

# LAKE HAVASU CITY WATER TREATMENT RESEARCH STUDY

Water Treatment Technology Program Report No. 8

September 1995

## **U.S. DEPARTMENT OF THE INTERIOR**

Bureau of Reclamation
Technical Service Center
Environmental Resources Team
Water Treatment Engineering and Research Group

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## LAKE HAVASU CITY WATER TREATMENT RESEARCH STUDY

Water Treatment Technology Program Report No. 8

by

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Water Treatment Engineering and Research Group Environmental Resources Team Technical Service Center Denver, Colorado

September 1995

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#### **ACRONYMS AND ABBREVIATIONS**

ADEQ Arizona Department of Environmental Quality

BPV back pressure valve
cfu colony forming units
ENR Engineering News Record

EPA Environmental Protection Agency

FCV flow control valve

HDPE high density polyethylene
HPC heterotrophic plate count
ICP inductively-coupled plasma

LHC Lake Havasu City

MCL maximum contaminant level

Mgal million gallons
NF nanofiltration

NPF normalized permeate flow NDP net driving pressure

ntunephelometric turbidity unitO&Moperations and maintenancePIDproportional integral derivativePLCprogrammable logic controller

RO reverse osmosis SDI silt density index

SEM scanning electron microscopy

SR salt rejection

TCF temperature correction factor

TDS total dissolved solids
TFC thin film composite
THM trihalomethane

THMFP trihalomethane formation potential

TOC total organic carbon

UV ultraviolet

#### **CHEMICAL FORMULAS**

Al<sup>+3</sup> aluminum ion

Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·14.3H<sub>2</sub>O aluminum sulfate (alum)

 ${
m Ba^{*2}}$  barium ion  ${
m Ca^{*2}}$  calcium ion  ${
m CaCO_3}$  calcium carbonate  ${
m Ca(OH)_2}$  calcium hydroxide (lime)

Cl chloride ion Cl<sub>2</sub> chlorine

ClO<sub>2</sub> chlorine dioxide
Cr chromium
Fe<sup>+2</sup> ferrous iron
Fe<sup>\*3</sup> ferric iron
H<sup>+</sup> hydrogen ion
HCO<sub>2</sub> bicarbonate ion

H<sub>2</sub>O water

## **CHEMICAL FORMULAS — CONTINUED**

$\mathrm{H_{2}SO_{4}}$	sulfuric acid
K*	potassium ion
KMnO <sub>4</sub>	potassium permanganate
Mg <sup>+2</sup>	magnesium ion
$Mn^{+2}$	manganese ion
MnO	manganous oxide
MnO <sub>2</sub>	manganic dioxide
MnO <sub>4</sub>	permanganate ion
Na <sup>+</sup>	sodium ion
Na <sub>2</sub> CO <sub>3</sub>	sodium carbonate (soda ash)
Ni	nickel
NO <sub>3</sub> -	nitrate ion
O <sub>3</sub>	ozone
	silica
SiO <sub>2</sub> SO <sub>4</sub> - 2	sulfate ion

## SI METRIC CONVERSIONS

From	То	Multiply by
ft	m	3.048 000 E-01
in	$\mathbf{m}$	2.540 000 E-02
$\mathbf{ft}^2$	${f m^2}$	9.290 304 E-02
kgal	$\mathbf{m^s}$	3.785 412
Mgal	$\mathbf{m}^3$	3.785 412 E+3
acre-ft	${f m^3}$	1.233 489 E+3
$lb/in^2$	kPa	6.894 757
$^{\circ}\mathbf{F}$	$^{\circ}\mathrm{C}$	$t_{\rm C} = (t_{\rm F} - 32)/1.8$

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#### 1. EXECUTIVE SUMMARY

The Bureau of Reclamation (Reclamation) and LHC (Lake Havasu City) conducted a jointly-funded research study to evaluate selected ground-water treatment options to assist the city in planning for water treatment expansion. The study included pilot testing at well S6, one of 14 city-owned wells. In this report, Reclamation is providing process recommendations, cost estimates, and design considerations for scale-up based on the results of this testing.

Based on available data, the city water supply is in full compliance with primary drinking water standards. However, concentrations of other constituents such as hardness, sulfate, TDS (total dissolved solids) and manganese exceed secondary MCLs (maximum contaminant levels). High manganese levels in several wells are of concern to the community because precipitates formed in the water from chlorination accumulate in the distribution system. Poor quality ground water is commonly made acceptable by blending with better quality wells. However, with established wells declining in quality, this approach is becoming impractical or unavailable.

Two processes were selected for pilot testing: KMnO<sub>4</sub> (potassium permanganate) oxidation followed by greensand filtration to remove Mn<sup>+2</sup> (manganese); and NF (nanofiltration) to reduce the concentrations of not only manganese, but sulfate, hardness, and TDS as well. Nanofiltration is capable of producing product water that meets *all* secondary MCLs.

KMnO<sub>4</sub> oxidation and greensand filtration were effective in reducing the concentration of  $Mn^{+2}$  in well S6 ground water to 0.05 mg/L, the secondary MCL. The optimum KMnO<sub>4</sub> dose was about 1.1 mg/L. Considering the average reduction of  $Mn^{+2}$  (0.62 mg/L [influent] - 0.05 mg/L [effluent]), this dose is equivalent to 1.93 mg/L KMnO<sub>4</sub> per mg/L  $Mn^{+2}$ , which is essentially the same as the stoichiometric requirement. The reaction is fast. About 90 percent of the  $Mn^{+2}$  is oxidized within the first 1-1/2 minutes. The initial reaction rate was determined to be second order dependent on both [ $MnO_4$ ] and [ $Mn^{+2}$ ] with a rate constant of -0.198/mol·sec.

Greensand effluent turbidity measurements were at or below 0.09 ntu (nephelometric turbidity units), indicating efficient filtration. The greensand filter media was also effective in controlling the over- and under-dosing of KMnO<sub>4</sub>. Interestingly, water samples collected just prior to filtration contained less  $Mn^{+2}$  (0.02-0.03 mg/L) than the filter effluent when operating near the optimum KMnO<sub>4</sub> dose of 1.1 mg/L (this observation is based on Hach analyses in the field). This result suggests that KMnO<sub>4</sub> could be removed more efficiently with conventional dual- or multimedia filtration, assuming an effective control could be employed for chemical dosing.

Jar test results showed that alum concentrations of 20 to 40 mg/L could be used to coagulate and settle  $MnO_2$  (manganic dioxide [oxidized  $Mn^{+2}$  particles]); however, the required settling time of 90 minutes was excessive and the floc produced was very fragile. Thus, this report concludes that alum is not effective for this application, and apparently is not necessary to produce acceptable filter effluent turbidities as described above.

The nanofiltration test system employed 18 FilmTec NF90-2540 elements in a 12:6 (2-stage) array. The feed flow rate and product recovery were maintained at 18.2 L/min (4.8 gal/min) and 80 percent, respectively. The initial feed pressure was 570 kPa (83 lb/in²). Hypersperse AF  $200^{\text{TM}}$  and sulfuric acid were added for scale control. The system was operated for about 1000 hours to allow time for any potential membrane degradation from fouling or scaling to develop. UV (ultraviolet) disinfection was used for the control of microbial contamination.

Nanofiltration effectively reduced the concentrations of all contaminants of concern to below MCLs. The average TDS rejection for the 984-hour test was 90.9 percent. The percent rejection for specific ions of interest was:  $Ca^{+2}$  (calcium) - 98.1;  $Mg^{+2}$  (magnesium) - 98.5;  $SO_4^{-2}$  (sulfate) - 97.4; and  $Mn^{+2}$  -  $\geq$ 90.3. The average TDS for the feed and permeate (product) was 826 and 75 mg/L, respectively. This low permeate concentration allows blending to obtain higher overall net product recoveries.

The test data show that biofouling probably occurred during the 6 weeks of operation. NPF (normalized permeate flow) dropped about 16 percent during this period, and feed pressure increased from 75 to 89 lb/in². A membrane autopsy was performed to determine the cause of the performance degradation. SEM (scanning electron microscopy) imaging was also used in an attempt to visually identify fouling or scaling components. The results of these procedures, along with high heterotrophic plate counts measured throughout the test system, all pointed to biofouling as the most likely cause. Because of high microbial populations present in the feed water (7900 cfu/mL measured at the wellhead sample tap), a more aggressive disinfection will be needed, e.g., the use of chloramines or chlorination followed by dechlorination. However, any Mn+² oxidized by these chemicals would have to be removed by media filtration prior to membrane desalting.

Lake Havasu City may choose a treatment plant for manganese removal alone, such as a potassium permanganate oxidation plant, or may elect treatment that would reduce the concentrations of all EPA-regulated primary and secondary drinking water parameters to below MCLs. This report concludes that several options are available to the city to provide the latter defined, full compliance, water quality. These options include nanofiltration alone or nanofiltration blended with water from either Lake Havasu or treated ground water. Lime-softened ground water can also be blended with treated water from either Lake Havasu or ground water that has received nanofiltration treatment.

If Lake Havasu City selects a nanofiltration treatment plant, the reject brine produced will have to be disposed of in a manner that is consistent with Arizona Department of Environmental Quality rules. Methods available to the city for reject brine disposal include evaporation ponds, reuse using irrigation of brine-tolerant plants or landscaped areas when the salt concentrations are appropriately adjusted (diluted), and creation of a wetlands environment.

#### 2. INTRODUCTION

#### 2.1 Purpose and Scope

In light of a rapid population increase over the last 15 years, Lake Havasu City is facing several challenges concerning their water supply and distribution system. One of these challenges is the need to provide a reasonable level of water treatment to ensure the delivery of safe and palatable drinking water to the residents of the city. To address this need, and to assist the city in planning for water treatment expansion, Reclamation and Lake Havasu City conducted a jointly-funded research study to evaluate selected ground-water treatment options.

The study included the pilot testing of selected water treatment processes to confirm their performance and efficiency. Based on the results of this testing, process recommendations and cost estimates are provided, along with design considerations for scale-up. The cost estimates include both capital and O&M (operation and maintenance) costs for a full-scale treatment plant with a capacity of 12 Mgal/day. Capital cost estimates are based on a combination of direct

quotes from manufacturers, plus allowances for installation, and cost curves prepared by the EPA (Environmental Protection Agency) using current indices of the ENR (Engineering News Record). O&M cost estimates include current prices for electricity, chemicals, and supplies, when available. Most estimates for materials, equipment, and labor are based on updated ENR values. Consequently, the estimates found herein are valuable for a comparison of the alternatives presented, and are not final construction estimates.

The final treatment process recommendations made in this report should be integrated with other design factors that address the city's comprehensive needs. It is recommended that consideration be given to such issues as capacity, water sources, desired level of treatment, and location, in an engineering analysis to arrive at a final determination of an appropriate treatment scheme.

#### 2.2 Background

Lake Havasu City receives all of its potable water supply from fourteen city-owned and operated wells located along the shoreline of Lake Havasu (fig. 1). The wells are grouped into three separate well fields: north, central and south. The total combined pumping capacity of the wells is about 24 Mgal/day (HDR Engineering, 1992).

Poor quality ground water in the study area is commonly made acceptable by blending with better quality wells. In the past, the city drilled wells to avoid poorer water quality areas. However, with established wells declining in quality, new wells must be located where only poor quality ground water is available and where blending is impractical or unavailable.

#### 3. CONTAMINANTS OF CONCERN

Based on available data, the city's water supply is in full compliance with all primary drinking water standards. However, the concentrations of other constituents such as hardness, sulfate, TDS (total dissolved solids), and manganese are in excess of secondary MCLs (maximum contaminant levels). High manganese (Mn<sup>+2</sup>) levels are of particular concern to the community because precipitates, formed in the water from chlorination, accumulate in the distribution system and cause discoloration of tap water. Elevated manganese concentrations are generally confined to the central and south well fields (wells C2, C9, S4 and S6).

#### 4. PILOT TEST OBJECTIVES

## 4.1 KMnO<sub>4</sub> (Potassium Permanganate) Oxidation

The principal objectives of the KMnO<sub>4</sub> (potassium permanganate) oxidation testing were to:

- Determine the concentration of KMnO<sub>4</sub> needed to reduce the Mn<sup>+2</sup> (manganese) level in well S6 ground water to below 0.05 mg/L (EPA secondary drinking water standard).
- Determine the required reaction time and attempt to develop an initial rate constant.
- Assess the ability of manganese-greensand filtration to compensate for over- and underdosing of KMnO<sub>4</sub>.

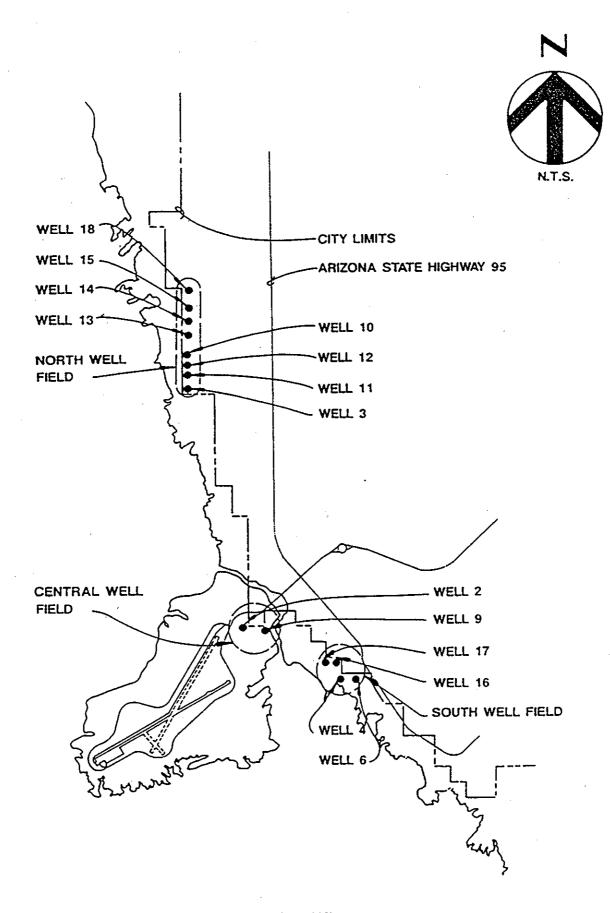


Figure 1. - Lake Havasu City well fields (HDR Engineering, 1992).

- Evaluate the efficiency of the greensand filter in removing MnO<sub>2</sub> (manganic dioxide [oxidized manganese particles]).
- Evaluate the effectiveness of alum (aluminum sulfate) for coagulating and settling  ${\rm MnO_2}$  precipitates.

#### 4.2 Nanofiltration

The principal objectives of the nanofiltration testing were to:

- Evaluate the performance of FilmTec NF-90 nanofiltration membrane elements for reducing TDS, hardness, sulfate, and manganese levels in the well S6 water.
- Assess blending opportunities (NF permeate with filtered well water) to achieve high overall net recoveries.
- Determine potential long-term adverse effects on the membranes from fouling or scaling.

#### 5. PILOT TEST DESCRIPTION

#### 5.1 Site Selection

Site selection for ground-water testing focused on the city-owned wells that contained manganese in 1993: C2, C9, S4, and S6 (fig. 1). Of these four wells, the 1992-93 average concentration for manganese and TDS indicated C2 and S6 as the leading candidates. The 2-year data base showed well C2 had an average of 0.28 mg/L manganese and 1227 mg/L TDS, and well S6 had an average of 0.50 mg/L manganese and 764 mg/L TDS. Because manganese is the primary contaminant of concern for Lake Havasu City, it was decided that testing at well S6 would yield the most useful data.

Table 1 shows the most recent chemical analysis (existing prior to this test program) for well S6. Corresponding primary and secondary federal drinking water standards are also included in the table for comparison. The constituents that exceed secondary MCLs are highlighted with an asterisk. Well S6, with a drilled depth of 160 feet, is rated at 900 gal/min and has a usable capacity of 550 gal/min.

#### 5.2 Process Selection

Two approaches to treatment were considered for evaluation in the field: the first, an oxidation and filtration process for the removal of manganese; and the second, a softening process (lime or membrane) to reduce the concentrations of not only manganese, but also sulfate, hardness, and TDS.

5.2.1 KMnO<sub>4</sub> oxidation. - The most effective oxidants available for the treatment of Mn<sup>+2</sup> (manganese) are O<sub>3</sub> (ozone), ClO<sub>2</sub> (chlorine dioxide) and KMnO<sub>4</sub>. Of the three, KMnO<sub>4</sub> is the more commonly used in the United States, and is generally favored for waters with high manganese content (Glase, 1990). Both O<sub>3</sub> and ClO<sub>2</sub> require on-site generation, which involves a greater equipment and capital investment. Cl<sub>2</sub> (chlorine) will also oxidize manganese, but care must be exercised when the THMFP (trihalomethane formation potential) of the water is high.

Table 1. - Available ground-water analysis for well S6.

				Feder	al MCL
Cations			Concentration	(Primary)	(Secondary
Calcium	Ca	mg/L	100		
Magnesium	Mg	mg/L	35		
Sodium	Na	mg/L	110		
Potassium	K	mg/L	1.8		
Strontium	Sr	mg/L	1.3		
Barium	Ba	mg/L	0.05	1	
Iron	Fe	mg/L	0.05		0.3
Manganese*	Mn	mg/L	0.55		0.05
Anions			•		
Bicarbonate	HCO <sub>3</sub>	mg/L	232		
Carbonate	CO <sub>3</sub>	mg/L	0		•
Sulfate*	SO <sub>4</sub>	mg/L	300		250
Chloride	Cl	mg/L	95		250
Nitrate	$NO_3$ (as $N$ )	mg/L	1	10	
Fluoride	F	mg/L	0.64	4.0	
Ammonia (N)	NH <sub>3</sub> (N)	mg/L	0.2		
Silica (soluble)	SiO,	mg/L	15		
Silica (total)	$SiO_2$	mg/L	17		
Alkalinity	as CaCO <sub>3</sub>	mg/L	190		
Hardness	as CaCO <sub>3</sub>	mg/L	390		
TDS (reported)*		mg/L	790		500
TDS (summation)*		mg/L	880		500
рН	•		7.62		6.5-8.5

A combination of KMnO<sub>4</sub> oxidation and manganese-greensand filtration was selected for testing. Manganese-greensand provides effective filtration and also controls under- and over-dosing of KMnO<sub>4</sub> (prevents the development of pink water breakthrough). Manganese (II) removal depends on the precipitation of  $MnO_2(s)$  (manganese [IV] [manganic dioxide]), as follows:

$$3Mn^{+2} + 2KMnO_4 + 2H_2O = 5MnO_2(s) + 2K^+ + 4H^+$$

Manganic dioxide is essentially insoluble over the entire pH range of interest in drinking water treatment. Also, the oxidation of both  $Mn^{+2}$  and  $Fe^{+2}$  (ferrous iron) using  $KMnO_4$  is reported to be quite rapid at pH 7 and higher (Glase, 1990).

The stoichiometry for manganese and iron oxidized with permanganate is:

- 1.92 mg/L KMnO<sub>4</sub> per mg/L of Mn<sup>+2</sup> removed
- $0.94~\text{mg/L}~\text{KMnO}_4~\text{per}~\text{mg/L}~\text{of}~\text{Fe}^{+2}~\text{removed}$

**5.2.2 Nanofiltration.** - Lime softening and membrane softening (nanofiltration) can be used to reduce hardness and TDS. Lime softening can remove carbonate hardness (calcium and magnesium bicarbonates) to the level of  $CaCO_3$  (calcium carbonate) solubility by stoichiometric additions of  $Ca(OH)_2$  (lime). Further reduction of noncarbonate hardness (calcium and

magnesium sulfates or chlorides) requires addition of  $Na_2CO_3$  (soda ash). Lime softening also causes effective chemical precipitation of manganese as MnO (manganous oxide) when the pH of the water is in the range of 9.5 to 10.0. No reduction in sulfate is achieved with this process.

RO (reverse osmosis) and NF are processes used for desalting water by the application of hydrostatic pressure to drive feed water through a semi-permeable membrane. Most of the water's impurity (dissolved salts) remain behind and is discharged as waste brine; relatively pure product water (permeate) emerges at near atmospheric pressure. A typical operating pressure range for RO is 200 to 400 lb/in² for brackish water and 800 to 1000 lb/in² for seawater desalination. Ion rejections achieved with RO usually are in the mid-to-high 90-percent range.

NF membranes are generally used to treat low TDS waters where the reduction of hardness ions is desirable. The rejection of divalent ions (Ca<sup>+2</sup>, Mg<sup>+2</sup>, SO<sub>4</sub><sup>-2</sup>) and organics having a molecular weight above 200 is very high, typically above 95 percent; thus the name "softening membranes." Monovalent ions (Na<sup>+</sup>, Cl<sup>-</sup>, HCO<sub>3</sub>) are rejected at around 60 to 70 percent. Typical applications for NF include the removal of TDS, hardness, color, THM (trihalomethane) precursors, TOC (total organic carbon), and radium. Operating pressures are in the range of 75 to 150 lb/in<sup>2</sup>, depending on temperature, feed-water constituents, salinity, and recovery (FilmTec Corporation, undated).

Both softening processes have disposal requirements: CaCO<sub>3</sub> sludge with lime precipitation; and reject brine with NF.

NF was selected for field testing because of its ability to meet *all* secondary MCLs, and because of blending opportunities to achieve high overall net recoveries (reducing brine disposal requirements).

### 5.3 Pilot Plant Equipment and Site Layout

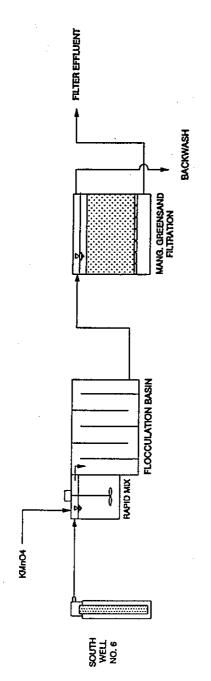
Reclamation's Mobile Water Treatment Pilot Plant was used at Lake Havasu City for the field testing described herein. This pilot plant incorporates skid-mounted equipment to test many unit treatment processes, including chemical precipitation, oxidation (ozone, permanganate), ion exchange, activated carbon, and membrane separation. Most of the process equipment is controlled using an Allen-Bradley SLC 500 programmable controller, and provisions are included for automatic data acquisition; however, data acquisition for this test program was performed manually. A 35-kW generator is available for remote-site operations where commercial power is not available.

Figure 2 presents schematic diagrams of the treatment processes that were pilot tested at Lake Havasu City:

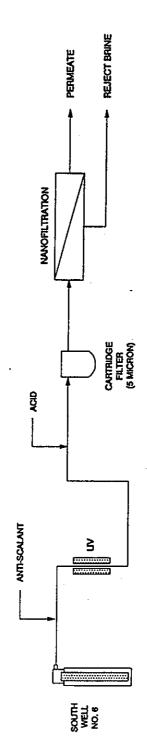
- KMnO<sub>4</sub> (potassium permanganate) oxidation followed by manganese-greensand filtration
- Nanofiltration preceded by anti-scalant addition and UV (ultraviolet) disinfection

Post-chlorination was not included in the testing, but would be incorporated as part of the prototype treatment plant(s).

Figures 3 and 4 show the equipment setups and interconnections (plumbing and level sensor) for the two processes.



Potassium permanganant (KMnO4) oxidation; greensand filtration



Anti-scalant addition; UV disinfection; nanofiltration

Figure 2. - Ground-water treatment processes selected for pilot testing.

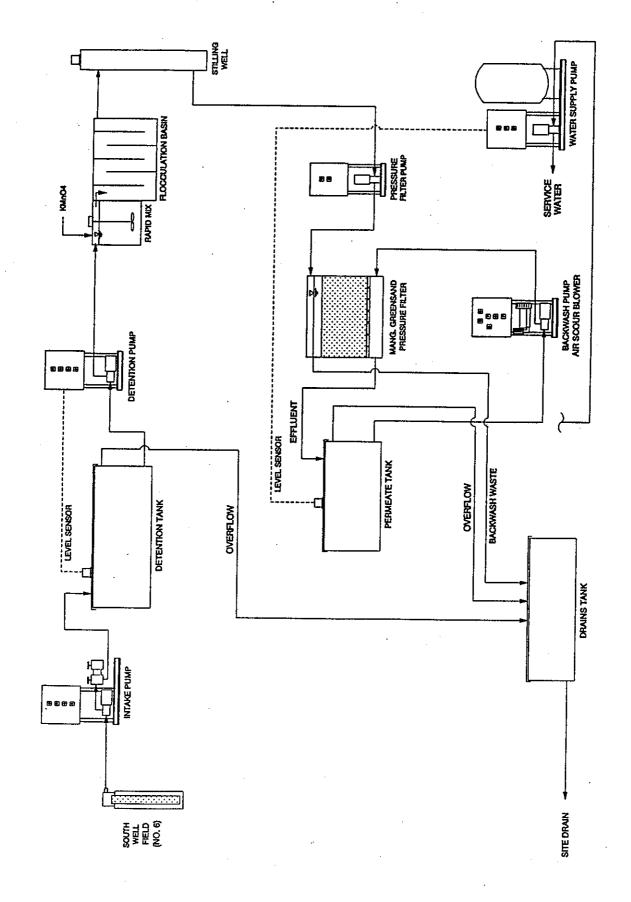


Figure 3. - Equipment setup for ground-water treatment process No. 1 (KMnO, oxidation and manganese-greensand filtration).

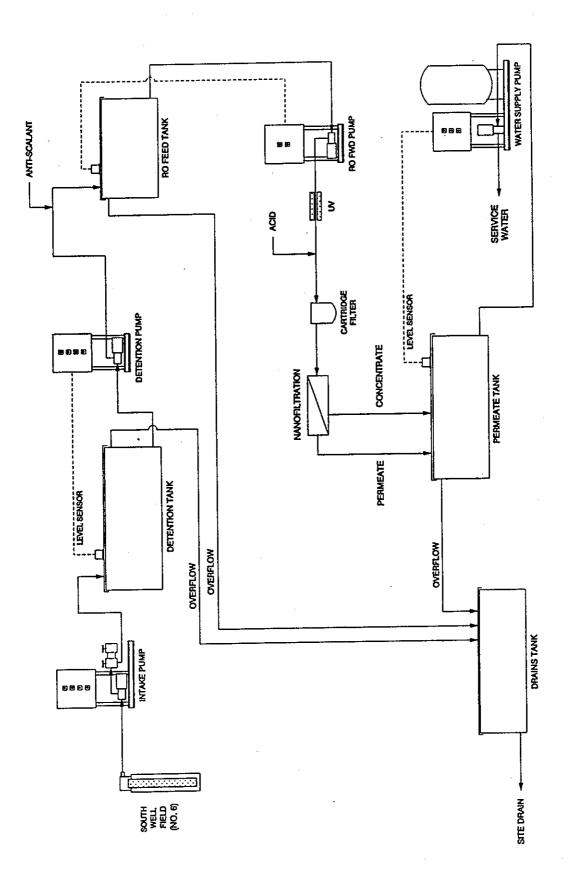


Figure 4. - Equipment setup for ground-water treatment process No. 2 (anti-scalant, UV disinfection, and nanofiltration).

Figure 5 shows the actual layout of the trailer, tanks, and exterior skids at well S6. The intake pump skid (not shown) was used to transfer feed water from the well to the intake pump connection on the trailer located just behind the stairway. All process effluent and drain flows were directed to the gravel drain shown next to the fence line behind the trailer.

#### 5.4 Test Procedures

5.4.1 KMnO<sub>4</sub> oxidation. - From the stoichiometry described earlier, the concentration of KMnO<sub>4</sub> required to completely oxidize 0.55 mg/L Mn<sup>+2</sup> and 0.05 mg/L Fe<sup>+2</sup> would be 1.10 mg/L. Based on this requirement, the range of KMnO<sub>4</sub> concentrations selected for evaluation in this test phase was 0.60 to 1.80 mg/L (this range represents 55 to 165 percent of the stoichiometric value).

Initially, each test was run at 22.7 L/min (6 gal/min), which provided about 55 minutes of reaction time before greensand filtration. Later, four additional tests were run at a reduced reaction time of 12 minutes. For each test, the KMnO<sub>4</sub> feed concentration was set in the morning and allowed to stabilize for about 10 hours. During this time, physical parameters (flow, pH, turbidity, and filter  $\Delta P$ ) were monitored and manually recorded on the operator's data sheet. Four recordings were made each day. At the end of each day's testing, two separate samples were collected of the raw feed and the greensand filter influent and effluent. One set was sent to a contract lab for manganese and iron analyses (refer to table 2), and a second set was analyzed in the field using Hach reagents. The two sets of data were later compared as a quality-control measure. Immediately after collection, the greensand filter influent samples were vacuum-filtered into a flask containing sodium bisulfite (1.5 times stoichiometric). This procedure was done to remove accumulated oxidation particles and to stop the reaction by neutralizing any residual KMnO<sub>4</sub>.

An LMI (Liquid Metronics, Inc.) pump was used to feed a 0.5-percent KMnO<sub>4</sub> solution directly to the rapid mix tank. The chemical feed rates for a process flow of 22.7 L/min were as follows:

KMnO <sub>4</sub> dosage (mg/L)	mL/min of 0.5% KMnO <sub>4</sub>
0.6	2.72
0.8	3.63
1.0	4.54
1.2	5.45
1.4	6.36
1.6	7.26
1.8	8.17

Permanganate dosages were checked daily by observing and recording the level change in the chemical feed tank. Also, periodic spot checks of the feed rate were made using a 10-mL graduated cylinder and stopwatch.

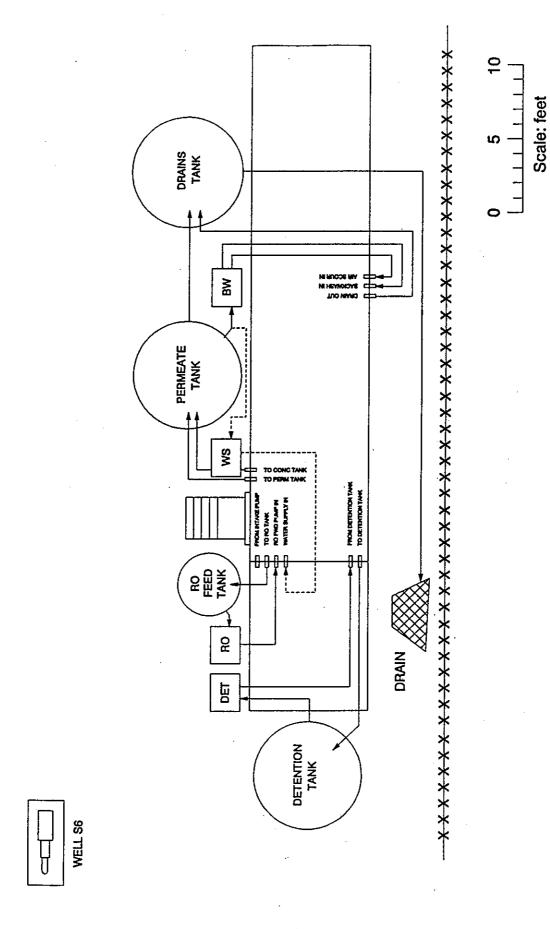


Figure 5. - Pilot plant equipment layout at the well S6 site.

Table 2. - Lake Havasu City water treatment study analytical requirements.

Maximum Holding Time	, , , , ,	28 days 28 days 28 days 28 days	6 months 6 months 6 months 6 months	28 days 28 days 28 days 48 hours 48 hours 6 months	28 days 7 days	8 hours
Minimum Volume (ml.)		500 500 500 500	250 250 250 250	300 300 100 100 100 100 100 100 100 100	100	0 <b>0</b> · ·
Container Type	· • • • • • •	Plastic Plastic Plastic Plastic	Plastic Plastic Plastic Plastic	Plastic, glass Plastic Plastic, glass Amber plastic, glass Amber plastic, glass Plastic, glass Plastic, glass	Plastic Amber glass; TFE cap	Sterilized glass, plastic
Preservation	• • • • • • • • • • • • • • • • • • •	Store at 4 deg C Store at 4 deg C Store at 4 deg C Store at 4 deg C	Nitric, < pH 2	Store at 4 deg C None required Store at 4 deg C	Store at 4 deg C 4 deg C; HCI, < pH 2 & 4 drops 10% S.T.	Store at 4 deg C
Responsibility for Testing/Recording	Operator Operator Operator Operator Operator Operator	2222	LHC LHC LHC/Operator, Hach LHC/Operator, Hach	222222 222222	CHC CHC	LHC Operator Operator
Ser of Readings GW#2	Many Many Many Many Many 40	<del>à</del> à à à à	5 5 5 5 5 5	8 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	16	4
Number Samples/Re GW#1	Many Many Many Many	0000	~ <b>4</b> 4 4 4	n		Мапу Мапу
Units	L/min deg C NTU uS/cm	т9/L т9/L т9/L т9/L	тg/L тg/L тg/L тg/L	пф. пф. пф. пф. пф.	mg/L mg/L	CFUs/mL psig hours
Parameter	Flow Temperature pH Turbidity Conductivity Sill Density Index (SDI)	Calcium, Ca Magnesium, Mg Sodium, Na Potassium, K Strontium, Sr (total)	Barium, Ba (total) Aluminum, Al (total) ** Iron, Fe (total) Manganese, Mn (total) Bicarbonate, HCO3	Chloride, C! Flouride, F Sullate, SO4 Nitrate, NO3 Phosphate, PO4 Hardness (as CaCO3) Alkalfinity (as CaCO3)	Silica, SiO2 Total Organic Carbon (TOC)	Standard Plate Count (SPC) Headloss Backwash (B/W) Frequency

GW#1 - Groundwater treatment process no. 1 ( KMnO4 oxidation; filtration) GW#2 - Groundwater treatment process no. 2 (Nanofiltration)

<sup>..</sup> Only if alum is being used in the process

The manganese-greensand filter was charged initially using a 3-percent KMnO<sub>4</sub> solution injected at a rate of 0.2 to 0.3 gal/min, until a total of 8 gallons had been pumped through the media (manufacturer's recommendation). During operation, the filter was backwashed when the  $\Delta P$  had increased by about 10 lb/in² over its original startup value.

Finally, a series of jar tests were performed to evaluate alum  $[Al_2(SO_4)_3\cdot 14.3H_2O]$  as a coagulant for the removal of suspended particles (oxidized  $Mn^{+2}$ ). Required settling time, supernatant turbidity and floc stability were used as the determining factors for judging the effectiveness of the coagulant.

## 5.4.2 Nanofiltration. - The NF test system design parameters were as follows:

• Array 12:6 (2-stage) (refer to appendix C for diagram)

Element FilmTec NF90-2540

Recovery 80 percent

Initial feed pressure
 Feed flow
 570 kPa (83 lb/in²) @ 25 °C
 18.2 L/min (4.8 gal/min)

• Projected permeate TDS 271 mg/L

The following chemical additions were added for scale control:

• Hypersperse AF 200<sup>TM</sup> @ 3.0 p/m

• Sulfuric acid to pH 7.00

The NF system was operated for nearly 1000 hours to allow time for any potential membrane degradation from fouling or scaling to develop. System startup was at operating pressures required to achieve 80-percent recovery at a feed flow of 18.2 L/min (4.8 gal/min). Thereafter, these same flows were maintained and system pressures were allowed to vary.

Process instrument data were manually recorded four times per day, with about 4 hours between observations. Just before data collection, the operator adjusted the system flows to the following:

• Feed 18.2 L/min (4.80 gal/min)

• Concentrate 3.6 L/min (0.96 gal/min)

These flows were achieved by adjusting the BPV (back pressure valve) on the high pressure pump recycle line and the FCV (flow control valve) on the concentrate line.

An SDI (silt density index) measurement of the cartridge filter effluent stream was made once a day. SDI is a measure of fouling potential of the feed from colloidal-size materials.

Samples of the feed, interstage, permeate, and concentrate (reject) streams were collected after 5, 362, 693 and 984 hours of operation. These samples were sent to a contract laboratory for the following analyses: major anions and cations, selected metals, hardness, alkalinity, and  $SiO_2$  (silica). In addition, feed samples were analyzed for TOC and standard (heterotrophic) plate count (refer to table 2 for the specific constituents).

The 5u cartridge filter elements on the NF skid were changed about every 3 to 4 days.

A PID-controlled LMI chemical feed pump was used to regulate the addition of a 10-percent  $H_2SO_4$  (sulfuric acid) solution for pH adjustment (fig. 4). Hypersperse AF 200 (anti-scalant) was added using a manually-controlled LMI pump. A fresh 5-percent solution was prepared about every 4 days.

#### 6. PILOT TEST RESULTS AND CONCLUSIONS

#### 6.1 Results

This section presents the test results for both the potassium permanganate oxidation and nanofiltration processes. Tabulations of the raw data and calculated parameters are presented in appendices A and B, respectively.

- 6.1.1 KMnO<sub>4</sub> oxidation. The testing described below was designed to determine the following:
- (1) The concentration of KMnO<sub>4</sub> needed to reduce the Mn<sup>+2</sup> level in well S6 ground water to below 0.05 mg/L (EPA secondary drinking water standard)
- (2) The required reaction time and an initial rate constant
- (3) The ability of manganese-greensand filtration to compensate for over- and under-dosing of  $KMnO_4$
- (4) The efficiency of the greensand filter in removing oxidized manganese particles (MnO<sub>2</sub>)
- (5) The effectiveness of alum for coagulating and settling MnO<sub>2</sub> precipitates
- 6.1.1.1 Optimization of KMnO<sub>4</sub> dosage. Figure 6 shows the variation in manganese ion concentration in the raw feed water, and in the greensand influent and effluent, with increasing KMnO<sub>4</sub> dosage. The greensand influent curve appears to verify the optimum dose of 1.10 mg/L KMnO<sub>4</sub> (based on stoichiometry). Permanganate additions above and below this level result in significantly higher residual Mn<sup>+2</sup> levels. Also, the greensand filter media seems to be effective in controlling the over- and under-dosing of KMnO<sub>4</sub>. It is interesting to note, however, that the water apparently picked up Mn<sup>+2</sup> from the greensand in the region of optimum KMnO<sub>4</sub> dosage. From figure 6, the overall process appears to have an effectiveness limitation of about 0.05 mg/L.
- **6.1.1.2** Permanganate reaction kinetics. Reaction kinetic experiments were conducted to optimize detention time prior to greensand filtration. Because of the pilot plant configuration, detention time was fixed (for a given process flow rate) by the combined capacities of the rapid mix tank, flocculation basin, and clearwell. At a process flow rate of 22.7 L/min (6 gal/min), this detention time was 55 minutes. The tests described below were designed to find out if a shorter detention time could be used.

Test water was taken from the detention tank. The optimum  $KMnO_4$  dose of 1.1 mg/L was added to water in a jar test apparatus. After a brief rapid mixing period, stirring was slowed to mimic the stirring in the flocculation basin. At a predetermined reaction time the contents of the jar were vacuum filtered, through a Whatman #40 filter paper, into a flask containing  $NaHSO_3$  to stop the oxidation. The  $Mn^{+2}$  concentration was then measured using the Hach DR/2000. This procedure was repeated for the following reaction times: 1, 1.25, 2, 5, 11, 21, 31, 41, 51, and 60 minutes. Results of this testing are shown on figure 7.

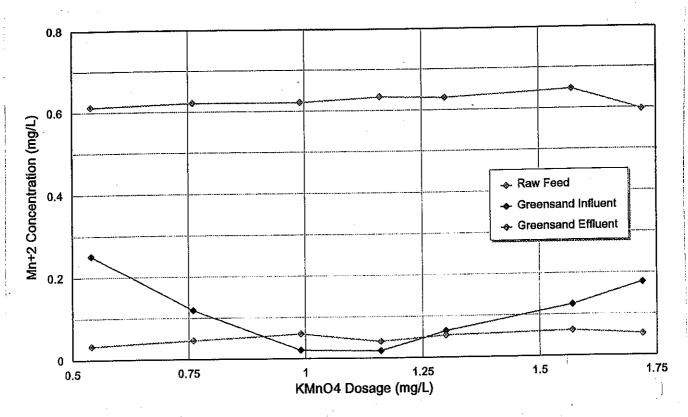


Figure 6. - Optimization of KMnO<sub>4</sub> dosage.

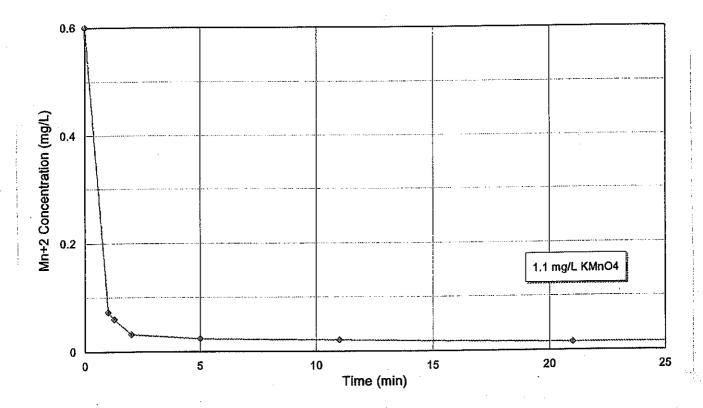


Figure 7. - Timed reaction test.

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About 90 percent of the Mn<sup>+2</sup> had been oxidized by the end of the first 1-1/2 minutes, and the reaction was complete within 20 minutes. By 40 minutes, distinct particles were forming. Shorter reaction times were attempted, but because the filtering process required about 2 minutes, obtaining a precise reaction stop time was difficult. Obviously, the permanganate reaction is not going to require significant detention time. The initial reaction rate was found to be second order dependent on both [MnO<sub>4</sub>] and [Mn<sup>+2</sup>] with a rate constant of -0.198/mol·sec.

6.1.1.3 Filtration efficiency. - Figure 8 shows turbidity data for the 22.7 L/min (55-minute detention) tests. The greensand effluent turbidities were all at or below 0.06 ntu, indicating very efficient filtration. The greensand influent turbidities ranged from 5 to nearly 20 ntu, generally increasing with KMnO<sub>4</sub>, suggesting a greater number of suspended MnO<sub>2</sub> particles.

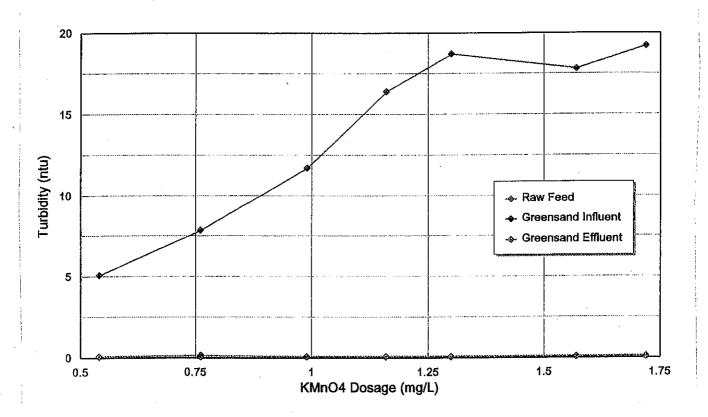


Figure 8. - Turbidity data.

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Although Mn<sup>+2</sup> is oxidized rapidly, it may still require time to form filterable particles. To check this possibility, additional pilot tests were run with a 12-minute detention time and the same optimum KMnO<sub>4</sub> dosage of 1.1 mg/L. This procedure was accomplished by bypassing the flocculation basin, i.e., flowing directly from the rapid mix tank to the clearwell. As expected, greensand filter effluent Mn<sup>+2</sup> concentrations were not significantly different from the earlier 55-minute detention tests (refer to data in appendix A). Interestingly, the filter effluent turbidities were also not that much different than before, ranging from 0.06 to 0.09 ntu.

**6.1.1.4** Alum clarification. Filter alum was evaluated for coagulating and settling  $\rm MnO_2$  precipitates. Jar tests were performed to determine optimum dosage, settling time, and supernatant clarity (in terms of turbidity). The best floc formation was achieved within the range of 20 and 40 mg alum/L. However, the required settling time of 90 minutes was excessive and the floc produced was very fragile. Slight agitation of the samples would tend to break up floc particles.

6.1.2 Nanofiltration. - The testing described below was designed to determine the following:

- (1) The performance of the FilmTec NF-90 membrane element in reducing TDS, hardness, sulfate, and manganese levels in well S6 ground water
- (2) The potential long-term adverse effects on the membranes from fouling or scaling
- (3) The blending ratio (NF permeate with filtered well water) to achieve high overall net recoveries

**6.1.2.1** Operational data. - A total of 984 hours of operation accrued on the NF elements during this test phase. The raw data collected by the plant operators and other calculated values are tabulated in appendix B. Flow, temperature, conductivity, and pressure data are also graphically depicted on figures 9 through 14.

Referring to figure 9, feed and reject flows were held constant at 18.2 L/min (4.8 gal/min) and 3.6 L/min (0.95 gal/min), respectively, yielding an 80-percent recovery of desalted water (permeate). The total amount of permeate recovered is the summation of the following three flows:

- Stage 1, vessel 1 permeate (orange symbols)
- Stage 1, vessel 2 permeate (yellow symbols)
- Stage 2 permeate (blue symbols)

Figure 10 shows diurnal and long-term variation in feed temperature. This measurement was taken at the feed end of the first stage. Temperature has a significant effect on membrane performance and is used in calculations of net permeate flow, which is normalized to 25 °C.

Figure 11 displays system conductivities as µS/cm (microSiemens per centimeter). Also, for better resolution, figure 12 shows an expanded view of the permeate conductivities. Note that the permeate conductivities show a gradual decrease throughout the test period, particularly in the second stage. This decrease may be the result of a "dynamic layer" developing on the membrane surface, either from biofouling or the deposition of colloidal-size particles (clays, silts). The greater effect in the second stage could be caused by the lower average brine velocity (about 16 percent less than the first stage). With reduced scouring action, microorganisms and colloidal particles settle out and attach to the membrane surface with greater ease. These contaminants are also present in greater numbers in the second stage.

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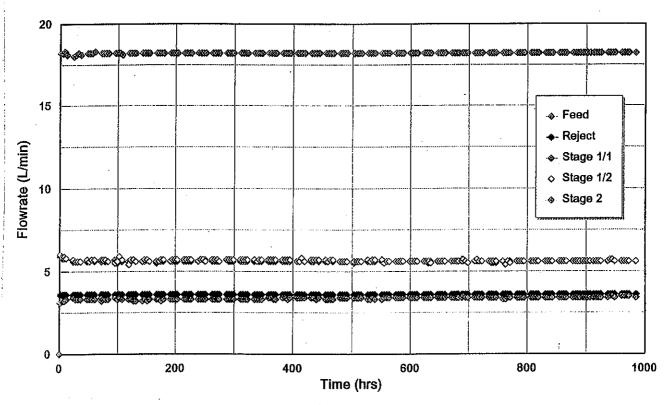


Figure 9. - System flow rates (nanofiltration).

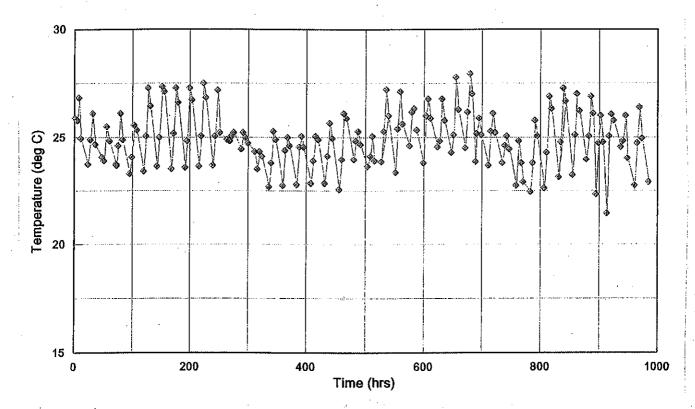


Figure 10. - Feed temperature.

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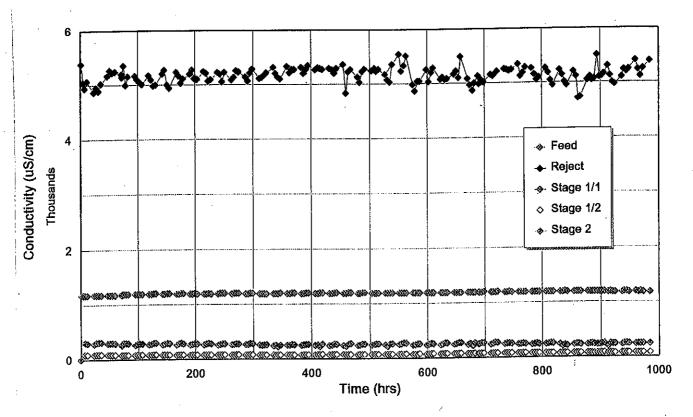


Figure 11. - System conductivities.

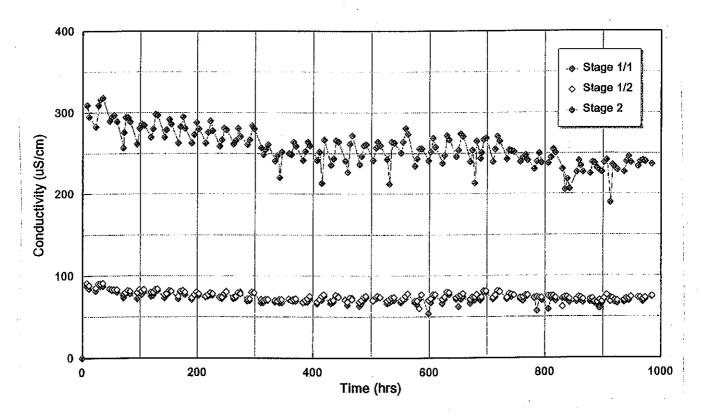


Figure 12. - Permeate conductivities.

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Figure 13 shows the feed, interstage, and reject operating pressures in lb/in². All three pressures are increasing with time, indicating again the possibility of membrane fouling or scaling (less likely). Also, referring to figure 14, the pressure drop across the first stage appears to increase at a slightly faster rate than across the second stage. The downward blip in the three data plots at about 30 hours elapsed time may have resulted from an initial instrument calibration problem (pressure or flow sensors). No other explanation is apparent from a review of the operator data sheets.

Chemical analyses were performed at 5, 362, 693, and 984 hours into the test program on four separate process streams (NF feed, interstage, permeate [combined], and reject), for the following constituents:

- Cations (Ca, Mg, Na, K)
- Anions (HCO<sub>3</sub>, Cl, SO<sub>4</sub>, NO<sub>3</sub>, F)
- Metals (Al, Ba, Sr, Fe, Mn, P)
- Silica

Results of these analyses are shown in appendix D.

Table 3 summarizes %SR (percent salt rejections) for selected ions and for TDS that were calculated from the concentration data in appendix D. The values shown as " $\geq$ " result from the permeate concentration falling below the detection limit. The %SR for other constituents (Al, Ba, Fe and NO<sub>3</sub>), where both the feed and permeate concentrations were below the detection limit, could not be calculated. As shown, the average reduction in TDS, sulfate, and manganese all exceeded 90 percent. Also, based on the Ca<sup>+2</sup> and Mg<sup>+2</sup> data in table 3, the average reduction in hardness exceeded 98 percent.

In addition to the analyses indicated above, TOC and standard (heterotrophic) plate counts were run on detention tank effluent. The plate counts taken at the 5- and 362-hour sampling times were high—5700 and 4000 cfu/mL (colony forming units per milliliter), respectively. Based on these values, it was decided that plate counts should be run at several locations within the process to better define what might be a potential problem. Samples collected on November 1, 1994, resulted in the following data:

•	Wellhead (sample tap)	7900 cfu/mL
•	Detention tank effluent	7200 cfu/mL
•	NF feed water (after cartridge filter)	1100 cfu/mL
•	Interstage	1200 cfu/mL
•	Permeate (second stage)	$790  \mathrm{cfu/mL}$
•	Reject	1900 cfu/mL

Because well S6 is fairly shallow (160 feet) and close to Lake Havasu (within a few hundred meters), the high bacterial populations could result from lake recharge. Deep inland wells generally have much lower plate counts. The city's practice of continually adding food-grade oil (Unocal White Oil) to the well as a lubricant may also be exacerbating the problem. This oil is biodegradable and was found at the surface and along the walls of the detention tank.

At least one SDI measurement was performed on the NF feed water (downstream from the  $5\mu$  cartridge filter) each day of testing. SDI measures fouling potential of the feed from colloidal-size materials. The recommended maximum SDI specified by the manufacturer for the NF-90

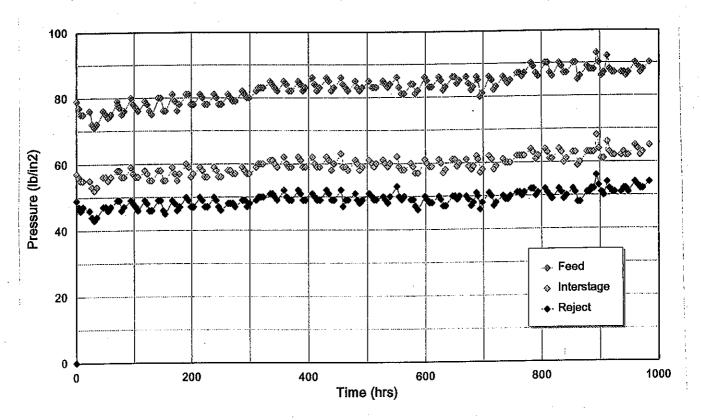


Figure 13. - System pressures.

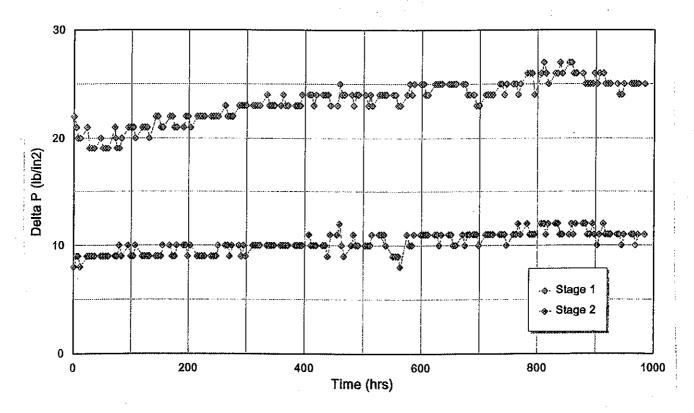


Figure 14. - Stage pressure drops.

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Table 3. - Percent salt rejection data.

Constituent	5-Hour	362-Hour	693-Hour	984-Hour	Average
Calcium	97.8	98.2	98.4	99.0	98.4
Magnesium	97.7	98.8	98.5	98.9	98.5
Sodium	80.0	84.5	81.8	85.8	83.0
Potassium	74.4	68.6	≥68.6	≥63.0	-
Manganese	≥90.0	≥90.0	≥90.7	. ≥90.6	≥90.3
Bicarbonate	83.5	91.8	89.4	91.2	89.0
Chloride	70.2	83.1	80.8	78.9	78.3
Sulfate	96.6	97.5	97.6	97.9	97.4
Fluoride	87.5	84.8	84.9	83.1	85.1
Silica	70.0	82.2	80.2	82.6	78.8
TDS	88.0	92.1	91.1	92.5	90.9

membrane is 5.0. During the 6-week test period, a total of 64 SDIs were performed with values ranging from 1.68 to 4.28. The average SDI was 2.98 with a standard deviation of 0.53.

**6.1.2.2** Performance degradation. - Figures 15 and 16 present the average NDP (net driving pressure) and NPF (normalized permeate flow) for this test phase. Average NDP is the pressure available to force water through the membrane, and is calculated as follows:

$$NDP = P_f - P_p - P_o$$

where:

 $P_f$  = average feed pressure (average of feed and reject pressures)

 $P_p =$  pressure in the permeate line (gauge pressure)

 $P_o$  = average osmotic back pressure of the feed water (estimated by averaging the feed and reject concentrations and dividing by 100)

NPF is the total permeate flow adjusted to standard temperature (25°C) and to normalized NDP at startup, and is calculated as follows:

$$NPF = NDP_{startup}/NDP_{today} \times TCF \times F_n$$

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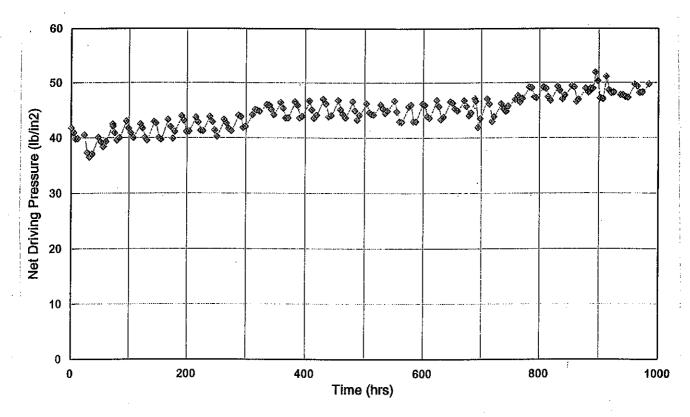


Figure 15. - Average net driving pressure.

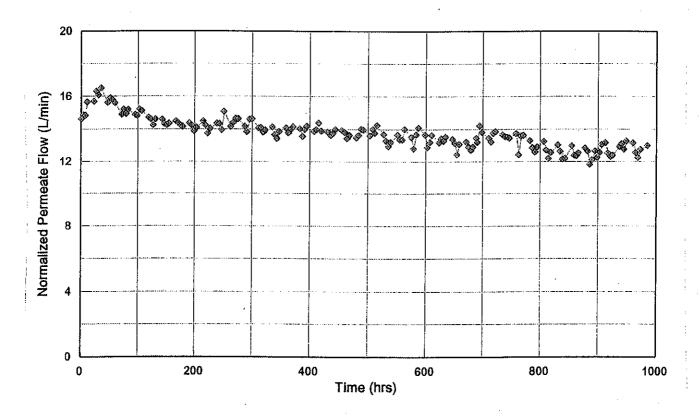


Figure 16. - Normalized permeate flow.

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where:

TCF = temperature correction factor

 $F_{\rm p} = {\rm permeate\ flow}$ 

The NPF graph (fig. 16) can be used to estimate the degree to which membranes are being fouled or if damage is occurring, and is commonly used to determine the time at which membranes should be chemically cleaned. Some drop in NPF with time is expected. For the TFC (thin-film composite) membranes used in this study, a 15- to 20-percent decline over a 3-to 5-year period would not be unusual. The roughly 16-percent drop in NPF experienced in this test program over a 1000-hour (6-week) test period is excessive by comparison.

One of two possible causes were considered for this decline in system performance: (1) the deposition of  $\mathrm{MnO}_2$  (oxidized manganese precipitates) on the membrane surface; and (2) biofouling.

**6.1.2.3** Membrane autopsy and SEM analysis. - On November 11, 1994, autopsies were performed on 2 of the 18 NF elements, one of the lead elements in stage 1 (serial # A2282494; refer to appendix E) and the trailing element in stage 2 (serial # A2282495). Initial observations of the lead element membrane surface revealed no obvious fouling or scaling, perhaps only a slight and fairly uniform discoloration (darkening). Also, the vexar (plastic feed water-brine spacer located between membrane envelopes) showed no signs of any buildup. After measuring the dimensions of the active membrane area of the two leaves, one of the membrane surfaces was irrigated with deionized water and thoroughly squeegeed to collect any adhering deposits. Surprisingly, this operation yielded a significant amount of light brownish-colored material (the total quantity of material collected was later determined to have a dry mass of 0.32 grams; this material was collected from a membrane active surface area of 0.52 m²). The uniformity of deposition, color, consistency, and strength of adherence was determined to be consistent with biofouling. Similar material was found in the trailing element of the second stage, but to a lesser degree (dry mass of 0.09 grams).

Two-inch-square membrane samples were cut from the second leaf of each element for SEM (scanning electron microscopy) imaging to determine if biological cell structure and morphology could be identified. Some of the samples were gold-coated to enhance the imaging resolution and detail. Several showed characteristic rod-shaped bacterial cell forms (refer to fig. 17 as a typical example). No attempt was made to classify specific bacterial types or strains.

The sample collected from the lead element squeegeeing operation (liquid with suspended material) was digested and analyzed for trace metals using ICP (inductively-coupled plasma) spectroscopy. Because the concentration of  $Mn^{+2}$  was high in the feed water, the presence of  $MnO_2$  (manganic dioxide) was suspected. A total of 25 metals were identified. Table 4 lists the metals found with a total mass  $\geq 0.030$  mg. Remember, these metals were collected from a membrane surface area of about 0.50 m<sup>2</sup>.

From these analyses, it appears unlikely that a problem exists with manganese. However, the presence of such a high level of iron is surprising. The concentration of iron (total) in the NF feed water was  $\leq 0.05$  mg/L, significantly less than the general RO/NF process limitation for Fe<sup>+2</sup> of 0.3 mg/L. Two possible explanations were considered:

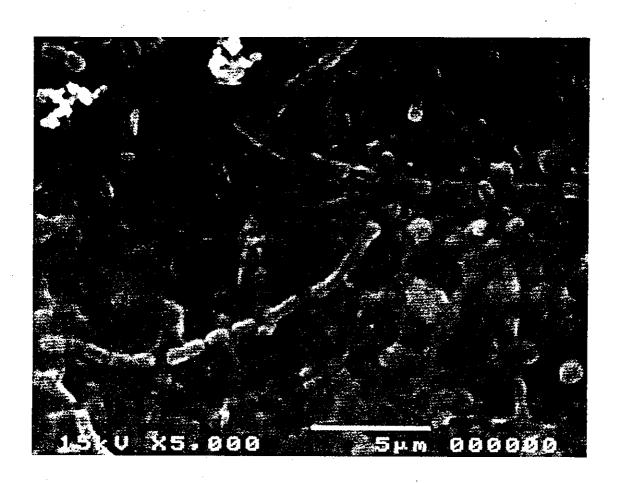


Figure 17. - SEM showing characteristic rod-shaped bacterial cell forms.

	16 . 95
Metal	Mass (mg)
Iron, Fe	8.1
Sodium, Na	2.5
Calcium, Ca	0.89
Chromium, Cr	0.56
Silicon, Si	0.55
Arsenic, As	0.30
Aluminum, Al	0.19
Zinc, Zn	0.15
Magnesium, Mg	0.14
Barium, Ba	0.037
Manganese, Mn	0.030
Lead, Pb	0.030

- Iron-fixing bacteria may have oxidized the available Fe<sup>+2</sup> to Fe<sup>+3</sup> (ferric iron) which deposited on the membrane, or may have concentrated iron within the biomass of the fouling layer (Bess, 1994).
- Some corrosion of system components may have occurred (both Cr [chromium] and Ni [nickel] were found on the membrane surface, which are major components of 304 and 316 stainless steel); all wetted components of the NF system were specified to contain only 316 stainless.

A final test was performed on the membrane surfaces (both lead and trailing elements) to determine if any physical degradation had occurred during operation. Congo red dye was applied to a small area of the membrane's surface and then wiped away after a few moments. By doing this, surface penetrations (pin holes, cracks, etc.) can be readily identified by a residual dye stain. No physical degradation was observed.

#### 6.2 Conclusions

# 6.2.1 KMnO<sub>4</sub> oxidation. - The following conclusions were reached based on this test phase:

• The combination of KMnO<sub>4</sub> oxidation and manganese-greensand filtration effectively reduced Mn<sup>+2</sup> in the well S6 ground water to an average concentration of 0.05 mg/L (fig. 6), which is the secondary MCL.

- The optimum KMnO<sub>4</sub> dose was found to be about 1.1 mg/L (fig. 6). Considering the amount of Mn<sup>+2</sup> removed (0.62 mg/L [average influent concentration] 0.05 mg/L [average effluent concentration]), this dose is equivalent to 1.93 mg/L KMnO<sub>4</sub> per mg/L Mn<sup>+2</sup>, which is essentially the same as the stoichiometric requirement (section 5.2.1).
- Greensand filtration was effective in controlling the over- and under-dosing of KMnO<sub>4</sub> (fig. 6).
- KMnO<sub>4</sub> oxidation could be used with conventional dual- or multi-media filtration, i.e., without greensand, if an effective control could be employed for chemical dosing. From figure 6, lower Mn<sup>+2</sup> concentrations appear to be achievable without greensand (near the optimum KMnO<sub>4</sub> dose of 1.1 mg/L).
- The reaction is extremely fast (fig. 7). About 90 percent of the Mn<sup>+2</sup> is oxidized within the first 1-1/2 minutes. The initial reaction rate was found to be second order dependent on both [MnO<sub>4</sub>] and [Mn<sup>+2</sup>] with a rate constant of -0.198/mol·sec.
- The greensand effluent turbidities were all ≤ 0.06 ntu for the 55-minute data and ≤ 0.09 ntu for the 12-minute data (appendix A), indicating efficient filtration.
- In jar testing for coagulating and settling MnO<sub>2</sub> precipitates, 20 to 40 mg/L alum was found to produce the best floc formation. However, the required settling time of 90 minutes was excessive and the floc produced was very fragile. It is therefore concluded that alum is not effective for this application.

## **6.2.2 Nanofiltration**. - The following conclusions were reached based on this test phase:

• NF effectively reduced the concentrations of all contaminants of concern (see section 3.0) to below MCLs. The average TDS rejection for the 984-hour test was 90.9 percent. Specific ions of interest were removed as follows:

Constituent	% Rejection	Average Conc. Permeate (mg/L)
Ca <sup>+2</sup> (hardness)	98.1	1.6
Mg <sup>+2</sup> (hardness)	98.5	0.5
SO <sub>4</sub> -2 Mn <sup>+2</sup>	97.4	7.7
Mn <sup>+2</sup>	≥90.3	< 0.05

- The average TDS (summation of ions [appendix D]) for the feed and permeate were 826 and 75 mg/L, respectively. By blending the NF permeate and filtered well water at an approximate ratio of 1:1.25, a net overall recovery of 90 percent (at 496 mg/L TDS) could be achieved. However, Mn<sup>+2</sup> would have be to removed from the well water, prior to blending, to meet its secondary MCL of 0.05 mg/L.
- NPF dropped by about 16 percent during the 6 weeks of testing (fig. 15). During this same period, system feed pressure increased from about 75 to 89 lb/in² (fig. 13). The membrane autopsy, SEM analysis, and high heterotrophic plate counts (measured throughout the NF system) all point to biofouling as the cause.

- UV (ultraviolet) disinfection was not effective in controlling microbial contamination (biofouling) of the NF system. Chloramines or chlorination-dechlorination would provide more effective control; however, Mn<sup>+2</sup> oxidized by these chemicals would have to be removed by media filtration prior to desalting. Also, because of the high microbial populations in the feed water, residual cell components of microorganisms killed during disinfection may have to be removed as well. These dead organisms can provide a food source for other live bacteria, and some evidence suggests that they may also deposit on and adhere to the membrane contributing directly to a biofouling layer (Ridgeway, 1984).
- The use of a food-grade oil (Unocal White Oil) as a well pump lubricant may have contributed to the biofouling. A plate count taken at the wellhead sample tap indicated 7900 cfu/mL (section 6.1.2.1). In addition, residues of the oil were found at the surface and along the walls of the detention tank.
- Very little manganese was found on the membrane surface of the autopsied first stage lead
  element (refer to section 6.1.2.3). However, a considerable quantity of iron was present.
  Because the feed-water concentration of iron was low (0.05 mg/L), it is suspected that ironfixing bacteria may have oxidized or concentrated Fe<sup>+2</sup> within the biomass of the fouling
  layer.

#### 7. FULL SCALE TREATMENT

#### 7.1 General

Lake Havasu City has several choices and decisions to make regarding the construction of a full-scale water treatment plant. Among these are plant location, level of treatment, and the amount of surface water versus ground water used to meet water demands. Full-scale treatment for Lake Havasu City is estimated to be accomplished over a phased expansion. The city and their consultant, HDR Engineering Inc., have concluded that water demands for the city are projected according to table 5 below.

Table 5. - City average day water demands.

Year	Water Demand (acre-ft\yr)	Water Demand (Mgal/d)
1994-95	14,562	12.99
2004-05	19,180	17.12
2015	24,454	21.8

City officials have stated a size preference for a full-scale treatment plant, initially at 12 Mgal/d (million gallons per day), with a projected expansion of 12 Mgal/d, for a total plant capacity of 24 Mgal/d. This report addresses several treatment plant options available to the city at 12 Mgal/d capacity and, in section 8, presents construction cost estimates that the city can consider in final design.

It is important to note that the current and recent past water quality levels of LHC drinking water meet all EPA primary drinking water standards, which are designed to protect the public health (if the current action level for lead of 0.015 mg/L had been in place in 1991, then southern wells S4, S6, and S17 would have exceeded the standard).

Secondary drinking water standards are in effect to protect the public welfare by providing guidelines regarding the taste, odor, color, and other aesthetic aspects of the water. A review of the city's ground-water quality from wells in the north, central, and southern well fields indicates that several secondary standards have been exceeded in recent years and currently still exceed the manganese, TDS, and sulfate MCLs. HDR Engineering, Inc., addressed these secondary water quality levels in their Comprehensive Water Master Plan (HDR Engineering, 1992) with a discussion of treatment options for each constituent. Based on this information, Reclamation was asked to perform field pilot testing at well S6, which has historically contained high levels of manganese, TDS, and sulfates.

Full-scale treatment alternatives presented in this report fall into two levels of treatment. A significant cost increase occurs from one level to the next. These levels can be described as:

- Manganese removal using the piloted process of permanganate oxidation followed by filtration. This alternative will remove manganese but will not affect total dissolved solids.
- A higher level of treatment which would provide, or nearly provide, full compliance with all secondary standards of the Safe Drinking Water Act. These processes of NF and lime softening are included for complete and partial compliance, respectively.

## 7.2 Manganese Removal (KMnO<sub>4</sub> Oxidation)

The results of pilot testing on ground water indicate that manganese removal can be accomplished by a water treatment plant employing potassium permanganate (KMnO<sub>4</sub>) oxidation followed by filtration. Full-scale ground-water treatment would consist of the following unit processes, as shown in the process flow diagram on figure 18: raw water pumping, permanganate feed system, 2-stage mixing, filtration, controls for monitoring and compensating for over- or under-dosing of KMnO<sub>4</sub>, clearwell, post-disinfection, and booster pumping.

Formation of the precipitate MnO<sub>2</sub> (manganic dioxide), from the oxidation of Mn<sup>+2</sup> (manganese), requires a rapid mix step followed by a slow mix, or flocculation step. The rapid mix step can be accomplished by an agitator-mixer designed to impart adequate energy for the size and shape of the rapid mix tank. A detention time of 1 minute is recommended for rapid mixing. From the piloting work, a detention time for flocculation of about 20 minutes is recommended.

The optimum amount of KMnO<sub>4</sub> required for the oxidation process was determined to be stoichiometric, based on the concentrations of soluble manganese and iron present (refer to section 6.1.1.1). Two control options could be considered for KMnO<sub>4</sub> addition: the use of greensand filtration for the compensation of over- and under-dosing; and the use of an effluent monitor and feedback loop to control the chemical feed pump. Test results indicated that although greensand filtration effectively controlled over- and under-dosing, the use of greensand apparently caused a slight increase in the soluble manganese level of the filter effluent at KMnO<sub>4</sub> doses that were determined to be near optimum (refer to section 6.1.1.1 and fig. 6).

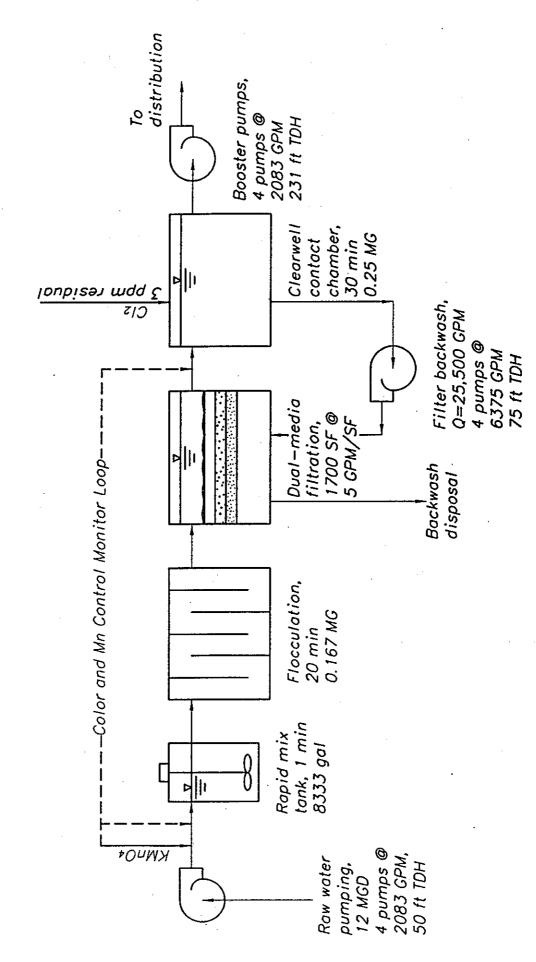


Figure 18. - Water treatment with potassium permanganate oxidation—Lake Havasu City water treatment study.

Because of this increase, and the higher cost of greensand filtration compared to conventional filtration, a multi-media (sand and anthracite) filter is recommended instead of greensand. This recommendation assumes that an effective  $KMnO_4$  feed control system is commercially available for incorporation in the process design.

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Following filtration, a post-disinfection step, consisting of chlorine addition, is recommended. The design includes a 30-minute chlorine contact basin, which also serves as a wetwell for the finished water booster pumping to the city's distribution system. Chlorine was selected because the final trihalomethane formation potential is low. Should this THMFP increase or more stringent limits be imposed for THMs, the city may have to use an alternate disinfectant, such as chloramine or ozone.

## 7.3 Full Compliance

7.3.1 General. - To achieve full compliance with secondary drinking water standards, contaminants which have repeatedly exceeded established MCLs in the city's ground water (manganese, sulfate, and TDS), must be reduced in concentration. Only a limited number of treatment methods can accomplish this reduction and, as shown in this report, the cost of treatment increases proportionately with levels of removal.

The nanofiltration option includes blending with partially treated ground water so that the volume of water treated by NF, and thus overall treatment costs, are reduced. The lime softening option does not remove all secondary drinking water contaminants to below the MCL, and therefore does not produce a product water that is in full compliance with all secondary MCLs. However, the lime softening option does have the potential to be blended with other higher quality waters, such as NF-treated ground water or surface water, to achieve full compliance.

**7.3.2** Nanofiltration. - As discussed earlier in section 5.2.2, NF is a separation process that desalts water by the application of hydrostatic pressure to drive feed water through a semi-permeable membrane. A major portion of the water's impurity (dissolved salts) remains behind and is discharged as waste brine; relatively pure water emerges at near atmospheric pressure. Operating pressures for NF are in the range of 75 to 150 lb/in². Typical ion rejections are 90 to 95 percent for divalent ions ( $Ca^{+2}$ ,  $Mg^{+2}$ ,  $SO_4^{-2}$ ), and 60 to 70 percent for monovalent ions ( $Na^+$ ,  $K^+$ ,  $Cl^-$ ,  $HCO_3^-$ ).

A proposed flow scheme for LHC using NF is presented on figure 19. As shown, the raw feed water is split, after pumping, with half of the flow directed to NF treatment, and the remaining half to KMnO<sub>4</sub> oxidation treatment for the removal of manganese. Because the NF product is of such high quality (far exceeds drinking water standards), the two product streams can be blended and still meet secondary MCLs. Also, by doing this blending, the overall treatment cost is lower than if the entire flow stream were to receive NF treatment.

Pretreatment is critical to protect membranes from scale deposits and colloidal and biological fouling. For LHC ground water, the following NF pretreatment is recommended: disinfection to destroy microorganisms; polymer addition and filtration to remove suspended colloidal materials; and the addition of an anti-scalant and acid to prevent scaling of the membranes.

Table 6 presents a summary of the anticipated ion concentrations in the proposed NF plant for the following flow streams: feed, reject brine, permeate (product), and blended (or final) product.

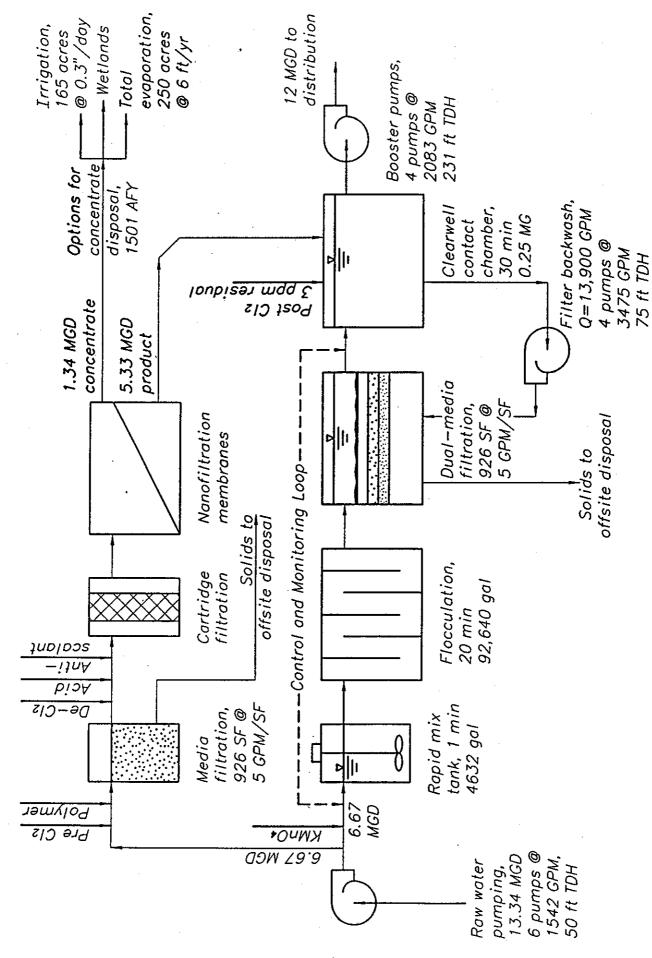


Figure 19. - Water treatment with nanofiltration—Lake Havasu City water treatment study.

Table 6. - Nanofiltration ion concentrations.

Constituent	Feed (mg/L)	NF Brine (mg/L)	NF Permeate (mg/L)	Blended Permeate (mg/L)
Calcium	84.4	400	1.39	48.1
Magnesium	29.5	120	0.46	16.7
Sodium	113	500	19	71.6
Potassium	3.33	11	≤1.0	<b>≤2.3</b>
Barium	< 0.05	0.16	< 0.05	
Strontium	1.35	7.4	<1.0	
Iron	< 0.05	< 0.05	< 0.05	
Manganese	<0.05	2.3	<0.05	
Bicarbonate	173	780	19	105
Chloride	108	430	23.3	70.7
Sulfate	293	1600	7.7	167
Nitrate	<0.80	0.7	< 0.50	
Fluoride	0.93	3	0.13	0.58
Silica (total)	18.3	77	3.88	12
Alkalinity	173	780	19	105
Hardness	333	1500	5.33	189
Total dissolved solids (TDS)	826	3932	75	496

**7.3.2.1** Brine production. - Reject brine, or concentrate, is the waste stream resulting from the NF desalting process. It contains most of the impurities (dissolved salts) originally present in the feed water. The estimated ionic makeup of this reject stream for the proposed LHC plant is shown in the second column of table 6 (NF Brine). As shown on figure 19, about 1.34 Mgal/d of reject brine will be produced from the nanofiltration process.

**7.3.2.2** Brine disposal. - Generally, brine disposal options include the following: surface water discharge, deep well injection, evaporation, spray irrigation, and constructed wetlands. However, for LHC, as with most other municipalities, brine disposal choices are restricted by State and Federal regulatory requirements.

By far, the least costly brine disposal option is discharge to a nearby surface water. Surface water discharges are regulated by the Clean Water Act and, as such, would be subject to permit restrictions. Arizona's permit program under the National Pollutant Discharge Elimination System program would probably not allow a high saline discharge to the Colorado River. Such cost-prohibitive treatment requirements make this option infeasible for the city.

Deep-well injection is possible in Arizona; however, an Aquifer Protection Permit is required with discharge limits which also are likely to make this option cost prohibitive. It would have to be shown that injection of the brine would not adversely impact sub-surface aquifers. This demonstration can be done with geohydraulic modeling of the aquifer, once the aquifer characteristics are known and understood.

The three remaining options for brine disposal are evaporation, spray irrigation, and the creation of a constructed wetlands. The final selection of the type of disposal depends on many factors. Combinations of these options are also possible and may satisfy several goals of the city.

- 1. Evaporation: An evaporative pond system of about 250 surface acres would evaporate the 1.34 Mgal/d of brine, based on an estimated net evaporation rate of 6 ft/yr. This land area can be separated into several ponds to suit the desired goals and objectives of the disposal option. The liner for the ponds would be PVC, HDPE, or compacted clay if locally available. A force main system from the plant to the pond is assumed, which includes a storage tank, sized at 5 days production (6.7 Mgal), and a pump station operating at 4653 gal/min for 24 hours every 5th day. This disposal option was assumed for preparation of the cost estimate (section 8), primarily because of its popularity.
- 2. Irrigation: In a desert environment, a high priority must be placed on both water conservation and the reuse of waste waters to lessen water demands where possible. The city's "Comprehensive Master Water Plan" devotes an entire chapter to the scenario of developing areas around the city which can use reclaimed water for the irrigation of landscaped areas, thereby lowering water demands. Specifically identified in this Master Water Plan are nine potential reclaimed water sites that could use an average monthly rate of 1.3 Mgal/d and a peak monthly rate of 2.9 Mgal/d of reclaimed water for irrigating landscaped areas. The report concludes that because the city's three reclaimed water plants produce only 1.4 Mgal/d, any additional reclaimed water would lower the city's potable water demand. Therefore, disposal of NF brine by landscape irrigation is an attractive option serving a dual purpose, reducing water demands while increasing the city's landscaping. The ADEQ (Arizona Department of Environmental Quality) must approve such a plan.

Using appropriate salt-tolerant grasses and foliage, irrigation of landscaped areas can be applied at a rate of about 0.3 in/d and require 160 irrigable acres. Such areas can include open space, green belts, golf courses, highway medians, and resort complexes. An irrigation system would require a storage tank or lined holding pond sized for at least 2 days of storage (6.7 Mgal), a pump station that would operate at 1860 gal/min for 12 h/d, and a force main-distribution network.

3. Wetlands: The creation of a wetlands in a desert environment is aesthetically pleasing in that it creates an environment where selected brine-tolerant plants can proliferate. At a pilot saline wetlands in Hemet, California, alkali bulrush, cattails, arrow grass and spikerush plants have survived and flourished in the reject brine from a reverse osmosis demonstration plant (Boegli and Thullen, 1994). In addition, the wetlands will attract waterfowl and animals, i.e., ducks, geese, frogs, and insects. The ADEQ must approve such a brine disposal plan, and there would likely be a permitted discharge from a wetlands system to surface water or to the ground-water aquifer. Alternately, the discharge brine would be further concentrated in an evaporation pond.

For the 12 Mgal/d water treatment facility being considered in this report, a wetlands area of 8.2 acres is required to dispose of 1.34 Mgal/d of brine produced. This area is based on an application rate of 6 in/d, which is the application rate in use at Hemet, California, a site similar in climate to LHC. The other main features of a wetland brine disposal system include a storage tank or lined holding pond sized at about 5 days of flow (6.7 Mgal), a pump station that can operate at 4653 gal/min for 24 hours every 5th day, and a force main.

**7.3.3 Lime softening.** - Lime softening is a widely used process for clarification, reduction of hardness and salinity, and heavy metals removal in natural waters. The process requires the presence of a significant concentration of  $HCO_3$  (bicarbonate ion) in the raw water.

Ca(OH)<sub>2</sub> (lime) is added to the water at ambient temperature. The following reactions occur:

$$Ca(OH)_2 + Ca^{+2} + 2HCO_3 - 2CaCO_3 + 2H_2O$$
 (1)

$$Ca(OH)_2 + Mg^{+2} \rightarrow Mg(OH)_2 \downarrow + Ca^{+2}$$
 (2)

In water containing a reasonable amount of bicarbonate, a voluminous precipitate, consisting primarily of calcium bicarbonate, is formed. This precipitate entraps suspended particles of silica and other materials which cause turbidity in the raw water. To enhance flocculation, a small amount of coagulant, say 10 p/m of ferric chloride, is added.

In waters containing heavy metals like iron and manganese, these metals are precipitated as hydroxides or oxides, which are caught up in the precipitate. Reaction zone pH and residence time can be adjusted to ensure satisfactory removal of manganese.

A lime softening ground-water treatment plant for LHC would consist of rapid mixing, flocculation, settling, and filtration. Figure 20 illustrates these unit operations along with the raw and finished water pumping, a clearwell, and post-disinfection using chlorine.

Rapid mixing, flocculation, and settling occur in one unit, the solids contact reactor. These steps may be separated for greater system flexibility, but at a higher construction cost. A slurry of

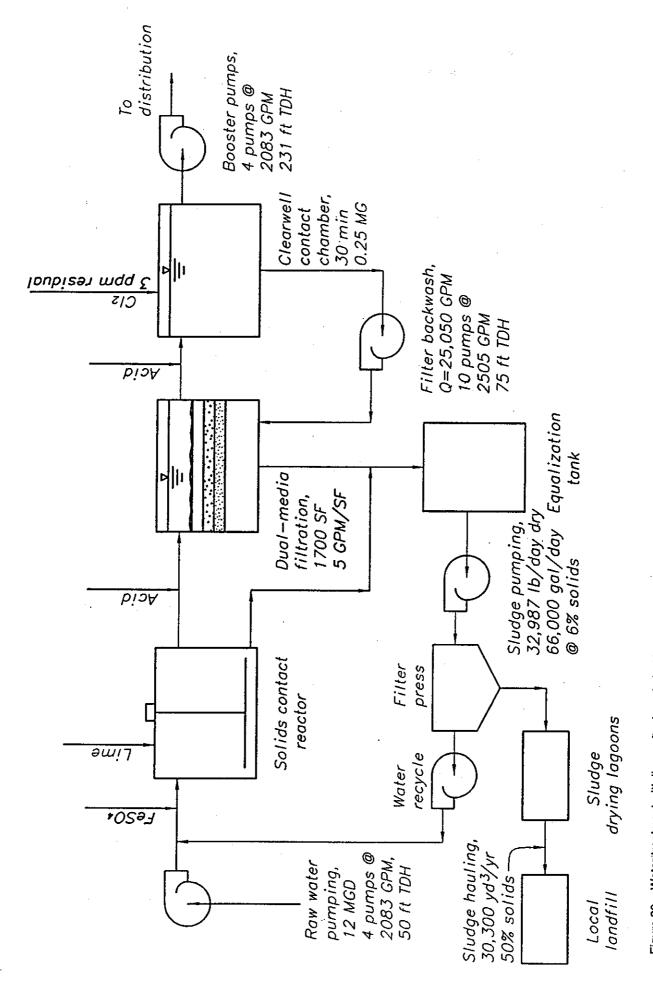


Figure 20. - Water treatment with lime softening—Lake Havasu City water treatment study.

lime in water is added to raw feed water in a high intensity mixing vessel or zone in which reactions (1) and (2) are carried out. Flocculation occurs under gentle agitation, and settling occurs in a zone with very low flow rates. The objective is to create large agglomerates of precipitate which will settle rapidly. Most of the solids are removed as a slurry from the bottom of the reactor. The clarified water is filtered, usually with a rapid sand and anthracite filter, to remove the balance of the solids and produce a clear, treated effluent. The filter is periodically backwashed with treated water to remove the solids. Backwash water is typically returned to the reactor.

The chemistry of this process is well understood. Sludge quantity and composition have been estimated, based on a process feed flow similar to well S6, and are included in appendix F.

**7.3.3.1** Sludge production. - Sludge from lime softening results from the initial raw water solids, plus the precipitate formed from the introduction of lime. At 12 Mgal/d, this volume is estimated to be 66,000 gal/d of a slurry that contains about 6 percent solids by weight. If this sludge was dried, it would weigh about 33,000 lb/d. These solids collect in the bottom of the solids contact reactor and are periodically pumped to a sludge holding tank and filter press for thickening.

**7.3.3.2** Sludge disposal. - It is recommended that the drying of thickened sludge at LHC be accomplished in a lined sludge drying bed. The final quantity of solids that would be hauled to a local landfill would be about 30,300 yd³/yr, at a solids content of 50 percent. For cost estimation purposes, the hauling distance is estimated to be 5 miles.

#### 7.4 Potential Alternatives Not Pilot-tested

Each summer, LHC experiences nationwide high temperatures and it struggles to provide enough water to its residents. To compound this problem, existing wells are declining in productivity as overall water levels in the vicinity recede. The quality of water found in Lake Havasu exceeds the quality of water in the wells. Both manganese and TDS in Lake Havasu are at levels that would not warrant the expense of their removal. Thus, a lower capital cost for treatment of surface water would be required than for ground water. To solve the city's imminent water shortage problem, a surface water treatment plant, sized to satisfy the city's critical summertime shortage, is recommended. The finished water quality data generated from such a plant would benefit the city in design of a larger ground-water treatment plant, both in terms of a lower capacity and for blending with ground water. By using the higher quality and less costly surface water, residents would pay less for their water because the volume of treated ground water would be lower.

Other ground-water treatment processes could not be tested during this pilot test, but they represent additional alternatives to the city to meet future water demands.

### 8. TREATMENT COSTS

#### 8.1 General

Construction, annual O&M (operations and maintenance), and life cycle cost estimates for a 12-Mgal/d plant capacity are provided for the following three levels of ground-water treatment:

- Manganese removal using potassium permanganate oxidation.
- Full compliance with both primary and secondary drinking water standards using nanofiltration.
- Complete removal of manganese and partial removal of sulfates and TDS using lime softening.

Capital cost estimates are based on a combination of direct quotes from manufacturers, plus allowances for installation, and a newly-developed Reclamation software program that uses cost curves prepared by the EPA (Environmental Protection Agency). This program uses the raw water quality from the site and current indices of the ENR (Engineering News Record), along with the Producer Price Index, to calculate both construction and O&M cost estimates.

Capital construction costs are for individual treatment units, including all equipment, but do not include costs for land ownership, rights of way, special site-work, easements, or yard and offsite piping. Also not included are costs for an intake structure, grit removal equipment, or buildings (chemical feed, storage, administration, or laboratory). Legal, administrative and engineering costs for permitting, water quality monitoring, testing, and modeling are not included, nor are general contractor overhead and profit, fees for engineering, legal and fiscal services, and interest during construction. For these reasons, the cost estimates found herein are valuable for a comparison of the alternatives presented, and are not final construction estimates.

The basis for the cost estimates is the EPA's Research and Development manual numbered EPA-600/2-79-162a, titled "Estimating Water Treatment Costs." Each unit process is defined in terms of the following eight subcategories: excavation and site work, concrete, steel, labor, pipe and valves, electrical equipment and instrumentation, and housing. These subcategories are linked to various cost indices and, for this report, have been updated to December 1994 or the Engineering News Record construction cost index for January 1995. Each unit's estimate also includes the cost of a standby or spare unit plus a 15-percent allowance for miscellaneous and contingency items. For O&M costs, the EPA curves are also used with updates for electrical energy costs, maintenance materials, chemicals, and labor. Citizens Electric provided a value of nearly \$0.06/kWh for plants under 1,000 kW. Chemical costs are estimated from recent contacts with chemical supply companies or from a chemical periodical. Labor has been estimated at \$20.00/h.

## 8.2 KMnO<sub>4</sub> Oxidation

A plant that uses potassium permanganate to oxidize manganese in the well S6 feed water is described in section 7.1, and is shown schematically on figure 18. Because little water is lost in this treatment scheme, the flow rate of 12 Mgal/d is used for all unit processes. This treatment plant will effectively remove manganese down to its MCL of 0.05 mg/L. However, other secondary drinking water parameters such as sulfates and TDS will remain above their maximum contaminant levels.

**8.2.1 Construction cost.** - The total estimated construction cost for a plant which removes manganese using potassium permanganate oxidation is \$3,018,700, as shown in table 7. This cost is equivalent to about \$0.25 per daily gallon.

**8.2.2 Operations and maintenance costs.** - The total estimated annual O&M cost for a plant which removes manganese using potassium permanganate oxidation is \$301,500, as shown in table 7. This cost is equivalent to about \$0.07 per thousand gallons treated.

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Table 7. - Construction and annual operations and maintenance costs (12 Mgal/d).

Option 1: Potassium permanganate oxidation

· .	Construction Cost (\$)	Annual O & M (\$)
Raw Water Pumping <sup>1</sup>	145,200	44,600
Potassium Permanganate Addition <sup>2</sup>	74,200	62,500
Rapid Mix <sup>3</sup>	72,200	30,000
Flocculation <sup>4</sup>	247,000	8,000
Dual Media Filter <sup>5</sup>	1,422,100	83,100
Filter Backwash Pumping <sup>6</sup>	455,000	10,800
Chlorination <sup>7</sup>	74,000	33,400
Clearwell <sup>8</sup>	393,000	8,000
Booster Pumping <sup>9</sup>	136,000	21,100
Total	3,018,700	301,500

<sup>&</sup>lt;sup>1</sup> Six pumps with intake screens, each rated at 2083 gal/min at 50 feet of TDH

## 8.3 Full Compliance using Nanofiltration

If the residents of LHC decide they want drinking water that meets all primary and secondary drinking water standards, then an NF plant is recommended. NF produces water of such high quality that blending with available raw water sources will still produce a combined product meeting all secondary MCLs. Data gathered during pilot testing suggest that a full scale treatment plant producing 12 Mgal/d of full-compliance water could treat 6 Mgal/d for manganese alone, and 6 Mgal/d with NF.

8.3.1 Construction cost. - The water treatment plant discussed above, described in section 7.3.2 and shown schematically on figure 19, will cost about \$16,910,100, as shown in table 8. This cost is equivalent to \$1.41 per daily gallon. This cost estimate includes brine disposal using total evaporative drying beds for half of the brine and spray irrigation of greenbelts for the other 50 percent of the brine flow.

<sup>&</sup>lt;sup>2</sup> Concentration of 1.2 p/m potassium permanganate

<sup>&</sup>lt;sup>3</sup> 1-minute detention time, G = 900/second

<sup>&</sup>lt;sup>4</sup> 20-minute detention time

 $<sup>^{5}</sup>$  1700 ft $^{2}$  filter area, includes dual media and housing

<sup>&</sup>lt;sup>6</sup> Four pumps, each rated at 6375 gal/min at 75 feet of TDH

<sup>&</sup>lt;sup>7</sup> Residual concentration of 3 p/m chlorine

<sup>&</sup>lt;sup>8</sup> 0.25-Mgal capacity

<sup>9</sup> Four pumps, each rated at 2083 gal/min at 231 feet of TDH

Table 8. - Construction and annual operations and maintenance costs (12 Mgal/d).

Option 2: Nanofiltration

	Construction Cost (\$)	Annual O & M (\$)
Raw Water Pumping <sup>1</sup>	217,100	66,700
Chlorination <sup>2,3</sup>	35,200	16,600
Polymer <sup>2,4</sup>	46,200	87,600
Dual Media Filter <sup>2,5</sup>	949,200	47,900
Filter Backwash Pumping <sup>2,14</sup>	269,300	8,300
Dechlorination <sup>2,6</sup>	46,200	87,600
Acid Addition <sup>2,7</sup>	778,100	11,200
Anti-Scalant <sup>2,8</sup>	87,900	285,900
Nanofiltration <sup>2</sup>	3,581,400	622,400
Brine Disposal <sup>2</sup>	8,735,000	436,800
Potassium Permanganate Addition <sup>9,10</sup>	46,200	61,400
Rapid Mix <sup>9,11</sup>	40,000	12,200
Flocculation <sup>9,12</sup>	186,000	5,600
Dual Media Filter <sup>9,13</sup>	949,200	47,900
Filter Backwash Pumping <sup>9,14</sup>	269,300	8,300
Chlorination <sup>3</sup>	78,500	35,700
Clearwell <sup>15</sup>	393,000	8,000
Booster Pumping <sup>16</sup>	202,300	31,500
Total	16,910,100	1,881,600

<sup>13.4</sup> Mgal/d, 6 pumps with intake screens, each rated at 1542 gal/min at 50 feet of TDH

<sup>&</sup>lt;sup>2</sup> Flow stream for nanofiltration process, 6.7 Mgal/d

<sup>&</sup>lt;sup>3</sup> Concentration of 1 p/m chlorine

<sup>&</sup>lt;sup>4</sup> Concentration of 2 p/m polymer

<sup>&</sup>lt;sup>5</sup> 930 ft<sup>2</sup> filter area

<sup>&</sup>lt;sup>6</sup> Concentration of 2 p/m sodium bisulfite

<sup>&</sup>lt;sup>7</sup> Concentration of 25 p/m of 93% sulfuric acid

<sup>&</sup>lt;sup>8</sup> Concentration of 7 p/m anti-scalant

<sup>&</sup>lt;sup>9</sup> Flow stream for potassium permanganate oxidation process, 6.7 Mgal/d

<sup>&</sup>lt;sup>10</sup> Concentration of 1.2 p/m potassium permanganate

<sup>&</sup>lt;sup>11</sup> 1-minute detention time

<sup>12 20-</sup>minute detention time

 $<sup>^{13}</sup>$  930 ft $^{2}$  filter area

<sup>&</sup>lt;sup>14</sup> Four pumps, each rated at 3475 gal/min at 75 feet of TDH

<sup>&</sup>lt;sup>15</sup> 0.25-Mgal capacity

<sup>&</sup>lt;sup>16</sup> Four pumps, each rated at 2083 gal/min at 231 feet of TDH

<sup>&</sup>lt;sup>17</sup> Assumes half of brine is spray irrigated and half is evaporated

**8.3.2 Operations and maintenance costs.** The total estimated annual O&M cost for a 12-Mgal/d ground-water treatment plant using NF is \$1,881,600, as shown in table 8. This cost includes a membrane cleaning apparatus, and the cost of membrane replacement every 3 years. This cost is equivalent to \$0.43 per thousand gallons treated.

## 8.4 Lime Softening

As described in section 7.3.3, lime softening of ground water will remove the manganese to levels below the MCL (as long as influent values do not differ much from well S6). Lime softening will not reduce TDS or sulfates to secondary drinking water standards. However, these standards could be achieved by blending with water of higher quality, such as surface or ground water that has been treated with NF.

**8.4.1 Construction cost.** - The 12-Mgal/d lime softening ground-water treatment plant, which is shown schematically on figure 20, will cost about \$7,818,400, as shown in table 9. This cost is equivalent to about \$0.65 per daily gallon.

Table 9. - Construction and annual operations and maintenance costs (12 Mgal/d).

Option 3: Lime Softening

	Construction Cost (\$)	Annual O & M (\$)
Raw Water Pumping <sup>1</sup>	145,200	44,600
Ferric Sulfate Addition <sup>2</sup>	488,900	723,100
Lime Feed Addition <sup>3</sup>	266,500	290,000
Solids Contact Reactor <sup>4</sup>	1,549,700	66,200
Dual Media Filter <sup>5</sup>	1,422,100	83,100
Acid Addition <sup>6</sup>	28,000	5,900
Filter Backwash Pumping <sup>7</sup>	455,000	10,800
Chlorination	74,000	33,400
Clearwell <sup>8</sup>	393,000	8,000
Sludge Disposal <sup>9</sup>	2,860,000	462,500
Booster Pumping <sup>10</sup>	136,000	21,100
Total	7,818,400	1,748,600

- Six pumps with intake screens, each rated at 1542 gal/min at 50 feet of TDH
- <sup>2</sup> Concentration of 10 p/m ferric sulfate
- <sup>3</sup> Concentration of 185 p/m of hydrated lime
  - Four pumps, each rated at 2083 gal/min at 231 feet of TDH
- <sup>5</sup> 1700 ft<sup>2</sup> filter area, includes dual media and housing
- 6 Concentration of 2.5 p/m of 93% sulfuric acid
- Ten pumps, each rated at 2505 gal/min at 75 feet of TDH
- 8 0.25-Mgal capacity
- Includes sludge pumping, storage, filter press thickening, drying beds, and off-site hauling up to 5 miles
- Four pumps, each rated at 2083 gal/min at 231 feet of TDH

**8.4.2 Operations and maintenance costs.** • The total estimated O&M cost for a 12-Mgal/d lime softening plant is \$1,748,600, as shown in table 9. This cost is equivalent to about \$0.40 per thousand gallons treated.

#### 8.5 Cost Analysis

Table 10 presents summarizes the ground-water treatment options presented in this study. It is important to note that these options are not equivalent in terms of the level of treatment each provides and that both construction and O&M costs increase with the increasing levels of treatment. These costs also do not represent final capital costs as explained in section 8.1. It is also important to note that final costs depend on the type of residual disposal option selected. For NF, the combined use of single-lined evaporation ponds and spray irrigation is assumed.

Table 10. - Cost summary of ground-water treatment options (12 Mgal/d).

	Option 1	Option 2	Option 3
Construction Cost (\$)	3,018,700	16,910,100	7,818,400
Annual O&M (\$)	301,500	1,881,600	1,748,600
Construction Cost (\$)/Daily Gallon	0.25	1.41	0.65
O&M Cost (\$)/1000 Gallons	0.07	0.43	0.40

A 20-year life cycle cost analysis is presented in table 11. The analysis is presented in terms of total present worth and total annual cost. A final cost/1000 gal of treated water is also shown.

## 8.6 Surface Water Treatment Costs

Assuming that lime or alum chemical treatment will remove suspended solids and the city can obtain all necessary water rights, a 12-Mgal/d water treatment plant for surface water would cost about \$9,300,000 or \$0.78/daily gallon to construct (HDR Master Plan, table 9-2, and figure 9-6). It would also cost about \$4,190,000 to operate per year or \$0.96/1000 gallons (interpolated from HDR Master Plan table A3-4).

#### 9. CONCLUSIONS

This report concludes the following:

- 1. Water treatment costs to remove contaminants of concern to the residents of LHC vary proportionately with the level of treatment provided. For a 12-Mgal/d (product) treatment plant, these costs range from \$0.25 to \$1.41 per daily gallon for construction, and from \$0.07 to \$0.43 per 1000 gallons treated for annual O&M, as detailed below:
  - a. For a 12-Mgal/d water treatment plant employing potassium permanganate oxidation and the unit operations displayed on figure 18, the construction cost estimate is \$3,018,700 and the annual O&M costs are \$301,500.
  - b. For a 12-Mgal/d water treatment plant employing nanofiltration and the unit operations displayed on figure 19, the construction cost estimate is \$16,910,100 and the annual O&M costs are \$1,881,600.

Table 11. - Life cycle costs for ground-water treatment options.

		Basic Assumptions	3
Study Period		20 years	
Annual Interest Rate		6.5%	
Capital Recovery Factor	•	0.0908	
Present Worth Factor		11.019	
	Potassium Permanganate	Nanofiltration	Lime Softening
Capital Cost	\$3,018,700	\$16,910,100	\$7,818,600
Present Worth of Annual Operating Cost <sup>1</sup>	\$3,322,200	\$20,733,400	\$19,267,800
Total Present Worth	\$6,340,900	\$37,643,500	\$27,086,400
Annualized Capital Cost <sup>2</sup>	\$274,100	\$1,535,400	\$709,900
Annual Operating Cost	\$301,500	\$1,881,600	\$1,748,600
Total Annual Cost	\$575,600	\$3,417,000	\$2,458,500
Cost/1000 Gal of Product³	\$0.13	\$0.78	\$0.56

Present worth of annual operating cost is annual O & M cost times the present worth factor

<sup>2</sup> Annualized capital cost is capital cost times capital recovery factor

3 Total annual cost/ $(365 \times 12,000)$ 

- c. For a 12-Mgal/d water treatment plant employing lime softening and the unit operations displayed on figure 20, the construction cost estimate is \$7,818,400, and the annual O&M costs are \$1,748,600.
- 2. Based on the assumptions made in this report and the life cycle cost analysis assuming 20 years at an interest rate of 6.5 percent, the total annual cost of a 12-Mgal/d plant for the three alternatives studied are:
  - 1) Potassium permanganate oxidation \$575,600 or \$0.13/1000 gal
  - 2) Nanofiltration \$3,417,000 or \$0.78/1000 gal
  - 3) Lime softening \$2,458,500 or \$0.56/1000 gal.

Refer to section 8.1 for an explanation as to what these life cycle costs include.

3. For a 12-Mgal/d treatment plant employing nanofiltration and the unit operations displayed on figure 19, potential benefits might be realized by using about 1.34 Mgal/d of reject brine in either a reclaimed water capacity that could lower irrigation demands on the city's water system, or to create a wetland environment.

### 10. RECOMMENDATIONS

Based on the conclusions noted above, the following recommendations are made:

- 1. LHC is experiencing a severe problem with manganese in their drinking water. Although manganese is only a secondary drinking water standard, and the quality of the water served to the residents of the city meets all Federal and State primary drinking water standards, manganese-related problems such as discolored water, stained clothing, and clogged waterlines are a continuing concern. It is recommended that the city use the economic and treatment process conclusions contained in this report in their planning for future water treatment expansion, and that consideration be given to achieving full compliance with secondary drinking water standards.
- 2. If LHC considers nanofiltration an affordable water treatment option, it is recommended that meetings be arranged with the State of Arizona to determine specific requirements for the brine disposal options presented in this report.
- 3. From a water shortage standpoint, the city is urged to proceed with the construction of a temporary surface water treatment plant sized to provide enough water to meet the critical summertime demands.

### 11. REFERENCES

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- Glase, William H., "Chemical Oxidation," chapter 12, Water Quality and Treatment A Handbook of Community Water Supplies, 4th ed., American Water Works Association, McGraw-Hill, Inc., New York, NY, 1990.
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# **APPENDIX A**

 $\ensuremath{\mathrm{KMnO_4}}$  oxidation and greens and test data

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KMnO4 Oxidation/Greensand Test Data

GS Effluent	9	90.0	S 6	9 6	9 6	00.0	0.03	90.0	ć	0.03	0.0	0.00
Raw Feed GS Influent GS	י ס	7.84	11 23	16.40	10.40	12.00	00.71	02.80	12 00	12.00	12.00	13.30
Raw Feed	000	0.15	10	2 G	800	0.00	<u>.</u>	21.0	. Q	5 5	0.12	0.10
Avg Temp. (deg C)	24.3	23.8	23.5	23.7	23.7	240	0.40	0.4.0	25.5	24.5	243	24.4
Min (mg/L) Westech	<0.05	<0.05	<0.05	<0.05	\$ C	\$ 00	30.0	60.00	900	90.0	0.07	0.08
Mn (mg/L) Hach	0.033	0.046	0.060	0.040	0.054	0.062	0.052	70.00	0.079	0.088	960 0	0.100
Sample	681.3	<b>GS2-3</b>	653-3	<b>GS4-3</b>	685-3	6.86-3	6.27.3		GS8-3	689-3	GS 10.3	GS11-3
Mn (mg/L) Westech	0.21	0.10	<0.05	<0.05	<0.05	0.10	0.15	2	<0.05	40.05	<0.05	<0.05
Mn (mg/L.) Hach	0.252	0.119	0.021	0.017	0.064	0.125	0.177	;	0.011	0.017	0.017	0.012
Sample	<b>GS1-2</b>	GS2-2	GS3-2	<b>GS4-2</b>	<b>GS5-2</b>	<b>GS6-2</b>	GS7-2	1	GS8-2	<b>GS9-2</b>	GS10-2	GS11-2
Mn (mg/L) Westech	0.57	0.58	0.57	0.52	0.53	0.57	0.55		0.56	0.59	0.59	0.58
Mn (mg/L) Hach	0.613	0.621	0.621	0.633	0.630	0.647	0.597		0.621	0.614	0.618	0.626
Sample	GS1-1	GS2-1	<b>GS3-1</b>	GS4-1	GS5-1	GS6-1	GS7-1		GS8-1	<b>GS9-1</b>	GS10-1	GS11-1
Res. Time (min)	55	55	55	55	55	55	55		12	. 12	12	4
KMnO4 (mg/L)	0.54	0.76	0.99	1.16	1.30	1.57	1.72		0.81	0.95	0.95	0.95
Feed (L/min)	22.7	22.7	22.7	22.7	22.7	22.7	22.7		15.1	15.1	15.1	15.1
Date	9/11/94	9/12/94	9/13/94	9/14/94	9/15/94	9/16/94	9/17/94		9/19/94	9/20/94	9/21/94	9/22/94

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# **APPENDIX B**

Nanofiltration test data

### Nanofiltration Test Data

Turbidity Feed - after cart. filter (ntu)	0.112 0.077 0.064 0.063	0.064 0.066 0.066	0.067 0.069 0.069 0.073 0.073 0.073	0.084 0.073 0.073 0.074 0.070	0.053 0.050 0.046 0.046	0.040 0.043 0.044 0.045 0.051 0.051 0.053	0.057 0.059 0.056 0.061 0.063 0.067 0.065	0.045 0.046 0.046 0.048 0.045 0.043 0.043 0.043
Feed (deg C)	25.9 25.8 26.8 24.9	24.9 24.1 24.1	25.5 24.8 23.7 23.7 24.6	26.1 23.3 24.1 25.6 25.6	23.4 27.3 28.4 23.6 23.6	25.2 27.1 27.2 27.3 27.3 27.3 27.3 27.3 27.3 27.3	24.8 26.7 23.6 23.6 27.1 26.8 26.8	25.2 24.9 24.9 25.1 25.1 25.2 25.2
Feed Feed (deg F) (deg C	78.6 78.4 80.3 76.9	76.8 78.4 76.4	75.0 77.9 76.7 74.7 74.6 76.3	79.0 76.8 73.9 75.3 78.0	74.1 77.1 81.1 79.6 74.5	808 808 743 773 799	76.7 81.1 77.1 80.3 74.5 74.5	77.1 80.9 77.4 76.8 76.1 77.4 76.0
Reject (lb/in2)	44 47 47 46	4 6 4 7 1	\$ 4 4 4 4 4	44 44 44 44 44 44 44 44 44 44 44 44 44	<b>3 2 2 3 4 5 5 5</b>	2444446 2444446	8444884448	0,4448888884444 0,44488884444
Pressure Interstage (Ib/in2)	55 55 55 55 55	88888	8 8 8 8 8 8 8	8 8 2 2 8 8	. 55 55 88 8	8 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	2 22 22 22 24 28 25 22 22 22 24 28 25 24 25 25 25 25 25 25 25 25 25 25 25 25 25	<b>21</b> 28 28 21 21 28 28 28 28 28 28 28 28 28 28 28 28 28
Feed (lb/in2)	75 75 75 75	2723	75 76 78 77	75 76 80 77 77	78 75 80 80 80 80	8	2 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	78 78 80 80 79 81 81 80 80
Permeate Stage 2 (uS/cm)	309 295 283	309 316 290 293	297 289 257 276 294	294 262 281 286 286	270 298 270 270	286 286 286 287 288 288 288 288 288 288 288 288 288	288 288 263 276 278 278 259	282 282 284 284 284 284 284 284 284
Permeate Stage 1/2 (uS/cm)	98.88 88.88	8822	882288	8 9 8 3 2 8 8	8888 888 78	82 82 82 83 84 85 87 87 87 87 87 87 87 87 87 87 87 87 87	987 77 78 74 74 74	. 5
Conductivity Permeate Stage 1/1 (uS/cm)	8 8 8 8 4 8 1	87 87 84 82	282726	87 22 28 72 87 88	<b>75</b> 88 87 82 83 85 85 85	22882424	28 84 75 76 76 76 76 76 76 76 76 76 76 76 76 76	75 72 73 73 70 80 80 70 70
Reject (uS/cm)	5380 5040 4940 5070 4870	4950 4890 5020 5160 5260	5220 5240 5140 5150	5150 5160 5100 5050	5170 5090 4980 5000 5200 5270	5000 4940 5230 5160 5110 5190	5240 5240 5240 5080 5080 5080 5180	5050 5220 5220 5130 5250 5230 5140 5060
Feed (uS/cm)	1170 1170 1170 1170	1170 1170 1170 1170	222523 22223	85 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	1190 1200 1200 1190	1200 1200 1190 1200 1200 1190	1250 1150 1150 1250 1250 1250 1250 1250	1200 1200 1200 1200 1200 1190 1200
Permeate Stage 2 (L/min)	0 0 0 0 0 0 - 0 0 0 0 4		0 0 0 0 0 0 0 0 0 4 0 0 0	, w w w w w w				,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
Permeate Stage 1/2 (Umin)	, 00 00 00 00 00 00 00 00 00 00 00 00 00	5.50 5.00 5.70 5.70 5.00 5.00 5.00 5.00		, 4 6 6 5 6 5 2 4 6 6 5 6 5	5.6 5.6 5.6 5.6 5.6	, , , , , , , , , , , , , , , , , , ,	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	
Flowrates Permeate Stage 1/1 (L/min)	က က လ လ လ လ တဲ့ တဲ့ ဆဲ့ ဆဲ့ ဆဲ့ တဲ့	, ry ry ry ry ra r			9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	ក្រស់សំសំសំសំ 7. សំសំសំសំសំសំសំ	ກ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ ດ <i>ດ</i>	
Reject (L/min)	<u>ლ</u>	, e,	က က က က က က က က က က က က က က က က	6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6				
Feed (Umin)	68 68 68 68 68 68 68 68 68 68 68 68 68 6	18.1 18.1 18.2 18.2 18.2 18.2 18.2 18.2	## ## ## ## ## ## ## ## ## ## ## ## ## ## ##	81 81 82 82 82 82 82 82 82 82 82 82 82 82 82	8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	85 85 85 85 85 85 85 85 85 85 85 85 85 8	48.2 48.2 48.2 48.2 48.2 48.2 48.2 48.2	48.5 48.5 48.5 48.5 48.5 48.5 48.5 48.5
Elapsed Time (hours)	1.0 4.8 7.6 23.0 23.0	200 35.3 50.6 6.6 6.6	54.6 59.4 71.0 74.6 78.6	92.5 94.7 98.6 102.6 118.8	122.8 126.8 130.6 142.5 146.1	150.9 154.6 166.9 171.0 175.1 190.6 194.6	198.6 202.6 214.6 222.6 226.6 238.6 242.6	246.6 246.6 262.6 262.6 270.6 274.6 290.7 294.7

# Nanofiltration Test Data - Continued

Turbidity Feed - after cart. fitter (ntu)	0.0554 0.0654 0.0656 0.0656 0.0445 0.0445 0.0445 0.0446 0.0446 0.0446 0.0446 0.0446 0.0446 0.0446 0.0446 0.0556 0.0566 0.0566 0.0566 0.05666 0	
:_ &	$\begin{array}{c} 2 & 2 & 2 & 2 & 2 & 2 & 2 & 2 & 2 & 2 $	
Feed Feed (deg C	6.84 4.84 6.85 6.85 6.85 6.85 6.85 6.85 6.85 6.85	
Reject (lb/in2)	######################################	
Pressure Interstage (Ib/fin2)	42888888888888888888888888888888888888	
Feed (Ib/in2)	8 7 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	
Permeate Stage 2 (uS/cm)	252 253 253 253 253 253 253 253 253 253	
Permeate P Stage 1/2 (uS/cm)	\$2.52.45.82.45.64.68.65.68.65.45.86.45.65.45.48.48.46.66.88.88.66.46.46.46.46.46.46.66.88.66.86.86.46.46.46.46.46.46.46.46.46.46.46.46.46	
Conductivity Permeate   Stage 1/1 (uS/cm)	5 9 9 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	
Reject (uS/cm)	5280 5210 5210 5210 5210 5210 5210 5210 5220 5230 5230 5230 5230 5230 5230 523	
Feed (uS/cm)	2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
Permeate Stage 2 (Umin)	യ ഒലെ ഒല്ലെ ഒ ഇന്ലെ ഒല്ലെ ഒ	
Permeate F Stage 1/2 (Umin)	\$\pi_\$\pi_\$\pi_\$\pi_\$\pi_\$\pi_\$\pi_\$\pi_	
Flowrates Permeate Stage 1/1 (L/min)	ου ο	
Reject (Umin)		
Feed (L/min)	######################################	
Elapsed Time (hours)	298. 298. 297. 298. 297. 297. 298.	

# Nanofiltration Test Data - Continued

± 9 8	(ulu)		0.055						0.062		. 0.067	0.057	0.056	0.055	0.052	20.0	0.057	0.058	0.057	0.052	0.053	0.073	0.078	0.096	0.081	0.079	2000	0.083	0.081	0.088	0.090	0.101	0.122	0.078	0.077	0.072	0.070	0.069	0.070	0.077	0,080	0.087	0.072	0.077	0.000	0.092	0.077	0.093
Feed Feed	(36eg C)	25.3	23.8	26.0	26.8	25.9	24.6	24.8	26.8	25.8	24.3	25.1	27.8	26.3	24.0 C. 80	27.02	27.0	23.8	25.2	25.9	25.1	23.7	25.3	26.1	25.2	27.0	25.1	24.4	22.7	24.8	23.8	22.9	22.4	25.6	25.1	22.6	24.3	26.9	20.5	24.8	27.3	26.7	33.1	24.8	26.7	23.2	25.1	27.0
Feed	(agg r)	77.6	74.8	78.8	80.2	78.6	76.2	76.7	80.2	78.4	75.7	77.2	82.0	2.67	70.	82.3	80.6	74.9	77.3	78.6	77.2	74.6	77.5	79.0	4.77	7.4.0	77.1	76.0	72.9	7.97	74.8	73.7	77.8	78.4	77.1	72.7	75.7	80.7 4.04	73.6	76.6	81.1	80.0	73.6	70.6 84 1	80.0	73.8	27.2	Q.U.O
Reject	(2)11(7)	46	8	64	82	<b>4</b> 6	2 5		; 4	₹ 5	3 8	3 \$	4 A	8 6	9	47	<del>4</del>	2	22	46	<b>4</b> 8	21	8 !	44	\$ £	3 9	49	2	51	51	និះ		2 6	8	51	52	5 21	8 8	25	25	49	<u>R</u> 1	25	40	200	25	5	ş
Pressure Interstage	(2011)	24	61	8	<b>3</b> 6	ñã	5 6	3 6	č ů	6 4	5	- Q	8 6	8 2	52	28	20	62	61	21	8	<b>79</b>	<u> </u>	2 6	9 6	<u>8</u>	8	8	8	62	3 8	7 9	E	5	82	<b>3</b> 8	2 4	5 E	8	æ	8	æ 3	\$ 8	8 8	6	æ	8 8	3
Feed (lb/in2)	(3,11,12)	82	98	S S	88	8 8	8 8	28	83	8 8	9 8	8 8	8	88	8	82	8	88	82	8	£ 8	8 8	88	7 6	8	80	84	85	87	87	00 6	5 6	8	87	98 1	8	S &	8	8	88	87	£ 6	8	8 6	87	8	8 %	}
Permeate Stage 2 (uS/cm)	,	256	241	707	268 258	238	248	27.0	267	246	254	274	271	240	254	213	265	244	251	797	592	240	274	265	243	254	253	252	240	245 248	243	231	240	250	239	738 74e	255	251	234	205	218	334	205	218	206	227	241	) } !
Permeate Stage 1/2 (uS/cm)		7.	21	7,7	- 1	7	74	6	62	73	75	76	78	7	74	74	77	٤;	4 2	5 3	5 £	2 2	2 2	8	2	78	77	92	2	4 K	2 92	2 2	7.	23	2 ;	C 7	2 22	23	62	74	<b>7</b> 5	2 6	7 7	74	2:	7.	4 E	
Conductivity Permeate Stage 1/1 (uS/cm)		12	9 2	3 6	72	99	7.1	77	12	7	62	7.	74	99	69	2	74	2;	- 8	3 8	7.5	74	62	9.	7	11	74	92	C 8	9 7	75	72	25	13	2 9	8 2	72	69	22	22	C 6	2	2	7.1	67	8 5	<b>2</b> 69	
Reject (uS/cm)	. ;	5040	5250	5200	5280	5070	5110	5070	5080	5160	5220	5100	5490	2080	4960	4860	2000	5130	4800	5050	5150	5130	5170	5210	5250	5240	5220	5240	5120	5180	5280	5260	5150	2020 2020	2080	5180	5040	4950	5230	5140	3010 4950	5230	5140	5010	4950	5110	4710	
Feed (uS/cm)		130	5 5	1190	1190	1190	1190	1190	1190	1190	1190	1190	1190	1190	1190	130	130	5 5	190	1200	1190	1190	1190	1200	1190	1190	1200	£ ‡	1190	12 26	1190	1190	1190	1180	8	1190	1190	1190	1190	255	1190	1190	1190	1200	38	5	198	
Permeate Stage 2 (Umin)		4.4	9.6	3,4	3.4	3.4	3.4	3,4	3.4	3.4	3.4	3.4	3.4	3.4	4.0	4.0	9.0	* tr	46	3.4	3.4	3.4	3.4	3.4	9.00 0.00	4.0	3. c	or en	, e.	3,4	3.4	3.4	3.4	4.4	3.4	3.4	3.4	e. e	4. c	t 7	, e	3.4	3.4	A. 4	, e	, e	3.4	
Permeate Stage 1/2 (Umin)		9 9	9.6	5.6	5.6	5.6	5.6	5.6	5.5	5.8	5.6	5.6	5.6	5.6	n D		ָ פּי	2 6	92	5,5	5.6	5.7	5.6	5.6	9.0	9.0	0 n	o co	5.4	5.6	5.5	5.6	D A	 	5.6	5.6	5.6	9.0	D C	9.69	5.6	5.6	5.6	ro ro ro	5.6	5.6	5.6	
Flowrates Permeate Stage 1/1 (L/min)	Ü	9 45	5,6	5.6	5.6	5.6	5.6	5,5	5.6	5.6	5.6	5.6	5.6	ο c	D 6	, c	5.0	5.6	5.6	5.5	5.6	5.6	5.6	5.6	5.6		ָ טיני	5.6	5.4	5.6	5.6	9.0	0. 4 0. 4	9.00	5.6	9.6	9.6	e S	9 6	5.6	5.6	5.6	5.6	0 0 0	5.6	5.6	5.8	
Reject (L/min)	9	9 6	3.6	3.6	3.6	3.6	3.6	3.6	3.6	3.6	3.6	3.6	9. G	0 0		) e	, e.	9	3.6	3.6	3.6	3.6	3.6	က တ	n e	O 0	9 6	9	3.6	3.6	9.6	20. c	9 60	3.6	3.6	3.6	တွင	9 60	9 69	3.6	3.6	3.6	9 9	ဂ ဂ ဂ	3.6	3.6	3.6	
Feed (Umin)	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	7 0	7.0	, e	18.5	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	10.Z	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2	18.2 4	18.2	18.2	18.2	18.2	18.2	10.2	18.2	18.2	18.2	18.2	
Elapsed Time (hours)	588.8	598.6	902.6	606.6	610.6	623.0	626.6 630.0	500.0 0.00	5.4.6	9,000	65U.6	024.0	670 B	874 A	678.6	682.1	688.3	690.1	694.2		710.1	714.1	718.1	734.3	738.2	7.42.2	746.2	757.9	762.5	766.4	797	786.1	790.1	794.1	806.1	810.1	848.	830.1	834.1	838.1	842.1	830.1	838.1	842.1	854.1	858.1	861.8	

# Nanofiltration Test Data - Continued

Turbidity	eed - after cart. filler	(nlu)	0.079	0.156	0.103	0.135	0.090	0.084	0.078	0.065	0.064	0.071	0.063	0.057	0.058	0.073	0.085	0.092	0.104	0.127	0.069	0.068	0.068	0.063
:	Feed	(deg C)	26.2	23.9	25.1	26.9	26.1	22.3	24.7	26.0	24.8	21.4	25.1	26.1	25.8	24.6	24.8	26.0	24.0	22.7	24.7	26.4	24.9	22.9
Temperature -	Feed	(deg'F)	79.2	75.1	77.1	80.4	79.0	72.2	76.5	78.8	76.6	70.6	77.1	78.9	78.4	76.2	1.91	78.8	75.2	72.9	76.5	79.5	76.9	73.2
	Reject	(lb/lin2)	48	2	51	25	25	8	53	2	જ	24	25	51	51	51	25	25	5	24	33	25	25	54
Pressure	nterstage	(lb/in2)	9	63	63	63	63	89	64	61	<del>.</del>	99	63	62	62	62	83	62	8	65	8	62	SS	65
:	Feed	(lb/ln2)	88	68 60	88	88	88	83 83	8	88	87	92	88	87	87	87	87	98	87	8	86	87	88	<b>8</b>
!	Permeate Stage 2	(mc/sn)	227	225	239	238	232	229	227	239	242	189	236	233	230	227	240	246	239	234	240	241	240	237
	Permeate P		71	71	72	99	67	20	89	72	92	88	72	02	69	02	72	72	74	73	73	71	73	75
Conductivity			29	89	69	99	64	8	99	02	92	73	20	29	99	67	69	69	52	73	72	89	71	74
0	Reject	(mS/cm)	4730	5040	5100	5040	2060	9200	5090	5110	5150	2300	5120	4990	4960	2030	5230	5190	5240	5400	5230	5110	5250	5390
	Feed	(mS/cm)	1190	1190	1190	1190	1190	1190	1190	1190	1190	1180	1190	1180	1180	1180	1190	1190	1180	1190	1180	1190	1180	1180
,	Permeate	(L/min)	3.4	4.60	9.6	3.4	3.4	3.4	3.4	3.4	3.4	3.4	3.4	3.4	3.4	3.4	3.5	3.5	4.6	(C)	10	6	(C)	3.4
,	-		9	6.6	9.5	5.6	5.6	5.6	5.6	5.6	5.6	95	5.6	92	25	9 6	5.7	5.7	5.6	92	22	6	5.6	5.6
Flowrates	Permeate	(Cmin)	η. 6	98	9 6	5.6	5.6	60	. C	5.6	η. 6	8	8.6	2.5	, r.	, rc	2.6	5.7	r.	i re		5.6	e co	5.6
	Reject	(Umin)	6	96	3.6	9 60	3.6																	3.6
	Feed	(L/min)	18.2	<u> </u>	18.5	18.5	18.2	182	i d	18.2	18.2	, d	18.0	i d	1 a	1 4	10.00	182	180	i at	 	182	182	18.2
	Elapsed	(hours)	998	877.8	2.5	88.5	889.2	803 0	0.000	6.00	0,400	042.9	046.5	0.00	0.04.2	0363	. C U 90	044.2	0.48.0	940.2	- 200		66	984.2

### Nanofiltration Calculated Values

Normalized Permeate Flow (Umin)	44446666666664446444664444444444444444	13.8 14.6
Average Net Driving Pressure (Ib/In2)	4 4 8 8 8 8 8 8 4 4 4 4 4 4 4 4 4 4 4 4	43.8
Average Osmosis Pressure (Ib/In2)	22222222222222222222222222222222222222	21.2
Average (Cf+Cr)/2 (mg/L)	2218 2013 2013 2013 2013 2013 2013 2013 2013	2116 2171
Concentration Reject Cr (mg/L)	3849 3418 3418 3405 3405 3514 3514 3514 3514 3514 3514 3514 351	3432 3534
Feed Cf (mg/L)	787 787 787 787 787 787 787 787 787 787	807 807
Average Feed Pressure (lb/ln2)	4000 P P 8000 P P 800	63.5
Inverse Temp. Correction Factor (1/TCF)	0.974 0.947 1.0034 1.0034 1.0036 1.0036 1.0036 1.0036 1.0036 1.0037 1.0037 1.0037 1.0037 1.0037 1.0037 1.0037 1.0038 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998 0.998	1.000
Temperature Correction Factor (TCF)	1.023 1.023 1.023 0.999 0.997 1.033 0.997 1.015 0.965 0.998 1.002 1.002 1.002 1.002 1.003	1.000
Stage 2 (Ib/in2)	ფით ღით ით ით ით ით ენი ენი ინი ით ით ით ინენი ინე იდ ით ით ით ინენი ინი ინე ით ენენი ინი ით ით ით ით ინენი ინი	. ¢
Stage 1 (lb/in2)	388888888888888888888888888888888888888	33
Elapsed Time (hours)	4.8 1.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2	294.7

Normalized Permeate Flow (L/min)	44 4 6 6 4 6 6 6 4 6 4 4 4 6 4 4 6 4 4 6 4 4 6 6 4 6 6 4 6 6 4 6 6 4 6 6 4 6 6 4 6 6 4 6	8 8 4 4 8 8 8 8 8 8 4 4 4 8 4 8 4 8
Average Net Oriving Pressure (Ib/in2)	24444444444444444444444444444444444444	64 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4
Average Osmosis Pressure (lb/in2)	22222222222222222222222222222222222222	28822222222222222222222222222222222222
Average (Cf+Cr)/2 (mg/L)	2198 2294 2294 2294 2294 2294 2294 2394 2394	
Concentration Reject Cr (mg/L)	3581 3468 3478 3534 3527 3527 3527 3527 3527 3528 3528 3528 3528 3528 3528 3528 3528	3568 3574 3574 3574 3574 3574 3574 3574 3574
Feed Cf (mg/L)	801108880101010101010101010101010101010	8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8
Average Feed Pressure (lb/in2)	\$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$	\$\$\$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$
Inverse Temp. Correction Factor (1/TCF)	1.007 1.044 1.044 1.045 1.089 1.089 1.080 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010 1.010	1.028 0.980 1.028 1.028 1.028 1.039 1.039 1.031
Temperature Correction Factor (TCF)	0.993 0.983 0.983 0.977 0.977 0.984 0.984 0.984 0.984 0.984 0.986 0.986 0.987 0.987 0.987 0.987	0.977 0.989 0.989 0.973 1.027 1.028 1.008 1.008 1.008 1.008 1.008 1.008 1.008 1.012 1.012 1.012 1.013 1.013
P Stage 2 (Ib/In2)		560110000110000001 <b>5</b> 6
Stage 1 (Ib/In2)		***************************************
Elapsed Time (hours)	98 3322.6 3322.6 3322.6 3322.6 3323.6 3323.7 3323.6 3323.7	4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4

# Nanofiltration Calculated Values - Continued

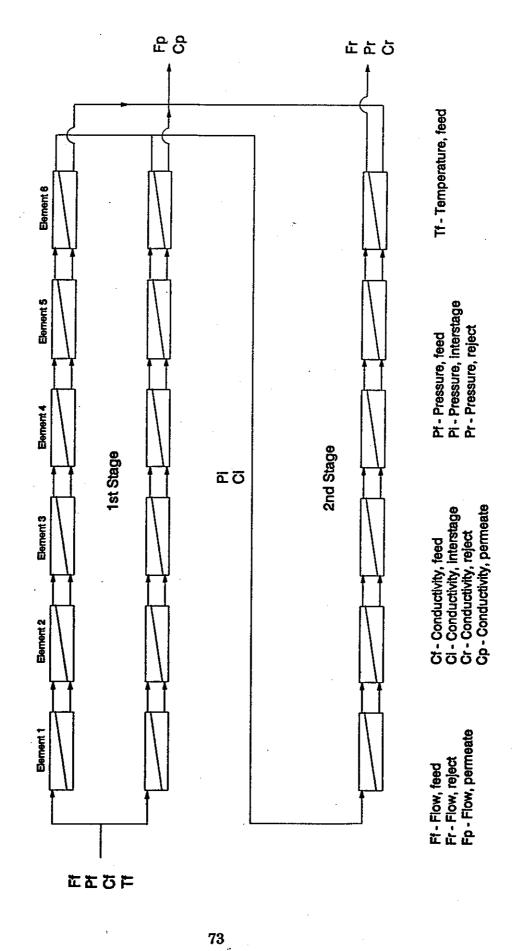
Normalized Permeate Flow (L/min)	1.4	12.9	13.2	13.6 43.6	13.5	. e	13.5	13.4	13.1	12.4	13.1	13.2	12.9	, C	12.5	. e.	14.5	13.8	13.4	13.2	13.8	13.8	13.7	0.5		5.5	12.4	13.6	13.6	13.3	12.9	12.6	12.9	5.5	12.5	126	130	12.6	12.2	12.2	13.0	12.6	12.2 43.3	13.0	12.4	12.4
Average Net Oriving Pressure (Ib/in2)	42.9	45.9	43.9	2. A. 2. A. 3. A.	45.7	43.3	43.8	46.5	46.3	45.2	44.9	46.8	45.7	2.44 0.44	47.5	46.6	419	43.5	47.0	46.1	43.0	43.8	46.2	40.2	4 4 6 4 7 4	46.0	47.6	46.4	47.1	49.2	49.0	47.4	7 6	787	47.4	46.7	49.3	48.6	47.0	47.7	49.3	48.ts	4 4 5 4	49.4	49.2	46.5
Average Osmosis Pressure (lb/in2)	21.1 21.1		21.6		2 <u>2</u>	21.2		21.5	21.7	21.3	22.6	21.2	8.00	2.0.0	2.5	20.9	21.1	21.0	21.5	21.4	21.5	21.7	27.8	6.1.6	- 4 - 4	, c , c	21.4	21.8	21.9	21.8	24.5	27.7	2 <u>2</u>	2 <u>2</u> 6 &	2.5	20.8	21.7	21.4	21.0	20.8	21.7	27.4	5. 6 5. 6	21.6	21.3	20.0
Average (Cf+Cr)/2 (mg/L)	2109	2108	3,5	200	2133	2120	2123	2150	2171	2130	2262	2000	2002	2008	2140	2089	2113	2104	2147	2140	2154	2171	2181	2476	247	22.	2137	2157	2191	2184	2147	2113	2450	2457	2109	2079	2174	2143	2103	2079	212	2143	2012	2164	2133	1998
Concentration Reject Cr (mg/L)	3418	3412	3527	3439	3466	3439	3446	3200	3541	3459	3724	3446	9300	3301	3479	3378	3425	3400	3493	3479	3507	3534	3551	35.44	9554	3622	3473	3513	3581	3568	3493	3425	2456 9468	3513	3418	3357	3547	3486	3398	3357	3547	0000	3357	3527	3466	3195
Feed Cf (mg/L)	801 801	804	8	8 8	8	801	801	8	80	5	5	5	5	3 2	801	8	801	807	108	8	8	200	500	<u> </u>	8	8	801	801	8	83	8	5 6	<u> </u>	8	9	801	801	801	807	56	5 6	- 20	8 8	801	8	801
Average Feed Pressure (lb/in2)	64.0 0.0	67.0	65.5 8.5 8.5	0.89	0.79	64.5	65.0	68.0	68.0	66.5	4.75 4.05	0.00	8 8 8 8 8	85.5	68.5	67.5	63.0	64.5	68.5	67.5	64.5	65.5	67.0	86.5 5.5	67.5	0.69	0.69	68.0	0.69	71.0	6.05	00.0	71.0	70.5	68.5	67.5	71.0	70.0	0.80	68.5	0.52	9	68.5	71.0	70.5	<b>6</b> 6.5
Inverse Temp. Correction Factor (1/TCF)	0.990	0.971	0.949	1.012	1.004	0.949	0.977	1.019	0.997	128.0	20.0	- C	0.933	0.943	1.031	0.995	0.974	0.997	1.036	0.992	0.968	0.893	555	0.998	1.015	1.063	1.004	1.033	1.058	1.071	1.033	78.0	1.086	1.019	0.946	0.961	1.052	1.006	0.935	798.0	200.1	0.00	0.952	1.048	0.997	0.943
Temperature Correction Factor (TCF)	.1.010 0.968	1.030	1.054	0.988	0.998	1.054	1.023	0.981	1.003	000.	200.E	1 035	100	1.061	0.970	1.005	1.027	1.003	0.965	1.008	1.033	7.00.0	066.0	1.002	0.986	0.941	966.0	0.968	0.945	0.934	0.800	1002	0.938	0.981	1.057	1.040	0.951	0.994	1.070	100.0	0.931	1.070	1.051	0.954	1.003	1,001
Stage 2 (Ib/In2)	<del>**</del> **	<b>=</b> :	= =	==	<del>=</del>	2	Ξ;	:3		2 \$	2 ∓	ç	<del>:</del> <del>-</del>	=	=	7	<del>-</del>	₽	Ξ	<b>=</b> :	= ;	_ +	===	==	9	=	Ξ	12	= 9	2.	= ;	: =	12	2		7	7	2;	Ξ:	- 5	4 5	-	Ŧ	Ξ	₽;	Ë
Stage 1 (Ib/In2)	52 52 52	25	2,4	25	25	<b>5</b> 2	52	52	2 20	2 6	3 5	25.55	24	24	24	54	23	83	24	55	\$ 2	* K	22 22	54	52	25	52	24	<b>5</b> 2	9 %	9 8	24	<b>5</b> 8	27	26	52	26	92	77	92	20 20 20	27	56	27	27	07
Elapsed Time (hours)	586.6 598.6	602.6	610.6	623.0	626.6	630.6	634.8	8.040 8.040	654.8	9	870.6 870.6	674.6	678.6	682.1	688.3	690.1				714.1	1.817	734.2	738.2	742.2	746.2	757.9	762.5	766.4	770.2	7884	58.5	794.1	806.1	810.1	814.1	818.1	830.1	634.1	920.	830.1	834.1	838.1	842.1	854.1	858.1 964.0	0.100

Normalized Permeate Flow	12.5	12.8 12.6	8.5	12.7	12.2	12.6	13.1	13.2	12.5	12.3	12.4	12.9	13.1	12.8	13.3	13.2	12.6	12.2	12.8	13.0
Average Net Driving Pressure (lb/in2)	47.0	48.9 48.2	48.9	51.8 51.8	50.2	47.2	47.0	51.1	48.6	48.1	48.2	47.8	47.8	47.4	47.3	49.7	49.3	48.2	48.2	49.8
Average Osmosis Pressure (lh/in2)	20.0	21.1 21.3	21.1	22.7	21.3	21.3	21.5	21.9	21.4	20.9	20.8	21.2	21.7	21.6	21.7	22.3	21.7	21.3	21.8	22.2
Average (Cf+Cr)/2	• ``	2109 2130	2109	2265	2126	2133	2147	2194	2137	2089	2079	2123	2174	2160	2174	2232	2171	2133	2177	2225
Concentration Reject Av Cr (Cf	3208	3418 3459	3418	3730	3452	3466	3493	3595	3473	3385	3364	3452	3547	3520	3554	3663	3547	3466	3561	3656
Feed Cf	. 2	8 8	80	88	801	8	8	794	801	794	794	794	801	801	794	801	794	801	794	794
Average Feed Pressure		70.0 69.5	70.0	74.5	71.5	68.5	68.5	73.0	70.0	0.69	0.69	0.69	69.5	0.69	0.69	72.0	71.0	69.5	70.0	72.0
Inverse Temp. A Correction Factor Pi	0.964	1.028 0.998	0.946	1.074	1.007	0.971	1.006	1.102	0,998	0.969	0.977	1.012	1.004	0.971	1.027	1.063	1.007	0.960	1.001	1.058
Temperature Correction Factor		0.973 1.002	1,057	0.931	0.993	1.030	0.994	0.908	1.002	1.032	1.023	0.988	0.996	1.030	0.974	0.941	0.993	1.042	0.999	0.945
Stage 2 (Ib/in2)	8	1 1 2 1	Ξ;	- 2	1	10	Ξ	4	<del>-</del> -	=	=	÷	Ξ	9	£	÷	=	9	Ξ	£
Stage 1	26	28 25	25	<b>52</b>	56	25	56	58	25	25	25	25	24	24	25	25	25	25	25	25
Elapsed Time (hours)	865.6	877.6 881.6	885.2	893.9	897.9	901.9	905.9	912.3	916.3	920.3	924.3	936.3	940.2	944.2	948.2	960.1	964.5	968.5	972.5	984.2

### APPENDIX C

Generalized NF process diagram for checking data reduction

Generalized NF Process Diagram for Checking Data Reduction



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### **APPENDIX D**

Analytical data for nanofiltration testing

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Analytical Data for Nanofiltration Testing

					5-Hour	Data			362-Hour	: Data	
ð	CATIONS			Eeed	Interstage	Permeate	Reject	Feed	Interstage	Permeate	Reject
ĬŐ.	Calcium .	Ö	mg/L	88		6.1	420	84	220	£.	390
Σ	Magnesium	Βġ	mg/L	တ္တ		0.69	130	29	65	0.34	120
<b>й</b> .	Sodium	g Z	mg/L	110		22	510	110	250	17	480
<u>~</u> :	Potassium	×	mg/L	3.0	∞	_	16	3.5	7.2	-	13
₹ 1	Aluminum	₹	mg/L	<0.05		<0.05	0.08	<0.20	<0.20	<0.20	<0.20
a a	Barium	Ba •	mg/L	<0.05		<0.05	0.17	<0.05	0.07	<0.05	0.15
เก	Strontium	ັດ	mg/L	1.2		۸ <u>۲</u> ۰	5.1	1.3	က	V.	60
<b>≝</b> ∶	Lon	Đ.	mg/L	<0.02		<0.02	0.03	<0.05	<0.05	<0.05	0.07
∑i	Manganese	S N	mg/L	0.50		<0.05	2.5	0.50	7	<0.05	2.4
ā	Phosphorus	Δ.	mg/L	0.11		<0.10	0.58	<0.10	0.26	<0.10	<0.10
A	ANIONS										
<b>商</b>	Bicarbonate	HC03	mg/L	170	400	28	770	170	410	14	790
ਹ 	Chloride	ច	mg/L	94	250	28	280	130	250	22	390
์ เ	Sulfate	804	mg/L	290	720	9	1112	290	730	7.2	1500
<b>Z</b> i	Nitrate (N)	(Z) (NO3 (NO3)	mg/L	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50
Ī	Flouride	ш	mg/L	<b>6</b> .	4.	0.2	ღ	0.66	1.6	0.1	က
ଅଁ	Silica (total)	SIO2	mg/L	18	42	5.4	74	18	99	3.2	69
۴	Total Organic Carbon	<b>10</b> C	mg/L	1.7	•	•	•	1.8	•	•	ı
が	Standard Plate Count	SPC	nJo	9200	•	•	•	4000	•	•	
₹ï	Alkalinity	, C	7/6m	170	400	28	770	170	410	4	790
=	00000	88 CgCC	mg/L	340	820	7.6	1600	330	770	5.1	1500
ပိ ပိ	Conductivity (lab)		uS/cm	1200	2600	160	4900	1200	2600	110	4700
3 É	Conductivity (ops)	(1) (A) (A)	Eo/sn	1170	• (	138	4900				
2	rotal Dissolved Solids	(mns) con	-1/BH	807	1979	64	3323	837	1977	99	3764
Ą	Anions (Ca,Mg,Na,K)		meq/L	11.74	28.42	1.13	54.25	11.45	27.39	0.87	50.55
3 &	Cations (ncOs,Ci,SO4) Ratio Anions Cations		meq/L	11.48	28.60	1.46	43.67	12.49	28.97	1.00	55.18
			•	1.02	9 9 9	. O	1.24	0.92	0.95	0.87	0.92

Analytical Data for Nanofiltration Testing - Continued

				693-Hour	Data			984-Hour	Data	
SNOTA			Feed	Interstage	Permeate	Reject	Feed	Interstage	Permeate	Reject
Calcium	Ca	mg/L	84	200	1.3	380	86	200	0.87	400
Magnesium	<b>5</b>	mg/L	ည ရှင်	69	0.46	130	500	ဗေ ဗိ	0.33	120
Dotagalim	 Z	mg/L mg/l	200	700	7 7	4 0 6 6	120	750 250	7,	200
Aluminum	۷ ح	mg/L	<0.20	<0.20	<0.20	-0.20	<0.20	0.00 0.00 0.00	0.10	- 00 - 00 - 00 - 00 - 00 - 00 - 00 - 00
Barium	Ва	mg/L	<0.05	0.08	<0.05	0.17	<0.05	0.08	<0.05	0.16
Strontium	S	mg/L	1.3	3.4	<1.0	7.1	1.6	3.7	۸.15 م	7.4
Iron	Fe	mg/L	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05
Manganese	Mn	mg/L	0.54	1.3	<0.05	2.6	0.53	1.2	<0.05	2.3
Phosphorus	۵.	mg/L	0.12	0.27	<0.10	<0.10	0.12	0.28	<0.10	<0.10
ANIONS										
Bicarbonate	HC03	mg/L	180	400	5	820	170	390	15	780
27 Chloride	ច	mg/L	120	220	23	380	96	230	20	430
	S04	mg/L	290	730	7	1500	300	720	6.4	1600
Nitrate (N)	NO3 (N)	mg/L	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50	0.7
Fiouride	L	mg/L	0.73	1.7	0.11	ო	0.71	<del>.</del> 6	0.12	က
Silica (total)	SiO2	mg/L	18	40	3.6	69	19	42	3.3	77
Total Organic Carbon	<b>TOC</b>	mg/L	1.7	•	•	•	ო	1	•	•
Standard Plate Count	SPC	cfu	1100	1200	790	1900				
Alkalinity Hardness	as CaCO3	mg/L mg/L	180 330	400 780	5.1	820 1500	170 330	390 760	15 3.5	780 1500
Conductivity (lab)		uS/cm	1000	2200	86	.4000	1100	2300	9	4300
Total Dissolved Solids	TDS (sum)	mg/L	835	1933	74	3795	825	1907	63	3932
Anions (Ca,Mg,Na,K) Cations (HCO3,CI,SO4) Ratio Anions:Cations		meq/L meq/L -	11.38 12.37 0.92	27.14 27.96 0.97	0.97 1.11 0.88	51.31 55.39 0.93	11.97 11.71 1.02	26.18 27.87 0.94	0.81 0.94 0.86	51.87 58.22 0.89

### APPENDIX E

Nanofiltration element serial numbers as loaded in pressure vessels

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•	•					نسب
•						
						Normal
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		•				

# NF Element Serial Numbers as Loaded in Pressure Vessels

	·	<u> </u>	]						1		,	
· ·		A2282495		A2282536		A2282515		A2282522		A2282501		A2282494
; Date 9/30/94			•									
; Date												
; Model NF70-2540		A2282531		A2282546		A2282481	(u	A2282524		A2282489		A2282466
Model	front)		rear)		from top)		from bottor		m rear)		n front)	
	4-6 (top		1-3 (top		-6 (2nd		-6 (2nd t		-3 (botto		3 (botton	
Filmtec	Elements 4	2539		2470	lements 4	2542	lements 4	2514	ements 1-	534	ements 1-3	535
Manufacturer	Stage 2, Vessel 1B, Elements 4-6 (top front)	A2282539	Stage 2, Vessel 1A, Elements 1-3 (top rear)	A2282470	Stage 1, Vessel 2B, Elements 4-6 (2nd from top)	A2282542	Stage 1, Vessel 1B, Elements 4-6 (2nd from bottom)	A2282514	Stage 1, Vessel 2A, Elements 1-3 (bottom rear)	A2282534	Stage 1, Vessel 1A, Elements 1-3 (bottom front)	A2282535

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### APPENDIX F

Lime softening process calculations

### COLD LIME PROCESS - Lake Havasu City - Well S6

```
FEED WATER COMPOSITION
                   mg/L
  Na+
            110.00
                                4.78
                                      meq/L
                                                   239.45
                                                           mg/L (as CaCO3)
  K+
              1.80
                   mg/L
                           =
                                0.05
                                      meq/L
                                            =
                                                     2.30
                                                           mg/L (as CaCO3)
  Ca++
            100.00
                   mg/L
                                4.99
                           =
                                      meg/L
                                            =
                                                   249.72
                                                           mg/L (as CaCO3)
  Mq++
             35.00
                                2.88
                                      meq/L
                   mg/L
                           ==
                                            ==
                                                   144.09
                                                           mg/L (as CaCO3)
S cations
                               12.70
                                      meq/L =
                                                   635.56
                                                           mg/L (as CaCO3)
  Cl-
            95.00
                   mg/L
                                2.68
                                      meq/L
                                            ==
                                                   134.10
                                                           mg/L (as CaCO3)
  HCO3-
            232.00
                                3.80
                   mg/L
                                                   190.29
                           =
                                     meq/L
                                            =
                                                           mg/L (as CaCO3)
                   mg/L
  SO4=
            300.00
                                6.25
                                     meq/L
                                            =
                           =
                                                   312.57
                                                           mg/L (as CaCO3)
  NO3-
              4.42
                   mg/L
                                0.07
                                     meq/L
                                            ==
                                                     3.57
                                                           mg/L (as CaCO3)
  CO3=
              0.00
                   mg/L
                           =
                                0.00
                                      meq/L
                                                     0.00
                                                           mg/L (as CaCO3)
                                                   0.00
  OH-
              0.00
                   mg/L
                           =
                                0.00
                                      meg/L
                                            ==
                                                           mg/L (as CaCO3)
S anions
                               12.80
                                     meq/L =
                                                   640.53 mg/L (as CaCO3)
 Cation/Anion Ratio
                               0.99 Balance is excellent.
 TDS =
           878.22
 pH =
             7.62
                                     R
                                                    20.89
 CO2
             9.11 mg/L (as CO2)
 Total Hardness (TH) =
                                                   393.81 mg/L (as CaCO3)
 Alkalinity (Alk)
                                                   190.29 mg/L (as CaCO3)
 COAGULENT-CORRECTED COMPOSITION
 Coaqulent:
                             Aluminum Sulfate (20 ppm)
                             Ferrous Sulfate (20 ppm)
                           4 Ferric Sulfate (10 ppm)
                             Sodium Aluminate (10 ppm)
 Total Hardness (TH)
                                                   393.81
                                                          mg/L (as CaCO3)
 Alkalinity (Alk)
                                                   184.29
                                                          mg/L (as CaCO3)
 Calcium Alkalinity (Ca Alk) =
                                                   184.29
                                                          mg/L (as CaCO3)
 Magnesium Alkalinity (Mg Alk) =
                                                          mg/L (as CaCO3)
                                                    0.00
 Sodium Alkalinity (Na Alk) = .
                                                          mg/L (as CaCO3)
                                                    0.00
 Calcium Noncarbonate Hardness (NCH) =
                                                   65.43
                                                          mg/L (as CaCO3)
 Magnesium Noncarbonate Hardness (MgNCH) =
                                                  144.09
                                                          mg/L (as CaCO3)
 Total Noncarbonate Hardness (NCH) =
                                                          mg/L (as CaCO3)
                                                  209.53
 Sulfate
                                                  318.57
                                                          mg/L (as CaCO3)
 Free Carbon Dioxide =
                                                   15.11
                                                          mg/L (as CO2)
 DOSAGE OF LIME REQUIRED
 Hyd. Lime - 93% Ca(OH)2 =
                                185
                                                    1.55
                                                          lbs/1000 gallons
                                       ppm =
 Chemical Lime - 90% CaO =
                                145
                                       ppm =
                                                    1.21
                                                          lbs/1000 gallons
 Theor. Lime - 100% CaO =
                                131
                                       ppm =
                                                    1.09
                                                          lbs/1000 gallons
        COLD LIME PROCESS - Lake Havasu City - Well S6
                                                          (continued)
 EFFLUENT COMPOSITION
                                                  114.84
 Calcium Hardness =
                                                          mg/L (as CaCO3)
 Magnesium Hardness =
                                                  129.68
                                                          mg/L (as CaCO3)
 Total Hardness =
                                                  244.53
                                                          mg/L (as CaCO3)
 Alkalinity =
                                                   35.00
                                                          mg/L (as CaCO3)
```

R

5011.87

10.0 (estimated)

= Hq

CO2 0.01 mg/L (as CO2)

DOSAGE OF ACID REQUIRED FOR PH ADJUSTMENT

```
4 93% H2SO4 =
                               2.46
                                       = mgg
                                                   0.02
                                                          lbs/1000 gallons
        35% HCl =
                               4.86
                                      ppm
                                                   0.04
                                                          lbs/1000 gallons
                                           =
              7.5 (target)
                                     R
                                                   15.85
 pH =
                                            =
  TREATED WATER COMPOSITION
           110.00 mg/L
                               4.78
                                                  239.45
                                     meq/L =
                          ==
                                                         mg/L (as CaCO3)
  K+
            1.80 mg/L
                          ==
                               0.05
                                     meq/L =
                                                   2.30
                                                         mg/L (as CaCO3)
                                    meq/L = meq/L =
                               2.29
            45.99 mg/L
                                                         mg/L (as CaCO3)
  Ca++
                          =
                                                  114.84
                                                         mg/L (as CaCO3)
  Mq++
            31.50 mg/L
                               2.59
                                                  129.68
S cations
                               9.72
                                     meg/L =
                                                  189.29
                                                         mg/L (as CaCO3)
  Cl-
            95.00
                               2.68
                   mg/L
                                     meq/L =
                                                  134.10
                                                         mg/L (as CaCO3)
  HCO3-
            39.82
                   mg/L
                          ==
                               0.65
                                     meq/L =
                                                         mg/L (as CaCO3)
                                                  32.66
                               6.41
  SO4=
           308.00
                   mg/L
                          =
                                     meq/L =
                                                  320.91 mg/L (as CaCO3)
  NO3-
             4.42 mg/L
                               0.14
                                     meq/L =
                                                   3.57 mg/L (as CaCO3)
  CO3=
             0.00 mg/L
                               0.00
                          =
                                     meq/L =
                                                   0.00 mg/L (as CaCO3)
  OH-
             0.00 mg/L
                               0.00
                          ==
                                     meg/L
                                           =
                                                   0.00 mg/L (as CaCO3)
S anions
                               9.89
                                     meq/L =
                                                 447.25 mg/L (as CaCO3)
 Cation/Anion Ratio
                               0.98 Balance is good.
 TDS =
         636.53
 CO2
             2.06 mg/L (as CO2)
```

### Mission

The mission of the Bureau of Reclamation is to manage, develop, and protect water and related resources in an environmentally and economically sound manner in the interest of the American Public. • .