Influence of the Microstructure on the Degree of Sensitization of a Duplex Stainless Steel UNS S31803 Aged at 650°C

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This work evaluates the phase transformations during aging of an UNS S31803 DSS at 650°C and its influence on the DOS. The material was solution treated at 1175°C and then aged at 650°C for times up to 360 h. SEM-BSC images indicate the formation of Cr,N, σ and χ phases in the samples aged at 650°C. The analysis of DL-EPR curves, obtained in a 1 M H₂SO₄ + 0.25 M NaCl + 0.01 M KSCN solution, shows an increase in DOS values for samples aged at 650°C. Probably, this increase observed in DOS values is mainly related to the presence of Cr- and/or Mo-depleted α, as a result of σ phase, χ phase and/or Cr,N formation at 650°C. Moreover, a possible healing up of Cr- and/or Mo-depleted areas is observed between 1 h and 4 h of aging at 650°C.

Keywords: duplex stainless steel, DL-EPR, degree of sensitization, sigma phase

1. Introduction

Duplex stainless steels (DSS) are those with a microstructure composed by two phases, ferrite (α) and austenite (γ), which appear in approximately equal volume fractions¹. Their favourable combination of properties makes this class of steels widely employed in oil and gas, petrochemical and pulp and paper industries².

Between −50°C and 250°C, a great combination of mechanical properties and corrosion resistance is offered by DSS. However, a large variety of secondary undesirable phases may form in the temperature range of 300°C and 1000°C, during isothermal aging or as a result of incorrect heat treatment procedures¹.

Between 300°C and 1000°C the most important secondary phases which can precipitate are alpha prime phase (α′) and sigma (σ) phase. Alpha prime phase is chromium enriched and, depending on ferrite composition, its formation may happen by nucleation and growth of precipitates or by spinodal decomposition³. Moreover, α′ phase formation is expected in temperatures below 500°C⁴, being mostly studied at 475°C. Sigma phase is chromium and molybdenum enriched if compared to the matrix alloy⁴, and it is mainly formed between 650°C and 900°C⁵.

Results presented by Borba and Magnabosco⁶ reinforce the possibility of σ phase formation in temperatures below 700°C, as it is observed the formation of this phase during aging of an UNS S31803 DSS at 600°C and 650°C, highly evidenced after 96 h of aging. Arikan and Doruk⁷, also working with an UNS S31803 DSS, observe the formation of σ phase from 1000 min (~16 h) of aging at 650°C on, with an increase of σ phase amount with aging time.

Between 650°C and 900°C σ phase may form from ferrite phase by an eutectoid decomposition mechanism leading to secondary austenite (γ₅) formation (α→σ+γ₅)⁷,⁸. However, at 850°C, some authors⁹,¹⁰ found that σ phase may form by different mechanisms depending on aging time. So, at 850°C, for aging times up to 10 min, σ phase may form from direct precipitation from ferrite phase, resulting in Cr and Mo depleted α, also denominated secondary ferrite (α₅) phase (α→σ+α₅). Moreover, between 30 min and 5 h of aging at 850°C the eutectoid decomposition of α, resulting in γ₅ (also depleted in Cr and Mo) is observed. Also, after total α phase consumption at 850°C, σ phase fractions continue to rise due to γ₅ phase consumption for its formation⁹,¹⁰.

Besides σ phase formation, chi (χ) phase and chromium nitride (Cr,N) formation may also result from isothermal heat treatments of DSS between 700°C and 900°C⁹. Usually, Cr,N precipitation occurs intergranularly at α/α grain boundaries or at α/γ interfaces¹¹,¹², together to the formation of γ₅, causing chromium depleted zones¹¹. Chi phase formation generally occurs before σ phase formation and is gradually consumed by σ phase precipitation¹¹,¹². As χ phase is enriched in Cr and Mo as well as σ phase (but with larger amounts of Mo than sigma phase¹¹,¹²), it is expected to promote depletion of the surrounding matrix in such elements¹¹. Besides some authors claim that χ phase formation occurs at α/γ interfaces¹¹, other authors indicate that its formation starts at α/α grain boundaries¹².

The formation of Cr and Mo depleted phases such as α₅ and/or χ₅ have a strong influence in localized corrosion resistance and such influence can be measured by the degree of sensitization (DOS) through Double-Loop Electrochemical Potentiokinetic Reactivation (DL-EPR) tests.
Although DL-EPR tests were at first developed in order to evaluate the DOS of austenitic stainless steels\textsuperscript{15}, recent studies use DL-EPR tests to evaluate the influence of isothermal aging on DSS and, consequently, of secondary phases precipitation, in the DOS of these steels\textsuperscript{7,16-23}.

As far as corrosion resistance is concerned, σ phase formation promotes an increase in the DOS\textsuperscript{7,16-18,20-23} probably as a result of etching to adjacent areas of formed σ phase, as these regions are likely to be depleted in chromium and/or molybdenum. Some authors, working with UNS S31803 DSS aged at 650°C, show that, after DL-EPR tests in a 2 M H\textsubscript{2}SO\textsubscript{4} + 0.5 M NaCl + 0.01 M KSCN solution, the DOS increases from 0.036236 (samples aged for 16 h) to 0.824235 (samples aged for 527 h), while others show that, after DL-EPR tests in a 0.3 M HCl + 0.002 M Na\textsubscript{2}SO\textsubscript{4} solution, the DOS increases from 0.075 (samples aged for 60 min) to 0.2574 (samples aged for 300 min)\textsuperscript{22}.

Recent studies related to isothermal heat treatments of DSS at 650°C are mainly based upon σ phase formation\textsuperscript{7,2}; however, the possibility of χ phase formation and even of Cr\textsubscript{N} formation may not be neglected, as there is lack of information about such phases as far as heat treatments below 700°C are concerned.

Moreover, although a great number of recent works relate DSS phase transformations due to aging between 650°C and 900°C\textsuperscript{7,16-20,22-23} and DOS evolution, some present an incomplete microstructural analysis, mainly concerning the Cr and Mo depleted phases and/or areas, and fail to correlate DOS values and even DL-EPR curves features with the obtained microstructures.

Therefore, the aim of this work is to study phase transformations and its influence on the DOS of an UNS S31803 DSS aged at 650°C.

2. Experimental Procedures

The studied material, UNS S31803 DSS, has the chemical composition given in Table 1, and was received as a 3 mm thick sheet, rolled at 1100°C and cooled by forced water jet and air blast.

Solution heat treatment was conducted in the as received material for 30 min at 1175°C, followed by water quenching. Then the samples were aged at 650°C for 20 min, 1 h, 4 h or 360 h in a tubular electric furnace which maintained aging temperature 2°C around the chosen temperature. All heat treatments were conducted in a 99.9% N\textsubscript{2} atmosphere to suppress oxidation of the samples surfaces. Following the heat treatments samples were water quenched.

After the heat treatments the specimens were ground using silicon carbide (SiC) papers to an 80-grit finish before mounting in thermosetting plastic, leaving an exposed surface area of approximately 0.4-0.6 cm\textsuperscript{2}, parallel to the rolling direction. The mounted samples were metallographic polished in a semi-automatic grinding and polishing machine, with final polishing provided by 1-μm diamond abrasive. Immediately after polishing, samples surfaces were washed with ethyl alcohol (C\textsubscript{2}H\textsubscript{5}OH) and dried with hot blown air.

X-Ray diffraction, using a Cu-Kα radiation, helped in phase identification. The scan was performed in the 2θ angular range of 30-120° with a scan speed of 1°/min and a step size of 0.04°.

Scanning electron microscopy (SEM), through backscattered (BSC) electron images, were also used for phase determination. BSC electron images were analysed considering that χ phase appears lighter than σ phase due to its higher molybdenum content\textsuperscript{11-14} and that Cr\textsubscript{N} appear darker due to its nitrogen content\textsuperscript{24}.

Quantitative metallography of σ and χ phases (showed as a sum and denominated as intermetallic phases) was performed in six fields of each sample with an automated image analysis system attached to an optical microscope, using SEM-BSC images of polished samples. The evaluation of ferrite phase percentage was obtained by magnetic measurements using a ferritscope (FISCHER MP30) calibrated with standard samples. In each sample 20 measurements were made.

Taking into consideration only the presence of α, γ and intermetallic phases, γ content can be calculated using Equation 1:

\[
\% \gamma = 100 - \% \alpha - \% \text{intermetallic phases.} \quad (1)
\]

DL-EPR tests were carried out in a 1 M H\textsubscript{2}SO\textsubscript{4} + 0.25 M NaCl + 0.01 M KSCN solution, exposed to air, at a controlled temperature of 30°C ± 2°C, using a saturated calomel electrode (SCE) as a reference electrode, a platinum wire as counter electrode and, as working electrode, samples polished until 1-μm diamond abrasive finish were used.

Samples were immersed in the test solution for 5 minutes to ensure both the stabilization of the open circuit potential (OCP) and the testing temperature of the cell, and then the anodic polarization was conducted from the OCP, at a scan rate of 1.67 mV/s. The scan direction was reversed at the potential of +300 mV\textsubscript{SCE} and the tests were finished at the OCP previously measured. After DL-EPR tests, samples surfaces were washed with distilled water and dried with hot blown air to be examined by optical microscopy. The electrochemical tests were repeated three times for each heat-treatment condition.

The DOS was determined by the ratio between the maximum anodic current density during the reactivation (i\textsubscript{a}) and the maximum anodic current density during the activation (i\textsubscript{c}), as described by Equation 2:

\[
\text{DOS} = \frac{i_a}{i_c}. \quad (2)
\]

### Table 1. Chemical composition (wt%) of the studied material.

<table>
<thead>
<tr>
<th>Cr</th>
<th>Ni</th>
<th>Mo</th>
<th>Mn</th>
<th>N</th>
<th>C</th>
<th>Si</th>
<th>Cu</th>
<th>Co</th>
<th>P</th>
<th>S</th>
<th>Nb</th>
<th>Fe</th>
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<td>22.07</td>
<td>5.68</td>
<td>3.19</td>
<td>1.38</td>
<td>0.17</td>
<td>0.017</td>
<td>0.34</td>
<td>0.15</td>
<td>0.06</td>
<td>0.020</td>
<td>0.001</td>
<td>0.01</td>
<td>Bal.</td>
</tr>
</tbody>
</table>
3. Results and Discussion

Figure 1 presents the equilibrium phase content as a function of temperature, calculated using Thermo-Calc® software and TC-Fe6 database. At 1175°C (solution temperature) the equilibrium consists of α and γ phases with approximate volume fractions of 62% and 38% respectively. Ferrite phase volume fraction obtained experimentally (57.3 ± 2.3%) is also plotted in Figure 1 and it is almost identical to the calculated equilibrium value (62%), considering the standard deviation of the experimental measurements. This may indicate that the solution treatment performed at 1175°C for 30 min is enough to reach equilibrium between α and γ phases and to develop a duplex microstructure. Still in Figure 1, at 650°C, the calculated equilibrium phases are α, γ, σ and Cr₂N, with approximate volume fractions of 12%, 54%, 32% and 2%, respectively.

X-ray diffraction patterns for the UNS S31803 DSS studied are presented in Figure 2. For the solution treated sample (Figure 2a) and for the sample aged 4 h (Figure 2b) at 650°C the X-ray diffraction patterns indicate only α and γ phases, while the X-ray diffraction pattern for the sample aged 360 h at 650°C (Figure 2c) indicates Cr₂N, σ and χ phases, besides α and γ phases. However, as all Cr₂N and χ phase peaks appear combined with other phases peaks, this technique cannot be used to clearly determine the formation of such phases.

In order to confirm the phases formed in the UNS S31803 DSS studied, SEM-BSC images were obtained (Figure 3). Solution treated sample (Figure 3a) indicates only α and γ phases. For samples aged 1 h (Figure 3b) and 4 h at 650°C (Figure 3c), besides α and γ phases, Cr₂N (darker regions, due to nitrogen content) formation mainly occurs at α/α phase boundaries. For the sample aged 360 h at 650°C (Figure 3d), besides α and γ phases, it is also observed the presence of σ phase (light grey, due to chromium and molybdenum contents), χ phase (white, as it is enriched in molybdenum if compared to sigma phase) and...
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Cr2N (darker points, due to nitrogen content). Therefore, BSC images confirm the evidences of Cr2N and χ phase formation previously pointed out by X-ray diffraction patterns (Figure 2c).

Comparing the microstructural observations of the present work with those of Arikan and Doruk it seems that, besides these authors make a brief comment on the possibility of χ phase or Cr2N formation after aging an UNS S31803 DSS, they claim it was not possible to observe χ or Cr2N in their optical micrographs. However, Arikan and Doruk would have probably observed such phases if they had not performed SEM analysis in their samples after electrolytic etching with oxalic acid, as they would have prevented the Cr enriched phases to be etched away before SEM analysis.

It is important to point out that the absence of Cr2N peaks in the diffraction pattern of the sample aged 4 h at 650°C (Figure 2b) may be related to the small amounts of Cr2N observed in the BSC images (Figure 3c). So, considering the small amounts of Cr2N, their volume fraction was neglected in phase quantification, and σ and χ phases will be designated as intermetallic phases, since they are both Cr- and Mo-rich phases, probably associated to areas depleted in those elements.

Figure 4 shows the volume fractions of phases obtained by quantitative metallography of BSC images. For samples aged for times up to 4 h at 650°C an increase in γ volume fraction as well as a decrease in α volume fraction is observed (Figure 4), but only small amounts of Cr2N are observed mainly at α/α phase boundaries (Figure 3). Intermetallic phase formation between 4 h and 360 h of aging at 650°C leads to a great drop in α phase volume fraction combined with an increase in γ volume fraction (Figure 4). The increase in γ volume fraction observed for samples aged up to 4 h at 650°C may be related to γ2 formation in cooperation with Cr2N, while between 4 h and 360 h of aging the increase observed in γ fraction is probably related to γ2 formation as a result of Cr2N, σ and χ phases formation. Moreover, the reduction observed in α phase fractions (Figure 4) indicates that the intermetallic phases and Cr2N are probably formed by the consumption of α phase.

A comparison between the experimental volume fractions obtained after 360 h of aging at 650°C (Figure 4) and the equilibrium volume fractions (Figure 1) shows that the remaining volume fraction of α phase (17.4 ± 1.0%) is higher than expected (12%) while the volume fraction of formed intermetallic phases (24.3 ± 1.1%) is lower than

Figure 3. SEM-BSC images of: (a) solution treated sample, (b) sample aged 1 h at 650°C, (c) sample aged 4 h at 650°C and (d) sample aged 360 h at 650°C. Ferrite (dark grey), austenite, sigma (light grey) and chi (white) phases can be distinguished, besides the presence of chromium nitrides (dark points at grain boundaries).
the value predicted for $\sigma$ phase at the equilibrium (32%). Moreover, after 360 h of aging at 650°C a slightly higher volume fraction value of $\gamma$ phase (58.3 ± 1.5) is observed than the expected at equilibrium (54%). So, it is important to highlight that as the equilibrium phases expected at 650°C are $\alpha$, $\gamma$, $\sigma$ and Cr$_2$N (Figure 1), the differences found in the comparison between experimental and calculated volume fractions of phases, as well as the presence of $\chi$ phase (Figure 3), indicate that aging at 650°C for time periods up to 360 h is not enough to establish the thermodynamic equilibrium in the UNS S31803 DSS studied.

DL-EPR tests in 1 M $H_2SO_4$ + 0.25 M NaCl + 0.01 M KSCN, for the solution treated sample and for samples aged at 650°C, produced the typical curves presented in Figure 5.

It can be observed in Figure 5 that there is an evident increase in $i_1$ and $i_2$ for the sample aged 360 h at 650°C. This increase may be related to the development of chromium and molybdenum depleted areas as a consequence of intermetallic phase formation.

Another important observation is the existence of a second current density peak during reactivation for the sample aged 360 h at 650°C. Magnabosco, Santos and Mélo suggest that the potentiodynamic polarization curves, in acid solution, for a solution treated sample of an UNS S31803 duplex stainless steel present two distinct anodic current density peaks, indicating that the one with a lower potential corresponds to $\sigma$ phase and the other with a higher potential represents $\gamma$ phase. Moreover, it is expected that a multiphase material presents an anodic polarization curve which represents a sum of the individual curves of existing phases20. So, the appearance of a second peak may be related to the existence of $\sigma$ and $\gamma$ phases, both with distinct electrochemical behaviours. Also, a high Cr and Mo depletion in the original $\alpha$ matrix in the sample aged 360 h at 650°C could explain why the $\alpha$ phase peak (lower potential) is highly evidenced in this sample, indicating the possibility of $\alpha$ formation.

However, it is important to point out that, as there is only one well defined peak in the activation portion of the curve, even though a second peak appears during reactivation, only the reactivation peak with higher potential was used to calculate the degree of sensitization for the sample aged for 360 h at 650°C in order to properly compare the values of $i_1$ and $i_2$.

Optical micrographs of samples surfaces after DL-EPR tests, for the solution treated sample and for the samples aged at 650°C, are presented in Figure 6. A more intense etching at $\alpha/\alpha$ grain boundaries, $\alpha/\gamma$ interfaces and inside $\alpha$ grains is observed when the solution treated sample and the samples aged at 650°C are compared (Figure 6).

While the etch at $\alpha/\alpha$ grain boundaries is enhanced from 1 h of aging at 650°C on (Figure 6c), at $\alpha/\gamma$ interfaces it is better observed from 4 h of aging at 650°C on (Figure 6d). Also, some areas inside $\alpha$ grains are slightly etched after 1 h of aging at 650°C and this etching becomes more intense with aging time, leading to a great material removal inside $\alpha$ grains in the sample aged 360 h at 650°C.

A comparison between SEM-BSE images (Figure 3) and samples surfaces after DL-EPR tests (Figure 6) shows that etching provided by DL-EPR tests promotes material removal at intermetallic phase adjacent areas. Therefore, samples surfaces behaviour after DL-EPR tests are absolutely consistent with the microstructural observations as the material removal is probably taking place at chromium and/or molybdenum depleted areas adjacent to intermetallic phase formation. It is important to point out that $\gamma$ phase remains with no signs of etching after DL-EPR tests, reinforcing that the secondary phases formation observed are mainly related to phase transformations taking place at $\alpha$ phase.

As the analysis of the volumetric fraction of phases (Figure 4) indicates not only an increase in $\gamma$ phase fractions but also a remaining $\alpha$ phase fraction (even after aging 360 h at 650°C), the chromium and/or molybdenum depleted areas may be related to $\alpha_1$ and/or $\gamma_2$ phases formation as a result of intermetallic phase formation. This statement is valid as it is not possible to separate the contributions of the original $\alpha$ matrix and a possible chromium and/or molybdenum $\alpha$ phase coexistence in the analysed samples.

Moreover, as the main precipitation observed for samples aged for times up to 4 h at 650°C is related to

![Figure 4. Volume fraction of phases as a function of aging time at 650°C.](image)

![Figure 5. Typical DL-EPR curves in a 1 M $H_2SO_4$ + 0.25 M NaCl + 0.01 M KSCN solution, for the solution treated sample and for samples aged at 650°C. The maximum anodic current density during the reactivation ($i_2$) and the maximum anodic current density during the activation ($i_1$) are indicated for the solution treated sample. Arrows indicate the scanning direction.](image)
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Cr$_2$N formation, probably, for such samples, the etched areas are depleted in chromium, although the slightly etched areas observed inside ferrite grains may indicate an early stage of some secondary phase formation, presenting particles yet too small to be distinguished by SEM-BSE images. However, when the sample aged 360 h at 650°C is analysed, the etched areas may be depleted in chromium and/or molybdenum as the intermetallic phases formed are chromium and molybdenum enriched.

DOS evolution as a function of aging time at 650°C is compared to the solution treated sample behaviour in Figure 7. It can be observed in Figure 7a that aged samples until 4 h at 650°C present DOS values a little higher than that of the solution treated sample, but values are still very low. However, for the sample aged for 360 h at 650°C it is observed a great increase in DOS values when compared to the solution treated sample or even other samples aged at 650°C (Figure 7a).

In addition, a better observation of DOS values of samples aged up to 4 h at 650°C is presented in Figure 7b. Besides the small increase in DOS values until 4 h of aging at 650°C, when compared to the solution treated sample,
apparently between 1 h and 4 h of aging, there is a slight reduction tendency in DOS values, already observed by other authors\(^{20,23,27}\). This reduction tendency in DOS values may indicate healing up of chromium/molybdenum depleted areas due to Cr and Mo redistribution between the surrounding matrix and depleted areas, as stated elsewhere\(^{27}\). This Cr and Mo redistribution is more likely to happen between the surrounding matrix and depleted areas than between intermetallic phases and depleted areas as intermetallic phases do not behave as solid solutions when it comes to alloying elements redistribution.

In addition, the great increase observed for DOS values for the sample aged 360 h at 650°C is consistent with the observations of higher intermetallic phase fractions values for this sample, showed in Figure 4, as higher intermetallic phase fractions may indicate the presence of more areas subjected to Cr and Mo depletion. Moreover, taking into consideration that the γ phase is not etched during DL-EPR tests and the fact that the highly evidenced α phase peak during reactivation (in the sample aged 360 h at 650°C) is probably due to α\(_2\) phase formation, the main Cr-/Mo-depleted phase responsible for the increase in DOS values seems to be α\(_2\).

Thus, this small increase in DOS values until 4 h of aging, besides the slight decrease observed between 1 h and 4 h of aging, as well as the great increase for sample aged for 360 h at 650°C point out that intermetallic phase formation at 650°C leads to the formation of chromium and/or molybdenum depleted regions, which are considerably deleterious to corrosion resistance of UNS S31803 duplex stainless steel studied in this work.

4. Conclusions

- Samples aged for times up to 4 h at 650°C mainly present Cr\(_2\)N formation at α/α\(_2\) grain boundaries and a possible γ\(_2\) phase formation;
- Aging at 650°C for times up to 360 h leads to Cr\(_2\)N, χ and σ phase formation probably accompanied by Cr- and/or Mo-depleted α\(_2\) and/or γ\(_2\) phase formation;
- A comparison between sample surfaces after DL-EPR tests and SEM-BSC images of aged samples shows that etching provided by DL-EPR tests promotes material removal at intermetallic phase adjacent areas, indicating that the material removal is probably taking place at chromium and/or molybdenum depleted areas, which may correspond to α\(_2\) and/or γ\(_2\) phases;
- A slight decrease in DOS values is observed between 1 h and 4 h of aging at 650°C indicating a possible healing up of the microstructure due to Cr and Mo redistribution between the surrounding matrix and depleted areas;
- The great increase observed for DOS values for the sample aged 360 h at 650°C may be associated with the existence of more areas subjected to Cr-/Mo-depletion as a consequence of higher intermetallic phase fractions values observed for this sample. Also, there is evidence that the main Cr-/Mo-depleted phase responsible for the increase in DOS values is α\(_2\).

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References