INVESTIGATION OF PIPELINE FAILURE IN A THERMAL POWER PLANT’S PROCESS WASTEWATER DISTRIBUTION SYSTEM

PREISKAVA POŠKODBE CEVOVODA V SISTEMU ZA OBDELAVO IN DISTRIBUTIJO ODPADNE VODE TERMEOLEKTRARNE

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In this paper, abundant, compacted, dust-like, corrosion scales from the interior of a corrosion-damaged pipeline were analyzed to determine the main reason for leakage and flow damping in the process-wastewater distribution system in the thermal power plant Plomin. The chemical composition, morphology and microbiological activity of the corrosion deposits present on the inner pipe wall were investigated and the physical and chemical analyses of the purified boiler water was made. The results show the presence of iron-related bacteria (IRB) in the corrosion deposits as well as in the distribution system water. XRD analysis shows exclusively magnetite and goethite with no calcium carbonate present in the layers, therefore indicating that no protective carbonate scales, which would protect the pipeline steel, had initially formed. It was concluded that the primary causes of intense corrosion were iron-oxidizing bacteria, that through their metabolism, support the redox cycling process, the formation of large tubercles as well as irregular thinning of the pipeline wall with separated anodic and cathodic areas.

Keywords: pipeline failure, microbiologically influenced corrosion (MIC), process wastewater distribution system, tubercles

1 INTRODUCTION

Process wastewater is generated as a consequence of the technological process of electricity production. Boiler water is part of process wastewater distribution system and is generated in an auxiliary boiler, stone pulverizers, wash residue of the regenerative air heater, condensate from the boiler area, the bilge pit and the pit of the bunker tract. Given that wastewater is very uneven in quality and quantity, with the purpose of a uniform treatment process, the equalization of wastewater takes place at the thermal power plant Plomin buffer pool with a capacity of 1200 m³. Further pH stabilization treatments were performed with the help of lime milk by sifting water from the neutralization tank to the sedimentation tank. Through the central pipeline, the wastewater from the flocculation tank is decanted into the settling tank. The clarified wastewater leaves the precipitator through the overflow gutter and goes to the pH control tank where the pH is adjusted with hydrochloric acid and filtered through a sand filter later. The average inflow of purified wastewater (chemical composition given in Table 2) is approximately 5 m³/h. The purified water from the sand filter is discharged into accumulation tank, 50 m³ volume and through the overflow into the surrounding channel. The main purpose of the accumulation tank is to accumulate treated wastewater for rinsing the droplet separator system during the technological process of desulphurization of the flue gases. The total average daily water needs for rinsing the droplet separator system is 55 m³, which means that rinsing can be done entirely using only accumulated wastewater in the thermal power plant Plomin. During daily operations over a period of 6 years, the treated wastewater recovery system
was showing a continuous decrease in the consumption of treated wastewater and an increase in the consumption of raw water for rinsing the droplet separator. By analyzing the values of pressures measured at the beginning and end of the pipeline, it was found that the pressure drop is almost 50% higher than the constructed values. Therefore, it was concluded that there is significant damping disrupting the pipeline flow and possible leakage. That required more frequent corrective and maintenance actions on the pressure pipeline. With a further visual examination in Figure 1, of the cut section of the steel pressure pipeline (DN125, PN 16, total length 820 meters), it was found that deposits were formed on the inner side of the pipe wall. The subjected section of the pressure steel pipe was given a detailed analysis in order to determine the cause of failure.1–9

Iron-oxidizing bacteria, also known as metal-depositing microorganisms, is causing microbiologically influenced corrosion (MIC) in many studies.10–11 MIC is a complex interaction between the environment, the microbial population and the metal substrate.11–13 It is estimated that microbial activity is causing annually about 20% of the corrosion damage of metals, of which a significant part is due to aerobic corrosion influenced by iron-oxidizing bacteria (IOB) and anaerobic corrosion influenced by sulfate-reducing bacteria (SRB).10–12 The reaction of carbon-steel pipes is considered to be a complicated process that is affected by bulk-water quality, including temperature, pH, alkalinity, oxygen concentration and the presence of sulfate and chlorine in the system, etc.13

XX Little and Wagner11 considered that the corrosion occurred in the presence of iron bacteria followed the crevice corrosion mechanism.11 The role of iron-oxidizing bacteria lies in the formation of condensed oxygen zones and the partition of the metal surface into small anodic sites and large surrounding cathodic area.11,12

Moreover, the deposit layers formed by iron-related bacteria could also create condensed oxygen zones and initiate crevice corrosion in water systems containing corrosion inhibitors. The corrosion rate beneath the rust deposits remains high due to the low penetrability of the corrosion inhibitors and declining of the local oxygen concentration where the site might turn anodic.

During the last decades in the cooling water system of the production industry, SRB and IOB were the most common groups of bacteria on tubercular corrosion and induced microbiologically influenced corrosion in the cooling circuit. They caused poor water quality and equipment clogging, which included flow blockages due to tubercle formation, choking of valves and strainers, pipe punctures and high corrosion rates, resulting in the serious pitting corrosion of carbon-steel equipment.14

Despite a large number of research papers reported on the MIC of carbon steels, the quantitative data and information about the real role of the IOB in the MIC process are still not clear, due to the lack of information and evidence of the mechanism describing MIC in water-distribution systems,10–15 because most of these studies were made in the laboratory. In the present paper, the focus is on understanding the real role of IOB in the MIC-process on the wastewater distribution system of thermal power plant Plomin.

2 MATERIALS AND METHODS

2.1 Materials

The corrosion-product samples were taken from a section of damaged steel pressure pipeline (total length 820 m, DN125, PN 16). Two main sample categories were visually identified and investigated. One sample unite was named “the black sediment” because it was visually black, and was taken from the bottom and central mass of corrosion scales that was attached to the inner wall of the pipeline. Another sample that was visually brown, was named “the brown sediment” and was taken from the surface of the corrosion scales turned towards the pipeline center. Basic information of the field conditions, as well as the material and year of installation in the power plant, is shown in Table 1. The tested wastewater sample came from the same water piping.
Table 1: Basic data on the sample of the steel pressure pipeline

<table>
<thead>
<tr>
<th>User:</th>
<th>HEP-Proizvodnja d.o.o</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plant location:</td>
<td>TE PLOMIN 2, Plomin Luka</td>
</tr>
<tr>
<td>Sample dimension:</td>
<td>Pipe ø125*3,6</td>
</tr>
<tr>
<td>Part of installations:</td>
<td>Boiler water pipeline</td>
</tr>
<tr>
<td>Working pressure:</td>
<td>7-10 bar</td>
</tr>
<tr>
<td>Overheating temperature:</td>
<td>20 °C</td>
</tr>
<tr>
<td>Year of construction:</td>
<td>2010.</td>
</tr>
<tr>
<td>Pipeline material:</td>
<td>EN St 35.8 (1.0061), DIN 17175</td>
</tr>
</tbody>
</table>

2.2 Methods

2.2.1 Microbiological analysis of wastewater distribution systems sediment and water

The microbiological test was performed on boiler water taken at the outlet of the system and on the corrosion-products sediment taken from the surface of the pipe. Sulphate-reducing bacteria (SRB) were tested using the Dalynn Biologicals kit, and iron-related bacteria (IRB) using the BTI Products kit. Sargent Welch lead acetate Test Paper and HCl p.a. by Kemika were used for the sulfide presence testing of the bottom, central and top layers of corrosion products.

2.2.2 XRD analysis of wastewater distribution systems sediment

Black and brown corrosion products were subjected to X-ray diffraction analysis (PXRD) on a Shimadzu XRD-6000 diffractometer with Cu–Kα radiation, with a voltage acceleration of 40 kV and current 30 mA, in the range 3–83° 2θ with a step of 0.02° 2θ and a retention time of 0.6 s. The results are graphically presented as a function of the intensity of diffraction maxima of the diffraction angle on the images appeared.

2.2.3 Chemical analysis of water with the calculation of corrosivity indices

Chemical analysis of the wastewater sample was performed in an accredited laboratory for water analysis. Calculation of the Corrosivity Indices was done according to the literature.16

2.2.4 Macro cut of a grinded section of the pipe wall

A macro cut was made from the sample of the steel pipe, on which the corroded wall thickness and its appearance was inspected visually.

3 RESULTS AND DISCUSSION

3.1 Microbiological analysis of sediments and wastewater

The control results of IRB and SRB-bearing samples are shown in Figures 2a and 2b, respectively. The wastewater and corrosion sediments tests yielded identical results shown in Figures 3a and 3b. Positive results were obtained for IRB and negative for SRB.

The control results of the sulfide analysis for the sulfide-free and sulfide-bearing samples are shown in Figures 4a and 4b, respectively. These were used for a visual assessment of the tested samples.

The results of the sulfide analysis on the central, bottom and top layers of the sediment are shown in Figure 5.

Comparison with control samples shows that all three samples were negative, i.e., sulphide-free. Since sulphides are formed as a result of SRB metabolism,17 this result confirms the previous observation that SRB
are not detectable. The microbiological analysis of the sediment samples shows the presence of IRB.

### 3.2 X-ray diffraction analysis

XRD analysis was performed on sediment samples named according to their color, i.e., the black and brown samples. The results are shown in Figures 6 and 7 as a function of the intensity of the diffraction maxima about the diffraction angle.

The black sediment sample consists of very well crystallized magnetite, Fe$_3$O$_4$, ICDD PDF (International Center for Diffraction of the Powder Diffraction File) # 65-3107. No additional diffraction maxima are observed on the diffractogram, thus it can be determined that magnetite is the only crystalline phase in the sample. Moreover, no amorphous halo is observed, so it can be determined that there is no amorphous phase in a larger share.

The diffraction maxima of the brown sediment sample are significantly lower and wider, indicating finer crystallites. No pronounced amorphous halo is observed, although the visual impression may be the opposite due to the low intensities. In this sample, the main crystalline phase is goethite, FeO (OH), ICDD PDF # 81-463, while the secondary phase is magnetite, Fe$_3$O$_4$, ICDD PDF # 65-3107. It is not excluded that another crystalline phase is present in a very small proportion in the sample.

The corrosion products are mainly magnetite and to a lesser extent goethite, which is found in a thin surface layer. There is no sulphate-reducing bacteria and sulphides in the composition of the sediment, and it is supported by the appearance of corrosion that indicates this result. Sulphate-reducing bacteria cause very localized damage and the tested sample shows the appearance of general corrosion over the entire surface of the pipe wall. The positive test on iron-related bacteria, probably indicates the presence of iron oxidizing bacteria which oxidize Fe$^{2+}$, formed by dissolution of the pipe material, into Fe$^{3+}$. Fe$^{3+}$ is an oxidant that significantly promotes the further corrosion of steel. Moreover, redox cyclization probably takes place so that the corrosion process continues. The formation of dense layers of products begins above the anodic site through the accumulation of deposits on the surface. The electrostatic forces gradually strengthen at the location and the surface of the accumulated deposits gradually passes into the cathode. This creates the optimal conditions for the formation of large tubercles.

### 3.4 Macro cut of a grinded section of the pipe wall

A section of the pressure steel pipeline was cut out, freed from loose corrosion products, ground along the profile, to make a macroscopic sample adequate for inspection of the steel pipe thickness, as seen in Figure 8.

Figure 9 shows the inner surface of the pipeline after removal of loose corrosion products. Distinctive, anodic (depressed) and cathodic areas are clearly visible at the surface.

Through a visual inspection it was concluded that the surface was extremely damaged and irregularly thinned. Since the entire surface is equally damaged by corrosion, more and more leaks can be expected to occur on the pipeline over time.
3.5 Physical and chemical analysis of wastewater with calculation of the corrosivity index

The physicochemical characteristics of the tested wastewater are shown in Table 2 and the ionic composition is shown in Table 3.

The Langelier saturation index (LSI) is defined as:

$$\text{LSI} = \text{pH} - \text{pH}_{\text{sat}}$$  \hspace{1cm} (1)

where \(\text{pH}_{\text{sat}}\) is calculated according to:

$$\text{pH}_{\text{sat}} = (9.3 + A + B) - (C + D)$$  \hspace{1cm} (2)

In Equation (2), A depends on the TDS, B depends on the temperature of the water, C depends on the water hardness and D depends on the water alkalinity.

The Larson Skold Index is defined as:

$$\text{L}&\text{Ski} = \left(\frac{[\text{Cl}^-]+[\text{SO}_4^{2-}]}{[\text{HCO}_3^-]+[\text{CO}_3^{2-}]}\right)$$  \hspace{1cm} (3)

where the anion concentrations are presented in Table 3.

The calculated values of the Langelier saturation index (LSI) and Larson-Skold index (L&Ski), calculated according to Equations(1-3, as well as their corrosivity assessment are shown in Table 4.

According to the Langelier index, the water is non-corrosive and tends to precipitate calcium carbonate. XRD analysis has shown that calcium carbonate is not present in the deposits, therefore indicating that no protective carbonate scales were formed at the pipeline sur-

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**Table 2: Physical and chemical analyses of purified wastewater**

<table>
<thead>
<tr>
<th>Properties</th>
<th>Units</th>
<th>Boiler water – output</th>
</tr>
</thead>
<tbody>
<tr>
<td>Appearance</td>
<td>Descriptive</td>
<td>Transparent</td>
</tr>
<tr>
<td>Color</td>
<td>Descriptive</td>
<td>Colorless</td>
</tr>
<tr>
<td>Smell</td>
<td>Descriptive</td>
<td>No smell</td>
</tr>
<tr>
<td>Temperature</td>
<td>°C</td>
<td>17.4</td>
</tr>
<tr>
<td>pH</td>
<td>–</td>
<td>7.4</td>
</tr>
<tr>
<td>Electric conductivity</td>
<td>25°C, μS/cm</td>
<td>3079</td>
</tr>
<tr>
<td>TDS</td>
<td>mg/L</td>
<td>443.3</td>
</tr>
<tr>
<td>Total hardness (Ca + Mg)</td>
<td>mg CaCO₃, mg/L</td>
<td>1665</td>
</tr>
<tr>
<td>Water alkalinity (p-, m-)</td>
<td>mg CaCO₃, mg/L</td>
<td>112</td>
</tr>
<tr>
<td>Total mineralization</td>
<td>mg/L</td>
<td>2089</td>
</tr>
<tr>
<td>Salinity</td>
<td>G NaCl/L</td>
<td>1.33</td>
</tr>
</tbody>
</table>

**Table 3: Ionic composition of purified wastewater**

<table>
<thead>
<tr>
<th>Ion</th>
<th>Concentration / μL/L</th>
<th>Molar mass /g mol⁻¹</th>
<th>Charge</th>
<th>Concentration / meq</th>
</tr>
</thead>
<tbody>
<tr>
<td>Na⁺</td>
<td>25.00</td>
<td>22.99</td>
<td>1</td>
<td>1.0874</td>
</tr>
<tr>
<td>K⁺</td>
<td>3.70</td>
<td>39.83</td>
<td>1</td>
<td>0.0929</td>
</tr>
<tr>
<td>Ca²⁺</td>
<td>339.00</td>
<td>40.08</td>
<td>2</td>
<td>16.9170</td>
</tr>
<tr>
<td>Mg²⁺</td>
<td>199.00</td>
<td>24.31</td>
<td>2</td>
<td>16.3752</td>
</tr>
<tr>
<td>NH₄⁺</td>
<td>0.58</td>
<td>18.04</td>
<td>1</td>
<td>0.0322</td>
</tr>
<tr>
<td>Fe²⁺</td>
<td>0.01</td>
<td>55.85</td>
<td>2</td>
<td>0.0003</td>
</tr>
<tr>
<td>toMn²⁺</td>
<td>0.04</td>
<td>54.93</td>
<td>2</td>
<td>0.0014</td>
</tr>
<tr>
<td>Cl⁻</td>
<td>804.00</td>
<td>35.45</td>
<td>–1</td>
<td>22.6779</td>
</tr>
<tr>
<td>HCO₃⁻</td>
<td>137.00</td>
<td>61.02</td>
<td>–1</td>
<td>2.2453</td>
</tr>
<tr>
<td>SO₄²⁻</td>
<td>581.00</td>
<td>96.06</td>
<td>–2</td>
<td>12.0966</td>
</tr>
<tr>
<td>Difference in anion and cation concentration</td>
<td>2.54</td>
<td>Percentage deviation</td>
<td>3.56</td>
<td></td>
</tr>
</tbody>
</table>

**Table 4: Corrosivity indices of the purified wastewater**

<table>
<thead>
<tr>
<th>Index</th>
<th>Value</th>
<th>Interval</th>
<th>Corrosivity assessment</th>
</tr>
</thead>
<tbody>
<tr>
<td>LSI</td>
<td>0.21</td>
<td>LSI &gt; 0</td>
<td>The water is supersaturated with calcium carbonate and there is a possibility of deposit formation.</td>
</tr>
<tr>
<td>L&amp;Ski</td>
<td>15.49</td>
<td>L&amp;Ski &gt; 0.8</td>
<td>Water tends to cause localized corrosion.</td>
</tr>
</tbody>
</table>
face. The Larson Skold Index indicates that wastewater is capable of causing serious localized corrosion. However, from the occurrence of corrosion along the entire wall of the pipeline, it can be concluded that sulphates and chlorides are not decisive corrosion agents, but only corrosion accelerators. The primary causes of the corrosion are likely the iron-oxidizing bacteria, which, in combination with sulphides and chlorides in the wastewater, cause intense corrosion, the formation of abundant, compacted dust-like corrosion products and irregular but general thinning of the pipe wall.

4 CONCLUSIONS

The following conclusions can be drawn:

• Corrosion products are made mostly of compacted dust-like magnetite with a thin surface layer of goethite.
• The presence of IRB was found in the sediment and the distribution system wastewater.
• According to the Langelier index, water should precipitate calcium carbonate, but it has not been identified in the corrosion sediment layer.
• Due to the general appearance of corrosion along the pipeline axes and circumference, sulphates and chlorides, despite the high Larson Skold index, are probably not decisive corrosion agents, but only corrosion accelerators.
• The primary causes of corrosion are likely the iron-oxidizing bacteria, which in combination with sulphates and chlorides in the water, cause intense corrosion and the formation of abundant, compacted dust-like corrosion products that are clogging the pipeline.
• The macro sanding of the cut pipe wall revealed that the entire pipe surface, longitudinally and circumferentially, was equally damaged by corrosion, showing irregularities in the form of overlapping wide pits so that over time, more and more leaks can be expected on the pipeline.

5 REFERENCES