

Analysis of Super Duplex Stainless Steel Properties as an Austenite-Ferrite Composite

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Abstract

Super duplex stainless steel (SDSS) is considered as a composite formed from a microstructure of an approximately equal mixture of two primary constituents (γ -austenite and α -ferrite phases) and the secondary precipitates (sigma, chi, alpha-prime, etc.). While the formation of these phases affects the properties of SDSS, however there are no rules that govern the relationship. In this work, the relationship between toughness as well as corrosion behavior of SDSS (UNS 32760) and the microstructure constituents has been experimentally investigated, and analyzed in view of the composite principles. Another two stainless steels namely; fully austenitic SASS (UNS N08367) and fully ferritic FSS (UNS S42900) are considered to simulate the constituent's primary components in the composite which are austenite γ and ferrite α phases respectively. Samples of the composite and constituent's steels are first subjected to solution annealing, where the composite steel has a microstructure of γ austenite and α ferrite grains. They were then subjected to similar different isothermal heat treatment cycles, for the formation of secondary phase precipitations within the transformation temperature ranges of each of γ and α primary grains. Impact toughness and corrosion (specific weight loss) tests were conducted on the annealed and isothermally treated samples. The composite rule of the mixtures (ROM) is used to analyze the relationship between the toughness and corrosion properties in the composite SDSS and the SASS and FSS constituent's steels. The analysis indicates that in case of toughness, ROM applies well on the composite and constituents' steels in the solution annealed and in isothermal treatment conditions, where better matching between experimental and calculated results is observed. When applying ROM for corrosion weight loss, a great difference is found between the experimental and calculated results, which is much reduced for solution treated samples ferritic and austenitic temperature ranges of 480°C - 500°C and 700°C - 750°C as for ferrite and austenite respectively.

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Keywords

Duplex Stainless Steel, Impact Toughness, Corrosion Resistance, Modelling

1. Introduction

Duplex stainless steels are formed from a microstructure of an approximately equal mixture of two prime phases, which are γ -austenite and α -ferrite phases, thus they are named austenitic-ferritic stainless steels. The merits of the two phases are granted, which are the high corrosion resistance and ductility of the austenitic phase, plus the high mechanical strength of the ferritic phase [1]-[7]. Their higher strength and corrosion resistance make them suitable for very demanding applications with very aggressive environments, such as oil and gas, seawater and marine engineering, chemical and food industries.

However, the disadvantage of austenite-ferrite mixture is the susceptibility to precipitation of secondary intermetallic phases when exposed to temperatures ranging from 350°C to 900°C [1] [2] [8]-[15]. These precipitates are dangerous intermetallic phases resulting in detrimental effects on impact toughness and corrosion resistance [1] [2] [9] [10]-[15].

Various intermetallic phases (σ -phase, χ -phase, carbides and nitrides) have been found to occur at the α/γ interface [1] [2] [9]-[15]. Other authors [16] [17] suggest that the precipitation of σ -phase is associated with the precipitation of χ -phase. If the alloy with 15 to 75 Cr % is heat treated or used in the temperature range 350°C - 550°C, spinodal decomposition occurs within α matrix as a decomposition into α and $\dot{\alpha}$ [15]. Due to the embritting effect of $\dot{\alpha}$, a serious decrease in toughness occurs, which is known as 475°C embrittlement [15]. Carbide and nitride precipitation in the ferrite-austenitic steels occur in the temperature range 550°C - 800°C. Chromiumrich precipitates, which are formed in the grain boundaries can cause intergranular corrosion and, in extreme cases, even a decrease in toughness [15]. However, after only short times in the critical temperature range, the risk of precipitation is very small for the low-carbon stainless steels.

In this work, super duplex stainless steel (SDSS) grade UNS S32760, which is one of the most superior upper duplex stainless steels, is considered. It has the chemical composition shown in **Table 1**, granting a high corrosion resistance, especially against pitting corrosion. This SDSS has higher tendency to precipitate intermetallic phases than other common duplex and austenitic stainless steels. This is due to the high Cr and Mo contents and high diffusion rates in the ferrite phase. In a previous research, microstructure variation and secondary phase precipitations were studied versus precipitation of UNS N08367 and UNS S42900 (super austenitic and ferritic stainless steels respectively). Secondary phases were identified by the microhardness and X-Ray Diffraction tests. An attempt to correlate between the type of phases versus toughness and corrosion resistance after aging was made.

The relation between the impact and corrosion behavior of solely phased stainless steels; austenitic UNS N08367 and ferritic UNS S42900 and the composite steel SDSS UNS 32760 was studied based on the composite rule of the mixtures (ROM). The ROM which is known as to enable the estimation of composite properties

mposition of tr	ie specifi	nens, F	e = ba	lance.							
	С	Si	Mn	Р	S	Cr	Ni	Mo	Cu	W	Ν
Standard ASTM A240	0.03	1	1	0.03	0.01	24 - 26	6 - 8	3 - 4	0.5 - 1	0.5 - 1	0.3
Feedstock	0.013	0.58	0.66	0.016	0.01	26	7	3.8	0.93	0.6	0.2 - 0.3
Standard ASTM A240	0.03	1	2	0.04	0.03	20 - 22	23.5 - 25.5	6-7	0.75	-	0.18 - 0.25
Feedstock	0.02	0.35	0.86	0.014	0.03	21.5	25.3	6.1	0.7	-	0.2
Standard ASTM A240	0.12	1	1	0.04	0.03	14 - 16	-	-	-	-	-
Feedstock	0.009	0.2	0.2	0.02	0.001	15.7	-	-	-	-	-
	Standard ASTM A240 Feedstock Standard ASTM A240 Feedstock Standard ASTM A240 Feedstock	CStandard ASTM A2400.03Feedstock0.013Standard ASTM A2400.03Feedstock0.02Standard ASTM A2400.12Standard ASTM A2400.12Feedstock0.009	CSiStandard ASTM A2400.031Feedstock0.0130.58Standard ASTM A2400.031Feedstock0.020.35Standard ASTM A2400.121Feedstock0.0090.2	mposition of the specimens, $Fe = ba$ CSiMnStandard ASTM A2400.0311Feedstock0.0130.580.66Standard ASTM A2400.0312Feedstock0.020.350.86Standard ASTM A2400.1211Feedstock0.0090.20.2	mposition of the specimens, $Fe = balance.$ CSiMnPStandard ASTM A2400.03110.03Feedstock0.0130.580.660.016Standard ASTM A2400.03120.04Feedstock0.020.350.860.014Standard ASTM A2400.12110.04Feedstock0.0090.20.20.02	CSiMnPSStandard ASTM A2400.03110.030.01Feedstock0.0130.580.660.0160.01Standard ASTM A2400.03120.040.03Feedstock0.020.350.860.0140.03Standard ASTM A2400.12110.040.03Feedstock0.020.350.860.0140.03Feedstock0.020.20.20.020.01	CSiMnPSCrStandard ASTM A2400.03110.030.0124 - 26Feedstock0.0130.580.660.0160.0126Standard ASTM A2400.03120.040.0320 - 22Feedstock0.020.350.860.0140.0321.5Standard ASTM A2400.12110.040.0314 - 16Feedstock0.0090.20.20.020.00115.7	CSiMnPSCrNiStandard ASTM A2400.03110.030.01 $24 - 26$ $6 - 8$ Feedstock0.0130.580.660.0160.01 26 7Standard ASTM A2400.03120.040.03 $20 - 22$ $23.5 - 25.5$ Feedstock0.020.350.860.0140.03 21.5 25.3 Standard ASTM A2400.12110.040.03 $14 - 16$ -Feedstock0.0090.20.20.020.001 15.7 -	CSiMnPSCrNiMoStandard ASTM A2400.03110.030.01 $24 - 26$ $6 - 8$ $3 - 4$ Feedstock0.0130.580.660.0160.01 26 7 3.8 Standard ASTM A2400.03120.040.03 $20 - 22$ $23.5 - 25.5$ $6 - 7$ Feedstock0.020.350.860.0140.03 21.5 25.3 6.1 Standard ASTM A2400.12110.040.03 $14 - 16$ $-$ Feedstock0.0090.20.20.020.001 15.7 $ -$	CSiMnPSCrNiMoCuStandard ASTM A2400.03110.030.01 $24 - 26$ $6 - 8$ $3 - 4$ $0.5 - 1$ Feedstock0.0130.580.660.0160.01 26 7 3.8 0.93 Standard ASTM A2400.03120.040.03 $20 - 22$ $23.5 - 25.5$ $6 - 7$ 0.75 Feedstock0.020.350.860.0140.03 21.5 25.3 6.1 0.7 Standard ASTM A2400.1211 0.04 0.03 $14 - 16$ Feedstock0.0090.20.20.020.001 15.7	CSiMnPSCrNiMoCuWStandard ASTM A2400.03110.030.01 $24 - 26$ $6 - 8$ $3 - 4$ $0.5 - 1$ $0.5 - 1$ Feedstock0.0130.580.660.0160.01 26 7 3.8 0.93 0.6 Standard ASTM A2400.03120.04 0.03 $20 - 22$ $23.5 - 25.5$ $6 - 7$ 0.75 $-$ Feedstock0.020.350.860.0140.03 21.5 25.3 6.1 0.7 $-$ Standard ASTM A2400.1211 0.04 0.03 $14 - 16$ $ -$ Feedstock0.0090.20.20.020.001 15.7 $ -$

Table 1. Chemical composition of the specimens, Fe = balance.

in terms of its constituent's properties and their volume fractions ROM and its modifications due to microstructural considerations has been approved for mechanical properties but also for some other physical properties such as electrical and thermal ones.

2. Methodology

SDSS UNS S32760 is considered as a composite material, which is formed from two prime phases (constituents); the γ austenite and α ferrite. The γ austenite is the solely primary phase in the super austenitic stainless steel (SASS) UNS N08367, while α ferrite is solely primary phase in the ferritic stainless steel (FSS) UNS S42900.ROM is applied to model the SDSS properties.

3. Experimental Work

3.1. Material

The standard (ASTM) and feedstock's chemical analyses, of the composite SDSS UNS S32760 steel and the constituent's, SASS UNS N08367 γ and FSS UNS S42900 α steels are given in **Table 1**. The feedstock dimensions are $50 \times 40 \times 15$ mm.

3.2. Heat Treatment

Two heat treatments with different targets were subsequently applied on the feedstock specimens, as follows:

1) Solution annealing, is mandatory to dissolve any retained secondary precipitates and to relieve any residual thermal stress in the feedstock specimens. It is used, for the purpose to measure toughness and corrosion weight (CWL) on samples with solely primary γ austenite or α ferrite phases without any secondary precipitates. Solution annealing was performed for the feedstock specimens of the SDSS UNS S32760 and SASS UNS N08367 materials, as per materials specification ASTM A240 [2]. In case of FSS ASTM A240, the feedstock is supplied in the annealed conditions, as mill's final treatment for the ferrite stainless [2].

2) Isothermal heat treatment is conducted to precipitate the secondary precipitates at their formation temperature ranges. This is for the purpose to test toughness and weight loss corrosion test on composite SDSS and constituent's SASS and FSS samples with secondary precipitates. The isothermal heat treatment was conducted on specimens cut from the feedstock pieces, after their solution annealing with size of $2.5 \times 10 \times 5$ mm.

The Isothermal heat treatment of the composite SDDS was conducted, at a low temperature range of 350° C - 500° C (α ferrite transformation) and at a high range of 700° C - 950° C (γ austenite transformation), as shown in **Table 2**. This is for the formation of secondary precipitates at the ferrite and austenitic transformation zones respectively. The treated samples are named hereafter as **SDSS**_{-F} and **SDSS**_{-A}, respectively as indicated in **Table 2**. Similarly, the specimens of FSS and SASS constituent' steels, were also isothermally heat treated for the formation of secondary precipitates at the lower and higher temperature ranges as shown in **Table 3** and **Table 4**, respectively. Based on TTT diagrams of the austenitic and ferritic steels [18] [19], a scheme for the isothermal

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Specimen No.	Temperature (°C)	Time (min)	Specimen No.	Temperature (°C)	Time (min)			
SDSS-I	F (ferrite temperature ra	nge)	SDSS-A (austenitic temperature range)					
10	350	116.0	2	700	1.7			
11	350	166.0	6	700	3.3			
12	350	333.0	7	700	4.7			
8	430	45.0	16	860	0.7			
1	430	55.0	17	860	0.8			
9	430	67.0	18	860	1.0			
13	500	59.0	3	950	1.7			
14	500	75.0	4	950	2.5			
15	500	92.0	5	950	3.3			

Table	2.	SDSS	Isothermal	heat	treatment	conditions
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Table 3. SDSS isothermal heat treatment conditions.							
Specimen No.	Temp (°C)	Time (min)					
1	350	5.0					
2	350	60.0					
3	350	180.0					
4	350	600.0					
5	475	5.0					
6	475	60.0					
7	475	180.0					
8	475	600.0					
9	550	5.0					
10	550	60.0					
11	550	180.0					
12	550	600.0					

Table 4. SASS isothermal heat treatment conditions.

Temp (°C)	Time (min)
750	100.0
750	300.0
750	700.0
860	70.0
860	200.0
860	500.0
925	100.0
925	300.0
925	700.0
	Temp (°C) 750 750 860 860 860 925 925 925

heat treatment conditions applied on the composite and the two constituent's steels is shown in Figure 1.

3.3. Tests

Impact toughness of the notched specimens with a cross section (2.5910 mm^2) was conducted according to ASTM A370. Corrosion weight loss test was conducted by use of ferric chloride solution according to ASTM G48 [20].

4. Results and Discussion

4.1. Measured Impact Toughness and Corrosion Wight Loss

Specimens of solution annealed and isothermal heat treated SDSS composite and both FSS and SASS constituent's steels have been tested for the impact toughness (according to ASTM A370) and specific weight loss corrosion (according to ASTM G48 practice A). This is given in **Table 5** and **Table 6** for solution annealed and isothermally heat treated specimens respectively.

4.2. Modeling of SDSS Properties Applying ROM

In this work, ROM is applied to model the impact toughness of SDSS UNS S32760 as a composite including



Figure 1. Isothmeral heat treatment scheme for SDSS, FSS and SASS.

Table 5. Measured toughness and corrosion specific weight loss (CWL) of solution annealed SDSS, FSS and SASS.

No.	Steel	Heat treatment	Toughness (J)	CWL (mg/cm ²)
19	SDSS	Ferrite temperature range—SDSS. _F Time = 1 min	23.0	0.0169
19	SDSS	Austenitic temperature range—SDSS _{-A} Time = 1 min	23.0	0.0169
13	FSS	Ferrite temperature range (as received)	21.0	11.7252
1	SASS	Austenitic temperature range	25.0	0.0000

 γ -austenite and α -ferrite as constituent phases. A trial is also made to use it for estimating the specific weight loss corrosion.

Since the volume fraction of each of the constituent's γ -austenite and α -ferrite phases in SDSS is 0.50%, then the basic ROM for the relationship between the composite SDSS property and the constituents α -ferrite and γ -austenite phases, can be described by the following equation:

$$Property_{SDSS} = 0.5 \times (Prperty_{FSS} + Property_{SASS})$$
(1)

In this equation, FSS and SASS denote the constituent's FSSUNS S42900 and SASSUNS N08367, respectively. The properties obtained from testing both steels under similar heat treatment conditions as for the composite steel SDSS UNS S32760 are considered representing α -ferrite and γ -austenite phases in the composite steel respectively. In Equation (1), "Property" denotes either toughness or weight loss corrosion. This is considered for three heat treatment conditions on SDSS composite and FSS and SASS constituent's steel. These correspond to solution annealing, similar isothermal heat treatment and different isothermal heat treatment with similar secondary precipitates.

4.2.1. Case of Solution Annealing (SA)

As shown in **Figure 2**, the microstructure obtained in the solution annealed (SA) SDSS is composed of primary austenite γ and primary ferrite α (ferrite matrix is darker), without any secondary precipitate [21].

In this case, ROM for calculating toughness is written as follows:

$$Toughness_{SDSS}^{SA} = 0.5 \times \left(Toughness_{FSS}^{SA} + Toughness_{SASS}^{SA} \right)$$
(2)

In the solution annealing conditions (SA), the measured toughness values of FSS and SASS are 21 J and 25 J



Figure 2. Composite structure obtained for SDSS UN 32760 ingots after solution annealing, showing γ and α phases with absence of any secondary precipitates.

Table 6. Measured impact toughnessand corrosion specific weight loss (CWL) of SDSS thermally treated on different specimen with different temperatures (T) and time (t).

No.	T (°C)	t (min)	Toughness (J)	CWL (mg/cm ²)	No.	T (°C)	t (min)	Toughness (J)	CWL (mg/cm ²)		
	SDS	S-A (austen	itic temperature ra	ange)		SDSS-F (ferrite temperature range)					
2	700	1.7	31.0	0.2769	10	350	116.0	16.0	0.1874		
6	700	3.3	17	0.0153	11	350	166.0	20.0	0.3561		
7	700	4.7	19.0	0.3939	12	350	333.0	27.0	0.1524		
16	860	0.7	22.0	0.1051	8	430	45.0	18.0	0.1218		
17	860	0.8	13.0	1.3540	1	430	55.0	18.0	0.0927		
18	860	1.0	13.0	0.9637	9	430	67.0	6.0	0.0153		
3	950	1.7	23.0	0.0000	13	500	59.0	27.0	0.0751		
4	950	2.5	14.0	0.0600	14	500	75.0	13.0	9.0932		
5	950	3.3	7.0	0.0326	15	500	92.0	22.0	2.4988		
	SAS	SS (austeni	tic temperature rar	nge)			FSS (ferri	te temperature range	e)		
2	750	100.0	24.0	0.1822	1	350	5.0	21.0	29.9938		
3	750	300.0	25.0	0.0267	2	350	60.0	21.3	10.0257		
4	750	700.0	24.0	0.2353	3	350	180.0	20.0	13.4866		
5	860	70.0	22.0	0.0391	4	350	600.0	20.6	7.3370		
6	860	200.0	17.0	0.0000	5	475	5.0	22.3	28.8082		
7	860	500.0	12.0	0.3535	6	475	60.0	22.0	16.7761		
8	925	100.0	16.0	0.0133	7	475	180.0	22.0	10.6719		
9	925	300.0	10.0	0.0363	8	475	600.0	21.6	12.8679		
10	925	700.0	7.0	0.0411	9	550	5.0	21.3	12.5479		
-	-	-	-	-	10	550	60.0	24.0	6.7079		
-	-	-	-	-	11	550	180.0	26.0	7.8159		
-	-	-	-	-	12	550	600.0	24.0	11.9562		

respectively (**Table 5**). These are used in Equation (2) where the calculated toughness of as-annealed SDSS composite, is found similar to the experimentally measured value of 23 J (**Table 5**). Thus the basic ROM is found to describe the measured toughness data in SA case, where there are no secondary phase precipitates in composite and constituent's steels.

4.2.2. Case of Isothermal Heat Treatment (IT)

Table 7 shows the main secondary precipitations detected in SDSS, UNS N08367 which was isothermally treated within the lower 350°C - 500°C, and higher 700°C - 950°C temperature ranges for α -ferrite γ -austenite transformations respectively. In order to investigate the impact of these phases on the toughness and corrosion, two concepts are used for applying ROM. First; considering the composite and constituent's steels as subjected to similar heat treatment conditions. This is although that in the case of isothermally heat treated samples, the composite and constituent's steels may differ in their secondary precipitations. Secondly, considering the composite and constituents steels having similar secondary precipitates after any heat treatment.

First Concept: Similar Isothermal Heat Treatment

In this case, ROM is applied considering composite and constituent's steels as subjected to similar heat treatment conditions, where the secondary precipitates may be different. The role of each precipitation group at low and high temperature transformation ranges is investigated.

Toughness

The role of precipitation group at the austenitic transformation range is investigated based on calculating toughness of SDSS which is subjected to isothermal treatment within the austenitic higher temperature range of 750°C to 925°C (Toughness_{SDSS-A}), in terms of toughness of SASS subjected to similar treatment (Toughness_{SASS}), while the toughness of FSS is that of the solution annealed (Toughness_{SASS}). This is expressed as follows:

$$Toughness_{SDSS-A} = 0.5 \times (Toughness_{SASS} + Toughness_{FSS}^{SA})$$
(3)

The experimental and calculated values are listed in **Table 8**, while they are represented for different heat treatment temperatures in **Figure 3**. The trend is that toughness decreases with increasing temperature, while both experimental and calculated data at any temperature are generally found within the same range. Calculated toughness is found to be 30% higher than the experimentally measured value. It is to be noticed that application of ROM ignored the detrimental effect of each of the formed secondary precipitate which may differ from each other as well as their variation in the similarly treated composite and constituent's steels. The σ phase and which forms at 700°C - 950°C upper temperature range is very harmful to toughness [15] [21]. With increasing isothermal treatment temperature and time, formation of σ phase is associated with χ -phase formation [21], which is even most harmful.Carbide and nitride precipitates occur in the temperature range 550°C - 800°C which are also harmful to toughness. Thus the decrease of toughness with temperature increase could be explained.

However, toughness should be considered as a function of isothermal heat treatment parameters; namely temperature and holding time. This could be represented by a polynomial function of the second degree correlating the toughness with temperature and time as represented by the following equation:

$$Toughness = a \times x^2 + b \times x \times y + c \times y^2 + d \times x + e \times y + f$$
(4)

where, x: temperature (in $^{\circ}$ C) y: time (in min) a, b, c, d, e and f are constants.

	Т٤	able	7.	Ma	ιin	seconda	iry	preci	oitate	s dete	ected	in	IT	samp	les	of	SD	SS	5
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Primary Phase Transformation Zone	Formation Temperature (°C)	Detected Phase	Average Microhardness (HV)
	700 - 950	σ-phase	664 - 674
γ-austenite 700°C - 950°C.	700 - 950	σ + γ_2 -phases	-
	700	σ , iron nitrides and carbide phases	1080
α-ferrite	350	ά	298
350°C - 500°C,	350 - 500	Aged å	372

Table 8	Experimental and calculat	ed Toughness _{SDSS-A} .	
No.	Isothermal Treatment Temperature, [°C]	Experimental Toughness _{sDSS-A} [J]	Calculated Toughness _{SDSS-A} [J]
6	700	17	23.0
7	700	19.0	22.5
17	860	13.0	19.0
18	860	13.0	16.5
4	950	14.0	15.5
5	950	7.0	14.0



Figure 3. Calculated and experimental Toughness_{SDSS-A} at austenitic high temperature (isotherml treatmnt times in min. is indicated between brackets).

The curve fitting process can be carried out to describe both the experimental and model results. Different fitting functions (power, trigonometric and polynomial) are tried and a polynomial function of the second degree is found to be fair for reaching a high regression of more than 97 %. This is shown in **Figure 4**, which presents the case of SDSS_A, where the experimental (a) and model data (b) data for Toughness_{SDSS-A} as well as the deviation % between them (c) are indicated. It is remarkable that each of the plotted curves follows a C-curve similar to the TTT diagram. This is taken as indication that toughness values are affected by the formation of secondary precipitations. At isothermal treatment conditions of 700°C and long holding times, **Figure 4(a)** and **Figure 4(b)**, show that high Toughness_{SDSS-A} values are obtained, while **Figure 4(c)** shows that the percentage deviation between the experimental and calculated results in this case is small.

The role of precipitation group at the ferritic transformation range is investigated based on calculating toughness of SDSS which is subjected to isothermal treatment within the ferrite lower temperature range of 350 to 500°C (Toughness_{SDSS-F}), in terms of toughness of FSS subjected to similar treatment (Toughness_{FSS}), while the toughness of SASS is that for solution annealing (Toughness_{SASS}). This is expressed as follows:

$$Toughness_{SDSS-F} = 0.5 \times (Toughness_{FSS} + Toughness_{SASS}^{SA})$$
(5)

The experimental and calculated values are listed in **Table 9**, while they are represented for different heat treatment temperatures in **Figure 5**. The trend is that toughness remains almost constant with increasing temperature, while both experimental and calculated toughness data at any temperature are generally found within



Figure 4. Second degree curve fitting for (a) experimental Toughness_{SDSS-A}- (b) modelled Toughness_{SDSS-A} (c) deviation% between the model and the experimental results.

Table 9.	Experiment and calculated	$Toughness_{{}_{SDSS-F}}.$	
No.	Isothermal Treatment Temperature, [°C]	Experimental Toughness _{sDSS-F} [J]	Calculated Toughness _{sdss-F} [J]
10	350	16.0	23.0
11	350	20.0	22.5
8	430	18.0	23.7
9	430	18	23.5
14	500	13.0	24.5
15	500	22.0	23.2



Figure 5. Calculated and experimental Toughness_{SDSS-A} at ferrite low temperature (isotherml treatmnt times in min. is indicated between brackets).

similar range. Calculated Toughness is found to be 30% higher than the experimentally measured value, similar to Toughness_{SDSS-A}. It seems that the α ' and the associated aged α ', which form at 350°C - 500°C low temperature range [15] [21], has similar harmful effect to Toughness_{SDSS-F} as Toughness_{SDSS-A}, while it is difficult to separate their effect from each other.

Similar to Toughness_{SDSS-A}, a polynomial function of the second degree is found to describe both the experimental and calculated results of Toughness_{SDSS-F} as shown in **Figure 6**. Again it can be seen that a skewed C-curve, similar to that in the TTT diagram, is obviously remarkable. For the calculated results in **Figure 6**, the C-curve is not complete and is less detrimental.

Corrosion

ROM is also used to estimate corrosion weight loss (CWL) of SDSS in terms of the experimental values of SASS and FSS. CWL is considered, since more accurate results are obtained compared to potentiostatic testing method as reported in a previous work on UNS S32760, UNS N08367 and UNS S42900 steels [21]. Similar to toughens, ROM is applied to investigate the role of each of the austenitic and ferritic *precipitation groups separately*.

The role of precipitation group at the austenitic transformation range is investigated, based on calculating CWL of SDSS which is subjected to isothermal treatment within the austenitic higher temperature range of 750 to 925°C (CWL_{SDSS-A}), in terms of CWL of SASS subjected to similar isothermal treatment (CWL_{SASS}), while the CWL of FSS is that of the as received(CWL^{SA}_{ESS}). This is expressed as follows:



Figure 6. Second degree curve fitting for (a) experimental Toughness_{SDSS-F}- (b) modelled Toughness_{SDSS-F}- (c) deviation% between the model and the experimental results.

$$CWL_{SDSS-A} = 0.5 \times \left(CWL_{SASS} + CWL_{FSS}^{SA} \right)$$
(6)

The role of precipitation group at the ferritic transformation range is investigated, based on calculating corrosion CWL of SDSS as isothermally treated within the ferrite transformation temperature range of 350°C to 500°C (CWL_{SDSS-F}), in term of CWL of FSS subjected to similar treatment (CWL_{FSS}), while CWL of SASS is that of solution annealing (CWL_{SASS}^{SA}). This is expressed as follows:

$$CWL_{SDSS-F} = 0.5 \times \left(CWL_{FSS} + CWL_{SASS}^{SA} \right)$$
(7)

Experimental and calculated CWL, using Equation (7) and Equation (8) for CWL_{SDSS-A} and CWL_{SDSS-F} are given in 10 and 11, respectively (Table 10, Table 11).

The data is presented as shown in **Figure 7** and **Figure 8**, for the cases of CWL_{SDSS-A} and CWL_{SDSS-F} respectively. Experimental and calculated data are generally far from each other. Corrosion is a surface related phenomenon depending on the protective chromium oxide layer [1]. Precipitated secondary phase plays also a detrimental effect on corrosion behavior of such alloys [1] [2] [9] [10]-[15]. However, the deviation between of the experimental and calculated CWL is much decreased by values up to 50% for isothermal treatment at or close to the nose of the austenitic and ferritic transformation ranges of 700°C - 750°C and 480°C - 500°C, respectively.

Table 10. Experimental and calculated $\text{CWL}_{\text{SDSS-A}}$.									
No.	Isothermal Treatment Temperature, [°C]	Experimental CWL _{SDSS-A} (mg/cm ²)	Calculated CWL _{SDSS-A} (mg/cm ²)						
2	700	0.2769	0.1080						
6	700	0.0153	0.0303						
7	700	0.3939	0.1346						
16	860	0.1051	0.0364						

Table 11. Experimental and calculated CWL_{SDSS-F} .

No.	Isothermal Treatment Temperature, [°C]	Experimental CWL _{SDSS-F} (mg/cm ²)	Calculated CWL _{SDSS-F} (mg/cm ²)
10	350	0.1874	9.151218
11	350	0.3561	0
12	350	0.1524	0.897638
8	430	0.1218	8.558423
1	430	0.0927	2.542368
9	430	0.0153	0
13	500	0.0751	0.428288
14	500	9.0932	0
15	500	2.4988	0
17	860	1.3540	0.0169
18	860	0.9637	0.1937
3	950	0.0000	0.0236
4	950	0.0600	0.0350
5	950	0.0326	0.0375



Figure 7. Comparison between the experimental and calculted $\text{CWL}_{\text{SDSS-A}}$ (isotherml treatmnt times in min. is indicated between brackets).



(isotherml treatmnt times in min. is indicated between brackets)

It seems that the associated χ and the aged α phases are much more harmful to corrosion than the earlier formed σ and α at each of the temperature ranges respectively. Moreover, although the volume fraction of the γ -phase is not usually very high, it consumes significant amounts of Cr and Mo from the parent matrix, and the formed γ_2 -phase becomes depleted of these elements. Hence, this usually decreases the pitting corrosion resistance [15] [21].

Second Concept: similar secondary precipitates

ROM is applied based on selection of samples of SDSS composite steel and SASS and FSS constituent's steels, with similar secondary precipitates in each of the primary austenite and ferrite phases. This is regardless of the isothermal heat treatment conditions conducted on each of these steels. This will help examining separately the influence of each group of secondary precipitates formed at lower and higher temperature transformation ranges, *i.e.* ferrite and austenitic transformation ranges respectively. ROM will consider the volume fraction (Vf) and toughness or CWL of each phase (property) as follows:

$$Property_{SDSS} = \left(Vf_{SASS} \cdot Property_{SASS} + Vf_{FSS} \cdot Property_{SASS} + \sum_{i=1}^{i=m} Vf_{Phase(i)} \cdot Property_{Phase(i)} \right)$$
(8)

Where; "i" and "j" are the numbers of phases at any heat treatment.

However, since it is difficult to measure the volume fraction of each of the secondary precipitates, ROM is applied considering secondary precipitates formed at each of austenitic or ferrite transformation zone as a group. **Toughness**

ROM is used to calculate the toughness of SDSS composite after isothermal treatment within the austenitic transformation range in terms of toughness of SASS after similar treatment $(SASS_A)$ and the toughness of FSS in the annealed conditions (FSS_{annealed}). This is in order to investigate separately the influence of secondary precipitates formed in γ -grains within this range. Table 12 gives an example of the results, in case of isothermal treatment at 860°C for 40 s and 200 min; on SDSS and SASS samples respectively where the secondary precipitates formed in γ -phase grains are σ -phase, carbides and nitrides. The annealed FSS has α -phase grains only without any secondary precipitates. The calculated SDSS_A is found to be 17 J, while the experimentally measured value is a higher value of 22 J.

ROM is also used to calculate the toughness of SDSS composite after isothermal treatment within the ferrite transformation range $(SDSS_F)$ in terms of toughness of FSS after similar treatment (FSS_F) and the toughness of SASS in the annealed conditions (SASS_{annealed}). This is in order to investigate separately the influence of secondary precipitates formed in α -grains within this range. Table 13, gives an example of the results, in case of isothermal treatment at 500°C for 59 min on SDSS and at 430°C for 5 min on FS, where α secondary precipitate is formed in α -grains. The annealed SASS has γ -grains only without any secondary precipitate. The calculated $SDSS_F$ is found to be 23.6 J, while the experimentally measured value is a higher value of 27 J.

It is to be noticed that the precipitates in the isothermally treated SASS, do not necessarily have the same size and volume fraction of the constituents steels subjected to the similar heat treatment temperature and time condi-

Table 12. Calcula	tted (ROM) and measured tou	ighness and CWL of $SDSS_A$		
Specimen	SASSA experimental (specimen 6, at 860°C for 200 min	FSS annealed experimental (specimen 13, annealed)	SDSSA experimental (specimen 16, heated at 860°C for 40 s)	SDSSA calculated
Microstructure	×400 equiaxed grains, Secondary precipitates Grain size differs	×400 equiaxed grains, No secondary precipitates Grain size differs	×400 Arrayed grains Secondary precipitates Grain size differs	
Phases	Grains: ≁phase, Precipitates: σ-phase, Carbides, nitrides	Grains: <i>a</i> -phase, equiaxed	Grains: γ-phase,α-phase Precipitates:σ-phase, Carbides, nitrides in γ grains	
Toughness (impact)	=17 J	=21 J	=22 J	$=0.5 \times (17 + 21)$ = 19 J with 14 % deviation
CWL	=Zero mg/cm ²	11.7252 mg/cm ²	0.1051 mg/cm ²	$=0.5 \times (0 + 11.7252)$ = 5.8626 mg/cm ² with almost 100 % deviation



tions. While the shape of γ - and α -grains are equiaxed in case of SASS- and SFSS-constituent's steels, it is in arrayed form in case of SASS. Such differences are not considered in ROM applied, which are reasons for the deviation between calculated and experimental results.

The calculated and measured toughness are more far from each other, in case of higher temperature isothermal treatment within the austenitic transformation range. This may indicate that the influence of σ -phase, carbides and nitrides formed in the γ -grains at this isothermal treatment is higher than the influence of α '-phase formed at the lower temperature range in the α -grains. ROM applied considering the precipitates formed as a group, while the influence of each one is different. Previous report has reported that both σ -phase and α' , formed at austenitic and ferrite transformation ranges respectively, have drastic influence on toughness [1] [2] [9] [10]-[15].

<u>Corrosion</u>

The experimental and calculated corrosion data(CWL), of SDSS with similar secondary precipitates in each of the primary γ - and α -grainsin SASS and FSS respectively was obtained as indicated in **Table 12** and **Table 13** for austenitic and ferrite isothermal treatments respectively. Generally, the calculated and experimental CWL values are greatly far from each other with a deviation of more than 100 %. As an example, in **Table 12**, for SDSS_A, isothermal treated at 860°C for 40 s, experimental CWL is 0.1051 mg/cm², while the calculated CWL is much higher with value of 5.8626 mg/cm². An example is given in **Table 13**, for SDSS_F, isothermally treated at 500°C for 59 min, and the constituent steel FSS_F isothermally heated at 430°C for 5 min while SAAS is annealed. In this case the experimental CWL of SDSS is 0.0751 mg/cm², while the calculated CWL is as high as 14.4041 mg/cm². The deviation between calculated and experimental CWL, is more than 100% in both cases, however it is much higher in case of ferrite transformation. This is an indication that a greater role is played by these condary precipitates on corrosion which is even higher in case of ferrite transformation α' -phase compared to austenitic transformation with σ -phase, carbides and nitrides.

Corrosion resistance in SDSS is not related to mechanical mixing of composite constituents, but depends mainly on resistance of the chromium oxide layer on the surface. Corrosion resistance is also dependent on the individual resistance of the austenite and ferrite primary phases in the duplex structure. It is concluded that ROM, although indicates the mechanical or toughness behavior, however it cannot be applied to calculate CWL.

5. Conclusions

- ROM can be applied to express the toughness of super duplex stainless steel due to precipitation of intermetallic phases. The calculated toughness is 30% greater than the experimental data, due to detrimental effect of the secondary precipitates formed with isothermal heat treatment.
- 2) In case of corrosion, ROM results in a great deviation of more than 100% between the calculated and experimental specific corrosion weight loss. However, this deviation is found to be greatly minimized in case of isothermal heat treatment at or close to the transformation temperature noses.

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