Successful treatment of genital ulcers with infliximab in Behçet’s disease

G Haugeberg, M Velken, V Johnsen

LETTERS

Recurrent oral and genital aphthous ulcerations are the hallmarks of Behçet’s disease. Up to now the various clinical manifestations in Behçet’s disease have been treated with limited success. Recently, clinical observations have reported dramatic responses on clinical signs and symptoms in patients treated with tumour necrosis factor α (TNFα) blocking drugs, including severe mucocutaneous, gastrointestinal and ocular manifestations.

We report the successful treatment with infliximab of a severe longstanding genital ulcer in a patient with Behçet’s disease.

CASE REPORT

The patient was a 29 year old woman with Behçet’s disease who had had recurrent oral ulcers from age 5 and genital ulcers, arthralgia, and uveitis from age 23. The outbreaks of uveitis were treated with local administration of dexamethasone only. From autumn 2000 the severity of the genital manifestations increased, and treatment with colchicine and prednisolone was started. However, during follow up she still had painful recurrent oral and genital ulcers. In December 2001 she had a new outbreak of a severe genital ulcer located at the left labium majus. No sign of spontaneous improvement was seen, and high dose prednisolone treatment (60 mg a day) was started in spring 2002. In summer, partial remission on oral corticosteroid treatment had been achieved, but the painful ulcer was still present. During the following months the genital ulcer deteriorated despite high dose prednisolone treatment (initially 60 mg, which had to be reduced to 40 mg because of mental symptoms). In December 2002 treatment with azathioprine, 50 mg twice a day, was started in addition to prednisolone 30 mg daily. Treatment with azathioprine was stopped after 2 weeks because of a respiratory infection. Although non-selective immunosuppressive drugs—for example, thalidomide, azathioprine, and cyclosporin A, had not been tested sufficiently as monotherapy or in combination, we decided to treat her with infliximab, after full discussion and consent. This decision was judged justified owing to, firstly, the severe longstanding physical and psychological burden for the patient, secondly, to reports on successful treatment of genital ulcers with infliximab in Behçet’s disease; and, thirdly, to lack of response to high dose long term prednisolone treatment. The infliximab dose 5 mg/kg body weight infusion was given on four occasions at baseline, and after 2, 6, and 15 weeks. The fourth infusion, planned at 12 weeks’ follow up, was postponed owing to sinusitis treated with penicillin. No other possible infliximab side effects occurred. During treatment her general well being and fatigue improved.

At the time of the first infusion the ulcer was deep and had a diameter of 19 mm (fig 1A). After one infusion, at 2 weeks’ follow up, a marked improvement was seen (fig 1B), and at 6 weeks’ follow up, 4 weeks after the second infusion, only a scar of the ulcer was left (fig 1C). At 2 months’ follow up after the fourth infusion the patient was still in remission.

DISCUSSION

Our case report supports the results of others1–3 and shows that treatment with infliximab is effective in inducing remission of genital ulcers in patients with Behçet’s disease.

Recently, preliminary results from a double blind, placebo controlled study were presented on the effect of etanercept, another TNF blocking drug, on mucocutaneous manifestations in Behçet’s disease.7 In this 4 week study eight (40%) of the 20 etanercept treated patients achieved clinical remission of mucocutaneous manifestations compared with one (5%) patient in the 20 placebo treated patients.7 Interestingly, in the case by Estrach et al.,3 the orogenital ulcerations in a patient with Behçet’s disease responded dramatically to treatment with infliximab after failure of etanercept. This raises the interesting question as to whether there is a difference in the efficacy in Behçet’s disease between etanercept (soluble receptor binding TNFα and -β) and...
infliximab (monoclonal antibody binding TNFα), drugs known to bind TNFα in different ways.

The experience so far with TNFα blocking treatment in Behçet’s disease is scarce as recently reviewed by Sfikakis. So far, significant side effects with the use of TNFα blocking drugs in Behçet’s disease have not been reported.

Further clinical studies, especially double blind, randomised controlled trials designed with sufficient power are warranted before final conclusions can be drawn about the efficacy of treatment with TNFα blocking agents compared with conventional immunosuppressive agents.

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Lupus myocarditis in children

CARDIOVASCULAR SYSTEM (CVS) disease is the third leading cause of mortality in systemic lupus erythematosus (SLE). Pathologically it is pancarditis and may affect coronary arteries. Most often, it is subclinical. Clinical manifestations, if present, are limited to pericardial and endocardial involvement. Clinically apparent myocarditis has been described in a few adult patients. To the best of our knowledge, no such case has been described in children. We describe two children with clinically manifest myocarditis in SLE.

CASE REPORTS
A 12 year old girl, who was known to have had SLE for 2 months, presented with fever, malar rash, and photosensitivity. She had stopped taking steroids for 2 weeks. On examination, she had tachycardia with normal blood pressure. She had pallor, alopecia, oral ulcers, and malar rash. An examination of the CVS showed gallop rhythm with no murmurs. The rest of the systemic examination was unremarkable. She had stopped steroids on her own. On examination, she was shown to have generalised oedema, malar rash, and oral ulcers. She had tachycardia with normal blood pressure. Examination of the CVS showed a soft systolic murmur at the apex with normal heart sounds. She had hepatosplenomegaly. The rest of the systemic examination was unremarkable. Investigations showed normocytic, normochromic anaemia (haemoglobin 54 g/l) and deranged renal functions (blood urea nitrogen 26.5 mmol/l, serum creatinine 220 μmol/l). There was gross albuminuria, with urine microscopy showing red blood cells and casts. Chest radiography showed cardiomegaly with normal lung fields. Echocardiography showed moderate mitral regurgitation, trivial tricuspid regurgitation, and global hypokinesia with an LVEF of 18%. No pericardial effusion or vegetations were present. ANF was strongly positive and anti-dsDNA titres were 1531 IU/ml. Treatment was started with pulse methylprednisolone. However, she developed cardiogenic shock on day 6 of admission. Pulse cyclophosphamide was started together with supportive treatment. She developed multiorgan dysfunction due to shock and died after 8 days in hospital. A necropsy showed diffuse proliferative glomerulonephritis, serositis, and myocarditis. Myocarditis was evident from the presence of focal collection of inflammatory cells with damage to adjacent myofibres, fibronoid necrosis, and interstitial oedema. Fibrinous pericarditis was also seen, but the coronary arteries and valves were normal.

DISCUSSION
Myocardial dysfunction in SLE is multifactorial; important contributors are immune injury, coronary vasculitis, valvulopathy, hypertension, and adverse effects of drugs. Endomyocardial biopsy remains the preferred method for the diagnosis of myocarditis, but carries a high risk of mortality in sick children.

REFERENCES
and was not carried out in our patients. In such a case, the presence of global hypokinesia with a low LVEF on echocardiography is a strong pointer towards the diagnosis. The diagnosis is even more definite in the second child as it was proved at necropsy. Lupus myocarditis has been associated with skeletal myositis and anti-Ro and anti-La antibodies.

Immunosuppressant drugs with decongestive agents remain the preferred treatment. Most patients respond dramatically. The second patient died despite immunosuppressive treatment and supportive care, probably because of delays in diagnosis and initiation of treatment. Finally, it is important to remember that acute myocarditis can be a fatal complication of SLE. Early diagnosis and prompt treatment may be very rewarding.

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Dose escalation of leflunomide (LEF) to 40 mg once daily in patients with rheumatoid arthritis and insufficient response to standard dose LEF

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Leflunomide (LEF) is a new disease modifying antirheumatic drug (DMARD) for the treatment of rheumatoid arthritis (RA) and it is recommended that it is given in a loading dose of 100 mg over 3 days followed by a stable dose of 20 mg/day. The dose of 20 mg/day LEF was recommended because in an initial study which compared 5, 10, and 25 mg/day LEF the highest dose tended to result in an increased rate of adverse events. However, in this study a clear dose-response relationship with best results in the 25 mg/day group was observed as well. Therefore, the aim of this investigation was to evaluate whether patients who tolerate LEF in the standard dose of 20 mg/day will tolerate a dose escalation of LEF to 40 mg/day and whether this improves the efficacy of the treatment.

PATIENTS AND METHODS

Eleven patients with RA were included in this prospective clinical investigation (table 1). All had received LEF in a dose of 20 mg/day for at least 3 months without adverse effects, but still had active disease with a mean (SD) disease activity score (DAS28) of 4.9 (0.6) (range 3.9–5.8). All of them had responded to the prior treatment with 20 mg/day LEF, but either the response was not sufficient or the disease activity increased again despite continuous treatment.

To obtain better control of the disease the dose of LEF was increased from 20 to 40 mg p.o. once daily. The patients were monitored by analysing the disease activity and the occurrence of adverse events after 2 months of treatment and after every subsequent 2 months for as long as the patient was receiving 40 mg LEF.

RESULTS

Table 1 summarises the data obtained. After a mean (SD) follow up of 4.4 (3.7) months (range 0.5–11) 5 of 11 patients were still receiving treatment with 40 mg/day LEF. Four patients had stopped treatment with LEF or reduced the dose to 20 mg/day again because of adverse events: one patient with concomitant MTX treatment developed increased liver enzymes more than twice the normal values and stopped LEF after 4 months because of glossitis, another patient after 7 months because of abdominal pain of unknown origin. Both adverse events were reversible after dose reduction of LEF. Six of 9 patients who were still receiving 40 mg/day LEF after 2 months responded to the dose escalation of LEF: two had a good response (reduction of DAS28 >1.2 and inactive disease) and four had a moderate response (reduction of DAS28 from 0.6 to 1.2 or active disease despite reduction of DAS28 >1.2) as defined by van Gestel et al 1996. Two patients showed no change in disease activity. In all five patients who continued to receive LEF in a dose of 40 mg/day, the response remained stable up to the last follow up.

CONCLUSION

In these preliminary findings the dose escalation of LEF to 40 mg/day seems to increase the effectiveness of the treatment in a number of patients with RA. The adverse events of the dose escalation which appeared in four of 11 patients were

REFERENCES

Polyclonal immunoglobulins for intravenous use induce interleukin 10 release in vivo and in vitro

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Polyclonal intravenous immunoglobulins (IVIg) are increasingly used in clinical practice, not only as substitutive agents but also in the treatment of immunologically mediated diseases. How IVIg modulate the immune system is not yet clear, but several immunoregulatory mechanisms have been postulated. We studied the production of the immunosuppressive cytokine interleukin 10 (IL10) as a potential explanation for the beneficiary effects of IVIg in autoimmune diseases.

IL10 concentrations were measured in six patients: one patient (aged 19) with X-linked agammaglobulinemia, two patients (aged 63 and 64) with common variable immunodeficiency, one patient (aged 15) with hyper IgM syndrome, two patients (aged 4) with IgG2 subclass deficiency. All patients or their parents gave informed consent. Blood was collected before and immediately after infusion of IVIg (Sandoglobulin; Novartis, Brussels, Belgium), and IL10 levels were measured with a sandwich enzyme linked immunosorbent assay (ELISA) technique (anti-IL10 Screening Line; Pharmingen, San Diego, CA). The serum IL10 levels were increased after infusion of IVIg in all six patients (Wilcoxon non-parametric test, p<0.01; fig 1A). No IL10 was detected in the IVIg.

Venous blood from healthy volunteers was collected and diluted 1:2 with culture medium. Final cell concentration was 1×10^6 white blood cells/ml. Cell culture was performed with or without endotoxin (1 ng/ml) or IVIg (3 mg/ml) for 15 minutes to 24 hours. An increase in IL10 levels as early as 1 hour after incubation with IVIg was consistently seen in all experiments performed (fig 1B). IL10 levels reached their mild and reversible. If confirmed in larger studies, the dose escalation of LEF might be especially valuable for patients with RA who tolerate LEF at the standard dose of 20 mg/day but show insufficient response. For them the dose escalation might be a way to further improve disease activity and an alternative to an early switch to DMARD combination treatment or treatment with biological substances.

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highest level after 4 hours and remained high over the next 4 days. The rapid increase in IL10 levels is remarkable because endotoxin induces IL10 only after 8 hours and induces levels comparable to IVIg only after 24 hours. Increases in IL10 were seen in all our experiments but varied in intensity between different donors.

We also studied the effect of IVIg on IL10 mRNA levels in peripheral blood mononuclear cells (PBMC) from healthy volunteers. 1 x 10^6 PBMC were stimulated with IVIg (3 mg/ml) in culture medium for 2 or 24 hours and IL10 mRNA content was determined using Taqman real time, quantitative, reverse transcriptase-polymerase chain reaction (Applied Biosystems, Lennik, Belgium; IL10 forward primer GTGATGCCCAAGCTGAGA, IL10 reverse primer CAGGGCCTTGCTCTGTTT, IL10 probe CCAAGACCCAGACA-TCAAGCGCA). Gene expression levels were normalised to the expression of the reference gene β-actin (ABI Prism Taqman Assay Reagents, Applied Biosystems). IVIg stimulation resulted in a mean 9.6-fold increase in IL10 mRNA levels after 24 hours, but not after 2 hours (data not shown). A rapid increase of IL10 protein levels in vivo has previously been reported after liver transplantation. Our data further corroborate the hypothesis that IL10 production is regulated by both transcriptional and post-transcriptional mechanisms.

In blocking experiments, cells were preincubated for 30 minutes in the presence of different anti-Fc receptors antibodies (anti-FcγRI (197), anti-FcγRIII (IV.3), and anti-FcγRIII (3G8), all from Medarex (Milpitas, CA)) at 10 ng/ml at 4°C. IL10 production was effectively inhibited by monoclonal antibodies (mAb) directed against the FcγRI and FcγRIII. These mAb did not stimulate IL10 production in controls. Anti-FcγRI was responsible for the strongest inhibition, followed by anti-FcγRIII. The combination of these mAb was more effective in inhibiting IL10 production than either of the mAbs alone, showing that triggering of both receptors increases IL10 release independently. Remarkably, recent data suggest that immune complexes or sera from patients with systemic lupus erythematosus (SLE) trigger IL10 production from PBMC mainly through FcRII. IL10 is a potent anti-inflammatory cytokine regulating the production of monokines such as tumour necrosis factor α and IL1 as well as shifting T cell responses towards a Th2 profile. Therefore, increased IL10 production by IVIg may provide an explanation for some of the remarkable immune suppressive properties in inflammatory and autoimmune diseases such as dermatomyositis.

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REFERENCES
Cevimeline gargle for the treatment of xerostomia in patients with Sjögren’s syndrome

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Irrigation of the salivary gland using a corticosteroid is one of the most successful approaches to improving xerostomia in patients with Sjögren’s syndrome, yielding longlasting effects associated with minimal side effects. However, the treatment procedures are complicated, and the extent of improvement in salivary function is inversely related to the clinical severity of the disease, with the patients with most advanced disease not responding to the treatment.

Recently, cevimeline hydrochloride hydrate (Evoxac or Saligren) has been clinically applied to patients with Sjögren’s syndrome for the treatment of xerostomia. Oral doses of cevimeline significantly improved subjective symptoms of dry mouth and dry eyes, and increased salivary flow. However, adverse effects were frequently reported, such as nausea and abdominal pain. Unless properly managed, these adverse effects would negatively affect the continued administration of cevimeline.

To overcome these disadvantages of oral doses of cevimeline, we evaluated the efficacy of cevimeline gargle for the treatment of xerostomia in patients with Sjögren’s syndrome. We first evaluated the effect of the gargle in 11 healthy female volunteers (mean (SD) age 39 (13) years, range 19–57), after approval from the ethical committee of our hospital. The volunteers were asked to gargle three times a day before meals using 30 mg cevimeline dissolved in 100 ml of water for each session. The effects of treatment were evaluated daily by a Saxon test performed at around 3 00 pm. Most of the healthy subjects responded well to the cevimeline gargle, and the salivary flow rate gradually increased, reaching an 81% increase on average by day 5. Some subjects showed more than twofold increases in salivary flow after cevimeline gargle. No adverse effect was seen. Simple gargles without cevimeline had no effect.

Given the satisfactory efficacy and safety of the cevimeline gargle in healthy subjects, we next tested whether the same treatment was effective in patients with Sjögren’s syndrome. These patients fulfilled the criteria proposed by Fox et al. and were also refractory to corticosteroid irrigation (table 1). Cevimeline gargle markedly increased salivary flow rates in two of the five patients. However, in the remaining three patients the effect of treatment was negligible. In three of these five patients clinical symptoms improved subjectively. No adverse effects were seen.

The exact mechanism of salivary flow stimulation by cevimeline gargle is not clear. Possibly, cevimeline binds directly to the muscarinic receptors of the minor salivary gland acini that communicate with the oral cavity. The observed inconsistency in treatment effects in the patients with Sjögren’s syndrome may be due to severely damaged salivary acini. Further studies are necessary to evaluate cevimeline gargle treatment in a large cohort including more mildly affected patients.

Effect of cevimeline gargle in patients with Sjögren’s syndrome

Table 1

<table>
<thead>
<tr>
<th>Patients</th>
<th>Age</th>
<th>Sex</th>
<th>Corticosteroid irrigation</th>
<th>Improvement in dry mouth</th>
<th>After cevimeline gargle</th>
<th>Saxton test (g/2 min)</th>
</tr>
</thead>
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<tr>
<td>1</td>
<td>73</td>
<td>F</td>
<td>0.33</td>
<td>+</td>
<td>0.45 (181)</td>
<td>0.16</td>
</tr>
<tr>
<td>2</td>
<td>61</td>
<td>F</td>
<td>0.58</td>
<td>+</td>
<td>0.45 (275)</td>
<td>0.12</td>
</tr>
<tr>
<td>3</td>
<td>71</td>
<td>F</td>
<td>1.80</td>
<td>++</td>
<td>1.81 (13)</td>
<td>1.60</td>
</tr>
<tr>
<td>4</td>
<td>58</td>
<td>F</td>
<td>0.40</td>
<td>–</td>
<td>0.06 (–40)</td>
<td>0.10</td>
</tr>
<tr>
<td>5</td>
<td>38</td>
<td>F</td>
<td>0.62</td>
<td>–</td>
<td>0.17 (21)</td>
<td>0.14</td>
</tr>
</tbody>
</table>

*These patients with Sjögren’s syndrome were refractory to corticosteroid irrigation; †corticosteroid irrigation was performed as described previously; ‡judged by verbal inquiry after cevimeline gargle.

References

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