



# THE CANADIAN ENVIRONMENTAL TECHNOLOGY VERIFICATION

*Enhancing the Credibility of Environmental Technologies*

## TECHNOLOGY VERIFIED: Ozonator NG-1000 Technology

### Performance Claim

The efficacy of the Ozonator NG-1000 for treating bio-hazardous and regulated medical wastes has been tested employing a surrogate bacterium, *Bacillus subtilis* (*B. atropheus*), and a surrogate waste composed of 10% glass/sharp, 30% plastic, 20% paper, 20% cloth/woven materials, and 20% liquids/organics.

The waste was shredded, loaded into the treatment chamber and ozonated for 900 seconds (or 15 minutes). The ozone concentration in the treatment chamber ranged from 4000 to 8000 ppm; temperature ranged between 17 to 24 °C; and humidity was between 70 and 100%.

Test results show the Ozonator NG-1000 was able to kill *B. subtilis* and reduce its viability by at least 6 orders of magnitude or 6 log<sub>10</sub> with 95% confidence.

### RENEWAL OF VERIFIED\* PERFORMANCE:

July 2016

**Renewal License Number:** ETV 2016-08

**Issued to:** Ozonator Industries Ltd.

**Expiration Date:** July 31, 2019

**John D. Wiebe, PhD**  
Executive Chairman



Canada

\* This verification conforms to the Canadian ETV Program's General Verification Protocol and the ISO/FDIS 14034:2015(E). Please refer to Technology Fact Sheet for additional information on the verification of this performance claim.

## Canadian ETV Verified



### OZONATOR NG-1000

Technology Fact Sheet for Ozonator Industries Ltd.

## Performance Claim

The efficacy of the Ozonator NG-1000 for treating bio-hazardous and regulated medical wastes has been tested employing a surrogate bacterium, *Bacillus subtilis* (*B. atrophaeus*), and a surrogate waste composed of 10% glass/sharp, 30% plastic, 20% paper, 20% cloth/woven materials, and 20% liquids/organics. The waste was shredded, loaded into the treatment chamber and ozonated for 900 seconds (or 15 minutes). The ozone concentration in the treatment chamber ranged from 4000 to 8000 ppm; temperature ranged between 17 to 24 °C; and humidity was between 70 and 100%. Test results show the Ozonator NG-1000 was able to kill *B. subtilis* and reduce its viability by at least 6 orders of magnitude or  $6 \log_{10}$  with 95% confidence.

## Performance Conditions

The performance claim evaluation is based on data and information provided by Ozonator Industries regarding the efficacy of Ozonator NG-1000 for inactivation of a surrogate bacterium, *Bacillus subtilis* (*B. atrophaeus*). Clifton Associates Ltd., an independent third party engineering and scientific firm, conducted tests to determine the effectiveness of the Ozonator NG-1000 for sterilization of bio-hazardous and regulated medical wastes. Ten test strips each containing at least  $1 \times 10^6$  spores were placed in the treatment chamber; eight on eight corners and two in locations along the central axis of the chamber. The chamber was then filled with shredded medical waste that covered all test strips and ozonated with a minimum of 2000 and typically 4000 to 8000 ppm of ozone. The relative humidity of the treatment chamber was between 70 and 100% and humidity was supplied by a water vapour generating system. The tests were conducted with and without ozone in the treatment chamber. The disinfection of *B. subtilis* was calculated using the following relation.

$$X_{uc} - X_t = X_r$$

Where  $X_{uc}$  and  $X_t$  are  $\log_{10}$  concentrations of untreated control (uc) and treated (t) spores and  $X_r$  is  $\log_{10}$  reduction in concentration of spores

The systems was operated in a batch mode. Test strips were located in specific locations inside the Ozonator NG-1000 treatment chamber covered with shredded waste comprising 10% glass/sharp, 30% plastic, 20% paper, 20% cloth/woven materials, and 20% liquids/organics and ozonated between 2000 and 8000 ppm of ozone for 900 seconds.

## Technology Application

Hospitals and health clinics generate bio-medical wastes that have to be managed and disposed of properly in order to protect the general public from pathogens and infectious diseases. Hospital waste can be managed by separating hazardous wastes and placing them in a specially designed container or by treating hazardous wastes and making them non-hazardous.

Treatment processes available for hospital wastes include incineration, steam sterilization, heat treatment, micro- and macro-wave processes, chemical treatment, ozonation, etc. The Ozonator Industries' NG-1000 employs ozone for inactivation of pathogens and infectious disease-causing agents after shredding the waste and reducing its volume by about 90%. The Ozonator NG-1000 is designed to treat bio-hazardous and regulated medical wastes either on-site or off-site.

# Environmental Technology Verification

## Technology Description

The Ozonator NG-1000 is designed to treat bio-hazardous and regulated medical wastes employing ozone. The treatment can be carried out on-site as well as off-site. It comprises a lift, an intake hopper, a shredding chamber, a treatment chamber, and a disposal tank. There are independently movable sealing interfaces on the top of the intake hopper, between the intake hopper and shredding chamber, and between the treatment chamber and the disposal tank.

In typical operation, hospital waste is brought to the device and placed on the cart lift by the operator. After confirming the load is within the weight (200 kg) and volume limits (1 cubic yard), the operator places the cart on the lift, pushes the START button, and the waste processing begins. From this point on, the entire process is computer controlled and the operator has no direct contact with the waste. After the contents of the cart are emptied into the hopper, the ozone generator will be activated, and ozone will be injected into the treatment chamber. Waste then falls onto shredder blades, becomes shredded, and shredded material falls down into the treatment chamber. A water vapour system ensures that the relative humidity within the system is maintained between 75 and 100%. The automated system opens the sealing interface and an exhaust fan draws external air into the hopper, forcing any residual ozone that may remain in the hopper through a Carulite® canister situated in the hopper wall.

## Verification

The verification was conducted by Amiri Clean Water Technologies, using the Canadian ETV Program's General Verification Protocol (February 2007). The verification was based on testing results generated by Clifton Associates Ltd. of Regina, Saskatchewan, and bacterial analysis by SunWest Food Laboratory Ltd., Saskatchewan. The renewal of this verification (July 2013) is issued by GLOBE Performance Solutions. The verification renewal completed in July 2016 also conforms to the ISO/FDIS 14034:2015(E), Environmental management -- Environmental technology verification (ETV).

## What is Canadian ETV?

Canadian Environmental Technology Verification (ETV) is delivered by GLOBE Performance Solutions under a license agreement from Environment Canada. Canadian ETV is designed to support Canada's environment industry by providing credible and independent verification of technology performance claims.

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