

Effect of aging on microstructure and tensile properties of ASTM F75 Co-based alloy containing different carbon contents

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Abstract

This study aims to investigate the effect of aging on microstructural evolution and mechanical properties of two ASTM F75 alloys, one without the carbon additive and the other one containing 0.21 wt.% C. Aging on the samples solution-annealed at 1225 °C for 1 hour was performed at 720, 760, and 800°C for 5 hours. The evaluation of the microstructure of both samples after aging showed that the $\gamma \rightarrow \epsilon$ transformation led to the formation of two various morphologies of the precipitates in the regions of the ϵ -phase. At the first stages of aging, the precipitates appeared as a series of straight bands in both samples, while with an increase in the temperature, pearlite-like regions were found at the grain boundaries of only the carbon-containing sample. In the sample without the carbon additive, the microstructure consisted of the σ -phase. In the carbon-containing sample, lamellar $M_{23}C_6$ carbides were formed during aging in the vicinity of the remaining blocky $M_{23}C_6$ carbides. The carbon-containing sample consisted of fine $M_{23}C_6$ carbides

after aging at 720°C, which were distributed inside the grains and at the grain boundaries. The appropriate precipitation rate and their optimum distribution in the structure led to an increase in the yield strength by up to 15%, along with an increase in the ultimate tensile strength by up to 6% compared with the as-cast state. However, no significant improvement was found in the ductility of the alloy.

Keywords: Co-Cr-Mo based alloy, aging, microstructure, carbide, tensile strength

1. Introduction

Medical grade casting ASTM F75 alloy is based on 27-30 wt.% Cr, 5-7 wt.% Mo, and a balanced amount of Co, along with a minor content of other elements such as C, N, Si, Mn, Al, Ti, Fe, Ni, and B. This alloy is broadly employed to fabricate orthopedic implants owing to its high strength because of the formation of carbides and its high wear resistance [1-3].

ASTM F75 alloy (in as-cast/annealed conditions) is widely used in the fabrication of prosthetics due to its high corrosion resistance in body fluid environments. The characteristics of the ordinary casting process, such as large grain size and casting defects, lead to both low yield strength and fatigue fracture in the hip prosthesis area. Therefore, to overcome these problems, investment casting and powder metallurgy methods are suitable alternatives. The investment casting process is usually used to reduce the grain size and also to improve the distribution of alloying elements and

carbides [4,5]. The defects formed in the as-cast condition have motivated researchers to study the effect of various heat treatments on the microstructure and mechanical properties of this alloy. Common heat treatments performed on this alloy include solution annealing and aging [6]. Controlling the carbon content of the solution, morphology, and the distribution of the carbides formed in the Co-Cr-Mo-based alloys are of great importance in enhancing the mechanical properties of these alloys [7,8].

The microstructure of ASTM F75 alloy in as-cast conditions, according to its chemical composition, can consist of matrix phases of γ from FCC cobalt or ϵ from HCP cobalt, $M_{23}C_6$ secondary carbides with blocky and lamellar morphologies, σ phase, and M_6C phase. The formation of $M_{23}C_6$ is considered the major strengthening mechanism in this alloy. The addition of carbon leads to an enhancement in both the stability of the γ -phase and mechanical properties. However, the addition of excess amounts of carbon causes the formation of $M_{23}C_6$ carbide [9-11]. When coarse $M_{23}C_6$ carbides are precipitated along the grain boundaries, they can act as preferred sites for the propagation of fatigue cracks. Therefore, obtaining an optimal microstructure through uniform distribution of precipitates, morphology, size, and volume fraction of these precipitates in the matrix phase is done to enhance the mechanical properties by the solution annealing heat treatment and aging [12].

Regarding their chemical composition, cobalt-chromium-molybdenum-based alloys possess two matrix phases, i.e., low-temperature ϵ with an HCP crystal structure and high-temperature γ with an FCC crystal structure. The $\gamma \rightarrow \epsilon$ transformation temperature for pure Co is 417 °C. However, by adding elements such as Cr and Mo, the transformation temperature reaches 870 °C since these elements are ϵ -stabilizing elements [13]. The ϵ -phase resulting from the $\gamma \rightarrow \epsilon$ transformation is known as the martensitic ϵ . The martensitic ϵ could be achieved in three ways: 1- Isothermal transformation, 2- Athermal transformation, and 3- Strain-induced transformation during the heat treatment [14].

Most of the studies on this subject investigated the effect of alloying elements such as Cr, N, Ni, Si, and Mn on the microstructure of ASTM F75 alloy, and only a few studies have paid attention to the microstructure and mechanical properties of the alloy, and the effect of heat treatments on the enhancement of mechanical properties has been rarely reported. Song [Lee et al. \[15\]](#) fabricated various samples of ASTM F75 alloy with various amounts of carbon from 0 to 0.18 wt.% and found an increase in the volume fraction of the γ -phase from 12 to 30. This led to an increase in the tensile strength and ductility (30%) with the presence of 0.1 wt.% C. In addition, carbon increased the stacking fault energy while decreasing the density of the stacking fault, which led to the stabilization of the γ -phase. In another study, [Dobbs et al. \[16\]](#) investigated the effect of the solution annealing heat treatment at 1240 °C

(for 0.5, 1, 2, 4, and 8 hours) and aging at 720 °C (for 6, 9, 16, 24, 48 hours) on tensile properties. They showed that heat treatment, in general, causes an improvement in the mechanical properties without compromising corrosion resistance. Also, the optimal properties are achieved in the sample undergoing partial solution annealing (at 1225 °C for 15 min and air cooling). [Herrera et al. \[17\]](#) studied the effect of carbon content on the mechanical properties of ASTM F75 alloy in as-cast and solution-annealed conditions and fabricated three alloys with low, medium, and high amounts of carbon to evaluate the effect of carbon content. They found that carbon content exhibited a noticeable effect on the mechanical properties of the heat-treated alloys, and the alloy with a medium amount of carbon showed the most optimal mechanical properties.

This study aims to investigate the microstructural evolution of ASTM F75 alloy, with and without the carbon additive, by changing the heat treatment cycle of the solution annealing to compensate for the poor mechanical properties of this alloy, such as its low strength and ductility in the as-cast state.

2. Experimental procedure

The ingots with different amount of carbon were melted in an induction melting furnace under Ar inert gas and cast into a Al₂O₃ crucible. Cast ingots were annealed at 1225 °C for 1 h followed by water quenching. 10×10×10 mm samples were exposed to aging test in temperatures of 720, 760 and 800 °C for 5 h and quenched

into water. The electrical resistance furnaces used in this research had accuracy about ± 5 °C. Chemical composition determined using quantometry method (Spectro2004) and the results are listed in table 1.

Table1. Chemical composition of ingots in accordance to standard.

element	Co	Cr	Mo	Si	Mn	Fe	Al	C
ASTM F75 [18]	Bal.	27-30	5-7	<1	<1	<0.75	<0.1	<0.35
Product 1	Bal.	28.16	5.49	0.35	0.53	0.3	0.016	0.01
Product 2	Bal.	28.24	5.63	0.32	0.43	0.24	0.077	0.21

Metallographic sections were prepared using standard mechanical polishing procedures and electroetched using 90% H₂SO₄-10% methanol solution for 30-240 second with the voltage of 6 Volt. A Vega Tescan scanning electron microscope (SEM) with an energy dispersive spectroscopy (EDS) analyser operating at 15 kV and optical microscopy (Olympus DP25) were used to observe the morphology and distribution of the phases.

A Rigaku Smart Lab XRD was used with Cu-K α radiation at 40 kV and 30 mA. The volume fraction of phases was calculated by Imag J analysis software.

Tensile tests at room-temperature were carried out on an Instron Model-8502 test machine. Tensile tests were conducted in accordance with ASTM E8 [19] of the tension testing of metallic material.

3. Results and Discussion

3.1. Microstructural investigations

Figures 1 and 2 present the optical micrographs of the samples with and without carbon in different annealing and aging conditions. In samples without carbon, some precipitates can be found, the EDS analysis of which is shown in Fig. 3 for the sample aged at 760 °C. As can be seen in point A, the matrix phase consists of the base alloy elements, i.e., Cr, Mo, and Co, with atomic percentages of 32.28, 67.77, and 2.95, respectively. In point B, Cr, Co, and Mo are present with atomic percentages of 42.97, 48.33, and 8.70. Thus, it appears that the precipitates present are of σ type. σ precipitates could also be found in annealed conditions, and with an increase in the aging temperature from 720 °C to 800 °C, the amount and size of σ precipitates increased. This can be explained by the fact that the aging temperature was close to the temperature range of the formation of σ , i.e., 540-980 °C [20]. σ exhibits a tetragonal structure, which is composed of the Co-Cr intermetallic phase. Due to its brittle nature, this phase is detrimental. By adding carbon, $M_{23}C_6$ carbides were formed, according to reports of other studies $(Co-Cr/M) + C \rightarrow M_{23}C_6$ [1], which can play a crucial role in precipitation strengthening.

The formation of $M_{23}C_6$ precipitates will be discussed in the carbon-containing alloy.

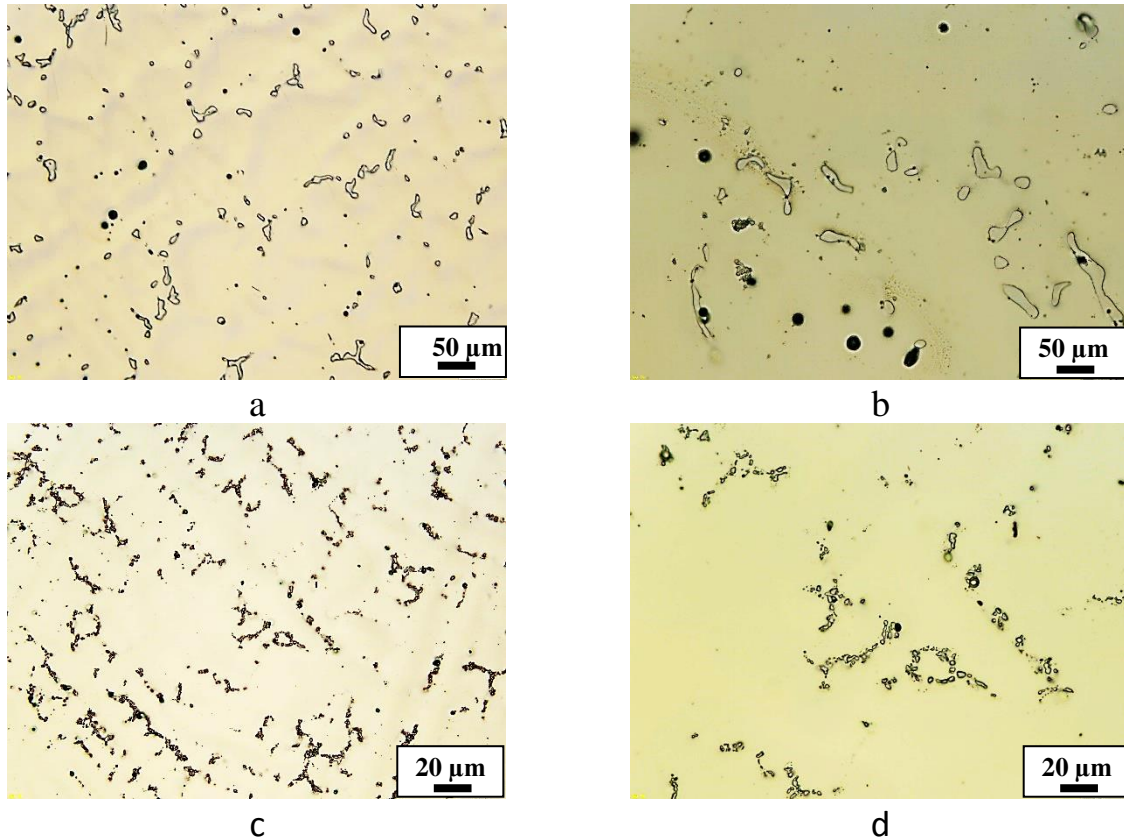


Fig 1. Optical microscope microstructure in the annealed state at 1225°C for 1 hour:

a,b) Samples without carbon c,d) Samples with carbon

At 800 °C and in the sample without the carbon additive, a series of precipitates were formed as straight bands around the initial precipitate of the σ -phase, which can be seen in the optical micrograph of Fig. 4 with a higher magnification. These regions are the ϵ -martensite phase with an HCP structure with numerous stacking faults, which are the preferred sites for the formation of precipitates. The reason for the formation of the precipitates in these regions is that Co present in the matrix reacted with Cr and Mo, which were segregated preferentially to dislocations and stacking faults, and the σ -phase was formed as straight bands.

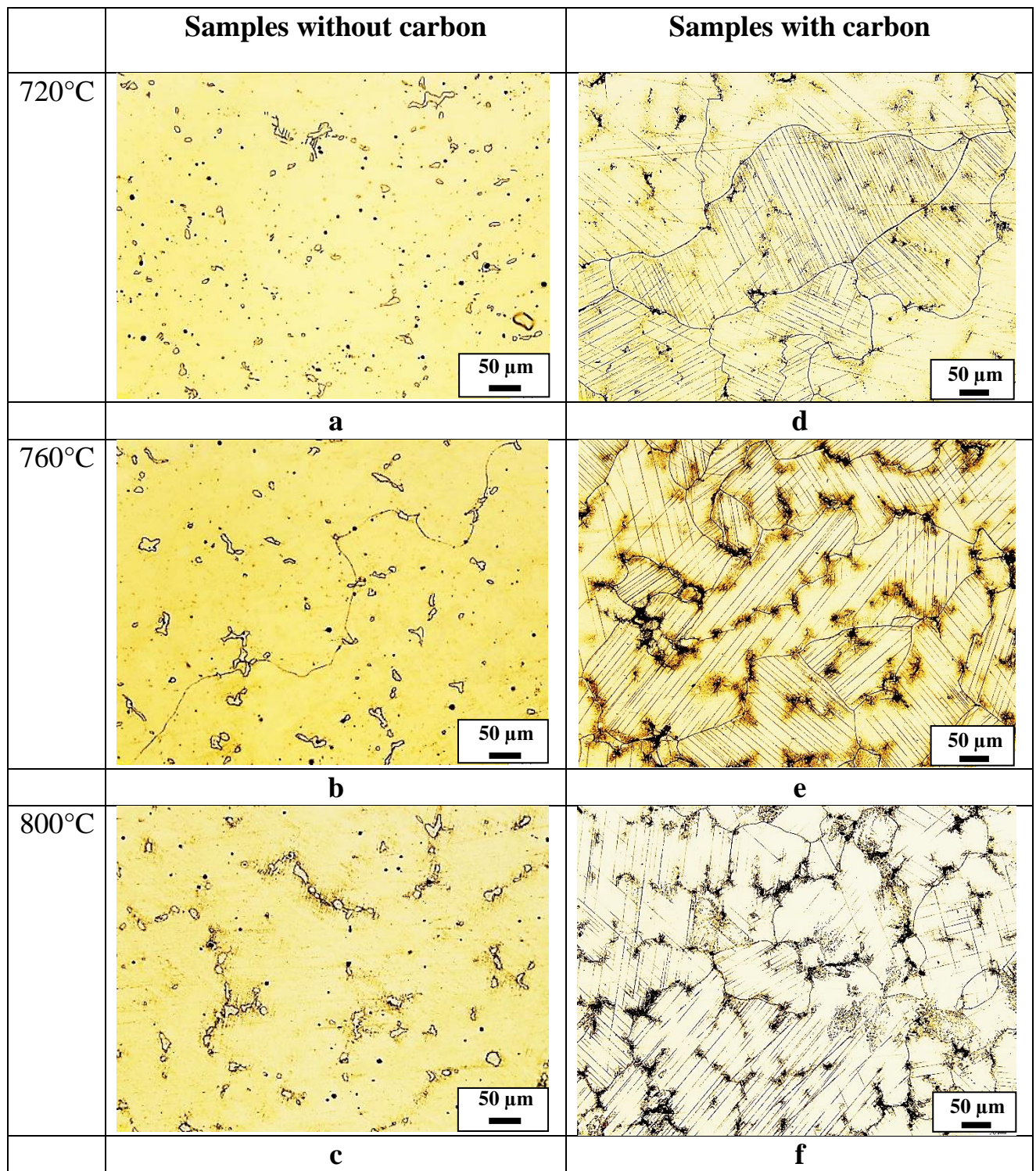


Fig 2. Optical microstructure images of a, b, c) carbon-free and d, e, f) carbon-containing samples aged for 5 hours at temperatures of a,d) 720°C, b,e) 760°C, c,f)

800°C

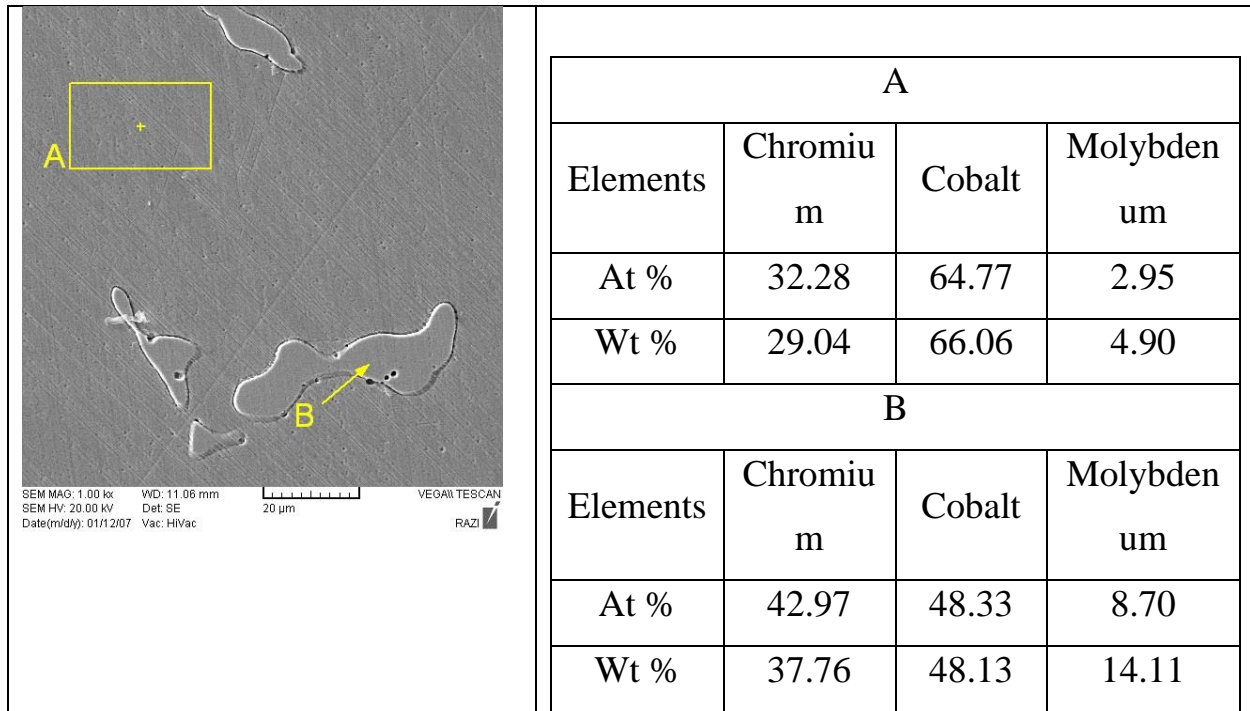


Fig 3. The electron microscope image of the microstructure of the carbon-free sample aged at 760°C for 5 hours along with the results of EDS analysis

According to the literature [21,22], by aging at temperatures lower than 925 °C, two mechanisms can be proposed for the $\gamma \rightarrow \epsilon$ transformation. The first one suggests that in the short times of the first stages of aging, the FCC to HCP₁ transformation with an orientation relationship of FCC{111} || HCP {0001} leads to the formation of the ϵ -martensite. The regions of the FCC phase are single-phase, while those of the HCP phase consist of carbides, as well. These carbides exhibit an orientation relationship of {0001} HCP || {111} M₂₃C₆ with HCP. Therefore, the carbide nucleation in HCP₁ occurs as a series of straight bands. The second mechanism, which was seen in the alloys that contain carbon, suggests that at longer times and higher aging

temperatures, the FCC to HCP₂ transformation occurred at the grain boundaries from the FCC matrix along with the carbide nucleation and results in the formation of a lamellar pearlite-like morphology, which will be discussed in the following.

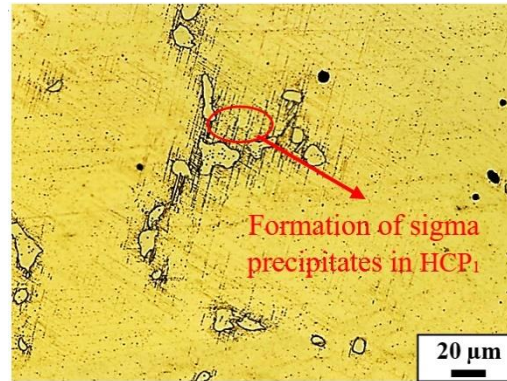


Fig 4. Image of the optical microstructure of the carbon-free sample aged for 5 hours at 800°C

Figure 2d-f shows the microstructures of the carbon-containing sample aged at 720 °C, 760 °C, and 800 °C, which consist of fine M₂₃C₆ precipitates at the grain boundaries and inside the grains along with parallel lines of the precipitation of carbides in HCP₁. The scanning electron microscope (SEM) micrograph and the EDS analysis for the formation of M₂₃C₆ in the regions of HCP₁ in the sample aged at 760 °C are displayed in Fig. 6. According to the EDS analysis of point A in Fig. 6d, the concentration of Cr is higher than that of Co and Mo. Since the Cr content in M₂₃C₆ carbide has been reported to be higher than that of Co and Mo [17], it appears that this precipitate was of M₂₃C₆ type. The reason for the absence of the σ-phase precipitates in the carbon-containing sample, as stated previously, is that these

precipitates reacted with carbon present in the alloy, and $M_{23}C_6$ carbides were formed. Since the stacking fault density in the HCP crystal structure is higher than that in the FCC crystal structure [23], the presence of the stacking fault in these regions led to a significant decrease in the activation energy of the precipitation. The solute atoms present in the alloy, which were dissolved by solution annealing, had a strong tendency to segregate in these regions. Therefore, parallel lines were the preferred sites for the nucleation of $M_{23}C_6$ carbides. With an increase in the aging temperature to 760 °C, as shown in Fig. 2e, the precipitation rate of $M_{23}C_6$ carbides inside the grains and especially at the grain boundaries increased, and coarser precipitates could be found at this temperature compared with those at 760 °C. In addition, it can be seen that the $\gamma \rightarrow \epsilon$ transformation has been done more at this aging temperature, and the parallel lines related to the precipitation of $M_{23}C_6$ carbides increased in the regions related to HCP_1 . In the early stages of aging, especially at low temperatures and short times, the formation of precipitates often occurs in the form of parallel lines in HCP_1 regions, and the formation of carbides with a pearlite-like morphology occurs at higher temperatures and longer times.

The XRD pattern of the carbon-containing sample after aging is seen in Fig. 5. With an increase in the aging temperature from 720 °C to 760 °C in the aged samples, the peaks corresponding to the ϵ -phase increased due to the occurrence of the $\gamma \rightarrow \epsilon$

transformation because of the increase in the stacking fault and distortion, whereas the peak corresponding to the γ -phase decreased.

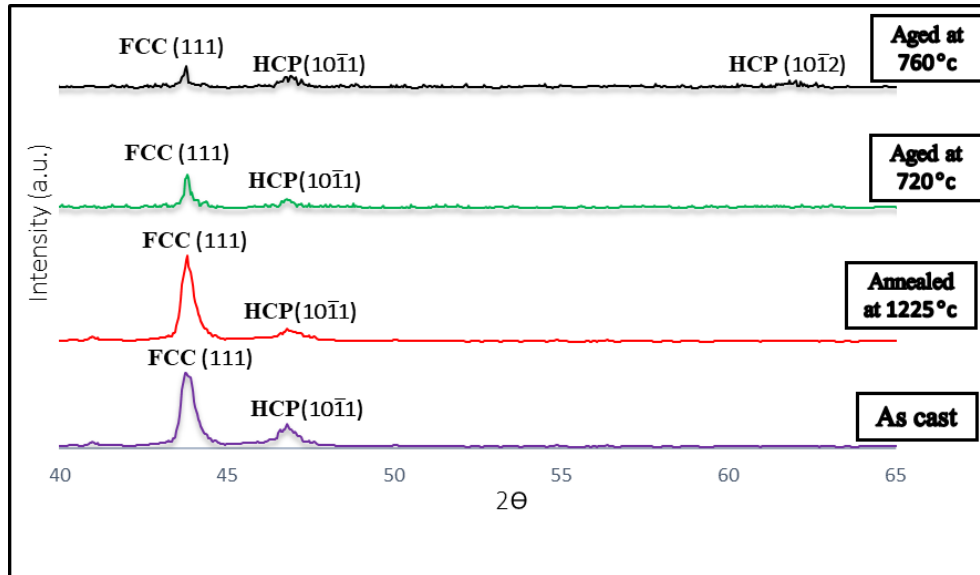


Fig. 5 X-ray diffraction pattern of ASTM F75 sample containing carbon in different conditions.

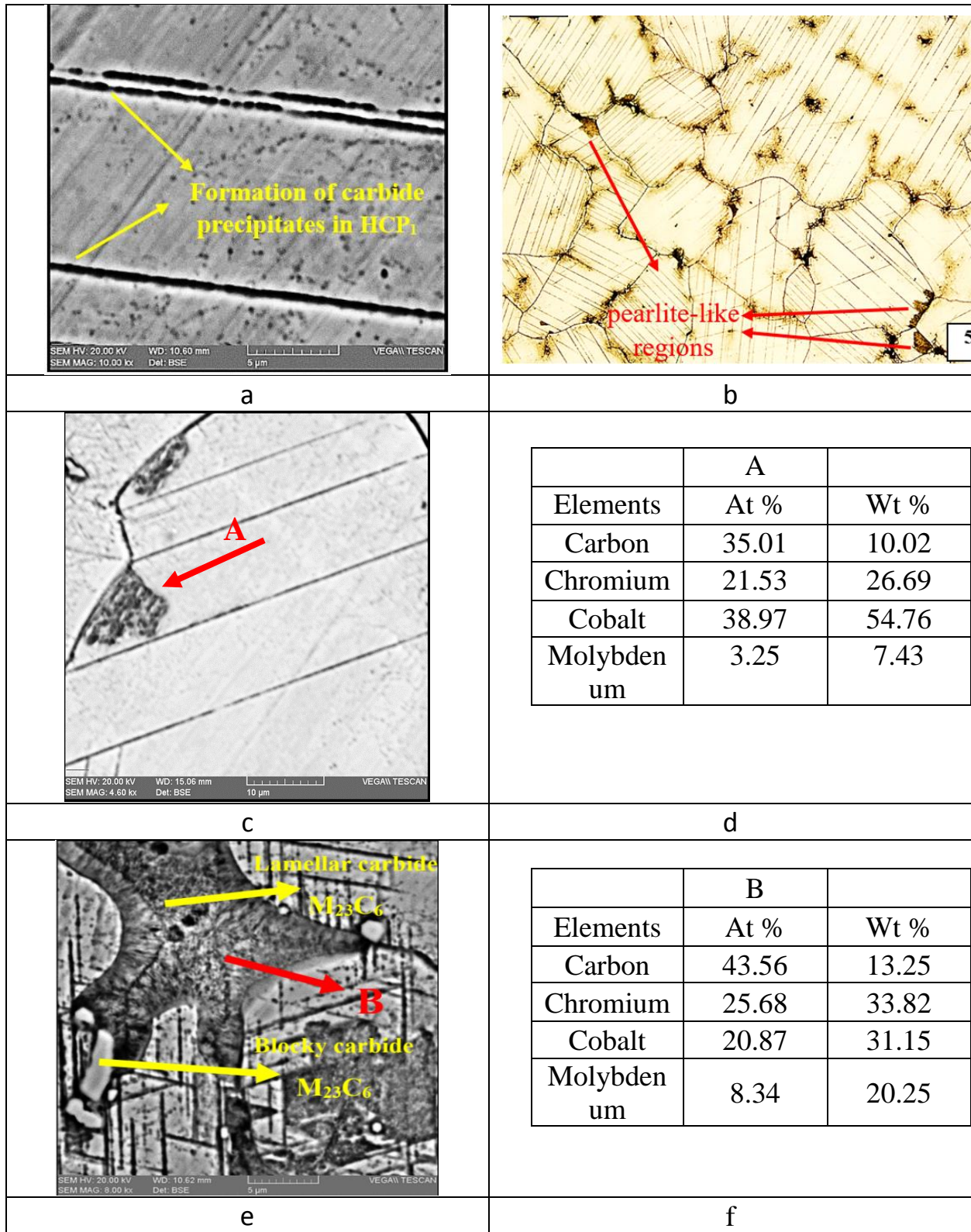


Fig. 6 Optical microstructure and electron microscope image of the sample containing carbon aged at 760°C for 5 hours with the results of EDS analysis.

It has been reported that [24] carbon atoms tend to be absorbed by dislocations and stacking faults and form segregated regions of carbon. These carbon atoms react with chromium and molybdenum atoms, which segregate preferentially to dislocations and stacking faults and lead to the creation of regions in the form of parallel lines. Another effect that was observed by adding carbon to ASTM F75 alloy aged at 760°C was the formation of a phase with a pearlite-like morphology at the grain boundaries. By increasing the aging temperature and time, the conditions for the transformation are provided in the form of a series of discontinuous reactions simultaneously with the formation of HCP₂ at the grain boundary, which has a low stacking fault, unlike HCP. This can be explained by the high mismatch at the grain boundaries and an increase in the growth rate of carbides in these regions due to an increase in the diffusion rate caused by increasing the aging temperature. The optical and SEM micrographs of the pearlite-like region formed at the grain boundaries can be seen in Fig. 6b and Fig. 6c, respectively. As shown, in the pearlite-like phase, the Co content is higher than that in the lamellar carbide, according to the EDS analysis result in Fig. 6d. The reason is that this region consisted of two phases, i.e., M₂₃C₆ and ε. The high content of Co in this region is related to the presence of the ε-martensite phase, which was formed from the γ→ε transformation during aging. Figure 6d shows M₂₃C₆ carbides with two different morphologies. In this figure, blocky carbides remaining from solution annealing can be observed around lamellar

carbide (black) formed in the aging conditions. Due to the dissolution of a part of blocky carbides during solution annealing at 1225 °C for 1 hour, the content of C, Cr, and Mo around these carbides increased. During aging, lamellar carbides were formed because of appropriate conditions provided for the diffusion and segregation of the elements, which are seen in black color in the three microstructures aged at 720 °C, 760 °C, and 800 °C, similar to the lamellar carbides present in the as-cast condition. With an increase in the aging temperature to 800 °C and according to Fig. 2f, an increase in the temperature led to the coarsening of lamellar $M_{23}C_6$ carbides and the formation of more consistent precipitates, especially at the grain boundaries. The morphologies observed at 720 °C as parallel lines and pearlite-like regions that appear at the grain boundaries became more prominent at this aging temperature (800°C).

According to Fig. 2f, lamellar $M_{23}C_6$ carbides (black regions) can be found inside the grains, especially at the grain boundaries. Another noteworthy point is that the volume fraction of the pearlite-like regions (gray regions) increased at the grain boundaries with an increase in the aging temperature compared with that of the samples aged at 720 and 760 °C since increasing the aging temperature led to an increase in the diffusion rate of the elements at the grain boundaries and as well as an increase in the mismatch of the grain boundaries. Also, the ϵ -martensite was formed with an increase in the stacking fault density.

3.3. Hardness measurements

Figure 7 presents the hardness values of the as-cast, annealed, and aged samples.

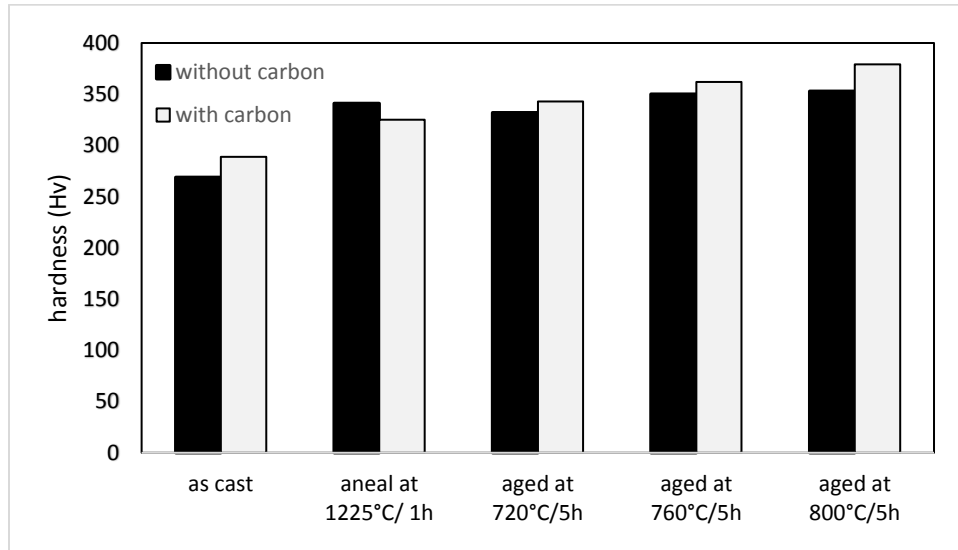


Fig 7. Hardness values of carbon-free and carbon-containing products in different conditions according to Vickers (VHN).

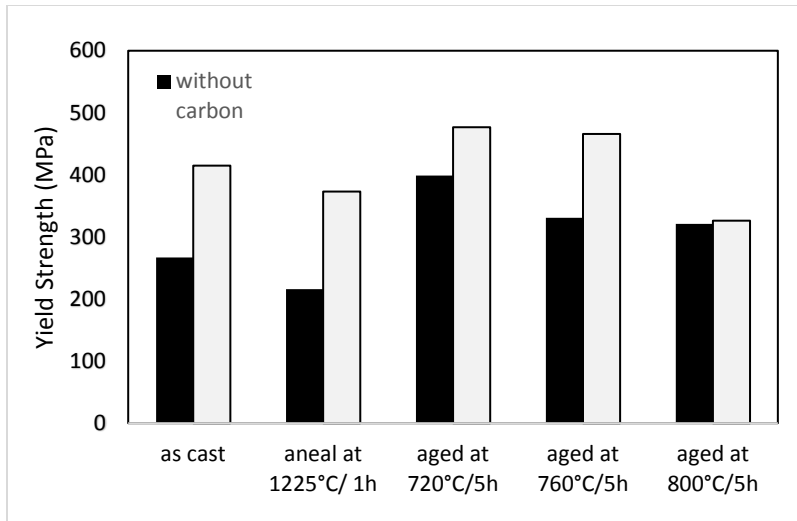
As can be seen, the hardness value increased owing to the addition of carbon, which led to the formation of carbides in the carbon-containing sample. By aging the sample without the carbon additive at 720 °C, the hardness value increased from 269 HV to 332 HV compared with the as-cast state because of the grain refinement that occurred in the aging state. In both samples, with an increase in the aging temperature from 720 °C to 800 °C, the hardness value increased due to an increase in the amount of the ϵ -phase by increasing the aging temperature, as could be seen in the XRD results. The reason is that by increasing the aging temperature from 720 °C to 800 °C, the $\gamma \rightarrow \epsilon$ transformation was done more, and the ϵ -phase content

increased. The other reason for the increased hardness of the carbon-containing sample was the formation of lamellar $M_{23}C_6$ carbides inside the grains and at the grain boundaries.

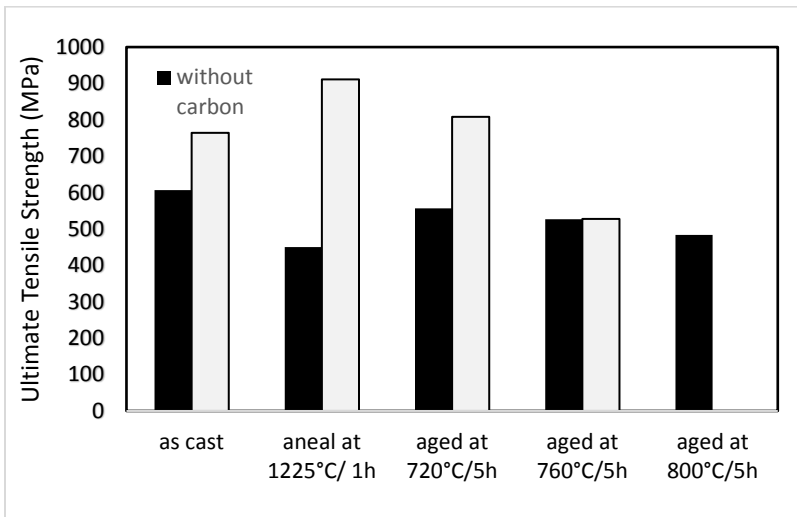
3.2. Evaluation of mechanical properties

The results of cold tensile tests of the samples aged at 720, 760, and 800 °C aged for 5 hours for both samples, i.e., without the carbon additive and with the carbon additive, along with the results of casting and annealing at 1225 °C can be seen in Fig. 7.

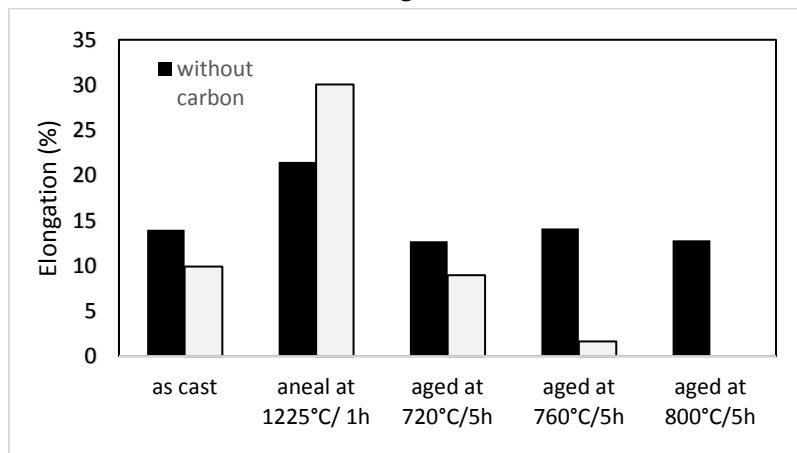
As shown in this figure, by aging the sample without the carbon additive at 720 °C, the yield strength values increased compared with those of the as-cast state but did not reach the standard level. Moreover, both the tensile strength values and ductility did not change noticeably compared with the values of the as-cast state. With an increase in the aging temperature, one could see a decreasing trend in yield and tensile strength as well as ductility with an increase in the volume fraction of the ϵ -phase. Thus, in the sample without the carbon additive, not only no improvement was observed in the mechanical properties due to aging but also, according to the precipitate type in this sample (the sigma phase) along with an increase in the volume fraction of the ϵ -phase, the



a



b



c

Fig 8.Changes in tensile properties according to aging temperature
a) Yield strength b) Tensile strength c) Elongation percentage

mechanical properties were degraded, and none of the parameters except ductility and reduction in area reached the standard level.

In the sample containing the carbon additive, an improvement in the mechanical properties compared with the as-cast state occurred due to the appropriate distribution of the carbides, which was an expected result. In addition, the yield strength of this alloy increased compared with the as-cast state, considering the precipitates formed in the stacking faults and the distribution of the precipitates inside the grains. However, the ductility decreased because of the formation of the ϵ -phase and carbides. Nonetheless, with an increase in the aging temperature, the sample became brittle, and the yield and tensile strength values decreased compared with those obtained in the sample aged at 720 °C owing to the $\gamma \rightarrow \epsilon$ transformation at 760 °C and formation of the ϵ -phase in the structure as well as the formation of interconnected lamellar carbides especially at the grain boundaries. Moreover, the ductility of the alloy reached 1.64%. Therefore, the tensile test results show that in the sample without the carbon additive, a decrease in the mechanical properties in both conditions, i.e., the as-cast and aged conditions, could be observed. This can be ascribed to the absence of carbon in this sample, which led to the absence of carbides in the matrix phase since carbides played a fundamental role in the strengthening of this alloy through precipitation-strengthening. However, in the carbon-containing

sample in aging conditions at 720 °C, the mechanical properties were enhanced compared with the as-cast state.

4. Conclusion

1. In the alloy without the carbon additive, due to the single-phase nature of the alloy (HCP), no significant changes were observed in the microstructure by aging at 720°C and 760°C, and the same sigma-phase precipitates that were present in the as-cast and solution annealing state, could be seen in the aging state. However, in the sample aged at 800 °C, the formation of sigma precipitates occurred in the form of parallel lines in HCP₁ created by the $\gamma \rightarrow \epsilon$ transformation.

2. In the carbon-containing alloy, the sample aged at 720 °C contained fine M₂₃C₆ carbides that were distributed inside the grains and at the grain boundaries. With an increase in the aging temperature to 760 °C, parallel lines of the precipitate formation in HCP₁ increased, and a limited number of regions with pearlite-like morphologies could be seen. In addition, the formation rate of lamellar M₂₃C₆ carbide, especially at the grain boundaries, increased. The formation of pearlite-like regions at the grain boundaries reached the maximum at the aging temperature of 800 °C.

3. After aging, the $\gamma \rightarrow \epsilon$ transformation increased in both samples with an increase in the aging time from 720 °C to 800 °C.

4. In the carbon-containing sample after aging at 720°C, both the ultimate tensile strength (808.9 MPa) and ductility (12.7%) were higher than those of the sample aged at 760 °C due to the presence of a lesser amount of the ϵ -phase and formation of finer carbides.

5. In the sample without the carbon additive, due to the fundamental role of carbides in strengthening through the precipitation-strengthening mechanism, the absence of carbon and consequently the absence of carbide in the matrix phase of this sample, lower mechanical properties in different conditions of the as-cast and aging were observed.

6. The carbon-containing sample showed the best mechanical properties when aged at 720 °C for 5 hours compared with the other aging temperatures.

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