurosporine, genistein and calphostin C are widely used as inhibitors of PKs (Tamaoki, 1991). However, as with any inhibitor, it is difficult to prove that the inhibition of attachment and invasion by these drugs results from an effect on protein phosphorylation rather than on some other, uncharacterized target (Ward et al., 1994). To address this issue, we attempted to overcome the effects of PK inhibitors with okadaic acid, a specific inhibitor of protein phosphatases 1 and 2A (Cohen et al., 1990; Hardie et al., 1991). We found that the addition of okadaic acid after treatment of cells with PK inhibitors could indeed partially alleviate the inhibitory effect of the drugs on the cells (Fig. 5). In most experiments, okadaic acid did not stimulate invasion by itself. The extent of recovery observed varied according to cell type. In macrophages, addition of okadaic acid completely reversed the effect of PK inhibitors on the EI (Fig. 5a). In epithelial cells, almost total reversal was also observed for all the drugs used (Fig. 5b). This reversion effect occurred in a

Fig. 3. Effect of pre-treatment of conidia with PK inhibitors for 30 min at 37 °C on the AI after an interaction of 1 h at 37 °C with IC21 cells (a) or parental Pro−/− CHO cells (b).

Fig. 4. Effect of pre-treatment of conidia with PK inhibitors for 30 min at 37 °C on the EI after an interaction of 1 h at 37 °C with IC21 cells (a) or parental Pro−/− CHO cells (b).
dose-dependent manner, as shown for staurosporine (Fig. 6). The same reversible behaviour was observed for both genistein and calphostin C (data not shown).

Antagonism of the effect of PK inhibitors on the fungus by okadaic acid

Incubation of conidia with PK inhibitors prior to the interaction in the presence of okadaic acid did not interfere with the interaction with macrophages. In contrast, when epithelial cells were used, the effects of PK inhibitors were partially reversed by okadaic acid; however, this effect was not evident for genistein (data not shown).

Immunofluorescence microscopy

We attempted to localize phosphotyrosine, phosphoserine, phosphothreonine residues and the actin cytoskeleton during the conidium–IC21 macrophage interaction using immunofluorescence techniques. Labelling with anti-phosphotyrosine antibodies was very intense on the macrophage surface, especially in those that were in contact with the fungus. Immunofluorescence micrographs are available as supplementary material in JMM Online (http://jmm.sgmjournals.org). Actin filaments, revealed using phalloidin–rhodamine, were concentrated at the regions of binding of conidia to cells, co-localizing with phosphotyrosine domains. Anti-phosphoserine antibody labelling was restricted to endocytic vacuoles containing conidia. No significant labelling of the cells was observed using anti-phosphothreonine antibodies, except for some conidia located inside macrophages.

Transmission electron microscopy of the interaction of F. pedrosoi conidia with host cells

Transmission electron micrographs of the interaction of conidia with macrophages showed emission of pseudopods as an initial step of conidium internalization (data not shown) and conidia inside a ‘tight’ phagosome, where the membrane of the vacuole was apposed directly over the fungal cell wall (Fig. 7a). When cells pre-treated with PK inhibitors were allowed to interact with the fungus, ingested conidia were observed inside a ‘loose’-type phagosome (Fig. 7b–d), with a wide space between the vacuole membrane and the fungal cell wall, similar to those observed in epithelial cells (Fig. 7e, f). In cells treated with staurosporine, the host membrane seemed to lose its natural shape and showed projections all over the surface (Fig. 7b). Pre-treatment with genistein (Fig. 7c) and calphostin C (Fig. 7d) also affected membrane continuity, showing disruptions and the presence of a significant number of vacuoles located near the phagosomes. The interaction of F. pedrosoi with epithelial cells showed that the ingested conidia are located inside a ‘loose’-type phagosome (Fig. 7e, f). When epithelial cells were pre-treated with calphostin, the host membrane seemed to lose its

Fig. 5. Influence of incubation with okadaic acid for 30 min at 37 °C after pre-treatment of IC21 cells (a) or parental Pro–/ CHO cells (b) with PK inhibitors for 30 min at 37 °C on the interaction with conidia of F. pedrosoi for 1 h at 37 °C. +, Okadaic acid added at 20 nM; DMSO, DMSO only; Gen, 200 μM genistein; Stau, 2·0 μM staurosporine; Calph, 100 nM calphostin C.

Fig. 6. Alleviation by okadaic acid at concentrations of 5, 10 or 20 nM of the effect of 2·0 μM staurosporine (Stau) on the interaction of F. pedrosoi with IC21 macrophages (a) and parental Pro–/ CHO cells (b).
natural shape and host cells did not show the emission of traditional pseudopods at the region of the initial contact with the fungus (Fig. 7f). CHO cells treated with genistein and staurosporine prior to the interaction showed effects similar to those observed with calphostin (data not shown).

DISCUSSION

Protein phosphorylation is generally accepted to play key roles in transducing signals involved in several processes such as cell adhesion, internalization and killing of pathogens (Reiner, 1994). Pharmacological agents that arrest invasion at a specific step are useful experimental tools that can provide clues about what is going on biochemically within the cell and the parasite as it attaches and enters the cell (Ward et al., 1994). Inhibitors of enzyme activity are valuable tools for probing kinase/phosphatase involvement in cellular processes; however, as the use of such compounds has inherent limitations due to non-specific reactions, it is important to bear in mind the properties of these compounds. Especially crucial is whether the inhibitor binds directly to the PK and with what affinity, how specific it is for a given kinase and how permeable the fungal cell is to a given inhibitor (Finlay et al., 1989). In this study, we used the PK inhibitors staurosporine, genistein and calphostin C to analyse the roles played by the PK activities of F. pedrosoi conidia and host cells in the interaction process. The inhibitors employed here presumably down-regulated cell PK activity, as treatments were performed prior to infection. We have shown here that pre-treatment of macrophages, epithelial cells and conidia with PK inhibitors decreased conidium invasion and that this could be overcome with okadaic acid in most cases. Our results suggest that not only is the mammalian cell PK activity important in the interaction process, but also that protein phosphorylation that occurs within the conidia may play a significant role in invasion.

Staurosporine is a potent inhibitor of several different Ser/Thr kinase classes, including PKC and cAMP-dependent PK and some PTKs (Tamaoki, 1991). Staurosporine inhibited the conidium adhesion and invasion processes in macrophages, suggesting that PKs are necessary for both steps. However, in the case of epithelial cells, only the invasion process was arrested. Similar results were obtained with genistein and calphostin C. It is important to point out another important difference between the response of
epithelial cells and macrophages to the drugs used here; when conidia were pre-treated with the drugs, no effects were observed on the interaction with macrophages. On the contrary, invasion of epithelial cells by the fungus was strongly inhibited. These results suggest that, in the case of phagocytic cells, the PK machinery of the fungus is not necessary for its entry, but, for non-phagocytic cells, the fungus has to use its own PKs to facilitate invasion of the cell, suggesting a truly active process of penetration.

It is intriguing to note that staurosporine blocks invasion in epithelial cells at a step that is morphologically similar to the arrest seen when the cells are treated with cytochalasins (Farbizaz et al., 1992). In both cases, the early stages of invasion (recognition and adhesion) appear to proceed normally, but the attached fungi are not internalized. Cytochalasin D binds to the barbed end of actin filaments and is a specific inhibitor of actin-based processes in intact cells (Ohmori et al., 1992). Entry of a particle into a cell by phagocytosis requires reorganization of the actin-based cytoskeleton underlying the region of the plasma membrane that contacts the particle. F-actin assembly in this region is initiated by signals that arise from the interaction of phagocytosis-promoting receptors on the cell surface with ligands on the surface of the particle (Greenberg, 1995). Farbizaz et al. (1992) have suggested the existence of two mechanisms used by \textit{F. pedrosoi} to penetrate vertebrate cells. One is a typical phagocytic process, with formation of pseudopods. The second occurs even in the absence of actin filaments and is probably the same mechanism used to infect epithelial cells, in a process in which the fungus may play an active role (Farbizaz et al., 1992).

The potential involvement of PTKs in regulating the interaction was verified by the use of genistein. In this study, we showed that this PTK inhibitor could also block fungus internalization after treatment of mammalian cells. Genistein inhibits the binding of ATP to the enzyme (Akiyama et al., 1987; Yaish et al., 1988) and is a highly specific PTK inhibitor; it exhibits a weak inhibitory effect against protein-Ser/Thr kinases and scarcely inhibits CAMP-dependent PK activity (Akiyama & Ogawara, 1991). Genistein treatment of the fungus did not interfere with the interaction with macrophages, but inhibited interaction with epithelial cells, suggesting that fungal PTK participates in entry into epithelial cells. This observation suggests the existence of tyrosine kinases in \textit{F. pedrosoi}, in contrast with the idea that no tyrosine kinases exist in filamentous fungi and that Ser/Thr kinases represent virtually all of the PKs found in these fungi (Dickman & Varden, 1999).

PKC is a protein-Ser/Thr kinase involved in the regulation of many cellular processes, including cellular growth, differentiation and tumour promotion (Nishizuka, 1986). To examine whether PKC activity is needed for invasion by \textit{F. pedrosoi}, we tested the effect of an inhibitor that is known to have a higher specificity for PKC. Staurosporine inhibits the proteolytically generated catalytic domain of PKC, while calphostin C interacts with the regulatory domain, being a highly specific PKC inhibitor (Tamaoki, 1991). Previous incubation of the host cells with calphostin C blocked fungal invasion. This effect was especially evident in CHO cells. The same effect was observed when conidia were pre-treated and allowed to interact with the cells. Calphostin C seems to be the only kinase antagonist that caused some effect on the adhesion of conidia to epithelial cells, suggesting that cellular PKC may participate in the adhesion of the fungus to epithelial cells, while cellular PTK and PKC participate in the adhesion of \textit{F. pedrosoi} to macrophages. Since the effects of PK inhibitors are, in most cases, alleviated by okadaic acid in a dose-dependent manner, and since okadaic acid is a specific inhibitor of protein-Ser-Thr phosphatases 1 and 2A, with little effect on protein-tyrosine phosphatases (Cohen et al., 1990; Hardie et al., 1991), it seems likely that protein-Ser-Thr kinases within the host cells and the fungus are possible targets for staurosporine. Okadaic acid reversed the effect of staurosporine in all the cells tested, as well as the effect of genistein and calphostin C in macrophages.

Kápozta et al. (1999) have shown that PKC activity is essential for phagocytosis of \textit{Candida}. The inhibitory effect of staurosporine on internalization of \textit{Candida albicans} by macrophages reflects the role of PKC in actin polymerization and in recruitment of cytoskeletal proteins to the binding sites of particles (Allen & Aderem, 1995, 1996). In addition, a tyrosine kinase was involved in tumour necrosis factor \(\alpha\) production by macrophages after stimulation by \textit{C. albicans} (Jouault et al., 1998). Shahan et al. (2000) demonstrated that fungal spores from different species activate macrophages differentially, especially by PTK activation. Pathogen invasion has recently sparked great interest, not only because its study may lead to better ways of controlling important pathogens but also because it may help to understand fundamental aspects of cell biology (Pace et al., 1993).

Immunofluorescence assays using monoclonal anti-phosphotyrosine antibodies revealed an accumulation of tyrosine-phosphorylated residues at the site of fungus attachment on the macrophage surface, although no labelling was observed after parasite internalization. It may be inferred that tyrosine phosphorylation during this process is an early, local and short-lived event. Simultaneous labelling of the cells with phalloidin showed that actin filaments also participate in this process. This is consistent with the transient nature of most PTK-mediated responses (Ulrich & Schlessinger, 1990). Phosphoserine staining was observed in macrophage phagocytic vacuoles containing fungi. This observation suggests a later and longer phenomenon, where protein-Ser kinases participate in the phagocytosis process itself. The fact that phosphothreonine staining was weaker, in fact almost indiscernible, does not exclude participation of protein-Thr kinases in the interaction process. Protein-Thr kinase activation could be a transient event, difficult to detect by the approach used in the present study.

Our observations by transmission electron microscopy showed that conidia infect macrophages by a typical phago-
cytic process, with formation of pseudopods and a ‘tight’ phagosome. When the cells were pre-treated with PK inhibitors, no pseudopods were observed and the ingested fungi were found in a ‘loose’ type of phagosome vacuole, similar to that found in epithelial cells and in cytochalasin-treated macrophages, thus indicating a process of interaction in which the fungus plays an active role.

The study of kinases and phosphatases in pathogenic fungi has become an active area of research in biology. The present results demonstrate that signal-transduction networks involving PK and protein phosphatase activities can modulate crucial events during F. pedrosoi infection.

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208
Protein kinases in F. pedrosoi–host cell interaction


