SOLID WASTE TREATMENT PROCESSES FOR SPACE STATION

By

N86-14091

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### ABSTRACT

The purpose of this study was to evaluate the state-ofthe-art of solid waste(s) treatment processes applicable to a Space Station. Previous studies, since the earlier 1960's, were collected to establish a project library and interviews were conducted of numerous personnel with NASA, from industry, and other researchers. From the review of available information a source term model for solid wastes was determined. An overall system is proposed to treat solid wastes under constraints of zero-gravity and zero-leakage. This study contains discussion of more promising potential treatment processes, including supercritical water oxidation, wet air (oxygen) oxidation, and chemical oxidation. A low pressure, batch-type treatment process is recommended. Processes needed for pretreatment and post-treatment are hardware already developed for space operations. The overall solid waste management system should minimize transfer of wastes from their collection point to treatment vessel.

Center Research Advisor: Chin H. Lin

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### INTRODUCTION

In this report potential processes are investigated for the treatment of solid wastes from the crew and tasks performed in a Space Station. A Space Station has been proposed by NASA as its next major program with an operational Station before the turn of the century. The Space Station would continuously orbit the earth for years, be operated (initially) by a six-man crew, and be resupplied by a Shuttle on about a ninety-day frequency. Resupply of the Station precludes the need for a closed environmental life support system, but the Station will have to treat wastes in order to at least reduce their volume prior to their return to Earth in the Shuttle. It is here assumed that wastes cannot be jettisoned to space from the Station; this is an important constraint to this evaluation of possible treatment systems. The Space Station treatment system will also need to function under zero-gravity conditions. In general, the Station environment will compare to a community with a high population density, limited available utilities (water, electricity, fuel, and air), and operate in a spacecraft with constraints on system size, weight, etc. There is no convenient place for waste disposal and no established recycle options for the waste materials.

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The Space Station solid wastes are: feces, urine, paper trash, food mastes including food containers (plastics), and other

solid wastes from routine maintenance operations and special biological, physical and engineering tasks. These materials might somehow be converted to substances needed on the Station. For example, wastes containing carbon could be oxidized to carbon dioxide that could be reduced to oxygen needed for life supprt. Water in wastes could be separated, filtered, and recycled. Food wastes and metabolic wastes contain nitrogen compounds that could be a source of nitrogen gas needed to maintain cabin atmosphere. These examples do not exhaust the possible uses for solid wates, but are indicative of more likely practical uses for wastes. Furthermore, solid wastes will need to be treated in order to maintain the habitability of the Space Station.

The primary purpose of this study was to evaluate the feasibility and potential of supercritical water oxidation and other waste incineration processes for use on a Space Station.

### THEORY

In order to achieve the project objectives information was retrieved by means of a literature search and by direct
discussions with knowledgeable personnel. At the start of the project only a few reports on supercritical water oxidation were in-hand. A computerized literature search was immediately initiated through the Technical Library of the Johnson Space Center. This search used the NASA/RECON data file. The useful reports were retrieved either in hard copy or microfiche form and

organized into an information file. References cited in the reports were checked and copies of applicable documents were added to the file. A limited "hand" search of current Chemical Abstracts and Science Citations was also done. By these means the solid waste treatment information file was developed as listed in table Footnotes and in References. References were classified into four main topics: Process Reviews, Supercritical Water Oxidation, Zimpro Process, and Treatment Processes.

In addition to the library search, many discussions were held with both NASA and non-NASA personnel. A list of people contacted is at the end of this report, after References. These discussions were very helpful towards the rapid establishment of major problems likely to be encountered and the state-of-the-art of solid waste treatment processes.

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The scope of this study did not include experimental measurements.

On the basis of the in-hand documents and interview data, the available information was evaluated and organized into two major areas; namely, Source-Term (solid waste(s) mass rates and composition) and Treatment Processes.

### RESULTS

The results of this evaluation of Space Station solid wastes and treatment processes are presented below. Results for the Source Term are presented first, then Treatment Proceses.

### A. Source Term

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Metabolic solid wastes from the crew include feces and urine, primarily. Wastes, hair, skin tissue, and nails are relatively small contributions. Toilet tissue paper and wipes are generated along with metabolic wastes but these items are discussed later.

### FECES

The rate, solids contents, and elemental composition of feces are listed in Tables 1 to 3. The reported total fecal rates range from 0.060 to 0.50 kg/man-day. This factor of eight difference can be due to many reasons. Variations in diet, particularly fiber content, is an important parameter. Another important factor is the duration of fecal rates, that is whether over 1-day or 1-monta, etc. There are many other factors to be considered. In Table 2 results are listed from reports of fecal measurements, not second-hand sources which were included in Table 1. The data in Table 2 indicates a range of 0.1 to 0.4 or a factor of four. The results for Skylab (1974) and the McDonnell-Douglas (1971) have the largest experimental basis. This author would weigh + ; data towards the Skylab results which were made in a zero-gravity

	COMMENT	NASA/AMES for CELSS model; Bioastronautics Data Rook	NASAVJSC. ettimated for actronauts	Boeing Aerospace Design Study; Bloastronau- tice hata Book	Hamilton Standard Design Study	Lockheed Design Study	USSR 1-year Test (ground) begun in 1967	Review Study Skylab Summary Data Space Station Design Study - 6 man crev	Operational 90 Day Manned Test (ground), average of 319 uses.	Rockwell; Space Station Waste Model	Design Study	Review for U.S. PHS. Basis 1956 Study (Ingram) of cabin cruiser wastes.	General Dynamics; Ignition ASH @ 1/16 total Mass.	Various diets, compilation of results from 9 sources. General Range.Feces solids contents 15 to 35 % Mixed diet - Wright - Patterson AFB study (1961)
MASS RATE, KG/MAN-DAY	VOTAL	0.104	0.25 to 0.30	0.20	ı	0.125	0.168	- - -	0.393	0.150	0.163	113)0.100 to 0.150	C. 150	0.060 to 0.370 0.35 to 0.50 0.150
MASS RA	WATER	(0.060)	,	0.0	•	ı	ı	0.116 0.12	0.260	0.113	0.113	0.025 to 0.037)(0.075 to 0.113)0.100 to 0.150	s) (0.1125)	0.020 to 0.075 - 0.0235 to 0.035 - 0.061
	SOLIDS	0.044	ı	0.11	0.032	¢	•	0.0338 0.0413 -	. 0.133	0.036	0.050	(0.025	(0.0375)	0.020 t 0.0235 0.061
	AUTHOR	Wydeven	Rapp	Guston and Vinopal	Brose	Jagow	Jones	Bioenvironmental Systems Sauer, R. Nelson and Cody	McDonnell-Douglas Astron.	Schaedle and Laubach	Hamilton Standard	Rich, Ingram and Berger	Dodson and Wallman	We Jb
	LAK	1983	1983	1982	1981	1976	1975	1975 1974 1972	161	1970	1969	1965	1964	1964

Unspecified diet - Wright - Patterson AFB study (1962)

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NOTE: Numbers in parenthesis calculated in this work.

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### REFERENCES

Dodson L. and Wallman, H., "Research on a Waste System for Aerospace Stations," Technical Documentary Report No. AMRL-TDR-64-33, 1964.

Webb, P., Ed., Bioastronautics Data Book, National Aeronautics and Space Administration, NASA-SP-3006, section 13 pp. 213-239, \*964.

Rich, L.H., Ingram, W.M., and Berger, B.B., "Waste Disposal on Space Craft and Its Bearing on Terrestrial Problems," U.S. Department Health, and Welfare, Public Health Service, PB 168787, 1965.

Hamilton Standard, "Alternate Mission Studies (AILSS)," NASA Contractor Report-66876, 1969.

Fairchild Hiller-Republic Aviation Division, "Housekeeping Concepts for Manner Space Systems," Document No. MS 124 Y0002, Vol. II Prepared for NASA, 1970.

Schaedle G.C. and Laubach G.E."An Introduction to the Waste Management Problem for Large Space Stations," American Society of Mechanical Engineers., Proceedings Space Systems and Thermal Technology for the 70's Part 1, Los Angeles, CA June, Paper 70-Av/SpT-24, 1970.

McDonnell-Douglas Astronautics "Test Results, Operational Ninety-day Manned Test of a Regenerative Life Support System," NASA Contractor Report 111881, 1971.

Nelson, W.G. and Cody J. "Life Support System Definition for a Low Cost Shuttle Launched Space Station," American Society of Mechanical Engineers, Publication 72-ENAv-17, 1972.

**Bioenvironmental** Systems Study Group of the Society of Automotive Engineers, "Evaluation and Comparison of Alternative Designs for Water/Solid-Waste Processing Systems for Spacecraf,," Final Report, NASA-CR-162492, 1975.

Jones, W.L., "Life-Support Systems for Interplanetary Spacecraft and Space Stations for Long-Term Use," in Foundations of Space Biology and Medicine, Calvin, M.and Gazenko, O.G., Eds., National Aeronautics and Space Administration, Vol. III p.270,1975.

Brose, H.F., "A Regenerative Life Support System for Space Operations Center (SOC) - A Probable First Flight Application," American Society of Mechanical Engineers, Paper 81-ENAs-12, 1981.

Guston, E. and Vinopal T., "Controlled Ecological Life Support System Transportation Analysis," NASA Contractor Report - 166420, pp. 48-51, 1982.

Rapp, R., Personal Communication, 1983.

Wydeven, T., "Composition and Analysis of a Model Waste for a CELSS," Preprint, to be published as a NASA Technical Memorandum, 1983.

Reference Addes in Proof Saver R.L., "Summary Sky Lab Intake-Output Cata Sheet, Personal Communication 1983.

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TABLE 2. FECAL MASS RATE - EXPERIMENTAL VALUES

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YEAR		AUTHOR	RATE, KG/MAN DAY			
	•		TOTAL	SOLIDS	SOLIDS %	
	1983	Rapp	0.25 to 0.30	-	-	
•	1981	Onisko and Wydeven <sup>a</sup>	-	-	20	
	1971	McDonnell-Douglas (90 dry ground study)	0.40	0.13	33.8	
	1975 1974 1964	Jones (USSR data of 1967) Sa <b>u</b> er Webb	0.17 0.16 0.35 to 0.50	).041 -	26 15 to 35	,
		Webb (Wright-Patterson AFB, 1961)	0.15	0.061	40.7	
		Webb (Wright-Pattersc AFB, 1962)	0.12	0.020	16.7	
•	1965	Rich, et al.(Ingram, 1956)	C.10 to 0.15	0.025 to 0.037	25	

a. Onisko, B. L. and Wydeven, T., "Wet Oxidation as a Waste Treatment Method in Closed Systems," NASA Conference Publication 2247, pp. 51-53, 1981.

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LEMENT		88	C	<u>D</u>	<u> </u>
C	41.92	41.92	26.3	67.73	-
0	-	-	34.6	11.71	11.71
N	8.26	8.26	3.92	4.00	6.0
н	6.59	6.59	6.98	11.71	-
x	2.8	2.8	-	-	0.88
Ca	2.5	2.5	-	-	3.5
α	2.1	2.1	-	•	-
Na	1.8	1.8	-	-	0.34
P	1.4	-	0.99	-	-
Mg	0.66	0.66	-	-	0.77
S	-	-	0.39	0.28	
Fe	0.043	0.043	-	-	0.087
Si	0.040	0.040	0.20	-	-
Zn	0.027	0.027	-	-	-
Mn	9.017	0.017	-	-	0.010
Cu	0.0040	0.0040	-	-	0.0032
8	0.0015	0.0015	-	-	-
۷	0.0006	0.0006	-	-	-
ASH	-	-	9.34 (DRY)	4.53 (DRY)	3.1

### TABLE 3. FECAL SOLIDS ELEMENTAL AND ASH COMPOSITION REFERENCE

A. Wydeven, T. 1983, op. cit.

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B. Carden, J.L. and Browner, R., "Preparation and Analysis of Standardized Waste Samples for Controlled Ecological Life Support Systems (CELSS), "NASA Contractor Report 166392, 1982.

C. Onisko, B. L. and Wydeven, T., 1981, op. cit.

D. Bioenvironmental Space Systems Study Group, 1975, op. cit.

E. Goldblith, S. A. and Wick, E.L., "Analysis of Human Fecal Components and Study of Methods for Their Recovery in Space Systems, Aerospace Medical Laboratory, Wright-Patterson Air Force Base, 1961.

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environment for about 120 days by nine different astronauts. The McDonnell-Douglas study was in a 1-g-gravity environment for 90 days by four men. The average of these two programs is 0.28 kg/man-day and for a weighted average take 0.25 kg feces per man per day. This average agrees well with Rapp's (1983) estimate of 0.25 to 0.30 kg/man-day.

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The total solids content of feces varies from about 15 to 40 percent. This author takes an average value of 25% which is above the Ames data, below the McDonnell-Douglas results, and in agreement with results by Ingram (1956) and Skylab (Sauer). Data for the elemental composition of feces was not readily available, see Table 3. Rough estimates of the carbon, hydrogen, oxygen, and nitrogen levels are 26%, 7%. 35%, and 4%, respectively. These concentrations are basically values reported by NASA/Ames (Onisko and Wydeven, 1981).

On the basis of the above average fecal mass rates and composition, the elemental and water mass rates can be calculated. The results are as follows:

		Rate, kg/man-day
Carbon		0.016
Hydrogen		0.0044
Oxygen	1	0.022
Nitrogen		0.0025
Water		0.19
		14.30

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The total energy available from the burning of feces, assuming a heat of combustion of 7700 Btu/lb (18,000 kJ/kg) and a fecal rate of 0.25 kg/man-day, was calculated to be 5 x  $10^{-5}$ kilowatts per man-day. Thus, feces is not a practical energy source by any incineration or oxidation process.

### URINE

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The urine mass rate and elemental composition data are presented in Tables 4 and 5, respectively. Variations in the urine rates are not as great as the fecal data. This author takes an average value of 1.6 kg per man per day. The Skylab average equals 1.58 kg/man day. During the Mercury-Apollo flights (6, 7 and 9) the urine rates varied considerably from 0.7 to 3.5 kg/man day, but the average value of these 3 flights is 1.8 kg/man day which is reasonable agreement with Skylab data. The solids content of urine is 5% based on the experimental data. Thus, from urine 1.52 kg of water per man day are available for recycle. The elemental composition of solids in urine is taken as follows:

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TABLE 4. URINE MASS RATE<sup>a</sup>

MASS RATE, KG/MAN-DAY TOTAL WATER SOLIDS YEAR AUTHOR 1.56 0.059 1.50 1981 Brose 2.18 1976 Jagow 1.28 1975 Jones 0.064 Bioenvironmental Systems Study 1975 1.58 1974 Sauer 1.64 1972 Nelson and Cody 1.5 McDonnell-Douglas Astronautics 1971 1.63 1.57 0.059 1970 Schaedle and Laubach 0.70 1.4 0.70 Hamilton Standard 1969 not available Shook and Thomas 1969 1.2 0.06 1965 Rich, Ingram and Berger 1.2 1964 Webb Avq. 1.3 Mercury-Apollo Flt. 6 3.5 Mercury-Apollo Flt. 7 0.7 Mercury-Apollo Flt. 9 1.5 1964 Dodson and Wallman 0.075 1.40 1.33 1962 Mattoni and Sullivan 0.070

a. Footnotes are listed in Table 1 plus the following:

Mattoni, R.H. and Sullivan, G.H., Sanitation and Personal Hygiene During Aerospace Missions," Wright-Patterson Air Force Base, Ohio, Technical Documentary Report No. MRL-TDR-62-68, 1962.

Dodson, J. and Wallman, H., "Research on a Waste System for Aerospace Stations," Wright-Patterson Air Force Base, AMRL-TDR-64-33, 1964.

Shook, R.E. and Thomas, E.C., "Urine Output Parameters for Space Cabin Environments," McDonnell-Douglas Astronautics Company, Paper 10069, April 1969.

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# TABLE 5. URINE ELEMENTAL COMPOSITION AND DRY WEIGHT

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- A. Tabulation excludes extensive data compiled by Webb, 1964; these data are rates (mg/24 hr) of numerous compounds as well as their range.
- B. Wydeven, T., 1983; op. cit.
- C. Carden, J. L. and Browner, R., 1982, op. cit.
- D. Onisko. B. L. and Wydeven, T., 1981, op. cit.
- E. Hoshizaki, T. and Hansen, B.D., "Generic Waste Management Requirements for a Controlled Ecological Life Support System (CELSS), American Society of Mechanical Engineers, Paper 81-ENAs-23, 1981.
- F. Bioenvironmental Systems Study Group, 1975, op. cit.
- G. Putnam, D.F., "Composition and Concentrative Properties of Human Urine," NASA Contractor Report 1802, 1971.

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carbon - 18%, hydrogen 4.9%, oxygen 25%, nitrogen 21%. Other elements are not included here, for additional details see Table 5. The elemental and water release rates from urine are calculated to be as follows:

	kg/man-day
Carbon	0.014
Hydrogen	0.0039
Oxygen	0.020
Nitrogen	0.017
Water	1.52

#### FOOD

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The wastes generated from food include uneaten foods and used food containers. On the basis of Shuttle data 3.6 pounds (1.64 kg) of food (dry food, water in food, and packaging) are provided per person per day. This includes 0.73 kg of dry food per person per day and 0.46 kg of water. Not all the food is eaten. Assume 10% of the food is wasted; then 0.046 kg of water per person per day are available for recovery. If food is assumed to approximate sugar  $C_6H_{12}O_6$ , which is a crude assumption, but useful for the determination of elemental release rates. The carbon, hydrogen and oxygen available from wasted food are estimated to be as follows:

	kg/man-day
Carbon	0.03
Hydrogen	0.005
Oxygen	0.04

The amount of carbon, etc. from wasted food containers is discussed under plastics.

### PAPER

Paper wastes would be generated on a Space Station due to personal hygiene requirements, used paper towels, wipes, gauze, Q-tips and midcellaneous other paper products. These wastes exclude washcloths, fecal-emesis collection bag, and a trash container liner spare. The total amount of cellulosic materials is estimated to be 1 kg per man-day. The empirical formula for cellulose is  $C_6H_{10}O_5$ , or 44% is carbon, 6% is hydrogen and 50% is oxygen. This estimate neglects th presence of water. The elemental release rates are calculated:

	kg/man-day
Carbon	0.44
Hydrogen	0.06
Oxygen	0.50

### PLASTICS

Wastes composed primarily of polyethylene are generated from food containers, packing materials, medical supplies, and from experimental work. Food containers are used at a rate of about 0.45 kg per man-day. Packing materials and other plastics could be triple this amount, or 1.4 kg per man-day. This author assumes that the plastic wastes consist mostly of polyethylene. From plastics the elemental release rates are:

kg/man-day Carbon 1.6 Hydrogen 0.26

The total wastes generated are summarized in Table 6, and a summary of the elemental release rates and water are presented in Table 7. The total solid waste feed rate is 5.4 kg per man day, plastics will constitute the largest amount followed by food and urine. The weight of paper wastes is less than these latter three wastes, but its bulk volume will be the largest. Feces amounts to about five percent of the total mass. These results indicate that water is available in wastes (1.8 kg per man-day) to provide the drinking needs for 2 astronauts per day. Carbon oxidation would require 7.5 kg of oxygen per man-day because the amount of

TABL	E 6
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# SOLID WASTE(S) TREATMENT SYSTEM FEED WASTES

WASTE		<u>RATE</u> kg/man-day
FECES		0.25
URINE		1.6
FOOD		1.6
PAPER		1.0
PLASTICS		1.9
	TOTAL	5.4

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Element/ <u>Waste</u>	Carbon	Hydrogen	<u>Oxygen</u> rate, kg per per	<u>Nitrogen</u> rson per day	Water	
Feces	0.016	0.0044	0.022	0.0025	0.19	
Urine	0.014	0.0039	0.020	0.017	1.52	
Food	0.03	0.005	0.04	-	0.046	
Paper	0.44	0.06	0.50	-	-	
Plastics	1.6	0.26				
TOTAL	2.1	0.33	0.58	0.020	1.8	

### TABLE 7

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### SUMMARY OF ELEMENTAL AND WATER WASTE RATES

oxygen in the wastes is insufficient for complete combustion. Relatively small amounts of nitrogen are available for utilization a Space Station.

Table 8 presents the heat of combustion of waste materials.

### **B.** TREATMENT PROCESSES

There are many possible treatment processes for Space Station solid wastes. Some are conventional technology, for example, incineration and others are high technology processes, like electric discharge plasma and supercritical water oxidation. This report will Jiscuss only a few of the processes. Information about others can be obtained from references cited in this report.

### SUPERCRITICAL WATER OXIDATION

In 1975 Modell discovered that supercritical water would oxidize specific organic compounds. Supercritical water is water at temperature and pressure levels above water's critical point, 374 degrees C and 218 atm, see Figure 1. In 1979, supercritical water oxidation (SCWO) was applied to mixtures of organic wastes. Tests were conducted to show its effectiveness in destroying organic halides which are now commonly known as hazardous wastes. The exploitation of the SCWO process was commercialized in 1980 by Modell who formed Modar, Inc.

# TABLE 8.

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HEATS OF	COMBUSTION		
	<u>Btu</u> Lb	<u>kJ</u>	
Cellulose	7500	17,000	
Feces (dry)	7700	18,000	
Polystyrene	17,200	40,000	
Polyethylene	19,000	44,000	

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In the SCWO both organic material and oxygen dissolve in supercritical water and then oxidation is carried out in supercritical water medium. Organic substances and gases are completely soluble in supercritical water but inorganic salts exhibit greatly reduced solubilities. This is just the opposite effects of solubilities at normal conditions. Thus oxidation may be carried out in a homogenous system and salts separated. Figure 2 is a schematic of the SCWO process. The supercritical water reactor is described in Figure 3.

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The SCWO process can be described in five steps:

(1) The solid waste is slurried with make-up water to provide a mixture of about 5 to 10 weight percent organics. The slurry is pressurized and heated to supercritical conditions. Heating is attained by mixing the feed with superheated SCW, which is generated in a subsequent step. During the period outside the oxidizer, organic materials in the feed are converted to combustible gases, low to intermediate molecular weight compound: (alcohols, aldehydes, furans) and inorganic salts.

(2) Oxygen or air is pressurized and mixed with the feed. In the adiabatic reactor organics are oxidized at residence times of less than one minute. The heat released by combustion is sufficient to raise the fluid phase to temperatures where all organics are rapidly oxidized, temperatures are about 550 degrees C.

(3) Effluent from the reactor/oxidizer is fed to a salt separator where inorganics originally present in the feed precipitate.
(4) A portion of the superheated SCW is recycled to an eductor upstream of the oxidizer. This recycle stream provides sufficient thermal energy to heat the feed to the oxidizer to supercritical conditions.

(5) The remainder of the superheated SCW (with some  $CO_2$  and  $N_2$ ) is available for power generation or use as high-pressure steam.

The available data from SCWO is presented in Tables 9 to 15.

For urea destruction, Figure 4 shows the pertinent reactions, including heats of reaction and temperatures. At certain temperatures all the nitrogen in urea  $((NH_2)_2CO)$  is not produced as N<sub>2</sub>, but can be in the form of ammonia NH<sub>3</sub> or nitrous oxide N<sub>2</sub>O. Figure 5 depicts the ammonia, nitrogen, and nitrous oxide weight ratios as a function of SCWO reactor temperature.

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The advantages and disadvantages of the SCWO process are as follows. The process can treat acqueous waste streams with a wide range of flows and composition. The streams may contain both inorganic and organic compounds. The reaction exotherm is sufficient to make the oxidation self-sustaining and to provide preheat for feed to about critical temperature. No catalysts are required for this oxidation process. Inorganic salts can be readily removed by precipitation. Finally, the necessary residence time in the reactor/oxidizer is from one-half to one minute, only. The disadvantages of the SCWO process are that it requires extremely high temperature and pressures. Large

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-	SCW	- U.S	S. PATER	VT 4,	113,446	(1978	)
	<u> </u>	SULTS NDITIO	NIS / 1.4	17K AND	218 AT/	m)	
Run	CARBON CLOSURE	% C AS VOLATILE GAS	<u>LUCOSE</u> H <sub>Z</sub>	CH4	CO2	СО	C2+
9(c)	97	10	30	1,5	42	27	-
10	86	8	25.8	1,3	34.4	38.5	-
11(c)	86	20	45.1	3,2	38.5	12.5	0.7
12 (c)	94.1	23.3	45,13	2.9	40,6	12.6	0.75

C DENOTES CATALYST RUN.

RUNS 9 AND 10 FOR ONE HOUR.

RUNS 11 AND 12 FOR 30 MINUTES.

\* TABLE III

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# TABLE 10

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# SCW - GLUCOSE AND WOOD

Reproducibility of Results; Glucope Peed and 60-Min. Residence Time

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GAS CAPOSICION									
Carbon in Feed (q)	Ave. Temp. (C)	H2	3	CH4	∞ <sub>2</sub>	c2+	Carbon Gasified (1)		
2	177	16.3	67.1	1.2	14.8	0.6	16.9		
				-		-	16.6		
	377		63.0	1.9	17.8	0.9	21.3		
2	377	11.5	70.8	1.5	15.5	0.7	22.9		
2	377	13.1	68.4	1.5	16.4	0.3	19.5		
2	371	19.9	53.5	2.3	23.3	1.1	25.9		
2	371	8.5	54.6	0.8	34.8	1.2	25.0		
2	371	14.2	54.1	1.6	29.1	1.1	25.5		
4	371	12.3	43.9	0.9	41.7	1.7	10.7		
4	371	25.8	38.5	1.3	34.4	-	8.2		
4	371	19.1	41.2	1.1	33.1	0.9	9.5		
	in Feed (c;) 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 4 4 4	in Feed Temp. (c) (C) 2 377 2 377 2 377 2 377 2 377 2 377 2 377 2 377 2 371 2 371 2 371 2 371 4 371 4 371	in Feed Temp. H <sub>2</sub> (c) (C) H <sub>2</sub> 2 377 16.3 2 377 10.2 2 377 14.4 2 377 11.5 2 377 13.1 2 371 19.9 2 371 8.5 2 371 14.2 4 371 12.3 4 371 25.8	Carbon in Feed (c)Ave. Temp. (C) $H_2$ CO237716.367.1237710.270.6237714.463.0237711.570.8237713.168.4237119.953.5237114.254.1437112.343.9437125.838.5	Carbon in Feed (c)Ave. Temp. (C) $H_2$ CO $CH_4$ 237716.367.11.2237710.270.61.5237714.463.01.9237711.570.81.5237713.168.41.5237119.953.52.3237114.254.60.8237114.254.11.6437112.343.90.9437125.838.51.3	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Carbon in Freed (c)Ave. Temp. (C)H2COCH4CO2 $C_2^+$ 237716.367.11.214.80.6237710.270.61.517.5-237714.463.01.917.80.9237711.570.81.515.50.7237713.168.41.516.40.3237119.953.52.323.31.1237114.254.11.629.11.1437112.343.90.941.71.7437125.838.51.334.4-		

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TABLE 11	
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## SCW - GLUCOSE AND WOOD

### Product Distribution in the Critical Region

	Reaction Time (min)	Time in Feed		Gas Corposition					$\leftarrow$ & Carbon as: $\rightarrow$		
Feed (Run No.)			Ave. Terp. (C)	H <sub>2</sub>	8	CH4	œ <sub>2</sub>	c2*	Gas	Extracted Liquid	Recovered Water (cm <sup>3</sup> )
Gluccase											
G-1-4	60	2	377	13.1	68.4	1.5	16.4	0.3	19.5	1.2	266
G-5,6	60	2	371	14.2	; 54.1	1.6	29.1	1.1	25.5	2.5	280
G-?	150	2	371	12.3	42.8	1.2	41.4	2.3	28.9	-	271
5-8,9	60	4	371	19.1	41.2	1.1	38.1	0.9	9.5	-	265
levulinic	7 .1 <b>d</b>										
<b>L-1</b>	60	4.82	377	4.8	87.9	0.6	6.6	-	10.8	-	251
Naple saud	list										
<del>%-</del> 1	5	.59	377	8.3	79.2	2.3	10.1	**	16.8	1.1	241
<del>2</del>	15	.96	377	17.9	69.4	3.7	9.1	-	18.1	1.5	258
<del>9-</del> 3	30	.10	377	15.9	65.5	5.1	13.0	0.5	88.3	-	249
<del>N-4</del>	30	.67	377	5.0	82.4	2.9	9.6	0.1	33.1	C.6	252
<del>. 1</del> -5	60	. 29	377	16.8	\$7.2	5.6	19.5	0.9	39.5	6.2	258

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50	W - U	U.S. PA	TENT	4.//3	3 446	[1978	)
Å		rs A <del>t</del> 647 K					
Run #		% CARBAN VOLATILE		GAS	COM PO	SITION	C2+
CELLULOSE 13	95,7	18.31	14.5	1.5	19.7	64.2	0,13
PULYETHYLEN 15	E 90,3	2.30	26.6	2.54	65. <b>8</b> 7	≤5,0	_
POLYF THYLENG 16	= 93,7	4.01	35.75	3.73	54.43	5.0	1.2
UREA 17	88.8	//.//	-	-	100		-
F	LE RU	INS WIT	H CA	TALXST			

TABLE 12

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SUPERCRITICAL WATER OXIDATION - UREA DESTRUCTION: (O2, 23 SEC, 25 cc/min)

(NH2)2CO + 3/2 O2 - N2 + CO2 + 2H2 REACTOR INFLUENTIO % UREA IN WATER

			RUN No.		
	<u>81A</u>	<u>815</u>	<u>81C</u>	<u>82</u>	225
Temperature (°C)	561	596	621	632	670
Hitrogen In (mg/l) (water + feed)	51422	50501	50825	50935	56679
Effluent Liquid Composit	tion (mg/l)				
Witrate W	100	85	29	57	- 04
Nitrite W	-	-	-	-	0.005
Ammonia N	17537	15798	3447	2719	1.65
Effluent pä	9.21	8.95	7.92	7.52	7.24
Effluent Ges Composition	(mole I)				
0 <sub>2</sub>	95.59	90.18	37.61	20.46	8.09 (3.63)*
αı	0.45	1.05	18.38	38.48	41.28 (18.53)
¥2	3.60	7.92	41.72	39.04	50.63 (22.73)
¥20	0.37	.85	2.33	0.51	0.00 ( 0.00)
Nitrogen in Liquid Efflu	ent (I)				
	26.7	26.7	5.5	4.5	0.0
Mitrogen in Ges Effluent	(1)				
	5.2	11.6	59.4	69.2	97.7
Mitrogen Recovery (I)	31.8	38.3	65.3	73.7	97.7

• Values in parentheses represent the actual measured quantities. For Run 225, sith 40% helium present, values are normalized for comparison purposes.

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# TABLE 14

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SCW - ORGANIC HALIDE DESTRUCTION

### SUMMARY OF RESULTS: OXIDATION OF ORGANIC CHLORIDES

			*********		
Eve No.	11	12	13	14	15
Besidence Time (min)	1.1	1.1	1.1	1.1	1.3
<u>Carbon Analysis</u>					
Organic Carbon In (ppm) Organic Carbon Out (ppm) Destruction Efficiency (I) Combustion Efficiency (I)	26,700. 2.0 99.993 100.	25,700. 1.0 99.996 100.	24,500. 6.4 99.975 100.	38,500. 3.5 99.991 100.	33,400. 9.4 99.97 100.
Ges Composition					
0, 00, CH4	25.58 59.02 -	32.84 51.03 -	37.10 46.86 -	10.55 70.89 -	19.00 70.20 -
<b>5</b>	-	-	-	-	-
Chloride Analysis					
Organic Chloride In (ppm) Organic Chloride Out (ppm) Organic Chloride Conversion (%)	876. .023 99.997	1266. .037 99.997	748. <.028 99.996	775. .032 99.996	481. .036 99.993
GC/HS Effluent Analysis					
Compound B (ppb Cl) C E F E K N N O	- - 18 - - - -	- 9 12 - 16 -	- - 18. <4. <5. 0.2 0.3	- 14 - 6 - 12	

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# TABLE 15 SCW - ORGANIC HALIDE DESTRUCTION

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#### TABLE 2. COMPOSITION OF FEED MIXTURES FOR SUNS 11 - 15

Rus 11		<u>vt 1</u>	vt I Cl
DOT	C14H9C15	4.32	2.133
NEK	C4H80	95.68	2.133
<u>Rum 12</u> 1,1,1-trichloroethese	C2H3C13	1.01	0,806
1,2-ethylene dichloride	C2H2C12	1.01	0.725
1,1,2,2-tetrachlorethylene	c <sub>2</sub> c1 <sub>4</sub>	1.01	0.866
e-chlorotoluene	C, H, C1	1.01	0.282
1,2,4-trichlorobenzene	C6H3C13	1.01	0.591
bipheayl	с <sub>12</sub> π <sub>10</sub>	1.01	-
o-xylene	C8810	5.44	-
MEK	C <sup>4</sup> B <sup>8</sup> O	<u>88.45</u> 100.0	3.284
<u>Run 13</u> bexachlorocyclohexane	C6H6C16	0.69	0.497
DDT	C14H9C15	1.00	0.493
4,4'-dichlorobiphenyl	C1288C1	1.57	0.495
hexachlorocyclopentadiene	с <sub>5</sub> с1 <sub>6</sub>	0.65	0.505
NEK	с <sub>цив</sub> о	<u>96.09</u> 100.0	1.99
Run 14 PCB 1242	C12 <sup>H</sup> x <sup>C1</sup> 4-6	0.34	0.14
PCB 1254	C12 <sup>R</sup> x <sup>C1</sup> 5-8	2.41	1.30
transformer oil	c10 <sup>-c</sup> 14	29.26	-
	C <sub>4</sub> H <sub>8</sub> G	<u>67.99</u> 100.0	1.44
<u>Run 15</u> 4,4'-dichlorobiphenyl	C <sub>12</sub> R <sub>B</sub> Cl	3.02	.96
HE K	C480	<u>96.98</u> 100.0	0.96

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temperature gradients apparently exist in the system which may make it difficult to duplicate measurements and to scale available data. The oxygen or air flow requires precise control along with the waste feed stream. The particle size in the waste stream, solids in water, must be reduced to less than a millimeter in diameter. The data available for the SCWO indicates that the gaseous effluents are not simply carbon dioxide and nitrogen but may contain significant amounts of carbon monoxide, methane, and hydrogen. The presence of these gases would require extensive post-treatment for Space Station application.

### ZIMPRO PROCESS

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The Zimpro process was invented about 50 years ago, in the mid-1930's. This process is also a wet air (oxygen) oxidation process but at temperatures and pressures well below water's critical conditions. Figure 6 is a schematic of a wet air oxidation (WAO) system. In WAO molecular oxygen reacts with suspended solid matter or dissolved organics almost complete (90 to 95%) reduction of the wastes to carbon dioxide and water. With a residence time of 90 minutes and temperatures/pressures below critical conditions the maximum amount of carbon monoxide is estimated to be 1000 ppm, average carbon monoxide concentrations are 200 ppm. The organic materials, if not converted to carbon dioxide, go to the water phase as alcohols, aldehydes, and ketones. The degree of oxidation can be controlled by WAO temperature, pressure, and residence time. For some applications catalysts have been used. The major advantage of WAO is that

contaminants tend to stay in the aqueous phase; effluent gas consists mainly of carbon dioxide and spent air (or excess oxygen).

As evident from Figure 6 the WAO system is similar to SCWO, Figure 2, except for the salt precipitation. In this report details of the WAO operating procedure are not described. However, the WAO operates at temperatures of 200 to 300 degrees C which are much less than SCWO. It is important to note that the WAO is commercially available with decades of operations in more than 200 units, world-wide. These units have been applied mostly to sewerage sludge treatment and manufacturing process wastewaters.

The wet air oxidation system can be used either on a batch or continuous basis.

### CHEMICAL OXIDATION

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Chemical agents such as potassium permangate, potassium dichromate, and many other alkali metal salts can oxidize solid wastes. The advantage of a chemical oxidation process is that it would operate at about 1 atm pressure, or be a low pressure system in contrast to the SCWO and WAO processes. A chemical oxidation system would need heating to temperatures from a minimum of 100 degrees C to several hundred degrees Celsius. About one kilogram (kg) of oxygen is needed to oxidize an equal mass of solid wastes. If a chemical oxidizer was provided than the mass of oxidizer

would be no more than twice that of pure (liquid) oxygen; however, the volume of chemical oxidizer would be about the same since the salts have a density more than twice that of liquid oxygen. The salts might also provide nitrogen needed for atmospheric control. With chemical oxidation system reactor materials of construction would have to be carefully considered. More expensive materials such as titanium and Hastelloy-C would probably be required. But, Hastelloy-C is often used in the WAO process.

Chemical oxidations could operate in a batch or continuous mode. Due to the nature of the solid wastes on a Space Station a batch type treatment process is suggested. This would allow the collection of materials for say about one week, then their oxidation.

#### OTHER PROCESSES

A few other processes were considered, namely the IT Enviroscience homogenous catalyst system, Figure 7 and the plasma torch under development at the Royal Military College, Ontario, see Figure 8. With regard to dry combustion, a study of General American Transportation (1968) of fecal matter incineration indicates that it is feasible but that such a system requires precise control of oxygen flow rate, and continuous monitoring of

temperature and oxygen concentration. This experimental study determined that dry feces will ignite at about 200 degrees C, and will burn very rapidly - under certain conditions will detonate. Thus, the collection and storage of dried fecal matter on a Space Station should be considered a potential fire hazard.

The burning of polyethylene (wastes) is difficult and would probably not be practical on a Space Station.

Biodegradation processes would probably require too much astronaut attention and too much space in the Station.

### SPACE STATION TREATMENT PROCESS

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A proposed solid waste management treatment system for a Space Station is presented in Figure 9. This schematic diagram indicates that the transfer of solid wastes from the collection point to the treatment system should be minimized. Furthermore, the maceration shredding, grinding of plastic and cellulosic wastes, if attempted, will pose severe maintenance problems. The specific type of treatment process needs to be developed. Prior to any chemical reactions the system would dry and compact the solid wastes. The treatment reactor is envisioned to be about 5 gallon capacity, with no penetrations between the reactor cavity and the outside. This could be done by means of a magnetically controlled stirrer, with a chain drive. Equipment other than the

tre int reactor has been used in each ting spacecraft or developed. The proposed system would exhibit zero-leakage except for spont filters and in-exchange adding. The amount of solid ash from the treatment process should be a few percent of the total feed mass.

Figures 10 to 13 summarize the principle reactions for oxidation of wastes, considered as a mixture of carbon, hydrogen and oxygen, and subsequent reduction of carbon dioxide to oxygen and either carbon or methane.

### CONCLUSIONS

C

As a result of this study the following general conclusion is made; a solid waste(s) treatment process for a Space Station needs further study to achieve a reliable operational system. The solid waste rates determined in this study, especially those due to food wastes, paper and plastics, need to be better acertained from at least Shuttle operational data. Processes for the conversion of solid wastes to carbon dioxide and other inert substances need additional experimental measurements.

There are a few feasible treatment processes, chemical or wet air oxidation; supercritical water oxidation does not appear suitable for a Space Station. In addition to the treatment system design needs, the problem of mechanical transfer of wastes from collection point(s) to the treatment vessel is critical. This

aspect along with total systems integration needs to be considered.

Finally, when design criteria for a Space Station are better "stablished and better data are available on treatment performance, a trade study should be conducted to evaluate alternatives.



### NAME LIST OF PERSONS CONTACTED REGAPDING

### TREATMENT OF SOLID WASTES OF A SPACE STATION

NASA Employees

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rion-NASA Personnel

Fullerton, G. Allen, L. (Excell) Grimaud, J. Johnson, C. (Ames) Barton, T. (Royal Military College) Kimzey, H. Bourland, C. (Technology) Langdoc, W. Engmann, R. (Zimpro) Martin, R. Howell, G. (U.S. EPA) Rapp, R. Lamparter, R. (Michigan Tech) Samonski, F. Modell, M. (Modar) Sauer, R. Murray, R. (GE) Winkler, E. Obert, J. (McDonnell-Douglas) O'Connor, T. (Univ. Missouri-Columbia) Schaefer, P. (Zimpro) Weitzmann, A. (Lockheed)

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#### -Treatment Processes-

Hurwitz, E., and Dundas, W.A., "Wet Oxidation of Sewage Sludge, "Water Pollution Control Federation Journal, Vol. 32 (9), pp. 918-929, 1960.

Pipes, W.O., "Waste-Recovery Processes for a Closed Ecological System," National Academy of Sciences-National Research Council, Publication 898, Washington, D.C., 1961.

Zämmerman, F.J., "Wet Air Combustion," Industrial Water and Wastes, Vol.6, pp. 102-106, 1961.

Quon, J.E. and Pipes, W.O., "Thermal Decomposition of Human Wastes," Proceedings American Society of Civil Engineers, Vol. 89, No. SA1, January, 1963.

Dodson, J., and Wallman, H., "Research on a Waste System for Aerospaces Stations," Technical Documentary Report No. AMRL-TDR-64-33, 1964.

Harding, J.C., and Griffin, G.E., "Sludge Disposal by Wet Air Oxidation at a Five MGD Plant," Water Pollution Control Federation Journal, Vol. 37, pp. 1134-1141, 1965.

Rich, L.G., Ingram, W.M., and Berger, B.B., "Waste Disposal on Space Craft and Its Bearing on Terrestrial Problems," U.S. Department Health, Education, and Welfare, Public Health Service, PB 168787, 1965.

Wheaton,R.B., Brown, J.R.C., Ramirez, R.V., and Roth, N.G., "Investigation of the Feasibility of Wet Oxidation for Spacecraft Waste Treatment," NASA Contractor Report No. 66450, 1966.

Elms, R.V., "Design Study of Integrated Life Support System for Aerospace Application Utilizing Radioisotopes for Thermal Energy," Final Report, LMSC 680 679, 1968.

Hurley, T.L., Rollo, E.J., and Remus, G.A., "Study for Evaluation of Incineration and Microwave Treatment of Human Fecal Matter for Spacecraft Operation," NASA CR-73247, 1968.

Hamilton Standard, "Alternate Mission Studies (AILSS)," NASA Contractor Report-66876, 1969.

"Huggett, C., "Combustion Processes in the Aerospace Environment," Aerospace Medicine Vol. 40, pp. 1176-1180, 1969.

Jagow, R.B., Jaffe, R.J., and Saunders, C.G., "The Processing of Human Wastes by Wet Oxidation for Manned Spacecraft," American Society of Mechanical Engineers, Paper 70-Av/SpT-1, Presented at Space Technology and Heat Transfer Conference, Los Angeles, CA, June 21-24, 1970; in Space Systems and Thermal Technology for the 70's, Part 1, Proceedings pp. 1-8, 1970.

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#### Zimpro Process

Hurwitz, E., Teletzke, G.H., Gitchel, W.B., "Wet Air Oxidation of Sewage Sludge," Water and Sewage Works, Vol. 112, pp. 298-305, 1965.

Pradt, L.A., "Developments in Wet Air Oxidation," Chemical Engineering Progress, Vol. 68 (12), pp. 72-77, 1972.

Randall, T.L., and Knopp, P.V., "Detoxification of Specific Organic Substances by Wet Oxidation," Reprint, Presented at 51st Annual Conference, Water Pollution Control Federation, Anaheim, CA, October, 1978.

Flynn, b.L., "Wet Air Oxidation of Waste Streams," Chemical Engineering Progress, • Vol. 75 (4), pp. 66-69, 1979.

Wilhelmi, A.R., and Knopp, P.V., "Wet Air Oxidation-An Alternative to Incineration," Chemical Engineering Progress Vol. 75 (8), pp. 46-52, 1979.

Randall, T.R., "Wet Oxidation of Toxic and Hazardous Compounds," Reprint, Presented at 13th Mid-Atlantic Industrial Waste Conference, June, 1981.

Schaefer, P., "Consider Wet Oxidation," Hydrocarbon Processing, Reprint, October, 1981.

Canney, P.J., and Schaefer, P.T., "Detoxification of Hazardous Industrial Wastewaters by Wet Air Oxidation," Presented at Spring National AIChE Meeting, Houston, Texas, March, 1983.

14-38

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#### -Process Reviews-

Presti, J.B., "Life Support System for Space Flights of Extended Time Periods," NASA-Contractor Report-168467, 1963.

Rich, L.G., Ingram, W.M., and Berger, B.B., "Waste Disposal on Space Craft and Its Bearing on Terrestrial Problems," U.S. Department of Health Education, and Welfare, Public Health Service, PB 168787, 1965.

Parker, J.F., and West, V.R., Eds, <u>Bioastronautics Data Book</u>, NASA SP-3006, Second Edition pp. 922-930, 1973.

Whitaker, C.F., Murray, R.W., and Scheikopf, J.D., "Shuttle Era Waste Collection," Institute Environmental Sciences, Proceedings 22nd Annual Technical Meeting, Philadelphia, PA, pp. 257-260, 1976.

Bioenvironmental Systems Study Group of the Society of Automotive Engineers, "Review of Technology Developments Relevant to Closed Ecological Systems for Manned Space Missions," NASA - Contractor Report-153000, 1977.

ţ

Modell, M., "Sustaining Life in a Space Colony," Technology Review, pp. 36-43, July/ August 1977.

13 H. Martin and the state

÷

Society of Automotive Engineers, "Space Technology Available for Transfer to Civil Environmental and Pollution Control," SAE Aerospace Information Report AIR 1246, 1977.

DeRenzo, D.J., Ed., <u>Unit Operations for Treatment of Hazardous Industrial Wastes</u>, (Noyes Data Corporation, New Jersey), 1978.

Spurlock, J.M., and Modell, M., "Technology Requirements and Planning Criteria for Closed Life Support Systems for Manned Space Missions," Final Report for NASA, Contract No. NASw-2981, 1978.

Meissner, H.P., and Modell, M., "Recycling plant, Human and Animal Wastes to Plant Nutrients in a Closed Ecological System," American Society of Mechnical Engineers, Publication 79-ENAs-29, 1979.

Spurlock, J., and Modell, M., "Systems Engineering Overview for Regenerative Life-Support Systems Applicable to Space Habitats," in Space Resources and Space Settlements, Billingham, J., Gilbreath, W., and O'Leary, B., Eds., NASA SP-428, pp. 1-11, 1979.

Basta, N., "Firms Avidly Seek New Hazardous-Waste Treatment Routes," Chemical Engineering, pp. 53-55, September 6, 1982.

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#### Supercritical Water Oxidation

Amin, S., Reid, R.C., and Modell, M., "Reforming and Decomposition of Glucose in an Aqueous Phase," American Society of Mechanical Engineers, Publication 75-ENAs-21, 1975.

Modell, M., "Reforming of Glucose and Wood at the Critical Conditions of Water," American Society of Mechanical Engineers, Publication 77-ENAs-2, 1977.

Modell, M., Gasification Process, United States Patent 4,113,446, Sept. 12, 1978.

Anon., "Using Supercritical Water to Destroy Tough Wastes," Chemical Week, Reprint, April 21, 1982.

Josephson, J., "Supercritical Fluids," Environmental Science and Technology, Vol. 16, pp. 548A-551A, 1982.

Modell, M., Processing Methods for the Oxidation of Organics in Supercritical Water, United States Patent 4,338,199, July 6, 1982.

Modell, M., Gaudet, G.G., Simson, M., Hong, G.T., and Biemann, K., "Destruction of Hazardous Waste Using Supercritical Water," United States EPA, Proceedings of **Eighth** Annual Research Symposium, pp. 202-212, March 8, 1982.

Modell, M., Gaudet, G.G. Simson, M., Hong, G.T., and Biemann, K., "Supercritical Water-Testing Reveals New Process Holds Promise," Solid Wastes Management, Reprint, August, 1982.

Timberlake, S.H., Hong G.T., Simson, M., and Modell, M., "Supercritical Water Oxidation for Wastewater Treatment: Preliminary Study of Urea Destruction," Society of Auto-Notive Engineers. Technical Paper Series 820872, Presented at Twelfth Intersociety Conference on Environmental Systems, San Diego, CA. July, 1982.

14-40

#### REFERENCES (CONT'D)

Cadotte, A.P., and Laughlin, R.G.W., "The Wetox (R) Process for Industrial Waste Treatment," in <u>Waste Treatment and Utilization</u>, Moo-Young, M. and Farquhar, G.J., Eds., Pergamon Press, Oxford, pp. 157-172, 1979.

Farquhar, G.J., "Biological Treatment of Industrial Wastes: Review of Principles, Methods and Applications," in <u>Wastewater Treatment and Utilization</u>, Pergamon Press, Oxford, pp. 373-393, 1979.

Othmer, D.F., "Earth + Water + Air = Fire," Mechanical Engineering, pp. 30-37, 1979.

Chow, C.L., and Verhoff, F.H., "Process for Power Generation from Wet Air Oxidation with Application to Coal Gasification Waste Waters," Industrial Engineering and Chemistry Process Design and Development, Vol. 20, pp. 12-19, 1981.

Onisko, B.L., and Wydeven, T., "Wet Oxidation as a Waste Treatment in Closed Systems," American Society of Mechanical Engineers, ASME-81-ENAs-22, 1981.

Shuler, M.L., Nafis, D., and Sze, E., "The Potential Role of Aerobic Biological Waste Treatment in Regenerative Life Support Systems," ASME, Paper 81-ENAs-20, 11th Intersociety Conference on Environmental Systems, San Francisco, CA, 1981.

Barton, T.G., "Plasma Destruction of Polychlorinated Biphenyls," Royal Military College, Kingston Ontario, Contract Report prepared for Ministry of Environment, Ontario, March 1982.

Barton, T.G., "Mobile Plasma Pyrolysis System," Presented at United States EPA, Cincinnati, 18 May 1983.

Excell, Inc., "Method for Destroying Toxic Organic Chemical Products," Personal Communication (Confidential Business Information) to T.R. Marrero, July 8, 1983.

IT Corporation, "Catalyzed Wet Oxidation Process Description and Summary of Stateof-the-Art," Personal Communication from R.W. Helsel to T.R. Marrero, June 20, 1983.

Johnson, C.C., and Wydeven,T. "Wet Oxidation for Waste Treatment in a Controlled Ecological Life Support System (CELSS)," National Aeronautics and Space Administration, Preprint, Presented at Thirteenth Intersociety Conference on Environmental Systems, San Francisco, July 11-13, 1983.

Miller, R.A., and Swientoniewski, M.D., "The Destruction of Various Organic Substances by a Catalyzed Wet Oxidation Process," Proceedings of Eighth Annual Research Symposium, March 8-10, 1982, Incineration and Treatment of Hazardous Waste, United States EPA, pp. 213-221, 1983.

#### REFERENCES (CONT'D)

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Schelkopf, J.D., Witt, F.J., and Murray, R.W., "Integrat 1 Waste Management - Water System Using Radioisotopes for Themal Energy," Summary Report. United States Atomic Energy Commission, Contract Number AT (30-1)-4104, 1970.

Jagow, R.B., "Design and Development of a Prototype Wet Oxidation System for the Reclamation of Water and the Dispositon of Waste Residues Onboard Space Vehicles," NASA Contractor Report 112151, 1972.

Labak, L.J., and Remus, G.A., "Integrated, Zero "G" Waste Incineration System," General American Research Division, General American Transportation, Research Report (Phase B), NASA Contract No. NAS 2-6386, 1972.

Schelkopf, J.D., Witt, F.J., and Murray, R.W., "Integrated Waste Management-Water System Using Radioisotopes for Thermal Energy," United States Atomic Energy Commision, Document No. 74SD4201, 1974.

Bioenvironmental Systems Study Group of the Society of Automotive Engineers, "Evaluation and Comparison of Alternative Designs for Water/Solid-Waste Processing Systems for Spacecraft," NASA-Contractor Report-162492, 1975.

Chowdhury, A.K., and Ross, L.W., "Catalytic Wet Oxidation of Strong Waste Waters," American Institute of Chemical Engineers, Symposium Series, No. 151, Vol. 71, pp. 46-58, 1975.

Budininkas, P., "Study of Removal of Ammonia from Urine Vapor by Dual Catalyst," NASA Contractor Report 151930, 1976.

Jagow, R.B., "Design, Fabrication and Testing of a Spacecraft Wet Oxidation System Including Trash Pulverization Studies," American Society of Mechanical Engineers, ASME-76-ENAs-15, 1976.

Katzer. J.R., Ficke, H.H., and Sadana, A., "An Evaluation of Aqueous Phase Catalytic Oxidation," Water Pollution Control Federation Journal, Vol. 48 (5), pp. 920-933, 1976.

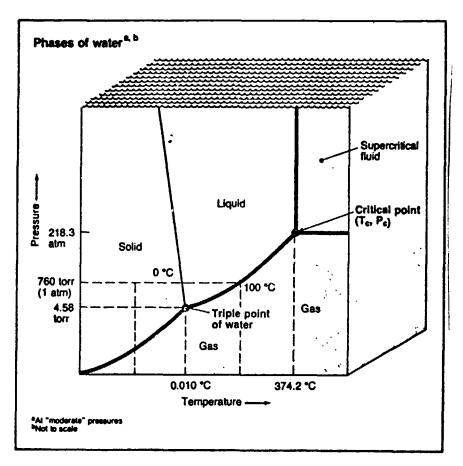
Budininkas, P., "Removal of Ammonia from Vapor by a Dual-Catalyst System," American Society of Mechanical Engineers, Publication 77-ENAs-48, 1977.

Heidemann, R.A., and Prausnitz, J.M., "Equilibrium Data for Wet-Air Oxidation. Water Content and Thermodynamic Properties of Saturated Combustion Gases," Industrial Engineering Chemistry, Process Design and Development, Vol. 16 (3), pp. -5-381, 1977.

Modell, M., "Sustaining Life in a Space Colony," Technology Review, July/August, pp. 36-43, 1977.

Weitzmann, A.L., "Development and Testing of a Wet Oxidation Waste Processing System," National Aeronautics and Space Administration, NASA-CR-151324, 1977.

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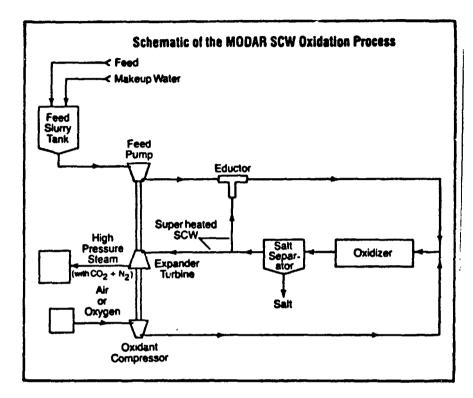
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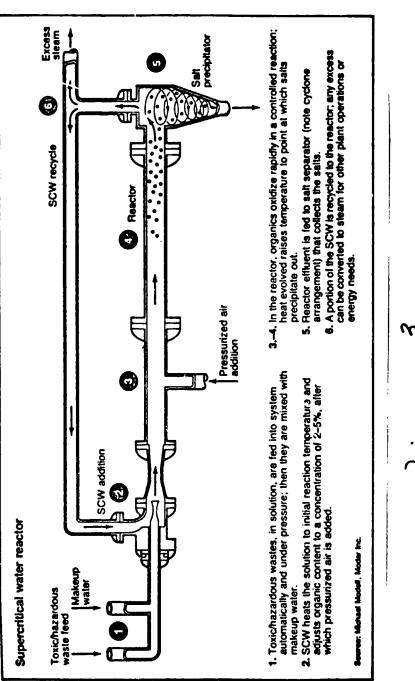
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# UREA REACTIONS IN SUPERCRITICAL REGION

<u>Overall Reaction at Nigh Temperature</u> (NN<sub>2</sub>)<sub>2</sub>CO + 3/<sub>2</sub>O<sub>2</sub> ------ N<sub>2</sub> + CO<sub>2</sub> + 2H<sub>2</sub>O  $\Delta$ H<sub>R,298</sub> = -632.2 kJ Urea

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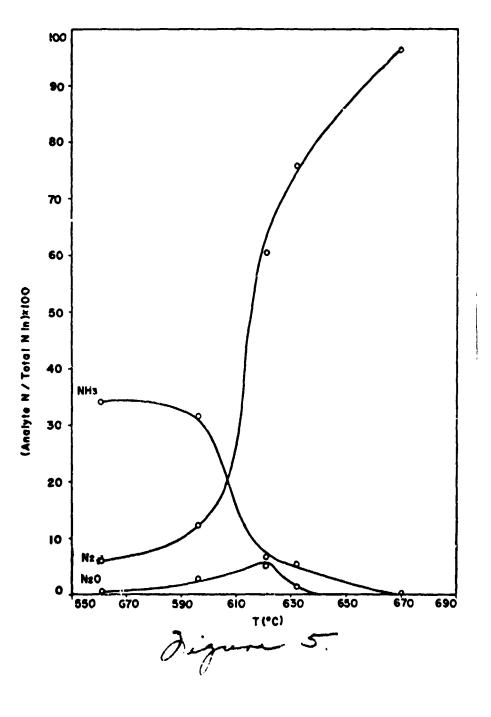
Step 1: Hydrolysis of ures to sumonis and carbon dioxide		WITIAL
(M12)2CO + H2O 2NH3 + CO2 Urea	ΔN <sub>R,298</sub> = 45.6 kJ	
Step 2A: Oxidation of amonia to nitrogen		
21013 + 3/202 N2 + 3H20	$\Delta H_{R,298} = -765.8 \text{ kJ}$ $\Delta 3_{R,873} = -631.5 \text{ kJ}$	z 561 °C
Step 28: Oxidation of amonia to nitrous oxide		
2NE3 + 202 N20 + 3H20	Δ8 <sub>R,298</sub> = -684.2 kJ Δ8 <sub>R,873</sub> = -549.7 kJ	2 621°C
Step 3: Decomposition of nitrous oxide to nitrogen and oxygen		7. 170%
2H20 2H2 + 02	$\Delta H_{R,298} = -163.3 \text{ kJ}$ $\Delta H_{R,873} = -162.5 \text{ kJ}$	≈ 670°C

Digure 4.

## SCW - UREA DESTRUCTION

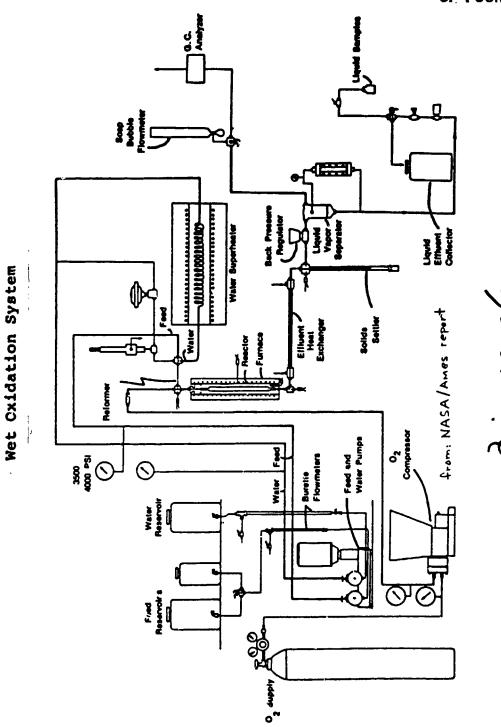
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DISTRIBUTION OF NITROGEN SPECIES IN THE LIQUID AND GASEOUS EFFLUENTS (TIMBERLAKE, S.H., ET AL., 1982)



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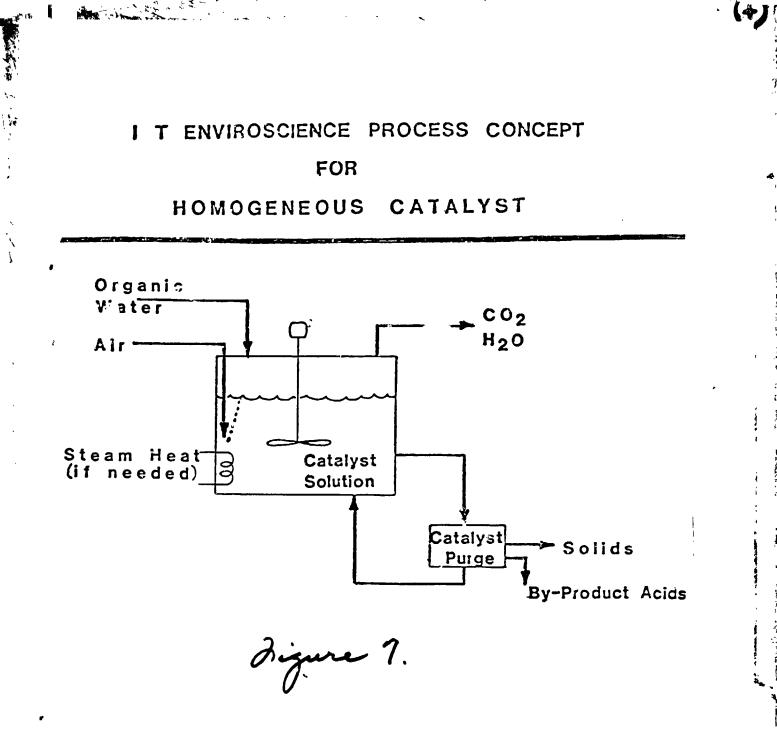
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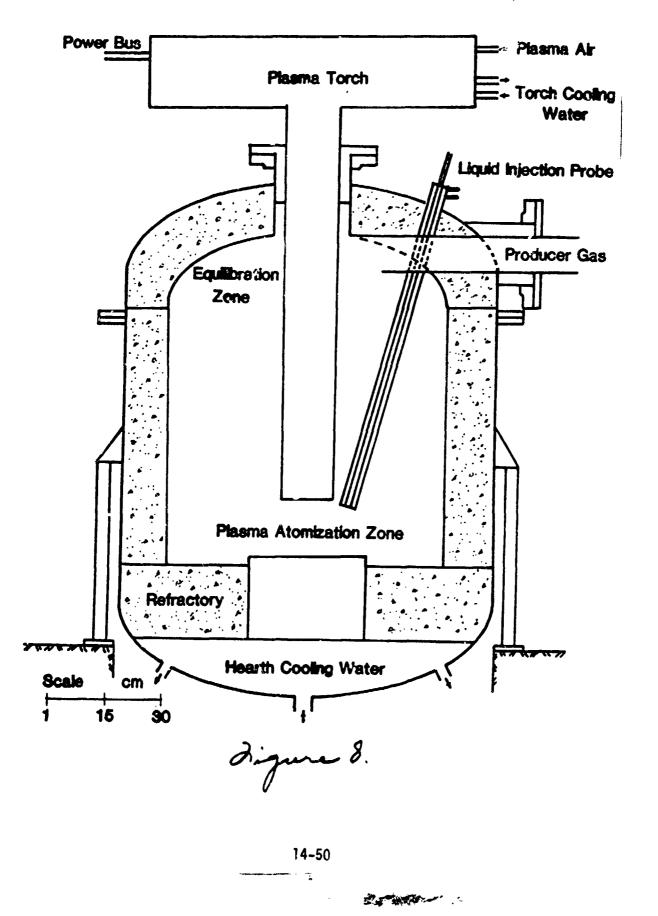
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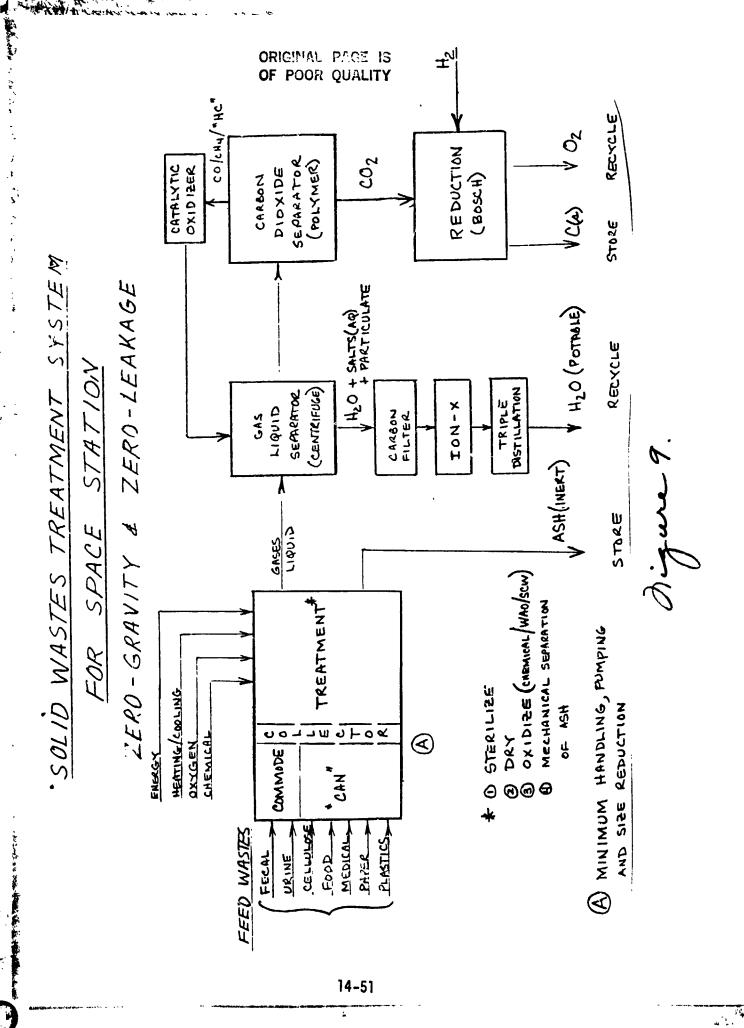


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### **REACTION VESSEL SCHEMATIC**

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OXIDATION REACTION(S)

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 $4c_{a}H_{b}O_{c} + (4a+b-2c)O_{z} \rightarrow 4aCO_{2} + 2bH_{2}O$ 

+ <u>b+84</u> H<sub>2</sub>0 + 2d Cu 34  $C_{a}H_{b}O_{c} + dC_{2}O_{7}^{-} + BdH^{+} \xrightarrow{\Delta} aCO_{2}$  $d = \frac{3}{3}\omega + \frac{b}{6} - \frac{2}{3}$ 

where

conversion efficiency = 5 h 8 % (carbon & water)  $\star CO_2 + \varkappa H_2 = a H_2O + bC + \kappa CO + dCH_4$ a,b,c,d = f (time, temperature, pressure, iniatial concentration of reactants) Carbon dioxide reduction by hydrogen = 20 to 12 Dique general 4 = 2X recycle ratio

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CARBON DIOXIDE REDUCTION

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 $2H_2 \longrightarrow C + 2H_2O$ ıron catalyst 530°C to 730°C

 $CO_2$ 

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CARBON DIOXIDE REDUCTION

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+ 20 Kutoules/gram Hz  $CH_4 + 2H_2O$ 

 $CO_2 + 4H_2$ 

Sabatier reactor 3 Diquere 1

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