Acyclovir in the prevention of duodenal ulcer recurrence

S J Rune, J Linde, O Bonnevie, H Draminsky Petersen, F Bendtsen, J H Wandall, C Gluud, J Eriksen, H Skovbjerg, M Vilien, M Olesen, S Larsen, P Schlichting, M Gormsen, H R Wulff, S Vadstrup, B F Vestergaard

Abstract

This study tests the hypothesis that reactivation of a latent herpes simplex virus infection may be a cause of recurrent duodenal ulceration. Patients with recently healed duodenal ulcer were entered into a double blind, randomised study of maintenance treatment with the antiviral drug acyclovir (400 mg bid) versus placebo, to determine if suppression of herpes virus infection would influence the natural history of the ulcer disease. One hundred and fifteen patients entered the trial and 76 patients completed it according to the protocol. Endoscopy was performed when ulcer symptoms recurred and at the end of the 25 week trial period. In the acyclovir group the cumulated relapse rate was 63% compared with 56% in the placebo group (NS). This result suggests that reactivation of herpes simplex virus is not a cause of recurrent duodenal ulcer.

The rapidly changing epidemiology of duodenal ulcer during this century6 points to an extrinsic factor as a possible cause of the disease, and 30 years ago it was proposed that infection with herpes simplex virus might be involved in the pathogenesis.7 Since then this hypothesis has been supported by demonstration of herpes simplex virus in human vagal ganglia8 and the finding of significantly higher titres of anti herpes simplex type 1 antibodies in patients with duodenal ulcer than in seropositive control individuals.9 Herpes simplex virus has also been found in the gastric mucosa, but only in patients being treated with immunosuppressive agents.10

As acyclovir was shown to be safe and effective in the prophylaxis against reactivation of herpes simplex virus5 we decided to use this drug to test the hypothesis that herpes virus is responsible for recurrent duodenal ulceration.

Methods

Patients

The patients included in this study had an endoscopically diagnosed duodenal ulcer which subsequently had been treated with an H2 blocker. Patients were considered eligible for inclusion if their duodenal ulcer was seen to have healed within eight weeks of treatment, if they had become symptom free, if their H2 blocker treatment was stopped, and if they were found to be seropositive for antip herpes simplex antibodies.

Patients who gave their informed consent entered the double blind study. They were allocated at random to take trial tablets containing either acyclovir 400 mg or placebo, one to be taken in the morning and one in the evening for 25 weeks. The patients were seen at five week intervals for clinical assessment and a tablet compliance count.

The patients contacted the clinic if ulcer symptoms recurred on three or more days within the same week, and an upper gastrointestinal endoscopy was then performed.

Treatment was stopped if duodenal ulcer recurrence was verified at endoscopy. All patients who completed the 25 week study without symptomatic ulcer recurrence were endoscoped at the end of the study. Patients who missed taking tablets for more than three days during a five week period and patients who developed a gastric ulcer without a simultaneous duodenal ulcer were withdrawn.

The efficacy variable was endoscopic recurrence of a duodenal ulcer and the data were subjected to both a 'per protocol analysis' - that is, including only patients who completed the study according to the protocol, and an ' intention to treat' analysis - that is, including all patients who entered the study, and considering all patients who were withdrawn as treatment failures - namely, those having relapsed.

Results

One hundred and fifteen patients entered the study and the treatment groups were found to be comparable. Details are shown in Table I. Thirty nine patients were withdrawn from the study for the following reasons: 19 because of non-compliance - that is, missed drug intake on more than three days during a five week period; seven because endoscopy was not performed at symptomatic relapse before treatment with an H2 blocker.
TABLE I  Patient characteristics

<table>
<thead>
<tr>
<th></th>
<th>Acyclovir n=55</th>
<th>Placebo n=60</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ratio Men:Women</td>
<td>37:18</td>
<td>36:24</td>
</tr>
<tr>
<td>Mean age (SD; years)</td>
<td>54.4 (11.8)</td>
<td>53.5 (14.5)</td>
</tr>
<tr>
<td>History of cold sore</td>
<td>10</td>
<td>17</td>
</tr>
<tr>
<td>Cigarette smokers (n)</td>
<td>35</td>
<td>37</td>
</tr>
</tbody>
</table>

NS=not significant, Student’s non-paired t test and Fischer’s exact test.

TABLE II  Cumulative recurrence rates for two analytical cohorts

<table>
<thead>
<tr>
<th></th>
<th>Acyclovir n</th>
<th>Placebo n</th>
</tr>
</thead>
<tbody>
<tr>
<td>‘Per protocol’ cohort</td>
<td>22/35 (63%)</td>
<td>25/41 (56%)</td>
</tr>
<tr>
<td>‘Intention to treat’ cohort</td>
<td>42/55 (76%)</td>
<td>42/60 (70%)</td>
</tr>
</tbody>
</table>

blocker was given; five patients did not attend at scheduled visits; four patients developed a gastric ulcer and one patient an oesophageal ulcer; and three patients wanted to leave the study.

The cumulative recurrence rate of duodenal ulcer after 25 weeks was 63% in the acyclovir group and 56% in the placebo group (standard error of difference = 11-0%), Table II. The ‘intention to treat’ cohort showed a similar result with 76% and 70% relapses in the acyclovir and placebo group respectively. The time of recurrence did not differ between the two groups (Figure). Only one asymptomatic ulcer was found at endoscopy at 25 weeks. Fourteen patients (six acyclovir, eight placebo) developed symptomatic relapse with no demonstrable ulcer at endoscopy.

No side-effects were recorded.

Discussion

The aim of this clinical trial was to test the hypothesis that reactivation of a latent herpes simplex infection is a cause of recurrent duodenal ulceration. The results show that longterm treatment with acyclovir does not reduce the recurrence rate of duodenal ulcer, and this is a strong argument against the hypothesis, provided that the chosen regimen effectively prevents reactivation of herpes simplex virus. It has in fact been shown in several studies** that the dose of acyclovir used in the present study reduces the risk of developing clinical or virological recurrence of herpes simplex infection by at least 75%. If acyclovir had prevented ulcer recurrence in, for example, 50 to 75% of the patients, the expected difference between the recurrence rates of the order of magnitude would be 30% to 50%. That hypothesis, however, is not compatible with this study as the calculation of the 99% confidence limits of the observed difference shows that we are unlikely to have missed a difference larger than 22%.

One of the arguments in favour of the hypothesis was that duodenal ulcer patients were found to have higher antiherpes virus antibody titres in their serum than matched controls. The immune response to a chronic viral infection such as herpes is, however, regulated in a highly complex fashion involving many elements. Raised antibody titres to herpes simplex virus may therefore be explained by a non-specific hyper-responsiveness of ulcer patients and not necessarily by activation of the virus. This explanation is supported by the finding that ulcer patients also have higher titres of antibodies against rubella virus.11

Acyclovir and placebo tablets were kindly provided by The Wellcome Research Labs, Beckenham, England.

11 Van der Merwe CF, Alexander JJ. Herpes simplex virus and duodenal ulceration. Lancet 1982; i: 762.
Acyclovir in the prevention of duodenal ulcer recurrence.

S J Rune, J Linde, O Bonnevie, H D Petersen, F Bendtsen, J H Wandall, C Gluud, J Eriksen, H Skovbjerg and M Vilien

Gut 1990 31: 151-152
doi: 10.1136/gut.31.2.151

Updated information and services can be found at:
http://gut.bmj.com/content/31/2/151

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections
Stomach and duodenum (1689)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/