

INFORMED CONSENT FOR OXYGEN/OZONE THERAPY

I,	, do voluntarily, knowingly, and willingly give my consent
to the administration of dental oxy	gen/ozone treatments. I seek this treatment at my own request.
the form of a gas with or withou jawbones, and teeth of the head, n as the creation of a therapeutic biochemical and physiologic cha following dental relevant and use circulatory stimulant, a wound-cl	cone therapy involves the injection of a mixture of oxygen and ozone in t local anesthetic, into the skin, mucous membranes, muscles, joints, neck and associated structures. Dental oxygen/ozone therapy is defined oxygen rich environment, which induces a multi-factorial positive ange in the affected tissues. Dental oxygen/ozone therapy has the eful properties: it kills bacteria, viruses, fungi and parasites. It is a eanser, an accelerant for wound healing, a hemostatic agent, and an may be other effects that at this time are unknown.
	doctor or staff if I have ever had an allergic reaction to any anesthetic, or to any treatment involving injections with anesthetics.
with it some risk of side effects temporary bruising. There may usually heals in a 1-5 day time per allergic reaction to the mixture of site. The most common patient c	th all types of dental treatments. Dental oxygen/ozone therapy carries, such as: pain and/or discomfort at the injection site, soreness and be a red, inflamed, blister-type area at the injection site. This area riod. All types of medications have some risk of allergic reactions. An oxygen/ozone would be unusual, and usually restricted to the injection omment is that there is a warm to burning sensation at the site of the erience flu-like symptoms for 2 to 3 days following treatment.
Patient/Legal Guardian	Date
Witness	Date
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Limitations of Treatment using Dental concentrations of Oxygen/Ozone for the Treatment of the Head, Neck, Face, TMJ, Teeth, and Associated Structures:

I understand with any treatment, there is no guarantee that I will obtain satisfactory results. I may achieve no results, satisfactory results, or unsatisfactory results. If I am currently under the care of a physician or dentist for a known or unknown condition(s), it is my responsibility to inform all practitioners that are providing treatment(s) for my condition(s), of ALL other courses of treatment that I am receiving. Dr. Wall has advised me that it is in my best interest to integrate all therapeutic modalities that are available to treat my health condition(s).

I understand that Dr. Wall is not my primary care physician. I understand that it is in my best interest to have a primary care physician advise me in regard to any treatment(s) that I may choose to receive.

INFORMED CONSENT TO RECEIVE TREATMENT WITH DENTAL OXYGEN/OZONE

I hereby authorize treatment with dental oxygen/ozone and certify that I understand the nature of this treatment, including risks of possible complications and other choices that may be available. I have had any questions concerning this type of treatment answered. I consider myself to be as completely informed as possible and hereby consent to treatment using dental oxygen/ozone. I represent that I am seeking treatment in order to further my own health and for no other reason. I do not represent a third party. I am aware that I may withdraw this consent at any time.

Patient/Legal Guardian	Date
Witness	Date

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Rationale for treatment using dental oxygen/ozone

Additional information and explanation for patients:

Dental oxygen/ozone has been shown to be an effective anti-bacterial, anti-fungal and anti-viral treatment agent. It increases circulation and oxygenation to the treatment area. It increases the immune response and creates an environment for the production of anti-oxidants.

Your disorder(s) may not respond to the treatment(s).	•	
You may experience pain, discomfort, soreness and b	oruising at and around the site of the injection.	
Transient small "bubble-like blisters" may occur at o	r around the site of the injection.	
All medications and treatments have some risk of alle	ergic reaction(s). This is an unusual event and	is
usually restricted to the local area of the injection site.		
The most common side effect of dental oxygen/ozo	one treatment is a warm to burning or stinging	ng
sensation at the injection site.		
This treatment of your head, neck, face, TMJs, tee	eth, and associated structures, will on occasion	or
produce flu-like symptoms, which last on average 2-5 da	ys.	
Any new treatment technique may produce unantic	ipated effects. All known effects to date ha	ve
been explained in this document. If you experience any	reaction not described in this document, plea	S
contact Dr. Judson B. Wall and/or his staff. The office p	hone number is: 801-298-1812.	
You will be notified of any significant new findings,	which are relevant to your treatment.	
Witnessing and Signatures: I		
have read the information and consent forms before sign	ning. I understand that oxygen/ozone therapy	ir
medicine has been used in the United States since 18	385. This therapy has been grandfathered f	ò
medical use prior to the formation of the Food and Drug	g Administration. The FDA has not reviewed	01
approved of statements made in this informed consent. F	Results may vary. No claims are made regarding	ng
the application of any particular therapy using any particular	cular products for any particular reason. I ha	v
been offered ample opportunity to ask questions and h		
satisfaction.		
Patient/Legal Guardian	Date	
Witness	Date	

EXPERIMENTAL SUBJECT'S BILL OF RIGHTS

medical /dental experimental procedure, or who is requested to consent on behalf of another, has the right to: Be informed of the nature and purpose of the experimental procedure. Be given an explanation of the protocol to be followed in the medical/dental experimental procedure and information on the substance or device being evaluated. Be given a description of any attendant discomforts and risks to be reasonably expected from the experimental procedure, when applicable. Be given a disclosure of any appropriate alternative procedures, substances, drugs or devices that might be advantageous to the subject, and their relative risks and benefits, if known. Be given an explanation of any benefits to the subject to be reasonably expected from the experimental procedure, if applicable. Be informed of the courses of medical/dental treatment, if any, available to the subject after the experimental trial if complications should occur. Be given the opportunity to ask any questions concerning the experimental trial or other associated procedures. Be instructed that consent to participate in the medical/dental experimental procedure may be withdrawn at any time, and that the subject may discontinue participating in the medical/dental experiment without prejudice. Be given a copy of the signed and dated written consent form when one is required. Be given the opportunity to decide to consent or not to consent to a medical/dental experimental procedure without the intervention of any element of force, fraud, coercion, or undue influence on the subject's decision. Patient/Legal Guardian Date Witness Date

Any person who is requested to consent to participate as a subject in a research study involving a