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**United States Patent** [19]

[11] E

**Patent Number: Re. 36,651**

**Hunniford et al.**

[45] **Reissued Date of Patent: Apr. 11, 2000**

[54] **PROCESS FOR REMOVAL OF DISSOLVED HYDROGEN SULFIDE AND REDUCTION OF SEWAGE BOD IN SEWER OR OTHER WASTE SYSTEMS**

4,148,726	4/1979	Smith .....	210/620
4,153,547	5/1979	McLean .....	210/749
4,297,216	10/1981	Ishida et al. ....	210/631
4,446,031	5/1984	List .....	210/916
4,501,668	2/1985	Merk et al. ....	210/749
4,505,819	3/1985	Barnes et al. ....	210/610
4,680,127	7/1987	Edmondson .....	210/749
4,681,687	7/1987	Mouche et al. ....	210/916
4,725,405	2/1988	Cassin et al. ....	422/7
4,760,027	7/1988	Sublette .....	435/264

[75] Inventors: **David J. Hunniford**, Tallevast; **H. Forbes Davis**, Sarasota, both of Fla.

[73] Assignee: **U.S. Filter Distribution Group, Inc.**, Tallevast, Fla.

**FOREIGN PATENT DOCUMENTS**

[21] Appl. No.: **08/437,874**

3414556	10/1985	Germany .
57-187079	11/1982	Japan .

[22] Filed: **May 9, 1995**

**OTHER PUBLICATIONS**

**Related U.S. Patent Documents**

Reissue of:

[64] Patent No.: **4,911,843**  
Issued: **Mar. 27, 1990**  
Appl. No.: **07/281,747**  
Filed: **Dec. 9, 1988**

“Chemical Control of Hydrogen Sulfide From Anaerobic Swine Manure”, Canadian Agricultural Engineering, vol. 17, No. 2, Dec. 1975.

“Effect of Nitrate on Biogenic Sulfide Production”, Applied and Environmental Microbiology, Jun. 1986, vol. 51, No. 6, pp. 1205–122.

[51] **Int. Cl.**<sup>7</sup> ..... **C02F 3/00**

[52] **U.S. Cl.** ..... **210/610; 210/631; 210/916; 435/264; 435/266**

“Sewage Works Journal”, published by Federation of Sewage Works Associations, Jul. 1946, vol. XVIII, No. 4, pp. 595–640.

[58] **Field of Search** ..... 210/610, 611, 210/620, 631, 916; 435/264, 266, 282

“Industrial Odor Technology Assessment”, Ann Arbor Science Publishers Inc., pp. 136–138, 1956.

[56] **References Cited**

“Odor and Corrosion Control in Sanitary Sewerage Systems and Treatment Plants”, EPA Design Manual, Oct. 1985, pp. 53, 60, 71, 75, 76.

**U.S. PATENT DOCUMENTS**

221,232	11/1879	Hirsh .....	423/210
1,701,825	2/1929	Seil .....	435/266
1,997,252	4/1935	Fischer .....	210/631
3,219,576	11/1965	Makabe .....	71/12
3,300,404	1/1967	Howe et al. ....	210/916
3,401,113	9/1968	Pruessner et al. ....	210/610
3,502,566	3/1970	Raymond et al. ....	210/610
3,607,736	9/1971	Miyaji .....	210/610
3,639,263	2/1972	Troscinski et al. ....	422/14
3,867,284	2/1975	Kappe et al. ....	210/631
3,930,998	1/1976	Knopp et al. ....	210/631
3,959,127	5/1976	Bartha et al. ....	210/610
3,959,130	5/1976	Kloster et al. ....	210/683
3,966,450	6/1976	O'Neill et al. ....	71/15
4,098,690	7/1978	Semmens .....	210/631
4,108,771	8/1978	Weiss .....	210/916
4,115,258	9/1978	Smith et al. ....	210/620

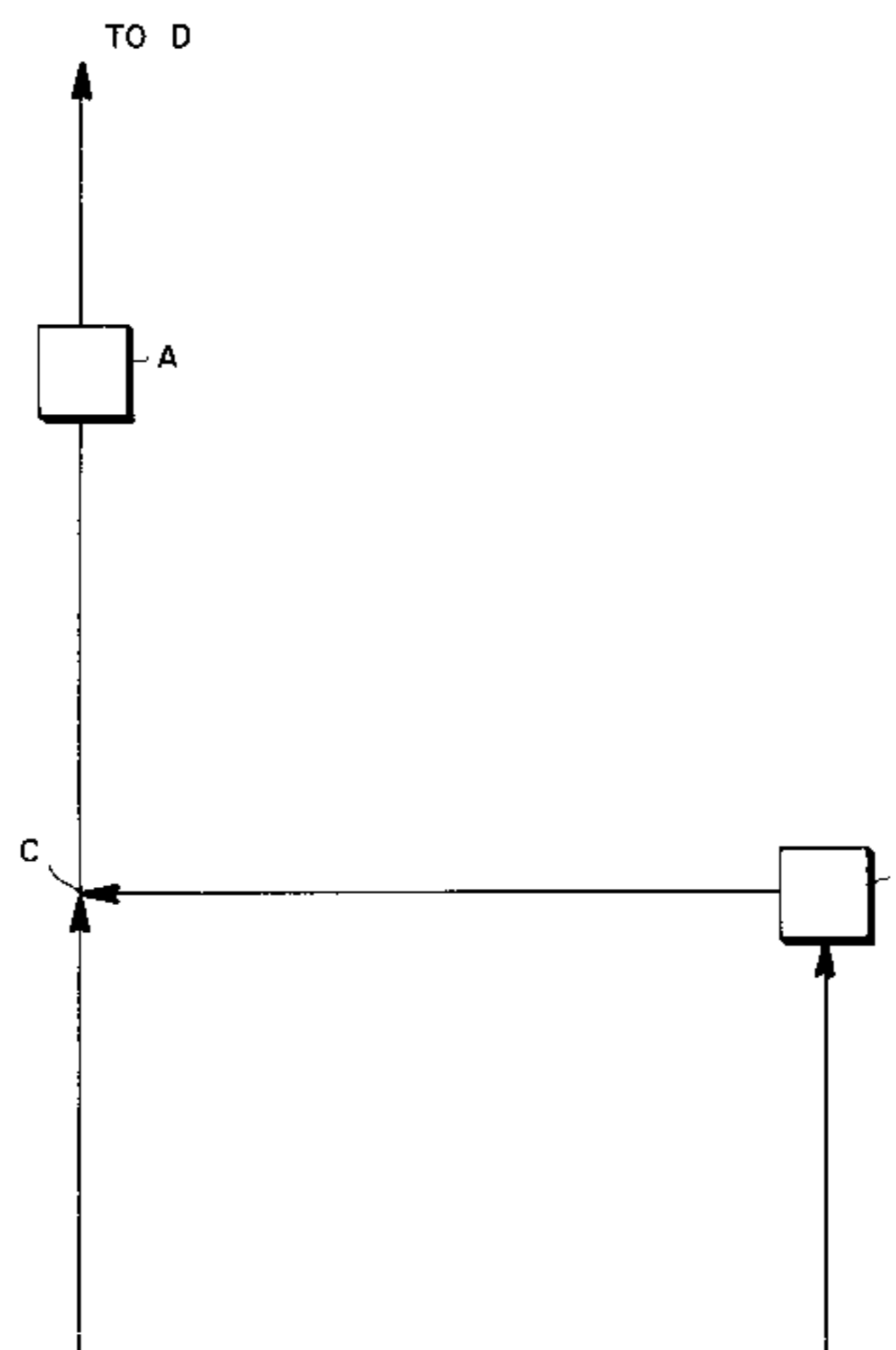
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*Primary Examiner*—Christopher Upton  
*Attorney, Agent, or Firm*—Nixon & Vanderhye P.C.

[57] **ABSTRACT**

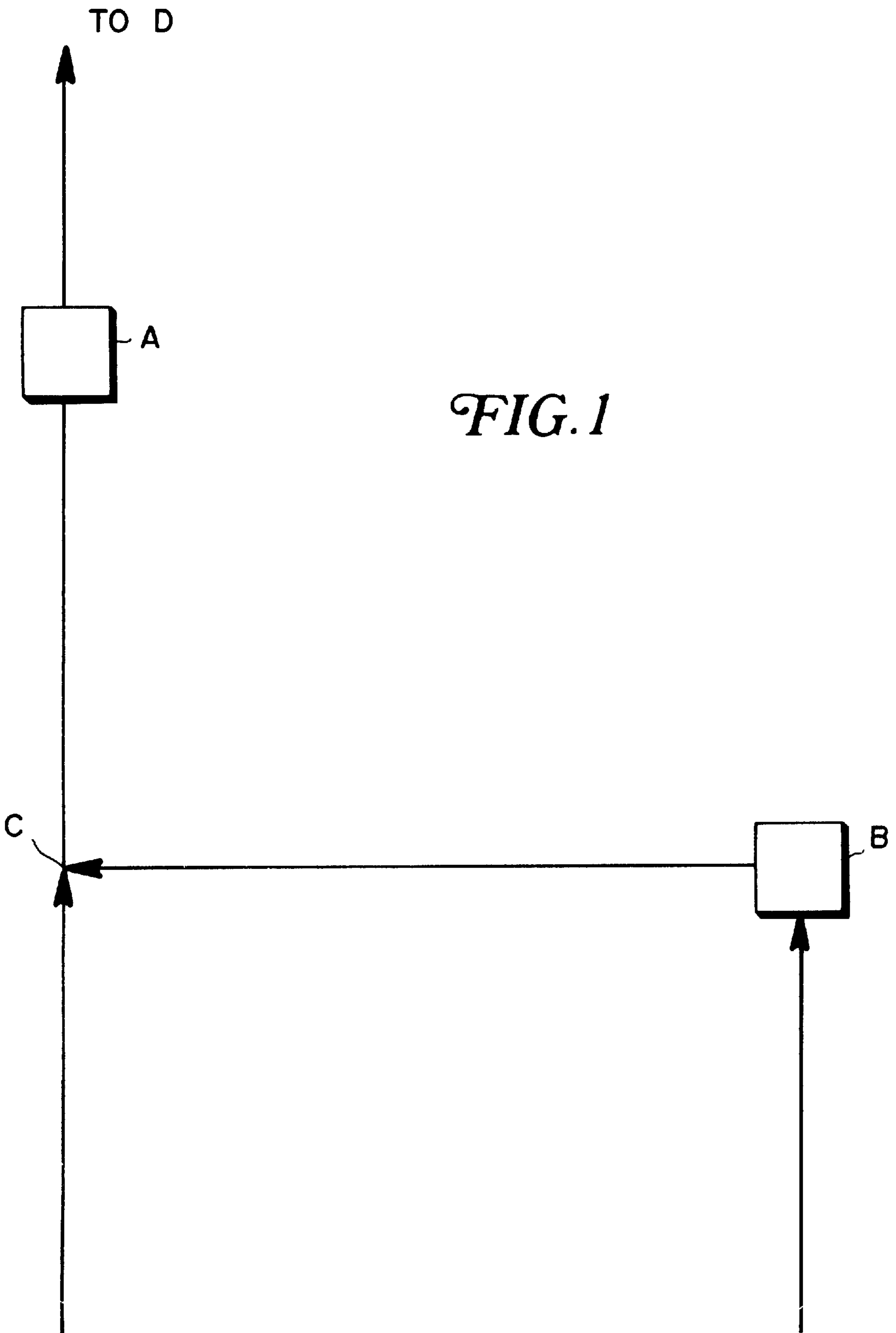
Removal of dissolved hydrogen sulfide and a reduction in BOD is achieved by the addition of nitrate ions to waste systems in an amount sufficient to stimulate growth of bacteria which utilize dissolved hydrogen sulfide in their metabolism. Specifically, about 2.4 lbs. *nitrate* oxygen per lb. of sulfide is required.

**2 Claims, 1 Drawing Sheet**



## OTHER PUBLICATIONS

- “Process Design Manual for Sulfide Control In Sanitary Sewerage Systems”, U.S. Environmental Protection Agency, Oct. 1974, pp. 5–34 and 5–35.
- “Odor Control and Olfaction”, Pollution Sciences Publishing Company, pp. 85, 454, 457, undated.
- Bryan, A.C., “Experiences With Odor Control at Houston Texas”, *Sew. & Ind. Wastes*, 28, 1512 (1956).
- Carpenter, W.T. “Sodium Nitrate Used to Control Nuisance”, *Water Works and Sew.*, 79, 175 (1932).
- Eliassen, R., et al., “The Effect of Chlorinated Hydrocarbons on Hydrogen Sulfide Production”, *Sew. Works Jour.*, 21, 457 (1949).
- Fales, A.L., “Treatment of Industrial Wastes from Paper Mills and Tannery on Neponset River”, *Jour. Ind. Eng. Chem.*, 21, 216 (1929).
- Heukelekian, H., “Effect of the Addition of Sodium Nitride to Sewage on Hydrogen Sulfide Production and B.O.D. Reduction”, *Sewage Works Journal* 15(2):255–261 (1943).
- Heukelekian, H., “Some Bacteriological Aspects of Hydrogen Sulfide Production from Sewage”, *Sew. Works Jour.* 20, 490 (1948).
- Lawrance, W.A., “The Addition of Sodium Nitrate to the Androscoggin River”, *Sew. and Ind. Wastes*, 22, 820 (1950).
- Lorgan, G.P., et al., “Nitrate Addition for the Control of Odor Emissions from Organically Overloaded, Super Rate Trickling Filters”, 33rd Ann. Purdue Ind. Waste Conf., West Lafayette, Ind., (1978).
- McKinney, R.E., “The Role of Chemically Combined Oxygen in Biological Systems”, *Jour. San. Eng. Div., proc. Amer. Soc. Civil Engr.*, 82 SA4, 1053 (1956).
- Poduska, R.A., “Operation, control, and Dynamic Modeling of the Tennessee Eastman Company Industrial Wastewater Treatment System”, 34th Ann. Purdue Indust. Waste Conf., Lafayette, Md. (1970).
- Pomery, R.D., “Controlling Sewage Plant Odors”, *Consulting Eng.*, Feb. 101 (1963).
- Pomery, R.D., et al., “Sulfide Occurrence and Control in Sewage Collection Systes”, U.S. Environmental protection Agency, EPA 600/X–85–052, Cincinnati, Ohio (1985).
- Prakasam, T.B.S., et al., “Microbial Dentrification of a Wastewater Containing high Concentrations of Oxidized Nitrogen”, Proceedings of the 31st Industrial Wste Conference, May 4–6, 1976, Purdue University.
- Price, E.C., et al., “Sewage Treatment Plants Combat Odor Pollution Problems”, *Water and Sew. Works*, 125, 10, 64 (1978).
- Reid, G.W., et al., “Sewer Odor Studies”, *Sew. and Ind. Wstes*, 28, 991 (1956).
- Ryan, W.A., “Experiences with Sodium Nitrate Treatment of Cannery Wastes”, *Sew. Works Jour.*, 17, 1227 (1945).
- Santry, I.W., Jr., “Hydrogen Sulfide Odor Control Measures”, *Jour. Water Poll. Control Fed.*, 38 459 (1966).
- Santry, I.W., Jr., “Hydrogen Sulfide In Sewers”, *Jour. Water Poll. Control Fed.*, 35, 1580 (1963).
- “Standard Methods for the Examination of Water and Wastewater”, 14th Ed., Amer. Pub. Health Assn., Wash. D.C., 499–509 1976).
- Peter R. Willenbring et al., “Calcium Nitrate” (incomplete title), Oct. 1988 or earlier.
- “Biochemical Oxidation of Polluted Lake Sediment with Nitrate—A new Lake Restoration Method”, 1976.
- R.R. Dague, “Fundamentals of Odor Control”, *Journal Water Pollution Control Federation*, 44(4): 583–594; Apr. 1972.
- William H. Moss et al., “Full–scale use of physical/chemical treatment of domestic wastewater at Rocky River, Ohio”, *Journal Water Pollution Control Federation*, 49(11): 2,249–2,254; Nov. 1977.
- Leon S. Directo et al., “Pilot plant study of physical–chemical treatment”, *Journal Water Pollution Control Federation*, 49(1):2,081–2,098; Oct. 1977.
- Richard A. Poduska et al., “Successful storage lagoon odor control”, *Journal Water Pollution Control Federation*, 53(3):299,310; Mar. 1981.
- Pomero, Johnston and Bailey, “Process Design Manual For Sulfide Control in Sanitary Sewage Systems”, Oct. 1974.
- “Basic Research On Sulfide Occurrence and Control In Sewage Collection Systems”, National Technical Information Service, a–5 (Feb. 28, 1969).
- Beardsley, C.W., et al., “Removal of Sewer Odors By Scrubbing With Alkaline Solutions”, *Sewage and Industrial Wastes*, vol. 30, 220 (1958).
- Lang, M., “Chemical Control of Water Quality In A Tidal Basin”, *Journal WPCF*, 1414–1416 (1966).
- Sanborn, N.H., “Nitrate Treatment of Cannery Waste”, *The Fruit Products Journal and American Vinegar Industry*, (1941).
- “Methods For Chemical Analysis of Water and Wastes”, U.S. Environmental Protection Agency, (1974).
- Painter, H.A., “A Review of Literature on Inorganic Nitrogen Metabolism in Microorganisms”, *Water Research, The Journal of the International Association on Water Pollution Research*, vol. 4, No. 6, (1970).
- Pomero, R.D., et al., “Feasibility Study on In–Sewer Treatment Methods”, Municipal, Environmental Research Lab, Chapter 6, “Chemical Treatment”, 77 (1977).
- Thistlethwayte, D.K.B., “The Control of Sulphides In Sewerage Systems”, *Ann Arbor Science Publishers Inc.*, Chapter 13, “Corrective Measures For Existing Systems”, 159 (1972).
- “Water Supply and Sewerage”, Ernest W. Steel, Chapter 27, pp. 600–601 (4th Ed. 1960).



**PROCESS FOR REMOVAL OF DISSOLVED  
HYDROGEN SULFIDE AND REDUCTION OF  
SEWAGE BOD IN SEWER OR OTHER  
WASTE SYSTEMS**

Matter enclosed in heavy brackets [ ] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

*Appl. Ser. No. 08/437,874, filed May 9, 1995, and Appl. Ser. No. 08/198,506, filed Nov. 24, 1998, are copending applications and are each reissues of U.S. Pat. No. 4,911,843, (which issued from Appl. Ser. No. 07/281,747, filed Dec. 9, 1988).*

**BACKGROUND AND SUMMARY OF THE  
INVENTION**

This invention relates to a process for the removal or reduction of dissolved hydrogen sulfide, and reduction of BOD in sewer systems, municipal waste treatment plants and in other industrial waste applications.

It is known to add nitrates or nitrites to sewage to effect reduction in BOD and even to suppress the formation of hydrogen sulfide gas via bacterial action. See, for example, U.S. Pat. Nos. 3,300,404; 4,446,031; and 4,681,687.

It is also known to add nitrates to sewage in order to control objectionable odors. See, for example, U.S. Pat. Nos. 3,966,450; 4,108,771.

There have also been attempts to remove hydrogen sulfide directly from waste. For example, in U.S. Pat. No. 4,680,127, the patentee adds amounts of glyoxal, or glyoxal in combination with formaldehyde or glutaraldehyde, in order to reduce or scavenge the amount of hydrogen sulfide in aqueous or wet gaseous mediums.

In U.S. Pat. No. 4,501,668, the patentee utilizes polycondensation products produced by the condensation of acrolein and formaldehyde to eliminate hydrogen sulfide present in aqueous systems, such as waste water clarification plants. Merk also mentions benefits relating to corrosion prevention and deodorization.

In U.S. Pat. No. 3,959,130, the patentee decontaminates sewage systems, waste water treatment plants and other industrial waste applications containing hydrogen sulfide by adjusting the pH of the sewage of a value over 7.0 and bringing the sewage into contact with an ash product.

It has now been discovered that the addition of nitrate, via an aqueous sodium nitrate solution, to sewage systems, waste treatment plants and other industrial waste applications containing dissolved hydrogen sulfide will result in the elimination or substantial reduction of the hydrogen sulfide, as well as the elimination of other "minor" odors associated with other sulphur-containing compounds.

It is believed that the addition of nitrate provides an oxygen source which promotes the growth of naturally occurring bacteria which utilize in their metabolism the sulfur tied up as hydrogen sulfide. It has been demonstrated both in lab jar tests and in an actual sewage collection system test, that dosing sewage containing over 50 mg/L of dissolved hydrogen sulfide with a sodium nitrate solution reduces the dissolved hydrogen sulfide to less than 0.1 mg/L. Along with this phenomena a significant reduction in sewage biological oxygen demand, BOD, of up to about 70%, and overall "sweetening", i.e., removal of other minor odors, of the sewage has been observed. These phenomena are believed to be the results of the biological process promoted by the nitrate addition.

More specifically, it has been found that 2.4 parts of nitrate oxygen ( $\text{NO}_3\text{---O}$ ) are necessary to remove 1 part dissolved sulfide ( $\text{S}^{2-}$ ). The source of nitrate to accomplish removal of the hydrogen sulfide is not specific, and aqueous solutions of both sodium nitrate and calcium nitrate appear to be suitable.

Because the necessary, reaction is biochemical, it will not occur within a sterile solution, i.e., naturally occurring bacteria must be present. Moreover, the removal of hydrogen sulfide is not instantaneous. According to applicant's tests, an "incubation" period of about 8 to about 96 hours and preferably about 24 to about 48 hours, is necessary to culture the bacteria, followed by about 1.5 to about 20 hours, and preferably about 3 to about 12 hours, for ongoing sulfide removal.

It has further been determined that the process in accordance with this invention achieves a significant reduction in sewage BOD due to the utilization of organic matter in the metabolism described.

Other objects and advantages will become apparent from the detailed description which follows.

**BRIEF DESCRIPTION OF THE DRAWING**

The FIGURE is a schematic diagram representing a sewage system employed in the Example described herein.

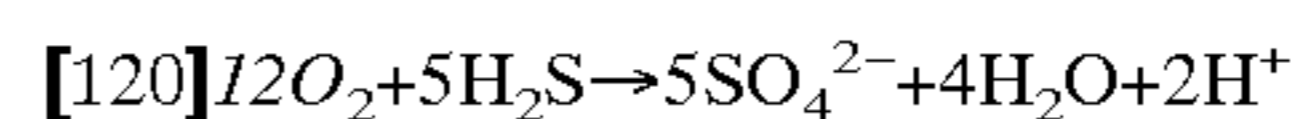
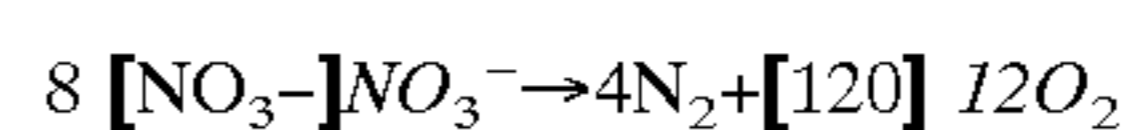
**DETAILED DESCRIPTION OF THE  
INVENTION**

Removal of dissolved hydrogen sulfide and a reduction in BOD in waste systems treated with sodium nitrate or calcium nitrate is believed to occur for the reasons described below.

The presence of dissolved hydrogen sulfide in sewage occurs as a result of a lack of dissolved oxygen. The addition of nitrate ions  $\text{NO}_3$  provides an oxygen source for certain bacteria already present in the waste or sewage to thrive.

The bacteria that grow as a result of the nitrate oxygen utilize the dissolved hydrogen sulfide as part of their metabolism. The dissolved hydrogen sulfide contains sulfur which the bacteria also require in their metabolism.

It is theorized that the biochemical reaction which occurs has the following half reactions:



Based upon the above it is calculated that 2.4 parts of nitrate oxygen ( $\text{NO}_3\text{---}[0]\text{O}$ ) are necessary to remove 1 part of dissolved sulfide ( $\text{S}^{2-}$ ).

$$\frac{8 \text{ moles } \text{NO}_3^-}{5 \text{ moles } \text{H}_2\text{S}} \times \frac{48 \text{ lb Oxygen/mole } \text{NO}_3^-}{32 \text{ lb Sulfide/mole } \text{H}_2\text{S}}$$

yields **2.4 [lb] lbs nitrate** oxygen/lb sulfide.

This ratio of oxygen to sulfide has been confirmed in both bench and field tests.

The source of nitrate to accomplish the sulfide removal is not critical, and both aqueous solutions of sodium nitrate and calcium nitrate have been used successfully.

This reaction is biochemical and it will not occur within a sterile solution, i.e., naturally occurring bacteria in sewage must be present. Additionally, the sulfide removal is not instantaneous; tests have shown that an "incubation" period

of 24–48 hours is necessary to culture the bacteria and thereafter 3–12 hours for ongoing sulfide removal. It is believed, however, that the incubation period may extend from about 8 to about 96 hours, and the ongoing removal period from about 1.5 to about 20 hours, depending on conditions.

The promotion of biological activity via nitrate addition as described also achieves a reduction in sewage BOD due to the utilization of organic mater in the metabolism described.

#### EXAMPLE

With reference to the FIGURE, sodium nitrate was added to a sewer system in Jacksonville, Florida at a master pump station, or feed point B, upstream of a second master pump station comprising a monitoring point A. The feed point B was at a point removed from an intersection C of the feed line and main sewage line, as indicated in the FIGURE.

The treated sewage continued to a downstream waste water treatment plant in Jacksonville, indicated as point D.

Average detention times (based on average daily flows, line sizes and lengths are as follows:

B→C 7 hours

C→A 3.3 hours

B→A 10.3 hours

In terms of the description provided above, the B→C distance and retention time of 7 hours constitutes the incubation period, coupled with the distance C→A and associated retention time of 3.3 hours comprises a total of 10.3 hours from addition of the nitrate station at point B to the monitoring at point A, thereby permitting sufficient time for the bacteria to culture.

The following table shows the change in dissolved hydrogen sulfide at point A, with addition of nitrate occurring at point B.

TABLE I

DATE	SODIUM NITRATE SOLUTION FEED - GPD	DAILY AVERAGE DISSOLVED H <sub>2</sub> S PPM AT POINT A
2/22/88	0	35–40
2/23/88	0	30–50
2/24/88	1800	30
2/25/88	1800	15–20
2/26/88	1800	0.1–15
2/27/88	1200	0.1–4
2/28/88	1200	0.3–4
2/29/88	1200	0.1–8
3/01/88	650	0.7–1.5
3/02/88	650	1.0–1.5

During the period of time, the average daily H<sub>2</sub>S at point B was 25–30 ppm.

It is readily apparent from the above chart that significant reduction in H<sub>2</sub>S was achieved over a nine day period of time, commencing about 24 hours after the addition of the sodium nitrate, with maximum reductions occurring after 48 hours.

Subjective sampling also indicated a significant reduction in sewage odors other than hydrogen sulfide.

It was also found that sewage BOD was also reduced or indicated as in the following table:

TABLE II

DATE	BOD (mg/L)		
	POINT B	POINT A	POINT D
03/02/88	165	112	138
03/03/88	145	55	135

It will thus be appreciated that the present invention provides for the removal of significant amounts of *existing* dissolved hydrogen sulfide and a corresponding reduction in sewage BOD. By properly feeding sodium nitrate into the sewage or waste, odor and corrosion problems can also be substantially eliminated.

While the invention has been described in connection with what is presently known to be the most practical and preferred embodiment, it is to be understood that the invention is not to be limited to the disclosed embodiment, but on the contrary, is intended to cover various modifications and equivalent arrangements included within the spirit and scope of the appended claims.

What is claimed is:

[1. A process for removing dissolved hydrogen sulfide from waste systems comprising the steps of:

(a) adding nitrate ions to the waste in accordance with a ratio of at least 2.4 parts nitrate oxygen for each 1 part dissolved sulfide in order to provide a source of oxygen for naturally occurring bacteria present in the waste which utilize dissolved hydrogen sulfide in their metabolism;

(b) providing sufficient time to culture said bacteria within said waste systems; and

(c) providing ongoing time sufficient to enable said bacteria to remove the dissolved hydrogen sulfide.]

[2. A process according to claim 1 wherein said nitrate ions are provided by the addition of sodium nitrate to the waste.]

[3. A process according to claim 1 wherein said nitrate ions are provided by the addition of calcium nitrate to the waste.]

[4. A process according to claims 2 or 3 wherein a period of from about 8 to about 96 hours is provided in the practice of step (b).]

[5. A process according to claims 2 or 3 wherein a period of from about 24 to about 48 hours is provided in the practice of step (b).]

[6. A process according to claims 2 or 3 wherein a period of from about 1.5 to about 20 hours is provided in the practice of step (c).]

[7. A process according to claims 2 or 3 wherein a period of from about 3 to about 12 hours is provided in the practice of step (c).]

[8. A process according to claim 1 wherein a period of from 8 to about 96 hours is provided in the practice of step (b).]

[9. A process according to claim 8 wherein a period from about 1.5 to about 20 hours is provided in the practice of step (c).]

[10. A process according to claim 8 wherein a period from about 3 to about 12 hours is provided in the practice of step (c).]

[11. A process according to claim 1 wherein a period of from about 24 to about 48 hours is provided in the practice of step (b).]

[12. A process according to claim 11 wherein a period of from about 1.5 to about 20 hours is provided in the practice of step (c).]

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[13. A process according to claim 11 wherein a period of from about 3 to about 12 hours is provided in the practice of step (c).]

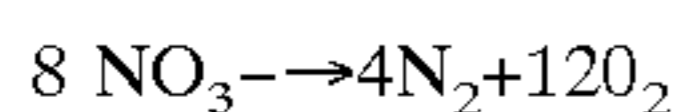
[14. A process according to claim 1 wherein a period of from about 1.5 to about 20 hours is provided in the practice of step (c).]

[15. A process according to claim 1 wherein a period of from 3 to about 12 hours is provided in the practice of step (c).]

[16. A process according to claim 1 wherein sewage BOD is also reduced by up to about 70%.]

[17. A process according to claim 1 wherein said process further eliminates minor odors associated with other sulphur-containing compounds.]

[18. A process for removing from waste systems dissolved hydrogen sulfide and other minor odors associated with other sulphur-containing compounds comprising the steps of: providing a source of oxygen in the form of nitrate in the form of nitrate in the waste in sufficient amount to cause naturally occurring bacteria in the waste which utilize dissolved hydrogen sulfide and sulfur in their metabolism to grow, and providing sufficient time for the bacteria to culture in said waste, to thereby initiate a biochemical reaction which has the following half reactions:



[19. The process according to claim 18 wherein the source of oxygen comprises sodium nitrate.]

[20. The process according to claim 18 wherein the source of oxygen comprises calcium nitrate.]

[21. The process according to claim 18 wherein about 8 to about 96 hours is provided for the bacteria to culture.]

[22. The process according to claim 18 wherein about 24 to about 48 hours is provided for the bacteria to culture.]

[23. A process for removing dissolved H<sub>2</sub>S and reducing sewage BOD in sewer systems comprising the steps of:

(a) adding a source of oxygen in the form of nitrate to the sewer system in an amount equal to about 2.4 lb. oxygen per lb. sulfide; and

(b) providing about 8 to about 96 hours to allow naturally occurring bacteria already present in the system to culture as a result of the addition of said source of oxygen; and providing about 1.5 to about 20 hours to effect ongoing H<sub>2</sub>S removal and sewage BOD reduction.]

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[24. The process according to claim 23 wherein said source of oxygen is sodium nitrate.]

[25. The process according to claim 23 wherein said source of oxygen is a calcium nitrate.]

[26. The process according to claim 23 wherein about 24 to about 48 hours is provided to allow bacteria present in the system to culture.]

[27. The process according to claim 23 wherein about 3 to about 12 hours are provided to effect ongoing H<sub>2</sub>S removal and sewage BOD reduction.]

[28. The process according to 23 wherein, during the process, additional minor odors associated with other sulphur-containing compounds are also eliminated.]

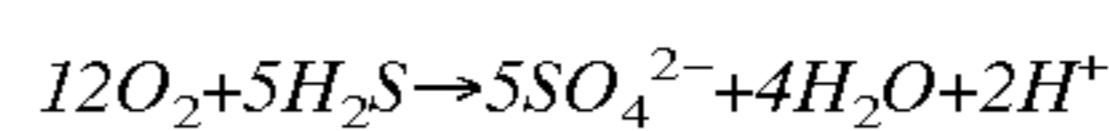
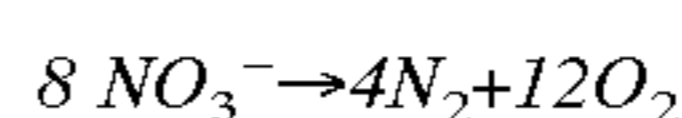
29. A process for removing existing dissolved hydrogen sulfide from waste systems wherein removal is achieved by a mechanism consisting essentially of:

(a) adding nitrate ions to the waste in accordance with a ratio of 2.4 parts nitrate oxygen for each 1 part existing dissolved hydrogen sulfide in order to provide a source of oxygen for naturally occurring bacteria present in the waste which utilize dissolved hydrogen sulfide in their metabolism;

(b) providing sufficient time to culture said bacteria within said waste systems; and

(c) providing ongoing time sufficient to enable said bacteria to remove substantially all said existing dissolved hydrogen sulfide.

30. A process for removing from waste systems existing dissolved hydrogen sulfide and other minor odors associated with other sulphur-containing compounds wherein removal is achieved by a mechanism consisting essentially of: providing a source of oxygen in the form of nitrate in the waste in accordance with a ratio of substantially 2.4 parts nitrate oxygen for each 1 part existing dissolved hydrogen sulfide to cause naturally occurring bacteria in the waste which utilize dissolved hydrogen sulfide and sulfur in their metabolism to grow, and providing sufficient time for the bacteria to culture in said waste, to thereby initiate a biochemical reaction which has the following half reactions:



to thereby remove substantially all of said existing dissolved hydrogen sulfide.

\* \* \* \* \*

UNITED STATES PATENT AND TRADEMARK OFFICE  
**CERTIFICATE OF CORRECTION**

PATENT NO. : RE 36,651  
DATED : April 11, 2000  
INVENTOR(S) : Hunniford et al.

Page 1 of 1


It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Column 2,  
Line 37, delete "NO<sub>3</sub>" and insert -- NO<sub>3</sub><sup>-</sup> --.

Signed and Sealed this

Nineteenth Day of March, 2002

Attest:



Attesting Officer

JAMES E. ROGAN  
Director of the United States Patent and Trademark Office