Persistent damage to *Enterocytozoon bieneusi*, with persistent symptomatic relief, after combined furazolidone and albendazole in AIDS patients

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Abstract

**Aim**—To investigate morphological changes in *Enterocytozoon bieneusi* and the duration of symptomatic relief after combination treatment with furazolidone and albendazole in AIDS patients.

**Methods**—Four severely immunocompromised AIDS patients with symptomatic *E bieneusi* infection of the gut received an 18 day course of combined furazolidone and albendazole (500 + 800 mg daily). All patients were monitored for parasite shedding in stool by light microscopy at the end of treatment and monthly during follow up. At the end of treatment, duodenal biopsy specimens obtained from three patients were studied by transmission electron microscopy by two pathologists blind to the patients’ treatment or clinical outcome. Duodenal biopsy specimens obtained from one of the patients two months after completion of treatment were also studied electronmicroscopically.

**Results**—All patients had long lasting symptomatic relief, with a major decrease—or transient absence—of spore shedding in stools from completion of treatment. After treatment, changes in the life cycle of the parasite; moreover, relapses rapidly occur after discontinuation of treatment. Also albendazole, which is considered the drug of choice for microsporidiosis caused by *Encephalitozoon* spp, was reported to be poorly and variably effective against *E bieneusi*, although changes in the parasite were described after treatment.

Recently we reported both symptomatic relief and light and electron microscopic changes in *E bieneusi*, with decreased parasite shedding in the stool, after treatment with furazolidone in six patients with AIDS. The infection, however, was not eradicated by the schedules of treatment used, and two patients had clinical and parasitological relapse two months after treatment was discontinued.

A search for more lasting results led us to use a combination of furazolidone and albendazole to treat four AIDS patients with symptomatic *E bieneusi* infection of the gut. The rationale of this approach lies in the fact that the mechanism of action of these drugs seems complementary, being directed at different metabolic pathways in the parasite. Furazolidone probably interferes with the synthesis of nucleic acids by an action on the Krebs cycle, while albendazole, which inhibits tubulin polymerisation into microtubules of parasites, may affect the precursors of polar tubes of *E bieneusi*.

In this study our aim was to confirm the results obtained using furazolidone alone, and to evaluate the ability of combined furazolidone and albendazole to produce persistent clinical and parasitological remission without the need for maintenance treatment.

**Methods**

We enrolled four HIV positive drug misusing patients (three male, one female) with symptomatic *E bieneusi* infection of the gut. The patients were aged between 31 and 35 years. All had AIDS at the time of study entry; according to the 1993 CDC revised classification system, two cases belonged to the C, stage, while the others to the B, stage. The mean CD4+T cell count was $7.7 \times 10^3$/litre (range 4 to 11). All patients had chronic diarrhoea (for at least two months), with a range of two to five watery stools a day (mean four). Weight loss ranged between 3 and 8 kg (mean 5.5).
Stool were processed according to the methods of Weber et al and of Van Gool et al to detect microsporidia.17–19 The amount of spore shedding was evaluated by a semiquantitative criterion on 100 microscopic fields: spores were defined as very frequent if present in every field, frequent if present in more than 50 but fewer than 100 fields, rare if present in fewer than 50 fields, and very rare if sporadically present.

Before study entry, frequent to very frequent spores of microsporidia were present in stools of all patients. Stool samples were also examined for bacteria, Clostridium difficile toxin, mycobacteria, eggs and parasites, adenoviruses, and rotaviruses, with negative results.

Transmission electron microscopy (TEM) was performed on endoscopic biopsy specimens taken from the third part of duodenum before treatment. At least two samples from each patient were processed. Specimens were fixed with glutaraldehyde 2.5% in cacodylate buffer (pH 7.4) for three to four hours at room temperature, washed in the same buffer, and postfixed in osmium tetroxide in 1% Milloning buffer (SIC, Rome, Italy) for one hour at room temperature. After dehydration through graded acetone series, the specimens passed through propylene oxide and were then embedded in Epon 812. Semithin sections (3 µm) were cut, stained with toluidine blue–sodium tetraborate, and studied by light microscopy. About 100 ultrathin sections were obtained from all the selected specimens and were stained with uranyl acetate and alkaline bismuth subnitrate. For each section at least three random fields were examined by TEM. Care was taken to evaluate only areas away from the borders of the biopsy specimen to avoid traumatic artefact.

All patients underwent a baseline evaluation that included full blood cell count, urine analysis, electrolytes, liver function tests, and abdominal ultrasonography.

Combination treatment with furazolidone (100 mg tablet five times daily) and albendazole (400 mg tablet twice daily) was given orally to all the patients for 18 days. Written informed consent was obtained before treatment. No concomitant antidiarrhoeal drugs were allowed. Clinical assessment and haematological and liver function tests were monitored after 10 days and at the end of treatment. Adverse events were evaluated.

Three days after completion of treatment, upper endoscopy was performed in three patients who gave their consent. One of these patients also underwent endoscopy two months after discontinuation of treatment. In all cases, at least two biopsy specimens were taken from the third part of duodenum, and were processed as described above. The ultrathin sections obtained were examined by TEM by two pathologists blind to the patients’ treatment or clinical outcome.

In all patients spore shedding in the stool was monitored by Weber’s method at the end of treatment and monthly during follow up (one stool sample a day for three days). Daily stool frequency and body weight were charted, and the CD4+T cell count was repeated every four months after discontinuation of treatment. The mean duration of follow up was eight months (range four to 15).

A complete response was defined as cessation of diarrhoea (stool frequency less than twice a day), stabilisation of body weight, and a major decrease (or clearance) of spore shedding in the stools for more than three months. A partial response was defined as improvement in baseline stool frequency, with continuing weight loss, and slight or no decrease in spore shedding in stool.

**Results**

CLINICAL RESPONSE

All patients showed a complete response, with remission of symptoms and a major decrease or absence of spore shedding in the stools from the time of completion of treatment. The drugs were well tolerated, although, probably as a side effect of furazolidone,20 there was a slight worsening of diarrhoea in all patients during treatment. This was promptly reversed when treatment was discontinued, and there were no alterations in the laboratory test results attributable to the drugs used.

Over the follow up period no patient had a relapse, either clinical or parasitological, despite the fact that no maintenance treatment was given and no improvement in CD4+T cell count was documented.

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Figure 1 Microsporidian spores (stool specimen after treatment). Many spores are evident, almost all showing a pronounced red stained clot. A normal spore is seen (arrow). (Magnification ×795.)
Case 1 regained his preinfection body weight and was asymptomatic during four months of follow up, although very rare to frequent microsporidian spores, most of which were dysmorphous, were intermittently excreted in the stools. The patient later moved to another city and was lost to the follow up.

Case 2 was symptom-free over 15 months of follow up, and regained his preinfection body weight. He showed clearance of parasite in stool, with occasional shedding of very rare to frequent spores, most of which were dysmorphous.

Case 3 had six months of follow up showing parasite clearance in stool or intermittent shedding of very rare spores, most of which were dysmorphous. The patient regained his preinfection body weight and was asymptomatic until he died of *Rhodococcus equi* pneumonia and *Staphylococcus aureus* septicemia.

Case 4 had no diarrhoea during seven months of follow up, although there was intermittent shedding in the stools of very rare to very frequent spores, almost all being dysmorphic. The patient showed body weight increase, and had neither clinical nor parasitological relapse until his death from severe intestinal bleeding.

### Parasite Changes in the Stools

Parasite changes were detected by Weber’s stain in all patients’ stools after treatment. In these cases a marked red stained clot was distinguishable within the spores (fig 1), and replaced the pinkish red belt-like stripe originally described by Weber et al. The mean number of altered spores was 10 in each microscopic field (range 1–50).

### Parasite Damage in Biopsy Specimens

Unaltered phases of the *E bieneusi* life cycle were found in all fields studied from biopsies...
taken before treatment (fig 2), \(^{11}\) and the parasites were present in almost all of the semi-thin and ultrathin sections obtained.

After treatment, ultrastructural changes at all stages of *E bienesi* were observed in the biopsy specimens, and were most pronounced in sporogonial plasmodia and spores. Most sporogonial plasmodia had disarranged nuclear chromatin, and many showed disruption of nuclear membranes, irregularity of nuclear profiles and diastasis between the nuclear membrane and the nucleoplasm (figs 3 and 4).

Several electron-dense discs also failed to develop into polar tubes and appeared disorganised, deformed, disrupted, and pale (figs 3 and 4). Most spores were very electron-dense, and showed enlarged electron-lucent vacuoles (fig 5).

These changes were observed both in biopsy specimens obtained just after completion of treatment and two months later, and were similar to changes reported after furazolidone alone.\(^ {13}\)

There was a substantial decrease in microsporidia, documented by both light microscopy and TEM, in all biopsies taken either just after completion of treatment or two months later, and the parasites were found in fewer than half of all semi-thin and ultrathin sections obtained. A broken spore and a giant spore containing two electron-lucent vacuoles and multiple coils of the polar tube were also observed in biopsies taken after treatment (figs 6 and 7).

These alterations were not found after treatment with furazolidone alone.\(^ {13}\) No abnormalities in host cell structures were seen in biopsy specimens obtained before and after treatment.

### Discussion

Combination treatment with furazolidone and albendazole was associated with a complete response in four severely immunocompromised AIDS patients with symptomatic *E bienesi* infection.

All patients had long lasting symptomatic relief with a substantial decrease—or transient absence—of protozoan shedding in the stools, and regained their preinfection body weight. The regimen was well tolerated by all the patients, and there were no alterations in the laboratory test results attributable to the drugs used. During long term follow up no patient had a relapse, either clinical or parasitological, despite there being no documented improvement in CD4+T cell count and no maintenance treatment. Such a prolonged effect was not seen after treatment with furazolidone alone,\(^ {13}\) nor has it been reported to date after any other treatment against *E bienesi*.

Combination treatment with furazolidone and albendazole resulted in persistent alterations in the parasite. Ultrastructural changes at all stages of *E bienesi* were documented not only in biopsy specimens taken at endoscopy just after completion of treatment, but also in biopsies obtained two months later. These changes were most pronounced in sporogonial plasmodia and spores, and were similar to changes observed after furazolidone alone,\(^ {13}\) including osmiophilic spores with enlarged vacuoles as revealed in a retrospective TEM study.

A substantial decrease in microsporidia was a persistent feature of all biopsies obtained, either immediately after discontinuation of treatment or two months later. We also found both a broken spore and a giant spore, suggesting incomplete division of the sporonts. These findings are of interest as they were not observed after furazolidone alone,\(^ {13}\) nor have they been reported with any other treatment, though giant spores have previously been found with albendazole treatment.\(^ {11,12}\) Furthermore, dysmorphous spores were persistently...

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**Figure 5** Mature spores (biopsy specimen after treatment). Osmiophilic spores are present, some containing a very enlarged vacuole. (Magnification ×9800.)

**Figure 6** Mature spores (biopsy specimen after treatment). Very electron-dense spores are seen. One spore is broken (arrow). (Magnification ×6380.)

**Figure 7** Giant spore (biopsy specimen after treatment). Two electron-lucent vacuoles, multiple polar tube coils, and a polaroplast are shown. (Magnification ×6570.)
Combined furazolidone and albendazole for E bieneusi
documented in all patients’ stool by Wéber’s method after combination therapy, and were
correlated to the post-treatment parasite alternate	ions seen by TEM in biopsy specimens. These
changes in faecal spores were identical to those
previously described after furazolidone alone,13
and were not seen in our four patients before
treatment. We hypothesise that these changes
could derive from furazolidone mediated damage
to the spore coat, with subsequent increasing
permeability, as suggested by osmiophilic spores seen by TEM in biopsy specimens.
We are unable to explain why dysmorphic spores persisted for long periods in all the
patients’ stools after combination treatment was discontinued. We can only hypothesise that this
might have been caused by the persistence of genomic damage to the parasite induced by
furazolidone and transmissible, as an inherited
defect, to the progeny during reproduction.
This hypothesis agrees with our TEM results of
persistent alterations in E bieneusi, at all stages
of the life cycle, in all biopsies obtained two
months after stopping treatment.

Both the decrease of parasite load and the parasite damage seen in post-treatment biop-
sies may account for the decrease in parasite load and the spore changes persistently seen in stool after treatment. These findings are in agreement with a recent report that closely
linked the quantitation of E bieneusi in tissue with quantitation in the stools,22
and may explain the long lasting symptomatic relief we observed during follow up in all the
patients.

Our preliminary results suggest that combined furazolidone and albendazole have a
direct action on E bieneusi, probably by inhibiting
parasite multiplication, and may be effective
in achieving persistent clinical and parasiti-
tological remission even in patients with severe
immunodeficiency. Whether our results can be interpreted as an enhancement of albendazole
activity secondary to parasite damage caused by furazolidone is difficult to assess at this
stage, although a complementary effect, as
suggested by clinical and parasitological re-
sponses, seems probable.

This study confirms the results reported after furazolidone alone,13 and emphasises the
importance of combining furazolidone with albendazole to achieve a substantial and
persistent reduction in the parasites, avoiding
the need for maintenance treatment. However,
although this combination greatly reduced the numbers of microsporidia, it did not com-
pletely eradicate the infection as some parasites
developed normally and gave rise to normal
spores. Nevertheless, our study supports the
efficacy and safety of a new drug regimen for
the treatment of E bieneusi microsporidiosis.
The long lasting symptomatic relief observed
after combined furazolidone and albendazole
was correlated with the persistent decrease in
parasite burden, both in the tissues and in the
stools, and with the morphological changes of
the protozoan life cycle.

The long term natural history of E bieneusi infection is not well understood, and spontane-
ous remission—as occurs with crypto-
sporidiosis—is quite often seen in clinical
practice. Randomised controlled trials are
required to validate the results of our un-
trolled pilot study and to dispel doubts about spontaneous remission or improvement result-
ning from the treatment of concomitant infec-
tions. These factors were unlikely to have been
operative in our cases, however, since all the
patients showed a pronounced response to the
combination treatment, while no spontaneous
remission, either clinical or parasitological, had
been observed before. Moreover, to our knowl-
edge, none of the drugs we used to treat
concomitant infections (table 1) has any
reported activity against E bieneusi.

Table 1. Concomitant infections

<table>
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<tr>
<th>Case</th>
<th>Disease</th>
<th>Treatment</th>
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<tbody>
<tr>
<td>1</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>2</td>
<td>Oral candidiasis</td>
<td>Fluconazole</td>
</tr>
<tr>
<td>3</td>
<td>Oesophageal candidiasis</td>
<td>Fluconazole</td>
</tr>
<tr>
<td>4</td>
<td>Rhodococcus equi</td>
<td>Vancomycin + rifampicin + clarithromycin</td>
</tr>
<tr>
<td>5</td>
<td>Staphylococcus aureus septicemia</td>
<td>Vancomycin</td>
</tr>
<tr>
<td>6</td>
<td>Oesophageal candidiasis</td>
<td>Fluconazole</td>
</tr>
</tbody>
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