Clinical clearing of moderate and severe onychomycosis with the Nd:YAG 1064nm laser and post treatment prevention with tolnaftate

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The Foot and Ankle Online Journal 10 (1): 3

Background: Many patients prefer local treatment to systemic therapy for onychomycosis but efficacy rates, especially in moderate and severe cases have been poor. The aim of this study was to assess laser therapy clearing in all degrees of severity.

Methods: Thirty-two patients with mycologically proven onychomycosis received three laser treatments with a long pulse ND:YAG 1064nm laser. The proximal clear zones (PCZ), onychomycosis severity index (OSI) and responses to standardized nail quality of life (Nail Q of L) questions were recorded. The patients were followed for one year at 9 week intervals while using a tolnaftate in oil topical antifungal after the laser treatments were completed.

Results: The PCZ improved an average of 3.46mms (p<.05), OSI improved an average of 54.6% (p<.05) while the 15.8% improvement in the Nail Q of L responses was found to be not statistically significant.

Conclusions: Severe total dystrophic type of onychomycosis improves significantly with laser therapy. Clinically measuring the onychomycosis severity index and proximal clear zone can help to guide management of an infection that can only improve one mm per month over 9-12 months. Satisfaction surveys should not be performed frequently but may yield significant results if limited to baseline and after one complete nail growth cycle. This study found improvement in both PCZ and OSI with the 1064 YAG laser and it encourages further research combining the laser and either pulsed oral or continuous topical antifungal therapies.

Keywords onychomycosis, ND:YAG 1064nm laser, onychomycosis severity index, nail quality of life.

Onychomycosis is a progressive fungal infection of the nail unit that leads to destruction and deformity [1,2]. Approximately 2%-26% of patients amongst various populations worldwide are affected [3]. Patients with concurrent diabetes mellitus and onychomycosis have an increased risk of subungual ulceration and gangrene that can lead to amputation [4]. Onychomycosis can result in significant psychosocial and emotional concerns that may have a significant impact on quality of life [5,6]. It accounts for about half of all nail disorders, and it is the most common nail disease in adults [7]. Although many different modalities are utilized for treatment, laser therapy offers a non-systemic option that many patients prefer.
Laser therapy is a FDA approved indication for the temporary clearing of nail for patients with onychomycosis [8]. The long pulsed Nd: YAG laser at the wavelength of 1064nm has shown improvements for patients with onychomycosis between 14% - 56% [9,10]. This 1064nm wavelength is best absorbed by darker fungal pigments resulting in fungal cell apoptosis [11,12]. We offer additional evidence of the successful partial clearing of toenail onychomycosis when treated with a 1064 nm diode Nd: YAG laser. The aim of this study was to assess the clinical clearing of moderate and severe onychomycosis of the great toes with the long pulsed Nd: YAG 1064nm laser and reduction of recurrence with tolnaftate topical solution.

Materials and Methods

The subjects for this study were recruited from the patient populations of the Cleveland Foot and Ankle Clinics. This study was approved by the Kent State University College of Podiatric Medicine Institutional Review Board. All subjects signed informed consent forms prior to participating. Thirty-five adult subjects with dermatophyte positive mycology determined by PAS stained nail plate biopsies, were enrolled over a 14-month period. Fungal cultures are done in FDA drug approval protocols but podiatric physicians often use PAS pathology which is more sensitive, timely and less costly than fungal cultures. Any subjects with less than 25% of nail involvement or negative mycology were excluded. Subjects were not excluded with severe onychomycosis. Due to the difficulty in measuring lesser toenail progress accurately, data collection was limited to only hallux toenails, however all ten toenails were treated. Subjects were excluded who had received oral antifungal therapy within the last 6 months or topical antifungal therapy within last 3 months. Patients with peripheral arterial disease, severe neuropathy, immunocompromised status, current cancer, or pregnancy were excluded (Table 1).

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>%</th>
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<tbody>
<tr>
<td>Male</td>
<td>58%</td>
<td>(18)</td>
</tr>
<tr>
<td>Female</td>
<td>42%</td>
<td>(13)</td>
</tr>
<tr>
<td>Caucasians</td>
<td>71%</td>
<td>(22)</td>
</tr>
<tr>
<td>African Americans</td>
<td>29%</td>
<td>(9)</td>
</tr>
<tr>
<td>Age Range</td>
<td>31-84 years</td>
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Table 1 Study cohort.

We used a Nd: YAG 1064 long pulse laser with a spot size of 5mm (Figure 1) [8]. Data was collected at baseline, 6, 12, 18, 36, and 52 weeks. The maximal extent of the infection was measured by determining the proximal clear zone (PCZ) by measuring the distance between the proximal extent of the infection and the central cuticle. Digital photos were taken and onychomycosis severity index (OSI) score was calculated. A nail quality of life (Nail Q of L) survey was administered at each visit. Lubeck’s 15 question nail q of l survey was administered and scored on a 0-100 scale where 0 equals never; 25 equals rarely, 50 equals sometimes, 75 equals often, and 100 equals always, at each visit (Figure 2) [5]. The validated OSI of Carney et al was determined at each observation (Table 2) [13].

<table>
<thead>
<tr>
<th>OSI points [13]</th>
<th>Baseline % (N)</th>
<th>Final % (N)</th>
</tr>
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<tbody>
<tr>
<td>Cured (0 pts)</td>
<td>0% (0)</td>
<td>1.7% (1)</td>
</tr>
<tr>
<td>Mild (1-5 pts)</td>
<td>0% (0)</td>
<td>21.0% (12)</td>
</tr>
<tr>
<td>Moderate (6-15 pts)</td>
<td>14% (8)</td>
<td>17.5% (10)</td>
</tr>
<tr>
<td>Severe (16-35pts)</td>
<td>86% (49)</td>
<td>59.6% (34)</td>
</tr>
<tr>
<td>All hallux nails</td>
<td>100% (57)</td>
<td>100% (57)</td>
</tr>
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Table 2 Distribution of Study Severities over Time.

Figure 1 Digital thermogram of an actual laser procedure as a pulse was delivered to a right third toe temporarily raising the temperature of the nail unit to 41.4C. Not a study patient.
1. My nail condition hurts.
2. I worry that my nail condition may be serious.
3. I am ashamed of my nail condition.
4. I worry that my condition may get worse.
5. The skin around my nails is irritated.
6. I am embarrassed about my nail condition.
7. I am frustrated with my nail condition.
8. The skin around my nails and or nail is sensitive.
9. I am annoyed by my nail condition.
10. I have felt frustrated by the lack of improvement of my nails with previous treatment.
11. My nails are difficult to cut.
12. I rely on other people to help cut my nail(s).
13. My nails make me feel less attractive.
14. My nails make me feel self-conscious.
15. I am worried I may give my nail fungus to other people.

**Figure 2** Nail Quality of Life Survey Questions.

At each visit, nails were debrided as needed utilizing a nail nipper and rotary burr as in standard clinical practice. The nails were cleansed with alcohol and allowed to dry. Each subject received 3 laser treatments at six week intervals. The Nd: YAG 1064 long pulse laser total fluence was set at 16 Joules/cm², 0.3 milliseconds duration and two pulses per second. Each hallux nail unit received 400 pulses at each laser treatment. Rest periods were allowed between lasing cycles to accommodate an individual subject’s heat tolerance. At week 18, after completing the 3 laser treatments, patients were dispensed a topical solution of tolnaftate in oil for daily application to prevent reinfection [14]. Treatment failure was defined as an actual reduction in the proximal clear zone rather than slow improvement. Post treatment mycology was not performed since mycological cure was not the goal of this study.

The PCZ, OSI and Nail Q of L data were statistically analyzed using either a One Way Repeated Measures ANOVA or a Friedman Repeated Measures ANOVA on Ranks, depending on whether the data was normally distributed or not. Significance was defined as $p<0.05$. When significance was detected, pairwise multiple comparison procedures were performed using the Holm-Sidak method for normally distributed data or the Tukey test for data that was not normally distributed. Again, significance was defined as $p<0.05$.

**Figure 3** Progress of typical study patient’s hallux nail. A: before laser treatment, B: 9 month follow-up, C: 12 month follow-up.

**Results**

Thirty-five patients were enrolled in the study over a period of 14 months (Figure 3). Thirty-one patients with 59 infected great toenails completed all three laser treatments. Four patients failed to complete the treatment phase. One patient was excluded because his baseline mycology failed to confirm onychomycosis. One enrolled patient elected oral therapy during the study and was excluded from the results enrolled. Two patients were lost to follow up after not completing all three laser treatments.

Both the PCZ ($p<.001$) and OSI ($p<0.05$) of the moderately and severely infected great toe nails (57) improved significantly and steadily over duration of the study ($p<.001$) (Figures 4 and 5).

Interestingly, the 49 severely infected great toenails, improved their PCZ and OSI scores over the course of the study as well ($p<0.001$). Although the Nail Q of L survey responses generally improved, the degree of improvement was not statistically significant ($p=0.245$) (Figure 6).

**Figure 4** The average PCZ improved 3.46mms in all patients ($p<.05$).
Figure 5 OSI score reduced an average of 15.1 points over one year (P<0.05).

Discussion

The FDA has approved Nd: YAG 1064nm lasers for the temporary increase of clear nail [8]. Several recent articles have studied the Nd: YAG 1064nm laser for mild to moderate onychomycosis while this study looked at the more difficult to treated severe toenail onychomycosis (59.6% N=34). Recently, Renner, et al., studied the 1064 YAG laser and measured the OSI [9]. In our study, we evaluated the OSI, as well as the Nail Q of L, over one year. Renner, et al., also allowed patients with recent oral or topical therapy to be enrolled while we excluded them. We did not perform fungal cultures because they are less sensitive, costlier and take longer. Many practitioners use clinical judgement, supplemented by PAS pathological studies, which are faster, more sensitive and less costly to diagnose onychomycosis [1,7,9]. In this clinical study, to help reduce the anticipated reinfection and recurrence rates that patients can incur, patients were given a preventative topical tolnaftate 1% solution after completing all the laser treatments. There is no evidence that tolnaftate solution alone improves cure rates and tolnaftate solution does not have an FDA indication for onychomycosis but is commonly used to decrease surface contamination [15].

This study found improvement in both PCZ and OSI with the 1064 YAG laser and it encourages further research with comparative studies combining laser therapy and pulsed oral or continuous topical antifungal therapies [16].

Acknowledgements

The authors would like to acknowledge the contributions of Jill Kawalec, PhD, Director of Research, and Joan Lannoch, the Senior Graphic Designer, at Kent State University College of Podiatric Medicine.
References


