



# EVALUATION OF AUSTEMPERING STUDY OF FERRITIC CAST IRON

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#### ABSTRACT

This study analyzed the effect of austempering temperatures on the structural characterization of ferritic spheroid cast irons. Austempering is the austenitization of the materials at 900 °C for an adequate period to obtain a complete austenitic structure and then quenching to an interval temperature for precipitation of ausferrite. Subcritical diffusion time was achieved as 1.5 h for 320, 420, and 520 °C ausferrite temperatures. Metallurgical description of samples was defined using optical microscopy, energy dispersive spectrometry, X-ray diffraction and microhardness. The austempering temperature had a remarkable effect on the phase variations of the ferritic ductile cast irons. The higher degree of austempering produced a coarser structure called ausferrite. *Anahtar Kelimeler: Austempering, Microstructure, Ausferrite, Hardness.* 

# FERRİTİK DÖKME DEMİRİN ÖSTEMPERLEME ÇALIŞMASININ DEĞERLENDİRİLMESİ

# ÖZET

Bu çalışmada, ferritik sfero dökme demirlerin yapısal karakterizasyonuna östemperleme sıcaklıklarının etkisi analiz edilmiştir. Östemperleme, tam bir östenit yapı elde etmek için malzemelerin 900 °C sıcaklıkta yeterli bir süre östenitlenmesi ve ardından bir aralık sıcaklığında su vermedir. Östemperleme 320, 420 ve 520 °C' de 1.5 saat süreyle elde edilmiştir. Numunelerin metalurjik karakterizasyonu, optik mikroskop, enerji dağılımlı spektrometri, X-ışını kırınımı, mikrosertlik kullanılarak belirlendi. Östemperleme sıcaklığı, ferritik sfero dökme demirlerin faz değişimleri üzerinde dikkate değer bir etkiye sahipti. Daha yüksek derecede östemperleme, ausferrit adı verilen daha kaba bir yapı üretti.

Keywords: Östemperleme, Mikroyapı, Ausferrite, Sertlik.

## 1. Introduction

Ferritic ductile cast iron (FDCI) consists of a ferritic matrix and a spheroidal graphite microstructure. After heat treatment, ausferrite, acicular ferrite and bainite can be observed in the structure [1-3]. The matrix with graphite is intended to create an ausferrite structure by isothermal heat treatments. FDCI has superiority such as lower density and cost, lubrication properties, high elongation and toughness [4,5]. Austempering is a method that produces austenite base structure in DCI. The

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method is a two-step process. The first stage is austenitization for 1 to 4 h in the temperature level of 815–920 °C. It is then quickly quenched in a salt bath. The casting is held isothermally at the detected isothermal annealing temperature in the salt bath. Depending on the desired properties of the casting, the austenite annealing temperature range is 230–450 °C. At higher austempering values, the ferrite nucleates and transforms into austenite [6-8]. A less isothermal annealing temperature will produce a finer, larger ferrite rate and higher yield rate. Austempering time and degree are effective in the occur of ausferrite structure. A higher degree of austempering produces a smaller proportion of coarse ferrite [9-10]. Mandal et al. [11] reported the impress of austempering on the strength values of high-Si cast material. Austempering significantly improved the strength and ductility of the casting.

The aim of this research is to detect the impress of austempering temperatures on the structural characterization of Ce inoculated ferritic ductile cast irons.

#### 2. Material ve Method

The chemical values of the samples included in the study was 3.10%C, Bal.%Fe, 4.30%Si, 0.24%Mn, 0.020%Cr, 0.12%Ce, 0.02%S, 0.025%P (wt.-%). Melting was done in a 300 kg induction furnace using steel scrap. Alloying elements were melted in an induction furnace at 1510 °C. The melt was transferred to the casting ladle at 1460 °C. The austempering parameters of test samples are presented in Table 1. In the first stage, the samples were austenitized at 900 °C for 1.5 h. Then, it was immersed in a salt bath for austempering and kept at this temperature for 1.5 h. Then the samples were cooled to room temperature. Austempering was applied at temperatures of 320, 420 and 520 °C. For metallographic studies, samples were sanded using 80-1200 mesh emery paper and polished a 2  $\mu$ m diamond solution. After etching with 2% nital solution, it was examined by optical microscopy (OM), energy distribution spectrometry (EDS). Phase contents were identified by Bruker brand X-ray diffraction (XRD) in Cu-K $\alpha$  radiation. Microhardness results was detected using a 100 g load at 0.5 mm intervals on the Qness Q10 microhardness machine.

No	Destabilization Temperature	Destabilization Time	Subcritical Diffusion Temperature	Subcritical Diffusion Time	Quenching
S-Ref.	As-Cast	-	-	-	-
<b>S1</b>	900 °C	1.5h	320	1.5h	Air
<b>S3</b>	900 °C	1.5h	420	1.5h	Air
<b>S4</b>	900 °C	1.5h	520	1.5h	Air

**Table 1.** The austempering parameters of test samples.

#### **3.** Experimental results

#### 3.1 Evaluation of microstructure

Fig. 1 shows OM micrographs of S-Ref, S1, S2, S3 specimens austenitized at 900 °C and austempered at 320, 420 and 520 °C. The ADI structure consisted of pearlite, graphite and ferrite. The microstructure of the austempered sample contained acicular dark ferrite and ausferrite. When the subcritical phase transformation temperature was low as 320 °C, the ferrite layers were unevenly oriented and thin. As the subcritical temperature increased from 320 °C to 420 °C, bainitic ferrite laths grew up. Austenite was sandwiched between adjacent ferrites. A thickness rich in austenite and carbon was obtained. In the first step of the transformation, ferrite nucleated at austenite grain boundaries and separated into plates or laths before transforming into carbon austenite. Therefore, ferrite and ausferrite

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emerged in the structure of the samples. The ausferrite formed in the first stage of the transformation had upper lateral growth, and the surrounding ferrite contained less carbon. The EDS values of the casting sample is displayed in Fig. 2. C, Fe, Ce, Si, Mn, S were the main elements of the casting sample. The white particles in Fig. 2 show that the primary precipitates are  $Ce_2O_2S$  with heterogeneous nuclei.  $Ce_2O_2S$  inclusions produced thinner primary ausferrites. It provided heterogeneous nucleation of ausferrites [12].



Figure 1. Optical micrographs of austempered samples at 320, 420, 520 °C at 1.5 h.

The silicon content in the produced ductile iron castings is desired in the range of 2.45-2.80%. It was sufficient to prevent melt cooling and at the same time facilitate the production of sufficient nodules [13]. The excessive silicon contained in the ferritic iron suppressed cementite precipitation during the austempering and retained a significant amount of stable high-carbon austenite.

Austenite and ferrite gradually coarsened as the austempering degree increased. Very slim ferrite and austenite were obtained at the lower austempering temperature. On the other hand, rough and hairy ferrite characteristics of upper bainite were obtained at higher austempering degrees. As austempering progresses, these ferrite needles increase larger and as a result the remaining austenite absorbs carbon, thereby increasing the carbon intensity of the austenite [14,15]. At higher temperatures, decomposition of austenite to ferrite occurred due to the free energy diversity. At higher temperature, austenite was precipitated by spread of carbon into existing graphite nodules along grain boundaries. At high subcritical diffusion temperatures, the ferrite structure nucleated and transformed

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into ausferrite. At higher subcritical diffusion temperatures such as 420–520 °C, the rate of ausferrite incremented. At 520 °C, large areas of austenite separated from each other and coarse ferrite formed. The increasing the subcritical diffusion temperature leaded the ausferrite spicules to thicken and the ferrite velocity to increase. Average lath and ausferrite lengths were the same [16, 17]. It produced finer and larger volume ferrite fractions at lower critical diffusion temperatures. The X-ray patterns of the produced ferritic ductile iron are presented in Fig. 3. The base phases created in the samples were ferrite ( $\alpha$ ), ausferrite, graphite.



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#### 3.2. Hardness

The effect of austenitizing temperature on the hardness is shown in Fig. 4. The hardness increased as the subcritical diffusion temperature decreased. Fine ausferrite created at low subcritical diffusion temperatures had higher hardness. The hardness of ausferrite appeared at higher temperatures was lower. As the isothermal annealing temperature increased, the extent and interval of austenite became more favorable for the transformation of austenite to martensite. Austenite was a softer structure and increasing austenite ratio reduced the hardness. In addition, grain coarsening caused a decrease in hardness [18-20].



Figure 4. The hardness amounts of as-cast and austempered samples.

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## 4. Conclusion

The results obtained are as follows:

The austempering degree had a remarkable action on the phase variations of the ferritic cast irons.

The higher degree of austempering produced a coarser structure called ausferrite.

Finer and larger bulk ferrite were obtained at the lower austempering temperature.

Hardness raised from 240 HV to 360-380 HV by austempering heat treatment. But, it declined due to the coarser ausferritic appeared with increasing austempering temperature.

The phases created in the samples were ferrite ( $\alpha$ ), ausferrite, graphite.

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# **Disclosure statement**

No potential conflict of interest was reported by the authors.

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