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Evaluation by the Double Loop Electrochemical Potentiokinetic Reactivation Test of Aged Ferritic Stainless Steel Intergranular Corrosion Susceptibility

H. SIDHOM, T. AMADOU, and C. BRAHAM

An experimental design method was used to determine the effect of factors that significantly affect the response of the double loop–electrochemical potentiokinetic reactivation (DL-EPR) test in controlling the susceptibility to intergranular corrosion (IGC) of UNS S43000 (AISI 430) ferritic stainless steel. The test response is expressed in terms of the reactivation/activation current ratio (I_r/I_a pct). Test results analysed by the analysis of variance (ANOVA) method show that the molarity of the H_2SO_4 electrolyte and the potential scanning rate have a more significant effect on the DL-EPR test response than the temperature and the depassivator agent concentration. On the basis of these results, a study was conducted in order to determine the optimal operating conditions of the test as a nondestructive technique for evaluating IGC resistance of ferritic stainless steel components. Three different heat treatments are considered in this study: solution annealing (nonsensitized), aging during 3 hours at 773 K (500 °C) (slightly sensitized), and aging during 2 hours at 873 K (600 °C) (highly sensitized). The aim is to find the operating conditions that simultaneously ensure the selectivity of the attack (intergranular and chromium depleted zone) and are able to detect the effect of low dechromization. It is found that a potential scanning rate of 2.5 mV/s in an electrolyte composed of H_2SO_4 3 M solution without depassivator, at a temperature around 293 K (20 °C), is the optimal operating condition for the DL-EPR test. Using this condition, it is possible to assess the degree of sensitization (DOS) to the IGC of products manufactured in ferritic stainless steels rapidly, reliably, and quantitatively. A time–temperature–start of sensitization (TTS) diagram for the UNS S43000 (France Inox, Villepinte, France) stainless steel was obtained with acceptable accuracy by this method when the IGC sensitization criterion was set to $I_r/I_a > 1$ pct. This diagram is in good agreement with the time–temperature–start of precipitation (TTP) diagram that delineates the domain of low dechromization consecutive to chromium carbide precipitation.

I. INTRODUCTION

FERRITIC stainless steels are usually used in the manufacturing of steam generators of power plants. These parts are very sensitive to the heat generated by welding, during the maintenance operations, and throughout service aging. The thermal cycles specific to these conditions induce structural changes, by a diffusion mechanism, *i.e.*, the precipitation of chromium carbides and intermetallic phases or without a diffusion mechanism, *i.e.*, the transformation of austenite into martensite during cooling. Such changes may lead, directly or indirectly, to the formation of relatively large depleted chromium zones that are able to sensitize the steel to intergranular corrosion.^[1] These phenomena have occurred in several ferritic stainless steel grades

such as AISI 430, AISI 430 Ti, AISI 430 Nb, AISI 434, AISI 444, and AISI 446 after solution annealing (1 hour at 1366 K (1093 °C) and water or air cooling) and after aging at 1033 K (760 °C) for 1 hour.^[2] They were also observed in the case of AISI 405 at the solution annealed state (0.5 hours at 1323 K (1050 °C) and water cooling) and at aged states at temperatures varying between 773 K and 1123 K (500 °C and 850 °C).^[3] The same phenomenon was observed for superferritic stainless steel X1Cr26Mo aged at temperatures varying from 723 K to 923 K (450 °C to 650 °C) for periods ranging from 2 minutes to 10 hours.^[4]

The frequently used methods to determine the degree of sensitization to the IGC are either destructive standard tests (ASTM A 262), *i.e.*, Strauss^[5] or Huey,^[6] which provide qualitative evaluations, or nondestructive tests (ASTM G 108) of the electrochemical double loop–electrochemical potentiokinetic reactivation (DL-EPR)^[2–4,7–27] type, which provide a quantitative assessment. However, the IGC assessment by the DL-EPR test is not absolute. Indeed, this evaluation is highly dependent on test operating conditions, such as the nature and the concentration of the electrolyte, the depassivator type, the potential scanning rate, and the

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temperature of the solution. In previous works, authors have optimized and validated the test operating conditions in the case of duplex^[8] and austenitic stainless steels.^[9] For these materials, the reproducibility of the results, the selectivity of the attack, and the ability of the test to detect the effects of weak dechromizations were clearly confirmed.

In order to continue investigating the optimal operating conditions of the DL-EPR test for a quantitative evaluation of the degree of sensitization (DOS) to the IGC of ferritic stainless steels, the experimental design method was used to determine the effects of the most significant parameters on the response of the DL-EPR test. Selected operating conditions on the basis of the attack selectivity and the ability of the test to detect the effect of low dechromization were performed to establish a time–temperature–start of sensitization (TTS) diagram for the UNS S43000 stainless steel.

II. MATERIAL, TREATMENTS, AND TESTS

A. Material and Treatments

The studied material is the most widely used ferritic stainless steel in the industry: UNS S43000 stainless steel type (AISI 430 or, according to the European standard, X8Cr17). Its chemical composition is reported in Table I. The material was received in the form of a 3-mm-thickness sheet.

1. Annealing treatments

Two conditions of solution annealing treatments were applied to the studied material:

- (1) “air solution annealing”: heating at 1373 K (1100 °C) for 1 hour followed by air cooling, and
- (2) “water solution annealing”: heating at 1373 K (1100 °C) for 1 hour followed by water cooling.

2. Aging treatments

Aging treatments were performed on samples of water-solution-annealed steel only, at temperatures ranging from 673 K to 1123 K (400 °C to 850 °C) during periods ranging from 5 minutes to 8 hours followed by air cooling.

B. Tests and Criteria

1. Microstructural observations

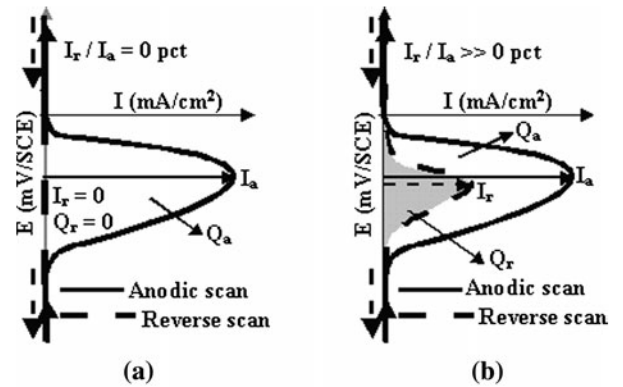
The phenomena of precipitation that occur during solution annealing or aging treatment were studied by metallographic examinations and by energy-dispersive X-ray microanalysis under scanning electron microscope (SEM).

2. DL-EPR test and sensitization criteria

The assessment of the DOS to the IGC of the annealed or aged steel was realized using the DL-EPR test. This electrochemical test consists of monitoring the anodic current (I_a) variation during a scan of the potential E (mV/s) from an active state of the sample to a passive state, and the reactivation current (I_r) during the return to the active state, using a three electrodes setup. Sensitization to the IGC is evaluated by the ratio (I_r/I_a) of anodic (I_a) and reactivation (I_r) current densities or by the ratio of charge densities (Q_r/Q_a). The steel is considered as sensitized when the ratio $I_r/I_a \geq 1$ pct or $Q_r/Q_a \geq 1$ pct (Figure 1).

3. Method and criteria for DL-EPR test operating conditions optimization

The DL-EPR test was performed using an experimental design that varies the concentration of the electrolyte [H_2SO_4] between 0.1 and 6 M, the concentration of the depassivator [NH_4SCN] between 0 and 10^{-2} mol/L, the rate (dE/dt) of the potential scanning (from -500 to $+200$ mV/ECS) between 0.5 and 5 mV/s, and the temperature of the electrolyte (T) between 293 K and 308 K (20 °C and 35 °C). The electrolyte temperature was controlled using a GRANT stirred thermostatic bath. The electrolyte was stirred without being deaerated.



Sensitization criteria [19].

	Non sensitized state	Sensitized state
I_r / I_a (%)	< 1	≥ 1
Q_r / Q_a (%)	< 1	≥ 1

I_a = activation current density
 I_r = reactivation current density
 Q_a = activation charge density
 Q_r = reactivation charge density

Fig. 1—Principle of DL-EPR technique and IGC sensitization criteria. (a) Non sensitized state. (b) Sensitized state.

Table I. Chemical Composition of UNS S43000 Stainless Steel (Weight Percent)

Steel Grade	C	S	P	Si	Mn	Cr	Fe
UNS S43000 (EN: X8Cr17)	0.079	0.001	0.019	0.34	0.211	16.24	balance

The tests were performed on three different microstructural states that are supposed to cover a wide range of sensitization degrees:

- (1) solution annealed with water cooling (expected to be nonsensitized),
- (2) aged at 773 K (500 °C) during 3 hours (supposed to be slightly sensitized), and
- (3) aged at 873 K (600 °C) during 2 hours (supposed to be highly sensitized).

Operating conditions optimization is made on the basis of the detection of significant value of the ratio I_r/I_a (pct) (proportional to the DOS), acceptable reproducibility (low scatter), good selectivity (intergranular

attack exclusively), and very high sensitivity to low DOS (low dechromization).

The DL-EPR test was performed on specimen surfaces that were polished using a 1200 mesh size abrasive paper.

a. Verification of the selectivity of the DL-EPR test. The selectivity of the test is evaluated by SEM examination of sites and forms of corrosion attack under different DL-EPR test conditions. The following criteria were adopted.

- (1) The test is considered as selective when the attack is intergranular exclusively.

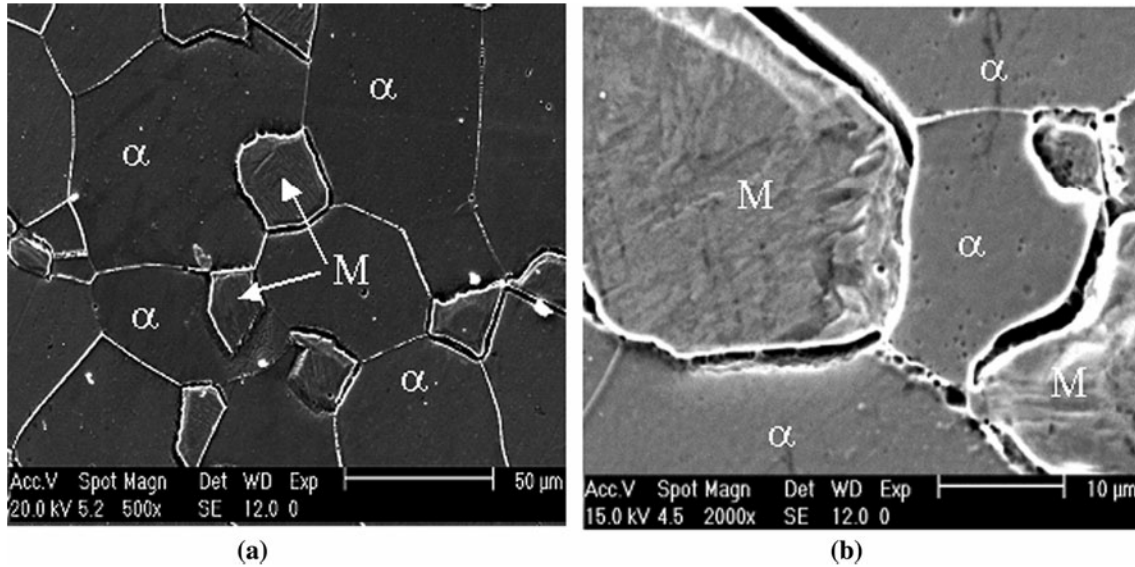


Fig. 2—Structure after water solution annealing of the UNS S43000 stainless steel. (a) Ferrite + aggregates of austenite transformed into martensite and likely small amount of residual austenite (γ_{res}). (b) Aggregates of martensite and likely small amount of residual austenite (γ_{res}) (higher magnification).

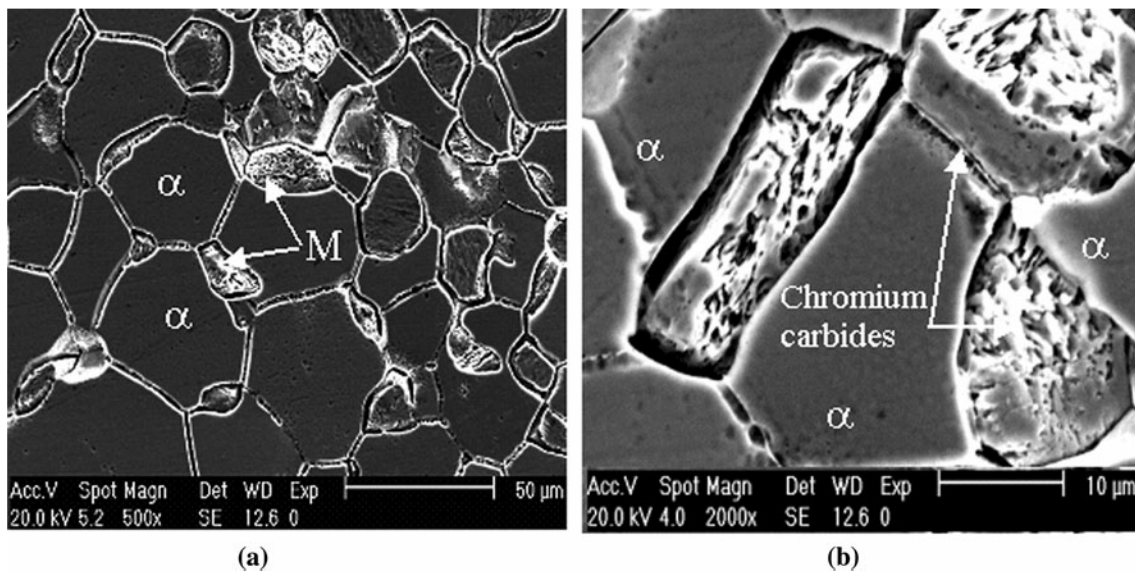


Fig. 3—Structure after air solution annealing of the UNS S43000 stainless steel. (a) Ferrite (dark) + aggregates austenite transformed into martensite (clear) and likely small amount of residual austenite (γ_{res}). (b) Precipitation of chromium carbides in aggregates of martensite and in ferritic grain boundaries (higher magnification).

- (2) The test is considered as nonselective when the attack is generalized. This kind of attack is characterized by uniform pitting distribution and additional intercrystalline dissolution.

b. Verification of the DL-EPR sensibility and response conformity. The test sensitivity to detect low dechromization is evaluated by the (I_r/I_a) pct ratio. Its conformity is assessed by the intensity and the depth of the intercrystalline attack, which is qualitatively evaluated by SEM examination. The following criteria were adopted:

- (1) step: no intercrystalline attack,
- (2) dual: slight intercrystalline attack, and
- (3) ditch: high intercrystalline attack.

III. RESULTS

A. Microstructure

1. Microstructure of solution-annealed steel

SEM metallographic examinations of samples annealed at 1373 K (1100 °C) cooled in water or in air reveal a mixed microstructure consisting of ferrite (α) and aggregates of austenite transformed to carbon supersaturated martensite (M) and likely a small amount of residual austenite (γ_{res}). Phase partitions are approximately 80 to 90 pct α and 10 to 20 pct aggregates (M + γ_{res}). These aggregates result from total or partial transformation of the austenite formed at high temperature in the intercritical area (α and γ) during rapid cooling. They appear mainly in the form of islets in the

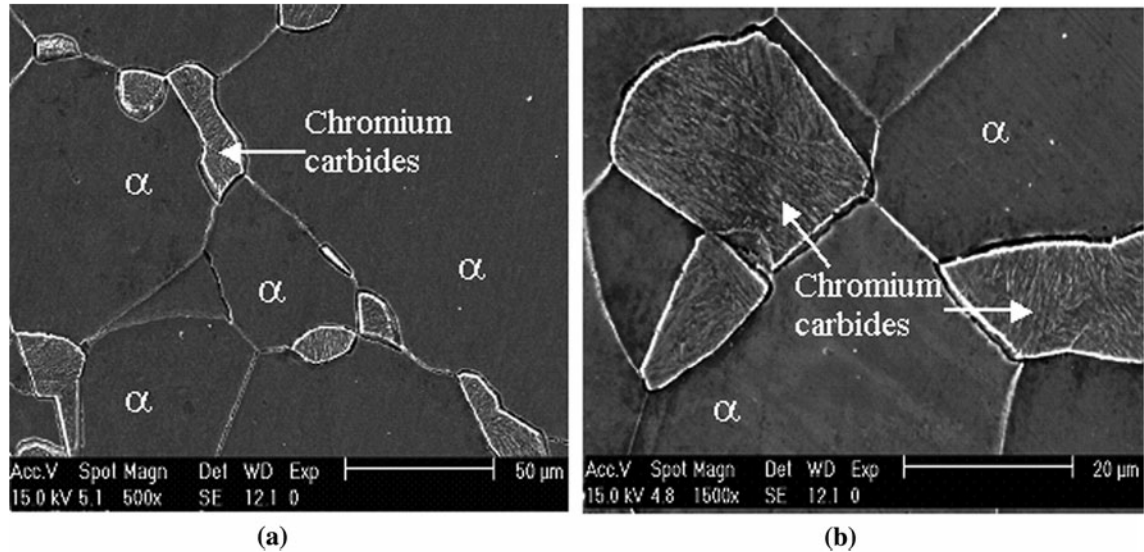


Fig. 4—Structure of the UNS S43000 stainless steel after aging at 773 K (500 °C) for 3 h: very fine decomposition of the quenching martensite into chromium carbides and tempered martensite. (a) Decomposition of quenching martensite into chromium carbides and tempered martensite. (b) Fine decomposition of the quenching martensite into chromium carbides (very weak continuous precipitation (white)) and tempered martensite (higher magnification).

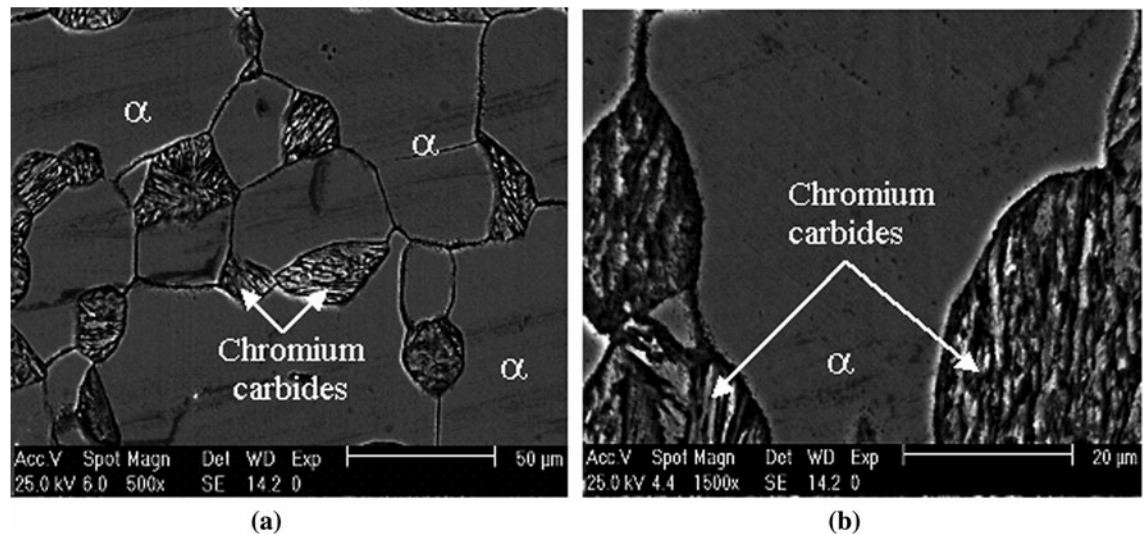


Fig. 5—Structure of the UNS S43000 stainless steel after aging at 873 K (600 °C) for 8 h: resolved decomposition of the martensite into chromium carbides and tempered martensite. (a) Resolved decomposition of aggregates (M + γ_{res}) into chromium carbides and tempered martensite. (b) Resolved decomposition of aggregates (M + γ_{res}) into chromium carbides and tempered martensite (higher magnification).

triple points and in the form of weak continuous interface precipitation in the ferritic grain boundaries (Figure 2(a)). At higher magnification, the aggregates of martensite islets and the ferritic grains of the water-solution-annealed steel appear without any perceptible precipitation (Figure 2(b)). This result conforms to the DL-EPR response ($I_r/I_a = 0.3$ pct), which suggests the absence of the chromium-depleted area. However, the air-solution-annealed microstructure ($I_r/I_a = 10.25$ pct) suggests the presence of chromium-depleted zone. This response conforms to the electrolytic attack morphology of martensite and residual austenite aggregates. This seems to be associated with chromium carbide precipitation, resulting from the annealing process during the slow cooling of the supersaturated martensite. Indeed,

very weak continuous precipitates, likely chromium carbides, are observed to surround the martensite needles in the aggregates. Additionally, very weak continuous precipitates in the ferritic grain boundaries are observed also (Figure 3).

2. Microstructure of aged steel

Aging treatments were performed on samples in the annealed state (annealed at 1373 K (1100 °C) cooled in water) without previous chromium carbides precipitation or chromium-depleted zones.

The evolution of the microstructure starts always at aggregates of the carbon-supersaturated martensite. Thereafter, it continues at the ferritic grain boundaries and within the ferritic grains when heating is prolonged.

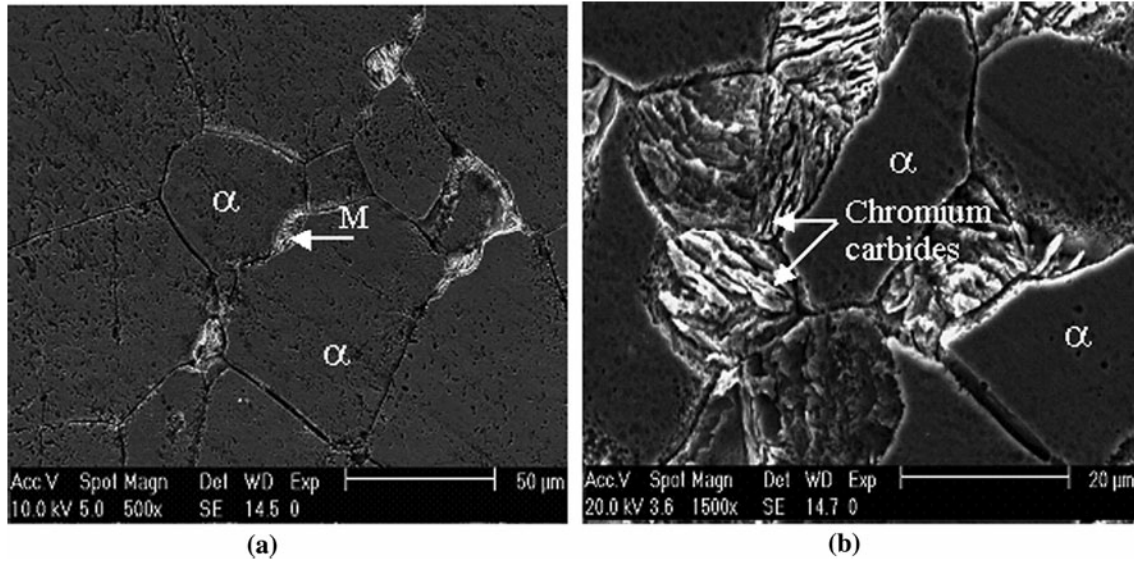


Fig. 6—Structure of the UNS S43000 stainless steel after aging at 973 K (700 °C) for 1 h. (a) Decomposition of the martensite and precipitation of intermetallic phases in ferritic structure. (b) Growth of chromium carbides in the aggregates.

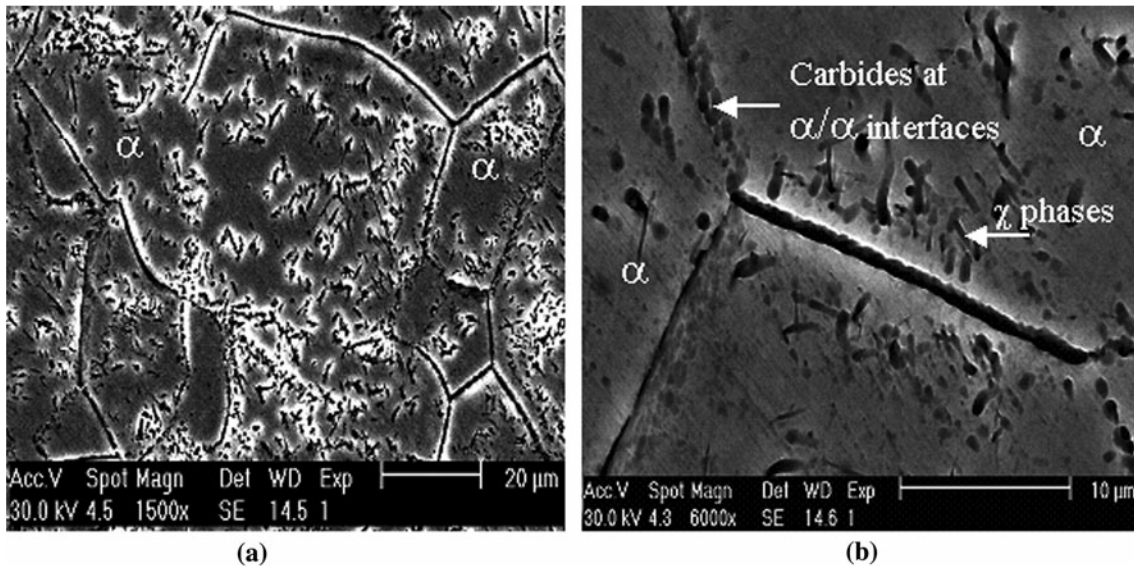


Fig. 7—Structure of the UNS S43000 stainless steel after aging at 1123 K (850 °C) for 30 min. (a) Uniform precipitation of intermetallic phases into ferritic matrix. (b) Total decomposition of aggregates and precipitation of carbides and intermetallic phases into ferritic structure.

The evolution of steel microstructure proceeds at a faster rate at elevated temperatures due to the higher solid-state diffusivity that is favorable for the atomic movement required to achieve the phase changes.

3. Aging at 773 K (500 °C)

Heating at 773 K (500 °C) for less than 3 hours duration did not show any apparent microstructural changes. However, for heating durations exceeding 3 hours, nucleation and growth of chromium carbides resulting from the decomposition of quenched martensite take place in the aggregates. Carbides appear in the form of very weak continuous precipitates (white) at martensite needle interfaces (Figure 4(a)). At higher magnification, this aspect is more obvious (Figure 4(b)).

4. Aging at 873 K (600 °C)

At this temperature, the decomposition of martensite into chromium carbides and tempered martensite is

faster. For a duration of 15 minutes, weak continuous chromium carbide precipitations at the martensite needle interfaces were revealed. These chromium carbides grow when heating is prolonged. However, no precipitation was observed at the boundaries or within ferritic grains during prolonged heating up to 8 hours at this temperature (Figure 5).

5. Aging at 973 K (700 °C)

Heating at this temperature leads to significant reduction of aggregates areas. This reduction occurs gradually to the benefit of chromium carbides and chromium-depleted ferrites. Moreover, uniform precipitation of carbides and intermetallic phases was clearly observed in the ferritic structure (Figure 6). χ phase precipitates after an aging duration of 10 minutes according to the time–temperature–start of precipitation (TTP) diagram given in Reference 1. For higher

Table II. Design of Experiments for Research of the Effect of Significant Factors on the DL-EPR Test Response (I_r/I_a Pct)

Test Number	Conditions of DL-EPR Test				Water Solution Annealing (1 h at 1373 K (1100 °C) and Water Cooling)	Aging (after Water Solution Annealing)					
	dE/dt (mV/s)	[H ₂ SO ₄] (mol/L)	[NH ₄ SCN] (mol/L)	T K (°C)		3 h at 773 K (500 °C)–Air Cooling			2 h at 873 K (600 °C)–Air Cooling		
						I_a (mA/cm ²)	I_r (mA/cm ²)	I_r/I_a (Pct)	I_a (mA/cm ²)	I_r (mA/cm ²)	I_r/I_a (Pct)
1	0.5	0.1	0	293 (20)	0	8.2	<0	0	15.41	1.41	9.14
2	5	0.1	0	293 (20)	0	8.6	<0	0	18.23	<0	0
3	0.5	6	0	293 (20)	78.87	48.34	52.44	108.48	59.21	54.51	92.06
4	5	6	0	293 (20)	36.98	48.27	13.92	28.83	54.31	21.26	39.14
5	0.5	0.1	0.01	293 (20)	25.83	82.71	23.53	28.44	71.25	15.01	21.06
6	5	0.1	0.01	293 (20)	14.86	96.69	10.82	10.85	73.27	14.39	19.64
7	0.5	6	0.01	293 (20)	96.58	118.58	78.3	66.03	174.58	180.33	103.29
8	5	6	0.01	293 (20)	31.67	160.01	93.24	58.27	203.86	162.85	79.88
9	0.5	0.1	0.01	308 (35)	27.02	77.13	19.07	24.72	71.12	21.42	28.51
10	5	0.1	0.01	308 (35)	20.46	76.88	7.75	10.21	80.53	12.34	15.32
11	0.5	6	0.01	308 (35)	57.45	220.19	162.34	73.73	207.53	216.45	104.29
12	5	6	0.01	308 (35)	29.35	357.17	166.44	46.59	510.95	297.18	58.16
13	0.5	0.1	0	308 (35)	8.63	21.93	3.53	16.09	8.99	3.06	34.03
14	5	0.1	0	308 (35)	0	42.4	1.91	4.5	45.04	0.24	0.53
15	0.5	6	0	308 (35)	36.4	266.62	229.72	86.16	211.8	127.90	60.39
16	5	6	0	308 (35)	16.51	218.22	104.71	47.98	152.56	49.23	32.26

Table III. Results of the Analysis by the ANOVA Method

State	Range of Validity	Model	Reliability (Pct)	Significant Factor*
1373 K (1100 °C)–1 h–water cooling (nonsensitized)	$0 \leq dE/dt$ (mV/s) ≤ 0.5 $0 \leq \text{H}_2\text{SO}_4$ (M) ≤ 6 $0 \leq \text{NH}_4\text{SCN}$ (M) ≤ 0.01 $293 \text{ K} (20 \text{ °C}) \leq T$ (°C)	$I_r/I_a = 38.4472 + 5.9794 \cdot \text{H}_2\text{SO}_4$ $- 5.0264 \cdot dE/dt + 1572.87 \cdot \text{NH}_4\text{SCN}$ $- 0.7414 \cdot T$	76.45	H_2SO_4 : 0.0008 dE/dt : 0.0144 NH_4SCN : 0.0687 T : 0.1814
773 K (500 °C)–2 h–air cooling (slightly sensitized)	$\leq 308 \text{ K} (35 \text{ °C})$	$I_r/I_a = 23.0997 + 8.7763 \cdot \text{H}_2\text{SO}_4$ $- 5.4561 \cdot dE/dt + 335.0 \cdot \text{NH}_4\text{SCN}$ $+ 0.0757 \cdot T$	83.22	H_2SO_4 : 0.0000 dE/dt : 0.0098 NH_4SCN : 0.6789 T : 0.8881
873 K (600 °C)–3 h–air cooling (highly sensitized)		$I_r/I_a = 28.7829 + 9.1925 \cdot \text{H}_2\text{SO}_4$ $- 5.7733 \cdot dE/dt + 2032.5 \cdot \text{NH}_4\text{SCN}$ $- 0.256 \cdot T$	88.67	H_2SO_4 : 0.0000 dE/dt : 0.0032 NH_4SCN : 0.0137 T : 0.5910

*Significant factor: determined by the probability of being on the null hypothesis (<0.05).^[28]

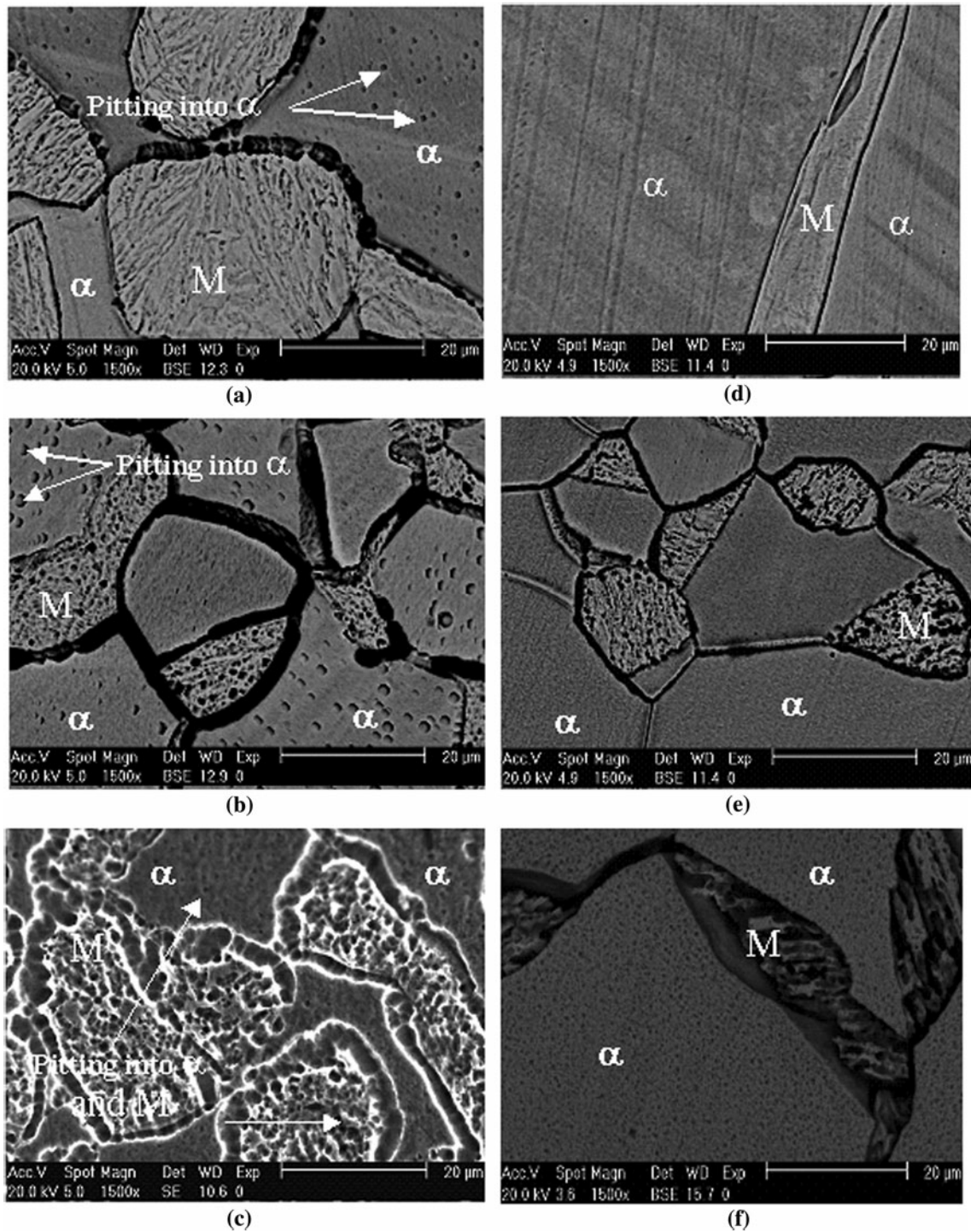


Fig. 8—Effect of the electrolyte concentration on the selectivity of the attack after DL-EPR testing under 2.5 mV/s and at 293 K (20 °C). (a) H₂SO₄ 6 M (uniform attack). Solution annealed steel (non sensitized). (b) H₂SO₄ 6 M (uniform attack). Steel aged at 773 K (500 °C) for 3 h (slightly sensitized). (c) H₂SO₄ 6 M (uniform attack). Steel aged at 873 K (600 °C) for 2 h (highly sensitized). (d) H₂SO₄ 3 M (selective attack). Solution annealed steel (non sensitized). (e) H₂SO₄ 3 M (selective attack). Steel aged at 773 K (500 °C) for 3 h (slightly sensitized). (f) H₂SO₄ 3 M (selective attack). Steel aged at 873 K (600 °C) for 2 h (highly sensitized).

temperature 1073 K (800 °C) and 5 hours duration, σ phase precipitates. Both phases are rich in chromium.

6. Aging at 1123 K (850 °C)

The decomposition of aggregates seems to be total at this temperature after durations as short as 30 minutes.

This decomposition leads to a microstructure characterized by carbides and intermetallic phases in the ferritic structure grain and at the ferritic grain boundaries (Figure 7(a)). Such continuous precipitation of carbides in the ferritic grain boundaries creates dechromized areas with a width equal or less than 4 μm (Figure 7(b)).

B. Optimization of Operating Conditions of the DL-EPR Tests

The DL-EPR test responses obtained with the operating conditions for solution-annealed state (nonsensitized) and aged state (sensitized) are clearly described by the values of the I_r/I_a (pct) ratio, as reported in Table II. These responses indicate aberrant detections of IGC sensitization of the solution-annealed state for operating conditions of test 7 ($I_r/I_a = 96.58$ pct) and of desensitization of the aged steel at 873 K (600 °C) for 2 hours corresponding to test 2 ($I_r/I_a = 0$ pct).

Quantitative analysis of these experimental results, reported in Table III, using the ANOVA method provides the main operating factors controlling the responses of the DL-EPR test. They reveal the significant effect of the electrolyte molarity and the potential scanning rate, regarding the degree of sensitization of the sample. These two factors have the most significant contributions in the DL-EPR test response expressed by the I_r/I_a (pct) ratio as their probability of being at the zero hypotheses, *i.e.*, variances is lower than 5 pct.

Depassivator concentration and electrolyte temperature seem to have much smaller effects on this ratio, as shown in Table III. The reliability of each relationship in the range of variation of each factor is given also in the same table. These values are below 100 pct because the effects of interactions between factors were not taken into account in these relationships. The linear model was selected because it is simple to develop (linear regression) and it offers acceptable reliability (≈ 80 pct) compared with the experimental error (Table III).

The effects of the significant factors on the selectivity of the test were established on the basis of SEM examinations of the attack forms resulting from the DL-EPR tests, for each operating condition. Similarly, the sensitivity of the test in detecting low dechromizations has been established on the basis of the conformity of the attack intensity with the I_r/I_a (pct) value and the microstructural changes. The following results were obtained.

1. Effect of the electrolyte (H_2SO_4) concentration

The results of the experimental design show that the (I_r/I_a pct) ratio increases significantly with the molarity of the electrolyte while other operating conditions are kept constant. The DL-EPR tests, carried out without the depassivator at constant potential scanning rate of 2.5 mV/s and constant temperature of 293 K (20 °C), show that the I_r/I_a ratio varies when the molarity of H_2SO_4 electrolyte varies from 3 to 6 M. The microstructure observations indicate the following.

- (1) For water-solution-annealed (nonsensitized) specimens, the I_r/I_a ratio increases from 0.3 (under optimal condition using 3 M) to 10 pct (under optimal condition using 6 M).
- (2) For aged specimens during 3 hours at 773 K (500 °C) (low sensitized), the I_r/I_a ratio increases from 2.6 (under optimal condition using 3 M) to 13 pct (under optimal condition using 6 M).
- (3) For aged specimens during 2 hours at 873 K (600 °C) (highly sensitized), the I_r/I_a ratio increases

from 33 (under optimal condition using 3 M) to 39 pct (under optimal condition using 6 M).

The rise of the current ratio coupled with the increase of the reactivation current density means that the repassivation kinetics in more concentrated electrolyte (6 M) are much slower and that the attack is uniform. This attack is characterized by a large distribution of corrosion pits for the solution-annealed state (Figure 8(a)) as well as for the low sensitized (Figure 8(b)) and highly sensitized (Figure 8(c)) states. However, the attack is selectively preserved at molarities of the electrolyte less than or equal to 3 M for the three states (Figures 8(d) through (f)). The attack appears in conformity with the microstructure (large chromium-depleted zones) and the response test:

- (1) step for ($I_r/I_a = 0.3$ pct) for the solution-annealed state (nonsensitized),
- (2) dual for ($I_r/I_a = 2.6$ pct) for the low sensitized state, and

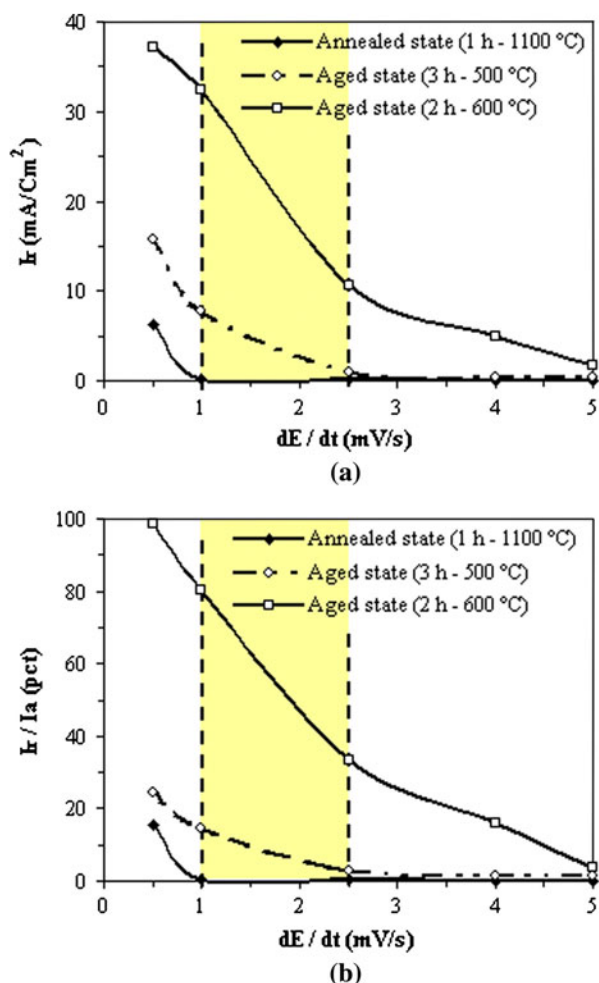


Fig. 9—Influence of the potential scanning rate on the repassivation kinetics and the DL-EPR test response (I_r/I_a pct); test carried out with a 3 M solution of H_2SO_4 at the temperature of 293 K (20 °C). (a) Effect of the potential scanning rate on the repassivation kinetics (I_r in mA/cm²). (b) Effect of the potential scanning rate on the DL-EPR test response (I_r/I_a pct).

(3) ditch for ($I_r/I_a = 33$ pct) for the highly sensitized state.

This concentration of sulfuric acid (3 M) will now be used for the assessment by the DL-EPR test of the degree of sensitization to the IGC of ferritic stainless steels.

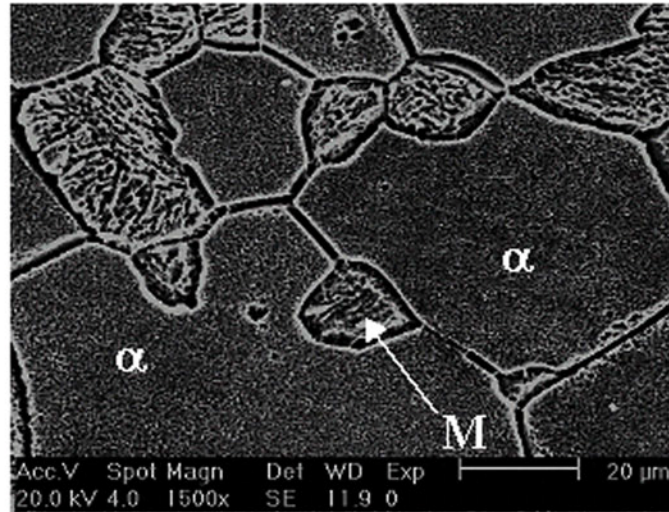
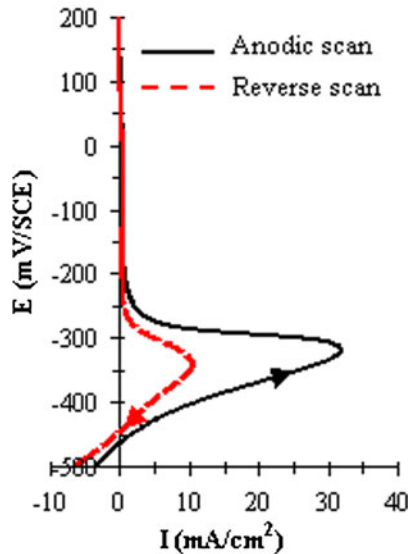
2. Effect of potential scanning rate

The reactivity of surfaces, expressed by the reactivation current density (I_r), for annealed solution state and aged states in the electrolyte without depassivator and composed of a 3 M H_2SO_4 solution at a temperature of 293 K (20 °C), increases when the potential

scanning rate (dE/dt) decreases from 5 to 0.5 mV/s (Figure 9(a)). The same evolution is observed for the DL-EPR test response, expressed by the I_r/I_a (pct) ratio (Figure 9(b)).

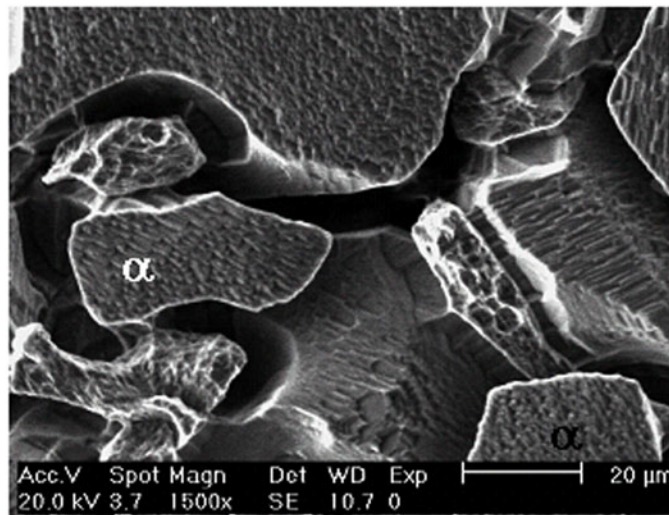
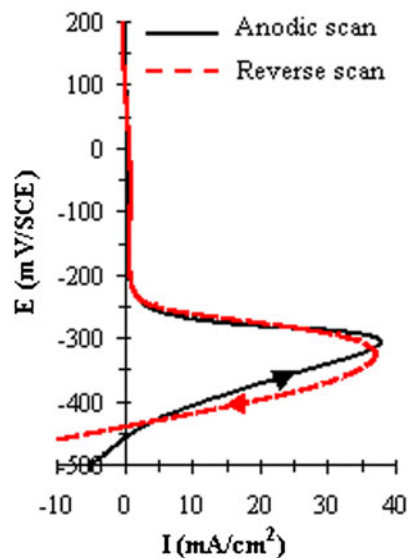
The results of attack form examinations and comparison with the change in the I_r/I_a ratio reveal the existence of three fields of potential scanning rates.

(1) Range of high scanning rates ($dE/dt > 2.5$ mV/s): The test is insensitive to the dechromization phenomena. In this case, the kinetics of electrochemical reactions were very slow compared to the kinetics of potential scanning. That is why the



Structure after DL-EPR test, selective attack

(a)



Structure after DL-EPR test, uniform attack of α phase and dissolution of martensite islets.

(b)

Fig. 10—Effect of the potential scanning rate on the DL-EPR test selectivity. (a) Steel aged at 873 K (600 °C) for 2 h, test carried out with a 3 M solution of H_2SO_4 under $dE/dt = 2.5$ mV/s and at the temperature of 293 K (20 °C). (b) Steel aged at 873 K (600 °C) for 2 h, test carried out with a 3 M solution of H_2SO_4 under $dE/dt = 0.5$ mV/s and at the temperature of 293 K (20 °C).

reactivation current is equal to zero ($I_r = 0$) for low sensitized state without attack.

- (2) Range of intermediate rates ($1 < dE/dt \leq 2.5$ mV/s): The test conserves the attack selectivity and sensitivity concerning the detection of the effect of low dechromizations (Figure 10(a)). In this range, good compatibility between the kinetics of electrochemical reactions and those of potential scanning rates is put in evidence.
- (3) Range of low rates ($dE/dt \leq 1$ mV/s): The DL-EPR test was not able to detect the presence of chromium-depleted areas, since I_r/I_a ratios exceed the sensitization criterion (1 pct) for the solution-annealed steel, which is supposed to be not sensitized. Moreover, under these conditions of very slow potential scanning, the surface reactivity (I_r) is sufficiently high in the H_2SO_4 electrolyte to induce uniform attack of the α phase with an orange skin

morphology and dissolution of the martensite islets (Figure 10(b)).

Accordingly, it can be stated that the rate of $dE/dt = 2.5$ mV/s is the most compatible with the kinetics of electrochemical reactions involved, in chromium-depleted areas. Therefore, this value will be retained for controlling the degree of sensitization to the IGC of ferritic stainless steels by the DL-EPR tests.

3. Effect of depassivator concentration (NH_4SCN)

Results of the experimental design reported in Table II show that the I_r/I_a (pct) ratios, corresponding to the solution-annealed steel that is supposed to be not sensitized, exceed 1 pct with the addition of the NH_4SCN depassivator even with concentrations as low as 0.01 mol/L. These ratios are incompatible with the excessive attacks revealed at grain boundaries without

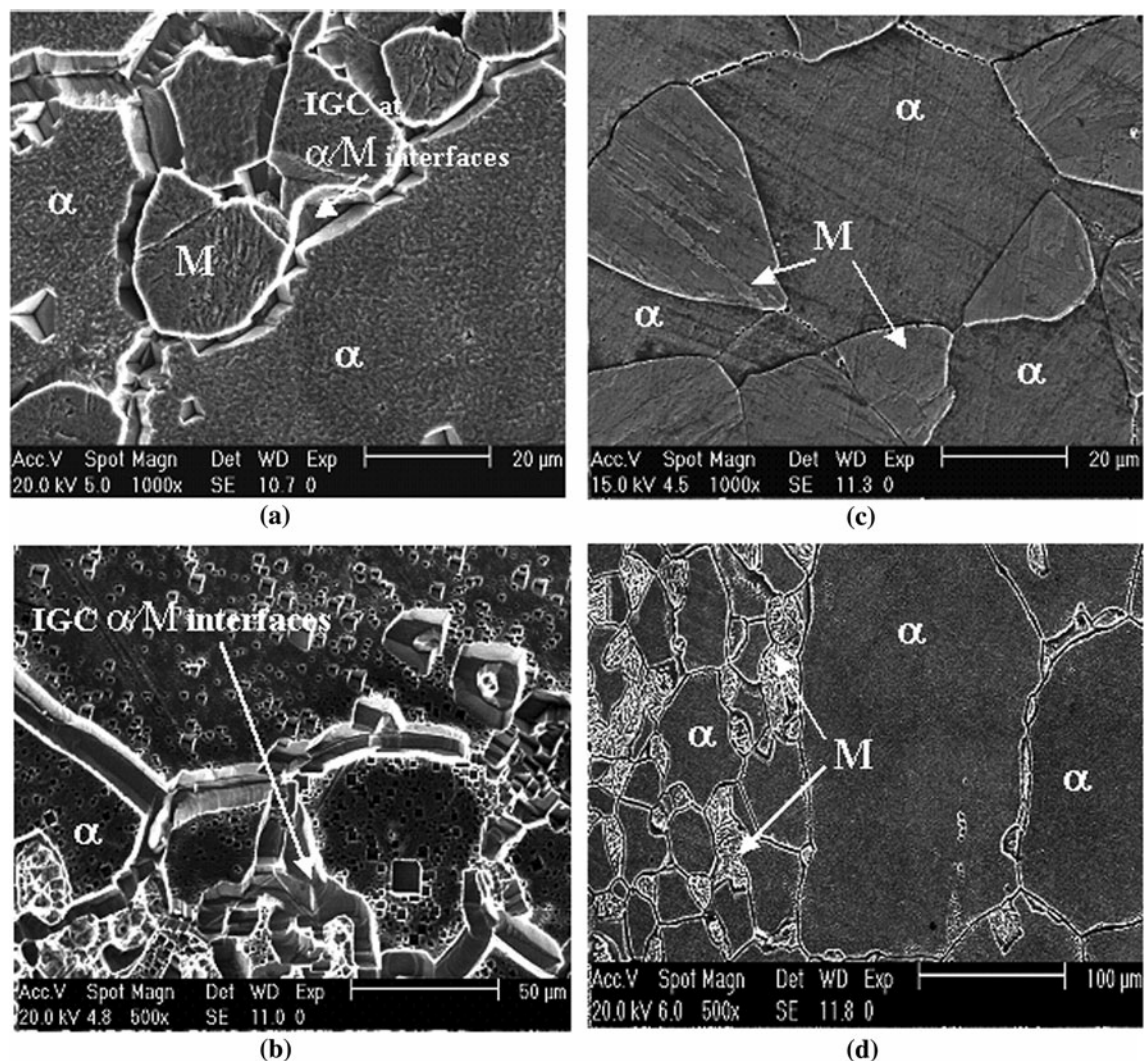


Fig. 11—Influence of the depassivator concentration on the selectivity of the DL-EPR test performed under 2.5 mV/s and at 293 K (20 °C). (a) Structure after DL-EPR test, non selective attack. Test with: H_2SO_4 0.1 M + NH_4SCN 0.01 M solution annealed steel (non sensitized). (b) Structure after DL-EPR test, uniform attack. Test with: H_2SO_4 0.1 M + NH_4SCN 0.01 M steel aged at 873 K (600 °C) for 2 h (highly sensitized). (c) Structure after DL-EPR test, selective attack. Test with: H_2SO_4 3 M solution annealed steel (non sensitized). (d) Structure after DL-EPR test, selective attack. Test with: H_2SO_4 3 M steel aged at 873 K (600 °C) for 2 h (highly sensitized).

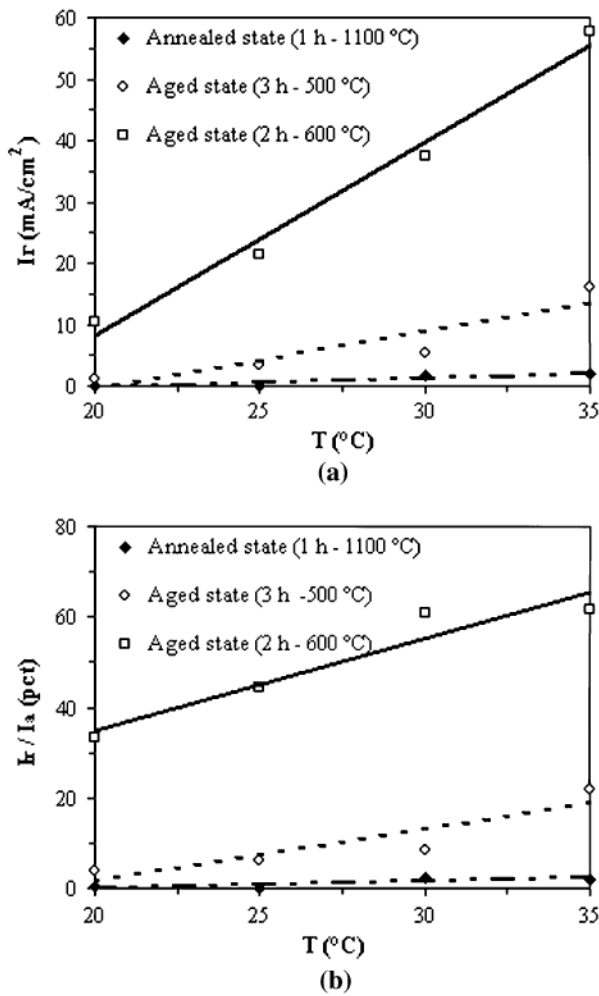


Fig. 12—Influence of the electrolyte temperature on the repassivation kinetics and the DL-EPR test response (I_r/I_a pct); test carried out with a 3 M solution of H_2SO_4 and a potential scanning rate of 2.5 mV/s. (a) Effect of the electrolyte temperature on the repassivation kinetics (I_r in mA/cm²). (b) Effect of the electrolyte temperature on the DL-EPR test response (I_r/I_a pct).

precipitation (Figure 11(a)). The same results are obtained for steel aged during 2 hours at 873 K (600 °C). In this case, the attack appears uniform with corrosion pit distribution (Figure 11(b)). In both cases, the selectivity of the attack was found in the electrolyte without the depassivator (Figures 11(c) and (d)). Accordingly, the depassivator is not recommended for controlling the degree of sensitization to the IGC of ferritic stainless steels by the DL-EPR method.

4. Effect of electrolyte temperature

The density of the current (I_r) and the current ratio (I_r/I_a) of the DL-EPR test, performed in H_2SO_4 electrolyte (3 M) without depassivator and at a potential scanning rate of $dE/dt = 2.5$ mV/s, are not significantly affected by temperature variation between 293 K and 308 K (20 °C and 35 °C) for the solution-annealed state (Figures 12(a) and (b)). However, the test response has changed significantly with increasing temperature for aged states, and the selectivity of the attack decreases in

accordance with the I_r/I_a ratio. Indeed, from 303 K (30 °C), the solution becomes sufficiently aggressive (decreased pH) to cause a uniform attack confirmed by the simultaneous increase of the reactivation current and an almost total dissolution of martensitic islets for aged states (Figure 13). Accordingly, maintaining the electrolyte temperature at 293 K (20 °C) is recommended for controlling the degree of sensitization to the IGC of ferritic stainless steels by the DL-EPR test (Figure 13(a)).

5. Optimal conditions of the DL-EPR test

The ratios of current densities (I_r/I_a pct) deduced by the DL-EPR test, performed on the solution-annealed state (nonsensitized) and states aged during 3 hours at 773 K (500 °C) (low sensitized) and during 2 hours at 873 K (600 °C) (highly sensitized), under optimal conditions are reported in Table IV. Under these conditions, the sensitivity of the DL-EPR test to detect the effect of low dechromization with good reproducibility and to conserve the attack selectivity are demonstrated for the UNS S43000 ferritic stainless steel.

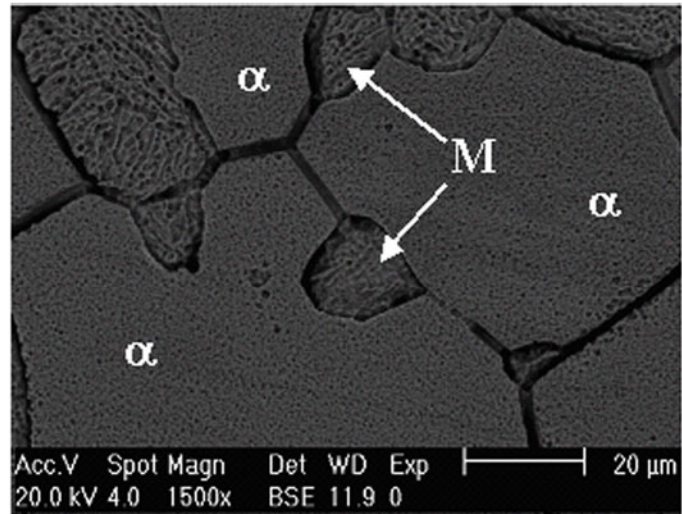
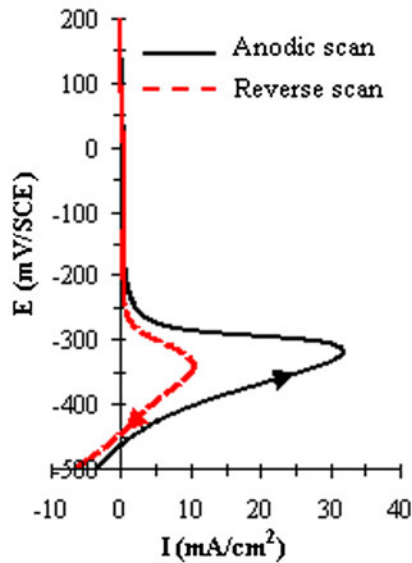
C. Quantitative Assessment of Sensitization to the IGC

The optimal operating conditions of the DL-EPR test are used to assess the degree of sensitization to the IGC of the studied steel aged at temperatures ranging from 673 K to 1123 K (400 °C to 850 °C). The I_r/I_a (pct) ratios and the resulting microstructures after the DL-EPR tests are reported in Table V. These ratios led to establishment of the TTS diagram for the UNS S43000 ferritic stainless steel. The DOS, expressed by the values of the (I_r/I_a pct) ratio, appear in good agreement with the TTP diagram derived from microstructural observations (Figure 14).

IV. DISCUSSION

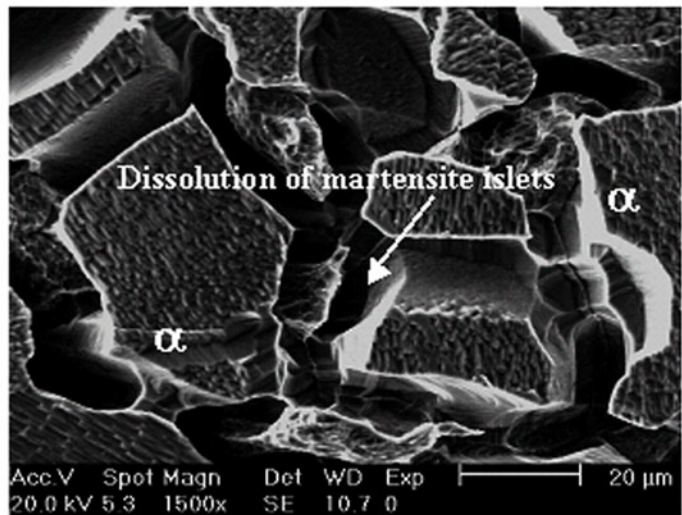
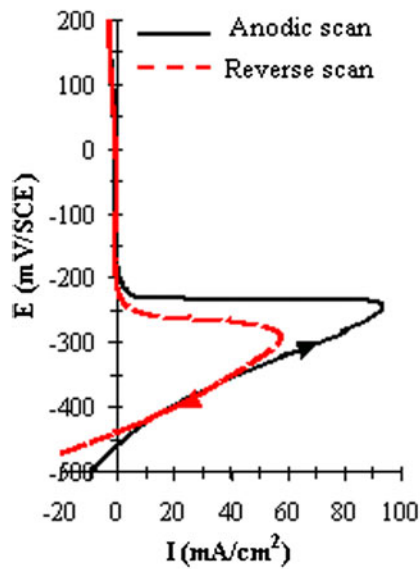
A. Optimization of Operating Conditions of the DL-EPR Test

Results of this study show that, in the same way as for austenitic^[9] and duplex stainless steels,^[8] the sensitivity of the DL-EPR test to detect the effect of low dechromization depends strongly on the operating conditions. It was shown also that the quantitative assessment of the DOS to the IGC, on the basis of the reactivation and anodic current ratio (I_r/I_a pct) values, requires selectivity of the attack that depends itself on the test operating conditions. In fact, the current ratio should reflect the responses of the chromium-depleted areas only (intercrystalline corrosion) during the potential scanning. In addition, taking into account the criteria of sensitivity and selectivity of the test, established by the ANOVA method, reveals the significant effects of various operating factors. This leads to the selection of optimal operating conditions for quantitative and reproducible control of the ferritic stainless steel IGC sensitization. The comparison given in Table VI of the results of DOS of tested specimens under the optimal operating



Structure after DL-EPR test, attack at α/α and carbides /M interfaces

(a)



Structure after DL-EPR test, dissolution of martensite islets

(b)

Fig. 13—Effect of the electrolyte temperature on the DL-EPR test selectivity. (a) Steel aged at 873 K (600 °C) for 2 h, test carried out with a 3 M solution of H_2SO_4 under $dE/dt = 2.5$ mV/s and at the temperature of 293 K (20 °C). (b) Steel aged at 873 K (600 °C) for 2 h, test carried out with a 3 M solution of H_2SO_4 under $dE/dt = 2.5$ mV/s and at the temperature of 308 K (35 °C).

conditions, with results of DL-EPR tests conducted using the operating conditions cited by the literature,^[2,7] highlights the following comments.

- (1) By using the Dowling operating conditions,^[7] the I_r/I_a ratios obtained for the solution-annealed state are abnormally high ($I_r/I_a = 52$ pct): according to the microstructure, this state is not sensitized. In addition, the I_r/I_a ratio, evaluated for the aged specimen during 3 hours at 773 K (500 °C), is quite low ($I_r/I_a = 39$ pct): according to the microstructure, the steel is supposed to be low sensitized in this case. Therefore, the value of the I_r/I_a ratio is expected to be much higher than the value

measured for the solution-annealed state. Under these operating conditions, it was verified that the high values of the reactivation current are the result of uniform attack. Accordingly, the selectivity of the test conducted under these conditions is not respected.

- (2) When the operating conditions selected by Lee^[2] were used, the solution-annealed state is found to be not sensitized ($I_r/I_a < 1$ pct), as revealed by the microstructure examinations. However, the values of I_r/I_a measured for the low sensitized state ($I_r/I_a = 5.5$ pct), corresponding to an aging duration of 3 hours at 773 K (500 °C), and the highly

Table IV. Optimal Conditions for Evaluation by DL-EPR Testing of the IGC Sensitization of the UNS S43000 Ferritic Stainless Steel

Operating Conditions				Sensitized State		
				Nonsensitized State WSA: 1 h at 1373 K (1100 °C) and Water Cooling	WSA + Aging for 3 h at 773 K (500 °C) and Air Cooling	WSA + Aging for 2 h at 873 K (600 °C) and Air Cooling
Electrolyte	T K (°C)	dE/dt (mV/s)	Range of Potential Scanning (mV/ECS)	I_r/I_a (Pct)	I_r/I_a (Pct)	I_r/I_a (Pct)
H ₂ SO ₄ 3 M	293 (20)	2.5	-500 ↔ 200	0.30	2.61	33.14

T: temperature; dE/dt : potential scanning rate; and WSA: water solution annealed.

Table V. Assessing the Degree of Sensitization to the IGC of the UNS S43000 Stainless Steel by the DL-EPR Test Performed in the Optimal Operating Conditions

	Heat Treatment			Microstructural Analysis		IGC Sensitization		
	Temperature K (°C)	Duration (Min)	Cooling Conditions	Interfaces, α/α	Interfaces Carbides, M	I_r/I_a (Pct)	Q_r/Q_a (Pct)	
WSA	1373 (1100)	60	water	step	step	0.30	0.14	
ASA	1373 (1100)	60	air	dual	ditch	10.25	10.98	
Aging (after WSA)	673 (400)	30	air	step	step	0	0	
		60		step	step	0.63	0.53	
	723 (450)	120		step	step	0	0	
		30		dual	dual	1.04	0.51	
		60		dual	dual	1.28	0.78	
		773 (500)	5		dual	dual	4.8	3.53
		10		dual	ditch	10.10	9.6	
		60		dual	ditch	10.12	11.37	
	823 (550)	120		dual	ditch	13.69	7.97	
		180		step	dual	2.61	2.39	
		873 (600)	5		dual	ditch	16.50	18.64
		5		dual	ditch	18.05	19.4	
		15		dual	ditch	18.22	20.52	
		120		dual	ditch	33.14	32.44	
973 (700)	240		dual	ditch	13.26	12.78		
	480		dual	dual	3.75	1.67		
	5		dual	ditch	16.38	12.81		
	60		dual	dual	1.75	1.59		
	180		dual	dual	1.85	2.64		
	5		dual	dual	2.06	2.46		
1073 (800)	30		step	step	0.42	0.31		
	60		step	step	0.02	0		
	120		step	step	0	0		
	1123 (850)	5		step	step	0	0	
	30		step	step	0	0		
	60		step	step	0	0		

ASA: air solution annealed; WSA: water solution annealed; α : ferrite; M: martensite; step: no intercrystalline attack; dual: slight intercrystalline attack; and ditch: high intercrystalline attack.

sensitized state ($I_r/I_a = 7.5$ pct), corresponding to an aging duration of 2 hours at 873 K (600 °C), are very close. This result does not conform to the obvious microstructure differences of the aged states. This test defect is due to the fact that the kinetics of potential scanning ($dE/dt = 4.17$ mV/s) are much faster than the kinetics of the repassivation of chromium-depleted areas. This explanation is supported by the low values of the reactivation current ($I_r = 3$ to 3.2 mA/cm²).

(3) Concerning the optimal operating conditions established in this study (H₂SO₄ 3 M, $dE/dt = 2.5$ mV/s, and $T = 293$ K (20 °C) without depassivator), the sensitivity and selectivity of the test are satisfactory. Indeed, nonsensitization of the solution-annealed state is confirmed ($I_r/I_a = 0.3$ pct). Additionally, sensitization to the IGC of aged states is well marked, with a value of $I_r/I_a = 2.6$ pct for the low sensitized state and a value of $I_r/I_a = 33$ pct for the highly sensitized state. Good results are also

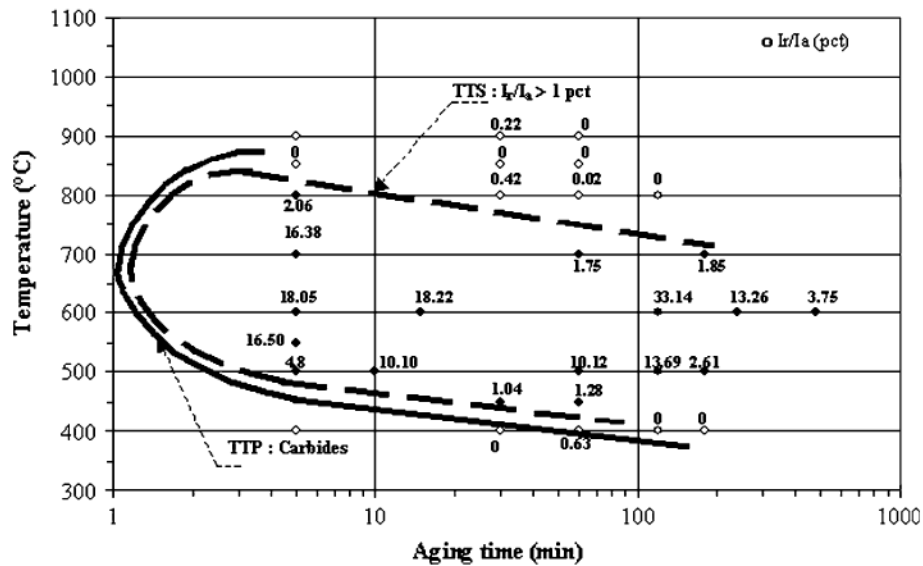


Fig. 14—Interaction precipitation-IGC sensitization of the ferritic stainless steel grade UNS S43000.

Table VI. Comparison of Operating Conditions of the DL-EPR Testing

Heat Treatment	Quantitative Evaluation of IGC Sensitization by the DL-EPR Test	Dowling <i>et al.</i> ^[7] H ₂ SO ₄ 2 M + NaCl 0.5 M; <i>T</i> = 293 K (20 °C); <i>dE/dt</i> = 2.5 mV/s	Lee ^[2] H ₂ SO ₄ 3 N; <i>T</i> = 303 K (30 °C); <i>dE/dt</i> = 4.17 mV/s	This Study H ₂ SO ₄ 3 M; <i>T</i> = 293 K (20 °C); <i>dE/dt</i> = 2.5 mV/s
Solution annealing 1373 K (1100 °C)–1 h– water cooling	<i>I_r</i> (mA/cm ²) <i>I_r/I_a</i> (pct)	154.8 52.1	<0 0	0.1 0.3
Aging 773 K (500 °C)– 3 h–air cooling	<i>I_r</i> (mA/cm ²) <i>I_r/I_a</i> (pct)	142.4 39	3 5.5	1 2.6
Aging 873 K (600 °C)– 2 h–air cooling	<i>I_r</i> (mA/cm ²) <i>I_r/I_a</i> (pct)	201 70	3.2 7.5	10.6 33.1

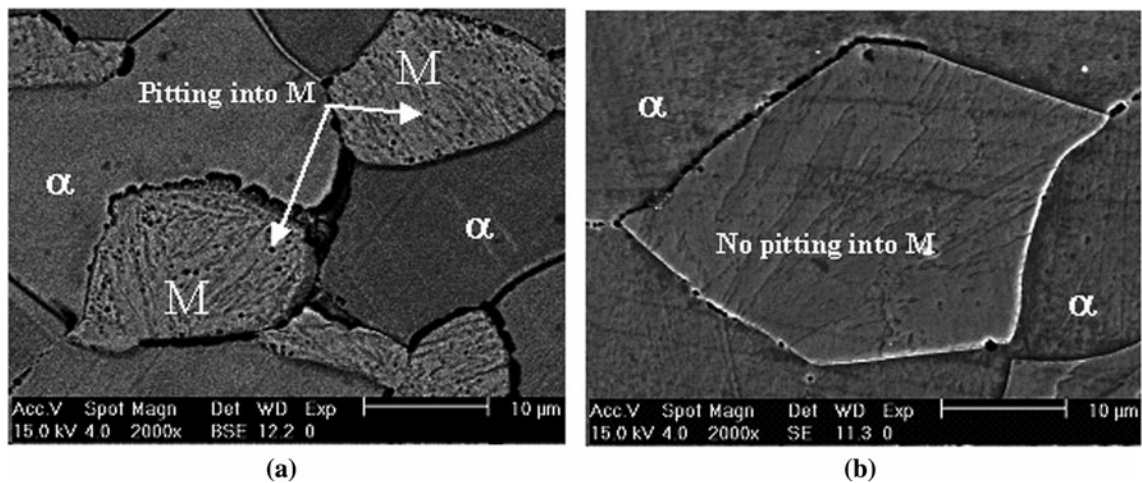


Fig. 15—Effect of the cooling rate on the selectivity of the attack after DL-EPR testing under 2.5 mV/s and at 293 K (20 °C). (a) Structure after DL-EPR test, H₂SO₄ 3 M (uniform attack). Solution annealed steel, air cooling. (b) Structure after DL-EPR test, H₂SO₄ 3 M (selective attack). Solution annealed steel, water cooling.

observed for the attack selectivity of the chromium-depleted areas. This selectivity may be affected when the potential scanning rate decreases

(<2.5 mV/s), or when the severity of the electrolyte increases (>3 M) or when its temperature is high (>303 K (30 °C)). Similarly, the selectivity of the

attack cannot be guaranteed in the presence of a de-passivator even with very low concentrations.

B. Effect of Solution Annealing and Aging Conditions on the IGC Resistance of the UNS S43000 Ferritic Stainless Steel

1. Effect of solution-annealing conditions

The difference in IGC sensitization was clearly observed between the two solution-annealed conditions (water and air cooling) by the DL-EPR test. The test detects a sensitization to the IGC for solution annealing and air cooling ($I_r/I_a = 10.25$ pct) compared to solution annealing and water cooling ($I_r/I_a = 0.3$ pct). This sensitization is attributed to low dechromization following a precipitation of chromium carbides, at the temperature range from 773 K to 1123 K (500 °C to 850 °C), in the aggregates of carbon oversaturated martensite (M), and likely a small amount of residual austenite (γ_{res}), during air cooling. Fine attacks, highlighted in the martensitic islets by metallographic observation, confirm this state of sensitization (Figure 15(a)). However, water cooling was sufficiently fast to prevent any precipitation and consecutive dechromization. This result is confirmed by the absence of any form of attack or selective dissolution after the DL-EPR test conducted on the water-solution-annealed steel (Figure 15(b)).

2. Effect of aging conditions

The superimposition of TTP and TTS diagrams puts in evidence the existence of significant interactions between precipitation-sensitization-desensitization in the IGC of the UNS S43000 ferritic stainless steel under various temperatures and durations due to its microstructural instability. It appears that the oversaturation of martensite by carbon and the fast diffusion of chromium in this phase control the dechromization mechanism and the rehomogenization phenomenon. The dechromization is mainly located in the martensitic islets, which may take place during heating at temperatures ranging from 773 K to 1123 K (500 °C to 850 °C). Indeed, at temperatures below 773 K (500 °C), the chromium diffusion kinetics is relatively slow, so it prevents the rehomogenization of the chromium-depleted zones created during the decomposition of martensite into chromium-depleted ferrite and chromium carbides. Therefore, the sensitization can start after aging for at least 3 hours duration at 773 K (500 °C). This duration coincides with the onset of martensite decomposition. However, aging at 873 K (600 °C) leads to a sensitization that occurs rapidly followed by desensitization after 8 hours when the depleted zones restore the critical chromium content (self healing) that is greater than or equal to 13 pct.^[29] At higher temperatures (>773 K (700 °C)), chromium diffusion is very fast and leads to the rehomogenization of the depleted zones, and the phenomenon of sensitization to the IGC is avoided.

V. CONCLUSIONS

It was verified that electrochemical DL-EPR tests, conducted in the H₂SO₄ 3 M electrolyte without

depassivator at temperature of 293 K (20 °C) with a potential scanning from -500 to +200 mV/ECS at a rate of 2.5 mV/s, can be used as a reliable nondestructive technique for quantitative assessment of ferritic stainless steel DOS. Using the criterion of $I_r/I_a > 1$ pct, the TTS diagram of the UNS S43000 ferritic stainless steel grade was plotted. It shows a good agreement with the results of microstructural evolution that occur during the aging at temperatures ranging from 673 K to 1123 K (400 °C to 850 °C).

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