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STORABILITY INVESTIGATIONS OF WATER

SECOND ANNUAL REPORT

AEROJET LIQUID ROCKET COMPANY

SACRAMENTO, CALIFORNIA

E. M. VANDER WALL

G. R. JANSER

DECEMBER 1975

APPROVED FOR PUBLIC RELEASE:

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FOREWORD

This report covers the work performed under Contract F04611-72-C-0062, "Storability Investigations of Water," performed by the Aerojet Liquid Rocket Company at Sacramento, California, and conducted under Air Force Project Task 305911 VD. The performance period covered from 15 August 1974 to 30 September 1975.

The program manager is Dr. S. D. Rosenberg; the project manager is Dr. E. M. Vander Wall. The experimental work was conducted by Dr. Vander Wall; R. L. Beegle, Jr., senior chemist; J. A. Cabeal, senior laboratory technician; and G. R. Janser, metallurgy specialist.

The program was administered under the direction of the Air Force Rocket Propulsion Laboratory, Mr. Oree Dyes, Project Engineer.

This report has been reviewed by the Information Office/DOZ and is releasable to the National Technical Information Service (NTSI). At NTIS it will be available to the general public including foreign nations.

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Chief, Liquid Rocke	Division					

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Abstract (cont.)

Inconel 718 (aged), 6Al-4V titanium (STA). Two types of water are used in the program; oxygen-saturated, deionized, filtered, and oxygen-free, deionized, filtered. Five-year storage tests have been initiated in 304 and 17-4 PH stainless steels, A-286 steel, Inconel 718, and 6Al-4V titanium (STA) containers using the filtered, deionized waters.

Evaluation of water and containers stored for eighteen months and twentyfour months has been completed. The data show that both oxygen-saturated and oxygen-free water can be stored in appropriate metal containers for the selected time periods without detrimental particulate matter formation or significant changes in the quality of the water. It is in excellent condition for transpiration coolant purposes.

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SECTION I

INTRODUCTION

Inherent in the concept of transpiration-cooling is the requirement that the coolant remain free of particulate matter which may clog the passages of the cooling surface. The object of the "Storability Investigations of Water" program is to gather data that will permit the Air Force to assess the long term storage characteristics of water particularly with regard to formation of particulate matter. This report is the second annual progress report to document the experimental results from the eighteen month and twenty four month storage tests of water in selected metal containers for the five year program being conducted under Contract Fi)4611-72-C-0062.

The five metallic materials used for container material in the program are:

304 stainless steel A-286 (aged) steel 17-4 PH (aged) stainless steel Inconel 718 (aged) 6A1-4V Titanium (STA)

Two types of water are considered in the program:

oxygen-saturated, deionized, filtered water and oxygen-free, deionized, filtered water

Three categories of tests are used for obtaining the data necessary for assessment of the storability of water. They are:

Water Characterization Biological Characterization Container Examination

The investigations which led to the selection of the candidate metallic materials for tankage for use in the five-year storage of waters are reported in AFRPL-TR-73-94 "STORABILITY INVESTIGATIONS OF WATER," VOLUME I: EXPERI-MENTAL STUDIES, FINAL REPORT, Aerojet Liquid Rocket Company, Sacramento, California, December 1973. The results from the first year of storage are reported in AFRPL-TR-74-76 "STORABILITY INVESTIGATIONS OF WATER, LONG TERM STORAGE EVALUATION," FIRST ANNUAL REPORT, Aerojet Liquid Rocket Company, Sacramento, California, December 1974.

This annual report is presented in three sections: (1) Introduction, (2) Experimental Results and Discussions, and (3) Conclusions. In addition, thre are two appendices provided for the convenience of the reader: Appendix A, Fabrication and Treatment Procedures for Water Containers which documents the history of the tanks; and Appendix B, Silting Index Measurement which describes a clogging tendency test.

SECTION II

EXPERIMENTAL RESULTS AND DISCUSSIONS

BACKGROUND INFORMATION

The purpose of the long-term storage tests is to demonstrate that water can be stored without formation of significant quantities of particulate matter and with insignificant corrosion of appropriate metal containers for time periods of at least five years in a controlled environment. The bacground information is presented for the convenience of the reader and documents the initial conditions of the selected containers and waters prior to the storage periods. The discussion is presented under the following headings: (1) Materials Selection, (2) Container Sterilization and Filling and (3) Storage Conditions.

A. MATERIALS SELECTION

1. Waters

Based on the data derived from the preceding experimental work (Reference 1) it was apparent that both oxygen-saturated and oxygen-free water were acceptable candidates for long-term storage tests. Further, the filtration of the water through a 0.22 micron pore size absolute filter was demonstrated to remove microorganisms effectively. Thus, the water used to fill the containers was passed through an activated charcoal bed to remove organic compounds and through two mixed-bed ion exchangers to obtain water that had an electrical resistance value of 1 megohm/cm or greater. The water was transferred through 0.22 micron filters into a 5 gallon stainless steel supply tank. To ensure the saturation of the water with oxygen, filtered oxygen was purged through the water in the supply tank for a minimum of 15 minutes. To obtain oxygen-free water, the water in the supply tank was heated to the boiling point of water for one hour while being purged with filtered, nitrogen obtained from the boil-off of liquid nitrogen. The tank was then pressurized with the filtered nitrogen, allowed to cool to ambient temperatures, and then repressurized with the filtered nitrogen. The outlet of the supply tank was fitted with a Twin 90 Filter* unit to assure the sterile characteristics of the water used to fill the storage containers.

2. Metals

The selection of the materials of construction for the long term storage test containers was based on the results of the laboratory investigations (Reference 1). The aluminum alloys were eliminated from

- Ref. 1 E. M. Vander Wall, R. E. Anderson, G. R. Janser, <u>Storability</u> <u>Investigations of Water, Volume I, Experimental Studies</u>, <u>AFRPL-TR-73-94</u>, Contract F04611-72-C-0062 (December 1973).
- * A 0.22 micron pore size, absolute filter pack available from Millipore Corporation, Bedford, MA.

A. Materials Selection (cont.)

consideration due to their introduction of insoluble corrosion products which are a source of particulate matter in the water. Because test results on the remainder of the seven candidate materials were not discriminatory, choice was made on the basis of selecting not more than one alloy from each class of material. The one class of material with more than one representative was the 18% chromium - 8% nickel austenitic stainless steels, i.e., 304L, 347 and Arde-form 301. Hence two of these materials were eliminated to provide the five materials required for container fabrication. The 304L stainless steel was selected due to its attractiveness as an expulsion bladder material. Hence, the selected materials are: 304L stainless steel, A-286, 17-4PH stainless steel, Inconel 718 and 6Al-4V titanium. During fabrication, some 304 stainless steel parts were incorporated into the 304L stainless steel containers and consequently the containers are identified as 304 stainless steel containers. The fabrication procedures, the heat treatment cycles, the cleaning procedures and the passivation procedures to which the containers were subjected are presented in Appendix A of this report.

B. CONTAINER STERILIZATION AND FILLING

Following the final rinsing with filtered, deionized water, and subsequent drying of the containers in a vacuum chamber, the containers were wrapped with reusable sterilization paper. The wrapped containers were then sterilized in an autoclave at 250°F with 15 psig steam for 30 minutes followed by a 30-minute drying period. The containers were then stored in the paper to maintain their sterile condition.

All the steps required to fill the containers with water were conducted in a sterile, laminar-flow bench. The tanks were removed from the wrapping paper in the laminar flow bench. The tanks were weighed empty; then weighed when filled completely with sterile water to determine the total volume of the tank. The water was drained out and the tank was rinsed once more with the sterile water. A sample of the rinse water was checked for pH, conductivity, and Silting Index (see Appendix B). If the values indicated that particulate matter and dissolved species were not present, the tank was considered ready for filling; if the values indicated that contaminants were present, the tank was rinsed until there was no evidence of contamination. A Silting Index value of 1 or less for the rinse water when using the filter with a cross-sectional area of 1.0 mm^2 was used as the criterion that no significant quantity of loose particulate matter remained in the containers.

Before the final filling with oxygen-saturated deionized water, the tank was purged with oxygen from a filtered supply. The tank was then filled with the water and a sample was withdrawn for pH, conductivity, and Silting Index measurements. The ullage was adjusted to the ten percent value by weighing the container and its contents; the ullage space was purged with the filtered oxygen; and the container was capped with a sterile, tapered plug made from the same material as the container. The plug was seated in B. Container Sterilization and Filling (cont.)

the fill-tube by use of a hammer. The containers were filled with the oxygenfree, deionized water in analogous manner except that filtered nitrogen was used instead of oxygen for purging and blanketing the container.

The final sealing of the containers was accomplished by GTA welding the fill-tube/plug interface. The welds were inspected visually for any apparent anomalies. None were found. Then the containers were labeled for the long-term storage test and placed in plastic bags.

The sampling plan for the long-term storage tests is to remove one container of each material with the two types of water for inspection and evaluation every six months for a period of five years. The contents will be characterized with respect to pH, conductivity, particulate content, and biological activity; and the containers themselves will be subjected to metallurgical examination if the other test data indicate that this is required.

C. STORAGE CONDITIONS

The storage area for the water containers is an air-conditioned room which is monitored continuously to document that the temperature is maintained at $70 \pm 10^{\circ}$ F and that the relative humidity is maintained at 50 ± 25 percent. The containers are stored in a closed metal cabinet to protect them from an accumulation of dust and the containers themselves are covered with plastic bags to prevent direct contact with foreign metal surfaces. The containers are visually examined on a weekly basis.

WATER CHARACTERIZATION

After six month intervals of storage at the conditions defined above ten containers are removed for evaluation. They consist of two containers of each selected material, one containing oxygen-free water and the other containing oxygen-saturated water.

A. PROCEDURES

After the six month storage period, the water containers were washed with deionized water and the placed in a sterile, laminar-flow bench for further handling to remove the stored water. The outlet of the container was rinsed repeatedly with filtered, deionized water to remove any contaminants and then briefly subjected to a torch flame to sterilize the exterior of the metal. After the twelve month, eighteen month and twenty four month storage periods, the water containers were immersed in a 95% ethanol bath prior to placement in the sterile, laminar-flow bench; and then after removal from the bath and placement in the flow bench, the residual alcohol on the tank surface was removed by burning. The outlet of the container was repeatedly

A. Procedures (cont.)

exposed to a torch flame to assure a sterile condition. All the containers were opened in an identical manner. A sterile tubing cutter was used to sever the fill tube. For the six month, twelve month, and eighteen month storage periods, the water was expelled from the containers by inserting into the fill tube of the container a sterile stainless steel capillary tube through which filtered, gaseous nitrogen was passed while the container itself was inverted. The first several ml of water were used to flush the tube and were discarded. Subsequent samples of water were collected for measurement of pH, electrical conductivity, dissolved solids, particulate matter, flow behavior, and for characterization with regard to possible biological contamination. For the twenty four month storage period, special cannula were fabricated so that the water could be expelled from the container while it was maintained in its normal upright position. The cyclic insertion and withdrawal of the cannula for the various samples was avoided by controlling the nitrogen flow rate used to pressurize the container during the water expulsion.

The measurement of the pH was made using a standard pH meter with a calomel reference electrode and a glass indicator electrode. The electrical conductivity of the water was measured using a Balsbaugh Conductivity Meter, Model No. 900-.01T with a standard dip cell. The dissolved solids content of the water was determined by evaporating 200-300 mI samples of the water to dryness and weighing the residue. In addition, any particulate matter which collected on the 0.8μ filter of the flow behavior device was examined microscopically and sized. The flow behavior of the water was evaluated using a Silting Index Apparatus (see Appendix B for description) which permits filtration of the liquid through a known area (1.0 mm^2) at a constant pressure so that the flow decay due to the presence of particulate matter may be recorded as a function of time. The standard method of the test is described in ASTM F52-69. The data are reported as a silting index values; the greater the value, the greater the degree of contamination by small particulate matter.

B. DISCUSSION OF RESULTS

The data obtained from the tests are presented in Table I. The data obtained during the initial loading of the containers as well as the data for the six and twelve month storage periods are included in the tabulation to facilitate comparison and identification of trends. The baseline data are labeled as initial and the data obtained after the storage periods are labeled as final.

The significant items to be noted from the data are as follows. First, the pH values of the waters change slightly during the storage periods. If the values are averaged one finds that for storage in 304 containers the pH value increased 0.2; for storage in A-286 containers the pH value decreased less than 0.1; for storage in 6A1-4V titanium the pH value increased 0.15; for storage in Inconel 718 containers the pH value increased 0.5; and for storage

DATA INDICATIVE OF THE STORABILITY OF WATER IN SELECTED METAL CONTAINERS

I to 3µ particles. few 50µ agglomerates l to 2µ particles agglomerated as a mat Particulate Matter Characteristics 7.2 x 7.2u + 5.4 x 5.4u platelets 24x24 Agglomerated Particle 32x16. Irregular Platelets 48x24. Irregular Platelet 4x4u Irregular Particles 4x4u Irregular Particles 4x4. Irregular Particles 8x4 Irreqular Farticles 7x7u Irrequiar Platelets 2x2µ Irregular Particles Mat of small particles Agglomerated particles 3.6 x 3.6. platelets 3.2 x 1.6 platelets Mat of 3. particles 4x4u Agglomerates 1 x 4 L platelets 1 x 3u platelets 3 x 3u platelets **P x 8** u platelets 8 x 8u platelets 3.6 platelets Activity Initial Final Biciogical Water Characterization Lice Total Solids Part Silting Index Content, mg/l 4.5 s.2 6.5 3.7 9 7 Ξ 7 7 Ţ 7 T 5 Y 1.20 0.68 2.35 0.99 06.0 1.1 0.65-0.06 0.24 ю. Л 0.16 0.69 0.73 0.00 0.23 0.49 1.06 0.44 0.73 0.57 0.67 8. 1.17 0.34 0.00 06.0 0.35 0.05 0.10 0.23 0.13 0.0 0.73 0.43 0.40 0.49 0.35 0.47 0.29 0.22 0.05 0.54 0.46 0.86 0.51 0 0 0 megohr/cm nitial Finai 0.88 0.83 1.5. 1.26 0.98 0.75 0.68 **8**.0 0.97 0.99 0.93 1.18 1.17 0.84 1.05 0.82 0.75 0.55 0.97 1.08 Resistance 1.02 0.8] 1.35 1.04 1.75 1.75 1.80 1.75 1.78 1.80 1.75 1.30 2.1 22. 1.75 1.80 8 8. .90 1.90 1.65 .95 1.75 8.1 8 ଞ 1.80 8. Final 7.0 7.2 6.9 7.6 7.5 6.5 6.9 7.2 7.1 7.3 7.1 6.8 7.4 7.2 6.6 7.5 7.3 7.2 7.2 7.2 7.7 6.7 ó.4 7.1 Initial 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 6.6 7.0 6.9 7.2 7.0 6.9 7.2 7.0 7.2 6.7 7.0 7.2 0₂-free 0₂-free 0₂-free 0₂-free 0₂-free 0₂-free 02-free C₂-free 0₂-free C2-sat. 02-free 02-sat. 0₂-free 02-sat. 02-sat. 0₂-sat. 02-sat. C₂-free Type of Water 02-5at. 02-sat. 0₂-sat. 02-sat. C2-sat. 0,-5at. Exposure Period Months و و 2 12 82 18 œ 81 γ. 8 24 2 2 2 2 24 24 2 212 24 6A1-4V T1, Ko. 3 6A1-44 Ti, No. 2 5A1-4V Ti, No. 5 64!-44 Ti, No. 7 6A1-4V T1, No. 4 6A1-4V T1, No. 8 641-4V T1, No. 6 A-286, Nc. A-10F 6A1-4V Ti, No. 1 A-286, NC. A-2 A-286, NC. A-3 A-286, Nc. 19 A-256, Nc. 13 30455, %c. 11 304SS, No. 3 A-285, Nr. 7 A-286, NC. 4 304SS, Vo. 5 30455, Nr. 6 A-286, Nr. 6 3045S, NC. 1 30455, %0. 2 32455, No. 7 304SS, Nc. 4 Container Material and Number 6

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TABLE I

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TABLE I (CONTINUED)

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DATA INDICATIVE OF THE STORABILITY OF WATER IN SELECTED METAL CONTAINERS

						Wate	er Chara	cterizat	fon				
Container	Exposure			ł	Resis	tance			Total So	lids	8101091	Cal	
Material and Number	Period	Type of Water	Initial	Final	Initial	m/cm Final	Silting Initial	ndex	Content	l/bu	Activi Initial	ty Final	Particulate Matter Characteristics
Inconel 718, No. 1	9	0,-free	7.0	7.6	1.70	0.58	o	0.42	ŗ	Ļ	•	•	l to 2º particles agglomerated to 30º
Incomel 718, No. 2	9	C ₂ -sat.	7.0	7.4	1.70	0.95	C.28	0.83	ç	4	٠	ı	l to 2 ^w particles agglomerated to 30 ^w
Inconel 718, No. 6	12	د 0 ₂ -free	7.0	7.8	1.75	0.88	0	0.14	ç	Ļ	•	ı	Mat of small particles, 18. applomerates
Incomel 718, No. 4	12	C ₂ -sat.	7.0	7.3	1.80	0.98	0	0.59	Ţ	ſ	•	×	1.3. particles, 50 x 90. agglomerate
Inconel 718, No. I-4	18		7.0	1.1	1.75	1.04	0.32	0.00	Ĩ	S	ı	•	2x2. to Bx16. agglomerated Irregular Particles
Inconel 718, No. I-1	18	0 ₂ -sat.	7.0	7.5	1.75	0.88	0.56	0.35	•	Ţ	·	ı	2x2µ to 8x8µ agglomerated Irregular Particles
Inconel 718, No. I-2	24	0 ₂ -free	7.0	7.3	1.80	1,00	0.15	11.0	ŗ	9	ı	ı	24x24. agglomerated Particle
Inconel 718, No. 7	24	- C ₂ -sat.	7.0	7.3	1.80	1.03	0.40	0.0	Ţ	3.5	ı	ī	48x112, zgglomerated Particle
17-4PH SS, No. P-11	9	_ 0 ₂ -free	7.0	7.4	1.80	0.82	0	0.66	7	۲	•	·	l to 4. particles agglomerated to 200.
17-4PH SS, No. P-1	ę	02-sat.	7.0	۲.٦	1.75	0.98	Ð	0.42	5	ŀ	•	1	l to 4. particles agglomerated to 300.
17-4PH SS, No. P-12	12	0 ₂ -free	7.0	7.9	2.00	1.02	0.27	0.49	4	Ţ	•	×	Mat of indeterminate sized particles
17-4PH SS, 10, P-2	12	C ₂ -sat.	6.7	7.6	1.80	0.70	0.03	0.62	ţ	ĉ	•	×	3 to 14µ platelets
17-4PH SS, No. P-13	18	0 ₂ -free	7.0	7.4	1.95	0.86	0.40	0°.05	ŗ	2.5	•	•	Aggiomerates Range from 2x8. to 2x24.
17-4PH SS, No. P-3	18	0 ₂ -sat.	6.9	۲.٦	1.70	1.02	0.19	6 0°0	-	5.5	8	+	Agglomerates Range from 8x8: to 8x24.
17-4PH SS, No. P-14	24	0 ₂ -free	7.0	7.0	1.0	1.07	0.40	0.0	7	1.5	•	ı	4x10u Irregular Platelet
17-4PH SS, No. P-4	24	0 ₂ -sat.	6.8	7.2	1.83	1.12	0.14	0.0	Ļ	4.5	ŧ	+	3.5 x 3.5° agglomerated particles

B. Discussion of Results (cont.)

in 17-4PH containers the pH value increased 0.4. Second, the resistance values of the waters have stabilized after the initial decrease in values due to an increase in concentration of ionic species, and the final resistance values still correspond to concentration levels which contain less than one part per million of dissolved metallic ions. Third, the Silting Index values indicate the presence of a slight amount of particulate matter in the size range of less than 5 microns in the water but the concentration levels are insignificant with regard to the quantities of particulate matter that are required to cause clogging in flow passages. In addition, the concentration level of particles in the less than 5 micron size is not increasing with increased periods of water storage. Fourth, there is apparently some increase in the total solids content of the waters as the storage period increases but the values are sufficiently low so that the quality of the water is not impaired. The level of detection in the method used corresponds to a mg lt. And fifth, the pH values, the resistance values, and the Silting Index values are not significantly different between the various storage periods.

The particulate matter that was collected on the filters appeared to be that which adhered to the container walls during the cleaning, pickling, passivation, and flushing procedures prior to filling. In summation, the waters were all suitable for use in transpiration-coolant devices after periods of storage of the water for twenty-four months.

BIOLOGICAL CHARACTERIZATION

A. PROCEDURES

200 ml samples of the water taken from the storage containers were filtered through pre-sterilized filter pads which were transferred directly to sterilized Petri dishes containing suitable nutrients for direct colony counting after a suitable culturing period. The procedures are described in <u>Standard</u> <u>Methods of Analysis of Water and Waste Water</u>, American Public Health Association, 13th Edition (1971) and <u>Biological Analysis of Water and Waste Water</u>, AM 302, Millipore Corporation, Bedford, Mass. (1973). In addition, any biological organisms present were washed from the filter surfaces with a sterile buffer solution and placed directly in sterile nutrient solutions for culturing so that adequate samples are available for identifying the genus and the specific species of micro-organisms that might be present in the stored water. Based on the lag-period prior to growth of micro-organisms which have been observed earlier in the program (Reference 1), the tubes containing the nutrient solutions were incubated for periods up to one month.

Bilological Characterization (cont.)

B. DISCUSSION OF RESULTS

The results obtained by culturing samples from the water containers are presented in Table I under the heading "Biological Activity". The lack of any indications of micro-organisms being present is denoted by a minus sign; if growth was indicated in either the culture tube or on the filter pad, but not on both an "X" is used; and if growth was found in both the culture tube and on the filter pad a plus sign is used as an indication of the positive result.

After a month of incubation of the samples from the six month storage tests there was no indication of micro-organism growth on the presterilized filter pads. Slight growth was observed in the culture tubes containing washings from the 304 stainless steel containers and one of the A-286 containers. The number of micro-organisms present was extremely small as indicated by the negative results with the filter pads and the slight amount of growth in the culture tubes. There was no evidence that any biological growth occurred during the six-month storage period. The micro-organisms found were identified as an Aeromonas species.

After a month of incubation of the samples from the twelve month storage period, one of the culture tubes containing the washings from an Inconel 718 container exhibited growth, but the filter pad was negative. The micro-organisms present were identified as most likely being <u>Pseudomonas</u> <u>aeruginosa</u>. The filter pad used for the culturing the contents of both of the 17-4 pH containers, one of the 304 stainless steel containers, and one of the A-286 containers exhibited growth, but the corresponding culture tubes were all negative. The micro-organisms were identified as <u>Pseudomonas species</u>. Again there was no evidence that biological growth occurred during the twelve month storage period in any of the containers.

After a month of incubation of the samples from the eighteen month storage period, there were indication of micro-organism growth on the filter pads and in culture tubes containing samples from one of the 304 stainless steel containers, one of the A-286 containers, and one of the 17-4PH containers. The micro-organisms were identified as a <u>Pseudomonas species</u>. The number of organisms present was extremely small and there was no evidence that growth occurred during the eighteen month storage period.

After a month of incubation of the samples from the twenty-four month storage period, there were indications of micro-organism growth on the filter pads and in the culture tubes containing samples from one of the 304 stainless steel containers, both of the A-286 containers, both of the 6A1-4V titanium containers, and one of the 17-4PH containers. The micro-organisms were all of the same species and were identified as most likely being <u>Pseudomonas</u> teslosteroni based on the classifications used in the 8th edition of Bergey's B. Discussion of Results (cont.)

"Manual of Determinative Bacteriology". In this edition, much of the former genus <u>Aeromonas</u> is now included in the genus <u>Pseudomonas</u> and therefore the micro-organisms observed in the two samples of the six month storage period may be the same species as those identified in samples from the twenty-four month storage period.

The number of organisms present in the samples from the twenty-four month storage period was extremely low, less than 100 per ml of water; and there was no evidence that growth occurred during the storage period. The fact that all the organisms are of the same species indicates that they were probably introduced during the fill procedure and the improved water removal procedure reduces the possibility of contamination during the sampling procedure.

In summation, the biological testing has shown that there is no evidence for any biological growth during the storage periods even though a slight number of micro-organisms were introduced during the filling procedures.

CONTAINER EXAMINATION

A. PROCEDURES

After removal of the water from the containers by draining, they were vacuum dried for a day to insure sectioning in a dry condition. The containers were then photographed to document their general appearance. Sectioning of the containers to expose the internal surfaces was done by sawing without coolant to prevent contamination. Subsequent handling of the container halves was carefully performed to avoid touching the interior surfaces. The internal surfaces were than photographed to document their general appearance.

Further general examination at magnifications from 5 to 10X were conducted on all interior surfaces to further define conditions found in the aforementioned visual examination and to reveal additional suspect areas. Selected discrepancies were then identified for additional examination at magnifications to 40%. All welds were examined at 40% magnification. Representative discrepancies were photographed at magnifications adequate for defect definition. Those defects requiring further definition were examined metallographically to establish their cause and extent. Sections taken either through or immediately adjacent to the affected area were mounted, polished, and examined. Photomicrographs were taken to document the condition. The contaminants or corrosion products capable of being sampled were identical in appearance to those chemically analyzed in the examination of the six and twelve month exposure tanks. Hence, additional analyses were not conducted. All interior surfaces were dye-penetrant inspected to determine whether any defects were undetected during the visual examinations. No additional indications of defects were found.

Container Examination (cont.)

B. DISCUSSION OF RESULTS

The results of the container examinations are discussed under three headings: (1) General Visual Examination, (2) Metallographic and Chemical Analyses, and (3) Implications of the Results of the Examinations.

1. General Visual Examination

The external appearance of the containers is documented photographically in Figures 1 through 5. The internal appearance of the containers is documented photographically in Figures 6 through 15. Examination of these surfaces without visual aids showed full penetration for the full length of all weldments. Other conditions resulting from fabrication and cleaning procedures were: (1) etching of A-286 containers during pickling; (2) isolated dark areas, a tightly adherent smut, and isolated shiny deposits of material in one of the 17-4PH stainless steel containers; (3) a tightly adherent smut in the Inconel 718 containers; and (4) isolated areas of residual titanium oxide, localized circular areas of attack, and a general mottled appearance including fingerprint contamination in the 6A1-4V titanium containers.

No difference could be determined between the six month, twelve month, eighteen month and twenty-four month exposure containers, or between those holding the oxygen-free and oxygen-saturated water.

2. Metallographic Examination

A summary of the analyses performed on the containers is presented in Table II. The results of macro- and microexamination are shown in Figures 16 through 24. The results of the analyses are discussed below for each container material.

a. 304 Stainless Steel

No discrepancies were found in the 304 stainless steel containers. One of the containers, No. 2, had a darker appearance than the remaining containers. This condition is attributable to inadequate hydrogen gas coverage in the tank during annealing after welding. The weld contaminants reported in the examination of the six and twelve month exposure tanks were not found in this examination.

b. A-286

One weld shrinkage crack found in container No. A-10F is shown in Figure 16. The crack is 0.005 in. deep. Slight localized etching from pickling was found on all tank interiors. No evidence of corrosion resulting from water storage could be found.



NO. 1

NO. 6



Figure 1. 304 Stainless Steel Containers, Magnification 1/3X



18 MONTHS

1



Figure 2. A-286 Containers, Magnification 1/3X







NO. I-1





Figure 4. Inconel 718 Containers, Magnification 1/3X



NO. 4

NO. 7







24 MONTHS

NO. 2

18 MONTHS

NO. 6







24 MONTHS

NO. 7

18 MONTHS

NO.]

Figure 7. Interior of 304 Stainless Steel Containers after Exposure in Oxygen Saturated Water, Magnification 3/5X



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ومحتر فالمطالبة ومحاودة والمتعارية والمتراجع والمحاري والمتحال والمتحا والمحالية والمحارك والمحارية والمعارية





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Figure 12. Interior of Inconel 718 Containers after Exposure in Oxygen Free Water, Magnification 3/5X

24 MONTHS

NO. 1-2

18 MONTHS

NO. 1-4



NO. 7 24 MONTHS

NO. I-1 18 MONTHS







والمتعادية والمتحدث والمحارب والمحارب

Contraction of the



NO. 8

24 MONTHS

18 MONTHS

NO. 7













MAGNIFICATION 11X



MAGNIFICATION 100X

Figure 16. Weld Crack in A-286 No. A-1 Container

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ECHANK OF CONTAINER ANALYSES

	Tank	Exposite				Metallo	tarion tarion
Material	Ident.	Period (mos.)	5-10X Inspection	40X Inspection	40X Weld Inspection	Micro	Macro
304 55	ø	18	ł	I	ł	1	ł
304 SS	N9	18	ł	ł	ł	ł	ł
304 SS	1	18	ł	ł	ł	I	ł
304 SS	M	16	ł	1	t	ł	ł
304 SS	8	24	Dull Finish	ł	1	1	1
304 SS	NZ	24	Dull Finish	ł	ł	ł	I
304 SS	٢	24	ł	1	ł	t	I
304 55	R	24	ł	ł	1	1	ł
A-286	A-10F	18	Slightly Etched from Pickling	1	Weld Crack	M	н
A-286	A-1074	16	Slightly Etched from Pickling	ł	I	ł	ł
A-286	A-2	18	Slightly Etched from Pickling	ł	ł	ł	I
A-286	NZ-4	18	Slightly Etched from Pickling	ł	1	ł	I
A-286	~	24	Slightly Etched from Pickling	ł	ł	I	I
A-286	Ŗ	24	Slightly Etched from Pickling	1	1	ł	I
A-286	4	24	Slightly Etched from Pickling	ł	ł	ł	I
A-286	3	24	Slightly Etched from Pickling	1	1	ł	1
17-4PB \$\$	P-13	18	Rough Surface where Heat Treat Scale Removed by Pickling	Metallic Granular Appearing Contamin- ation	Two Weld Cracks	ł	м
SS HJ4-/T	P-13W	18	Rough Surface where Heat Treat Scale Removed by Pickline	Metallic Granular Appearing Contamin- erion	:	ł	1

TABLE II (cont.)

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1.

	Material	Tank Ident.	Exposure Period (mos.)	5-10% Inspection	40X Inspection	40% Weld Inspection	Metallo Docume Micro	Ngraphi Intatio Nac
	17-4PH SS		18	ł	1	Weld Crack	м	м
	SS HJY-LT	180 - A	18	1	:	Weld Crack	ł	1
	17-4PH SS	P-14	24	I	I	3 Weld Cracks	Ħ	1
	17-4PH SS	P-14H	24	ł	ł	2 Weld Cracks	ł	ł
	17-4PH SS	P-4	24	ł	ł	3 Weld Cracks	ĸ	м
	17-4PH SS	P-4W	24	I	ł	Weld Crack	I	I
	Inconel 718	4	18	Dark Area at Edge of Weldments	t	Porous Appearance at Edge of Welds. Weld Crack.	м	ĸ
	Inconel 718	1-44	18	Dark Area at Edge of Weldments.	ł	Porous Appearance at Edge of Welds. 2 Weld Cracks.	M	1
	Incomel 718	1-1	16	ł	ł	I	ł	ł
29	Inconel 718	NI-1	16	I	ł	1	ł	ł
	Inconel 718	1-2	24	I	ł	1	ł	ł
	Incomel 718	HZ-I	24	I	I	Weld Crack	н	×
	Inconel 718	~	24	1	ł	Weld Crack	н	ł
	Incomel 718	R	24	ł	1	ł	I	ł
	Titanius 6Al-47	~	16	Spote of Oxide, Stains from Contam- intants during Heat Treatment	Fite and Circular Knifeline Attack in Stained Areas	ł	I	ł
	Ticanium 6Al-4V	2	18	Spote of Oxide, Stains from Contam- inants during Heat Treatment	Fite and Circular Enifeline Attack in Stained Areas	ł	I	I
	Titanium 6A1-4V	4	16	Small Amounts of Oxide and Stain	ł	ł	ł	ł
	Titanium 6A1-4V	44	16	Small Amounts of Oxide and Stain	1	I	ł	1

TABLE II (cont.)

Material	Tank Ident.	Exposure Pericd (mos.)	5-10X Inspection	40X Inspection	40X Weld Inspection	Metallo Documen Micro	rephic tation Macro
Titanium 6Al-4V	æ	24	Spots of Oxide. Stains from Contam- inants during Keat Treatment. Finger- print stains.	Pits and Circular Ruifeline Attack in Stained Areas. Fingerprint Stains.	ł	×	ł
Ticanium 6Al-4V	84	24	Spots of Cxide. Stains from Contar- inants during Heat Treatment. Finger- print stains.	Pits and Circular Knifeline Attack in Stained Arcas. Fingerprint Stains.	1	ł	ł
Titanium 6Al-4V	Ś	24	Small Amounts of Oxide. Slight Staining.	ł	ł	1	ł
Titanium 6Al-4V	3	24	Small Amounts of Oxide. Slight Staining.	ł	ł	ł	ł



MAGNIFICATION 11X



MAGNIFICATION 100X

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Figure 17. Weld Crack in 17-4PH Stainless Steel No. P-3 Container. Crack is Enlarged Due to Crevice Corrosion During Pickling



MAGNIFICATION 11X



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MAGNIFICATION 8X





MAGNIFICATION 11X



MAGNIFICATION 50X

Figure 21. Weld Crack in Inconel 718 No. I-4 Container



MAGNIFICATION 11X



MAGNIFICATION 8X

Figure 22. Weld Cracks in Inconel 718 No. I-4 (Top) and No. 7 (Bottom) Containers



MAGNIFICATION 8X



MAGNIFICATION 100X

Figure 23. Discontinuity in Weld of Inconel 718 No. I-2 Container



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B. Discussion of Results (cont.)

c. 17-4PH (H1025) Stainless Steel

One of the four containers, No. P-13 had the same condition found in all six and twelve month storage containers; i.e., localized roughing of the surface due to scale removal during pickling. The tightly adherent metallic particles found in all previously examined containers and identified as 17-4PH stainless steel was also found in the No. P-13 container. Since this contaminant is identical in appearance to those previously analyzed, a chemical analysis was not performed. All four containers contained weld cracks. Representative photomacrographs and photomicrographs of these cracks are shown in Figures 17 through 20. These cracks occurred during welding and are not the result of water storage.

d. Inconel 718

As with the six and twelve month storage containers, examination of the interior surfaces at moderate magnification revealed a dark, porous area at the edge of all closure weldments. It could not be definitely established whether the condition is selective attack during pickling, alone, or in combination with entrapment of oxides due to inadequate back-up gas coverage during welding. The condition is not attributable to water storage. Three of the four tanks contained weld cracks. Representative photomacrographs and photomicrographs are shown in Figures 21 through 23. The network devolute pattern of the crack illustrated in Figure 21 is either shrink porosity or areas where entrapped oxides from welding were removed by pickling. Metallographic examination of the weld crack shown in Figure 23 indicated that the defect was not a crack but a discontinuity formed by the removal of entrapped oxides by pickling. Porosity similar to that shown in Figure 21 was also found.

e. 6A1-4V Titanium

All containers exhibited small isolated areas of titanium oxide from the heat treatment (aging) that were not removed during pickling. Examination of mottled areas at small magnifications revealed that these areas are primarily stain. Portions of these stained surfaces contained crater-like areas outlined by knifelike corrosion. Metallographic examination of sections taken through these areas revealed no detectable depth to the corrosion effects associated with the stained areas. Areas where stains appeared in the form of fingerprints, Figure 24, were given closer scrutiny to detect possible hot-salt stress corrosion effects, since the containers were aged in the contaminated condition. No defects attributable to water storage were found.

B. Discussion of Results (cont.)

3. Implications of the Results of the Examination

Discrepancies found in the internal surfaces of the containers were due to fabrication and cleaning procedures, and not due to the stored water, The discrepancies were the same as those found in the six and twelve month storage containers with no new defects being observed. The most serious of these discrepancies, with regard to influencing the function of the containers for storing transpiration coolant water, is the generation of a smut on the Inconel 718 and 17-4PH stainless steel and the deposition of parent metal granules on one of 17-4PH stainless steel during pickling. However, neither of these irregularities have produced particles of sufficient size and quantity to functionally degrade the water quality. The primary cause of these conditions is attributed to poor inert gas coverage on the container interiors during heat treatment and the resulting excessive oxidation. The presence of small amounts of oxide on the titanium alloy cannot be readily explained, because the containers were aged at 1000°F in air and the interiors were pickled for longer times than the exteriors from which all oxides were removed.

The presence of contaminants in the titanium alloy containers prior to heat treatment and the weld cracks found in the A-286, Inconel 718, and 17-4PH stainless steel containers are attributable to inadequate process control on the part of the container supplier. In spite of the presence of the contaminants, the water quality was not significantly impaired.

All discrepancies found on the internal surfaces of the containers were due to fabrication and cleaning procedures and not due to the presence of water in the containers.

SECTION III

CONCLUSIONS AND RECOMMENDATIONS

A. CONCLUSIONS

The following conclusions may be drawn from the twenty-four month storage evaluation program.

1. All five container materials, 304 and 17-4PH stainless steels, A-286, Inconel 718, and 6A1-4V titanium, are suitable for the storage of water for transpiration cooling purposes.

2. Both the oxygen-free and oxygen-saturated waters demonstrated favorable storability characteristics with regard to pH changes, ionic contamination as evidenced by electrical resistance measurements, and particulate formation as evidenced by Silting Index values and microscopic inspection of filters.

3. Based on the biological tests, there is no evidence that biological growth has occurred in the containers during storage periods up to twenty-four months.

4. Discrepancies in the containers such as cracks, smut and etched surfaces were produced during fabrication, cleaning, and passivation of the containers. Yet with adequate flushing prior to storage, the stored water is suitable for use as a transpiration coolant.

B. RECOMMENDATIONS

1. Long-term storage tests should be conducted with the selected waters in containers in which dissimilar metals are present to demonstrate which combinations of metallic materials can be used as suitable components in a transpiration coolant system.

2. The ease of fabrication, cleaning, and passivation of containers should be given proper priority in the selection and design of container materials for transpiration coolant devices.

APPENDIX A

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FABRICATION AND TREATMENT PROCEDURES FOR WATER CONTAINERS

FABRICATION AND TREATMENT PROCEDURES FOR WATER CONTAINERS

The fabrication procedures, the heat treatment cycles, the cleaning procedures, and the passivation procedures for the water containers prior to filling are presented below.

A. CONTAINER FABRICATION

Drawings of the long term storage container and its associated weld tooling are shown in Figures A-1, A-2, and A-3. The sequence of operations in the container fabrication is as follows:

1. Cylindrical Tube, -1

a. Fabricate -1 cylinder tube by rolling the sheared sheet stock into the desired diameter.

b. Weld the longitudinal joint using the automatic GTA* welding process. Full weld penetration must be obtained using gas backup for the welds.

c. Dye penetrant inspect welds then trim tubes to required length. No cracks allowed.

d. Machine tube ends for weld joint preparation.

e. Clean tubes and store in appropriate container while awaiting for next assembly.

2. <u>Fill Tube, -2</u>

a. Section fill tubes to 6 + 0.12 in. lengths.

b. Deburr fill tube ends and inspect.

c. Clean fill tubes and then package individually and store for next assembly.

3. Tank Head, -3

a. Blank tank heads to the desired diameter by punching or sawing and then machining.

b. Machine weld joint preparation at outer edge as required.

c. Drill the fill tube hole to mate with the fill tube.

*Gas Tungsten Arc

Page A-1



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Figure A-1. Long Term Storage Container

Page A-2









Page A-4

A, Container Fabrication (cont.)

d. Counter bore the tank head on one side at the fill tube hole for weld joint preparation.

e. Inspect tank heads, clean and package.

4. Tank End, -4

a. Perform operations 3, a, 3, b, and 3, e as above.

5. Assembly Sequence

a. Assemble -2 fill tube into -3 tank head using appropriate weld fixture and tooling.

b. Weld root pass on grooves side without weld filler wire by the automatic GTA welding process.

c. Clean root pass by rotary wire brushing using a clean stainless steel wire brush.

d. Weld the cover pass using the appropriate weld filler wire.

e. Reposition the part in the weld fixture and then place a fillet weld at the fill tube to tank head junction.

f. Inspect welds and clean part.

g. Fixture tank head assembly with -1 tube for welding and purge tank with Ar or He.

h. Weld root pass without the use of weld filler wire. Full penetration must be obtained at the ID of the joint.

i. Dye penetrant inspect and clean root pass. No cracks allowed.

j. Fill weld groove using the appropriate weld filler wire.

k. Inspect and clean the inner tank surfaces.

1. Locate the tank end to the tank for welding, then purge the closed tank with Ar or He.

Page A~5

A, Container Fabrication (cont.)

m. Make the root pass without the use of weld filler wire to insure full penetration.

n. Clean root pass surface.

o. Inspect the internal weld penetration using a borescope. Full weld penetration is required.

wire.

p. Run weld cover pass using the appropriate weld filler

q. Clean, inspect, and package unit for shipment.

Heat treatment of the containers was performed in an argon atmosphere and with one exception, as noted below, utilizing the thermal cycles listed in Table A-I.

Upon receipt of the tanks, the certifications were examined and it was found that some of the material used to fabricate the 304L stainless steel containers was actually 304 stainless steel. With the concurrence of the Air Force, the containers were subjected to a heat-treatment at 1925°F in hydrogen, followed by rapid cooling to preclude sensitization in the weld heat affected zone.

B. PREPARATIVE PROCEDURES

1. Container Cleaning and Passivation

All the containers were degreased by submerging and agitating the tanks three times in fresh isopropyl alcohol. The containers were then purged with dry, filtered nitrogen and placed in a vacuum chamber for final drying. The A-286, 17-4PH stainless steel, and Inconel 718 containers were then subjected to an alkaline descaling treatment for 60 minutes with Kelite No. 235 at a concentration level of 32 oz per gal at 190°F. The containers were then rinsed with water at 150°F for 2 to 5 minutes, followed by a rinse at ambient temperatures with deionized water for 2 to 5 minutes, then purged with dry, filtered nitrogen, and finally dried in a vacuum chamber.

The 304, A-286, and 17-4PH stainless steels and Inconel 718 were descaled in a pickling solution of 20% HNO_3 , 5% HF, and 75% H_2O at 130°F for 30 minutes per immersion until the last traces of scale were removed as confirmed by examination with a borescope. Following the acid treatment, the tanks were flushed with tap water for 2 to 5 minutes, then flushed with deionized water for 2 to 5 minutes, then subjected to ultrasonic vibration in a water bath for 15 minutes, flushed with deionized water for 5 minutes, purged with dry, filtered nitrogen, and finally dried in a vacuum chamber.

Page A-6

TABLE A-I

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HEAT TREATMENT PROCEDURES FOR WATER CONTAINERS

	uald Uive			
Material	Material	Heat Treatment	<u>Surface Finish</u>	
304L Stainless Steel	ı	·	Alkaline Clean, Pickle an Passivate	ğ
304L Stainless Steel	308L	t	Alkaline Clean, Pickle an Passivate	P
A-286	ı	Solution Treat and Age - MIL-H-6875	Alkaline Clean, Pickle an Passivate	ğ
A-286	A-286	Solution Treat and Age - MIL-H-6875	Alkaline Clean, Pickle an Passivate	2
17-4 PH Stainless Steel	ı	Age to H-1025 Condition - MIL-H-6875	Alkaline Clean, Pickle an Passivate	P2
17-4 PH Stainless Steel	17-4	Solution Treat and Age to H-1025 Condition - MIL-H-6875	Alkaline Clean, Pickle an Passivate	P L
Inconel 718	ı	Age (1400 - 1200°F)	Alkaline Clean, Pickle ar Passivate	ž
Inconel 718	Inconel 718	Solution Treat (1950°F) and Age (1400-1200°F)	Alkaline Clean, Pickle ar Passivate	Ĕ
6A1-4V-Titanium	6Al-4V Titanium	Age (1000°F - 4 Hours) - Weld-Stress Relieve (1000°F - 4 Hours)	Pickle	

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B, **Preparative** Procedures (cont.)

The 304 and A-286 stainless steels, and Inconel 718 were passivated with a 30% $HNO_3/3$ % $Na_2Cr_2O_7$ aqueous solution at 130°F for 25 minutes, rinsed with tap water for 2 to 5 minutes, rinsed with deionized water for 2 to 5 minutes, subjected to ultrasonic vibration in water for 5 minutes, rinsed with deionized water for 2 to 5 minutes, purged with dry, filtered nitrogen, and dried in a vacuum chamber. The 17-4PH stainless steel received the same treatment with the exception of a 10 minute passivation time.

The 6A1-4V titanium was degreased in isopropyl alcohol as described previously for the other tankage and then descaled in a pickling solution of 33.2% HNO3, 1.6% HF, and 65.2% water at 140°F for 3 minutes. The titanium containers were rinsed with 130°F tap water for 2 to 5 minutes, followed by a rinse in deionized water for 2 to 5 minutes, then subjected to ultrasonic vibration in water for 5 minutes, rinsed with deionized water for 2 to 5 minutes, purged with dry nitrogen, and then dried in a vacuum chamber.

2. Container Sterilization, Filling, and Sealing

Following the final rinsing with deionized water and the drying of the container in a vacuum chamber, the containers were wrapped with reusable sterilization paper. The wrapped containers were then sterilized in an autoclave at 250°F with 15 psig steam for 30 minutes followed by a 30-minute drying period. The containers were then stored in the paper to maintain their sterile condition.

All the steps required to fill the containers with water were conducted in the sterile laminar flow bench. The tanks were removed from the wrapping paper in the laminar flow bench. The tanks were weighed empty; then weighed when filled completely with sterile water to determine the total volume of the tank. The water was drained out and the tank was rinsed once more with the sterile water. A sample of the rinse water was checked for pH, conductivity, and Silting Index. If the values indicated that particulate matter and dissolved species were not present, the tank was considered ready for filling; if the values indicated that contaminants were present, the tank was rinsed until there was no evidence of contamination.

Before the final filling with oxygen-saturated deionized water, the tank was purged with oxygen from a filtered supply. The tank was then filled with the water and a sample was withdrawn for pH, conductivity, and Silting Index measurements. The ullage was adjusted to the ten percent value by weighing the container and its contents; the ullage space was purged with the filtered oxygen; and the container was capped with a sterile, tapered plug made from the same material as the container. The plug was seated in the fill-tube by use of a hammer. The containers were filled with the oxygen-free, deionized water in an analogous manner except that filtered nitrogen was used instead of oxygen for purging and blanketing the container. B, Preparative Procedures (cont.)

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The final sealing of the containers was accomplished by GTA welding the fill-tube/plug interface. The welds were inspected visually for any apparent anomalies. None were found. Then the containers were placed in plastic bags and labeled for the long-term storage tests.

The sampling plan for the long-term storage tests is to remove one container of each material with the two types of water for inspection and evaluation every six months for a period of five years. The contents will be characterized with respect to pH, conductivity, particulate content, and biological activity; and the containers themselves will be subjected to metallurgical examination if the other test data indicate that this is required. APPENDIX B

فالخطفين فالاطراب فتر وتغديزا فنافعا فالعماقات مطالب والنجاب فكالمحافظ فكالاجرمامي تسايد ووجعه الالاحتلاف والم

- Contraction

SILTING INDEX MEASUREMENT

SILTING INDEX MEASUREMENT

In order to obtain a measurement of the clogging tendencies of the particulate matter which may be produced during the storage of water in the presence of metals and non-metals for prolonged periods of time, a flow test of the water samples through nominal 1 micron size pores was required. Due to the limited quantity of the water available for each sample, approximately 20 ml, the ASTM Method F 52-69, "Silting Index of Fluids for Processing Electron and Micro-electronic Devices", was selected as being appropriate for the program.

The method determines the silting or clogging tendency of a fluid containing fine particles and gelatinous materials suspended in the fluid. The fluid is filtered through a membrane filter having a uniform pore size of 0.8 micron at a constant differential pressure. Particles lerger than 5 microns form an open network above the filter and do not affect the clogging tendency; particles smaller than 5 microns tend to block the flow passages of the filter and cause a decay in the flow rate. The rate of flow decay is expressed in terms of a Silting Index value; the greater the value, the greater the clogging tendency. A schematic diagram of the apparatus is shown in Figure B-1.

The total volume of fluid passed through the filter is 12 ml. The flow of the last 10 ml is timed incrementally as V_1 (1 ml), V_2 (5 ml), and V_3 (10 ml) with T_1 , T_2 , and T_3 the times required to flow the respective volumes. The Silting Index value is calculated from the equation:

S.I. =
$$\frac{T_3 - 2T_2}{T_1}$$

The tests in the program were conducted with three silting heads, No. 1 with an effective area of 1.0 mm^2 , No. 2 with an effective filter area of 4.3 mm^2 , and No. 3 with an effective filter area of 18.5 mm^2 . The procedure used in the tests was that prescribed in ASTM Method F52-69 except that samples were tested in triplicate only when sufficient sample was available. A photograph of the apparatus* used is shown in Figure B-2.

Water which had passed through the 0.22 micron pore size absolute filter and which was used in preparation of the tests, always produced a Silting Index value of less than 1 using the No. 1 silting head which has a crosssectional area of 1 mm^2 . The effect of particle size on the Silting Index was evaluated using latex spheres with a mean particle size of 1.25 microns with a range from 0.5 to 2.0 microns and puff ball spores with a size range from 3 to 4 microns. These data are presented for silting heads Nos. 1, 2, and 3 which have cross-sectional areas of 1 mm^2 , 4.3 mm², and 18.5 mm², respectively, in Figures B-3 and B-4. The particle concentration is given in mg/l because the suspensions were prepared on a weight basis; the actual

*Available from Millipore Corp., Bedford, MA.



Figure B-1. Schematic of Silting Index Apparatus

Page B-2



Figure B-2. Silting Index Apparatus

Page B-3



Figure B-3. Concentration Effect of 1.25 Micron Average Diameter Latex Particles on the Silting Index Values





Figure B-4. Concentration Effect of 3-4 Micron Diameter Puff Ball Spores on the Silting Index Value



densities of the particles are comparable. The significant item to note from the comparison of the data is that the Silting Index values are comparable at concentration levels that differ by at least one order of magnitude, the smaller particle producing the higher Silting Index value.