



RESEARCH ARTICLE

F.U. Vet. J. Health Sci.
2026; 40 (3): 209 - 215
<http://www.fusabil.org>

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Effects of *Nigella sativa* and *Aloe vera* on Burn Healing: Clinical and Histopathological Evaluations

In this study, clinical and histopathological effects of *Nigella sativa* (NS) and *Aloe vera* (AV) on the healing of experimental skin burns were evaluated. Thirty-six rats were divided into three equal groups: control (cold cream, C), NS, and AV. A 1.5 cm diameter second-degree burn was created on the dorsal thoracic region and cold cream, NS, and AV were applied topically twice daily. The burn and epithelialization areas were measured on days 0, 4, 7, 10, and 14. On days 7 and 14, six rats from each group were euthanized for histopathological examination.

Macroscopic examinations revealed that the healing was better in the AV group on days 4, 7, and 10; on day 14, healing was better in both the AV and NS groups compared to the control group ($p < 0.0001$).

Histopathological examinations showed severe necrosis on day 7 and severe mononuclear (Mn) cell infiltration between days 7 and 14 in the control group. Oedema was severe in the AV and NS groups on day 7. Epithelialization was incomplete in the control group on day 7, moderate in the NS group, and advanced in the AV group. Epithelialization was complete in the AV and NS groups on day 14. Granulation tissue formation was advanced in the AV group.

Clinical and histopathological findings demonstrated that AV enhances the healing of skin burns by preventing necrosis and promoting epithelialization. In conclusion, the obtained data indicate that NS and AV can be safely used in the treatment of thermal skin burns.

Key Words: *Aloe vera*, *Nigella sativa*, burn, rat

Nigella sativa ve *Aloe vera*'nın Yanık İyileşmesi Üzerine Etkileri: Klinik ve Histopatolojik Değerlendirmeler

Bu çalışmada, *Nigella sativa* (NS) ve *Aloe vera*'nın (AV) deneysel deri yanıklarının iyileşmesi üzerindeki klinik ve histopatolojik etkileri değerlendirildi. Otuz altı rat kontrol (Cold krem, C), NS ve AV olmak üzere üç eşit gruba ayrıldı. Dorsal torasik bölgede 1.5 cm çapında 2. derece yanık oluşturuldu ve günde iki kez cold krem, NS ve AV topikal olarak uygulandı. Yanık ve epitelizeasyon alanları 0., 4., 7., 10. ve 14. günlerde ölçüldü. Yedinci ve 14. günlerde histopatolojik inceleme için her gruptan 6 rat ötenazi edildi.

Makroskopik incelemelerde, iyileşmenin 4., 7. ve 10. günlerde AV grubunda daha iyi olduğu; 14. günde ise AV ve NS gruplarında kontrol grubuna göre daha iyi olduğu gözlemlendi ($p < 0.0001$).

Histopatolojik muayenelerde, kontrol grubunda 7. günde şiddetli nekroz ve 7.-14. günlerde şiddetli düzeyde mononükleer (Mn) hücre infiltrasyonu saptandı. Ödem, 7. günde AV ve NS gruplarında şiddetli düzeydeydi. Epitelizasyon 7. günde kontrol grubunda tam olarak şekillenmemişken, NS grubunda orta, AV grubunda ileri düzeydeydi. Ondördüncü günde AV ve NS gruplarında epitelizasyon tamamlanmıştı. Granülasyon dokusu oluşumu AV grubunda ileri düzeydeydi.

Klinik ve histopatolojik bulgular, AV'nin nekroz oluşumunu önleyerek ve epitelizasyonu destekleyerek deri yanıklarının iyileşmesini artırdığını göstermiştir. Sonuç olarak, elde edilen veriler NS ve AV'nin termal deri yanıklarının tedavisinde güvenle kullanılabileceğini ortaya koymaktadır.

Anahtar Kelimeler: *Aloe vera*, *Nigella sativa*, yanık, rat

Received : 27.04.2026
Accepted : 02.06.2026

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Introduction

In the treatment of thermal burns, ointments containing antibiotics, antimicrobials, and antiseptics, as well as those that promote epithelialization and possess antioxidant properties, are used with the aim of achieving rapid tissue repair (1-3). Research is still ongoing to identify the most effective treatment method for burns (4).

Nigella sativa (NS) is composed of approximately 36-38% fixed oils along with proteins, alkaloids, and saponins, and it also contains 0.4-2.5% essential oils. In addition, it includes several bioactive constituents such as thymoquinone, dithymoquinone (5), and thymol (6). Through these compounds, it exhibits antineoplastic, antibacterial, antifungal, anthelmintic, antidiabetic, and antioxidant effects (7). NS has been reported to show therapeutic efficacy in conditions including inflammation, bronchitis, asthma, and eczema (8). It is also reported to support wound healing by stimulating angiogenesis, increasing fibroblast proliferation and collagen

synthesis, and reducing leukocyte count, tissue damage, and bacterial contamination (9).

Aloe vera (AV) possesses topical anesthetic, tissue penetration, antimicrobial, anti-inflammatory, and immunomodulatory properties (10). AV, containing vitamins C and E, essential amino acids, and polysaccharides, is a powerful moisturizer that protects the skin barrier (11). It has been reported that AV shortens the treatment duration by supporting vascularization and accelerating healing and epithelialization (12). Additionally, it has been reported to be effective against various bacterial pathogens that cause infection in burn wounds, as well as certain *Candida* species (3).

This study aimed to systematically assess the therapeutic effects of NS and AV on burn wound healing in an experimental rat model and to clinically and histopathologically compare the efficacy of these two herbal treatment agents.

Materials and Methods

Research and Publication Ethics: The present study was performed following approval from the Firat University Animal Experiments Local Ethics Committee (FU-HADYEK) (approval date: 30.06.2021; decision no: 2011/11) and was financially supported by the Scientific Research Projects Coordination Unit of Firat University (FÜBAP) (Project No: VF.21.20).

Animals and Experimental Groups: This study utilized 36 female Wistar albino rats aged 2–3 months. The animals were randomly assigned to three main groups (n=12 each), and each group was further divided into two subgroups based on experimental duration (7 and 14 days), with six rats per subgroup. Before the experimental procedures, all animals were acclimatized for at least one week in plastic cages under controlled environmental conditions (temperature and humidity), with unrestricted access to standard laboratory chow and water.

Anesthesia: General anesthesia was induced by intramuscular administration of 6 mg/kg Xylazine hydrochloride (Rompun, Bayer, 23.32 mg/mL), followed 10 minutes later by 85 mg/kg Ketamine hydrochloride (Ketalar, Parke-Davis, 50 mg/mL).

Creation of the Burn Model: After shaving the dorsal thoracic regions of the rats, the areas were disinfected with a 10% povidone-iodine solution. A standardized full-thickness second-degree burn lesion with a 1.5 cm diameter was induced in each animal. For this purpose, a brass probe of the same diameter was heated in boiling water (100°C) until thermal equilibrium was achieved, then gently applied to the dorsal thoracic region without pressure and maintained in contact for 15 seconds. Following the procedure, all rats received intraperitoneal resuscitation with Lactated Ringer's solution at a dose of 2 mL/100 g body weight (6, 9, 13–15). A total of 36 rats were then allocated equally into three experimental groups. Cold cream (control, K) was applied topically twice daily to the 12 rats in the first

group, NS to the 12 rats in the second group, and AV to the 12 rats in the third group.

Clinical Examinations: Digital photographs of the burn areas were taken from a distance of 8 cm under standard conditions on days 0, 4, 7, 10, and 14 following burn induction. The wound healing process was evaluated using these images. A quantitative assessment of wound healing was carried out using ImageJ software (National Institutes of Health, Bethesda, MD, USA) (16). In this process, wound areas were quantified from digital images acquired at each time point using ImageJ, and the wound closure rate was calculated over time.

Histopathological Examinations: After burn induction, six animals from each experimental group were euthanized on days 7 and 14. Skin specimens, including the burn area and adjacent normal tissue, were harvested for histopathological evaluation.

The skin specimens were fixed in 10% neutral buffered formalin. After fixation, routine tissue processing was performed through graded alcohols and xylene, followed by paraffin embedding. Sections of 4 µm thickness were cut and placed on poly-lysine-coated slides. Hematoxylin and eosin staining was applied for routine histological evaluation, while Masson's trichrome staining was used to assess granulation tissue formation. All slides were examined under a light microscope for necrosis, mononuclear (Mn) cell infiltration, oedema, and epithelialization. Histopathological findings were semi-quantitatively graded as absent (–), mild (+), moderate (++), or severe (+++) (Table 1).

Table 1. Histopathologic scoring

Necrosis	–	:	None
Mn cell infiltration	+	:	1/3 of the wound length
Epithelialization	++	:	2/3 of the wound length
	+++	:	Along the entire length of the wound
Granulation tissue formation,	–	:	None
Oedema	+	:	5–20% of the field
	++	:	20–40% of the field
	+++	:	40% or more of the field

Statistical Analysis: All collected data were analyzed using IBM SPSS Statistics software (version 22). The Shapiro-Wilk normality test was performed to determine whether the raw values of all measured parameters followed a normal distribution. For parametric data showing a normal distribution, a one-way analysis of variance (ANOVA) was performed to determine differences between groups, and the *post hoc* Tukey test was used for pairwise comparisons. Differences in histopathological findings between groups were determined using the Kruskal-Wallis test, a nonparametric test, and the group showing the difference was identified using the Mann-Whitney U test. A *p*-value less than 0.05 was considered statistically significant.

Results

Clinical Findings: Macroscopic examinations revealed marked inflammation and erythema in the burn area on day 1. On days 4 and 7, healing in the AV group was observed to be better than in the other groups ($p < 0.0001$). While scab detachment occurred earlier in the AV group compared to the NS group, no scab detachment was observed in the control group for 14 days. On the 10th day, it was determined that healing in the NS and AV groups was better than in the control group, with the best healing observed in the AV group ($p < 0.0001$). By the fourteenth day, healing in the treatment groups was better than in the control group ($p < 0.0001$); although there was no statistical difference between the NS and AV groups, it was determined that healing in the AV group was numerically better than in the NS group.

Macroscopic images illustrating the effects of NS and AV on cutaneous burn healing in rats are presented in Figure 1, while data on the remaining wound area according to the post-burn period are presented in Table 2.

Histopathological Findings: On days 7 and 14, significant differences were detected among the experimental groups regarding necrosis, Mn cell infiltration, oedema, epithelialization, and granulation tissue formation (Table 3).

On day 7, severe necrosis was observed in the control group, while mild necrosis was observed in the NS group. On day 14, mild necrosis was observed in the NS group, while no necrosis was detected in the control and AV groups.

On day 7, Mn cell infiltration was assessed as severe in the control group and moderate in the NS and AV groups. In the AV group, Mn cell infiltration decreased to a mild level on day 14. Oedema was evaluated as severe in both NS and AV groups on day 7, while it decreased to a moderate level in the NS group and a mild level in the AV group on day 14. Epithelialization was not yet fully established in the control group. On day 7, epithelialization was moderate in the NS group and more advanced in the AV group. By day 14, complete epithelialization was observed in both treatment groups.

Although granulation tissue had not yet fully begun to form in the control group by day 7, mild formation was observed in the NS and AV groups. By day 14, granulation tissue formation was graded as moderate in both the control and NS groups, whereas the AV group exhibited advanced granulation tissue development (Figure 3).

Table 2. The remaining burn wound area relative to the elapsed time

Day	Groups			p-value*
	Control	<i>Nigella sativa</i>	<i>Aloe vera</i>	
0	176.625	176.625	176.625	-
4	122.62±13.21 ^{a, A}	115.89±10.03 ^{a, A}	90.97±9.52 ^{b, A}	<0.0001
7	81.79±5.67 ^{a, B}	76.04±10.12 ^{a, B}	57.51±9.16 ^{b, B}	<0.0001
10	41.36±6.15 ^{a, C}	32.21±5.36 ^{b, C}	19.77±3.09 ^{c, C}	<0.0001
14	12.02±1.88 ^{a, D}	5.12±2.71 ^{b, D}	2.14±1.89 ^{b, D}	<0.0001
p-value**	<0.0001	<0.0001	<0.0001	-

* The difference between groups (a–c) with different letters on the same row is statistically significant ($p < 0.05$).

** The difference between time points labeled with different letters (A–D) in the same column is statistically significant ($p < 0.05$).

Table 3. Statistical analysis of histopathological findings

	Necrosis	Mn cell infiltration	Oedema	Epithelialization	Granulation tissue
Day 7 Control	2.83±0.51 ^a	2.66±0.40 ^a	0.33±0.51 ^a	0.12±0.40 ^a	0.33±0.51 ^a
Day 7 <i>Nigella sativa</i>	1.12±0.40 ^b	1.83±0.51 ^b	2.66±0.40 ^b	0.12±0.40 ^a	1.12±0.40 ^b
Day 7 <i>Aloe vera</i>	0.33±0.51 ^c	1.83±0.51 ^b	2.83±0.51 ^b	1.66±0.40 ^b	1.00±0.00 ^b
Day 14 Control	0.12±0.40 ^c	2.66±0.40 ^a	0.33±0.51 ^a	0.33±0.51 ^a	1.66±0.40 ^c
Day 14 <i>Nigella sativa</i>	0.33±0.51 ^c	1.66±0.40 ^b	1.83±0.51 ^c	2.83±0.51 ^c	1.83±0.51 ^c
Day 14 <i>Aloe vera</i>	0.12±0.40 ^c	0.83±0.40 ^c	1.00±0.00 ^d	2.66±0.40 ^c	2.83±0.51 ^d

a,b,c,d, Different letters in the same column indicate differences between groups ($p < 0.05$).

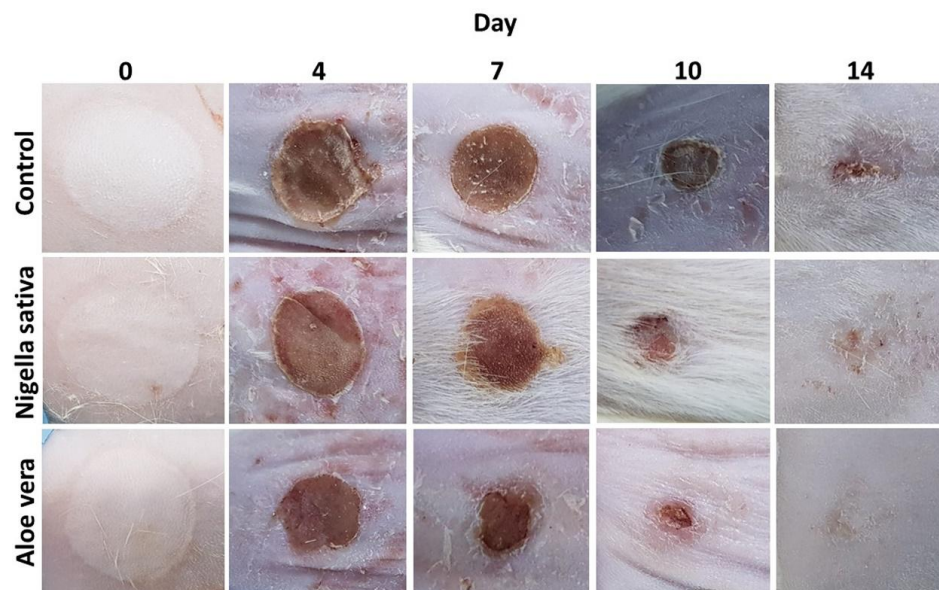


Figure 1. The effects of *Nigella sativa* and *Aloe vera* on the healing of cutaneous burns in rats. The figure shows representative images of the wounds, and rats treated with *Nigella sativa* and *Aloe vera* exhibited more pronounced healing compared to the control group. Time-dependent healing was observed in all groups.

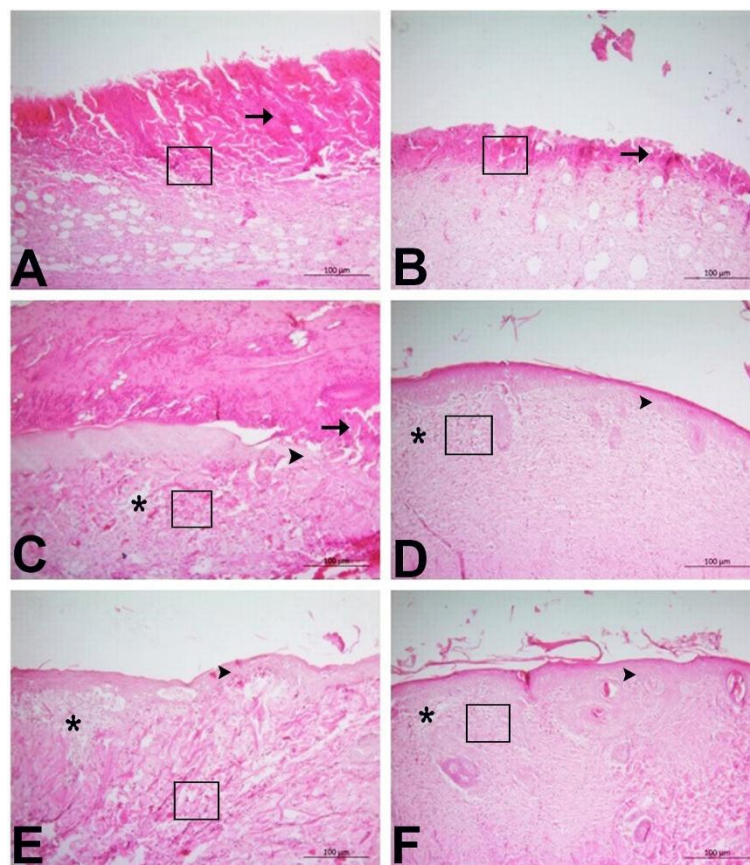


Figure 2. Microscopic appearance of burn wound healing on days 7 and 14. H&E. **A.** Control (Day 7): Severe necrosis (arrow) and Mn cell infiltration (□). **B.** Control (Day 14): Mild necrosis (arrow), severe Mn cell infiltration (□). **C.** *Nigella sativa* (Day 7): Mild necrosis (arrow), moderate Mn cell infiltration (□), moderate epithelialization (arrowhead), severe oedema (*). **D.** *Nigella sativa* (Day 14): Moderate Mn cell infiltration (□), severe oedema (*) and severe epithelialization (arrowhead). **E.** *Aloe vera* (Day 7): Moderate Mn cell infiltration (□), severe oedema (*) and severe epithelialization (arrowhead). **F.** *Aloe vera* (Day 14): Mild Mn cell infiltration (□), mild oedema (*), and severe epithelialization (arrowhead).

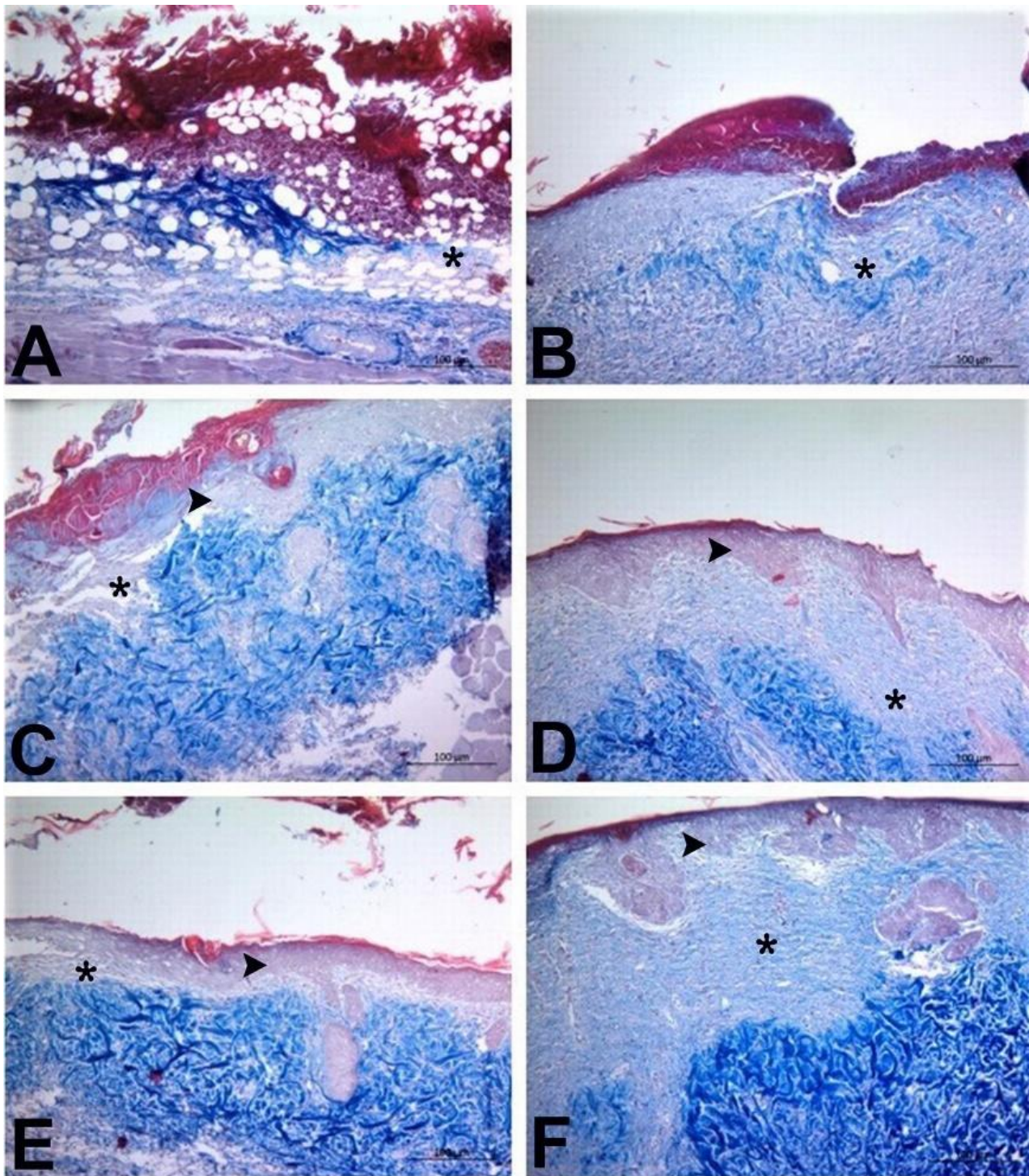


Figure 3. Microscopic appearance of burn wound healing on days 7 and 14. MST. **A.** Control (Day 7). Minimal granulation tissue formation (*), **B.** Control (Day 14). Moderate granulation tissue formation (*), **C.** *Nigella sativa* (Day 7). Mild granulation tissue formation (*) and moderate epithelialization (arrowhead), **D.** *Nigella sativa* (Day 14). Moderate granulation tissue formation (*) and severe epithelialization (arrowhead), **E.** *Aloe vera* (Day 7). Mild granulation tissue formation (*) and severe epithelialization (arrowhead), **F.** *Aloe vera* (Day 14). Severe granulation tissue formation (*) and severe epithelialization (arrowhead).

Discussion

Burn healing is a dynamic, multi-stage biological process that begins following tissue damage and involves phases of inflammation, proliferation, and remodeling. The primary clinical goal in this process is to ensure that the wound heals as quickly as possible with minimal scar formation by preventing the development of complications, particularly infection (6, 9, 13, 15, 17). The effects of plant-derived agents on wound healing have long been studied in both traditional and modern medicine (17). The aim of this study was to investigate the clinical and histopathological effects of NS and AV on an experimental burn model established in rats and to determine any differences in efficacy between the groups.

In this study, the burn model was created by applying a 1.5 cm diameter brass probe heated to 100°C to the dorsal region of rats for 15 seconds, in accordance with similar methodologies reported in the literature (6, 9, 13-15). The full-thickness second-degree skin burn created by this method allowed for the establishment of a standard tissue injury and the objective monitoring of the healing process. Wound area measurement is of critical importance for the reliability of results in experimental studies. It has been reported that individual variations and measurement errors in drawings made on live animals can lead to deviations in area calculations (6, 9, 13, 14, 17). In this study, digital measurements were performed using ImageJ software instead of traditional acetate paper and millimeter-based calculation methods, with the aim of minimizing measurement errors and enhancing the objectivity of the results.

In this study, macroscopic evaluations revealed that wound area reduction in the AV group on days 4, 7, and 10 was statistically significantly greater than in all other groups ($p < 0.0001$). By day 14, significant improvement was observed in both the NS and AV groups compared to the control group ($p < 0.0001$). However, the data indicate that the AV group's rate of healing is higher than that of the NS group, suggesting that AV may be a more effective alternative in burn treatment.

In the first few days following a burn injury, the tissues surrounding the central necrotic area may progress to necrosis due to inflammation and oxidative stress. This can lead to an expansion of the area requiring surgical intervention and increased scar formation. In this study, the fact that necrosis remained at a lower level in the treatment groups compared to the control group demonstrates that NS and AV have the capacity to limit tissue loss in the stasis area.

It has been reported that NS possesses antimicrobial, antioxidant, anti-inflammatory, and immunomodulatory properties and accelerates the histopathological healing process in burn models (6). It is stated that the positive effects of NS on wound healing are associated with the induction of angiogenesis,

increased fibroblast proliferation, and stimulation of collagen synthesis. It is also noted that it contributes to reducing tissue damage and bacterial infection by regulating leukocyte activity (18).

AV is widely used due to its antibacterial, antiviral, and anti-inflammatory properties (19, 20). It has been reported that AV is more effective in chronic wounds than in acute wounds and accelerates the healing process of burn wounds due to its anti-inflammatory effects, enhanced immune activity, antibacterial and antiviral effects, and reduced histamine activity (21).

In the histopathological evaluation, collagen deposition and granulation tissue formation during the proliferative phase (key indicators of wound healing) were taken into account (22, 23). By day 7, a significant increase in granulation tissue was observed in the NS and AV groups compared to the control group. By the fourteenth day, it was notable that granulation tissue levels were higher in the AV group compared to the other groups.

When evaluated in terms of epithelialization, the fact that the epithelialization rate in the AV group was significantly higher than in all other groups on day 7 demonstrates the significant positive effect of AV on early-stage epithelialization. The higher levels of necrosis observed in the control group on day 7 and of mononuclear cell infiltration on days 7 and 14; in contrast, the more advanced epithelialization observed in the NS and AV groups on day 14, indicate that NS and AV positively influence the healing process.

One limitation of the present study is that repeated measurements obtained from the same animals were evaluated primarily through time-point-based group comparisons. In addition, the relatively limited sample size in histopathological evaluations should be considered when interpreting the findings. Future studies using longitudinal statistical models and larger sample sizes may provide additional insight into healing dynamics over time.

In conclusion, the findings of this study demonstrate that both AV and NS significantly accelerate burn healing at both the clinical and histopathological levels. However, AV exhibited more pronounced efficacy compared to NS in terms of limiting necrosis, initiating early-stage epithelialization, and achieving faster wound bed reduction. While this study supports the potential of herbal agents in burn treatment, it also highlights the need for comprehensive research evaluating different dosages, application forms, and long-term outcomes to fully integrate these findings into clinical practice and veterinary medicine.

Acknowledgements

We thank Prof. Dr. Mustafa ÖZKARACA and Ali Sefa MENDİL for their contributions to the histopathological evaluations.

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