

#### Diagnosing and Solving Corrosion Problems in Jordanian Industry (Case Study)

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By

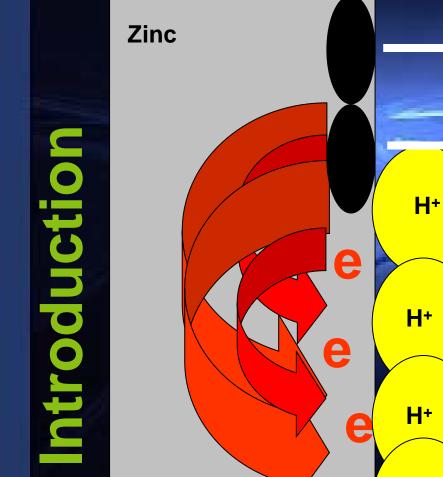
Introduction

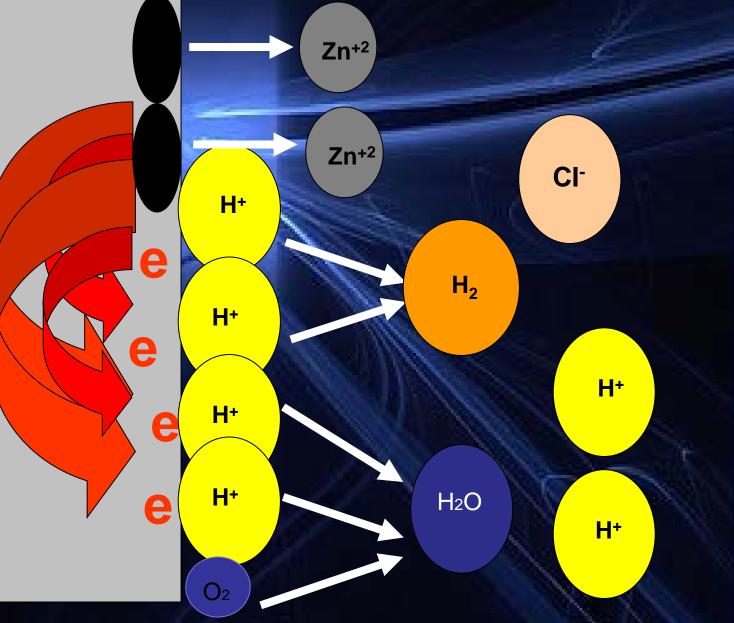
CORROSION is defined as the deterioration of a material because of reaction with its environment



#### **HCI Solution**









**Loosing Mechanical Properties Direct Economical Losses: results** from replacing corroded structures **Indirect Economical Losses:** shutdown, loss of product, loss

of efficiency, over design

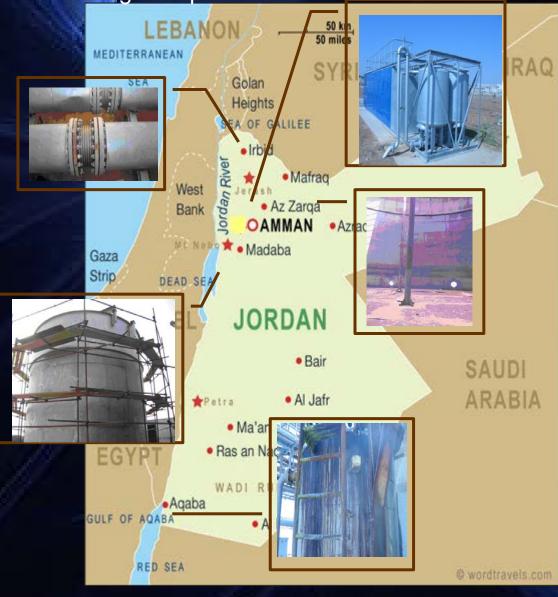
Appearance

Contamination of Products Loss of Safety



ntroduction

After accomplishing more than 60 consultancy studies to solve corrosion problems, It is obvious that many industries in Jordan are facing this phenomena :





## Introduction











#### Bhopal Disaster / on Carbide India Limited (UCIL)



INDIA, 1984, 17000 Died, 558,125 permanent Injuries

water entered a side pipe that was missing its slip-blind plate and entered Tank E610, which contained 42 metric tons of MIC (<u>methyl</u> isocyanate). A <u>runaway reaction</u> started, which was accelerated by contaminants, high temperatures and other factors. The reaction was sped up by the presence of iron from corroding non-stainless steel pipelines. The resulting <u>exothermic reaction</u> increased the temperature inside the tank to over 200 °C (392 °F) and raised the pressure.

This forced the emergency venting of pressure from the MIC holding tank, releasing a large volume of toxic gases. About 30 metric tons of MIC escaped from the tank into the atmosphere in 45 to 60 minutes



The secret of effective engineering generally lies in controlling the corrosion process through the following:

How to recognize the form of corrosion



How to measure its severity



- Inspection techniques
- Methods of prevention

HIGH INVESTMENT IN MATERIAL = LOW MAINTENACE COST

LOW INVESTMENT IN MATERIAL = HIGH MAINTENACE COST



0 **Orros** 

**General Corrosion Galvanic** Corrosion **Intergranular** Corrosion **Crevice** Corrosion **Selective Leaching Pitting Corrosion Stress corrosion cracking Erosion Corrosion** 



#### **General Corrosion**

Uniform attack occurs with equivalent intensity over the entire exposed surface.



#### **General Corrosion**



#### **Galvanic** Corrosion

Occurs when two metals or alloys having different compositions are electrically coupled while exposed to an electrolyte.

Aluminum helicopter blade has corroded because of galvanic or bimetallic action where it was in contact with a steel counterbalance



**Vpes of Corrosion** 



#### Intergrannular Corrosion

of **Corrosion** vpes

Occurs along grain boundaries for some alloys and in specific environment .Some stainless steels when heated to temperatures between 500 – 800 C for sufficiently long time periods ,these alloys become sensitized to intergrannular attack. It is believed that this heat treatment permits the formation of small precipitate particles of chromium carbide by reaction between the chromium and carbon in the stainless steel .Both the chromium and the carbon must diffuse to the grain boundaries to form the precipitates ,which leave a chromium depleted zone adjacent to the grain boundary. Consequently ,this grain boundary region is now highly susceptible to corrosion



#### **Crevice** Corrosion

Occurs as a consequence of concentration differences of ions in the electrolyte solution.



**Crevice Corrosion** 



#### **Selective Leaching**

Occurs when one element is preferentially removed as a consequence of corrosion processes.

Many brass plumbing fittings have electrochemical

cells between their constituent phases. This leads to rapid corrosion cracks which have a coppery color due to dezincification where the zinc content is leached out.





**Vpes of Corrosion** 

#### **Pitting Corrosion**

It is a form of localized corrosion attack in which small pits or holes form. They penetrate from the top of a horizontal surface downward in a nearly vertical direction.



#### **Pitting Corrosion**



#### Stress corrosion cracking

Results from the combined action of an applied tensile stress and a corrosive environment.

The deep drawn brass cup on the right shows stress corrosion cracking under the influence of the residual manufacturing stresses and a mildly corrosive environment. The cup on the left has been annealed before putting it into service which solves the problem.



#### **Erosion Corrosion**

Arises from combined action of chemical attack and mechanical abrasion or wear as a consequence of fluid motion.



**Erosion Corrosion** 



### Miscellaneous

### Case Studies



**Case Studies** 

## First Case Study



Failure Due to Corrosion Phenomenon in The Fire Tubes of The Main Boilers

**Case Studies** 



Verifying the root causes of corrosion failure in the fire tubes was an ultimate goal of this study:



Fig.(1) : Fire tubes of the boilers



# **Case Studies**

Fig.(2) : Corroded fire tube with full deterioration in the material as shown in the red arrows



Therefore a thorough investigation was conducted on the following:

**Studies** ase

1.Corroded fire tubes2.Corrosion products and ash3.Diesel fuel4.Water before and after treatment



#### **Corroded fire tubes**

**Case Studies** 

**Chemical analysis** of the corroded fire tubes' material showed that it is Carbon Steel grade AISI 1010 with C % = 0.083 and S % = 0.0142.

Mechanical tests on the failed tube showed the following : Yield strength = 245 MPa Tensile Strength = 341 MPa Elongation = 45%

**Chemical analysis using EDX** of the corrosion products for several metallic samples revealed the following:



#### Table (1) Sample #1 (row # 1, tube # 8), position #1 from boiler #3

	Wt%
Compound	
FeO	42.91
SO <sub>3</sub>	47.23
	n a sitism #0 frame h silen #0
Table (2) Sample #1(row # 1, tube # 8),	position #2 from boller #3
	Wt%
Compound	
FeO	49.29
SO <sub>3</sub>	50.71
Table (3) Sample #2(row #5, tube #7), p	osition #1 from boiler #3
	Wt%
Compound	
FeO	68.26
SO3	28.21
the second se	ALL ALL ALL AND
Table (4) Sample #2(row #5, tube #7), p	osition #2 from boiler #3
	Wt%
Compound	
FeO	47.72
SO3	48.70
-	



#### Table (5) Sample #3(row #1, tube #4), position #1 from boiler #3

	Wt%
Compound	
FeO	42.88
SO <sub>3</sub>	55.98

#### Table (6) Sample #3(row #1, tube #4), position #2 from boiler #3

	Wt%
Compound	
FeO	47.56
SO3	49.98





Fig.(3): (a) Part of the failed fire tube showing the corrosion products on the internal surface as shown in the red arrow
(b) Corrosion products near the opening of the fire tube external surface located at the rear side of one of the boilers.



3

4

**Micro-hardness analysis** was conducted on random selected spots on the failed fire tube surface given in fig.(3a) and near a deep pit adjacent to seam weld line as shown in the red circles of fig.(5) and table (7) :

Fig.(5) : Part of the failed fire tube surface magnified using optical microscope (200 X ) showing the random selected spots to measure the microhardness near a shallow and deep pit.



Table (7) : Measured Vickers Micro hardnesson the random selected spots on the failed firetube surface

Position No.	Measured Vickers Hardness
1	118.656
2	145.83
3	82.4
4	126.6



#### **Microstructural analysis** of several specimens revealed the existence of deep and large "WORM HOLES" pits.

case Studies

Fig.(6) : Clear pits propagated into large "WORM HOLES" as shown in the red arrows





Fig.(7) : Clear pits propogated into large "WORM HOLE" as shown in the red arrow



#### Non Destructive Tests (NDT) :



Fig.(8) : Reduction in thickness in the lower part of the fire tube as shown in the red arrow



**Case Studies** 



Fig.(9) : Reduction in thickness in the lower part of the fire tube as shown in the red arrow



**Fig.**(10) : **Reduction in thickness in the lower part of the fire tube with complete deterioration in the material as shown in the red arrow.** 



**X-Ray** test on one of the failed tubes revealed heavy damage due to the corrosion phenomena on the internal surface of the tube with severe reduction in thickness as shown in the following figs.(-11 & 12):



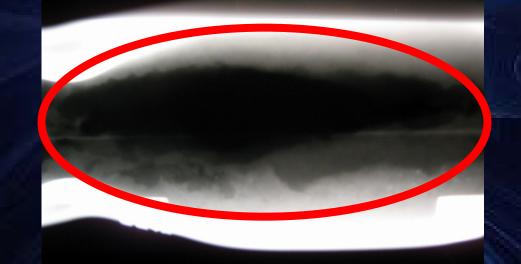


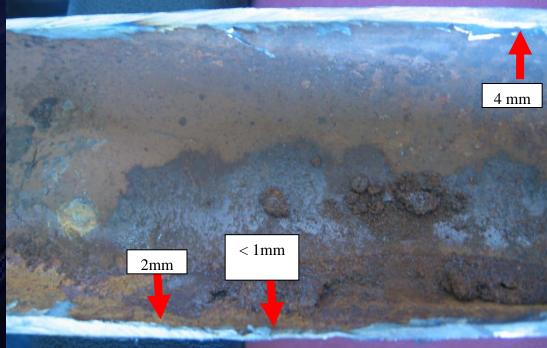
Fig.(11) : X –Ray photo of the failed tube revealed heavy damage due to the corrosion phenomena on the internal surface of the tube with severe reduction in thickness as shown in the red circle.



Fig.(12) : X –Ray photo of the failed tube revealed heavy damage due to the corrosion phenomena on the internal surface of the tube with severe reduction in thickness as shown in the red circle.



In order to measure the actual reduction in thickness of the fire tube given in figs.( 11 and 12) , longitudinal cutting was implemented as shown in fig.(13) .Thickness measuments results of the fire tube given in in fig.(13) deviated between ( <1 mm in the lower part of the tube and 4 mm in the upper part of the tube).



**Fig.(13) : Red arrows reveals severe reduction in thickness in** the lower part in comparison with the upper part.



# **Corrosion products and ash**

Chemical analysis of the corrosion products and ash fetched from four random positions located in boiler # 3 revealed the following as shown in the following table:

Wt.%	<b>S1</b>	<b>S2</b>	<b>S3</b>	<b>S4</b>
Moisture Content	1.4	4.43	3.31	0.6
sulfur content	15.18	13.04	14.53	0.3
Iron content	16.5	17.5	13.9	33.3

# Diesel fuel

Chemical analysis of the diesel fuel samples from main storage tank and the burner revealed higher sulfur content in ppm the following as shown in the following

table:

ррт	Main Storage Tank	Burner
Total Sulfur Content	10000	9780



## Water before and after treatment

Water analysis before and after treatment revealed the following:

**Case Studies** 

Parameter	Water before treatment	Water after treatment
рН	8.28	9.28
TH (as CaCO <sub>3</sub> ) mg/L	132	8.83
Cl mg/L	159	129
SO <sub>3</sub> mg/L	<2	<2
TDS mg/L	582	476
Iron content mg/L	<0.039	<0.039



### **Conclusion:**

The above results showed the following :

**Mechanical tests** (Yield Strength (YS) = 245MPa, Tensile Strength(TS)=341MPa and Elongation (E)= 45 % which showed a deviation from standards for carbon steel AISI 1010 (i.e. YS = 305 MPa, TS = 365MPa and E = 20%) due to the corrosion phenomena.

**EDX surface analysis** revealed the existence of high percentages of SO<sub>3</sub> which deviated between (28.21 to 55.98 %) in combination with high percentages of iron oxide which deviated (42.88 to 68.26%) for various fire tube samples. **Micro hardness test results revealed** an obvious increase in the hardness at the points near the pits (i.e.118.656, 145.83& 126.6 Hv) in comparison with the points far from the pits (i.e. 82 Hv).

**Microstructural analysis revealed** the existence of deep and large pits propagated into "WORM HOLES".

Ultrasonic and X-ray tests revealed heavy damage due to the corrosion phenomena on the internal surface of the fire tubes with severe reduction in thickness.
 Thickness measurement revealed a deviation between ( <1 mm to 4 mm) due to the corrosion phenomena.</li>

**Chemical analysis of the corrosion products** and ash showed percentages of sulfur deviated between (0.3 to 15.18 %) and moisture content deviated between (0.6 to 4.43 %).

**Diesel fuel analysis showed** extremely high percentages of sulfur content deviated between (9780 to 10000 ppm).



So it is obvious that the scenario of the failure in the fire tubes of the three main boilers occurred due the existence of large quantities of sulfur in the diesel forming sulfurous gases (i.e.  $SO_2$  and  $SO_3$ ), which as a result  $SO_3$  reacts with moisture formed from the condensation of air near the outlet of the fire tubes (i.e. 20 cm before the outlet) due to deviation in temperature between (water side and fire side across the fire tube wall) causing the formation of diluted sulphuric acid (i.e.  $H_2SO_4$ ) according to the following chemical reactions and the sketches :

 $S + O_2 \rightarrow SO_2$  $SO_{2} + 1/2O_{2} \rightarrow SO_{3}$  $SO_3 + H_2O \rightarrow H_2SO_4$ 



## Flue gases (i.e. SO<sub>2</sub>, CO<sub>2</sub>)

 $SO_3$  reacts with moisture content forming diluted sulphuric acid  $(H_2SO_4)$ 

Forming a pit that propagates into worm holes causing full deterioration of the material



Because this grade of steel (AISI 1010) doesn't resist sulphuric acid, therefore oxidation of iron occurred causing the initiation of deep and large pits which propagated into WORM HOLES and as a result thinning of the wall or full deterioration in the material of the fire tubes occurred. On the otherhand the accumulation of corrosion products and ash with high percentages of sulfur compounds will cause the initiation of what is called localized concentration cells which plays also an additional role in increasing the aggressiveness of the corrosion attack.



### Recommendations

Replacement of the material of the existing fire tubes (i.e. AISI 1010) with new material made of iron base alloy containing Chromium (Cr) and Molybdenum (Mo) in order to resist the saturated sulfur and sulfur dioxide (SO<sub>2</sub>) gaseous environment and other compound formed as a secondary reaction ( i.e. Dilute  $H_2SO_4$ ).

Monitoring the percentage of sulfur content in the diesel fuel .It should be with minimal.

The boilers should be cleaned three times a year at least in order to remove any materials that might be generated at the internal surface of the fire tubes within the corregations. Taking into consideration that improper cleaning will reduce the lifetime of the fire tubes and the boiler as a result due to corrosion phenomenon as well as increasing the consumption of fuel.

Regulating and monitoring the fuel- to- air mixture ratios in the burner by an expert because it has a direct effect on the efficiency of combustion in order not to have partial combustion conditions that generate combustion products which might generate localized concentration cells and initiating the corrosion phenomenon. Therefore it is worthy of mention in this regard that the maintenance technician shouldn't build his judgment on the efficiency of the burner just by visual evaluation of exhaust smoke and the flame.



There should be a log book for each boiler in order to have a historical record that reflects both normal and abnormal operating condition as well as shut downs, periods of preventive maintenance and what kind of notifications were recorded during maintenance or during overhaul. In this regard a preventive maintenance management software's could be utilized in an efficient way.

Having special devices to monitor the water quality (i.e. digital pH meter, TDS meter , Oxygen meter and TH meter ) in order to have a snap shot on the conditions of the water before and after treatment. Because water analysis conducted at RSS laboratories showed an obvious deviation from the analysis conducted by the water treatment company .On the other hand the type of inhibitors used by the water treatment company are working as anti scale and anti corrosion (i.e. Oxygen scavenger) inhibitors, and it has no effect against the formation of biological corrosion due to aerobic and anaerobic Bacteria as well as Algae. Therefore biocide should be added to the scale and corrosion inhibitors.



# Second Case Study





The following study was conducted on the four Stainless Steel tanks (Water, Concentrate 1, Concentrate 2 & Syrup) as well as on a sample of the stainless steel piping network.

# **1.Stainless Steel tanks (Water, Concentrate 1, Concentrate 2 & Syrup):**

Fig.(1) represents the external surface of the water tank which showed an extensive corrosion phenomena underneath the surface of the external stainless steel sheets as shown in fig.(2).This phenomena was also obvious in the other three tanks (Concentrate 1, Concentrate 2 & Syrup).



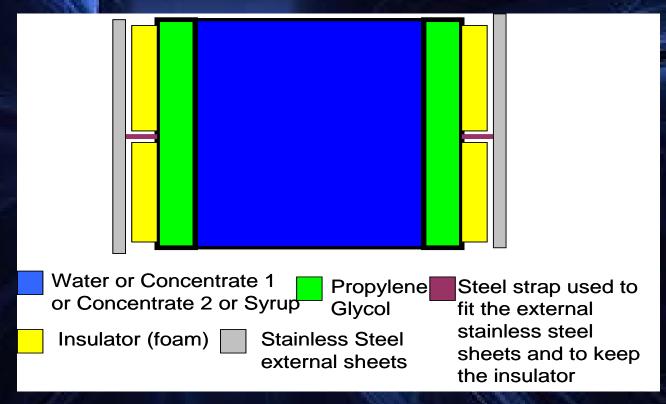
Fig.(1) : External surface of the water tank.





Fig.(2) : Corrosion products are very obvious below the external surface of the stainless steel sheets which appeared in the four above mentioned tanks.





# Fig.(3) : Crrossectional area of the tanks



The following table (1) shows the chemical analysis by EDX of the material of each species used to construct the tanks, which revealed that the type of stainless steel was of grade 304.The straps were of carbon steel.

# Table (1) Type of the material used to construct the tanks

No.	Туре	Material
T Sapt	Inner surface of the tank attached to the solution (Water,Conc.1,Conc.2 & Syrup ) from one side and the propylene glycol from the other side	Stainless Steel 304
2	Outer surface of the tank attached to the propylene glycol from one side and the insulator from the other side	Stainless Steel 304
3	External Sheets	Stainless Steel 304
4	Straps	Carbon steel
5	Welds	Stainless Steel 304



Chemical analysis of the Propylene Glycol revealed that the pH was 7.1 ,while biological analysis revealed traces ( < 1 CFU/ml) of aerobic and anaerobic bacteria which have no significant effect from corrosion stand point of view on the material of the tank.

- Chemical analysis of the Water showed that the pH was equal to 3.05 with TDS equal to 39 mg/L and TH as  $(CaCO_3)$  equal to 9 mg/L.

-Chemical analysis of the Syrup and the Concentrate revealed pH values of (4.3 & 4.4) respectively.

-Corrosion tests were conducted using the cell given in fig.(4) in order to implement accelerated corrosion tests on the metallic specimens in various environments (Water , Conc.1,Conc.2 & Syrup) using the Potentiostat given in fig.(5) .Tests results shown in table (2) below.





# Fig.(4) : Corrosion Cell



Fig.(5) : Potentiostat

# **Case Studies**



# Table (2) Corrosion rate results using potentiostatic test

No.	Working Electrode	Electrolyte	Time hour	Corrosion Rate (mm/y)	Comments
197	Inner surface material (SS 304 ) of the Syrup tank 1	Syrup	ALL ALL	3.6 x 10 <sup>-3</sup>	Very good corrosion resistance to the electrolyte
2	Inner surface material (SS 304 ) of the Syrup tank 2	Syrup		3.0 x 10 <sup>-2</sup>	Good corrosion resistance to the electrolyte
3	Inner surface material (SS 304 ) of the Water tank	Water	A NEW	3.323 x 10 <sup>-3</sup>	Very good corrosion resistance to the electrolyte
4	Inner surface material (SS 304 ) of the Concentrate tank	Concentrate		1.335 x 10 <sup>-2</sup>	Good corrosion resistance to the electrolyte
3	External sheets (SS 304)	Water		5.396 x 10 <sup>-3</sup>	Very good corrosion resistance to the electrolyte

**Studies** Jase



The mechanism given in fig.(6) was verified using the Galvanic Corrosion Monitoring & Analyzing System which indicated that the carbon steel straps shown in figs.(2 & 3) were galvanically corroded due to the coupling with stainless steel sheets (SS 304) and stainless steel (SS 304) of the outer surface of the jacketed tank using stainless steel welding rods (SS 304). This was obvious in fig.(7) which showed the graphs of (E1 (stainless steel sheet) and E3 (stainless steel outer surface of the jacketed tank) with negative galvanic currents against time (1 hour), while graph (E2 (carbon steel strap)) revealed positive galvanic current against time (1 hour ) which means that it was sacrificially corroded in order to protect E1&E3.



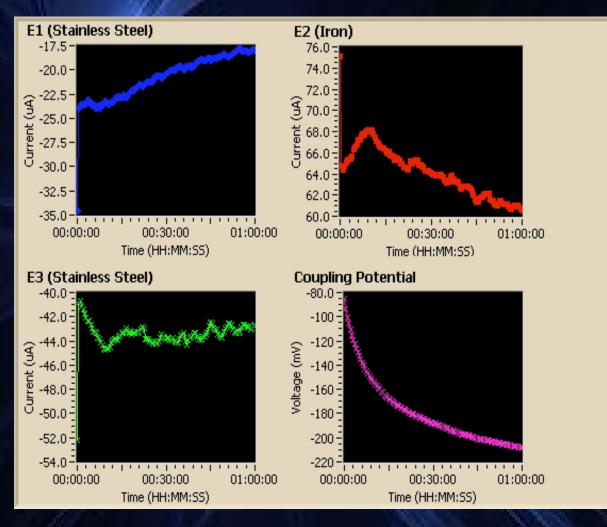


Fig.(7) :Galvanic corrosion result due to the coupling of stainless steel sheets (SS 304), stainless steel (SS304)of the outer surface of the jacketed tank and carbon steel straps.

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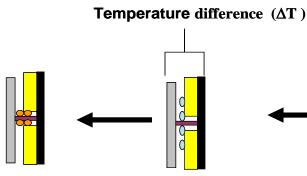


Studies

**SSE** 

Corrosion rates given in table (2) deviated between (Good to excellent) corrosion resistance to the environment at stagnant conditions.

But what actually caused the corrosion to occur beneath the stainless steel sheets shown in fig.(2) is the mechanism shown in fig.(6).



Due to the temperature difference between the ambient temperature and the coolant (propylene glycol) temperature, Condensation process will occur followed by initiation of water droplets over the T connection among the carbon steel straps, stainless steel sheets and stainless steel outer surface of the jacketed tank. Water droplets work as an electrolyte to conduct GALVANIC current and causing Galvanic Corrosion to occur

Insulator (foam)

Stainless Steel external sheets

Corrosion Products

) Water droplet

Steel strap used to fit the external stainless steel sheets and to keep the insulator Fig.(6) :sketch shows the major cause of corrosion beneath the stainless steel

sheets



# Third Case

Study

**Case Studies** 



This study is based on verifying the causes of cracking due to corrosion in the expansion joints shown in fig.(1) below:



Fig.(1): Expansion joints



Chemical analysis of the of the expansion joint material sample showed the following:

Table(1):Chemical analysis of the expansion joint material

Element	Chemical Composition, %
The state of the states	The state of the state
Carbon	0.04
Sulfur	0.006
Silicon	1.042
Chromium	17.464
Nickle	10.112
Manganese	1.548
Iron	69.834

Studies **Jase** 



Chemical composition given in table(1) showes that the expansion joint material of Stainless Steel type 304 according to the ASTM (A666) as follows:

Element	Chemical Composition, %
Carbon max.	0.08
Sulfur max.	0.03
Silicon max.	0.75
Chromium	18.00-20.00
Nickel max.	8.00-10.50
Manganese max.	2.00



Cutting the sample of the expansion joint given in fig.(2) showed multiple layers of stainless steel sheets compressed together as shown in fig.(3).



Fig.(2):Sample of expansion joint





Fig.(3):Sample of expansion joint showing the stainless steel 304 layers compressed together.



Analyzing the chemical composition using SEM of the corrosion products given in fig.(4) showed the following:

Table(2):Chemical analysis of the corrosion products given in fig.(4)

Corrosion Product	Wt%
Cr <sub>2</sub> O <sub>3</sub>	22.86
FeO	61.071
SO <sub>3</sub>	0.486
Cl	0.264





Fig.(4):Sample of one of the corroded layers in the expansion joint given in fig.(3).

**Case Studies** 



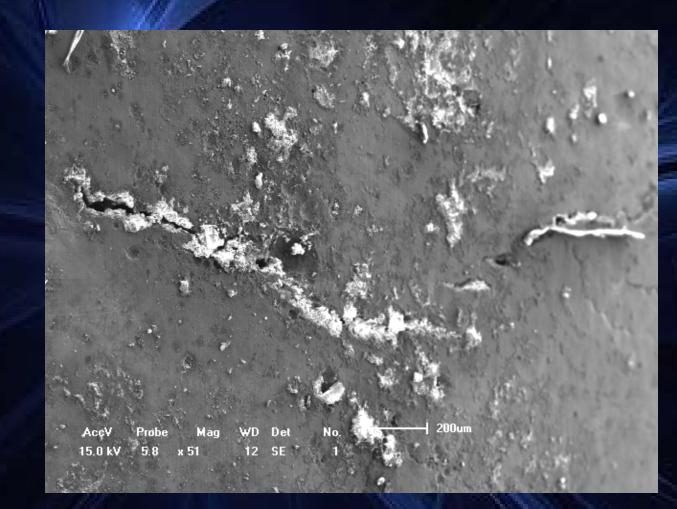


Fig.(5):SEM photo of one of the corroded layers in the expansion joint showing a crack occurred due to the Stress Corrosion Cracking (SCC).



Analyzing the chemical composition using SEM of the corrosion products in the cracked area given in fig.(5) showed the following: Table(3):Chemical analysis of the

corrosion products given in fig.(4)

Corrosion Product	Wt%
Cr <sub>2</sub> O <sub>3</sub>	15.983
FeO	45.13
SO <sub>3</sub>	3.782
Cl	2.943

Tables (2&3) showed percentages of  $(SO_3 \& CI)$ which give the evidence that these elements came from the surrounding environment and Acidic Rain which is mainly diluted sulphuric acid  $(H_2SO_4)$ .



These percentages of  $(SO_3 \& Cl)$ , played an important role in the initiation of what is called "Concentration Cells" after a normal evaporation process, adding to it the thermal stresses which in combination with the concentration cells started the corrosion phenomena causing a crack in what is called the Stress Corrosion Cracking (SCC) in the first layer of the expansion joint, taking into consideration that the material of the expansion joint given in table(1) doesn't resist corrosion, which opened the path for the acidic rain to enter to the second layer and initiating again the "Concentration Cells" repeating what is happened in the first layer, then the same process will occur to the third ,fourth etc...Fig.(6) give a schematic hypothetical assumption of the given process process



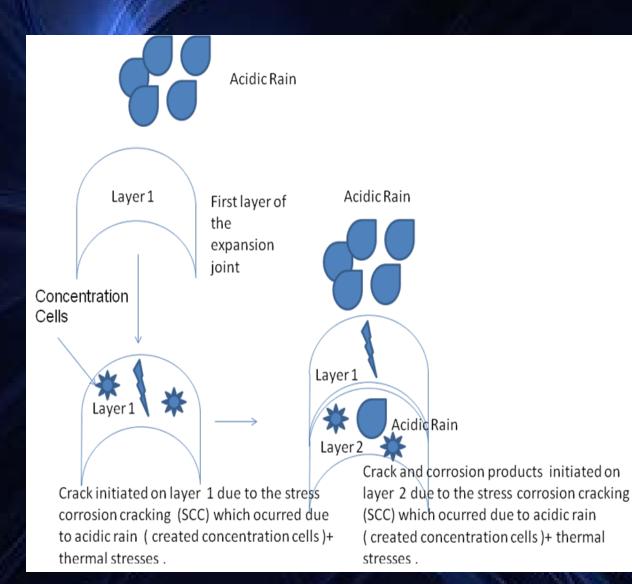


Fig.(3):Schematic representation to what have happened to the expansion joint.



# **Case Studies**

# Fourth Case Study



# Fig.(1) below represents the steam drum which showed pitting corrosion on the internal suface.



# Fig.(1) : Steam drum



**Case Studies** 

Standard replica procedure according to ASTM E 1351 was used to obtain metallurgical information from selected locations (1,2 &3) at the internal surface of the steam drum. The replication procedure involved initial surface preparation by grinding discs. The grinded areas were polished successively with Alumina followed by diamond polishing pastes(30,12 & 1 micrometer). Microstructural features were revealed by etching the surface with Nital (3% Nitric acid in 97%) Ethanol).Cellulose acetate film moistened with acetone was used to replicate the microstructure. At least four replicas were taken from each position to ensure optimum results. Figs (2,3,4, 5,6&7) show large pits of different dimensions obtained from replicas.



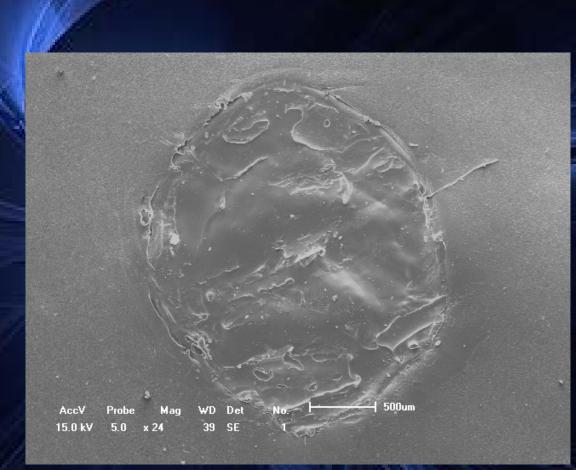
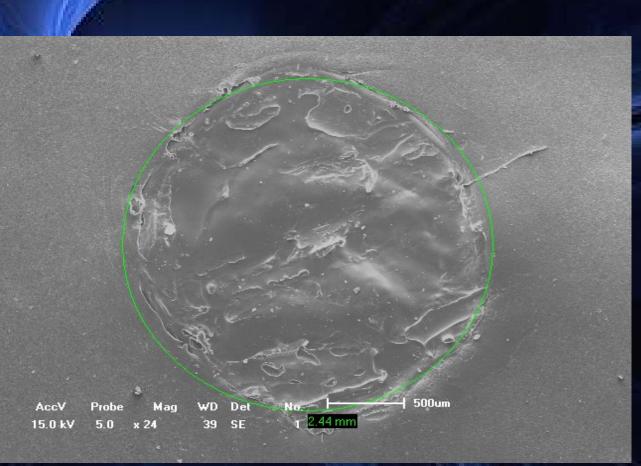


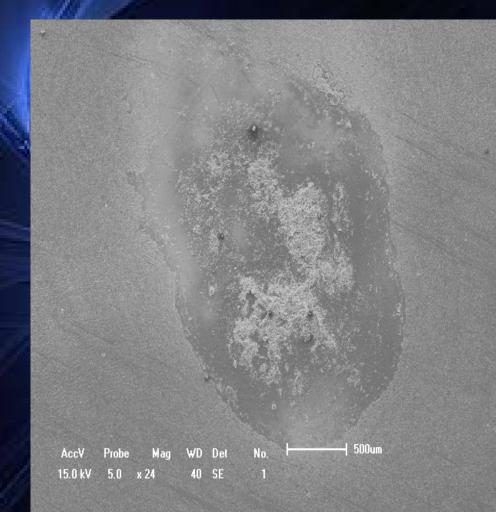
Fig.(2) SEM photo shows a large pit in location 1 of the internal surface of the steam drum





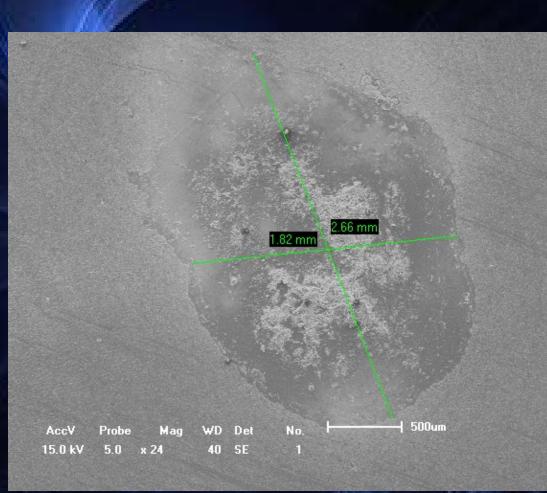
#### Fig.(3) SEM photo shows the diameter of the pit in location 1





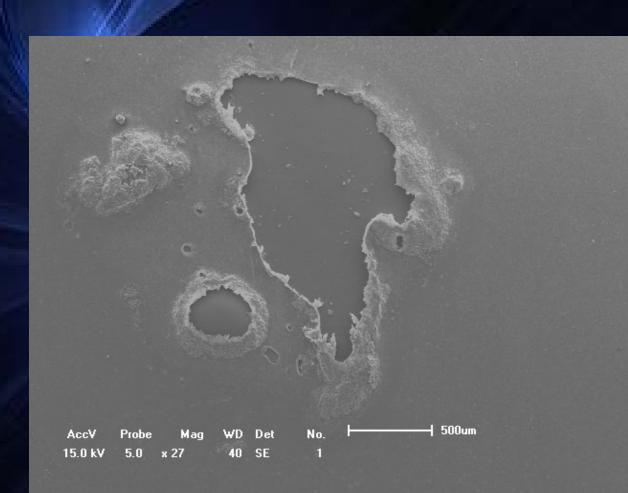
#### Fig.(4) SEM photo shows a large pit in location 2 of the internal surface of the steam drum.





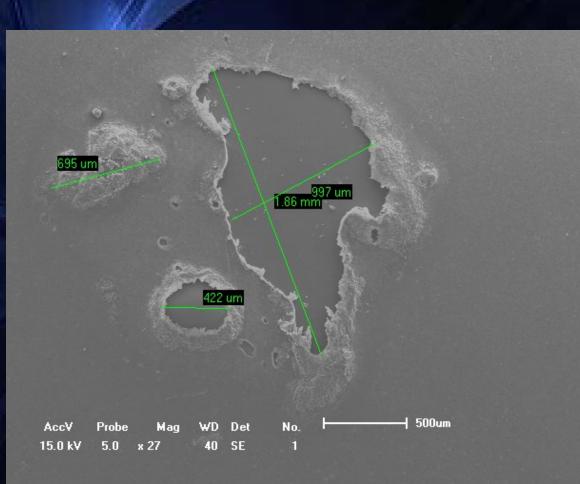
#### Fig.(5) SEM photo shows the dimensions of the pit in location 2





#### Fig.(6) SEM photo shows the dimensions of the pit in location 3





#### Fig.(7) SEM photo shows the dimensions of the pit in location 3



#### <u>Conclusion</u>

Studies Jase

It can be concluded from the above figs( 2 to 7) that phosphate induced corrosion as well as formation of Caustic Soda have initiated the pits shown in the above figures according to following reaction mechanism :

 $Na_3PO_4 + H_2O \rightarrow Na_2HPO_4 + NaOH$ 

 $2Na_2HPO_4 + Fe + \frac{1}{2}O_2 \rightarrow NaFePO_4 + Na_3PO_4 + H_2O$ 

 $2NaOH + Fe \rightarrow Na_2FeO_2 + H_2$ 





Studies

Case

It is clear from the above chemical reactions that the formation of maricite (i.e. NaFePO<sub>4</sub>) is due to the attack of the acidic phosphate salt (i.e. NaHPO<sub>4</sub>) to the internal surface of the steam drum at elevated temperature (i.e.250 C), because phosphate is known to affect the crystalline structure of the steel structures. Formation of Caustic soda (NaOH) causes Caustic corrosion (gouging) which results from the reaction of caustic soda with the internal surface of the steam drum. In order to minimize this phenomena continuous monitoring of the PO4 concentation, pH and Na/PO4 ratio which should remain constant in the steam drum



### Fifth Case

Study



#### In order to verify the root causes of the failure in bearing no.1 of the steam turbine given in fig.(1, 2) below :



Fig.(1) : Steam Turbine of the National Sugar Company.



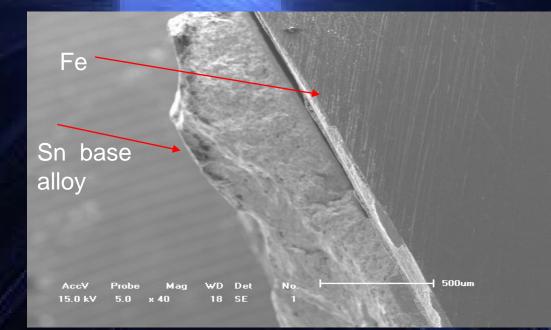


#### Fig.(2) : Part of the failed bearing no.1



A thourogh investigation was conducted on the failed parts of the given bearing as follows:

Chemical analysis using EDX on the part given in fig.(2) showed that the failed part was consisted of two main materials : Iron (Fe = 97.4 wt.%) & Tin base alloy (Sn = 94.81 wt.%) known commercially as babbitt metal as follows :



Fig(3) : Part of the failed bearing no.1 magnified using SEM ( 40 X ) showing the layer of Tin base alloy covering the base metal constructed of Iron (Fe)



Studies

Case

Surface analysis using SEM showed that a layer of Tin base alloy  $\cong$  1834  $\mu$  m in thickness was combined to a base metal made of Iron as mentioned in the above chemical analysis using cladding process without heating and this is clear in fig.( 4 ) in which the layer of Iron is very well distinguished from the layer of Tin base alloy at the metal – metal interface shown in the red elliptical shape or in other word there is no interaction between the materials of the both faces ( i.e. negligible bonding force ) .

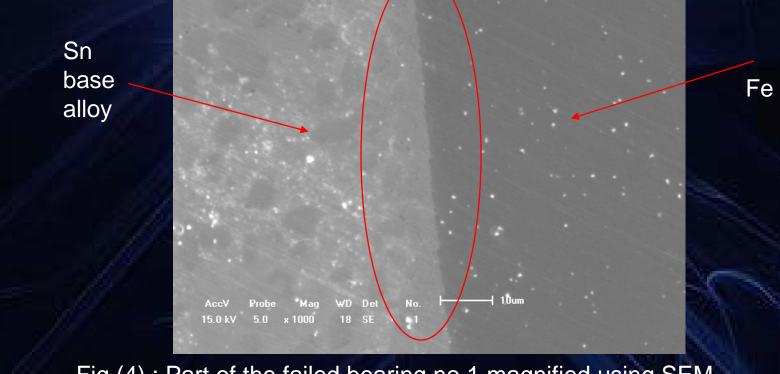


Fig.(4) : Part of the failed bearing no.1 magnified using SEM (1000 X) showing the Iron-Tin base alloy interface.



Hardness analysis for the failed bearing was conducted on random selected spots on the Iron –Tin interface from the Iron side as shown in the red circles in fig.(5) :

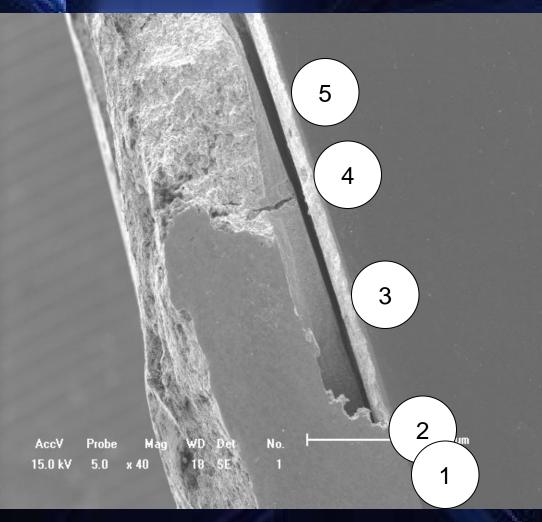


Fig.(5): Selected positions at the metal - metal interface to conduct Hardness testing.



Table (1) : Measured Vickers Hardness on the random selected spots on the Iron –Tin base alloy interface from the Iron side

Position No.	Measured Vickers Hardness
in the first first for a	228
22222	230
3	240
4	242
X245 X24	245



Finite Element Method to determine the maximum and minimum stresses subjected on the failed bearing using VON MISES method revealed the following results given in fig.(6) below:

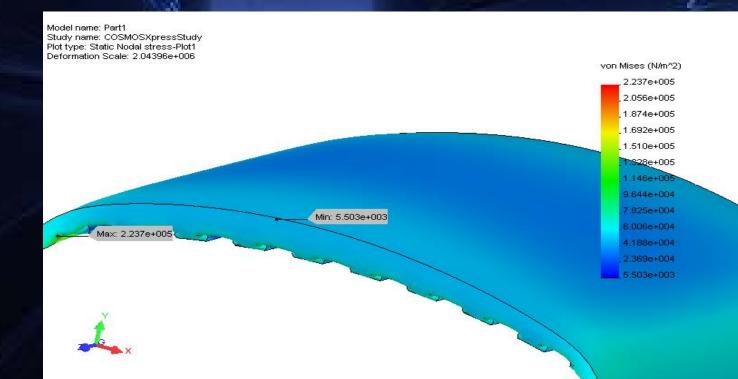


Fig.(6) : Finite Element Method using VON MISES method to determine the stresses subjected on the failed bearing revealed that the maximum stresses were concentrated at the left and right edges of the bearing while minimum stresses were concentrated near the middle of the bearing .

Thickness analysis of the deteriorated Tin base alloy layer in the failed bearing revealed the followings as shown in table (2) below:



Fig.(7) : Part of the failed bearing no.1 showing the random selected spots in order to measure the thickness of the Tin base alloy layer



Table (2): Thickness measurement on the random selected spots of the Tin base alloy layer

Tin layer thickness, μ m
1 4 1 K What I 4 1 K What I
782
to and when a wind when
704
794
1 - 14 TEL - 14 TEL
794
you ster you ster you you
The house in the skill
42.2
The main with main with
100.0
100.0
a Strand Strand
52.3
A BOAR PARA BOAR PARA
622
NEN ENGENEN ENGEN
and the second second second
757
I I HARLIN HARL
735
18 11 135 11 144 S
The A hard



6.Chemical analysis on the used oil supplied by your side to identify the wear metals as follows:

Type of metal	Unit	Result
Fe	ppm	0.12
Sn	ppm	< 0.1

#### **Conclusion:**

The above results showed the following :

A.Surface analysis using SEM showed that the cladding process used to combine the Tin base alloy layer to the base metal (i.e. Iron) was done without heating. In other word this process has left weak points at the metal – metal interface.

B.Hardness results revealed an obvious increase in the hardness at the points near the failure (i.e. 240,242 & 245) in comparison with the points far from the failure (i.e. 228-233).

C.Stress analysis on the failed bearing showed that maximum stresses were concentrated at the left and right edges of the bearing while minimum stresses were concentrated near the middle of the bearing.



D. The lowest thickness measurement of the Tin base alloy layer were recorded near the middle of the bearing (i.e. 42.2,100,52.3  $\mu$ m) while the maximum thickness measurment were recorded near the right and left edges (i.e. 1438,794,757  $\mu$ m). So it is obvious from (A,B,C & D) that the scenario of the failure in the given bearing happened due to the de-attachment of the layer of Tin base alloy at the Middle points of the bearing due to manufacturing insufficiency causing an increase in the friction forces which enabled an accelerated wear and tear process to occur with increasing the temperature of the Tin base alloy layer to the melting point of the alloy ( $\cong$  232 ° C) and as a result the layer dissolved partially at the middle points.



## Sixth Case Study











## Botteom Inside 5 6 7 1 (113 0



#### 9 60 5 6 9 30 1 2 3 4 5 6 8 9 40 1 9 50



### Seventh Case Study





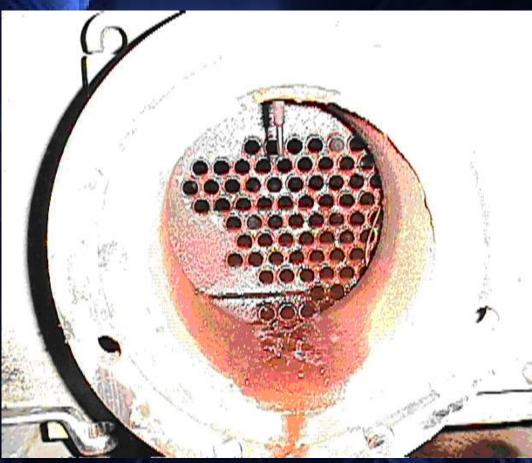










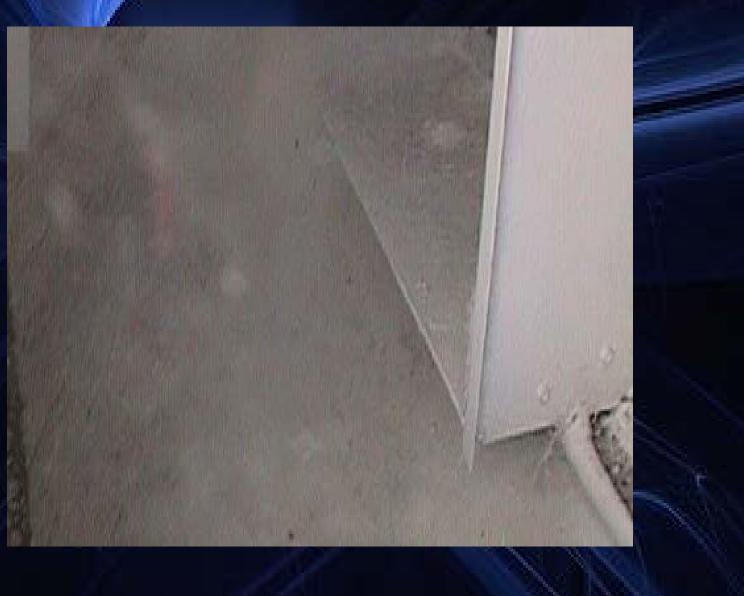












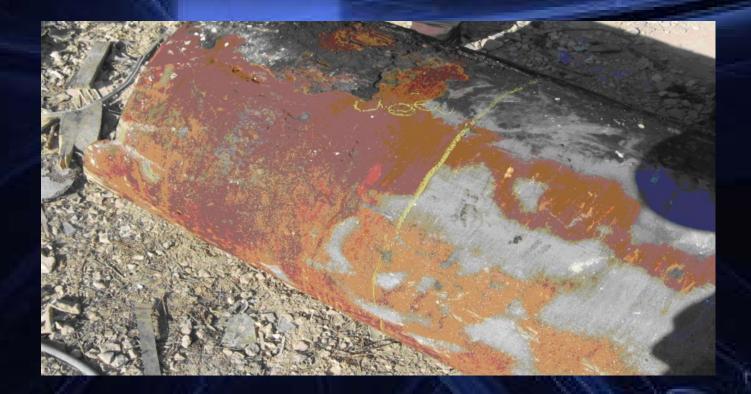


## Eighth Case Study











## NinthCase Study



#### **Biological Corrosion**

Biological corrosion is not a form of corrosion; it is the deterioration of a metal by corrosion processes which occur directly or indirectly as a result of the activity of living organisms. These organisms include microforms such as bacteria and macro types such as algea. Microscopic and macroscopic organisms have been observed to live and reproduce in mediums with pH values between 0 to 11, at temperatures between 30 and 180 F, under pressure up to 15000 lb/in 2.



Aerobic Bacteria: Sulfur oxidizing bacteria, which oxidize elemental sulfur to sulfuric acid 5 % in conc.

2S+3O<sub>2</sub>+2 H2-O ----- 2 H<sub>2</sub>SO<sub>4</sub>

Macrorganisms such as Fungus produces quantities of organic acids such as oxalic,lactic,acetic and citric acids which can initiate crevice attack of metal surfaces.

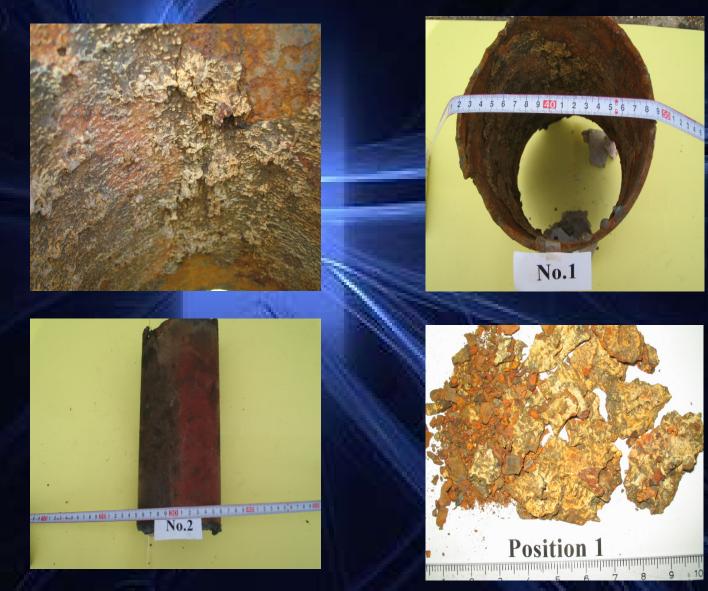
Algae also attach themselves to metallic structures causing corrosion



Living organisms are sustained by chemical reactions. These processes can affect corrosion behavior in the following ways: 1.By directly influencing anodic and cathodic reactions 2.By influencing protective surface films. 3.By creating corrosive conditions. 4.By producing deposits.

These effects may occur singly or in combination ,depending on the environment and the organism involved.





**6inch Pipe** 





Shows the installation of the Corrosion Coupon and Scale Coupon in 12 inch pipeline



#### Corrosion Coupon before exposure







**Corrosion Coupon after exposure for six months** 



Studies Case

SEM photo of the surface of Corrosion Coupon after removing products from the surface

Corrosion rates of the Corrosion Coupon equal to 0.09 mm/y or 3.54 mpy





#### Corrosion Coupon after removing products from the surface



