The inhibitory effect of the antifungal agents and the HIV proteinase inhibitors on Sap activity was analysed using proteinase-containing culture supernatants from three C. albicans strains (hereafter named strains 1–3) isolated from the oral mucosa of three HIV-infected patients. Results were compared to the inhibition of pepstatin A. Isolates were identified as C. albicans by their colony morphologies, their ability to form germ tubes and their biochemical patterns, as assigned using the API system ATB 32 C (bioMérieux). The HIV proteinase inhibitor saquinavir was kindly provided by Roche; amprenavir was obtained from GlaxoSmithKline and pepstatin A was obtained from Sigma. Terbinafine (Novartis), ketoconazole (Janssen–Cilag), amphotericin B (Bristol Myers Squibb) and ciclopiroxolamine (Aventis) were obtained as reagent-grade powders from their respective manufacturers. Each C. albicans strain was grown in Sabouraud/glucose broth (Difco) in an incubator (Heraeus) for 48 h at 27 °C. The induction of C. albicans Saps was performed as described previously (Korting et al., 1999). Briefly, 100 μl of C. albicans suspension was added to 10 ml Remold medium [2 % glucose, 0·1 % KH₂PO₄, 0·5 % MgSO₄, 1·25 ml 100 × sterile filtered minimum essential medium vitamins (Sigma) and 1 % BSA]. The mixture was incubated for 7 days at 27 °C in a shaker at 150 r.p.m. Thereafter, titres (c.f.u.) were determined and the yeast cells were removed by centrifugation at 1500 g for 30 min. Supernatants were adjusted to pH 6·5 with NaOH to limit auto-degradation and frozen at −20 °C after filter sterilization (500 ml Stericup, pore size 0·22 μm, Millipore) to give the final crude enzyme preparation.

In the last decade, it has been demonstrated that secreted aspartyl proteinases (Saps) are important virulence factors for several types of Candida albicans infections and that inhibition of these proteinases have a protective effect for the host (De Bernardis et al., 2001; Hube & Naglik, 2001). Based on the observation that antifungal drugs may have broad modes of action, we questioned whether certain antifungal components may also influence the activity of Saps, which, in turn, may enhance the antifungal activity of a particular drug. For example, Wu et al. (1999) showed that the natural antimicrobial agent lysozyme not only showed a candidacidal effect at higher concentrations but also decreased the extracellular concentration of Saps significantly without affecting cell growth or viability of C. albicans. In addition, lysozyme also directly caused degradation of purified Sap protein (Wu et al., 1999). Other studies showed that certain inhibitors designed to inhibit human immunodeficiency virus (HIV) proteinase also had a direct effect on the activity of Saps (Korting et al., 1999; Cassone et al., 1999; Borg-von Zepelin et al., 1999). To investigate whether recently designed HIV proteinase inhibitors, such as amprenavir, or antifungal agents, such as members of the allylamines, azoles, polyenes or pyridone antimycotic components, inhibit Sap activity, we measured the effects of selected HIV inhibitors and antifungal drugs on the proteolytic activity of C. albicans using a spectrophotometric assay.
Stock solutions were prepared for amprenavir, saquinavir and pepstatin A by dissolving in absolute methanol at a concentration of 1.0 mM for saquinavir and 100 mM for pepstatin A and amprenavir. Amprenavir and saquinavir were diluted with 0.2 M sodium citrate/HCl buffer (pH 4.5) (Merck) to 1.0, 0.2 and 0.1 mM; pepstatin A was diluted with sodium citrate/HCl buffer to 0.5, 0.75 and 1.0 mM. Terbinafine was diluted in distilled water to 100 µM. Ketoconazole and amphotericin B were diluted in dimethyl formamide (Sigma) to 1.0 mM. Clicopiroxolamine was diluted in dimethyl formamide to 100 µM. Dilutions were 0.5, 1 and 2 µM for terbinafine and clicopiroxolamine, 0.2, 0.5 and 1 µM for amphotericin B and 0.5, 0.75 and 1 µM for ketoconazole.

Studies were carried out using bovine haemoglobin (Sigma) as substrate (Korting et al., 1999). Test tubes were each filled with 750 µl 0.2 M sodium citrate/HCl buffer, 750 µl fresh substrate solution (1% substrate in 0.2 M sodium citrate/HCl buffer), 250 µl each sample and 250 µl amprenavir, saquinavir, pepstatin A, terbinafine, ketoconazole, amphotericin B or clicopiroxolamine. Control experiments included assays without the addition of antifungal agents or inhibitors. Control experiments also included assays with dimethyl formamide or sodium citrate/HCl buffer alone without addition of antimycotics or protease inhibitors. Test reactions were incubated at 37 °C for 60 min (T60) in a shaker. The reaction was linear with time for up to 60 min. Three triplicate reactions were used for each experiment. Reactions were stopped with 500 µl trichloroacetic acid (TCA) and stored on ice. For each reaction mixture, an additional control was prepared by adding all ingredients plus 20% TCA simultaneously prior to incubation (T0). After the addition of TCA, all specimens were centrifuged at 3000 g for 30 min at 4 °C. A 160 µl sample of each cleared supernatant was then added to 40 µl dye reagent (Coomassie brilliant blue G-250, Bio-Rad). Peptides produced by proteolytic activity are not precipitated by TCA and bind to the dye. The amount of peptides in the supernatant can therefore be measured in a spectrophotometer (MR 4000, Dynatech) as a shift in the maximum absorbance value of the dye (measured at a wavelength of 595 nm) and correlated with proteolytic activity. Activity was calculated as the change in absorbance value using the following formula: sample (T60) – control (T0). One unit of activity was defined as an increase in 0.100 per 60 min at 595 nm. Activities were calculated for 1 l of Remold medium at a yeast density of 10^8 cells ml^-1. The least significance difference (LSD) test was used to determine differences between means. P values of < 0.05 were considered to be statistically significant.

Sap activity of the three C. albicans strains was inhibited by all protease inhibitors. The results of the present study confirmed the data published previously for pepstatin A and saquinavir (Korting et al., 1999). Pepstatin A, at concentrations ranging from 0.5 to 1.0 µM, inhibited Sap by approximately 18–78% (P < 0.001). Saquinavir, at concentrations ranging from 0.1 to 1.0 µM, inhibited Sap by approximately 15–79% (P < 0.001). Amprenavir was tested at the same concentrations and inhibited Sap by approximately 28–84% (P < 0.001). Statistical analysis of Sap activity with and without a protease inhibitor showed a high degree of significance for each of the three agents (Fig. 1). Sap activity of the three C. albicans strains was not significantly inhibited by ketoconazole, terbinafine and amphotericin. In contrast, in the presence of clicopiroxolamine, a significant inhibition (P < 0.001) was seen and ranged from 40 to 86% (Fig. 1).

The present study confirms the data published previously of Sap inhibition by pepstatin A and saquinavir (Korting et al., 1999; Borg-von Zepelin et al., 1999). In addition, a marked and highly significant inhibition of Sap activity by amprenavir could be demonstrated for the first time. These results provide further evidence for a direct effect of HIV proteinase inhibitors on one of the most relevant virulence attributes of C. albicans. In addition, we also observed that treatment of C. albicans proteins with the hydroxypyridone antimycotic agent clicopiroxolamine caused reduced proteolytic activity. Effects were not dose related. A possible explanation for this observation might be an interference of the antifungal agent/protease inhibitor at higher concentrations. One possible function of Sap activity is the attachment of C. albicans to epithelial cells and invasion into deeper tissue (Ollert et al., 1993; Watts et al., 1998; Schaller et al., 1999). Braga et al. (1992) investigated the effects of subinhibitory concentrations of clicopirox on the adherence of C. albicans to human buccal and vaginal epithelial cells. These authors found a significant reduction in adhesion at concentrations from 1/2

![Fig. 1. Effects of pepstatin A (○), amprenavir (●) and clicopiroxolamine (▲) on Sap activity of C. albicans strain 1 (a) and 2 (b). Each point represents the mean ± SD for three triplicate determinations. Differences in Sap activity between untreated and inhibitor-treated samples were highly significant (P < 0.001), as determined by the LSD test.](image-url)
to 1/16 of MIC₅₀ and explained this effect by a reduced intracellular uptake of essential substrates and ions necessary for the ability of *C. albicans* to express its adherence mechanisms. Our study suggests that ciclopiroxolamine also directly effects the activity of Saps of *C. albicans*, which, in turn, may cause reduced adherence *in vivo*.

**References**


