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TREATMENT OF DISASSEMBLY BASIN WATER

F. A. Locke
Pile Engineering Division

October 1954

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Explosives Department — Atomic Energy Division
Technical Division — Savannah River Laboratory

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TREATMENT OF DISASSEMBLY BASIN WATER

by

F. A. Locke
Pile Engineering Division

October 1954

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ABSTRACT

Corrosion of steel piping and basin equipment was found to be the main cause of the deposition of a brown flocculent material on the floor and walls of the disassembly basin at R Pile.

A combination of sodium dichromate and chloramine, the best of seven water treatments tested, reduced steel corrosion 50 per cent, zinc corrosion 80 per cent, and met the standards for control of turbidity and bacteria in the water of the basin.

TREATMENT OF DISASSEMBLY BASIN WATER

INTRODUCTION

A few weeks after water was charged to the disassembly basin of the R Pile, deposits of material containing silica, iron, aluminum, and manganese began to accumulate on the walls and floor of the basin.

The objectives of this study were to determine the cause of the deposition and to evaluate possible corrective measures. This work led to a laboratory investigation of the effect of various water treatments on water turbidity, bacteria control, and corrosion.

SUMMARY

Corrosion of the supply pipe line and the basin equipment is an important cause of deposition in the disassembly basin of the R Pile. None of the water treatments tested will completely eliminate this deposition, but a combination of sodium dichromate and chloramine, the best of the water treatments tested, reduced steel corrosion 50 per cent, zinc corrosion 80 per cent, and afforded complete protection from bacteria.

In low concentrations, only chloramine and a combination of sodium dichromate with chloramine met the turbidity and bacteria control standards for basin water. As a result of these tests, chlorine will be replaced by chloramine on a trial basis as a bactericide for basin water treatment.

Water additives tested were chlorine, chloramine, Calgon, Arquad T-2C, Santobrite, and sodium dichromate. Deionized water was also investigated.

DISCUSSION

BACKGROUND

The storage, disassembly, and transfer of irradiated material discharged from a Savannah River reactor are conducted under water in a concrete basin. The necessity of viewing underwater operations requires that the turbidity of this water be not greater than 0.5 ppm (as SiO_2). Therefore, coagulated and filtered water is used. This water is pumped from the water purification station through steel pipe at a maximum rate of 2000 gpm. A bactericide is added to the water to assure that bacteria which could increase turbidity do not grow in the basin.

A few weeks after water was charged to the disassembly basin of the R Pile, deposits of material containing silica, iron, aluminum, and manganese began to accumulate on the walls and floor of the basin. This condition reduced visibility by reducing the amount of reflected light. Also, manganese is a scavenger of fission product activity and its presence greatly increases the difficulty of decontamination in the event the basin must be drained for repairs.

EQUIPMENT

The experiments consisted of feeding filtered water treated by various methods into Amercoated 55-gallon drums at a rate (66 cc per minute) such that the turnover in the drums was comparable to the maximum turnover in the basin. A small section of carbon steel pipe was included in each of the feed lines to simulate basin supply pipe (see Figure 1).

PROCEDURES AND RESULTS

Three drums were charged with filtered water, filtered water and chlorine, and filtered water and chloramine⁽¹⁾ to determine if the addition of chlorine were causing the deposition. Within a week, a brown precipitate, similar in appearance to that deposited in the disassembly basin at R Pile, was noticed on the bottoms of the drums. The greatest deposition occurred in the drum fed with chlorinated water and the least occurred in the drum fed with filtered water.

Turbidity measurements were made with a Hellige turbidimeter on samples taken immediately before and after the water passed through the steel supply pipes to the drums. For each line it was found that the turbidity of the water increased while passing through this section of pipe. This strongly suggested that the precipitation in the drums and in the disassembly basin was caused by deposition of the products resulting from the corrosion of the pipeline and basin equipment.

To prove or disprove this theory, filtered water was charged to a drum without being passed through a carbon steel pipe. There was no deposition in the drum and the turbidity remained constant at a value of less than 0.1 ppm (as SiO_2) during the two weeks of the test. This is in contrast to the first three experiments in which the water in the drums leveled off at an average turbidity of 0.2 ppm. Data typical of these initial experiments are given in Table I. All chlorine and chloramine residuals were measured using a comparator and the ortho-tolidine-arsenite method(2).

The preceding experiments indicated that the deposition is caused by corrosion and not by inefficient filtration, after-precipitation, or bacterial action.

Chlorine was not completely efficient as a bactericide for basin water because chlorine is absorbed by organic material present in the water. In addition, an effective chlorine residual can not be maintained in the basin because of the slow rate of water turnover.

Therefore, further experiments were set up to find a water treatment that would inhibit the corrosion and at the same time meet the turbidity and bacteria control standards for basin water. Water additives tested were chlorine, chloramine, Calgon, Arquad T-2C, Santobrite, and sodium dichromate. Tests were also made with deionized water.

WATER TREATMENTS

The results obtained with each of the water treatments are discussed in the following paragraphs.

Chlorine

In the existing water treatment, 0.5 ppm of chlorine was added to the filtered water to inhibit the growth of bacteria in the basin. However, during the tests the Cl_2 residual of process water decreased from an average value of 0.5 ppm to 0.35 ppm while passing through the supply pipe to the test drum. Upon reaching the drum the chlorine residual decreased further to less than 0.1 ppm throughout the drum through absorption of chlorine by the organic matter present in the water and escape of chlorine from the water. Bacteria counts showed a steady increase in the number of bacteria present in the drum, indicating that chlorine is not an effective bactericide for basin water treatment unless larger amounts of chlorine which increase corrosion, are used. Actual conditions in the disassembly basin paralleled these test results.

Chloramine

When 0.5 ppm of NH_4OH was added to process water

containing 0.5 ppm of chlorine to form a chloramine, the Cl_2 residual of the water did not decrease while passing through the supply pipe as it did with chlorine alone. The chlorine residual throughout the drum remained steady at a value of 0.35 ppm and bacteria counts showed that there were no bacteria growing in this drum.

Taylor⁽³⁾, supports these observations as he states, "Chloramines are less active oxidizing and germicidal agents than chlorine but they are more stable, and hence an active residuum remains for a longer period (several days) in the treated water. This is an advantage where the water is retained for long periods in reservoirs and mains before delivery to the consumers, since a high standard of bacterial purity can be preserved and after-growth of bacteria prevented".

Corrosion of steel, aluminum coupled with stainless steel, and zinc coupled with stainless steel, was reduced slightly. Relative corrosion rates are listed in Table II.

Chlorine and Calgon

Two ppm of Calgon added to the process water reduced steel corrosion 20 per cent but increased aluminum corrosion twenty-fold. This accelerated corrosion of aluminum eliminated Calgon from further consideration.

Arquad T-2C

Five ppm of Arquad T-2C, the required concentration for effective protection against bacteria, had a turbidity of 0.4 ppm (as SiO_2). After corrosion samples were placed in the drum, a high turbidity, ranging up to 1.25 ppm, developed. This is much higher than the maximum allowable turbidity of 0.5 ppm.

Santobrite

Thirty ppm of Santobrite, the concentration necessary for control of bacteria, had an unacceptably high turbidity of 2.5 ppm.

Chloramine and Sodium Dichromate

The use of 2.0 ppm sodium dichromate and 0.5 ppm chloramine reduced steel corrosion by 50 per cent and the corrosion of zinc coupled with stainless steel by 80 per cent, and at the same time afforded complete protection against bacteria. The turbidity of filtered water treated in this manner was less than 0.1 ppm.

Deionized Water

The corrosion of steel in deionized water was approximately four per cent as great as that observed in chlorinated filtered water. The corrosion of zinc coupled with stainless steel was reduced by 70 per cent while the corrosion of aluminum was very low. However, tests showed that no protection is offered against bacteria by deionized water exposed to the atmosphere.

RELATIVE CORROSION RATES

Comparative results of the corrosion tests after one month are reported in Table II. These results were obtained by measuring the weight loss of metal coupons after exposure to the treated water for one month.

None of the water treatments reduced the corrosion of steel enough to eliminate deposition of the products resulting from corrosion of the pipe line. However, sodium dichromate coupled with chloramine reduced steel corrosion by 50 per cent and at the same time afforded excellent protection from bacteria. Chloramine was the only bactericide tested that met the turbidity and bacteria control standards for basin water. As a result of these tests, chlorine will be replaced by chloramine on a trial basis at R Pile.

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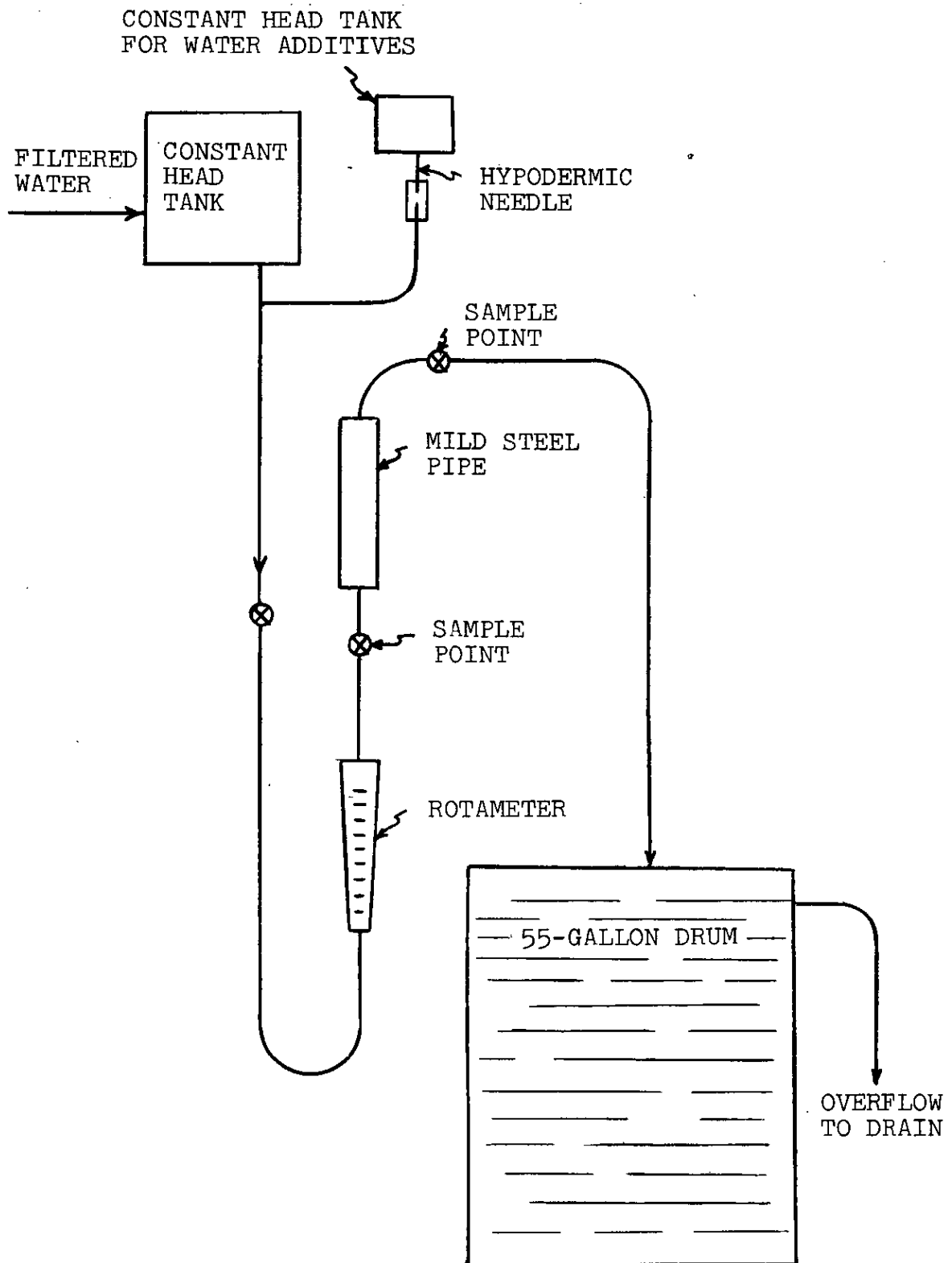


DIAGRAM OF EQUIPMENT

TABLE IDP - 81
Page 13RESULTS OF INITIAL EXPERIMENTS

	<u>Filtered H₂O</u>	<u>Chlorinated H₂O</u>	<u>Chlorinated H₂O / NH₄OH Chloramine</u>	<u>Filtered H₂O (No Pipe)</u>
<u>Turbidity, ppm as (SiO₂)</u>				
Before pipe	<0.1	<0.1	<0.1	--
After pipe	0.4	0.5	0.4	--
In drum	0.2	0.2	0.2	<0.1
<u>Chlorine residual, ppm</u>				
Before pipe	--	0.5	0.5	--
After pipe	--	0.35	0.5	--
In drum	--	<0.1	0.35	--
<u>Deposition</u>	Occurred in one week, grew heavier	Occurred in one week, grew heavier	Occurred in one week, grew heavier	None
<u>Bacteria</u>	Present	Present	None	Present

RELATIVE CORROSION RATES

(Relative to corrosion rate in chlorinated water)

<u>Water Treatment</u>	<u>Al (Coupled with Stainless Steel)</u>	<u>Zn (Coupled with Stainless Steel)</u>	<u>Steel</u>	<u>Al</u>
Chlorine	1.0	1.0	1.0	1.0
Chloramine	0.8	0.5	0.8	1.0
Calgon & Chlorine	1.2	0.8	0.8	20.0
Arquad T-2C	1.2	0.3	1.0	1.0
Chloramine & Sodium Dichromate	1.0	0.2	0.5	~0
Deionized H ₂ O	--	0.3	0.04	~0