



Ozone Therapy for Dermatological Conditions: A Systematic Review

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BACKGROUND: Known in the past for its toxic aspect as the main urban pollutant, in the last few decades, ozone has been gaining greater visibility for its possible antimicrobial, antiviral, and antioxidant effects when used in human dermatological pathologies. Despite the reports of clinical benefits, the standard dosage for clinical efficacy and safety are yet not clear, nor are its means of application and its true acting mechanism. **OBJECTIVE:** We conducted a review to determine the efficacy and safety of ozone therapy for a variety of dermatological conditions. **METHODS:** We considered clinical trials (both randomized and non-randomized) published between December 2020 and March 2021 as long as they provided some PICO information, i.e., population (P), intervention (I), and study design. The skin dermatological conditions researched were: acne, dermatitis, psoriasis, systemic sclerosis, herpes, aging, ulcers, and skin scarring. **RESULTS:** A total of 326 articles were identified and 150 remained after duplicates were removed. After titles, abstracts and full articles were read, 17 articles were included in the systematic review (with 643 patients). **CONCLUSION:** Ozone therapy seems promising for some dermatological conditions; however, the articles included in this review had methodological limitations and did not sufficiently demonstrate sound evidence for safe therapy. Therefore, more studies with better methodological standards and longer-term assessments of side effects should be conducted to achieve better standards and safety in ozone therapy for dermatological conditions. **KEY WORDS:** dermatological affections, dermatological pathologies, ozone, ozone therapy

Ozone has been used in medicine for over 100 years in an alternative and empirical way; however, in the last decade, there have been great advances in the therapy due to new technological resources, with greater precision in gas concentration and a better understanding of its possible action mechanisms. Thus, ozone therapy has turned out to be an open research field, in which studies aim to clarify its effectiveness and biological performance, as to ascertain the control of its therapeutic actions.^{1,2}

Ozone is defined as an oxygen (O₂) tri-atomic molecule. Its first report occurred in 1840, when Christian Friedrich Schonbien found that, via electron sharing, the properties of atoms allowed them to originate ≥1 simple and different substances, like ozone (O₃).^{2,3}

In the mid-19th century, the first ozone generator was developed following the concept of transformation of pure oxygen via a high-voltage gradient, generating as final product a mixture of gases that was 95% oxygen and 5% ozone. However, these first devices did not always possess a precise photometer to calculate the percentage of

ozone in the mixture.^{3,4}

The gas is colorless, with a sour odor and explosive in its liquid or solid form. Its half-life is 40 min at 20°C and about 140 min at 0°C. O₃ is hazardous to humans, especially when inhaled, but for therapeutic purposes the gas must be produced from medical grade oxygen and administered in precise therapeutic doses, never by inhalation.^{1–3,5}

It is believed that ozone therapy, together with many other therapies, was methodically eliminated from the educational process in the United States; however, this suppression was milder in Europe. Hence, in Germany and other countries, ozone therapy is common. More than 7,000 doctors in Germany use ozone therapy daily. Nowadays in the United States, only 13 states practice ozone therapy. In countries like Cuba, Spain, Italy, Russia, Ukraine, and Greece, the practice is a little more widespread.^{4,5}

As the gas is extremely unstable, there are some forms used in ozone therapy in dermatology. They are mainly categorizable as topical application with ozonated water or oils, ozone bags,

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TABLE 1. Administration routes of ozone therapy in dermatology

ADMINISTRATION	PREPARATION	METHOD	ADVANTAGES	DISADVANTAGES
Topical	Ozone in water or oil	Immersion or compress	Therapy dosage maintenance/No age restrictions/Low adverse risk	Rapid ozone degradation when in water
Ozone bag	Ozone and oxygen	Topical with containment of gases with appropriate plastic.	Action on bigger lesions	Rapid ozone degradation
Subcutaneous	Ozone and oxygen	Subcutaneous	Local action	No systemic action
Rectal insufflation	Ozone and oxygen	Rectal	Immediate stimulus to cell metabolism	Rapid ozone degradation
Major and minor autohemotherapy	Ozone with individual's own blood	Intravenous	Immediate stimulus to immune system	Contamination risk

subcutaneous applications, rectal insufflation, and major or minor autohemotherapy.^{1,6} (Table 1).

As ozone is a potent oxidative physiological response, therapy has been used to disinfect and treat diseases for over 150 years, being recommended to treat infections, wounds, and multiple diseases. During World War I, doctors conversant with the antibacterial properties of O₃ applied it topically to infected wounds and discovered that not only did O₃ heal the infection, it also had hemodynamic and anti-inflammatory properties.⁵

Starting from its physiological effects, researchers began to assess the use of ozone therapy in skin conditions. The first evidence of beneficial effects of ozone on skin diseases was given by Shpektorova in 1964 and soon made apparent by Białoszewski et al. in 2003, when patients with chronic infected wounds that were resistant to antimicrobial therapy underwent topical ozone therapy. Results showed inhibition of the infections and a faster healing process.⁷⁻⁹

Other types of dermatological conditions, such as acne, dermatitis, psoriasis, herpes, systemic sclerosis, aging, ulcers, and skin scarring, have been being included in ozone therapy treatment protocols; nonetheless, there is no clear evidence of the real efficacy or safety nor guidelines to such interventions.^{1,6,7,8}

Hence, this systematic review is justified as an evaluation of whether ozone therapy has efficacy and safety and if it may be recommended as therapy for those

dermatological conditions.

METHODS

Protocol and registration. A systematic review was carried out, following the Preferred Reporting Items for Systematic Reviews and Meta-Analyses 2020 checklist. The research was registered in the International Prospective Register of Systematic Reviews (PROSPERO no. CRD42021238665).

Eligibility criteria. Eligibility criteria followed some PICO elements, i.e., population (P), intervention (I), and study design. The studies included children, men, and women with skin dermatological conditions such as acne, dermatitis, psoriasis, systemic sclerosis, herpes, aging, ulcers, and skin scarring. Eligible trials of intervention were those that assessed the efficacy, safety, and application benefits of ozone therapy in the aforementioned types of dermatological conditions.

Randomized, non-randomized, and quasi-randomized clinical trials were considered. This research was conducted without date, sample size, or language restrictions for the evaluation of publications. Trials that associated ozone with other therapies in the same area of treatment were excluded.

Information sources and search strategy. Scientific databases assessed for the identification of relevant trials were the Cochrane Central Register of Controlled Trials (CENTRAL), National Center for Biotechnology Information (PubMed), ScienceDirect, Biblioteca Científica Eletrônica

Online (SciELO), and Embase. The following medical scientific terms (Medical Subject Headings) were considered to develop the research strategy: [("ozone therapy") AND ("acne") AND ("dermatitis") AND ("psoriasis") AND ("herpes") ("systemic sclerosis") AND ("aging") AND ("ulcers and scarring skin")].

Trial selection. The search strategy in scientific databases was conducted by 2 independent researchers (D. A. O. M. and R. C. F.), and any disagreements on whether to include a trial in the systematic review were resolved by a third senior researcher (K.C.R.), who was invited to aid in the analysis and inclusion of the article. Randomized, non-randomized, and quasi-randomized clinical trials were included in this systematic review. Studies were primarily selected by their title, then secondly by their abstract, underwent verification of the type of trial, and, in case they contained the proposed subject matter, were accessed in full form for final evaluation and inclusion.

Moreover, the references present in the articles were also checked, with the purpose of identifying any other study missed by the search strategy. After reading the full versions of selected studies, in cases deemed necessary, authors were contacted and requested to provide the data presented in the articles. Data extraction from the articles was done via a data-collection form and tabulated in an electronic spreadsheet, with the following data for each study: database, type of research, article, author/year, recommendation, aim of study, studied

population, intervention, results, and conclusions.

Data collection process. Results were expected to demonstrate if ozone therapy is safe, effective, and present benefits in healing of skin dermatological conditions, such as acne, dermatitis, psoriasis, herpes, systemic sclerosis, aging, ulcers, and skin scarring.

Risk of bias in individual studies. The risk of bias was assessed according to the checklist by Downs and Black, which was already used in another review study previously published.¹⁰

Synthesis of results. Trials were divided according to their recommendations: acne, dermatitis, psoriasis, systemic sclerosis, herpes, aging, ulcers, and scarring. For trials that compared ozone therapy to other techniques, only the arm treated with ozone was considered.

RESULTS

Search history. Initially, 326 articles from January to March 2021 were identified by searching the databases for the key words. After the removal of duplicate articles, 150 were selected. After examination of the titles, 49 were chosen and, after reading of abstracts, 24 articles remained. After reading of full articles, 17 were included in this systematic review (643 patients, Figure 1).

Study characteristics. Table 2 describes the main characteristics and results of the 17 studies with the objective of enhancing the clinical state of dermatological conditions, subdivided by conditions included in the review. As for study design, 3 trials were non-randomized¹⁰⁻¹² and 14 were randomized.¹³⁻²⁶ Moreover, considering the ozone therapy recommendation, 2 articles were on acne,^{10,13} 3 were on atopic dermatitis,¹⁴⁻¹⁶ 2 on psoriasis,^{11,17} 2 were on systemic sclerosis,^{12,18} 1 was on herpes-zoster,¹⁹ 6 were on ulcers,²⁰⁻²⁵ and 1 was on scarring.²⁵ No study on aging was found during the data search.

Ozone therapy application routes. Choosing the route of application for ozone therapy is an important factor for the results. In this review, 12 trials used ozone therapy as a topical agent (ozonated oil) or immersion in hydrotherapy with ozone^{10-17,19-22,25}; 5 trials used only ozone bags,^{18,20,23,24,25}; 1

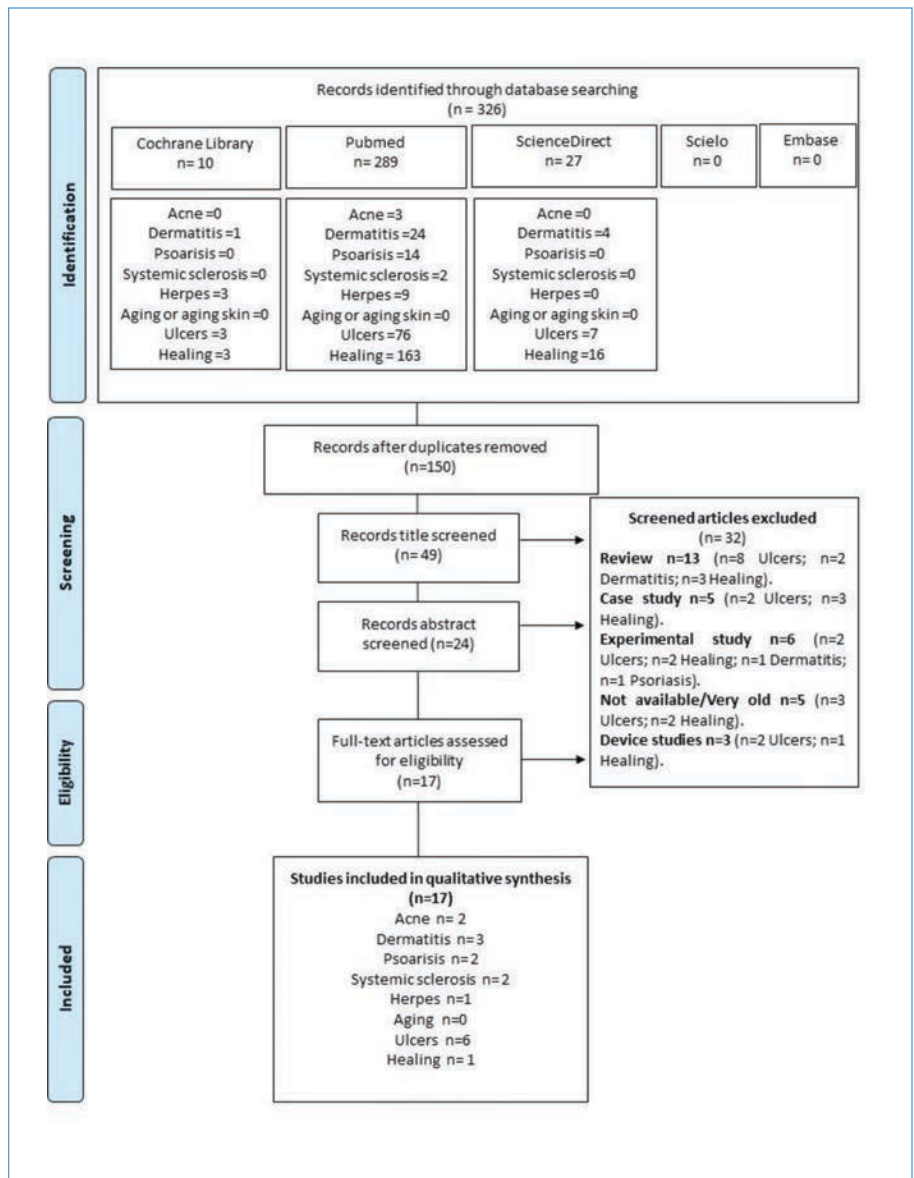


FIGURE 1. Flowchart of the studies included in the systematic review.

trial combined topical use with ozone bag, subcutaneous, and rectal insufflation^{21,25}; and 1 trial used autohemotherapy.¹³ Ozone concentrations varied according to the route of application; however, in 4 studies, information on the concentration used was not found.

Efficacy, benefits, safety, and adverse events in ozone therapy.

Only Gloor and Lipphardt¹¹ reported ozone therapy as non-effective for acne treatment, because the therapy did not present the expected effect on *Propionibacterium acnes* and other microorganisms. The other included studies reported a possible safety

of the therapy, through the use of adequate ozone concentrations for the treatment of each type of dermatological condition, with the dose used not being sufficient to cause damage and adverse effects to the research participants.

Benefits reported for acne were reduction in the inflammatory process and, for dermatitis, were reduction of size of lesions and prurigo.¹²⁻¹⁴ In psoriasis, the therapy reduced painful states, enhanced circulation, and the immune system^{15,16}; in systemic sclerosis, it aided in early healing and enhanced joint mobility,^{17,18}; for herpes-zoster, it significantly reduced duration of

TABLE 2. Characteristics of the studies included in the systematic review

AUTHOR/YEAR	DERMATOLOGICAL CONDITIONS	TYPE OF STUDY	NUMBER OF PATIENTS	GROUP CONTROL	INTERVENTION GROUP
Davatdarova and Kazimov, 2008	Acne	RTC	72	Standard drug treatment	Autohemotherapy and topical ozone
Gloor and Lipphardt, 1976	Acne	Not RTC	16	NA	Topical ozone
Zeng et al., 2020	Dermatitis	RCT split sides	12	Routine care	Ozonated water immersion and topical ozonated oil
Guizhi et al., 2018	Dermatitis	RTC	60	Common water immersion	Ozonated water immersion and topical ozonated oil application
Jianyun et al., 2018	Dermatitis	RCT split sides	12	Common water wash and common oil application	Ozonated water immersion and ozonated camellia oil
Gao et al., 2019	Psoriasis	Not RCT	20	-	Ozonated oil
Lina et al., 2018	Psoriasis	RCT	40	Methone ointment with 0.2 mg/g of flumetasone	Ozonated oil
Hassaniem et al., 2018	Systemic sclerosis	RCT	50	Drugs (Epilat Retard 40 mg/day)	Drugs (Epilat Retard 40 mg/day) associated with ozone bags
Nowicka, 2017	Systemic sclerosis	Not RCT	42	-	Ozonated water immersion
Jian et al., 2018	Herpes zoster	RTC	60	Routine care: drugs, low-intensity laser, and topical ointment	Routine care associated to ozonated water compress and topical ozonated oil
Kasmawati et al., 2020	Diabetic foot ulcer	RTC	27	Antimicrobial dressings	Antimicrobial dressings associated to ozone bags
Izadi et al., 2019	Diabetic foot ulcer	RTC	100	Routine care (debridement and dressings)	Association of ozone bags and ozonated gel; subcutaneous application around the wound and systemic use by rectal insufflation
Solovăstru et al., 2015	Chronic Venous Ulcers	RTC	29	Routine care (debridement and dressings)	Routine care associated to topical ozonated sunflower oil
Zhang et al., 2014	Diabetic foot ulcer	RTC	50	Routine care (debridement and dressings)	Ozone bag
Wainstein et al., 2011	Diabetic foot ulcer	RTC	32	Routine care (debridement and dressings) and placebo ozone therapy (inactive form)	Routine care associated with ozone bag
Martínez-Sánchez et al., 2005	Diabetic foot ulcer	RTC	51	Only drug therapy	Ozone bag, ozonated oil and rectal insufflation
Campanati et al., 2013	Healing	RTC split sides	30	Topical hyaluronic gel	Ozonated oil

TABLE 2 (continued). Characteristics of the studies included in the systematic review

AUTHOR/YEAR	OZONE CONCENTRATION	NUMBER OF SESSIONS/TIME	CONCLUSION	RISK OF BIAS
Davatdarova and Kazimov, 2008	100 mL of blood with 150 mL of ozonated saline solution/topical not reported	Not reported/topical 3–4 times/week	Forms of ozone therapy were effective when compared to drug therapy	Not reported
Gloor and Lipphardt, 1976	Not reported	Once daily for 1 week	Ozone therapy was ineffective against <i>P. acne</i> and other microorganisms	Not reported
Zeng et al., 2020	Ozonated water (3.0 ± 1.5 mg/L); oil not reported	Once daily for 15 minutes and topical oil twice daily for 3 days	Compared to the control group, ozone therapy may significantly mitigate lesions	Not reported
Guizhi et al., 2018	Ozonated water (3.0 ± 1.5 mg/L). Oil not reported	3–5 times/week for 15 min for 2 weeks	Ozone therapy is safe and efficient	Not reported
Jianyun et al., 2018	Ozonated water (3.0 ± 1.5 mg/L). Oil not reported	Twice daily for 7 days	Reduced effects of <i>S. aureus</i> colonization	Not reported
Gao et al., 2019	13.5% ozone	Twice daily for 4 weeks	Ozonated oil may be a safe and efficient treatment	Not reported
Lina et al., 2018	Not reported	Twice daily for 4 weeks	Ozonated oil is safe and effective, equivalent to glucocorticoids topical ointments	Not reported
Hassaniem et al., 2018	Bag: 52 mg/mL of ozone (total volume, 20–50 mL)	Once daily for 30 min for 20 days	Stimulated early wound healing	Not reported
Nowicka, 2017	Ozone at 2.42 mg/min	Twice daily for 10 min for 10 days	Increased joint mobility and skin temperature and decreased its thickness	Not reported
Jian et al., 2018	Ozonated water (3.0 ± 1.5 mg/L); oil not reported	Once daily for 5 days	Reduced pain in herpes-zoster acute phase and shortened virus duration	Not reported
Kasmawati et al., 2020	Bag: 70 µg/mL of ozone	Once every 3 days for 21 days; bag time not reported	Standard treatment combined with ozone therapy reduced the number of bacterial colonies	Not reported
Izadi et al., 2019	Not reported	30 min of ozone bag; topical ozonated oil every 12h; subcutaneous and rectal routes not reported; sessions conducted until complete healing of ulcer: average, 69.44 ± 36.055 days	Efficient as complement to conservative management	Not reported
Solovâstru et al., 2015	15% ozonated sunflower oil (9.6% O ₃)	Once daily for 30 days	Promising therapeutic option with calming and anti-inflammatory effects	Not reported
Zhang et al., 2014	with 52 µg/mL ozone (total volume: 20–50 mL)	Bag 30 min once daily for 20 days	Ozone therapy promoted early healing of ulcer	Not reported
Wainstein et al., 2011	Step 1: 96% oxygen and 4% (80 lg/mL) ozone Step 2: 98% oxygen and 2% (40 lg/mL) ozone.	Step 1: 4 times/week for 4 weeks; step 2: 2 times/week for 12 weeks	Clinical benefit when combined with conservative management	Selection bias; study data collection logistics
Martínez-Sánchez et al., 2005	Bag: 60 mg/L of ozone; oil not reported; systemic use by rectal insufflation: 10 mg ozone and 50 mg/L concentration	Bag: 1 h every day; rectal insufflation for 20 sessions	Therapy considered efficient and may be used as alternative treatment	Not reported
Campanati et al., 2013	Not reported	Once daily for 12 weeks	As effective as hyaluronic acid	Not reported

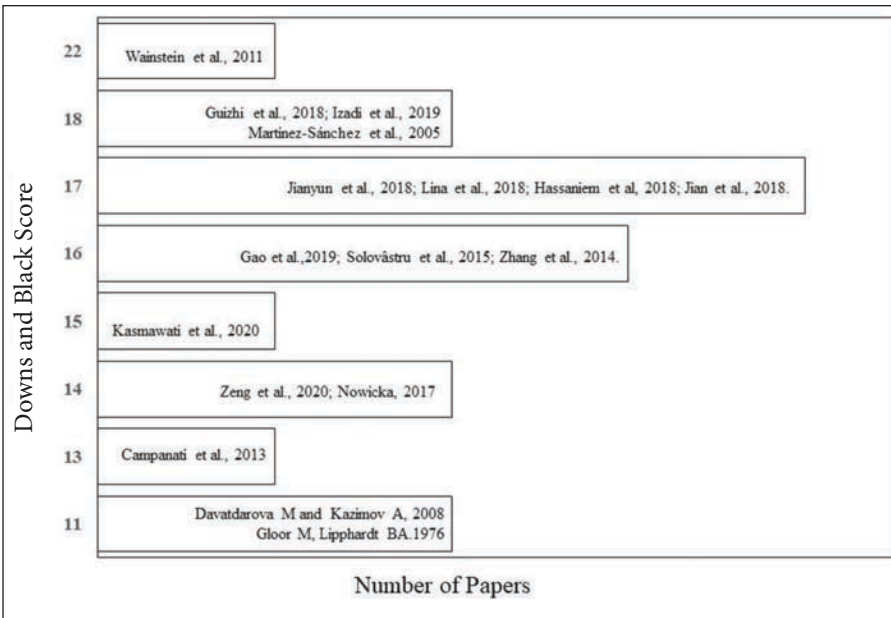


FIGURE 2. Downs and Black scores of methodological quality for the systematically reviewed articles.

the pathological condition¹⁹; and, for ulcers and scarring, it helped with the repair and regeneration processes, increasing blood circulation and combating bacteria and infectious agents.^{20–26}

Methodology quality assessment.

The 17 trials included in this review were assessed to discern the methodological quality using the Downs and Black checklist, which is used for the assessment of methodological quality of randomized and non-randomized trials. The checklist categorizes the quality of trials according to the following points: 26–28 points, excellent; 20–25 points, good; 15–19 points, regular; and 14 points, poor.²⁷

In evaluating the included studies, 5^{12,15,17,25,28} scored 11–14 points and were considered methodologically poor, 11^{12,13,4,16,19–23,25} scored 15–19 points and were considered to have regular methodological quality, and only 1 study²⁴ scored 22 points and was thus considered to have good methodological quality (Figure 2).

The greatest bias risks were observed concerning the absence of information on confusion factors, dispersion estimation, reports of adverse events, loss to follow-up, probability values, alternatives to blind patients and researchers, reports of post-hoc exploratory analysis, types of randomization, and reports of possible protocols of

treatment intent.

DISCUSSION

In spite of being an old therapy, little or no evidence on its use in dermatological conditions can be found nowadays. Through our review research, we observed a somewhat greater foundation for the therapy in other health areas, such as rehabilitation and odontology.^{29–32}

In 2006, Bocci already forewarned of the scarcity of studies on ozone therapy and their concern on the therapy dissemination by unqualified professionals, for whom the most important information is the ozone dosage, instead of understanding how it acts on tissues and what we can do to avoid its toxicity. Moreover, the lack of a precise ozone generator and of dosage standards lead up to the absence of sound scientific and clinical data published under adequate methodological conditions. All this favors the skepticism towards the possible recommendations for ozone therapy.³³

According to Wang, ozone therapy is recent in dermatology and can be a complementary therapy for skin conditions caused by bacteria, viruses, fungi, yeasts, and/or parasites because it acts on the inactivation of these micro-organisms. In such cases, ozone oxidizes the compound phospholipids and lipoproteins of the

cell walls of bacteria; hence, they lose their integrity, and their growth and multiplication are inhibited.³⁴

On skin conditions caused by viruses, such as in herpes, ozone damages the virus structure known as the “viral capsid” via its lipid peroxidation mechanism, which prevents the viral proliferation in the host cells, breaking its reproductive means, while causes cell cytotoxicity in the already infected cells, which may lead to cell death by apoptosis or necrosis.^{32–36}

In the present review, we found studies that reported the use of ozone in treating acne, atopic dermatitis, and ulcers, in which therapy aided in reducing bacterial colonization, such as of *Staphylococcus aureus*, showing the shortening on the duration of bacterial presence, or virus in the case of herpes-zoster, contributing to the repair process and the regeneration of dermic lesions, especially in cases of patients with diabetic ulcers.^{11,12–14,19,28}

For treatment of venous or diabetic ulcers, stimulus of lesion healing can be triggered in varied ways, and, for wounds that do not heal with conventional treatments, ozone therapy may become an alternative.^{20–25}

According to Kushmakov et al., besides its antibacterial and antifungal effects, ozone has the capacity to enhance platelet aggregation. In non-hemostatic functions, platelets are important in inflammation and wound healing; they interact with leukocytes and release vasoactive amines; cytokines; mitogens; and growth factors such as vascular endothelial growth factor, transforming growth factor β , and platelet-derived growth factor. In such a manner, they promote chemotaxis, cell proliferation and differentiation, neovascularization, and extracellular matrix deposition. This process stimulated by ozone aids in the faster healing of wounds/lesions.³⁸

In cases like diabetic ulcers, ozone has an even more specific actuation. When the glycemic state of these patients is not under control, it causes mitochondrial damages in repair cells and even their apoptosis. There is an increase in free radicals and a lower level of antioxidants, leading to vascular damage that hinders the wound healing. In these cases, the ozone’s power of induction for an antioxidant response is beneficial,

for it acts on the balance between glycemic level and free radicals and stimulates a rise in antioxidant levels, enhancing the blood microcirculation and thus increasing oxygen delivery to tissues.^{20–23,37}

Since ozone is a potent inducer for the antioxidant system, this effect helps the physiological process of damaged cells and tissues, for it contributes to the modulation of oxidant and antioxidant systems of the organism, since it also activates the nuclear factor erythroid-derived 2, enhancing the activity of antioxidant enzymes and reducing the levels of pro-inflammatory cytokines. The oxidant molecules, among which is oxygen, play an important role in physiological processes. The reactive oxygen species, also known as free radicals, induce oxidative stress in cells and activate lymphocytes and monocytes and the release of cytokines in circulation and tissues, even stimulating the production of Interferon, which has powerful antiviral activity.^{33–37,38}

Atopic dermatitis and psoriasis both usually present with an imbalance in the skin microbiota that affects immune and anti-inflammatory responses, thus contributing to the severity of the disease. In many cases, there is an association of bacterial colonization that adds to the exacerbation and relapse of these diseases. We have included in this review 3 articles that have used ozone therapy as an alternative treatment for atopic dermatitis and 2 articles that reported its use in psoriasis. Authors have reported that, compared to conventional therapy, ozone may be a beneficial treatment alternative for both diseases; nonetheless, it must be used as complimentary to drug therapy. The reason for its success suggested is that its antibacterial action is associated with the ability to stimulate the balance of the immune system and its potent antioxidant action alters the epigenetics of the disease and generates skin restoration via microbiological regulation of the damaged skin.^{12–14,16}

Another skin condition for which ozone therapy has been being used is systemic sclerosis. This disease has a high morbidity rate, since, besides damaging the skin, it also affects internal organs. Its main characteristic is skin fibrosis with vascular

diseases, which causes ulcers, mainly digital. Researches report that ozone therapy enhances endogenous growth factors, stimulating fibroblasts, increasing then in the microcirculation via the release of nitric oxide and rebalancing the immunological function, favoring the skin restoration and revitalizing the repair and regeneration processes of ulcers and wounds.^{17,18,33,34}

Generally speaking, the articles report that ozone therapy may aid in certain types of dermatological conditions of the skin, for it favors the increase in blood flow with consequent increase in oxygen flow, through stimulus of the oxygen absorbing capacity of the erythrocytes, enhancing their metabolism. Moreover, ozone may balance the immune system and prompt growth factors, cytokines, and chemokines involved in the inflammatory response, modulating pain and altering the cell biology in benefit of its initial characteristic, all due to its antioxidant effect.^{29–35}

However, authors report that evidence on ozone use for the dermatological conditions included in this review is yet extremely doubtful because most trials do not present follow-up studies and do not report in a trustworthy fashion the doses used during therapy. Besides, adverse effects are seldom taken into account.^{24,25,33}

Furthermore, the biological effects of ozone may be dubious, depending on its concentration and site of application. Very low concentrations are practically useless—at best, they act as placebo. Very high concentrations may provoke a negative effect, such as ill-being and fatigue, and even severe adverse effects, like lowering the pulmonary function and dyspnea, among others. In the articles included in this review, some did not report the concentration of ozone when used with water or in the form of ozonized oil. Therapy forms of administration also differed among studies. For the most part, ozone therapy was used as a topical agent, as ozonated oil, immersion in hydrotherapy, or simply with oxygen via the ozone bag. Rectal insufflation, autohemotherapy, or subcutaneous applications were used in few studies. A lack of information on forms of administration and concentration and doses of ozone hinder the efficacy assessment of the resource and

efforts to standardize its use.^{33–36,41}

Our review has shown that evidence on the use of ozone therapy for skin dermatological conditions is scarce. Regarding methodological quality, the findings demonstrate there are few randomized studies and, among the 17 included, all presented diverse methodological aspects. The Downs and Black checklist was used to assess randomized and non-randomized trials. It offers a score for quality of the study and a profile for the research external validity in order to enable understanding on the adequacy of the selected studies.²⁷

Only Wainstein et al.²⁴ scored 22 points on the Downs and Black checklist—that is, only this 1 study was considered to have good methodological quality. The objective of the study was to assess the effect of ozone therapy on the healing of ulcers on diabetic feet. The methodological quality of 11 studies was considered regular and the rest had poor methodological quality.

Studies included in this review presented an apparent lack of information regarding confusion factors, dispersion estimate, adverse effects reports, loss of follow-up, probability values, and patient and researcher blinding alternatives. The data not reported by the studies are considered of utmost importance, and their absence hinders an assessment of the studies regarding therapy recommendation and the qualification of their indication associated with the evidence level.

Considering the information and methodological evaluation of the studies included in this review, it is still not possible to support an evidence level of ozone therapy for treatment of dermatological conditions such as acne, dermatitis, psoriasis, herpes, systemic sclerosis, and aging. This indicates the need for new, methodologically adequate trials with follow-up assessments. For ulcers and scarring dermatological conditions, we have observed that ozone therapy presents greater evidence for its efficacy and safety. However, more studies in these areas are still important.

It is necessary to stress the pertinence of this literature systematic review, in which a yet low number of studies on the subject matter were found. Among the studies

included here, some presented adequate and similar methodology in the application of ozone therapy. It was evident that short-term therapy may even present beneficial results for the dermatological conditions cited in the present study, especially ulcers and scarring processes. However, we point out the limitations of this systematic review related to the lack of information in the follow-up period of the studies after the end of treatment, to assess possible long-term adverse effects, and the methodological quality of the studies, which for the most part did not present standards for ozone therapy application.

CONCLUSION

Based on the above, this review concludes that evidence on ozone therapy for treatment of dermatological conditions of the skin seems promising for short term use, especially for ulcers and scarring. Nonetheless, long-term efficacy and safety still present limitations since the studies do not assess adverse effects or long-term results after the end of treatment. Moreover, all studies included show a scarcity of data regarding therapy, which makes setting standards for ozone therapy for the treatment of the studied conditions more difficult. More research is needed, under better methodological standards, with dosage and ozone concentration reports, equal forms of application for each treatment, and a follow-up period that covers a longer time frame so as to assess the possibility of permanent adverse effects.

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