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Abstract

In this second part of the investigation, room temperature mechanical properties and hardness evolution of cast irons with silicon contents ranging from 2.29 wt.% to 9.12 wt.% have been studied and related to structural results from the first part. Increasing silicon content increases ultimate tensile strength and yield stress until a maximum value of 719 MPa at around 5.0-5.2 wt.% silicon for the former and 628 MPa at 5.2-5.4 wt.% silicon for the latter. Brinell hardness remains increasing with silicon content with a maximum value of 396 at 9.12 wt.% silicon. Elongation at rupture shows an opposite evolution and gradually decreases to zero at 5.3 wt.% silicon. This evolution is related to chemical ordering of ferritic matrix (embrittlement effect). Chunky graphite shows apparently no significant effect on the ultimate tensile strength and yield strength in cast irons with silicon contents higher than 4.0 wt.%. However, it has a negative effect on elongation. This result contrasts with the negative effect of chunky graphite on mechanical properties of ductile irons reported in the literature for alloys with silicon contents lower than 3 wt.%. It is suggested that this difference is due to the matrix strengthening effect of high silicon contents which overtakes the detrimental effect of chunky graphite. This study suggests that cast irons with silicon content as high as 5.0 wt.% could be considered for industrial applications when high resistance and some ductility are requested.

Introduction

The interest for increasing silicon content in cast irons for better mechanical properties and higher corrosion resistance has been recognized for long as reviewed by Fairhurst and Röhrig [1]. This has led to the development of the SiMo spheroidal graphite cast irons and there is still strong interest in further improving these latter alloys as shown with recent works [2-4]. Although further increase in silicon content is detrimental for impact properties of cast irons [5], there is a renewed interest for such high-silicon alloys. This is because they show a good combination of tensile properties, a homogeneous microstructure and an expected excellent machinability with low tools wear when compared to conventional ferritic or ferritic-pearlitic alloys at similar levels of tensile strength [3, 6-8]. Also, high silicon contents improve the corrosion resistance of cast irons against various environments [9-10]. However structural characteristics and mechanical properties of high-silicon ferritic cast irons are still unclear for the highest silicon contents, i.e. above 3.5–4.0 wt.%. These uncertainties are contributing to make it difficult the development of this group of ductile iron alloys for different applications as customers commonly require highly controlled and low scattered casting properties.

Solid solution hardening with silicon is well-known in ferritic cast irons and is associated with increased hardness, rupture stress and yield strength while elongation at rupture is progressively

reduced [6, 8, 11-12]. Impact resistance of ferritic ductile irons sharply decreases at increasing silicon content [13-16] though ductility is not reduced as much as it is commonly observed in ductile irons with increasing pearlite contents. When silicon content is further increased above most common practice, Stets et al. [12] and Glavas et al. [17] reported the existence of a maximum value for tensile strength and for yield strength at 4.2–4.3 wt.% silicon in agreement with previous fragmentary knowledge [1]. Above this critical content they reported that both properties rapidly decrease. As reviewed by Wittig and Frommeyer [18], there is a similar decrease in ductility in soft magnetic steels at about 4 to 5 wt.% silicon. There is thus a clear interest in further studying this transition in cast irons and making it clear if the sharp drop in mechanical properties in this range of silicon contents has similar characteristics as those known for silicon steels.

One of the reasons that could make difficult the analysis of the effect of silicon on mechanical testing is that this element is known to favor graphite degeneracy, i.e. decrease in nodularity and also appearance of chunky graphite [12, 19-23, 24]. Besides the influence on the matrix constituents, the nodule count, the size and the roundness of the graphite particles are determining factors with respect to mechanical properties [11, 25-31]. For alloys with silicon content lower than 3 wt.%, it has been reported that chunky graphite decreases elongation at rupture and ultimate tensile strength without affecting yield strength [25, 32-33]. However, the most problematic effect for engineering applications is certainly that chunky graphite does also decrease fatigue resistance [25, 34-39].

In the first part of this study was presented the microstructure of 30 ferritic cast irons containing 3.88 to 6.11 wt.% silicon, and one alloy at 9.12 wt.% silicon. Chunky graphite formation could be observed and antimony was added to some of these alloys to limit the extent of this graphite degeneracy. An index denoted Ω_{Si} that is based on the content of the alloys in silicon, cerium, magnesium and antimony was proposed that shows a critical value around 7 wt.%, over which the amount of chunky graphite increases steadily from zero. In this second part of the study, we report room temperature hardness and tensile properties of the alloys presented in part 1 and also from an additional set of 21 high silicon cast irons prepared similarly for reproducibility check in the highest silicon content range. The present data is also complemented with previous results on nodular cast irons with lower content in silicon, and is compared to literature data. In the discussion of these results, emphasis is put on the role of silicon on hardening the matrix and on the impact of chunky graphite.

Experimental details

In this second part of the study, the room temperature mechanical properties of the 31 ductile iron alloys presented in the first part and of the 21 additional alloys are characterized. These additional alloys were prepared following the same procedure as that described in the first part of this work with some antimony addition to decrease chunky graphite formation. Data from the 25 ferritic alloys reported by de la Torre et al. [8] and from the three Ni-free ferritic alloys reported by Lacaze et al. [15] have been also considered in the present study. The tensile parameters, ultimate tensile stress (UTS), yield strength (Y) and elongation (A), were measured using a Zwick Z250 tensile testing equipment at a controlled strain rate of 0.90 mm/min in the range where Y was determined. This rate was then increased to 24.12 mm/min to determine UTS and A according to the standard ISO 6892-1 A224. Brinell hardness (HBW) was measured with a Instron Wolpert apparatus with a 10 mm diameter sphere and a load of 3000 kg. Vickers micro-hardness (HV) measurements were performed using a Leica WMHT Auto workstation with a diamond pyramid and loads of 10 and 5 g for 10 and 5 s, respectively.

Scanning electron microscopy (SEM) characterization was carried out on the fracture surface of a few representative alloys using a Zeiss Ultra Plus microscope.

Results

Table 1 shows tensile tests and Brinell hardness values together with chunky graphite fractions and the relevant amounts of significant elements for the same 31 alloys than in the first part of this work. All alloys were fully ferritic but alloy #26 which showed 3–5% pearlite because of its low Si content. Table 2 lists the tensile mechanical properties and composition of the 21 additional alloys. Values of the Ω_{Si} parameter that was defined in the first part of this study has been also included in tables 1 and 2 to evaluate the risk of chunky graphite appearance.

Figure 1 shows the tensile strain-stress curves recorded on five alloys with silicon content in between 4.84 wt.% and 5.70 wt.%. For readability, the curves have been shifted along the abscissa as indicated between brackets. In the high silicon range illustrated in figure 1, it is observed that silicon does not significantly affect the Young's modulus, i.e. the slope of the curves in the elastic or pseudo-elastic regime. Increasing the silicon content does increase the UTS value up to 5.21 wt.% (see alloy #25 in Table 1) while it decreases at higher silicon contents. In figure 1, this decrease is clearly related to a marked reduction of A up to a point where there is no plastic deformation for the highest silicon contents. When there is no plastic deformation at rupture, the stress at 0.2% deformation cannot be determined so that the yield stress was here set to zero. This happened for silicon content higher than 5.34 wt.% during the present study (alloy 18-2 in table 2).



Figure 1. Effect of silicon content on the room temperature tensile strain-stress curves for alloys of the first series. For clarity, the curves have been shifted along the abscissa by a value indicated between brackets.

Allow	UTS	Y	Α		f ^A	C	Si	Sb	$\Omega_{ m Si}$
Alloy	(MPa)	(MPa)	(%)	нвм	1 _{CHG}	(wt.%)	(wt.%)	(wt.%)	(wt.%)
1	541	442	10.8	200	0.19	3.15	3.88	< 0.0005	10.15
2	566	470	8.7	208	0.34	3.16	4.11	< 0.0005	10.09
3	595	502	6.2	225	0.31	3.16	4.34	< 0.0005	10.88
4	614	520	7.2	225	0.39	3.10	4.45	< 0.0005	10.96
5	637	544	3.9	234	0.34	3.08	4.66	< 0.0005	11.10
6	565	456	16.7	203	0.00	3.13	3.94	0.0038	7.15
7	587	485	10.2	217	0.30	3.13	4.25	< 0.0005	10.66
8	631	516	10.9	228	0.04	3.10	4.45	0.0037	8.01
9	673	578	2.2	253	0.18	2.93	4.93	0.0028	9.46
10	701	592	2.5	265	0.02	2.93	5.11	0.0035	8.38
11	671	549	4.8	242	0.00	2.95	4.84	0.0040	7.41
12	659	577	1.4	256	0.04	2.91	5.04	0.0036	8.19
13	679	609	1.0	271	0.23	2.69	5.32	0.0040	8.70
14	526	0	0.0	282	0.03	2.72	5.55	0.0044	7.83
15	482	0	0.0	295	0.15	2.75	5.70	0.0039	9.70
16	681	603	2.1	263	0.04	2.71	5.15	0.0031	10.07
17	605	0	0.2	265	0.21	2.65	5.42	0.0031	9.93
18	661	625	0.5	269	0.15	2.75	5.36	0.0034	9.79
19	536	0	0.0	313	0.26	2.76	5.39	0.0029	10.42
20	397	0	0.0	285	0.33	2.77	5.56	0.0025	9.96
21	0	0	0.0	315	0.65	2.64	6.11	< 0.0005	13.29
22	0	0	0.0	310	0.11	2.71	6.14	0.0042	10.16
23	615	517	10.1	225	0.43	2.96	4.61	< 0.0005	10.89
24	636	518	15.5	220	0.00	2.90	4.60	0.0059	5.95
25	706	628	1.3	266	0.06	2.31	5.21	< 0.0005	11.87
26	417	289	22.3	146	0.00	3.67	2.29	< 0.0005	8.14
27	481	0	0.0	295	0.88	2.26	5.75	< 0.0005	11.82
28	0	0	0.0	396	0.17	2.41	9.12	< 0.0005	14.44
29	642	517	14.0	221	-	2.85	4.63	< 0.0005	10.89
30	671	548	8.2	232	0.14	2.93	4.74	< 0.0005	11.53
31	676	563	6.3	240	0.57	2.95	4.87	< 0.0005	11.40

Table 1. Tensile test results (UTS, Y and A), hardness values HBW, fraction of chunky graphite f_{CHG}^{A} (see part I), and carbon, silicon, antimony and Ω_{Si} contents of the same 31 alloys as in part I of this study.

Figure 2 shows the evolution with silicon content of UTS, Y and HBW for the 31 alloys listed in Table 1 and for alloys with 0.92–1.85 wt.% Si [15] and 2.28–3.81 wt.% Si [8] previously investigated. In the graph, the solid vertical line at 3.85 wt.% Si separates present from previous results, with the exception of alloy #26 in table 1 which is on the left of this line. In figure 2, it is observed a good agreement between the various series of alloys and it is seen that HBW increases in the whole range of silicon contents, namely up to 9.12 wt.%. Further, it is noticed that increasing silicon content leads to a continuous increase of UTS and Y up to about 5.2 wt.% Si for the former and 5.2–5.3 wt.% Si for the latter. It thus appears a maximum in both UTS and Y which relates to the decrease in A as stressed in the presentation of figure 1 and not to a decrease of the strength of the matrix. The vertical interrupted line in figure 2 corresponds to the maximum in UTS, thus emphasizing that the maximum in Y could lie to slightly larger Si content. However, the scattering

of the data plotted in figure 2 makes it difficult to ensure such result and this was the reason for preparing the second series of alloys.

Alloy	UTS	Y	Α	С	Si	Sb	Mg	Ce	$\Omega_{ m Si}$
	(MPa)	(Mpa)	(%)	(wt.%)	(wt.%)	(wt.%)	(wt.%)	(wt.%)	(wt.%)
1-2	719	576	6.2	2.91	4.98	0.0025	0.037	0.0060	9.25
2-2	695	592	2.0	2.85	5.20	0.0032	0.034	0.0055	8.46
3-2	709	585	3.0	2.85	5.12	0.0041	0.035	0.0060	8.00
4-2	651	605	0.6	2.60	5.27	0.0047	0.031	0.0055	7.11
5-2	681	590	1.5	2.71	5.24	0.0035	0.038	0.0071	9.24
6-2	622	613	0.2	2.62	5.24	0.0033	0.033	0.0074	8.91
7-2	671	617	0.7	2.69	5.42	0.0037	0.036	0.0070	9.02
8-2	708	574	5.4	2.88	4.93	0.0028	0.037	0.0057	8.88
9-2	707	595	2.5	2.89	5.11	0.0030	0.036	0.0054	8.71
10-2	687	583	2.2	2.86	5.14	0.0038	0.033	0.0056	7.89
11-2	681	607	1.1	2.66	5.32	0.0045	0.029	0.0051	6.95
12-2	699	597	2.1	2.69	5.15	0.0033	0.034	0.0070	8.81
13-2	681	617	0.9	2.59	5.29	0.0032	0.034	0.0068	8.96
14-2	651	623	0.4	2.64	5.36	0.0039	0.033	0.0066	8.35
15-2	705	558	6.4	2.88	4.99	0.0027	0.037	0.0060	9.11
16-2	672	577	1.7	2.82	5.24	0.0032	0.034	0.0057	8.57
17-2	688	559	4.5	2.86	4.84	0.0042	0.035	0.0058	7.59
18-2	467	0	0.0	2.59	5.34	0.0049	0.033	0.0057	7.32
19-2	648	584	0.9	2.70	5.23	0.0035	0.037	0.0071	9.12
20-2	617	602	0.3	2.67	5.35	0.0037	0.036	0.0068	8.89
21-2	603	0	0.0	2.60	5.38	0.0035	0.035	0.0076	9.20

Table 2. Tensile test results (UTS, Y and A), carbon, silicon and Sb contents and value of Ω_{Si} for the 21 additional alloys.



Figure 2. Evolution of UTS, Y and HBW with Si content for alloys prepared in this work (right side from the solid vertical line) and for other ferritic ductile iron alloys (left side from the solid vertical line) [8, 15]. The vertical interrupted line corresponds to the maximum value in UTS.

Notice that the evolutions of tensile strength and yield strength with silicon content are not parallel below the limit defined by the interrupted vertical line. The rate of increase in yield strength appears slightly larger than that of UTS, leading to a Y/UTS ratio which achieves a maximum value of 0.89 for the Si content corresponding to the maximum in UTS. Beyond their respective maximum, UTS and Y decrease sharply to zero as silicon content is further increased.

A continuous increase in tensile strength and yield strength with silicon content in ductile cast irons was described by Björkegren et al. [6] but their study was limited to a maximum silicon content of 4.2 wt.%. On the other hand, a maximum value of tensile strength for high silicon ductile irons was reported by Stets et al. [12] at 4.3 wt.% in silicon for alloys cast in both Y2 and Y4 keel-blocks. These authors also reported a maximum value of yield strength at a silicon content around 4.6 wt.%. Such a difference between the maximum strength and yield stress seems to be similar to the effect observed in figure 2 though at lower silicon content.

Values of tensile strength and yield strength in figure 2 have been plotted again in figure 3 together with data from Stets et al. [12] and from Glavas et al. [17] for standard Y2 blocks and from Björkegren et al. [6] for cast bars 25 mm in diameter. It is observed an excellent agreement among the various sets of data for silicon contents up to 4.6 wt.% (UTS) and 5.0 wt.% Si (Y). However, data from the present work clearly shift the maximum in UTS and Y at a silicon content significantly higher than reported by Stets et al. and Glavas et al. In all studies, UTS and Y decrease anyway sharply to zero when the silicon content is further increased beyond the respective maximum. Note that it seems Stets et al. [12] set Y equal to UTS when the yield stress could not be evaluated while it was set to zero in the present work as indicated above.



Figure 3. Comparison of UTS and Y values from the present work with data by Stets et al. [12], Glavas et al. [17] and Björkegren et al. [6].

As mentioned previously, the additional group of 21 alloys was used to further characterize the range where the UTS and Y maximum values have been found with the first series of alloys. The results of both series are compared in figure 4. It is seen that the UTS and Y maximum values are in the range 5.0–5.2 wt.% and 5.2–5.4 wt.% Si, respectively. In the range 5.2–5.4 wt.% Si, UTS decreases slowly for most of the values but some very low values are also observed. These

evolutions are in line with what could be expected from the description of the strain-stress curves in figure 1, i.e. strongly related with the coupling between A and UTS values in this high silicon range.



Figure 4. Comparison between UTS results from first and second series of high silicon alloys prepared in this work.

Figure 5 presents the evolution of the elongation at rupture (A) with the silicon content for the same three series of data than those used for figure 2. In agreement with the so-called embrittling effect of silicon in cast irons, it is seen that A starts decreasing at a quite low amount of about 2.2 wt.% Si and is zero at about 5.3 wt.% Si. In the low range of silicon contents, the values in figure 5 agree very satisfactorily with those reported by Björkegren et al. [6] who reported a decrease in A from 20% at 2.2 wt.% Si to 15% at 4.2 wt.% Si. In the high range of silicon contents, the evolution of A seen in figure 5 totally agrees with values reported by Friess et al. [4] with a decrease from 17% at 4.0 wt.% Si to 0% at a Si content of 5.3 wt.%. Note that the critical silicon content at which A becomes zero relates exactly with the position of the maximum of Y in figure 4.

Such a continuous transition between fully ductile and fully brittle materials strongly suggests it depends on ordering of the ferritic matrix associated with silicon addition. In the Fe-Si system, the bcc phase (ferrite) becomes B2 (FeSi) and then DO₃ (Fe₃Si) ordered at increasing contents of silicon, see figure 6. The transition between the disordered ferrite and the start of B2 ordering occurs at a silicon content that increases with temperature, up to the liquidus. It is at about 10 at.% Si at 500°C and 15 at.% at 1000°C. The DO₃ ordering starts at slightly higher silicon contents. For temperatures lower than 700°C, the DO₃ ordering leads to the appearance of a two-phase B2+DO₃ domain which extends to a maximum of 14 at.% Si.



Figure 5. Evolution with silicon content of the elongation at rupture A. The dotted curve represents the evolution of the quantity $25 \cdot (1-S)$ as calculated at $300^{\circ}C$ (see text). The full symbols are for samples with no chunky graphite and the partially filled symbols are for samples with some chunky graphite as indicated in the caption (see discussion section).



Figure 6. Schematic of the Fe-Si phase diagram in the composition range of ordering of the bcc ferrite (redrawn from Wittig and Frommeyer [18]).

The B2 ordering has been described in the assessment of the Fe-C-Si system [40] with a twosublattice model. At low silicon content, the bcc phase is disordered and the two sublattices have the same composition. Above a certain content in silicon which increases with temperature their compositions start differing and the maximum ordering is achieved when Si atoms fill one sublattice and Fe atoms the other one (FeSi composition). An ordering parameter can be defined as $S = (y_{Fe}^1 - y_{Fe}^2)/(y_{Fe}^1 + y_{Fe}^2)$, where y_{Fe}^i is the fraction of sites occupied by Fe in sublattice "i", with S varying from 0 to 1 when ordering increases. The evolution of S has been calculated with Thermocalc and the TCFE8 database in the binary Fe-Si system as no effect of carbon can be expected owing to its very low solubility in ferrite. Also, the calculation was performed at 300°C as it is not expected that any atomic movement needed for ordering could take place at lower temperature (as a matter of fact, most of the experimental data in the literature are limited to temperatures above 500°C). The quantity 25·(1-S) has been plotted with a dotted line in figure 5, where the constant 25 is for scaling the curve with the value of A at low silicon content. It is seen that the predicted evolution of 25·(1-S) follows satisfactorily the change in A, thus confirming the close relation of the decrease of A with ordering. At high silicon content, the abrupt decrease to zero of the experimental values at 5.3 wt.% Si must be related to DO₃ ordering which was not modelled. Similarly, the embrittling effect of silicon has been reported for steels when the silicon content exceeds 3.0–3.5 wt.% [41] where it leads to the same abrupt decrease of UTS and Y values at the same time as A drops to zero.

Ordering is also known to be related to an increase in hardness, and this is what the measurements of HBW in figure 2 show. To further investigate this relation, it was decided to perform microhardness measurements. Preliminary investigation with micro-hardness Vickers measurements were carried out under a load of 5 g during 5 s on alloys #6 (3.94 wt.% Si) and #15 (5.70 wt.% Si). Such a load is large enough to obtain measurable length of the indent diagonals (around 5 μ m) and small enough to let room for further prints. Because build-up of silicon microsegregation during solidification, it is expected that the local hardness of the ferritic matrix changes from place to place. More precisely, the silicon content and hence hardness should be higher close to graphite nodules than away from them as already reported by Alhussein et al. [16]. Figure 7 shows examples of HV measurements performed along lines between two nodules for alloys #6 and #15. Notice that HV evolution is similar for both samples, achieving the highest values in locations close to graphite nodules as expected. However, HV differs between these two samples, being higher in case of the alloy with the highest silicon content.

HV micro-hardness measurements have then been made systematically on all alloys of the first series using a load of 10 g applied during 10 s. In each sample, ten prints were randomly performed in areas away from graphite particles and ten others in areas close to them. Figure 8 shows the evolution with silicon content of the average HV values, with gray and open symbols corresponding to measurements respectively close to and away from graphite nodules. The standard deviation on individual measurements increases from 10-15 HV at low silicon content to about 30 HV for 6 wt.% silicon. As expected, it is observed that HV values increase as silicon content does and that they are higher in areas close to graphite nodules than in those far from them. It is noticeable that the difference in HV values between areas close to nodules and those remote from them nearly vanishes at silicon contents larger than 5.3 wt.%, and this is seen to relate to a slope change (pointed with the arrow in figure 8) has been also reported by Wittig and Frommeyer [18] in their study of Fe-Si ribbons at exactly the same silicon content. These authors related it to the beginning of DO₃ ordering.



Figure 7. Micro-hardness indent lines (column to the left) and corresponding evolution of HV (column to the right) in alloy #6 (3.94 wt.% Si, top row) and in alloy #15 (5.70 wt.% Si, bottom row).



Figure 8. Evolution of HV micro-hardness with silicon content in the 31 alloys of the first series. Gray dots were measured in areas close to nodules and open ones correspond to areas away from them. The arrow points on a slope change related to the start of DO₃ ordering (see text).

Discussion

This discussion is mainly intended to find out if chunky graphite has any effect on room temperature tensile properties of high silicon spheroidal graphite cast irons and to analyze the effect of silicon on these properties. In these analyses, it is considered that the very small amount of Fe-Si-C and Ti-C particles reported in part I of the present study could not account for the changes in these properties which are closely related to the increase in silicon content per se.

Figure 9 shows a number of strain-stress curves to discuss the effect of silicon and of chunky graphite. The curves for alloy #26 with 2.29 wt.% Si and alloy #24 with 4.60 wt.% Si were plotted to illustrate the effect of silicon in samples without chunky graphite. It is seen that an increase of silicon content leads to important increase of UTS and Y while elongation at rupture is reduced from 21–22% to 15–16%. SEM micrographs of the rupture surface of these samples are shown in figure 10. For alloy #26, figures 10a and 10b show two views of a typical ductile fracture that could be expected from an alloy with comparatively low silicon content. In the case of alloy #24, figures 10c and 10d show quite different features: i) figure 10c shows a mainly ductile rupture with some cleavage facets in the upper right of the micrograph; ii) figure 10d shows mainly intergranular rupture (note that several intergranular cracks may be observed) with however some cleavage facets in the right side of the micrograph. These observations suggest that the ductile area in figure 10c accounts for the large rupture strain showed by alloy #24 while the intergranular and brittle features in figure 10d accounts for the decrease in elongation at rupture as compared to alloy #26.





For investigating the effect of chunky graphite at intermediate silicon content, alloy #1 with 3.88 wt.% Si and 0.19 area fraction of chunky graphite was selected. The corresponding strain-stress curve in figure 9 shows that UTS is increased while A is significantly decreased with respect to alloy #26. Figure 5 shows in fact that this A value is the smallest one of the alloys with 3.8–4.0 wt% silicon and that there is a close relationship with the presence of chunky graphite. The rupture surface of alloy #1 is shown at two magnifications in figure 10 (e and f) where it is seen a ductile rupture in areas containing nodules and those containing chunky graphite. A higher content in

chunky graphite would have quite certainly further decreased the elongation at rupture and thus the UTS value as reported in the literature.



Figure 10. SEM micrographs of the fracture surface of tensile samples of alloys #26 (a and b), #24 (c and d) and #1 (e and f).

Finally, the tensile strain-stress curves for alloys #23 (4.61 wt.% Si) and #24 (4.60 wt.% Si) in figure 9 allow studying the effect of chunky graphite for alloys with very similar and high silicon contents. These curves show the yield stress is the same for both alloys while the UTS values are very similar. The weak effect seen for the UTS value is much lower than the one reported in studies on ductile irons with comparatively low silicon contents [35-37]. Thus, the main difference between

these two alloys is the value of A which is significantly lower for alloy #23 containing chunky graphite than for alloy #24 with no graphite degeneracy.

SEM observation of the fracture surface of alloy #23 is illustrated with the micrographs in figure 11 where areas with nodules mainly show cleavage facets and intergranular rupture (see the upper right zone of figure 11a) which appear similar to those observed in alloy #24. Areas with chunky graphite show surfaces with some level of deformation of the metallic matrix (figure 11b) that could indicate a ductile fracture.



Figure 11. SEM micrographs of the fracture surface of tensile sample of alloy #23.

It appears somehow paradoxical that areas with chunky graphite show a ductile-like fracture when they have certainly solidified early during the eutectic reaction, thus at high silicon content as this element segregates negatively during solidification. It has been verified that the fracture surface in areas with chunky graphite in alloys containing slightly more than 5 wt.% Si also shows characteristics similar to those illustrated in figure 11 while elongation was lower than 2.6%. With further increase in silicon content, observations made on alloy #27 with 5.75 wt.% Si show a fully cleavage fracture in the extensive areas with chunky graphite (see figure 12) that is likely the reason for the lack of any measurable value of Y and A in this alloy and in those with the highest silicon contents.

As expected, increasing the silicon content of the alloys promotes a change in the fracture surface from ductile to fragile in areas with nodules. However, there is no such change in areas containing chunky graphite at silicon contents lower than 5.5–5.7 wt.%. This suggests that the small distance between graphite strings in the chunky graphite cells leads to early damage (at low strain, i.e. in the elasto-plastic regime) of the matrix during tensile testing. Because of this damage, the actual stress on the loading surface increases at lower overall strain than for samples without chunky graphite. This schematic would explain the role of chunky graphite on tensile properties of cast irons: i) for low-Si ductile irons this leads to a decrease of the UTS and A values because the matrix is ductile and highly sensitive to early damage; ii) for high-Si ductile irons, the high strength of the matrix in areas with nodules delays the propagation of the damage initiated in the chunky graphite areas, leading to a much lower sensitivity to graphite degeneracy.



Figure 12. SEM micrographs of the brittle fracture observed in areas with chunky graphite of alloy #27.

That chunky graphite has apparently no effect on UTS and Y was confirmed by plotting the same data as in figure 2 but differentiating the results with grey shade according to the amount of chunky graphite. This is shown in figure 13 where it is effectively seen that the increase of UTS and Y with increasing silicon content up to 5 wt.% does not seem to be affected by the amount of chunky graphite. This result is made evident in case of alloys #23 and #31 which contain high level of chunky graphite and are represented with black solid symbols well within the range of values for other alloys with lower chunky graphite fractions. In the same area of the graph, it can also be seen dots corresponding to alloys #24 and #11 marked with arrows and plotted with open circles, i.e. in which no chunky graphite was detected. Finally, it may be noticed that alloys defining the maximum in UTS contain a chunky graphite area fraction equal to or lower than 0.06 (palest gray solid dots in figure 13).



Figure 13. Values of UTS and Y for alloys with different amounts of chunky graphite.

The apparent lack of effect of chunky graphite on tensile strength is also observed when comparing UTS data from the first and second series of alloys represented with open circles and crosses in figure 4. The 21 alloys of the second series show Ω_{Si} values in the range 6.95–9.25 wt.% (see table

2) while this parameter ranges from 7.41 wt.% to 11.87 wt.% for alloys from the first series that define the maximum in UTS. As it has been indicated in the first part of this study, the critical value of Ω_{Si} for standard Y2 keel-blocks is around 7 wt.%, below which chunky graphite formation is not expected. By considering figure 11 in part I, alloys from the second series show Ω_{Si} values that would lead to chunky area fractions of about 0.00 to 0.18 while 0.02 to 0.65 would be expected for the corresponding alloys from the first series. SEM inspections made on fracture surfaces of alloys #4-2 ($\Omega_{Si} = 7.11$ wt.%) and #6-2 ($\Omega_{Si} = 8.91$ wt.%) led to estimate f_{CHG}^{A} values lower than 0.10 which agree with the chunky graphite prediction made by means of Ω_{Si} . A more detailed structural study of the whole set of alloys from second series will be the subject of future work.

Concerning elongation at rupture A, the results reported in figure 5 were already showing the amount of chunky graphite. It could be seen that there are alloys with low chunky graphite content or without this defect at high silicon content (alloys #10 to #12, #16 and #25) which show low and very low values of A. This is an indication that chunky graphite may not be prevalent in determining elongation at rupture for alloys with high silicon contents. In other words, even though the effect of chunky graphite was confirmed for alloys with lower silicon content and high A values, the intrinsic effect of increasing silicon content on the decrease of A seems to overtake the effect of chunky graphite at silicon content higher than 4.60 wt.%. As already demonstrated, the embrittlement effect is most probably related to the long-range ordering of ferrite and it has been stressed that there should be a close relation between UTS and A near the maximum of UTS and Y. This is further illustrated in figure 14 where the data for the first series of alloys have been differentiated depending on the amount of chunky graphite. Thus, increase in silicon content makes the matrix tougher, overtaking the detrimental effect of chunky graphite.



Figure 14. Relation between A and UTS around the maximum in UTS.

Conclusions

In this second part of the study, the room temperature mechanical properties of the alloys investigated in the first part have been characterized and complemented with results from an additional series of high silicon alloys and from previous results of low silicon alloys. In this way, a detailed evolution of the mechanical properties of cast irons with silicon ranging from 1.3 wt.% to

6.15 wt.% was performed. As expected, HBW hardness increases continuously up to 310-315 at 6.15 wt.% Si, and even to 396 in an alloy containing 9.12 wt.% silicon. In contrast, the ultimate tensile strength UTS and the yield strength Y present a maximum at 5.0–5.2 wt.% Si for UTS and 5.2–5.4 wt.% Si for Y. Both parameters rapidly decrease to zero as silicon content is increased. The values of UTS and Y reported in the present study agree with literature results up to 4.6 wt.% Si, but show a maximum at higher silicon contents than previously indicated. This opens up the possibility to design cast irons with higher silicon contents than thought until now.

However, the increase in UTS and Y is also associated with a decrease of the elongation at rupture A which is zero at about 5.3 wt.% Si. This decrease is associated with the chemical ordering of the ferritic matrix which is the reason for the so-called embrittlement of cast irons as their silicon content is increased. The critical silicon content is exactly the one at which the maximum values of UTS and Y were found. The decrease of A is quite sharp between 3 and 5.3 wt.% Si. In turn, this means that alloys at 5.0–5.2 wt.% Si content that correspond to the maximum in UTS do present an elongation at rupture of a few percents and could thus be considered for industrial applications.

As presented in the first part of the study, many of the high silicon alloys contain chunky graphite in an amount that could be decreased by adding antimony. For silicon contents up to 4 wt.%, it was found that chunky graphite affects significantly both UTS and A as reported in the literature for common low silicon ductile irons. However, this sensitivity is much less prevalent at silicon contents higher than this value. The present results suggest that as silicon makes the matrix harder, this overtakes the detrimental effect of chunky graphite at silicon contents higher than 4.60 wt.%.

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References

[1] W. Fairhurst, K. Röhrig, High-silicon nodular irons, Foundry Trade J., vol 146, 1979, 657–681.
[2] G. Liao, D. Li, G. B. Burger, R. N. Logan, Cast iron with improved high temperature properties, Patent application publication US 2008/0274005 A1.

[3] P. Weiβ, J. Brachmann, A. Bührig-Polaczek, S. F. Fischer, Influence of nickel and cobalt on microstructure of silicon solution strengthened ductile iron, Mat. Sci. Technol., vol 31, 2015, 1479–1485. DOI: 10.1179/1743284714Y.0000000735.

[4] J. Friess, B. Pustal, P. Weiss, M. Riebisch, A. Kutz, A. Bührig-Polaczech, CASTECH 2016, Darmstadt, <u>http://www.casttec2016.de/uploads/CastTec_2016_Tagungsband_Proceedings.pdf</u>, accessed July 23, 2017.

[5] W. L. Bradley, M. N. Srinivasan, Fracture and fracture toughness of cast irons, Inter. Mater. Rev., vol 35, 1990, 129–161. DOI: 10.1179/095066090790324028.

[6] L. E. Björkegren, K. Hamberg, B. Johannesson, Mechanical properties and machinability of Sisolution-hardened ferritic ductile iron, AFS Trans., vol 104, 1996, 139–145.

[7] L. E. Björkegren, K. Hamberg, Silicon alloyed ductile iron with excellent ductility and machinability, Keith Millis Symp. proceedings, 2003.

[8] U. de la Torre, A. Loizaga, J. Lacaze, J. Sertucha, As cast high silicon ductile irons with optimised mechanical properties and remarkable fatigue properties, Mat. Sci. Technol., vol 30, 2014, 1425–1431. DOI: 10.1179/1743284713Y.0000000483.

[9] A. Reynaud, Corrosion of Cast Irons, in Shreir's Corrosion, T.J.A. Richardson, Ed. vol 3, cap. 2, Elsevier, Amsterdam, The Netherlands, 2010, 1737–1788.

[10] M. A. Arenas, A. Niklas, A. Conde, S. Méndez, J. Sertucha, J. J. de Damborenea, Corrosion behaviour of ductile cast irons partially modified with silicon in 0.03 M NaCl, Rev. Met., vol 50, 2014, e032. DOI: 10.3989/revmetalm.032.

[11] J. Sertucha, J. Lacaze, J. Serrallach, R. Suárez, F. Osuna, Effect of alloying on mechanical properties of as cast ferritic nodular cast irons, Mat. Sci. Technol., vol 28, 2012, 184–191. DOI: 10.1179/1743284711Y.0000000014.

[12] W. Stets, H. Löblich, G. Gassner, P. Schumacher, Solution strengthened ferritic ductile cast iron properties, production and application, Int. J. Metalcast., vol 8, 2014, 35–40. DOI: 10.1007/BF03355580.

[13] G. N. J. Gilbert: BCIRA J., 1960, vol 8, 401–421.

[14] H. T. Angus, Cast iron: physical and engineering properties, vol 210; 1976, London, Butterworths.

[15] J. Lacaze, P. Larrañaga, I. Asenjo, R. Suárez, J. Sertucha, Influence of 1 wt-% addition of Ni on structural and mechanical properties of ferritic ductile irons, Mat. Sci. Technol., vol 28, 2012, 603–608. DOI: 10.1179/1743284711Y.0000000100.

[16] A. Alhussein, M. Risbet, A. Bastien, J. P. Chobaut, D. Balloy, J. Favergeon, Influence of silicon and addition elements on the mechanical behavior of ferritic cast iron, Mat. Sci. Eng. A, vol 605, 2014, 222–228. DOI: 10.1016/j.msea.2014.03.057.

[17] Z. Glavas, A. Strkalj, A. Stojakovich, The properties of silicon alloyed ferritic ductile irons, Metalurgija, vol 55, 2016, 293–296.

[18] J. E. Wittig, G. Frommeyer, Deformation and fracture behavior of rapidly solidified and annealed iron-silicon alloys, Metall. Mater. Trans A, vol 39A, 2008, 252–265. DOI: 10.1007/s11661-007-9436-7.

[19] S. I. Karsay, E. Campomanes, Control of graphite structure in heavy ductile iron castings, AFS Trans., vol. 78, 1970, 85–92.

[20] B. Prinz, K. J. Eschborn, T. Schulze, R. Döpp, E. Schurmann, Untersuchung von Ursachen von Graphitentartungen bei Gusseisen mit Kugelgraphit in Form von Chunky-Graphit, Giessereiforschung, vol 43, 1991, 107–115.

[21] Z. Jiyang, W. Schmitz, S. Engler, Untersuchung der Gefügebildung von Gusseisen mit Kugelgraphit bei langsamer Erstarrung, Giessereiforschung, vol 39, 1987, 55–70.

[22] K. Hamberg, L. E. Björkgren, Z. X. Sun, Chunky graphite in ductile iron, a literature survey, Svenska Gjuteriföreningen, 1997, 16–49.

[23] J. Lacaze, L. Magnusson-Åberg, J. Sertucha, Review of microstructural features of chunky graphite in ductile cast irons, Proc. Keith Millis symposium, Nashville, AFS, 2013, 360–368.
[24] U. de la Torre, J. Lacaze, J. Sertucha, Chunky graphite formation in ductile cast irons: effect of silicon, carbon and rare earths, Int. J. Mater. Res. (formely Z. Metallkd.), vol 107, 2016, 1041–1050. DOI: 10.3139/146.111434.

[25] G. K. Bousse, J. Parolini, N. Rojek, C. Zhou, Quality requirements for General Electric ductile iron castings exceeding 10T used for wind and gas turbine components, Proceedings of the WFC congress, 2010.

[26] G. M. Goodrich, R. W. Lobenhofer, Effect of cooling rate on ductile iron mechanical properties, AFS Trans., vol 110, 2002, 1003–1032.

[27] G. M. Goodrich, R. W. Lobenhofer, Effect of cooling rate on pearlitic ductile iron mechanical properties, AFS Trans., vol 115, 2007, paper 07–045.

[28] G. H. Hsu, M. L. Chen, C. J. Hu, Microstructure and mechanical properties of 4% cobalt and nickel alloyed ductile irons, Mater. Sci. Eng. A, vol A444, 2007, 339–346. DOI: 10.1016/j.msea.2006.09.027.

[29] J. Serrallach, J. Lacaze, J. Sertucha, R. Suárez, A. Monzón, Effect of selected alloying elements on mechanical properties of pearlitic nodular cast irons, Key Eng. Mater., vol 457, 2011, 361–366. DOI: 10.4028/www.scientific.net/KEM.457.361.

[30] J. M. Tartaglia, R. B. Gundlach, G. M. Goodrich, Optimizing structure-property relationships in ductile iron, Int. J. Metalcast., vol 8, 2014, 7–38. DOI: 10.1007/BF03355592.

[31] L. Ceschini, Al. Morri, An. Morri, E. Salsi, R. Squatrito, I. Todaro, L. Tomesani, Microstructure and mechanical properties of heavy section ductile iron castings: experimental and numerical evaluation of effects of cooling rates, Int. J. Cast Met. Res., vol 28, 2015, 365–374. DOI: 10.1179/1743133615Y.000000022.

[32] M. Gagné, D. Argo, Heavy section ductile iron castings – Part I: structure and properties, Int. Conf. "Advanced Casting Technology", J. Easwaren ed., ASM Int., 1987, 231–244.

[33] R. Källbom, K. Hamberg, L. E. Björkegren, Chunky-graphite – Formation and influence on mechanical properties in ductile cast iron, in Competent design by castings – Improvements in a Nordic project, ed. By J. Samuelson, G. Marquis and J. Solin, VTT, Helsinky, Finland, 2005, 63–86.

[34] P. Minnebo, K. F. Nilsson, D. Blagoeva, Tensile, compression and fracture properties of thickwalled dustile cast iron components, J. Mater. Eng. Perform., vol 16, 2007, 35–45. DOI: 10.1007/s11665-006-9005-z.

[35] A. Mourujärvi, K. Widell, T.Saukkonen, H. Hänninen, Influence of chunky graphite on mechanical and fatigue properties of heavy-section cast iron, Fatigue Fract. Eng. Mater. Struct., vol 32, 2009, 379–390. DOI: 10.1111/j.1460-2695.2009.01337.x.

[36] P. Ferro, P. Lazzarin, F. Berto, Fatigue properties of ductile cast iron containing chunky graphite, Mat. Sci. Eng. A, vol 554, 2012, 122–128. DOI:

http://dx.doi.org/10.1016/j.msea.2012.06.024.

[37] P. Ferro, A. Fabrizi, R, Cervo, C. Carollo, Effect of inoculant containing rare earth metals and bismuth on microstructure and mechanical properties of heavy-section near-eutectic ductile iron castings, J. Mat. Process. Technol., vol 213, 2013, 1601–1608. DOI:

10.1016/j.jmatprotec.2013.03.012.

[38] E. Foglio, D. Lusuardi, A. Pola, G. M. La Vecchia, M. Gelfi, Fatigue design of heavy section ductile irons: Influence of chunky graphite, Mater. Des., vol 111, 2016, 353–361. DOI: 10.1016/j.matdes.2016.09.002.

[39] T. Borsato, P. Ferro, F. Berto, C. Carollo, Mechanical and fatigue properties of heavy section 3 solution strengthened ferritic ductile iron castings, Adv. Eng. Mater., vol 18, 2016, 2070–2075. DOI: 10.1002/adem.201600256.

[40] J. Lacaze, B. Sundman, An assessment of the Fe-C-Si system, Metall. Mater. Trans. A, vol 22, 1991, 2211–2223. DOI: 10.1007/BF02664987.

[41] F. González, Y. Houbaert, A review of ordering phenomena in iron-silicon alloys, Rev. Met., vol 49, 2013, 178–199. DOI: 10.3989/revmetalm.1223.