

Efficiency of gaseous ozone in reducing the development of dry socket following surgical third molar extraction

Jehona Ahmedi¹, Enis Ahmedi², Osman Sejfiija¹, Zana Agani¹, Vjosa Hamiti³

Correspondence: Dr. Jehona Ahmedi
Email: jehona.ahmedi@uni_pr.edu

¹Department of Oral Surgery, Faculty of Medicine, School of Dentistry, Prishtina, Kosovo,
²Department of Prosthodontics, Faculty of Medicine, School of Dentistry, Prishtina, Kosovo,
³Dental Clinic "Identity" Prishtina, Kosovo

ABSTRACT

Objective: The objective of this study was to assess the efficacy of ozone gas (O₃) on the reduction of dry socket (DS) occurrence following surgical extraction of lower jaw third molars, influence of the indication for the extraction, and the difficulty of extraction on the incidence of DS. **Materials and Methods:** This study included thirty patients with bilaterally impacted third molars of mandible requiring surgical procedure for extraction. Following extraction, in the control group, saline solution was used for irrigation of extraction sockets and in the experimental group, intra-alveolar O₃ was applied for 12 s (Prozone, W and H, UK, Ltd.). The surgeries were performed by the same oral surgeon. The follow-up visits were performed at 48 h and on day seven, postsurgery where the symptoms of DS were evaluated and intensity of pain has been recorded using visual analog scale 0–100. **Results:** In this pilot study, DS was present in 16.67% and 3.33% of cases in the control and experimental groups, respectively ($P = 0.20$). **Conclusion:** The application of O₃ may reduce the incidence of DS and accelerates the recovery period after the surgery. Prophylactic use of O₃ may be suggested in all patients, especially in the patients at a risk of development of DS.

Key words: Dry socket, ozone gas, third molar

INTRODUCTION

Dry socket (DS) is one of the most common complications following third molar surgery. Its exact etiology and pathogenesis, despite various approaches to this pathology, are still not known. DS was originally described as a consequence of disintegration of blood clot during days 2–4 after the tooth extraction. Its appearance has been described based on the clinical symptoms, such as a dry alveola covered with a layer of necrotic, yellow-gray tissue, halitosis, and high intensity of pain that radiates to the neck and ear.^[1,2] Although rare, other symptoms

are lymphadenitis, headache, insomnia, and trismus, as described in the literature.^[3,4]

Ozone gas (O₃) is an agent with powerful antimicrobial action.^[5,6] It has helpful use as a disinfectant in the field of medicine and dentistry and has been used widely in all fields of dentistry. Atabas D found that treatment with ozone combined with re-mineralizing solution or either alone is very effective to the initial fissural caries lesions.^[7] In clinical terms, O₃ can be applied in three basic forms such as gaseous, water, and oil for treating

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

For reprints contact: reprints@medknow.com

How to cite this article: Ahmedi J, Ahmedi E, Sejfiija O, Agani Z, Hamiti V. Efficiency of gaseous ozone in reducing the development of dry socket following surgical third molar extraction. *Eur J Dent* 2016;10:381-5.

DOI: 10.4103/1305-7456.184168

Access this article online	
<p>Quick Response Code:</p> 	<p>Website: www.eurjdent.com</p>

various pathologies. Prozone (W and H, UK, Ltd.) is a new technology that produces ozone in a gaseous state for use in various dental applications. It generates O₃ and regulates its flow of concentrations, so they do not exceed the permitted therapeutic values. O₃ has several actions in the human body as an antibacterial, anti-inflammatory, and immunostimulatory agents by utilizing the body's oxygen metabolism and also stimulates the antioxidant humoral system of human organisms.^[8] Its antibacterial effect is based on its ability to form oxidizing free radicals and destruction of microorganisms.^[8] It promotes its antioxidant potential in the destruction of cytoplasmic membranes and cell walls of bacteria, blocking the system enzyme of the cell. This action results in increased permeability of the cell membrane, causing immediate termination of all of its functions and death of microorganisms. This action is selective in microorganisms' cells, but not in human cells because of the human body's greater antioxidant capacity of cells and inhibition, which is vital to stave off the uncontrolled activity of free radicals.^[9,10]

The objectives of our study were as follows:

1. To determine the effect of gaseous O₃ in reducing the presence of DS following bilateral surgical lower jaw third molar extraction compared to the control group
2. Influence of the indication for the extraction on the incidence of DS by groups
3. Influence of the difficulty of extraction on the incidence of DS by groups.

MATERIALS AND METHODS

This pilot study was performed in the Department of Oral Surgery in the University Dental Clinical Centre of Kosovo (UDCCK) which included thirty patients addressed in our clinic for surgical extraction in the period of January 2014–June 2014. The Ethics Committee of UDCCK approved the research protocol. Before the surgery, all patients signed an informed consent for participation in the study.

Inclusion criteria were the following: Age 18–30 years; bilateral impacted lower third molars in similar position verified by orthopantomography; and an indication for surgical extraction (pericoronitis, caries, pulpitis, or orthodontic indication). Exclusion

criteria included the following: Any systemic disease; smokers; immunocompromised patients; pregnant women; and those who were taking contraceptives.

Surgical procedure

Randomization was used to determine which side would comprise the control group (Gr1) and which would comprise the experimental (O₃) group (Gr2). The patients were not aware of these designations. All the surgeries were performed by the same oral surgeon. Local analgesia was achieved using 4 ml of 2% lidocaine with 1:80,000 adrenalin (Alkaloid, Skopje, Macedonia). A triangular, full-thickness buccal flap was elevated for bone exposure. Bone osteotomy was performed for the section of the tooth if required. Afterward, 5 ml of 0.9% saline solution (Gr1) or O₃ gas (Gr2) was used to irrigate the socket. We used Prozone equipment to supply the O₃ gas, which enabled us to introduce the gas into the socket using plastic attachments for 12 s calibrating the therapeutic dose. This procedure was performed using a surgical suction unit to avoid respiratory aspiration and related complication. Suturing was performed with 3/0 absorbable suture (Ethicon, Somerville, NJ, USA).

The patients were then given postoperative instructions and prescribed ibuprofen 400 mg as an analgesic/anti-inflammatory drug. Postoperative visits were scheduled for 48 h after the surgery and on postoperative day 7. Additional appointments were made as needed. Clinical examination was performed to find evidence of one or more of the following main objective signs of a DS: Absence of a blood clot, bone exposure, or necrotic blood clot. We also recorded the number of analgesic tablets ingested.

Subjective clinical findings included patient data of persistent postoperative pain after the surgery. The degree of pain was recorded on a visual analog scale.

Statistical analysis

Data analysis was performed using Pearson's χ^2 test, Fisher's exact test (*P*), and McNemar's test. *P* < 0.05 was considered to indicate statistical significance.

RESULTS

The mean age of the patients was 21.87 ± 4.11 years (95% confidence interval [CI]: 20.33–23.40) [Table 1].

Table 1: Age of all participants in the study Group 1/Group 2

Age	Valid n	Mean	Confidence–95.00%	Confidence+95.00%	Minimum	Maximum	SD
Years	30	21.87	20.33	23.40	18	33	4.11

SD: Standard deviation

The incidence of DS in Gr1 ($n = 5$; 16.67%) was higher than that in Gr2 ($n = 1$; 3.33%), but with no significant difference ($P = 0.20$) [Table 2].

In the Gr1, the incidence of DS was higher in those who underwent very difficult surgery and in those requiring osteotomy with resection during the tooth extraction compared with those without tooth or root resection. The only case of DS in the Gr2 was recorded in a patient whose tooth extraction was very difficult (osteotomy, root, and tooth resection required) [Table 3].

No adverse reactions were recorded during the study implementation.

According to indications for removing the tooth, the frequency of DS was higher in cases of orthodontic versus pericoronitis indications [Table 4].

Table 2: Incidence of dry socket in the Group 1/Group 2

Group	DS, n (%)		Total
	Yes	No	
Group 1	5 (8.33)	25 (41.67)	30 (50.00)
Group 2	1 (1.67)	29 (48.33)	30 (50.00)
Total	6 (10.00)	54 (90.00)	60

DS: Dry socket

The number of consumed analgesics within 48 h after surgery was 3.07 ± 1.96 tablets (95% CI: 2.33–3.80) in Gr1 and 1.20 ± 1.16 tablets (95% CI: 0.77–1.63) in Gr2. The minimum number of consumed analgesics was 0 and the maximum number was 6 in Gr1 and 0 and 4, respectively, in Gr2 [Table 5].

DISCUSSION

Overall complication following third molar surgery rates of about 10%^[11] including pain, secondary infection of the head and neck region, dislocation of third molars, excessive bleeding, temporary or permanent damage to the cranial nerves, and mandibular fracture was reported by Yadav *et al.*^[12] DS is one of the most challenging complications after surgical extraction of lower third molars. The role of bacteria in the development of DS has been discussed in many clinical and scientific studies. This concept is based on research results that confirmed the increased DS frequency in patients with pericoronitis, poor oral hygiene, and periodontal disease.^[13] Potential interference of *Actinomyces viscosus* and *Streptococcus mutans* in DS has been studied by Rozanis *et al.* and it is demonstrated in

Table 3: Influence of the difficulty of extraction on the incidence of dry socket Group 1/Group 2 Difficulty of extraction DS, n (%) T

Difficulty of extraction	DS, n (%)				Total	
	Yes		No		Group 1	Group 2
	Group 1	Group 2	Group 1	Group 2		
Osteotomy	2 (6.67)	0 (0)	18 (60.00)	20 (66.67)	20 (66.67)	20 (66.67)
Osteotomy and root or tooth resection	2 (6.67)	0 (0)	6 (20.00)	8 (26.67)	8 (26.67)	8 (26.67)
Very difficult extraction (osteotomy, tooth, and root resection)	1 (3.33)	1 (3.33)	1 (3.33)	1 (3.33)	2 (6.67)	2 (6.67)
Total	2 (6.67)	1 (3.33)	25 (83.33)	29 (96.67)	30	30

DS: Dry socket

Table 4: Influence of the indication for extraction on the incidence of dry socket Group 1/Group 2

Indication for extraction	DS, n (%)				Total	
	Yes		No		Group 1	Group 2
	Group 1	Group 2	Group 1	Group 2		
Pericoronitis	1 (3.33)	0 (0.00)	9 (30.00)	9 (30.00)	10 (33.33)	9 (30.00)
Caries	0 (0.00)	0 (0.00)	4 (13.33)	4 (13.33)	4 (13.33)	4 (13.33)
Pulpitis	0 (0.00)	0 (0.00)	1 (3.33)	1 (3.33)	1 (3.33)	1 (3.33)
Orthodontic indication	4 (13.33)	1 (3.33)	15 (50.00)	15 (50.00)	15 (50.00)	16 (53.33)
Total	5 (16.67)	1 (3.33)	29 (96.67)	29 (96.67)	30	30

DS: Dry socket

Table 5: Number of tablets consumed within 48 h after surgical extraction Group 1/Group 2

Number of tablets in 48 h	Valid n	Mean	Confidence-95.00%	Confidence+95.00%	Minimum	Maximum	SD
Group 1 - saline group	30	3.07	2.33	3.80	0	6	1.96
Group 2 - ozone group	30	1.20	0.77	1.63	0	4	1.16

SD: Standard deviation

experimental models of delayed socket recovery after inoculation of these microorganisms.^[14] Increasing the number of pyrogenic bacteria as activators of indirect fibrinolysis creates ideal conditions for blood clot disintegration and DS development. Interference of bacteria in the development of DS by introducing preventive antibiotics explains the reduced incidence in many clinical research studies.^[15-18] Akota *et al.* demonstrated a significant reduction in the incidence of DS after the application of chlortetracycline-impregnated gauze compared with a control group.^[19] Nitzan *et al.* observed increased fibrinolytic activity by *Treponema pallidum* which is present in periodontal disease,^[20] explaining why children, who are not colonized by this bacterium, do not develop DS.

The results of our research showed the efficacy of gaseous O₃ in reducing the incidence of DS after surgical extraction of lower third molars compared with the saline solution. Our results also showed that the control patients consumed a large number of analgesic tablets 48 h after tooth extraction compared with the patients who were given O₃, with the difference reaching statistical significance ($P < 0.01$).

Based on our results, intensity of pain following third molar surgical extraction in cases where gaseous O₃ was applied was significantly less than in cases where only saline solution was used for irrigation. This can be explained by the fact that O₃ helps in the synthesis of biologically active substances as leukotrienes, interleukins, and prostaglandins, which are beneficial in reducing inflammation and pain.^[21,22] O₃ activates angiogenesis in inflamed tissue^[23,24] by reacting with blood elements (erythrocytes, leukocytes, platelets, and vascular system). It also has a positive impact on oxygen metabolism, cellular energy, and the antioxidant defense system in the blood microcirculation, thereby improving oxygen delivery to hypoxic tissues by stimulating its metabolism. In addition, it has been found that ozone could accelerate wound healing and bone reparative processes.^[25]

Applying O₃ to the socket following tooth extraction promotes faster healing of the wound without complications.^[26] O₃ also reduce the period of recovery by forming a pseudomembrane in the socket that protected it from mechanical and physical insults.^[27] It has also been proved to be useful in the treatment of refractory osteomyelitis as a complementary therapy to antibiotics and oxygen hyperbaric application^[8,28,29] and promoting the bone healing.^[30]

CONCLUSION

According to our findings, O₃ gas has a positive effect on reducing the development of DS and pain following third molar surgery. The microbiological and metabolic capabilities of O₃ for promoting hemostasis, increasing the supply of oxygen, and inhibiting bacterial proliferation increase the opportunities for its use in all fields of surgery. Further clinical trials are needed with larger samples to support our conclusion.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

REFERENCES

- Swanson AE. A double-blind study on the effectiveness of tetracycline in reducing the incidence of fibrinolytic alveolitis. *J Oral Maxillofac Surg* 1989;47:165-7.
- Fazakerley M, Field EA. Dry socket: A painful post-extraction complication (a review). *Dent Update* 1991;18:31-4.
- Sasaki T, Okamoto T. Topical treatment of infections of alveolar socket infections following dental extraction. *Rev Bras Odontol* 1968;25:82-92.
- Calhoun NR. Dry socket and other postoperative complications. *Dent Clin North Am* 1971;15:337-48.
- Tuncay Ö, Dinçer AN, Kustarci A, Er Ö, Dinç G, Demirbuga S. Effects of ozone and photo-activated disinfection against *Enterococcus faecalis* biofilms *in vitro*. *Niger J Clin Pract* 2015;18:814-8.
- Boch T, Tennert C, Vach K, Al Ahmad A, Hellwig E, Polydorou O. Effect of gaseous ozone on *Enterococcus faecalis* biofilm *in vitro* study. *Clin Oral Investig* 2015 Dec 4. [Epub ahead of print].
- Atabek D, Oztas N. Effectiveness of ozone with or without the additional use of remineralizing solution on non-cavitated fissure carious lesions in permanent molars. *Eur J Dent* 2011;5:393-9.
- Loncar B, Mravak Stipetic M, Matosevic D, Tarle Z. Ozone application in dentistry. *Arch Med Res* 2009;40:136-7.
- Ozone Therapy in Dentistry. Available from: <http://www.absoluteozone.com/ozone-therapy-in-dentistry.html> [Last accessed on 2016 Jun 06]. *Ozone Therapy in Dentistry. J Nat Sci Biol Med* 2011;2:151-3.
- Makkar S, Makkar M. Ozone treating dental infections. *Indian J Stomatol* 2011;2:256-9.
- Valmaseda-Castellón E, Berini-Aytés L, Gay-Escoda C. Inferior alveolar nerve damage after lower third molar surgical extraction: A prospective study of 1117 surgical extractions. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2001;92:377-83.
- Yadav S, Tyagi S, Puri N, Kumar P, Kumar P. Qualitative and quantitative assessment of relationship between mandibular third molar and angle fracture on North Indian population: A clinico-radiographic study. *Eur J Dent* 2013;7:212-7.
- Rud J. Removal of impacted lower third molars with acute pericoronitis and necrotising gingivitis. *Br J Oral Surg* 1970;7:153-60.
- Rozanis J, Schofield ID, Warren BA. Is dry socket preventable? *Dent J* 1977;43:233-6.
- Contar CM, Oliveira P, Kanegusuku K, Silva BR, Azevedo Alanis LR, Machado MA. Complications in third molar removal: A retrospective study of 588 patients. *Med Oral Patol Oral Cir Buccal* 2010;15:74-8.
- Curran JB, Kennett S, Young AR. An assessment of the use of prophylactic antibiotics in third molar surgery. *Int J Oral Surg* 1974;3:1-6.
- Halpern LR, Dodson TB. Does prophylactic administration of systemic antibiotics prevent postoperative inflammatory complications after third molar surgery? *J Oral Maxillofac Surg* 2007;65:177-85.

18. Ritzau M, Hillerup S, Branebjerg PE, Ersbøl BK. Does metronidazole prevent alveolitis sicca dolorosa? A double-blind, placebo-controlled clinical study. *Int J Oral Maxillofac Surg* 1992;21:299-302.
19. Akota I, Alvsaker B, Bjørnland T. The effect of locally applied gauze drain impregnated with chlortetracycline ointment in mandibular third-molar surgery. *Acta Odontol Scand* 1998;56:25-9.
20. Nitzan D, Sperry JF, Wilkins TD. Fibrinolytic activity of oral anaerobic bacteria. *Arch Oral Biol* 1978;23:465-70.
21. Moezizaden M. Future of dentistry, nanodentistry, ozone therapy and tissue engineering. *J Dev Bio Tissue Eng* 2013;5:1-6.
22. Sujatha B, Manoj Kumar MG, Pratap Gowd MJ, Vardhan R. Ozone therapy – A paradigm shift in dentistry. *Health Sci* 2013;2:3.
23. Seidler V, Linetskiy I, Hubálková H, Stanková H, Smucler R, Mazánek J. Ozone and its usage in general medicine and dentistry. A review article. *Prague Med Rep* 2008;109:5-13.
24. Gupta G, Mansi B. Ozone therapy in periodontics. *J Med Life* 2012;5:59-67.
25. Adeyemo WL, Ogunlewe MO, Ladeinde AL, Abib GT, Gbotolorun OM, Olojede OC, *et al.* Prevalence and surgical morbidity of impacted mandibular third molar removal in the aging population: A retrospective study at the Lagos University Teaching Hospital. *Afr J Med Med Sci* 2006;35:479-83.
26. Arita M, Nagayoshi M, Fukuizumi T, Okinaga T, Masumi S, Morikawa M, *et al.* Microbicidal efficacy of ozonated water against *Candida albicans* adhering to acrylic denture plates. *Oral Microbiol Immunol* 2005;20:206-10.
27. Haensler RV. *The Use of Ozone in Medicine*. 4th Edition. Heidelberg: Karl F. Haug Publishers; 2002.
28. Agapov VS, Shulakov VV, Fomchenkov NA. Ozone therapy of chronic mandibular osteomyelitis. *Stomatologiia (Mosk)* 2001;80:14-7.
29. Steinhart H, Schulz S, Mutters R. Evaluation of ozonated oxygen in an experimental animal model of osteomyelitis as a further treatment option for skull-base osteomyelitis. *Eur Arch Otorhinolaryngol* 1999;256:153-7.
30. Alan H, Vardi N, Özgür C, Acar AH, Yolcu Ü, Dogan DO. Comparison of the effects of low-level laser therapy and ozone therapy on bone healing. *J Craniofac Surg* 2015;26:e396-400.