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PATENT

Attorney Docket No. 8177-2

METHODS FOR TREATING INFECTIOUS WASTES

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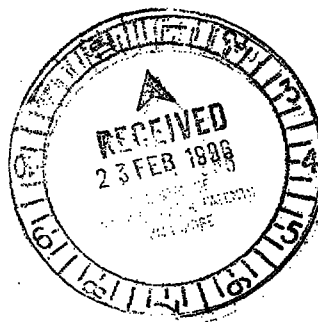
Field of the Invention

The present invention is directed to methods for the treatment of biomedical and other infectious wastes. More specifically, the present invention is directed to methods for processing wastes containing pathogenic
10 microorganisms to substantially reduce or eliminate the amount of pathogens present therein and to generate safer, industrially useful products.

Background of the Invention

Various methods for treating or processing waste
15 containing infectious material, i.e., waste containing various pathogenic microorganisms, are known in the art, and various technologies have been tested in attempts to effectively sterilize such wastes. However, the known, conventional processes have inherent disadvantages which
20 limit their use.

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The currently most widely used method of sterilizing or treating infectious wastes utilize thermal degradation as the primary treatment mechanism. The elevated temperatures used in these processes are
5 generally achieved through dry air, steam or flame in order to sterilize infectious microorganisms by partial molecular degradation at lower temperatures or total molecular destruction at high temperatures.

An example of this type of method is the
10 autoclave, which uses a combination of vacuum and pressurized steam to treat clinical products, such as instruments and containers, as well as to decontaminate post-patient care materials prior to disposal. The limitations of such sterilization systems include the
15 possible escape of aerosolized and liquid borne pathogens through the drain and exhaust ports of the system; the incomplete sterilization of materials containing pathogens resistant to the temperatures used; possible inadequate steam penetration to the center of the waste load; and the
20 continuing requirement of storing, transporting and disposing the treated material, generally utilizing further incineration or landfills. Thus, this technology creates a secondary environmental problem relating to the disposal of the treated waste and is not designed to
25 create any useful by-products from the treated materials.



Another conventional process utilizing thermal technology is incineration, which is presently the most commonly used infectious waste treatment method in the world. However, this technology is now coming under
5 severe environmental criticism, and is subject to strict regulatory standards. In many instances, incineration is being banned entirely because of the likelihood that hazardous products are generated thereby. Thus, new guidelines regarding incineration involve higher standards
10 relating to emissions, such as hydrogen chloride and carbon monoxide, and new requirements relating to dioxins and furans produced from burning complex plastic materials, such as polyesters. In order to comply with these standards, conventional incineration technology must
15 be re-designed significantly, making these treatment methods economically unfeasible. The problems resulting from potential toxic emissions from incineration processes are an additional limitation to the traditional problems of intermediate storage and transport associated with this
20 process, as well as the disposal of the residual ash in landfills.

A third conventional method of treating infectious waste which employs thermal technology comprises heating the waste in large scale microwave
25 systems. Microwave systems are particularly effective for



sterilizing water-based tissue and materials which allow the absorption of microwave frequencies. However, microwave systems have been found to be ineffective for achieving thermal sterilization of dry matter. Moreover, microwave methods do not penetrate materials shielded by metallic enclosures, such as needles, syringes, etc. Such processes are also expensive for large volume treatment and produce disinfected waste which requires further disposal, with its accompanying costs and problems.

10 Waste treatment processes employing the use of chemicals are also known and used in the art. In general, there are two major chemical methodologies utilized in waste treatment systems, utilizing both gas and liquid mechanisms. An example of a gaseous system comprises
15 contacting the waste load with an appropriate amount of a gaseous chemical agent such as ethylene oxide, formaldehyde, peracetic acid and beta-propyl acetone in an appropriate treatment vessel, such as an atmospheric chamber. Ethylene oxide gas is widely used in the
20 sterilization of thermolabile materials, which would otherwise be damaged by exposure to heat and moisture. However, these treatment processes have fallen into disfavor since the chemicals used are considered to be probable human carcinogens and require extreme care and
25 containment during use. These materials also remain



hazardous after treatment and require special detoxification procedures so that their concentration in the treatment area meets safety standards.

Waste treatment methodology employing liquid chemicals comprises the use of chlorine which is a very strong oxidizing agent and reacts in water to form hypochlorite ions. An example of a treatment methodology employing liquid chlorine comprises granulating the waste material and then subjecting the granulated particles to a chlorine spray or bath treatment. The chlorine treatment disinfects the granulated waste material by oxidizing the pathogens. However, such systems have limitations relating to the drying and final disposal of the chlorinated disinfected wastes, the safe disposal of the residual chlorinated treatment solution and the comprehensiveness of the exposure and subsequent inactivation of all pathogens embedded in the waste mass.

More recent, emerging infectious waste treatment processes employ advanced technologies originating from the fields of electronics and physics. Such processes achieve the destruction of infectious microorganisms within the waste by bombarding the waste load with electron beams or electromagnetic radiation. Examples of



such methodology presently in use employ gamma radiation, electron beam radiation and ultraviolet radiation to destroy the pathogens present in the waste load.

For example, gamma irradiation systems utilize
5 powerful radiation originating from radioactive sources
such as Cobalt-60 and Cesium-137. Such systems are
primarily used for the sterilization of medical supplies
and food. However, recent high power gamma irradiation
systems have been designed to treat large volumes of
10 infectious waste materials in continuous conveyerized
facilities. The disadvantages of such systems include the
obvious hazards to personnel which require extra safety
measures, such as shielding of the radiation units, which
ultimately decreases the cost efficiency of these methods.
15 Moreover, the effectiveness of such processes are
dependent on the continual adjustment of exposure
durations in order to accommodate for the continuous decay
of the radioactive material. A more serious problem of
these systems relates to the disposal of the spent
20 radiation sources and the treated waste due to the
existence of trace radioactivity therein.

Another treatment comprises subjecting the waste
material to electron beam radiation utilizing electron
energies exceeding 10^7 electron volts. Commercial linear
25 accelerators are used extensively for the small scale



sterilization of surgical bandages and other disposable medical products. However, the use of electron beam energy for the sterilization and treatment of infectious waste material requires a large scale installation with attendant problems relating to costs and occupational safety procedures. For example, the possibility of workers absorbing low level secondary x-ray radiation remains and the disposal of the treated waste requires full secondary storage, transport and final disposition. Moreover, while this method is generally effective in destroying existent pathogens, the penetration efficiency of the electron beam energies diminishes with distance, density and the presence of metallic shielding in the waste material caused by needles, syringes, etc., thus limiting the guarantee of total disinfection of the treated material.

A third type of radiation technology now being used to treat infectious waste material utilizes ultraviolet radiation. However, it has been found that in general, ultraviolet wavelengths are effective only for the surface treatment of the waste materials and therefore, are not appropriate for the processing of infectious waste materials requiring subsurface



penetration to accomplish sterilization. This characteristic limits the use of ultraviolet radiation for treating large volume mixed infectious waste.

Processes for eliminating or reducing the concentration of pathogens within waste materials to produce fertilizer materials are known in the art. For example, U.S. Patent No. 3,953,191 discloses a process for ridding cotton gin waste of detrimental pathogens and weed seeds to produce a fertilizer. This process comprises chopping or grinding the gin waste and steaming the material by the temperature of 215°F and at constant pressure of 30 psig. U.S. Patent No. 4,743,287 discloses a method of treating waste organic materials to produce a humic acid base fertilizer formulation which comprises granulating the material and then reacting the granulated material with water, acid and a base. A temperature of about 110 to 280°F and a pressure of up to 30 psi is maintained in the reaction vessel. Also, U.S. Patent No. 5,021,077 discloses a method of preparing natural nitrogenous particles useful as plant food by granulating waste material and then heating the material at a temperature of about 50 to 100°C. However, this patent discloses that heating at temperatures higher than 100°C disadvantageously causes the denaturing of proteins in the



material. In contrast to the present method, none of these processes are directed to the treatment of infectious waste.

Accordingly, there is a need for a method for
5 treating infectious waste material to substantially reduce or eliminate the concentration of pathogens therein, which is safe, effective, economical and which produces useful products.

Summary of the Invention

10 The present invention has as its objective to provide a method for effectively treating infectious waste material to substantially reduce the concentration of pathogens therein and which results in the provision of useful products. According to the present invention,
15 waste material containing pathogenic microorganisms is processed by a method comprising: granulating the waste material; treating the granulated waste material by heating at a temperature of about 160 to about 200°C, at a pressure of about 90 to about 226 psi in a non-isotonic
20 atmosphere. The granulated waste material is treated for a period of time sufficient to substantially reduce or eliminate the concentration of pathogenic microorganisms in the waste material. After treatment, the treated material is separated into solid and liquid phases. The



present invention also encompasses useful liquid fertilizers and industrial composite materials which are products of the present method.

Brief Description of the Drawings

5 Fig. 1 shows the main steps of the process according to the present invention in diagramatic form.

Detailed Description of the Invention

The present invention is directed to a method for processing and disinfecting waste containing
10 pathogenic microorganisms. More specifically, the present invention is directed to a method for treating infectious waste materials which contain pathogens, such as those found in hospitals, veterinary facilities, and the like, so as to substantially reduce or totally eliminate the
15 amount of pathogenic microorganisms present therein. By "substantially reduce" it is meant that the concentration of the pathogens is reduced to non-detectable levels using conventional assay methods. In many cases, the pathogens are entirely eliminated from the waste. The thus treated
20 material may then be further processed into useful materials, such as liquid fertilizers, industrially useful composite feedstock material, etc.



The pathogenic microorganisms treatable with the present method appear to be unlimited and include all known human pathogenic microorganisms, generally including viruses, bacteria, fungi, protozoa, etc. Specific

5 examples of pathogens which may be treated with the present method include, but are not limited to:

Brucellosis spp., Campylobacter spp., Clostridium spp., diphtheria, Haemophilus spp., Listeria spp., Meningococcus spp., Bordetella spp., Pneumococcus spp., Salmonella spp.,
10 Shigella spp., E. Coli spp., Yersinia spp., hepatitis A virus, hepatitis B virus, hepatitis C virus, hepatitis D virus, human immunodeficiency virus (HIV), measles virus, mumps virus, rubella virus, varicella-zoster virus, cytomegalovirus, Epstein-Barr virus, herpes simplex virus
15 type 1, herpes simplex virus type 2, human herpes virus type 6.

Although not wishing to be limited to any particular theory, it is believed that the present method substantially reduces or totally eliminates the pathogenic
20 microorganisms present in the waste material through the specific combination of temperature, pressure and atmosphere used. That is, it is believed that the treatment of the waste material at high temperatures and pressures in the presence of a non-isotonic atmosphere
25 denatures the pathogenic microorganisms by altering their



molecular structure so that their outer protective covering is either broken open or altered through osmotic shock and/or biochemical reaction. This acts to either functionally disintegrate the microorganisms and/or

5 structurally alter the genetic information within the microorganisms, so that further multiplication of the pathogens cannot occur. Thus, the original pathogenic entities are broken down to discrete chemical elements, consequently losing their infectious character.

10 The process for treating infectious waste according to the present invention can best be described with reference to Fig. 1. The main steps of the present process comprises granulating the waste material 1; treating the granulated waste material by heating at a

15 temperature of 160-200°C at a pressure of about 90-226 psi in a non-isotonic atmosphere 2; and separating the treated material into solid and liquid phases 3.

In accordance with the present method, the waste material is first mechanically prepared for treatment by

20 granulating the material into particles having a particle size which will facilitate treatment of all of the material with the non-isotonic atmosphere. This is represented by step 1. The particles should be granulated to an extent sufficient to raise the effective surface

25 area of the material to an appropriate degree to allow for



the effective processing of all portions of the material. However, generally, the waste material should be granulated into particles having a particle size of about 0.0625 to about 0.25 inches and preferably about 0.0625 to
5 about 0.125 inches.

The granulation step may be carried out by any suitable device known and used in the art which will granulate the material into appropriate sizes. An example of such a device is a Ball and Jewell granulator
10 manufactured by Sterling, Inc. of Milwaukee, WI. An example of a particularly preferred granulator device useful in the present process consists of knives mounted on rotors which shear the material against knives mounted on the interior walls of a cutting chamber. The size of
15 the particles may be controlled by passing the granulated material through an appropriate base screen upon granulation. The waste material should be granulated for a time period sufficient to process the waste material into fragments and particles of the desired size.

20 After the granulation step, the granulated waste material is treated by heating at elevated temperatures under high pressures in a non-isotonic atmosphere for a suitable period of time. Such treatment of the granulated waste material may be conducted in a suitable pressure
25 vessel. Appropriate pressure vessels should be prepared



in accordance with ASME standards based upon their size and the amount of pressure to be reached in the process. This is represented by step 2 in Fig. 1. Specifically, the granulated waste material is heated at a temperature
5 of about 160 to about 200°C at a pressure of about 90 to about 226 psi in a non-isotonic atmosphere. This treatment is continued for a period of time sufficient to substantially reduce or totally eliminate the amount of pathogenic microorganisms present in the material to non-
10 detectable levels as determined by conventional assay technology, such as the Infectious Unit Assay method.

The treatment of the granulated waste material includes subjecting the material to temperatures which are sufficiently high to inactivate or destroy pathogenic
15 microorganisms present therein. Generally, it has been found that heating the material at a temperature of about 160 to about 200°C will accomplish this objective. Heating the waste material at this temperature is also advantageous as it is below levels at which dioxins and
20 other toxic chemicals may be produced from the waste. Preferably, the material is heated at a temperature of about 170 to about 180°C. The heat can be supplied to the reaction chamber by any appropriate means evident to one skilled in the art. However, an example of an appropriate



heat generating device is a heater band such as that manufactured by Fast Heat Element Manufacturing Co., Inc. of Elmhurst, IL.

The treatment of the waste material in accordance with the present method must take place at a pressure which is elevated to a sufficient degree to ensure the destruction of the outer covering of pathogens present in the material. Generally, it has been found that treating the waste material at a pressure of about 90 to about 226 psi will accomplish the stated objective. Preferably the waste material should be treated at a pressure of about 115 to about 146 psi.

In accordance with the present process, the treatment of the granulated waste material at the temperatures and pressures stated above takes place in a non-isotonic atmosphere. That is, the atmosphere in which the waste is treated has a solute concentration which differs from that known to occur within the microorganisms in the waste. This makes the pressure on one side of the cell membrane different from that on the other side creating a pressure differential across the cell membrane. The non-isotonicity of the atmosphere further aids in destroying or altering the protective outer covering of pathogenic microorganisms present in the waste. To accomplish this result, the atmosphere may be



either hypertonic or hypotonic with respect to the organisms being treated, but in any event must be non-isotonic with respect thereto.

The atmosphere must be non-isotonic with
5 reference to the microorganisms being treated. The
tonicity of the atmosphere is believed to be an important
factor in the destruction or alteration of the integrity
and infectious nature of the pathogenic microorganisms
which require a zero pressure differential across the cell
10 wall membrane to maintain their integrity.

The non-isotonic atmosphere may be generated in
any suitable manner evident to one skilled in the art
based upon the present disclosure and is not limited.
However, it is preferable that the non-isotonic atmosphere
15 comprise the steam from a non-isotonic salt solution.
When employing the steam from a non-isotonic salt
solution, the atmosphere may be generated, e.g., by
loading a hypertonic salt/water solution onto the base of
the treatment vessel prior to heating. Once heating is
20 begun within the vessel, the salt/water solution will
evaporate causing steam from the salt solution to form.
The steam will also be hypertonic, thus providing a non-
isotonic atmosphere (i.e., specifically hypertonic) within
the reaction vessel.



An alternative method of generating the non-isotonic atmosphere in accordance with the present method utilizes an external high pressure steam generator loaded with a non-isotonic salt solution attached to the

5 treatment vessel. Examples of appropriate steam generators useful in the present method are, e.g., those manufactured by Foster-Wheeler, Inc. of Perryville, NJ. The steam generator injects the treatment vessel with preheated steam arising from the non-isotonic salt
10 solution thus providing the necessary atmosphere. This alternative has advantages in that the steam may be injected at appropriate pressures and temperatures to aid in achieving and maintaining the desired pressures and temperatures within the vessel as discussed above.

15 The non-isotonic salt solution which is used to form the non-isotonic atmosphere within the treatment vessel in accordance with the above methods may be prepared from any salt which can provide an atmosphere, which is non-isotonic with respect to the pathogenic
20 microorganisms present in the waste. For example, potassium chloride may be used to form the non-isotonic atmosphere. Other salts which are also believed to be effective as a component of the non-isotonic salt solution to produce the non-isotonic atmosphere include the
25 chloride, sulfate, phosphate, nitrate and carbonate salts



of potassium, sodium, ammonium, magnesium and calcium. However, it is preferred that potassium chloride, or another salt having plant nutrient properties be used.

The non-isotonic salt solution should contain
5 about 20 to about 30 weight % and preferably, about 25 weight % of the salt. Solutions should be provided which will provide an atmosphere having about 1 to about 7% and preferably about 5%, of the salt. The non-isotonic salt solution may be prepared by any standard procedure which
10 will be evident to one skilled in the art.

In order to insure maximum penetration of the granulated waste material by the non-isotonic atmosphere, it is preferred that the infectious waste material be constantly agitated during the entire treatment period.
15 The agitation of the material may be effected by any appropriate mechanical device evident to one skilled in the art from the present disclosure and is not limited. Examples of appropriate agitation devices are those manufactured by Vibration Products Inc. of Wyoming, RI.

20 The granulated material is treated for a period of time sufficient to substantially reduce (i.e., non-detectable levels) or totally eliminate the pathogenic microorganisms within the material. The duration of treatment may vary dependent upon the make-up of the
25 material being treated, particularly the amount of



pathogens contained therein and the amount of material being treated. Appropriate treatment durations for specific samples will be evident to one skilled in the art from the present disclosure. However, it is believed that
5 generally, treatment for a time period of about 20 to about 60 minutes, and preferably about 30 minutes, would be sufficient to disinfect a 500 lb. or greater load of waste material and achieve the objectives of the present method.

10 At the end of the treatment period, the treatment vessel is depressurized. At this time, any accumulated liquids in the treatment vessel may be pumped into a holding tank while the main treatment chamber is opened to allow for the removal of the disinfected,
15 treated waste material. This is represented by step 4 in Fig. 1. These accumulated liquids may be retained for mixture with additional liquids arising from the material due to the further treatment discussed below or may be directly processed as a liquid fertilizer or a component
20 thereof.

As the treatment vessel is depressurized, steam may be generated from the treated material and the treatment system. In order to capture and utilize this steam, a condenser may be attached to the treatment vessel
25 at an appropriate location. The condenser acts to convert



the steam to liquid which may then be pumped into the holding tank discussed above and mixed with other liquids arising from the process. As the condensed steam generally contains at least the salt used to form the non-isotonic atmosphere, it is useful in the liquid fertilizer formed by the present method.

The treated waste material is then further processed to effect the separation of the material into solid and liquid phases. This is represented by step 3 in Fig. 1. This separation may be carried out by various suitable devices as will be evident to one skilled in the art from the present disclosure. For example, conventional centrifuge devices may be used to effectively separate the treated material into solid and liquid phases. Examples of appropriate centrifuges are those manufactured by Bird Machine of South Walpole, MA. This treatment separates the treated material into a solid phase comprising the disinfected solid waste material and a liquid phase generally comprising condensation from the salt-steam atmosphere and an aqueous phase from the waste material comprising highly concentrated organic, inorganic and mineral elements.

The liquid phase may then be removed and combined with the accumulated liquid from the treatment vessel previously removed and discussed above. The



combination of the residual liquids may be performed in an appropriate vessel, such as a blender, as will be recognized to one skilled in the art. This is represented by step 6 in Fig. 1. The resulting liquid blend contains
5 a number of valuable plant nutrients, such as the electrolytic component from the non-isotonic atmosphere and/or any residual non-isotonic salt solution and therefore, is useful for direct application to the soil as a liquid fertilizer. The liquid fertilizer thus produced
10 may be used directly in concentrations correlated to a specific fertilizer analysis, or may be blended with other commercial water soluble chemical fertilizers to achieve a target NPK combination liquid fertilizer.

The remaining solid phase may then be safely
15 discarded or further processed. If the material is to be discarded, it can be done so safely without further treatment. The treated solid phase is environmentally safe and non-toxic and thus, may be discarded with conventional waste.

20 If the solid phase is to be used as an industrially functional composite feedstock product, it may first be dehydrated in order to remove any remaining moisture present therein. The dehydration is represented as step 5 in Fig. 1. Generally, the material should be
25 dehydrated to a moisture content of no more than 10%,



although the final moisture content will vary with the ultimate use of the material. The solid phase may be dehydrated in conventional dehydrators or extruders as would be evident to one skilled in the art from the present disclosure. If the solid material is to be dehydrated, it is generally appropriate to attach a scrubber to the dehydration unit in order to remove any odorous organic materials contained therein.

Once dehydrated, the solid material may then be utilized as industrial composite feedstock and processed into any of various articles of manufacture by known extrusion techniques. This solid phase will generally contain variable percentages of polymeric materials, paper, glass, etc., dependent upon the profile of the original waste being treated. Preferably, the dehydrated solid phase material is treated in accordance with Applicant's methods for preparing composite materials which is the subject matter of copending U.S. Patent Application Serial No. 07/727,176.

Infectious or biomedical waste materials which may be processed with the present method are generally any waste material containing pathogenic microorganisms, such as wastes originating from medical, veterinarian, laboratory and transportation sources which are classified by regulatory bodies as "infectious". The waste material



containing pathogenic microorganisms may include mixtures of all types of common waste including plastics, paper, metal, glass, etc. Examples of the type of material contained in the waste created by the present process

5 include plastic and composite paper packaging; plastic and glass tubes; syringes and utensils; aluminum packaging components; hypodermic needles; paper stationery; fabric materials; bedding textiles and fibers; absorbency products (such as diapers and sanitary napkins);

10 disposable paper; composite drapery; bandages; rubber gloves, tubes and other polymeric materials; surplus medical liquids and organic tissue; residual metal fragments from packaging; waste from diagnostic and surgical procedures and cultures; stocks of infectious

15 agents, etc. Generally, the only materials which are presently believed to be unsuitable for treatment with the present method are radioactive and toxic chemicals whose admixture to the waste material to be treated would result in product materials exceeding regulatory safety

20 standards.

The waste material may be treated directly with the present method without the need for presorting. Thus, the present method can be used to treat materials which are specifically packaged in accordance with regulatory

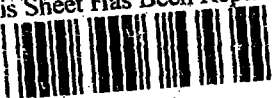
25 guidelines, such as in the form of prescribed coded boxes



and plastic pouches, without separation of the packaging from its contents. This provides a unique advantage over conventional methods of treating infectious waste-containing material.

5 Although the apparatus used to carry out the present process may vary and different appropriate configurations thereof will be evident to one skilled in the art from the present disclosure, it is preferred that the granulator device and treatment vessel be maintained
10 within the same sealed system. That is, the granulator device is preferably maintained inside a sealed vessel designed to insulate the infectious material and associated airborne particles so that no contamination of the external environment can occur. To accomplish this
15 objective, the sealed granulator device should be designed as a subsystem of the total treatment system. Thus, following granulation, the sized particles are transferred to the treatment stage within the same vessel. The entire vessel may thus remain sealed and pressurized during
20 granulation and treatment to ensure disinfection prior to opening the system to reload subsequent batches of waste material therein.

 The present invention will now be illustrated with reference to the following specific non-limiting
25 examples.



Examples

The following examples illustrate the method of the present invention and its efficacy in substantially reducing or totally eliminating the amount of pathogenic
5 microorganisms present in infectious waste material. Each of the examples used the following apparatus.

An apparatus was provided which comprised, as a reaction chamber, a custom pressure vessel designed in accordance with ASME requirements, specifically ASME
10 Boilers and Pressure Vessels Code Section VIII, and which was tested to a maximum pressure of 1200 psi. The reaction chamber was prepared from a stainless steel cylinder having a 1/2 inch thick wall and a 3.5 inch internal diameter. The cylinder was 10 inches high.

15 The reaction chamber further comprised a lid which was also prepared from 1/2 inch stainless steel. The lid was attached to the body of the reaction chamber with eight pressure bolts. A tetrafluoroethylene (TFE) O-ring served as a pressure seal between the lid and the
20 chamber. A heater band manufactured by Fast Heat Element Manufacturing Co., Inc. of Elmhurst, IL was attached to the base of the pressure vessel and electrical power was continuously supplied to the heater band.



A pressure gauge was mounted on the lid to allow the continuous monitoring of the pressure within the chamber. A thermocouple was mounted on the lid of the chamber and served as a means of monitoring the
5 temperature within the chamber and also as a controller for the electrical power required to maintain a constant temperature within the chamber. A pressure relief valve was also provided on the lid of the chamber to guard against the accidental over-pressuring of the reaction
10 chamber.

The reaction chamber further comprised a loading basket to hold the test samples. The basket was made from stainless steel screen having a mesh opening of 1/16 inch. The basket had a diameter of 3.5 inches and was 4 inches
15 in height. The basket was attached to a steel ring allowing it to be maintained at the top of the chamber.

Example 1

Example 1 illustrates the efficacy of the present method in inactivating certain representative
20 bacteria, specifically *Bacillus Stearothermophilus* and *Mycobacteria tuberculosis*. These pathogens were chosen due to their high resistance to most conventional decontamination and sterilization procedures. Moreover,



it is believed that conditions known to inactivate or destroy these pathogens should also inactivate or render innocuous virtually all other known pathogens.

Samples B1 to B6 employed a test procedure required by the Canadian Standards (CAN 3-Z314.3-M79) as a sterility test for all steam sterilization processes. According to this procedure, a standard concentration of *Bacillus stearothermophilus* is contained in nutrient media-based solution in sealed glass or plastic containers. These standard samples contain approximately 1.2×10^5 c.f.u./ml of bacillus. The test vials containing the bacilli were manufactured by Medical Laboratory Systems, 3M Co., MN. Following incubation at 370°C for 72 hours, the color change of the nutrient media based solution from purple to any other color indicates the inactivation of the bacilli present in the invention.

In Samples M1 to M6, *Myrobacteria* was cultured to a concentration of >1000 col./ml in Middlebrook media solution. The cultured solutions were placed in glass Bijou bottles and treated with the present process. After treatment, the samples were tested for the presence of *Mycobacteria* by a culture test as set forth in the Manual of Clinical Microbiology, Ballows A. et al., American



Society of Microbiology, Chapter 34 (1991). All samples (B1 to B6 and M1 to M6) were treated in the manner set forth below.

The reaction chamber was first thoroughly
5 cleaned and rinsed with deionized water. The vials
containing the pathogens (i.e., Samples B1-B6 and M1-M6)
were then placed separately in the treatment apparatus
described above. The vials were placed in either the
loading basket or on the bottom of the reaction chamber as
10 indicated in Tables I and II. Then, 50cc of a 25%
potassium chloride solution in water was poured onto the
bottom of the reaction chamber. The chamber was then
heated to a temperature of 160°C and the pressure within
the chamber was maintained at 90 psi. This treatment was
15 conducted for 20 minutes. After 20 minutes, the chamber
and its contents were allowed to cool to room temperature
and the chamber was depressurized. After treatment,
Samples B1-B6 were observed for color change and then were
incubated at 37°C for 72 hours and observed for color
20 change again. After treatment, Samples M1 to M6 were
incubated at 37°C for 21 days. Cultures of each sample
were taken after the 21 day period and the concentration
of Mycobacteria was determined. The results are set forth
in Tables I and II below.



Table I

Sample	B1	B2	B3	B4	B5	B6
Container Type	Glass	Glass	Plastic	Plastic	Glass	Glass
Initial Color and Turbidity	Clear Purple	Clear Purple	Clear Purple	Clear Purple	Clear Purple	Clear Purple
Location in Chamber	Basket	Bottom	Basket	Bottom	Basket	Bottom
Color and Turbidity After Treatment	Clear Brown	Hazy Brown	Clear Brown	Hazy Brown	Clear Brown	Hazy Brown
Color and Turbidity After Incubation for 72 hours at 37°C	Clear Brown	Hazy Brown	Clear Brown	Hazy Brown	Clear Brown	Hazy Brown

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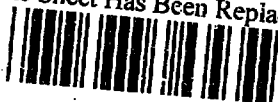


Table II

Sample	Control		M1		M2		M3		Control		M4		M5		M6	
	Location in Chamber	Basket	Basket	Basket	Basket	Basket	Basket	Basket	Bottom	Bottom	Bottom	Bottom	Bottom	Bottom	Bottom	Bottom
Initial Concentration (colonies/ml)	0	>1000	>1000	>1000	>1000	>1000	>1000	>1000	0	0	>1000	>1000	>1000	>1000	>1000	>1000
Concentration After Treatment	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0

From the results set forth in Tables I and II, it can be seen that treatment of the samples containing both the Bacillus and those containing the Mycobacterium with the present method resulted in the destruction of the pathogens to non-detectable levels. The effectiveness of the present method is clearly evident for the Bacillus and both Mycobacteria species.

Example 2

Example 2 sets forth an actual working example of the present process. The reaction chamber of the treatment apparatus was first thoroughly cleaned and rinsed with deionized water. Then, 50cc of a 25% potassium chloride solution in deionized water was poured onto the bottom of the reaction chamber.

The base material specified in each of samples P1 to P9 set forth in Table III was pre-granulated to a particle size of 0.25 inches in a Ball and Jewel granulator manufactured by Sterling Inc. of Milwaukee, WI. Standard stock solutions of Polio Virus in phosphate buffer saline solution were analyzed in order to determine the concentration (in plaque forming units per milliliter) of Polio Virus present in the solutions. The granulated base material was then innoculated with an amount of the Polio Virus solution appropriate to provide the



concentration (pre-treatment) as indicated in Table III. Excess liquid was drained from the granulated base material and the wet, inoculated base material was then placed in the reaction chamber. The material was placed
5 either in the loading basket or on the bottom of the chamber as indicated.

The reaction chamber was then sealed and heated to a temperature of 160°C and maintained at a pressure of 90 psi. During the entire treatment period, the
10 temperature was maintained within 1°C and the pressure was maintained within 1 psi. The examples were treated in this manner for 20 minutes.

At the end of 20 minutes, the chamber and the material were allowed to cool to room temperature and the
15 vessel was depressurized. Once cooled, existent residual liquid and the treated solid material were removed from the reaction chamber and separated. The solid material was then washed in a minimal amount of phosphate buffer solution. The washed solution was then combined with the
20 residual liquid earlier removed.

The combined liquid from each sample was then tested by the standard Infectious Unit Assay procedure which is a standard test for viruses of this type. The full procedure is set forth in detail in the Manual of

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Basic Virological Techniques, G.C. Rovoza and C.M. Burke,
Prentice-Hall, pp. 64-93 (1973). The results are set
forth in Table III.

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Table III

Sample	P1*	P2	P3	P4	P5	P6	P7	P8	P9
Initial Conc. PFU/ml	10^7	10^6	10^8	10^8	10^9	10^9	10^8	10^8	10^7
Base Material		20g Mixture of Metal/Plastic/ and Fiber and Paper**	10g Mixture of Newspaper and Fiber and Paper**	10g Mixture of Newspaper and Fiber and Paper**	10g Mixture of Newspaper and Fiber and Paper**	20g Cloth	20g Cloth	Syringes	Syringe
Measureable Concentration 10^7 of Virus After Addition to Base Material		10^5	10^6	10^5	10^8	10^7	10^7	10^8	10^7
Concentration of Virus in Residual Liquid After Treatment	0	0	0	0	0	0	0	0	0
Concentration of Virus in Solids After Treatment	-	0	0	0	0	0	0	0	0

*Sample P1 was the control. **Tetrapak® Packaging.



As can be seen from the results set forth in Table III, treatment of the granulated waste material inoculated with the Polio Virus results in the destruction of the virus at least to non-detectable
5 levels.

The present invention may be embodied in other specific forms without departing from the spirit or essential attributes thereof and, accordingly, reference should be made to the appended claims, rather than to the foregoing
10 specification, as indicating the scope of the invention.

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CLAIMS

1. A method of processing waste material containing pathogenic microorganisms comprising:

granulating said waste material;

5 treating said granulated waste material by heating at a temperature of about 160 to 200°C at a pressure of about 90 to 226 psi in a non-isotonic atmosphere for a period of time sufficient to substantially reduce or totally eliminate the amount of pathogenic microorganisms in said material; and
10 separating said treated material into solid and liquid phases.

2. A method as in claim 1 wherein said non-isotonic atmosphere comprises the steam from a non-isotonic salt solution.

15 3. A method as in claim 1, wherein said waste material is granulated into particles having a particle size of about 0.0625 to 0.25 inches.

4. A method as in claim 3, wherein said waste material is granulated into particles having a particle size of
20 about 0.0625 to 0.125 inches.





5. A method as in claim 1, wherein said granulated waste material is heated at a temperature of about 170 to 180°C.

6. A method as in claim 1, wherein said granulated waste material is treated at a pressure of about 115 to 146 psi.

7. A method as in claim 1, wherein said material is treated for a time period of about 20 to 60 minutes.

8. A method as in claim 7, wherein said material is heated for a period of about 30 minutes.

9. A method as in claim 2, wherein said non-isotonic salt solution comprises a salt selected from the group consisting of chloride sulfate, phosphate, nitrate and carbonate salts of potassium, sodium, ammonium, magnesium and calcium.

10. A method as in claim 9, wherein said salt is potassium chloride.





11. A method as in claim 2, wherein said non-isotonic salt solution has a salt concentration of about 20 to 30 weight %.

12. A method as in claim 11, wherein said non-5 isotonic salt solution has a salt concentration of about 25 weight %.

13. A method as in claim 2, wherein said atmosphere has a salt concentration of about 1 to 7%.

14. A method as in claim 13, wherein said atmosphere 10 has a salt concentration of about 5%.

15. A method as in claim 1, wherein said separation of said treated material comprises centrifugation.

16. A method as in claim 1, further comprising dehydrating the separated solid phase.

15 17. A method as in claim 1, wherein said solid phase is useful as an industrial composite feedstock material and said liquid phase is useful as a liquid fertilizer.





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18. A liquid fertilizer produced by the process of claim 1.

19. A composite feedstock material produced by the process of claim 1.

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