Histoplasmosis is an important systemic fungal infection, particularly among immunocompromised individuals, who may develop a progressive disseminated form which is often fatal if it is untreated. In such patients, the detection of antibody responses for both diagnosis and follow-up may be of limited use, whereas the detection of Histoplasma capsulatum var. capsulatum antigens may provide a more practical approach. We have recently described an inhibition enzyme-linked immunosorbent assay (ELISA) for the detection in patients’ sera of a 69- to 70-kDa H. capsulatum var. capsulatum-specific antigen which appears to be useful in diagnosis. To investigate its potential for the follow-up of histoplasmosis patients during treatment, antigen titers in the sera of 16 patients presenting with different clinical forms of histoplasmosis were monitored at regular intervals for up to 80 weeks. Sera from four of five patients with the acute form of the disease showed rapid falls in antigenemia, becoming antigen negative by week 14 (range, weeks 10 to 16). Sera from four patients with disseminated histoplasmosis showed falls in antigen levels; three of them became antigen negative by week 32; the fourth patient became negative by week 48. In contrast, antigen titers in four of six AIDS patients with the disseminated form of the disease remained positive throughout follow-up. Sera from only one patient who presented with the chronic form of the disease were analyzed, and this individual’s serum became antigen negative by week 9. The inhibition ELISA is shown to be of particular use in the monitoring of non-AIDS patients with the acute and disseminated forms of the disease and may complement existing means of follow-up.

Histoplasmosis is a systemic mycosis with a world wide distribution and is one of the most common systemic mycoses in North, Central, and South America (1, 12, 15, 16, 17, 20). The clinical spectrum varies from asymptomatic infection to severe, disseminated, and often fatal forms of disease (3, 8, 11, 22). Progressive disease is particularly severe in immunocompromised individuals (2, 8, 21). The definitive diagnosis of histoplasmosis relies on the isolation and identification of Histoplasma capsulatum var. capsulatum by culture or the direct visualization of the fungus (3, 19). However, both approaches may be problematic (10, 21). Antibody detection methods offer a rapid alternative to microbiological techniques, and the detection of antibodies to H. capsulatum var. capsulatum by immunodiffusion and the complement fixation (CF) test is often used (3, 13, 14, 17). However, the use of such tests for the follow-up of patients is problematic since anti-H. capsulatum var. capsulatum antibody titers remain elevated months or even years after successful therapy (3, 7, 8). In addition, false-negative results are often obtained for immunocompromised patients, since antibody titers may be low or absent in such individuals (24). The same may also be true in certain patients with the chronic, disseminated forms of disease (20, 21).

A more practical approach to both the diagnosis and the follow-up of patients with histoplasmosis may be the detection of H. capsulatum var. capsulatum antigen in body fluids. H. capsulatum var. capsulatum polysaccharide antigen has successfully been detected by radioimmunoassay (RIA) (4, 21, 25, 27), particularly in patients with AIDS, who develop disseminated histoplasmosis (21, 22, 24). In these patients, it has been shown that falls in antigenemia correlate well with effective therapy, making it feasible for clinicians to monitor treatment responses (21–23). However, there are problems associated with the use of RIA, notably relating to cross-reactivity with other dimorphic fungi such as Blastomyces dermatitidis (5, 21), Coccidioides immitis, Paracoccidioides brasiliensis, and Penicillium marneffei (26).

We have recently developed a novel H. capsulatum var. capsulatum antigen detection test (6) using a species-specific murine monoclonal antibody in an inhibition enzyme-linked immunosorbent assay (ELISA) system. We used this test to monitor the follow-up of 16 patients under treatment for different clinical forms of histoplasmosis, and it appears to be useful in this context.

MATERIALS AND METHODS

Patients and serum samples. Sixteen histoplasmosis patients with different forms of the disease were studied. Five patients presented with the acute pulmonary form, one patient presented with the chronic form, and four patients presented with the disseminated form: six of the patients had AIDS and the disseminated form of histoplasmosis (Table 1). The diagnosis for all individuals was confirmed either by cultural identification of H. capsulatum var. capsulatum or by direct identification of intracellular yeast cells in samples from the patients. CF and immunodiffusion (ID) tests were also performed. A total of 86 serum
samples taken both at the time of diagnosis and subsequently at regular intervals were analyzed; samples were collected between November 1991 and March 1998 at the Mycology Laboratory, Corporación para Investigaciones Biológicas, Medellín, Colombia. Details relating to the time of follow-up, medication, and length of therapy are included in Table 1. The ages of the patients varied from 1 to 62 years, with a mean age of 27.8 years; there were 12 males and 4 females. Fifty serum samples from healthy volunteers (normal human sera [NHS]) from areas where histoplasmosis is endemic were included as negative controls. All sera were aliquoted and stored at −20°C until use.

**Clinical evaluation.** Patients were evaluated clinically and radiographically at the time of diagnosis and subsequently at each follow-up visit. Clinical symptoms and signs were recorded, as were the results of a physical examination and chest X-rays. The serological tests described below were also carried out at similar intervals.

**Antigen detection by inhibition ELISA.** The inhibition ELISA was performed as described previously (6). In brief, inhibition standards were produced by adding increasing concentrations (from 4 to 66.5 μg/ml) of *H. capsulatum* var. *capsulatum* Hc 1980 cytoplasmic yeast antigen (CYA) to a pool of NHS (6). Constant aliquots of monoclonal antibody HIC were mixed with the inhibition standards, sera from patients with histoplasmosis, and control NHS, and these were incubated overnight at 4°C on a previously blocked microtiter plate (inhibition plate) to allow the occurrence of the inhibition reaction. The reaction plates were coated with *H. capsulatum* var. *capsulatum* Hc 1980 CYA and were incubated under the same conditions described above. On the next day the reaction plates were blocked and samples were transferred from the inhibition plate to the respective wells in the reaction plate, and after further incubation at 37°C, the plates were washed and incubated with goat anti-mouse immunoglobulin G peroxidase conjugate. The reaction was developed with o-phenylenediamine as described previously (6). The optical densities (ODs) were read at 490 nm; the values obtained with the inhibition standards were plotted to produce a standard inhibition curve. A regression model constructed with the reciprocal values of fixed concentrations of CYA and the ODs were used to calculate the antigen concentration in the samples tested. The cutoff point was established as the upper limit of the 90% least significant difference (LSD) confidence interval of the OD values obtained from the negative controls (2.2 μg/ml; the values obtained with the inhibition standards were plotted to produce a standard inhibition curve). The regression coefficient of the inhibition standard curve was 0.97; it accounted for 97.4% of the variations in the antigen concentrations in the samples. The inhibition standard curve was used to determine the concentration of *H. capsulatum* var. *capsulatum* antigen in each sample tested. The cutoff was defined as the upper limit of the 90% confidence interval (CI) for the negative controls (2.2 μg/ml). Thus, samples with an antigen concentration greater than this value were considered positive.

Patients were classified into four groups according to their clinical presentations (Table 1). The mean antigen concentration at the time of diagnosis is expressed as the means ± 90% CIs as follows: acute form (n = 5), 7.3 ± 3.1 μg/ml; disseminated form (n = 4), 10.37 ± 3.45 μg/ml; disseminated form with AIDS (n = 6), 11.57 ± 2.83 μg/ml; chronic form (n = 1), 6.97 μg/ml; and healthy controls (n = 50), 1.23 ± 0.75 μg/ml. Patients with the acute form, the disseminated form, and the disseminated form with AIDS had significantly higher levels of antigenemia compared to those for uninfected controls (P < 0.0001). It is noteworthy that patients with disseminated histoplasmosis (both patients with AIDS and patients without AIDS) had higher concentrations of circulating antigen compared to those in patients with other clinical forms (P < 0.01, based on the LSD by the multiple-range test).

Four of the five patients with the acute form of histoplasmosis had falls in antigen levels and became negative by week 16 (Fig. 1a). Two of them (patients 2 and 4) subsequently became briefly antigen positive before becoming negative again. The remaining patient (patient 5) had a more variable progression of antigenemia.
antigenemia with a peak at week 24, followed by a subsequent decline toward negativity. Overall, in this group of patients there was a significant reduction ($P < 0.01$) in the level of circulating antigen by week 13 after the initiation of therapy. This reduction was maintained until the end of the follow-up period. All patients in this group improved clinically and mycologically with itraconazole therapy, but certain residual abnormalities persisted. Generally, the antibody titers determined at the time of diagnosis by the CF test were low (Fig. 1b). In most individuals the titers subsequently rose and remained persistent.

The non-AIDS patients with disseminated histoplasmosis had decreasing titers of circulating antigen, and three of them became antigen negative by week 32 (Fig. 2a). The fourth patient (patient 8) was lost to follow-up at week 30, when a low antigen concentration (4.8 µg/ml) was detected; this individual returned for follow-up at week 68, at which point the antigen levels were undetectable (Fig. 2a). By taking these patients as a group, a significant ($P < 0.01$) fall in the circulating antigen level was observed by week 2 after the initiation of treatment. Clinical evaluation revealed that these patients improved markedly shortly after the initiation of therapy; this improvement was maintained over the period of observation. In this group underlying illness was not apparent, although the two youngest patients (patients 7 and 8) were severely undernourished. Antibody levels in this group at the time of diagnosis

![Graphs showing antigen and antibody levels for patients with acute and disseminated histoplasmosis](http://jcm.asm.org/...)

FIG. 1. Follow-up of patients with acute form of histoplasmosis. (a) Measurement of circulating 70-kDa antigen (Ag) levels as determined by inhibition ELISA. ■, patient 1; ○, patient 2; △, patient 3; ◊, patient 4; ●, patient 5. The dashed line represents the cutoff point (2.2 µg/ml) of the inhibition ELISA. (b) Measurement of antibody titers as determined by CF test. Symbols are the same as those in panel a. The patient numbers are listed in Table 1. NR, nonreactive.

FIG. 2. Follow-up of patients with the disseminated form of histoplasmosis. (a) Measurement of circulating 70-kDa antigen (Ag) levels as determined by inhibition ELISA. ■, patient 6; ◊, patient 7; △, patient 8; ○, patient 9. The dashed line represents the cutoff point (2.2 µg/ml) of the inhibition ELISA. (b) Measurement of antibody titers as determined by CF test. Symbols are the same as those in panel a. The patient numbers are listed in Table 1. NR, nonreactive.
were variable, and they remained so and did not predict the clinical outcome.

Patients with AIDS and disseminated histoplasmosis typically showed initial increases in circulating antigen levels (Fig. 3a), and only two patients (patients 10 and 15) became antigen negative during the course of follow-up. The remaining four patients maintained high levels of antigen during the follow-up period, with values that were, in some cases, greater than those detected at the time of diagnosis. Over the whole follow-up period there was a decrease in the concentration of circulating antigen which was significant ($P < 0.01$) only several weeks after the initiation of treatment. The clinical responses of this group were complex. The two patients (patients 10 and 13), in whom major clinical improvements were observed, had been treated not only with itraconazole but also with a combination of three antiretroviral drugs. In contrast, patient 14 did not respond to treatment, despite combined antimycotic and antiretroviral therapies. Finally, the three patients (patients 11, 12, and 15) who received only antimycotic treatment had poor clinical responses. The antibody responses in these patients were generally low (Fig. 3b), and only one patient had detectable antibody levels at the time of diagnosis, with two other patients developing antibody responses by the first follow-up appointment.

In this study we were able to monitor only one patient (patient 16) with the chronic pulmonary form of histoplasmosis. This individual demonstrated an increase in detectable antigen concentration from 6.97 to 8.73 $\mu$g/ml by the first follow-up appointment (data not shown). Subsequently, by week 10 of follow-up, this patient became antigen negative. Clinically, the patient was reported as being cured. The antibody response in this patient increased after the first follow-up visit (1:64) and remained positive (1:16) during the whole follow-up period.

**DISCUSSION**

The inhibition ELISA described previously has now been applied to the follow-up of patients undergoing treatment and has proved to be useful in this context. The assay methodology required no modification from that published originally (6), and the mean antigen concentrations at the time of diagnosis in patients with different clinical forms of histoplasmosis were highly reproducible. The correlation coefficient and the antigen cutoff point reported in this study were also similar to those described previously (6). In addition, samples which tested negative in the former study were also negative in this study (data not shown).

The inhibition ELISA appeared to be very useful for monitoring of the response to therapy in patients presenting with acute pulmonary histoplasmosis. Overall, these patients showed decreasing levels of circulating antigen, a fall which became statistically significant by week 13. The reductions in antigen levels closely mirrored the clinical improvements. In contrast, the detection of antibody levels was less useful since, in some patients, titers were low and remained so during the follow-up period, while in other patients they remained elevated even after an apparent clinical cure. The progress of patients with the acute pulmonary form of the disease has historically been difficult to monitor by antibody detection techniques (3, 7).

All of the non-AIDS patients with disseminated histoplasmosis demonstrated a statistically significant fall in antigen levels 2 weeks after the commencement of treatment, a fall which was more rapid than that seen in patients with the acute form of histoplasmosis. Circulating antigen was completely cleared from all patients with the disseminated form of the disease. This correlated well with complete clinical cure. In contrast, the antibody levels in these patients were variable and did not appear to predict the clinical outcome. Similar variabilities in antibody levels in patients with this form of histoplasmosis have been documented previously (8, 10).

In general, there was no clear pattern in terms of antigenemia during the follow-up of AIDS patients presenting with the disseminated form of histoplasmosis. However, all six patients exhibited some increase in the concentration of circulating antigen at the first follow-up visit after the induction of
therapy. Similar observations have previously been reported by Wheat et al. (22, 23), who used an RIA which detects a polysaccharide antigen. It is unclear why this should be so, and this question awaits further investigation. Following the rise in antigenemia, there was then a trend toward reductions in antigen levels; such reductions attained statistical significance by week 8 after the initiation of treatment. However, in the majority of the patients (four of six), high levels of circulating antigen persisted up to the last follow-up visit. The persistence of \textit{H. capsulatum} var. \textit{capsulatum} polysaccharide antigen during follow-up has also been observed by RIA (18, 22, 23), although apparently not to the same extent seen here. All of the patients included in this study received itraconazole therapy at various doses (200 to 400 mg/d), and Wheat et al. (18) reported slower rates of clearance of antigenemia after treatment with itraconazole than after treatment with amphotericin B. The persistence of antigenemia is also seen in AIDS patients with cryptococcosis (9). Antigen levels may remain detectable in AIDS patients due to the persistence of low-level infection that is partially controlled by maintenance therapy but which is not completely eradicated due to the impairment of the cellular immune response. Five of six of these patients had CD4 counts of less than 150 cells/mm$^3$ at the time of diagnosis; unfortunately, CD4 counts were not determined at subsequent time points, making it impossible to correlate cellular immune status with a fall in antigenemia.

Three patients in this group were also receiving antiretroviral treatment (patients 10, 13 and 14); only one of them (patient 10) became antigen negative and at the same time showed a marked clinical improvement. Overall, there would appear to be no clear correlation between the use of combined antifungal and antiretroviral treatment and a fall in antigenemia in our study. However, the situation is made more complex by the fact that some of the patients included in this group were severely immunocompromised and presented with up to four concomitant opportunistic infections (patients 11 and 12). As has been reported elsewhere (3), antibody levels in these patients were low and were not generally useful for follow-up.

A fall in antigen titers was also seen in the patient with the chronic pulmonary form of disease. This correlated well with clinical cure, but because the group had only one patient, it would be unwise to draw conclusions as to the general effectiveness of the inhibition ELISA for such patients.

In conclusion, the inhibition ELISA previously described as a diagnostic tool has also proved to be useful for the monitoring of the progress of patients under treatment for acute pulmonary and disseminated forms of histoplasmosis. This is of particular significance given that there are no previous reports about the follow-up of these groups of patients by antigen detection techniques. Its effectiveness in AIDS patients with the disseminated form of histoplasmosis is less clear, as is the case with regard to patients with the chronic pulmonary form of the disease; there is a need for an expanded study to encompass a larger number of such individuals. The results of our work indicate that regular assessment of antigen titers by the inhibition ELISA is useful in the determination of the response to therapy. In general, on the basis of the results of this study, we suggest that human immunodeficiency virus-positive patients be tested for antigenemia at 2-week intervals during the first 2 months of therapy, followed by more sporadic testing according to clinical judgment. Patients not infected with the human immunodeficiency virus but with the disseminated form of histoplasmosis should be tested at least twice during the first month of therapy and monthly thereafter; patients with the chronic form should be tested monthly for a period of up to 6 months after diagnosis.

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