Treatment of periodontal infections due to anaerobic bacteria with short-term treatment with metronidazole

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Abstract. In the present report, five selected periodontal patients were treated for 1 week with metronidazole. Two of the patients had their teeth scaled and root-planed the week they received metronidazole. Prior to treatment, B. asaccharolyticus accounted for 41% of the cultivable isolates and the spirochetes averaged 29% of the microscopic count in plaque removed from each of four pockets per patient. The presence of these elevated proportions of periodontopathic bacteria combined with the presence of periodontal pockets and attachment loss suggested that the patients were in a state of an active infectious process involving primarily anaerobic bacteria. If this be the case, then antimicrobial therapy directed against these anaerobes with metronidazole was indicated. The 1-week treatment with metronidazole significantly reduced the proportions of these organisms for up to 6 months after treatment. Coincident with these findings was an improvement in the clinical parameters, especially in those sites that initially had greater than 5 mm pocket or attachment loss. These sites showed a 2 mm or more reduction in pocket depth and an almost 2 mm gain in apparent attachment that was evident 6 months after treatment.

The results obtained were in only five patients. However, the magnitude of improvement suggests that antimicrobial therapy directed against anaerobic organisms may be a valuable adjunct to periodontal therapy.

Bacteriological investigations of plaques taken from various periodontal disease entities indicate that absolute and/or relative increases in presumably indigenous organisms are associated with the disease state (Loesche 1976a, Socransky 1977). An increase in spirochetes, especially of the intermediate and large sizes, was seen in acute necrotizing ulcerative gingivitis (ANUG) (Listgarten & Lewis 1967) and in periodontitis (Listgarten & Helldén 1978, Keyes et al. 1978); an increase in Bacteroides asaccharolyticus was observed in advanced periodontitis (Slots 1977); an increase in gliding saccharolytic rods, including Capnocytophaga species, was found in juvenile periodontitis (Newman & Socransky 1977); collagenase-producing Bacillus cereus strains were isolated from periodontitis in trisomy 21 patients (Loesche et al. 1974); Actinomyces viscosus increased in experimental gingivitis (Loesche & Syed 1978) and possibly in chronic periodontitis (Williams et al. 1976); Bacteroides melaninogenicus ss. intermedius increased during pregnancy gingivitis (Kornman & Loesche 1980); and Actinobacillus
actinomyceetemcomitans and Eikenella coro-
dens increased in active periodontitis (Tan-
ner et al. 1979). All of these species, with
the exception of the untested spirochetes,
cause periodontal pathology in the germ-
free rodent (Socransky 1977).

These findings individually and collective-
ly lend support for the specific plaque
hypothesis which states that only certain
plaques cause "periodontal infections" due
to the presence of one or more pathogens
and/or to a relative increase in the levels
or proportions of one or more potentially
periodontopathic organisms (Loesche 1976a).
The presence of discrete microbial profiles
associated with many of the above clinical
entities suggests that definitive treatment
should be aimed towards the elimination
and/or suppression of the suspected perio-
dontopathic organism(s). The nature of this
definitive treatment is in a state of evolu-
tion, but would include intensive mechanical
plaque control procedures (Lindhe & Ny-
man 1975, Rosling et al. 1976a, Rosling et
al. 1976b, Axelsson & Lindhe 1978, Knowles
et al. 1979), as well as the judicious usage
of antimicrobial agents. Topical antimicrobi-
als will improve the gingival health and reduce
plaque accumulations (Löe & Schiott 1970,
Loesche & Nafe 1973), but apparently, are
not able to effectively penetrate the pocket
sites and alter the levels of periodontopathic
organisms contained therein. This reservoir
of periodontopathic organisms conceivably
could be eliminated by systemic antibiotics
and/or by delivery of an antimicrobial agent
directly into the pocket either by irrigating
devices (Keyes et al. 1978), or by hollow
fiber device (Goodson et al. 1979).

Several studies suggest that systemically
administered tetracyclines either alone or in
conjunction with mechanical treatment can
change the profile of the pocket flora from
a predominantly Gram-negative flora to a
predominantly Gram-positive flora (List-
garten et al. 1978, Osterberg et al. 1979,
Slots et al. 1979, Williams et al. 1979). The
clinical findings "revealed that the adminis-
tration of tetracycline had only a minor
effect on the parameters examined" (Held-
dén et al. 1979). Other systemic anti-
microbials have not received similar clinical
trials. One potentially useful systemic agent
is metronidazole (Flagyl®), an antiprotocoal
agent that has a unique spectrum of activity
against anaerobic bacteria, including such
periodontopathic organisms as B. asaccharo-
lyticus and the various spirochete species
(Sutter & Finegold 1977, Chow et al 1977,
Davies et al. 1964). Metronidazole is re-
markably effective in the treatment of
ANUG (Shinn 1962, Duckworth et al. 1966),
but apparently has not been used in humans
for the treatment of the various forms of
periodontitis. In beagle dogs, a 28-day treat-
ment with metronidazole decreased plaque,
prevented the development of gingivitis and
maintained a plaque flora dominated by
coccal and rod forms, i.e. a nondisease-
associated flora (Heijl & Lindhe 1979).

In the present report, five selected perio-
dontal patients were treated for 1 week with
metronidazole. Various bacteriological and
clinical parameters were monitored periodi-
cally for 6 or more months following the
initial treatment.

Material and Methods

Patients. All patients were medically
healthy, and all but one had no recollection
of recent antibiotic usage. Patient D had
received antibiotics several times in the im-
mmediately preceding 6-month period for
recurrent periodontal abscesses. All patients
were advised of the mutagenicity of
metronidazole in bacterial systems (Ros-
kranz & Speck 1977), and of its adverse
reaction with alcohol consumption. They
were told that the usage of metronidazole
in periodontal infections was experimental
and that we were investigating this usage
under an investigational new drug exemption from the Food and Drug Administration (FDA).

Treatment schedule. Patients D, L, and H were given 21 tablets of metronidazole (Flagyl, 250 mg, G. D. Searle Co.), and advised to take one tablet three times a day for 7 days. Patients E and R were given the same tablets and instructions, but had one-half of the dentition scaled and root-planed on the first day of metronidazole therapy, and the other half scaled and root-planed on the last day of metronidazole therapy. No oral hygiene instructions were provided, nor were patients given any additional mechanical therapy during the 6- to 8-month period of observation, with one exception. Patient H received a thorough scaling and root-planing of the maxillary dentition approximately 22-25 weeks after she finished her 1 week of metronidazole therapy.

Bacteriological procedures. Plaque was removed from one periodontal pocket per quadrant in each patient for a total of four samples per patient. The sites chosen appeared from X-ray examination to be the most severely involved in each quadrant and were usually about molar teeth. The marginal plaque about the site was removed with a curette and discarded. The curette was introduced into the pocket and extended as far apically as possible. The root surface was then scaled, and the adherent plaque on the scaler tip was transferred to a vial containing 4 ml of reduced transport fluid (RTF) without EDTA (Loesche et al. 1972). A second plaque sample from the same site was added to 0.3 ml physiological saline with 1% gelatin (Listgarten & Hell-dén 1978), and used for the microscopic enumeration of spirochetes. The first plaque sample was immediately placed within the anaerobic chamber, sonically dispersed for 20 sec with a Kontes sonifier (Vineland, NJ), serially diluted in RTF and plated automatically with a spiral plater (Spiral Systems, Inc., Cincinnati, Ohio), on a variety of nonselective and selective agar media. This dispersal procedure appears to give optimal recoveries of Gram-negative organisms from plaque samples (Syed & Loesche 1978). Concurrent studies with this methodology revealed that the percent recovery of viable organisms from periodontal samples averaged about 65% of the microscopic count (Kornman, unpublished data). The total count and the *B. asaccharolyticus* counts were obtained from growth on enriched Trypticase soy agar (ETSA) (Syed et al. 1980). Representative brown-black colonies were Gram stained and tested for glucose fermentation. There was no difficulty in distinguishing *B. asaccharolyticus* by colony morphology from the red-brown pigmented colonies of *Actinomyces odontolyticus*.

The second plaque sample was dispersed by vortexing, and aliquots were viewed by dark-field microscopy. Either 200 organisms or the number of organisms in 20 high power fields (hpf) were enumerated, depending on which event occurred first. The number of spirochetes per 10 hpf are reported, as well as the percent of the spirochetes in the microscopic sample. The detailed cultural and microscopic findings for each site will be presented in a separate report.

Clinical examinations. The number of papillary bleeding sites and the magnitude of the bleeding were assessed so as to give a papillary bleeding score (PBS) (Loesche 1980). The method used was a modification of the sulcular bleeding index of Mühlemann & Son (1971), and consisted of a subjective estimate of the amount of bleeding which occurred from a papillary site after the insertion of a Stim-U-Dent® toothpick. All teeth, including molars with furcation involvement, were scored for pocket depth.
and loss of attachment using procedures previously described (Ramfjord 1967). All readings were obtained by one individual (EM).

Statistics. The various measurements were repeated at intervals following the 1-week treatment. These were compared with the pretreatment values by the paired t-test.

Results
The findings will be presented in the form of a case report for each patient.

Table 1. Effect of 1 week of metronidazole treatment on clinical and bacteriological parameters: Patient D, age 48: Terminal periodontitis

<table>
<thead>
<tr>
<th>Clinical (values given in mm)</th>
<th>Pretreatment</th>
<th>0</th>
<th>6</th>
<th>19</th>
<th>26</th>
<th>38</th>
<th>72</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pocket reduction (16)</td>
<td></td>
<td>1.6±0.4</td>
<td>1.8±0.4</td>
<td>1.8±0.4</td>
<td>1.6±0.4</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gain in attachment (25)</td>
<td></td>
<td>1.5±0.3</td>
<td>1.5±0.3</td>
<td>1.6±0.3</td>
<td>1.2±0.3</td>
<td>0.9±0.3</td>
<td></td>
</tr>
</tbody>
</table>

Bacteriological

<table>
<thead>
<tr>
<th>% B. asaccharolyticus</th>
<th>35±5</th>
<th>6±6</th>
<th>5±3</th>
<th>9±4</th>
<th>12±6</th>
<th>8±3</th>
<th>20±11</th>
</tr>
</thead>
<tbody>
<tr>
<td>B. asaccharolyticus</td>
<td>7.8±0.9</td>
<td>0.8±0.8</td>
<td>0.3±0.1</td>
<td>0.7±0.6</td>
<td>2.5±1.5</td>
<td>2±1</td>
<td>13±9</td>
</tr>
<tr>
<td>CFU × 10⁶</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Microscopic

<table>
<thead>
<tr>
<th>% spirochetes</th>
<th>15</th>
<th>4</th>
<th>5</th>
<th>3</th>
<th>4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spirochetes/10 hpf</td>
<td>9.4±6</td>
<td>2±2</td>
<td>0.4±0.3</td>
<td>11.3±6</td>
<td>3.5±2</td>
</tr>
</tbody>
</table>

* Number in parentheses is number of sites which initially exhibited greater than 5 mm pockets or 5 mm attachment loss
* Average ± standard error
* Values significantly different from pretreatment value, paired t-test, *P* < .05
* Percentage of total cultivable count
* hpf = high power fields

Patient D was a 48-year-old white female whose periodontal health had progressively deteriorated in the preceding 5-year period. Upper arch extractions had been recommended, but the patient hesitated to have this done. She had a history of periodontal abscesses that were managed by local debridement and systemic antibiotics. At the initial examination, 16 sites had pockets ranging from 6 to 9 mm in depth and another 51 sites had pockets ranging from 3 to 5 mm in depth. Twenty-five sites had 6 to 8 mm attachment loss, and another 49 sites had 3 to 5 mm attachment loss. The
molar teeth had furcation involvement. The papillary bleeding score was 22, indicative of mild gingivitis. *B. asaccharolyticus* accounted for 35% of the cultivable flora from the four sample sites. Spirochetes were present, but were outnumbered by other organisms observed in the dark-field examination (Table 1).

The 1-week metronidazole treatment significantly reduced either the proportions or levels of *B. asaccharolyticus* for at least 19-38 weeks after treatment (Table 1). Seventy-two weeks after treatment the *B. asaccharolyticus* proportions had increased to 20%, but CFUs of *B. asaccharolyticus* were slightly elevated compared to the pretreatment values. The levels of spirochetes were reduced at 0 and 6 weeks, but returned to pretreatment values by 19 weeks. The decrease in pocket depth, as well as the change in apparent attachment distance from the pretreatment values, were calculated for each recall visit. The results achieved in the most severely involved sites are shown in Table 1. The metronidazole treatment was associated with a 1.6-1.8 mm reduction in pocket depth during the 72-week period following treatment. There was an apparent gain in attachment of 1.5 mm during the first 26 weeks, a 1.2 mm gain at 38 weeks, and a 0.9 mm gain after 72 weeks (Table 1). This patient did not receive any additional periodontal therapy during this 72-week period, but had taken antibiotics on one occasion for medical reasons.

*Patient L* was a 17-year-old black female who had had all first molars and maxillary incisors extracted in the previous year. X-rays taken at that time indicated that this patient exhibited classic molar-incisor ju-

<table>
<thead>
<tr>
<th>Table 2. Effect of 1 week of metronidazole treatment on clinical and bacteriological parameters. Patient L, age 17; Residual juvenile periodontitis</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Clinical (values given in mm)</strong></td>
</tr>
<tr>
<td>Pocket reduction (9)*</td>
</tr>
<tr>
<td>Gain in attachment (4)*</td>
</tr>
<tr>
<td><strong>Bacteriological</strong></td>
</tr>
<tr>
<td>% <em>B. asaccharolyticus</em></td>
</tr>
<tr>
<td>% <em>B. asaccharolyticus</em></td>
</tr>
<tr>
<td>CFU X 10^6</td>
</tr>
<tr>
<td><strong>Microscopic % spirochetes</strong></td>
</tr>
</tbody>
</table>

* Patient already had all first molars and maxillary incisors extracted
* Number in parentheses is number of sites which initially exhibited greater than 5 mm pockets or 5 mm attachment loss
* Average ± standard error
* Percentage of total cultivable count
* Pretreatment value significantly different from all other values, paired t-test, *P* < .05
Table 3. Effect of 1 week of metronidazole treatment on clinical and bacteriological parameters. Patient H, age 28; Advanced periodontitis

<table>
<thead>
<tr>
<th>Pre-treatment</th>
<th>Time after metronidazole treatment in weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
</tr>
<tr>
<td><strong>Clinical (value given in mm)</strong></td>
<td></td>
</tr>
<tr>
<td>Pocket reduction (49)*</td>
<td>1.7±0.1°</td>
</tr>
<tr>
<td>Gain in attachment (18)*</td>
<td>1.1±0.3</td>
</tr>
<tr>
<td><strong>Bacteriological</strong></td>
<td></td>
</tr>
<tr>
<td>% <em>B. asaccharolyticus</em></td>
<td>34±12</td>
</tr>
<tr>
<td>% <em>B. asaccharolyticus</em> × 10^6</td>
<td>15±9</td>
</tr>
<tr>
<td><strong>Microscopic</strong></td>
<td></td>
</tr>
<tr>
<td>spirochetes/10 hpf</td>
<td>190±83</td>
</tr>
<tr>
<td>% Spirochetes</td>
<td>37±9</td>
</tr>
</tbody>
</table>

* Number in parentheses is number of sites which initially exhibited greater than 5 mm pockets or 5 mm attachment loss

° Average ± standard error

Teeth were scaled and curetted during weeks 22–25.

Values are significantly different from pretreatment value, paired t-test, P < .05

Percentage of total cultivable count

hpf = high power field

venile periodontitis. When seen at the initial examination, the premolars and molars abutting the edentulous molar space exhibited bone loss and deep pockets, prompting us to make a diagnosis of residual juvenile periodontitis. *B. asaccharolyticus* accounted for 33% of the cultivable flora and averaged about 46 million colony-forming units (CFU) per pocket. One week of metronidazole treatment significantly reduced the proportions and levels of *B. asaccharolyticus* over the next 27-week period (Table 2). The spirochetes averaged about 37% of the microscopic count prior to treatment. The 1-week therapy with metronidazole caused an immediate 10-fold decrease in their proportions followed by a return at 6 and 27 weeks to about 25% of the microscopic count. The nine deep pockets showed an average reduction of 3.8 mm, 27 weeks after therapy. The four sites which initially had 6 mm or more attachment loss had an apparent gain of 3.8 mm in attachment at the 27-week recall visit (Table 2). This patient neither received periodontal therapy nor took antibiotics for medical reasons during this 27-week period.

Patient H was a 28-year-old black female who had had maxillary first molars and maxillary incisors extracted while in her teens. She believed that the extractions were because of "gum disease". When seen by us, she had advanced periodontitis with furcation involvement of some molars. Forty-nine sites had pockets ranging from 6 to 10 mm, and 18 sites exhibited attachment loss of 6 to 9 mm. *B. asaccharolyticus* accounted for 34% of the cultivable flora.
Table 4. Effect of 1 week of metronidazole treatment and scaling on clinical and bacteriological parameters. Patient E, age 21; Advanced periodontitis

<table>
<thead>
<tr>
<th>Clinical (values given in mm)</th>
<th>Pre-treatment</th>
<th>Time after metronidazole treatment in weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>0</td>
</tr>
<tr>
<td>Pocket reduction (31)</td>
<td></td>
<td>2.3±0.2</td>
</tr>
<tr>
<td>Gain in Attachment (4)</td>
<td></td>
<td>3.0±0.4</td>
</tr>
</tbody>
</table>

| Bacteriological               |               |     |     |     |     |     |     |
| % B. asaccharolyticus         |               | 46±11°  | 0   | 0   | 1.2±0.1 | 1.1±0.8  | 0   |
| B. asaccharolyticus          |               | 14±3°  | <.004  | <.004  | <.03  | 0.4±0.2  | <.06  | <.004  |
| CFU × 10⁶                    |               |     |     |     |     |     |     |

| Microscopic                  |               |     |     |     |     |     |     |
| spirochetes/10 hpf           |               | 16±10°  | 0   | 0   | 0.1±0.1  | 0   | 1.5±1  |
| % Spirochetes                |               | 28±15°  | 0   | 0   | 0.5±5   | 0   | 9±5   |

* Number in parentheses is number of sites which initially exhibited greater than 5 mm pockets or 5 mm attachment loss

* Average ± standard error

* Pretreatment values significantly different from all other values, paired t-test, P < .01

* Percentage of total cultivable count

* hpf = high power fields

and the various spirochetes for about 37% of the microscopic count (Table 3). Metronidazole treatment significantly decreased the proportions and levels of these organisms for at least 5 weeks. Thereafter, the proportions of both organisms gradually increased (Table 3). However, the absolute levels of spirochetes remained reduced by about 90%, compared to the pretreatment value (Table 3). The pockets were reduced by 1.5–2.3 mm during the first 20 weeks, and there was an apparent increase in attachment of 1.1 mm (Table 3). The patient had received systemic tetracycline at week 11 for a medical infection. At weeks 22–25, the patient received a thorough scaling and root-planing of her maxillary teeth which could account for the further pocket reduction and attachment gain noted at week 33. This mechanical therapy did not appear to affect the B. asaccharolyticus values, but might have lowered the spirochetal values (Table 3).

Patient E was a 21-year-old black male who presented with a diffuse gingivitis and periodontitis. Four premolars had been extracted for orthodontic purposes. This patient had 31 pockets ranging from 6 to 10 mm in depth, but only in four of these sites was attachment loss greater than 5 mm. B. asaccharolyticus averaged 46% of the cultivable flora, and the spirochetes accounted for 28% of the microscopic count (Table 4). An occasional amoeba was seen by dark-field microscopy. The patient was started on metronidazole and had the teeth...
Table 5. Effect of 1 week of metronidazole treatment and scaling on clinical and bacteriological parameters. Patient R, age 23; Early periodontitis

<table>
<thead>
<tr>
<th>Clinical (Values given in mm)</th>
<th>Pre-treatment</th>
<th>0</th>
<th>5</th>
<th>17</th>
<th>21</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pocket reduction (72)*</td>
<td>61 ± 9</td>
<td>2.4 ± 1.6</td>
<td>6.3 ± 4</td>
<td>13 ± 13</td>
<td>19 ± 10</td>
</tr>
<tr>
<td>Gain in attachment (7)*</td>
<td>151 ± 80</td>
<td>0.4 ± 0.3</td>
<td>1.5 ± 1.2</td>
<td>0.2 ± 0.2</td>
<td>7.4 ± 6.4</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Bacteriological</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>% B. asaccharolyticus</td>
<td>61 ± 9</td>
<td>2.4 ± 1.6</td>
<td>6.3 ± 4</td>
<td>13 ± 13</td>
<td>19 ± 10</td>
</tr>
<tr>
<td>× 10*</td>
<td>151 ± 80</td>
<td>0.4 ± 0.3</td>
<td>1.5 ± 1.2</td>
<td>0.2 ± 0.2</td>
<td>7.4 ± 6.4</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Microscopic</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>% Spirochetes</td>
<td>5.6 ± 2</td>
<td>1.4 ± 0.9</td>
<td>15 ± 11</td>
<td>1 ± 0.6</td>
<td>4.8 ± 3</td>
</tr>
<tr>
<td>Spirochetes/10 hpf*</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* Number in parentheses is number of sites which initially exhibited 3-5 mm pockets or 3-5 mm attachment loss
b Average ± standard error
c Values are significantly different from pretreatment value.
d Percentage of total cultivable count
* hpf = high power fields

on the right side scaled and root-planed. One week later the teeth on the left side were similarly treated. B. asaccharolyticus and the various spirochetes were essentially undetectable during the following 30-week period (Table 4). The 31 pockets that initially were greater than 5 mm were reduced by about 3 mm at week 30. In the four sites that initially had greater than 5 mm attachment loss, there was an apparent gain in attachment of 2-3 mm (Table 4). No mechanical treatment was given during this 30-week interval, nor did the patient receive antibiotics for medical purposes during this time.

Patient R was a 23-year-old white male who presented with gingivitis and a diffuse periodontitis that included 72 sites that had 3-5 mm pockets, and two sites with 6 mm pockets. When the cultural studies showed that B. asaccharolyticus averaged 61% of the cultivable flora (Table 5), a diagnosis of active periodontitis was made. The patient was placed on metronidazole and the teeth were mechanically scaled in the same manner as described for patient E. The B. asaccharolyticus levels were reduced by over 90% and its proportions by over 60%, 21 weeks after treatment (Table 5). There was a 0.7 mm pocket reduction and a 0.7 mm apparent gain in attachment, 17 weeks after treatment in the sites that initially exhibited 3-5 mm pockets or attachment loss (Table 5).

The mean values for the papillary bleeding score (PBS) in the time after metronidazole treatment for all patients are shown in Table 6. The 1-week metronidazole treatment significantly reduced the PBS by about 50-75% during the period of observation. Note that in the two older patients
Table 6. Mean change in papillary bleeding score following 1 week of metronidazole treatment

<table>
<thead>
<tr>
<th>Patient</th>
<th>Treatment</th>
<th>Time after metronidazole treatment in weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>0</td>
</tr>
<tr>
<td>Patient D</td>
<td>22</td>
<td>10</td>
</tr>
<tr>
<td>Patient L</td>
<td>50</td>
<td>29</td>
</tr>
<tr>
<td>Patient H</td>
<td>22</td>
<td>7</td>
</tr>
<tr>
<td>Patient E</td>
<td>58</td>
<td>26</td>
</tr>
<tr>
<td>Patient R</td>
<td>38</td>
<td>21</td>
</tr>
<tr>
<td>Average/patient</td>
<td>38 ± 7</td>
<td>19 ± 4</td>
</tr>
</tbody>
</table>

* Teeth were scaled and root planed during weeks 22-25
b Teeth were scaled and root planed while on metronidazole
c Pretreatment value significantly higher than all succeeding posttreatment values, paired t-test $P < 0.05$, with the exception of the 4-10 week interval

(D and H), whose periodontitis included furcation involvement, the initial PBS was relatively low. In these patients the infection was confined mostly to the apical areas of the pockets.

The profile of changes in the various

Table 7. Effect of 1 week of metronidazole treatment on various clinical and bacteriological parameters ($\bar{x} \pm$ standard error)

<table>
<thead>
<tr>
<th>Pretreatment</th>
<th>Time after metronidazole in weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
</tr>
<tr>
<td>% B. asaccharolyticus</td>
<td>$41 \pm 11^a$</td>
</tr>
<tr>
<td>% spirochetes</td>
<td>$29 \pm 5^a$</td>
</tr>
<tr>
<td>Papillary bleeding score</td>
<td>$38 \pm 7^a$</td>
</tr>
<tr>
<td>Reduction in pocket depths in mm</td>
<td></td>
</tr>
<tr>
<td>$\bar{x}$/patient (4)</td>
<td>2.2</td>
</tr>
<tr>
<td>$\bar{x}$/pocket (105)</td>
<td>1.9</td>
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<tr>
<td>Gain in apparent attachment</td>
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</tr>
<tr>
<td>$\bar{x}$/patient (4)</td>
<td>2.1</td>
</tr>
<tr>
<td>$\bar{x}$/pocket (51)</td>
<td>1.6</td>
</tr>
</tbody>
</table>

* Pretreatment value significantly higher than succeeding values $P < 0.05$, paired t-test
b Number of patients
c Number of pockets
clinical and bacteriological parameters for the five patients is summarized in Table 7. Although the two patients who received the metronidazole during the period of scaling and root planing showed the greatest reduction in the PBS (Table 6), and responded optimally in regards to the other parameters, the numbers involved were too few to perform tests of significance between this group and those who received only metronidazole.

As all five patients showed a similar improvement in the various parameters (Tables 1-5), and as the mechanical treatment was minimal in the context of the patients' total treatment needs, the data for the five patients were grouped together and compared over time. The 1-week treatment with metronidazole with or without scaling significantly reduced the proportions of *B. asaccharolyticus* and spirochetes and reduced the PBS for at least 6 months after treatment (Table 7). The pockets that were initially greater than 5 mm were reduced by about 2.5 mm. This was true if the average reduction was calculated per patient, or per pocket (Table 7). The apparent gain in attachment in those sites which initially exhibited greater than 5 mm attachment loss was about 1.8 mm when calculated per patient and 1.4 mm when calculated per pocket.

**Discussion**

The patients in the present study presented with clinical disease coincident with elevated proportions and levels of *B. asaccharolyticus* and spirochetes in plaques removed from periodontal pockets. *B. asaccharolyticus* accounted for 41% of the cultivable isolates, and the spirochetes averaged 29% of the microscopic count (Table 7). These proportions are high compared to what has been reported for plaques from periodontal patients (Socransky et al. 1962, Loesche et al. 1972), but are consistent with values found when active disease is suspected (Slots 1977, Listgarten & Helldén 1978). These bacteriological findings combined with the clinical appearance of the periodontium suggested that the patients, or at least the sampled pockets, were in the state of an active infectious process involving primarily anaerobic bacteria. If this is the case, antimicrobial therapy directed against these anaerobes was indicated according to the specific plaque hypothesis (Loesche 1976a). In recent years, several antimicrobials have been recommended for treatment of anaerobic infections (Finegold 1977), but only one, metronidazole, has been used in the treatment of oral infections (Shinn 1977, Ingham et al. 1977). Metronidazole interacts with low-redox-potential electron transport proteins of the type normally found in anaerobes to form a derivative which exerts a killing action in the cell (Müller et al. 1977). Microaerophilic organisms such as *A. viscosus*, and the *Capnocytophaga* and *Eikenella* species that exhibit periodontal pathology in animal models (Socransky 1977) would be expected to be, or are resistant to this agent. Thus, the usage of metronidazole in these patients provided a probe that would selectively discriminate against the anaerobic species while leaving the various facultative and microaerophilic species unaffected. In this manner the relative contribution of the anaerobes to the observed periodontal pathology could be gauged. The efficacy of metronidazole against the plaque anaerobes in the present patients was assessed by following the proportions of *B. asaccharolyticus* and spirochetes in the plaques before and after treatment. These organisms were chosen as indicator organisms because both can be easily identified in plaque samples, both were present in high numbers, and both are suspected of being periodontopathic. Metronidazole should be without an appreciable effect in those instances in which
the microaerophilic organisms are primarily responsible for the concurrent periodontal pathology.

The 1-week treatment with metronidazole caused an obvious improvement in the periodontal health of the most diseased sites in each of these five patients. This was manifested by the greater than 50% reduction in the PBS (Table 6), a 2 mm or more reduction in pocket depth (Table 7), and an almost 2 mm gain in apparent attachment (Table 7) that were evident 6 months after treatment. These deep pockets would be expected to show the greatest improvement following a therapeutic regimen (Knowles et al. 1979). The magnitude of the improvement is surprising, considering that no efforts were made to instruct the patients in oral hygiene procedures and a minimal amount of mechanical debridement was used. This latter consisted of scaling and root planing in patients E and R during the week they were on metronidazole and scaling and root planing the maxillary teeth of patient H, 22 to 25 weeks after she received metronidazole. These mechanical procedures contributed to the improvement of the periodontal condition, as patient E exhibited the best response to treatment of all the patients, and patient H showed reductions in the PBS and pocket depth as well as an attachment gain in the interval following scaling and root planing.

The persistent decline in numbers and proportions of *B. asaccharolyticus* and spirochetes coincident with the sustained improvement in periodontal health argues that these bacterial types were contributing to the observed periodontal pathology. This is consistent with the long-known association of spirochetes (Loesche 1976b) and *B. melaninogenicus* (MacDonald et al. 1962) with periodontal disease. However, this finding is not synonymous with saying that these organisms initiated the disease or are the only organisms contributing to the disease. Clearly in a complex anaerobic flora of the type found in periodontal pockets, other organisms or microbial interactions not detected by the present methodology could be of etiologic significance. The results suggest that the microaerophilic and facultative species present in these plaques were not primarily responsible for signs of disease observed.

Other investigators have monitored the clinical and bacteriological response to systemic tetracycline combined with various scaling and root-planing schedules (Listgarten et al. 1979, Heildén et al. 1979, Slots et al. 1979), or to locally administered tetracycline (Lindhe et al. 1979). The magnitude of clinical improvement attained appears to be that which can be achieved by repeated scaling (Heildén et al. 1979, Slots et al. 1979). The degree of improvement observed at 6 months in the five patients given metronidazole, especially as it relates to apparent attachment gain, is beyond that observed with tetracycline. The apparent superiority of metronidazole relative to tetracycline probably resides in metronidazole’s unique spectrum of activity against anaerobes.

Some mention should be made concerning the significance of the apparent gain in attachment. In the absence of evidence that bone had regenerated and/or that anatomical reattachment had occurred, this measurement can only imply an apparent gain in attachment. It most likely reflected that the inflammation in the apical portion of the pocket had been reduced to the extent that the epithelium formed a tight cuff about the root surface. Accordingly, the periodontal probe that was used in a standard manner (Ramfjord 1967) could not penetrate as deeply in the pocket after treatment as before treatment (Armitage et al. 1977, Listgarten et al. 1976). The measurement thus gives some indication of tissue response at the apical extent of the
pocket. The observation that metronidazole caused an apparent gain in attachment of about 1.5 mm in the sites with the most advanced signs of periodontal disease is encouraging.

The treatment and/or management of periodontal disease has, as its fundamental principle, the control of the bacterial flora in the periodontal pocket. Surgical procedures, if supplemented with periodic scaling and root planing, will effectively control this flora and restore periodontal health (Rosling et al. 1976a, b; Knowles et al. 1979). However, if these surgical procedures are not supplemented with rigorous oral hygiene and maintenance therapy, periodontal health will relapse and actually result in attachment loss which can be detected as early as 6 months after surgery (Nyman et al. 1978). The requirement for maintenance therapy can be explained by the regrowth of a periodontopathic flora that needs to be periodically disrupted. The present results indicate that when this periodontopathic flora comprises anaerobes, it may be effectively reduced by a systemic antimicrobial agent. If this is so, then metronidazole or other agents with a similar spectrum of activity, when incorporated into a regular periodontal treatment schedule, may enhance and/or prolong the periodontal health seen at recall visits. With this in mind, a double-blind clinical trial of metronidazole or placebo in conjunction with a periodontal treatment schedule that would include root planing, curettage, and surgery has been initiated. The metronidazole, or placebo, is administered during the hygienic phase of therapy, approximately 6–8 weeks prior to the initiation of surgery in patients with advanced periodontal disease.

If the current documentation that certain forms of periodontal disease represent specific plaque infections can be sustained, then some form of antimicrobial therapy with chemical agents will be an inevitable future treatment possibility. The choice of drugs available in the United States will probably be limited to those which already have FDA approval. Currently, neither metronidazole nor tetracyclines have FDA approval for usage in periodontal infections, simply because this usage was never described in the original drug application. This does not prevent the prescription usage of these drugs in dentistry. The practitioner should thus be aware of the known risks of each drug. Tetracyclines cannot be tested in the standard bacterial mutagenicity assay because the indicator organisms are sensitive to it. However, the high percentage of tetracycline-resistant organisms found in plaques following tetracycline therapy (Hawley et al. 1980, Williams et al. 1979) raise the possibility that some of these resistant organisms could arise by mutation. Metronidazole is mutagenic in the Ames test (Rosenkranz & Speck 1977). Lifetime ingestion of very high dosages of metronidazole, ie. 0.3 and 0.5 % of the diet, increased the incidence of malignant lymphoma in female mice (Rustia & Shubik 1972). The dosages used were about 3500 times the single course of treatment used in the present patients (Roe 1977). This study was complicated by the fact that the high-dose animals received oxytetracycline in the drinking water during two periods, each of 2 weeks' duration (Roe 1977). Subsequent mouse, rat and hamster studies have not shown metronidazole to be carcinogenic (Roe 1977), and to be "probably no more than a weak tumoHgen in mice after prolonged high-level exposure" (Rust 1977). A retrospective human study could find no evidence of carcinogenicity in female patients who had been prescribed metronidazole for the treatment of trichomonal vaginitis (Beard et al. 1979). This study encompassed over 8200 patient-years and included for some patients, 10-year follow-up data. These
animal and human studies indicate that metronidazole should be used discriminately. Metronidazole’s unique spectrum of activity against *B. asaccharolyticus*, spirochetes and other anaerobes involved in periodontal disease, coupled with the favorable clinical response to short-term treatment, indicate that this agent will have increased usage in dentistry. Efforts to find antimicrobials with a similar anaerobic spectrum such as nitrimidazine (Lozdan et al. 1971), but minus the mutagenic potential, should be pursued (Rosenkranz & Speck 1977). Until that time metronidazole should be reserved for those instances in which the anaerobic nature of the infection is documented.

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Zusammenfassung
Parodontale Infektionen durch anaerobe Bakterien – Behandlungsresultate mit Kurzzeitgaben von Metronidazol
Der vorliegende Bericht beschreibt 5 ausgewählte parodontalerkrankte Patienten, die eine Woche lang mit Metronidazol behandelt wurden. Bei 2 dieser Patienten wurde in der Woche, zusammen mit Metronidazolgaben, der Zahnstein entfernt und Wurzelplanung vorgenommen. Vor der Behandlung machte der Anteil des *B. asaccharolyticus* 41% der kultivierten Isolate aus, und der Anteil der Spirochäten betrug etwa 29% bei mikroskopischer Zählung der Mikroorganismen in der Plaque einer jeden der 4 Zahnfleischtaschen, die bei jedem Patienten untersucht wurden. Die erhöhten Anteile an der parodontopathischen Bakterienflora, bei gleichzeitigem Vorhandensein parodontaler Taschen und dem Vorkommen von Attachmentverlust, liessen die Folgerung zu, dass diese Patienten sich im aktiven Stadium eines infektiösen Prozesses bei gleichzeitiger Anwesenheit primär anaerober Bakterien befanden. War diese Folgerung richtig, war antimikrobielle Therapie mit Metronidazol gegen diese Anaerobier angezeigt. Die Behandlung mit Metronidazol während einer Woche reduzierte den proportionalen Anteil dieser Organismen bis zu 6 Monaten nach der Behandlung. Gleichzeitig damit wurde eine Verbesserung klinischer Parameter gesehen, und das vor allem in den Regionen mit Zahnfleischtaschen, die tiefer als 5 mm waren und bei denen Attachmentverlust vorlag. In diesen Regionen verringerten sich die Taschentiefen um 2 mm oder mehr und 6 Monate nach der Behandlung war ein Gewinn an Attachment von 2 mm offenbar. Die hier erhaltenen Resultate wurden bei nur 5 Personen erreicht. Das Ausmass der Verbesserung lässt jedoch vermuten, dass antimikrobielle Therapie, gezielt auf anaerobe Organismen eingestellt, ein wertvolles Adjuvans zu parodontaler Therapie bedeuten kann.

Résumé
Le traitement à court terme au moyen du métronidazole dans les cas d’infections parodontales dues à des bactéries anaérobies. Compte-rendu de cinq cas
Le présent travail rend compte de cinq cas de parodontopathies que l’on a choisis et traités pendant une semaine au métronidazole. Chez deux des patients, les dents ont subi un détartrage et un polissage des surfaces radiculaires pendant la semaine où ils recevaient le traitement au métronidazole. Avant le traitement, *B. asaccharolyticus* représentait 41% des colonies isolées pouvant être cultivées et les spirochètes se montraient en moyenne à 29% dans les numérations microscopiques faites sur la plaque prélevée dans quatre culs-de-sac par patient. La présence en proportions élevées de ces bactéries parodontopathiques, associée à la présence de culs-de-sac parodontaux et de perte de l’attachement, semble indiquer que les patients étaient sujets à un processus infectieux actif où participaient principalement des bactéries anaérobies. En supposant que ce soit le cas, un traitement antibactérien au métronidazole, dirigé contre ces anaérobies, était indiqué. Le traitement au métronidazole institué pendant une semaine a réduit significativement les proportions de ces organismes pendant une période allant jusqu’à six mois après le traitement. En concordance avec ces résultats, il se produisait une amélioration des paramètres cliniques, surtout au niveau des localisations.
où les culs-de-sac ou la perte de l'attache ment étaient initialement de plus de 5 mm. Ces localisations présentaient une réduction de la profondeur des culs-de-sac de 2 mm ou plus et une amélioration de presque 2 mm de l'attache ment apparent, visible six mois après le traitement. Les résultats obtenus ont été constatés chez cinq patients seulement. Cependant, l'ampleur de l'amélioration semble indiquer qu'un traitement antimicrobien dirigé contre les organismes anaérobies peut représenter une aide de grande valeur pour le traitement parodontal.

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