Randomized Trial of Aromatherapy
Successful Treatment for Alopecia Areata
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Objective: To investigate the efficacy of aromatherapy in the treatment of patients with alopecia areata.

Design: A randomized, double-blind, controlled trial of 7 months' duration, with follow-up at 3 and 7 months.

Setting: Dermatology outpatient department.

Participants: Eighty-six patients diagnosed as having alopecia areata.

Intervention: Eighty-six patients were randomized into 2 groups. The active group massaged essential oils (thyme, rosemary, lavender, and cedarwood) in a mixture of carrier oils (jojoba and grapeseed) into their scalp daily. The control group used only carrier oils for their massage, also daily.

Main Outcome Measures: Treatment success was evaluated on sequential photographs by 2 dermatologists (I.C.H. and A.D.O.) independently. Similarly, the degree of improvement was measured by 2 methods: a 6-point scale and computerized analysis of traced areas of alopecia.

Results: Nineteen (44%) of 43 patients in the active group showed improvement compared with 6 (15%) of 41 patients in the control group (P = .008). An alopecia scale was applied by blinded observers on sequential photographs and was shown to be reproducible with good interobserver agreement (k = 0.84). The degree of improvement on photographic assessment was significant (P = .05). Demographic analysis showed that the 2 groups were well matched for prognostic factors.

Conclusions: The results show aromatherapy to be a safe and effective treatment for alopecia areata. Treatment with these essential oils was significantly more effective than treatment with the carrier oil alone (P = .008 for the primary outcome measure). We also successfully applied an evidence-based method to an alternative therapy.

Arch Dermatol. 1998;134:1349-1352

With the recent resurgence of interest in alternative medicine, aromatherapy (aroma, from the Greek meaning spice) has attracted great public interest in its health-promoting or medicinal properties. With herbalism as its basis, aromatherapy involves the use of essential oils and essences derived from plants, flowers, and wood resins, which are generally massaged into the skin. These essential oils have been used to complement traditional medicine with some benefit. As with other forms of alternative medicine, scientific bases for the claims made are few, but in dermatologic studies, physiological and psychological benefits were found after treatment of psoriasis with aromatherapy. Significant, consistent improvement in behavior occurred in 1 of 4 patients with dementia in 10 repeated experiments. Aromatherapy is also useful in hospice care.

As examples of proven therapeutic benefit, use of sandalwood oil has been shown to significantly inhibit skin papillomas in mice. Tea tree oil is an effective bactericide and fungicide, especially for Malassezia furfur. Use of essential oils can also alter the barrier functions of the skin and induce contact dermatitis.

Cedarwood, lavender, thyme, and rosemary oils have hair growth-promoting properties. These oils have been anecdotally used to treat alopecia for more than 100 years. To date, there have been no controlled trials to evaluate this treatment. Our experience using aromatherapy provides anecdotal evidence that several patients have had marked improvement with this form of therapy. Our aim in this study was to test the hypothesis that pharmacologically active stimulants for hair growth are present in these oils and that use of these oils can be therapeutic in patients with alopecia.

Alopecia areata is a common condition affecting 1% of the Western world. It can cause substantial social and psychological distress and is often highly detrimental to the patient's well-being and self-esteem. In 1 study, alopecia areata...
PATIENTS, MATERIALS, AND METHODS

PATIENTS

Eighty-six patients diagnosed as having alopecia areata were invited to take part in this randomized, controlled trial. These patients were interviewed, and they completed a questionnaire. Patients with a medical history of hypertension, epilepsy, or pregnancy were excluded. Two of 86 patients were excluded because they had androgenic alopecia. Topical medication and intralesional corticosteroid therapy for the alopecia were discontinued before the trial.

Eighty-four patients entered the trial; 28 (68%) of the patients in the control group and 35 (81%) of the patients in the active group completed the trial (Figure 1). The distribution of patients by the 4-point scale was similar in active and control groups, as follows: 1 indicates vellus hair or no hair; 2, sparse pigmented or nonpigmented terminal hair; 3, terminal regrowth with patches of alopecia areata; and 4, terminal regrowth in all areas.

2. A standardized professional photographic assessment of each volunteer was taken at the initial interview and after 3 and 7 months. Changes in these photographic assessments formed the primary outcome measure, with improvement as the most important factor. These changes were scored independently by 2 dermatologists (I.C.H. and A.D.O.) who were unaware of the therapy administered. Improvement in photographic assessment was graded using a numerical scale (Figure 1).

3. A further secondary outcome measure was performed. A map was traced onto transparent film wherever the alopecia occurred in patches. These tracings were then transferred onto flat acetate sheets. A computerized image analyzer was used to calculate the areas of alopecia at the initial assessment and after 3 and 7 months.

STATISTICAL METHODS

We calculated that if improvement occurred in 20% more patients with active treatment than in the control group it would require 47 active and 47 control patients to detect improvement at a 5% significance level with a power of 80%. Statistical analysis was performed on an intention-to-treat basis. A pooled variance estimate Student t test for independent samples was used to test improvement in the patients’ alopecia with the map-tracing method. The chi^2 test was used to detect improvement in the active and control groups. A Mann-Whitney U test, corrected for ties, was used to assess the significance of the degree of improvement in scored photographic assessment. The level of agreement of the alopecia scoring scale between 2 assessors was examined using the Cohen weighted κ statistic.

The trial was approved by the Joint Ethical Committee for the Grampian Health Board and by the University of Aberdeen, Aberdeen, Scotland.

MEASUREMENT AND EVALUATION

Initial and 3- and 7-month assessments were made by 3 dermatologists (I.C.H. and A.D.O.) who were unaware of the therapy administered. A.D.O.) who were unaware of the therapy administered. Weighted κ statistic was 0.84 for agreement between scorers on the assessment of the photographic scale, showing good interrater correlation.

RESULTS

Eighty-four patients entered the trial; 28 (68%) of the patients in the control group and 35 (81%) of the patients in the active group completed the trial (Figure 2). The patients were well matched for the important demographic indicators that might affect response to therapy (Table 1). The distribution of patients by the 4-point scale was similar in both groups.

The improvement was statistically significant in all assessments undertaken. The primary outcome measure of improvement vs no improvement showed improvement with essential oils (P = .008, χ^2) (Table 2). The degree of improvement shown in the photographs was assessed by the Mann-Whitney U test and was significant (P = .05). The results of the alopecia scale, which scored the degree of improvement (from 1-6), are illustrated in Figure 3. The measurement of traced areas, which could be performed in only 32 patients, showed a mean±SD reduction in area affected of 103.9 ± 140.0 cm^2 compared with −1.8 ± 155.0 cm^2 in the control group (Figure 4). This was significant, with P = .05 (Student t test). A relative risk of 2.6 (95% confidence limits, 1.2, 5.6) was calculated for the likelihood of improving on the active therapy. Weighted κ statistic was 0.84 for agreement between scorers on the assessment of the photographic scale, showing good interrater correlation.

This indicates that this is a reproducible method of assessment. One patient who received active treatment and had an excellent response (ie, a score of 6 on our scale) is pictured in Figure 5.
The responses were variable but showed a clear and statistically significant advantage to treatment with this standardized regimen of aromatherapy. Although the tradition of aromatherapy is to combine several oils, it seems likely that 1 of these agents has a stimulatory effect on hair growth. One male patient also had severe androgenic alopecia, which was not included in the assessment of efficacy, and within this area there was some moderate regrowth of hair and improvement in alopecia areata.

There was a higher dropout rate in the control group, which could be explained by the fact that the volunteers became discouraged with the 7-month protocol. The control oil was not odorless because the carrier oils have some smell, and patients did not know what aroma to expect. However, they may have surmised that their treatment was inactive and withdrew from the trial. The control group’s relative lack of response again suggests a pharmacoactive property of the topically applied therapy as opposed to an effect arising from the comforting, relaxing effect of massage and of the application procedure, which was the same for both groups.

Previous studies of alopecia therapies have used only subjective scales of improvement, such as those described by MacDonald Hull and Norris. We found these scales less helpful because there are large intervals between the points in the alopecia scale. We validated the method of sequential photography with a standardized approach using a professional photographer’s studio. Images were judged by blinded observers and showed good agreement. This proved to be the most blinded and unbiased approach because there was no possibility of smells indicating which treatment had been applied.

We encountered no significant adverse events from this treatment, which makes the therapeutic ratio high.
compared with other therapies, such as diphencyprone,14,16 squaric acid dibutylester,19 and systemic or intralesional corticosteroid injections.

Populations in other trials may differ in prognostic factors. Shapiro et al20 and Gordon et al14 found a 38% success rate in producing cosmetically acceptable regrowth in patients with alopecia using diphencyprone. A review of psoralen–UV-A therapy for alopecia areata by Taylor and Hawk21 revealed that phototherapy was disappointing, with minimal benefit. Therefore, this aromatherapy trial, with an improvement rate of 44%, is comparable to and possibly of more benefit than trials of conventional therapies for alopecia areata. Compared with these other treatments, its safety is also greater, offering a better therapeutic ratio.

Accepted for publication April 16, 1998.

The Soropatomists International of Aberdeen, Scotland, provided financial support for the trial.

We thank Jill Mollison, BSc, for statistical advice and the members of the Aberdeen Alopecia Self Help Group for their participation.

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