Photodynamic Therapy of Necrobiosis Lipoidica – A Multicenter Study of 18 Patients

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**Introduction**

Necrobiosis lipoidica (NL) is a rare granulomatous skin disease of unknown origin, first described by Oppenheimer in 1929 in association with diabetes mellitus. In 1935, Goldsmith reported the first case in a nondiabetic patient [1]. Clinically, NL appears as irregular or circular lesions ranging in color from red brown, dark purple to yellow. Ulcerations and telangiectasia may occur [2]. Most commonly the anterior portions of the lower legs are involved, but other localizations have been described [3].

Since the lesions of NL are not only aesthetically troublesome, but may also be accompanied by pain or secondarily infected ulcerations, treatment of this disease is necessary. Several therapeutic approaches have been described for NL, including topical, intrasional and systemic application of corticosteroids, topical retinoids, PUVA, cyclosporine and chloroquine [4–9]. However, these treatment modalities are of limited success, and new approaches are wanted to handle this often frustrating and cosmetically disturbing disease.

In 2006, Heidenheim and Jemec [10] successfully treated a diabetic patient with a 10-year history of NL by topical photodynamic therapy (PDT). They observed a...
complete remission of the NL lesion after 6 cycles of PDT using methyl aminolevulinate (MAL, 160 mg/g) and red light. This single case report published so far motivated us to perform a study in a series of patients in order to evaluate the overall efficacy of this therapeutic modality to control this often therapy-resistant and uncontrolled disease.

Patients and Methods

This study was conducted in 3 European university departments of dermatology: Roskilde Hospital (Roskilde, Denmark), Ludwig Maximilian University (Munich, Germany) and the Faculty Hospital Královské Vinohrady (Prague, Czech Republic). Eighteen patients (14 female, 4 male) with NL of the lower legs and resistant to conventional therapies were included (table 1). The average age of the patients was 38 years with a range of 16–62 years. Comorbidity of diabetes mellitus was present in 28% of the patients (5/18). The duration of NL was 1–8 years in 16 patients and 19 and 26 years in 2, respectively. In 4 patients, lesions were partially ulcerated. A diagnostic biopsy was performed in 12 patients, while in 6 there was no histological confirmation of the disease, but the clinical appearance was clearly NL. Each patient gave informed consent for the planned procedure.

PDT was initiated by the topical application of a photosensitizer onto the lesions. We used MAL (160 mg/g) in 15 patients and 5-aminolevulinic acid (200 mg/g) in 3. The lesions were then covered with occlusive and light-protective dressing for 3 h. After removal of the dressing the MAL-treated areas (n = 15) were illuminated with red light at a dose of 37 J/cm² (Aktilite lamp, Photocure ASA, Norway) and the 5-aminolevulinic-acid-treated areas (n = 2) at a dose of 75 J/cm². The third 5-aminolevulinic-acid-treated patient received 75 J/cm² for the first 2 cycles and 40 J/cm² further on. During illumination, the respective areas were cooled in 11 patients to relieve burning sensations. Three patients received analgesics (NSAIDs), 1 patient received local anesthesia with bupivacaine and 1 with mepivacaine. With respect to pain during illumination, patients tended to react differently during each session. On a 10-cm visual analog scale from 0 (no pain) to 10 (unbearable pain), the level ranged from 2 to 10 with a median of 5.

### Table 1. Overview of 18 patients with NL treated with PDT

<table>
<thead>
<tr>
<th>Patient</th>
<th>Sex</th>
<th>Age (years)</th>
<th>Institution</th>
<th>Duration of disease (years)</th>
<th>Diabetes</th>
<th>Biopsy</th>
<th>Treatment modalities prior to PDT</th>
<th>PDT cycles</th>
<th>Type of PDT/energy (J/cm²)</th>
<th>Pain score</th>
<th>Cooling</th>
<th>Interval between treatment and evaluation</th>
<th>Outcome of therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>46</td>
<td>DK</td>
<td>6</td>
<td>no</td>
<td>yes</td>
<td>allopurinol</td>
<td>9</td>
<td>MAL/37</td>
<td>4</td>
<td>yes</td>
<td>3 months</td>
<td>CR</td>
</tr>
<tr>
<td>2</td>
<td>F</td>
<td>17</td>
<td>CZ</td>
<td>2</td>
<td>no</td>
<td>n.a.</td>
<td>TS, OTS, ILS</td>
<td>3</td>
<td>MAL/37</td>
<td>7</td>
<td>yes</td>
<td>4 weeks</td>
<td>PR</td>
</tr>
<tr>
<td>3</td>
<td>M</td>
<td>32</td>
<td>D</td>
<td>8</td>
<td>no</td>
<td>yes</td>
<td>TS</td>
<td>6</td>
<td>MAL/37</td>
<td>6</td>
<td>yes</td>
<td>6 weeks</td>
<td>PR</td>
</tr>
<tr>
<td>4</td>
<td>F</td>
<td>62</td>
<td>D</td>
<td>26</td>
<td>no</td>
<td>yes</td>
<td>TS, cryo.</td>
<td>6</td>
<td>MAL/37</td>
<td>5</td>
<td>yes</td>
<td>6 weeks</td>
<td>PR</td>
</tr>
<tr>
<td>5</td>
<td>M</td>
<td>31</td>
<td>DK</td>
<td>1</td>
<td>no</td>
<td>no</td>
<td>TS</td>
<td>5</td>
<td>MAL/37</td>
<td>4</td>
<td>no</td>
<td>8 weeks</td>
<td>PR</td>
</tr>
<tr>
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<td>F</td>
<td>25</td>
<td>DK</td>
<td>2</td>
<td>yes</td>
<td>no</td>
<td>OTS</td>
<td>14</td>
<td>MAL/37</td>
<td>5</td>
<td>no</td>
<td>4 months</td>
<td>PR</td>
</tr>
<tr>
<td>7</td>
<td>M</td>
<td>57</td>
<td>DK</td>
<td>n.a.</td>
<td>n.a.</td>
<td>n.a.</td>
<td>TS, tacrolimus</td>
<td>2</td>
<td>MAL/37</td>
<td>n.a.</td>
<td>no</td>
<td>1 year</td>
<td>PR</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>57</td>
<td>D</td>
<td>4</td>
<td>no</td>
<td>yes</td>
<td>TS</td>
<td>8</td>
<td>MAL/37</td>
<td>2</td>
<td>yes</td>
<td>1 week</td>
<td>NR</td>
</tr>
<tr>
<td>9</td>
<td>F</td>
<td>60</td>
<td>D</td>
<td>19</td>
<td>no</td>
<td>yes</td>
<td>TS, PUVA</td>
<td>6</td>
<td>MAL/37</td>
<td>7</td>
<td>yes</td>
<td>3 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>10</td>
<td>F</td>
<td>35</td>
<td>D</td>
<td>4–5</td>
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<td>yes</td>
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<td>6</td>
<td>MAL/37</td>
<td>2</td>
<td>yes</td>
<td>4 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>11</td>
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<td>38</td>
<td>CZ</td>
<td>4</td>
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<td>yes</td>
<td>TS, OTS, ILS</td>
<td>3</td>
<td>ALA/75</td>
<td>4</td>
<td>no</td>
<td>4 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>12</td>
<td>F</td>
<td>27</td>
<td>CZ</td>
<td>5</td>
<td>DM1</td>
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<td>TS, OTS, ILS, PUVA</td>
<td>5</td>
<td>ALA/40–75</td>
<td>4</td>
<td>no</td>
<td>6 weeks</td>
<td>NR</td>
</tr>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>13</td>
<td>MAL/37</td>
<td>4</td>
<td>yes</td>
<td>8 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>14</td>
<td>F</td>
<td>43</td>
<td>D</td>
<td>4</td>
<td>no</td>
<td>yes</td>
<td>TS</td>
<td>8</td>
<td>MAL/37</td>
<td>3</td>
<td>yes</td>
<td>8 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>15</td>
<td>F</td>
<td>52</td>
<td>CZ</td>
<td>2–3</td>
<td>no</td>
<td>no</td>
<td>TS, OTS, ILS</td>
<td>4</td>
<td>ALA/75</td>
<td>5</td>
<td>no</td>
<td>8 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>16</td>
<td>M</td>
<td>16</td>
<td>DK</td>
<td>8</td>
<td>DM1</td>
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<td>pentoxifylline</td>
<td>2</td>
<td>MAL/37</td>
<td>10</td>
<td>yes</td>
<td>8 weeks</td>
<td>NR⁵</td>
</tr>
<tr>
<td>17</td>
<td>F</td>
<td>17</td>
<td>DK</td>
<td>3</td>
<td>no</td>
<td>yes</td>
<td>TS</td>
<td>6</td>
<td>MAL/37</td>
<td>4</td>
<td>no</td>
<td>8 weeks</td>
<td>NR</td>
</tr>
<tr>
<td>18</td>
<td>F</td>
<td>42</td>
<td>DK</td>
<td>2</td>
<td>no</td>
<td>yes</td>
<td>none</td>
<td>6</td>
<td>MAL/37</td>
<td>n.a.</td>
<td>yes</td>
<td>1 year</td>
<td>NR</td>
</tr>
</tbody>
</table>

D = Germany; DK = Denmark; CZ = Czech Republic; DM1/DM2 = diabetes mellitus type 1/2; n.a. = not assessed; TS = topical corticosteroids; OTS = occlusive topical corticosteroids; ILS = intralesional corticosteroids; cryo. = cryotherapy; PUVA = psoralen and ultraviolet A; SS = systemic corticosteroids; ALA = 5-aminolevulinic acid; CR = complete response; PR = partial response; NR = no response.

⁵ Stopped due to pain.

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Intervals between separate PDT cycles ranged from 1 to 3 weeks. In 4 patients the intervals exceeded 1 month. The number of PDT cycles performed ranged from 2 to 14 per patient with an average of 6. These variations in the treatment protocol were either due to a complete lack of effect, unacceptable side effects or the refusal of the patient to continue the treatment.

Results

The interval between PDT and evaluation of therapy outcome was 1 week to 4 months in 16 patients and 1 year in 2. One patient showed complete response after 9 cycles, 6 patients showed partial response after 2–14 cycles and 11 patients showed no response (table 1, fig. 1). Adverse events included erythema of the treated areas in 9 patients, crusting in 6 patients, pustules, exudation and blistering, respectively, in 1 patient. The only patient with complete response to therapy was a 46-year-old female without diabetes comorbidity and with a 6-year history of NL located on the shins of both legs. Allopurinol (200 mg daily for 3 months) had been given prior to MAL-PDT without effect. This patient’s lesions had shown ulcerations and a slight hyperpigmentation at the edges. Complete response was observed after 9 cycles of PDT over a period of 8 months. The patient reported no adverse events during the treatment period, and the average pain score during illumination was 4/10.

The 6 patients with partial response were all treated with MAL-PDT: 3 females and 3 males with an age of 17–62 years. The duration of NL was 1–3 years in 4 patients, and 8 and 26 years in 1 patient, respectively. Diabetes comorbidity was present in 2 of the 6 patients. Lesions were ulcerated in 2 patients and hyperpigmented in 1. All of them had been treated with topical corticosteroids prior to PDT. One patient each had previously received intralesional corticosteroids, tacrolimus, PUVA and cryotherapy. Five of the 6 patients received 2–6 cycles of PDT, while 1 received 14 cycles. The pain score during PDT ranged from 4/10 to 10/10. After therapy, 3 patients had transient erythema, 1 had a bullous reaction and 1 pustules. The partial response to PDT was observed after a period of 1 month to 1 year after therapy.

Discussion

NL is a cosmetically disturbing disease which may cause pain or itching. Therapeutic efforts to treat NL have been frustrating in the past. Topical and intralesional corticosteroids seem to be the most promising drugs, while other cases may respond to systemic therapy [5, 8].

The etiology remains subject to discussion. It has been proposed that diabetic microangiopathy is the main etiological agent, because similar changes can be seen in diabetic retinas and kidneys. Other theories about the development of NL suggest deposits of abnormal collagen or of immunoglobulins. Furthermore, metabolic changes and trauma have been discussed as etiological factors [3].

This exploratory study focused on a noninvasive approach for the therapy of NL with the use of a technique which offers the possibility of repeated applications. The overall response rate was 39% (7/18). Whereas partial cure was observed in 6 of 18 patients, dramatic improvement was noted in 1 patient only.

The mechanisms responsible for the response to therapy are not yet understood. PDT has been shown to elicit immunomodulatory effects that eventually may lead to tumor destruction. These effects include the modification of cytokine expression, induction of immune-specific responses, production of interleukin 1β, interleukin 2, granulocyte colony-stimulating factor and tumor necrosis factor α [10, 11]. Kim et al. [12] have used PDT to successfully cure granuloma annulare, a granulomatous disease with certain similarities to NL. Since the actual photosensitizer – protoporphyrin IX – has been shown to accumulate in lymphatic infiltrates and inhibit T-cell proliferation [13], it can be hypothesized that this mech-
anism was also effective in NL lesions treated with PDT.

With respect to the characteristics of the responders, there seemed to be no association with the duration of the disease. The group of responders included patients with ulcerations, hyperpigmented lesions, with and without diabetes comorbidity and with adverse events like erythema. Regarding comorbidity, clinical appearance and adverse events, no association with treatment outcome was observed. The only common variable we could identify was the use of MAL as the sensitizing agent.

One limitation of PDT for NL may be the dermal and subcutaneous localization of the disease, which is too deep to be reached by topical PDT which is most effective at a depth of 2–3 mm. Other reasons may be an inadequate penetration into altered tissue or the granulomatous character of the disease preventing cellular uptake of the photosensitizer.

Another limiting factor of PDT for NL could be pain. Even though it was not observed in a vast number of our patients, a few did report very strong discomfort during illumination. It has been shown previously that the intensity of pain significantly increased at the second PDT treatment [14]. We also observed an increase in pain with the number of PDT sessions. This may be due to an improved penetration of the topical agent because of excretion of the epidermis after the first PDT treatments. Alternatively, the nociception in the dermis may have changed by the initial PDT itself. This may limit the acceptance by the patient and therefore the feasible number of PDT sessions needed for cure.

In conclusion, in this multicenter study PDT was shown to be effective in the treatment of NL in few cases only with a response rate of 39%. We did not find specific clinical features of NL which would predict a response to PDT. A prospective and controlled study should now be started on the basis of the presented results to further evaluate the efficacy of PDT in NL.

Acknowledgement

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References