Corrosion in your distribution system. the silent pipe killer, Clean and Safe **Drinking Water Workshop** Gander, NL 2004



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Overview of Presentation

What is corrosion?
Why do we care?
How to determine if your water is corrosive
How to control corrosion
Example where corrosion control measures are being implemented



What is Corrosion?

 Gnawing away or attack on a material, usually by some chemical or electrochemical means

Thinning of pipe walls or,

 Formation of a precipitate on pipe walls









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Why Should We Care?

Corrosion leads to two different kinds of problems
Pipe killer - failure of pipe
Water quality changes (aesthetic and health)
Economics - investment in buried infrastructure (pipes, valves, etc)

> Greater than the cost of the treatment plant



Source: EPA's 1999 Drinking Water Infrastructure Needs Survey



Failure of Pipes

Impact on carrying capacity of a pipe

- Thinning of pipe
 - Pipe leakage
 - Catastrophic pipe failure
- Formation of a precipitate
 - Loss of hydraulic capacity
- Economic impact
 - Early replacement of piping, plumbing and water heaters
 - Increased pumping costs





Water Quality Changes

Otential to remove metals (copper, lead and iron) from pipes in distribution system

 Health concerns - toxic properties of metals such as lead

Regulatory compliance

Staining of laundry and plumbing fixtures

Aesthetic objectives

Red/brown from iron, blue/green from copper

Creates areas for microorganisms to live

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Factors Impacting Corrosion

Dissolved oxygen

PH - acidic or caustic nature of water

- Alkalinity capacity of water to neutralize an acid, expressed as mg/L of CaCO₃
- Calcium concentration
- Suspended solids and organic matter
- Free chlorine residual
- ♦ Temperature

How to Measure Water's Potential for Corrosion

- Many methods can be used
- Visual signs

Dirty water, staining, corroded pipe, pipe failures
 Measurement and observation in distribution

system

 Compare copper, iron or lead levels at the plant with first flush concentration at consumer taps

- True indication of what is happening in distribution system
 - Associated analytical costs

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How to Measure Water's Potential for Corrosion

Calculation using a corrosiveness index
 Based on pH, alkalinity and calcium

- Aggressive Index
- Ryznar Index
- CCPP (Calcium Carbonate Precipitation Potential)
- Langelier Saturation Index (LSI)

Computer programs simplify the calculations
 Rough guide - there is no perfect index
 Corrosion chemistry is very complex.

Corrosiveness Index

Langelier Saturation Index (LSI) - most commonly used

- Estimates the theoretical tendency of a water to form a protective coating of calcium carbonate (CaCO₃) on a pipe wall
 - Under-saturated water is corrosive (negative LSI)
 - Most raw surface waters in Newfoundland
 - Over-saturated water will deposit calcium carbonate (positive LSI)



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LSI can be interpreted as the pH change required to bring the water to equilibrium

- Indicator (not an absolute measurement) of the potential for corrosion or deposits
- ♦ Recommended LSI range +0.5 to -0.5



LSI Calculation

\diamond LSI = pH - pH_s

PH = measured water pH

PH_s = pH at which a water is saturated with CaCO₃

= (9.3 + A + B) - (C + D)

▲ A = $(Log_{10} [Total Dissolved Solids in mg/L] - 1)/10$ ▲ B = -13.12 x Log₁₀ (Temperature in °C + 273) + 34.55 ▲ C = Log₁₀ [Calcium hardness in mg/L as CaCO₃] - 0.4 ▲ D = Log₁₀ [Alkalinity in mg/L as CaCO₃]



LSI Calculation

\diamond LSI = pH - pH_s

- PH = measured water pH
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 - = (9.3 + A + B) (C + D)
 - $A = (Log_{10} [Total Dissolved Solids in mg/L] 1) /10$
 - ▲ B = -13.12 x Log₁₀ (Temperature in °C + 273) + 34.55
 - \land C = Log₁₀ [Calcium hardness in mg/L as CaCO₃] 0.4
 - \blacktriangle D = Log₁₀ [Alkalinity in mg/L as CaCO₃]
- AWWA website LSI calculator/program
- Sample calculation provided at end of handout



Factors Affecting LSI

LSI will increase in value if
Calcium hardness is increased
pH is increased
Alkalinity is increased
Temperature is increased
A change in pH has greatest impact on LSI



Mitigating Corrosion

Use only pipe and materials that can withstand corrosive water

PVC, stainless steel, lined steel

Replace existing pipes/plumbing material/solder

• Expensive!

 Adjusting the water chemistry to make the water non-corrosive

Change pH and alkalinity of water
 Add a chemical to line the pipe
 Corrosion inhibitor

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 Alter the water quality such that chemical reactions between the water supply and pipe material favour the formation of a protective layer on the interior of the pipe walls

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Adjust the pH and alkalinity of the water to make it non-aggressive and stable

Increase pH (ideally > 8.5)

- Increase alkalinity to 30 to 40 mg/L as CaCO₃
 - Maintain a constant water quality in the distribution system

Various chemicals can be used
Increase alkalinity and pH differently
Chemical form varies
Application method varies
Poorly buffered waters (low alkalinity and pH) may require a combination of chemicals



	Chemical	Use	Composition	Alkalinity Change per mg/L of Chemical	
6	Caustic Soda NaOH	Raise pH	 93% purity liquid Storage at <50% purity to prevent freezing 	1.55 mg/L CaCO₃	
Ş	Lime Ca(OH) ₂	Raise pH	95-98% purityDry storage with slurry feed	1.21 mg/L CaCO ₃	
	Sodium Bicarbonate NaHCO ₃	Little increase in pH	98% purityDry storage with solution feed	0.60 mg/L CaCO₃	
	Soda Ash Na ₂ CO ₃	Moderate increase in pH	95% purityDry storage with solution feed	0.90 mg/L CaCO ₃	
9	From AWWA Lead and Copper Rule Guidance Manual, Volume 2: Corrosion Control				

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Liquid Chemical



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Adjusting the Water Chemistry

 LSI can be used to estimate the impact of water chemistry adjustment on corrosion control
 Example

Raw water characteristics

- pH = 6.2
- Alkalinity = 3 mg/L as CaCO₃
- Calcium hardness = 5 mg/L as CaCO₃
- Temperature = 4°C
- ◆ LSI = -4.5

• Water is corrosive - tendency to remove metals from pipes

Example (continued)

- Add 12 mg/L of soda ash to water
- Treated water characteristics
 - ◆ pH = 8.5
 - Alkalinity = 14 mg/L as CaCO₃
 - Calcium hardness = 5 mg/L as CaCO₃
 - ◆ LSI = -1.3
 - Treated water is less corrosive than raw water (LSI = -4.5)

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Adjusting the Water Chemistry

Selection of chemical depends on various factors
 Cost of chemical and associated equipment
 Impact on water quality

- Lime addition can increase aluminum concentration
- Sodium based chemicals will increase sodium concentration of water
- Addition of a batched chemical (lime, soda ash, etc.) may increase turbidity of water
- O&M and safety

Type of treatment provided at facility

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Example - Conventional Treatment Process Train

Coloured waterHigh organic content

Coagulant (Alum) Raw Water Coagulation Flocculation/Clarification Filters Clearwell

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Items to Consider

Do you have sufficient alkalinity for coagulation?

Coagulant consumes alkalinity

- 1 mg/L of alum consumes ~ 0.5 mg/L as CaCO₃
- 1 mg/L of polyaluminum chloride (SternPAC) consumes
 ~ 0.18 mg/L as CaCO₃

Recommended to have > 20 mg/L as CaCO₃ prior to coagulation for complete coagulation reaction

Minimize aluminum residuals

Optimize organics, turbidity and colour removal

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Items to Consider

Is the pH optimal for coagulation?

- Alkalinity adjustment chemicals typically increases pH
- PH reduction will most likely be required
 - Minimize aluminum residuals
 - Optimize organics, turbidity and colour removal
- PH adjustment chemical
 - Sulphuric acid
 - Carbon dioxide (CO₂)
 - Does not consume alkalinity

Items to Consider

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Is your water non-corrosive following filtration?

- Low pH required for good coagulation and effective disinfection
- Higher pH is required to make water less corrosive
- Preference to use caustic soda (NaOH)
 - Does not impact turbidity to the water
- If insufficient alkalinity prior to NaOH adjustment, pH will drastically increase
 - Important to have > 20 mg/L of alkalinity as CaCO₃ after coagulation

 Recommended treated water alkalinity = 30 to 40 mg/L to maintain water quality stable in distribution system















Example - Conventional Treatment Process Train

Coloured waterHigh organic content





Example - Classic Conventional Treatment Process Train

Coloured water
High organic content



Sodium silicates, polyphosphates and orthophosphates Proprietary chemicals ANSI/NSF 60 approval or equivalency Cover potentially corrosive sites by passivating the metal surfaces in contact with water or by forming a protective film or scale Long time to form film Long time to assess effectiveness



Example - Classic Conventional Treatment Process Train

Coloured water
High organic content

Lime Coagulant CO_2 (Alum) Cl₂ To Distribution \mathbf{X} Raw Water Clearwell Coagulation **Filters** Flocculation/Clarification **CH2MHILL**



Example - Classic Conventional Treatment Process Train

Coloured water
High organic content





Example - Membrane Filtration Plant

- Low turbidity water
- Low colour
- Low organics



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Items to Consider

Location of water chemistry adjustment
 Turbidity impact of chemical
 Membrane supplier preferences
 May not require pH adjustment with carbon dioxide

 Depends on selection of alkalinity adjustment chemical

Depends on raw water alkalinity

Dosage determine by trial

Assistance from chemical supplier

Site specific (source water, material and condition of pipes)

Want to maintain slight phosphate or silica residual

Water sampling - metals, phosphate or silica



Addition of corrosion inhibitor does not change LSI

Performance is difficult to assess without obtaining samples in distribution system
 May require pH correction to effectively work
 May require a constant pH
 Issue for poorly buffered water



 Selection of inhibitor is dependent on target metal

 May cause microbial growth in distribution system

Addition of nutrients

 Phosphate based corrosion inhibitors may result in additional wastewater treatment





Do I Need to Implement Corrosion Control Measures?

Determine corrosivity of water
 LSI < -1
 Recommended to perform a full-scale corrosion monitoring study in distribution system

Exceed MAC for lead, iron, copper?

Pipe material

Selection of Corrosion Control Approach

Approach is site specific Performance Chemical availability Chemicals currently used on site Operability (ease of use, reliability, maintenance, staffing, training) Current treatment process Cost (capital, chemical and O&M)

Chemical	Dosage (mg/L)	Chemical Cost* \$/1000 m ³	O&M Cost** \$/1000 m ³
Caustic Soda NaOH	2 to 30	0.70 – 10.50	0.15 – 0.73
Lime Ca(OH) ₂	2 to 30	0.45 – 6.75	0.38 – 1.39
Soda Ash Na ₂ CO ₃	10 to 30	4.70 – 14.10	0.38 – 1.39
Carbon Dioxide (CO ₂)	5 to 20	2.80 – 14.00	N/A
Blended Orthophosphate	0.3 to 1.3	6.00 – 25.00	0.15 – 0.73
Sodium Silicate	4 to 10	10.00 – 25.00	0.15 – 0.73

** From Lead Control Strategies, AWWARF, 1990 – adjusted to CAN\$ and 2004 prices

Selection of Corrosion Control Approach

Approach is site specific (continued)

- Implementation (ease, time, public opinion)
- Secondary impacts (disinfection efficiency and maintenance of a residual in distribution system, THM levels, wastewater processing capabilities)



Conclusions

You should care about corrosion Silent pipe killer - extend the life of distribution system Maintain water quality - health and aesthetic Various tools can be used to measure the corrosivity of your water Various methods can be implemented to control corrosion - site specific Corrosion control costs \$ but you will save much more \$\$\$ in the long run



Local Example of Corrosion Control Implementation

Town of Bonnavista Surface water supply from Long Pond Industrial water supply built in 1967 Designed for additional capacity for the Town System started in 1970 Water quality satisfactory with exception of ♦ pH = 6.1 Colour = 30 TCU

Local Example of Corrosion Control Implementation

- Water is pumped from supply main to 2 distribution reservoirs
- Pumping is controlled to avoid conflict with peak industrial demand, i.e. generally 6 p.m. to 6 a.m.
- In early 1980's, leaks appeared as a result of corrosion in copper pipes
- Initially, recommendation to monitor the occurrences of corrosion

Local Example of Corrosion Control Implementation

In early 1990's, start of complaints of staining of plumbing fixtures, discoloration of laundry and frequent leaks in copper service lines
In 1993, water analyses were carried out
Source water was unchanged
Tap water indicated

pH levels were 4.7 to 5.7
Iron levels were 0.47 to 1.2 mg/L

Copper levels were 4.7 to 6.7 mg/L

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Local Example of Corrosion Control Implementation

In 1996, year long source sampling program
pH = 5.78 to 6.51
Colour = 10 to 40 TCU
LSI = -4.3 to -5.1
Recommendations
Add hydrated lime to increase pH and alkalinity and decrease corrosiveness



Local Example of Corrosion Control Implementation

- Design carried out in 1997
- Tenders called in 1998
- System installed and operational in late 1998
- System operated for 2 years using hydrated lime
- Due to operational problems, Town switched to soda ash in late 2000
- Operating range for system

PH 6.8 to 7.0

Local Example of Corrosion Control Implementation

Operating cost of system = \$12,000/year

 Operating and maintenance expenditures works department

1994 to 1998 - \$295,220/year

1999 to 2003 - \$248,120/year

Net saving - \$47,100/year

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Water main leaks due to corrosion

1994 to 1998 - 83 leaks/year

4 1999 to 2003 - 17 leaks/year

Questions and Answers





Given - Water Plant A
Temperature = 18°C
pH = 8.0
TDS = 64 mg/L
Calcium = 42 mg/L as CaCO₃
Alkalinity = 47 mg/L as CaCO₃

Given - Water Plant A ♦ Temperature = 18°C ♦ pH = 8.0 TDS = 64 mg/L \diamond Calcium = 42 mg/L as CaCO₃ • Alkalinity = 47 mg/L as $CaCO_3$ Solution \bullet LSI = pH - pH_s \bullet pH_s = (9.3 + A + B) - (C + D)



♦ Solution (continued)
★ A = (Log₁₀ [TDS in mg/L] - 1) /10
= (Log₁₀ [64] - 1) /10
= 0.08



Solution (continued)
A = (Log₁₀ [TDS in mg/L] - 1) /10 = (Log₁₀ [64] - 1) /10 = 0.08
B = -13.12 x Log₁₀ (Temp. in °C + 273) + 34.55 = -13.12 x Log₁₀ (18 + 273) + 34.55 = 2.22



♦ Solution (continued)
♦ C = Log₁₀ [Ca²⁺ in mg/L as CaCO₃] - 0.4
= Log₁₀ [42] - 0.4
= <u>1.22</u>



Solution (continued)

C = Log₁₀ [Ca²⁺ in mg/L as CaCO₃] - 0.4
Log₁₀ [42] - 0.4
1.22
D = Log₁₀ [Alkalinity in mg/L as CaCO₃]
Log₁₀ [47]
1.67



Solution (continued) • $C = Log_{10} [Ca^{2+} in mg/L as CaCO_3] - 0.4$ $= Log_{10} [42] - 0.4$ = 1.22 \bullet D = Log₁₀ [Alkalinity in mg/L as CaCO₃] $= Log_{10} [47]$ = 1.67 $PH_{s} = (9.3 + A + B) - (C + D)$



Solution (continued)
 pH_s = (9.3 + A + B) - (C + D)
 = (9.3 + 0.08 + 2.22) - (1.22 + 1.67)
 = 8.71



Solution (continued)
pH_s = (9.3 + A + B) - (C + D)

= (9.3 + 0.08 + 2.22) - (1.22 + 1.67)
= 8.71

LSI = pH - pH_s

= 8.0 - 8.71
= -0.71



Solution (continued)

◇ pH_s = (9.3 + A + B) - (C + D)
= (9.3 + 0.08 + 2.22) - (1.22 + 1.67)
= 8.71
◇ LSI = pH - pH_s
= 8.0 - 8.71

= <u>-0.71</u>

Water is still slightly corrosive

- pH will not easily fluctuate once in distribution system (good alkalinity level in water)
- a slight tendency to remove metals from pipes

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LSI Calculation

If calcium hardness is not known, assume calcium hardness = total hardness

 If total dissolved solids (TDS) is unknown but conductivity is known, estimate TDS with table shown on next page

 Conductivity is easily measured using a lab pH analyzer



Conductivity vs TDS

Conductivity	TDS	
(µmho/cm)	(mg/L as CaCO ₃)	
1	0.42	
10.6	4.2	
21.2	8.5	
42.4	17.0	
63.7	25.5	
84.8	34.0	
106.0	42.5	
127.3	51.0	
148.5	59.5	

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