Structure and Solidification Process Observation of Fe-C-Cr Alloy Using Confocal Laser Scanning Microscope

Shin Ito1*, Hideki Kawai1, Yoshihiko Oishi1, and Hiroshi Nogami2

1Department of Production Systems Engineering, Faculty of Engineering, Muroran Institute of Technology, 27-1, Mizumoto, Muroran, Hokkaido, Japan
2Department of Metal Frontier Engineering, Faculty of Engineering, Tohoku University, 2-chome 1-1, Aoba-ku Katahira, Sendai, Miyagi, Japan

Abstract In this study, we investigated the growth process of M7C3 carbide by a Confocal Laser Scanning Microscope. Fe-C-Cr alloy was prepared using an arc furnace. The composition was determined to be Fe:C:Cr=55:5:40 (wt%). From the results of XRD and EPMA, the Fe-C-Cr alloy contained M7C3 carbide in the internal structure. In condition of two cooling rates 60°C/min and 30°C/min observed with a Confocal Laser Scanning Microscope. Corners of the Hexagonal structure grew in the preferred orientation direction of M7C3 carbide at 60°C/min. After the growth of preferred orientation direction, the M7C3 carbide formed to connect between the corners. At a cooling rate of 30°C/min, M7C3 carbides grew in oblique direction to the observation area. At the beginning, the structures were hidden in the fluid. But the structures started to grow, the shape of the hexagon became eventually. After the melting experiment, the samples were observed by Optical Microscope and Scanning Electron Microscope. As a result, since hexagonal structure arose a difference in height between the outside and the inside. M7C3 carbides grew from the outside in the solidification process.

1 Introduction

The wear-resistant material Fe-C-Cr alloy is generally a ternary white cast iron containing 7% or more of Cr, and forms a large amount of hard carbides M23C6, M7C3, M7C3 (M=Fe, Cr) (Sato et al., 1958). Hence, it is used for a slurry pump that transfers slurry liquid, and rolling rolls used for rolling. However, the rolling conditions are becoming more severe in terms of energy saving and quality improvement in rolling processing, and the slurry pump has problems of wear and corrosion due to high concentration of slurry liquid. In order to cope with the current situation, it is important to change the solidification structure by adjusting the Cr content and C content to improve the low toughness and to improve the corrosion resistance and wear resistance. Changes in composition and cooling rate are related to the mechanical properties of the alloy and affect the solidified structure. Tang et al. (2011) show that the mass separation of M7C3 carbides increases with the increase of C content, and the corrosion resistance and wear resistance increase accordingly. In addition, Tong et al. (1990a) revealed that the area fraction of carbides increased with the increase of Cr. The commonly used Fe-C-Cr alloy has a Cr content of about 12 to 30 wt%. The alloy of this composition is hypoeutectic, Tong et al. (1990b) Fe-C-Cr alloy in the hypoeutectic composition range crystallizes austenite into primary crystals, It is clarified that Fe-C-Cr alloy in the hypereutectic composition range crystallizes M7C3 carbide in primary crystals. At present, the study of M7C3 carbides in Fe-C-Cr alloys mainly focuses on its performance. Although there are few studies on the growth mechanism of M7C3 carbides, Liu et al. (2016) observed in situ Cr cast iron at a cooling rate of 0.1°C/sec, and proposed a theory on the growth characteristics of hollow and unsealed shell structures present in the center of M7C3 carbides. However, the formation process of solidification structure and the growth process of single carbide by the change of cooling rate and composition are not investigated. It is important to clarify the actual growth of the formation structure from direct observation. Direct observation of the solidification process in which carbides and matrix structure appear leads to elucidation of the growth mechanism of M7C3 carbides, and it can be expected to control the size, amount, and formation position of crystallized materials by changing the cooling rate. In addition, in order to elucidate the growth mechanism of the metal structure, it is necessary to directly observe the metal under high temperature. Therefore, it is essential to use a Confocal Laser Scanning Microscope (CLSM) that is not affected by the radiation emitted from the molten metal. Most common Fe-C-Cr alloys contain small amounts of elements such as Si and Mn. In this study, in order to observe the crystallization behavior of pure Fe, Cr, C ternary system,
samples were prepared using pure metals. In this study, after examining the internal structure using pure ternary samples, melting experiments with a CLSM Microscope were conducted to investigate the growth process of M7C3 carbide.

2 Experimental Method

2.1 Experimental sample

Materials used to make the Fe-C-Cr alloy were pure iron, pure chromium, and graphite plates, respectively. All materials have a purity of 99.9% or more. It is assumed that the composition of the sample to be prepared melts in the high temperature image furnace of the CLSM. From the liquid phase area projection of the Fe-C-Cr system and the Fe-Cr system phase diagram (Baniff et al., 2001), the weight ratio is set to Fe: C: Cr = 55: 5: 40. This composition is in the hypereutectic composition range, and it is assumed that the sample is completely melted at 1600°C. Arc furnace (Nisshin Giken: model NEV-AD03) was used for sample preparation. Since the arc furnace can be heated up to 3000°C, it is suitable for melting materials with high melting points of Cr and C. Furthermore, since the inside can be replaced with an inert gas, the samples do not react with air during melting.

2.2 Measurement Principle of Confocal Laser Scanning Microscope

Structures of samples passing the process of melting and solidification were observed Confocal Laser Scanning Microscope (Lasertech YONEKURA VL2000DX - SVF17SP). Figure 1 shows schematic diagrams of the CLSM and infrared image furnace. The CLSM is capable of real-time observation of molten metal under high temperature by combination of a microscope using a violet laser light source and a high temperature image furnace. A wavelength of light source of the microscope used the violet region at a wavelength of 405 nm. Therefore, compared with the Optical Microscope which irradiates light uniformly, reflected light with less scattered light is obtained. The light reflected from the surface of the sample follows the same path and is split by the beam splitter. Then it is collected by the light detector. A pin hole is placed in front of the

light detector, which makes it possible to detect light only at a focused position. Therefore, observation can be performed without the influence of radiation (Shibata and Emi, 1997). The high-temperature image furnace has an elliptical inside and is coated with gold. Infrared light emitted from a halogen lamp installed at the bottom reflects inside the furnace and it is heated by collecting it on the installed sample. Since the inside is replaced with Ar gas, the molten metal does not react with air. A thermocouple is attached to the sample holder on which the sample is placed, so the furnace temperature can be measured in real time.

2.3 Experimental procedure

XRD (Rigaku Ultima IV) was used for the identification of the compound inside a sample. The XRD is an apparatus for performing an analysis by irradiating a crystalline substance with X-rays and reading diffracted X-rays. The phase state used Cu spheres at a speed of 2.0°/min in the range of 10° to 100°. EPMA (JEOL JXA 8900R) was used for elemental analysis and surface analysis of samples.

The experimental conditions using CLSM are shown in Table 1. The Fe-C-Cr alloy was heated to 1550°C in a high temperature imaging furnace and held for 5 minutes. The melted alloy was cooled at 60°C/min or 30°C/min, and the solidification process was observed. When M7C3 carbides began to crystallize out, they were quenched at a cooling rate of 2600°C/min to maintain the state of the structure. Optical Microscope (OM, OLYMPUS BX51) and Scanning Electron Microscope (SEM, JEOL JCM-5000) were used to observe the sample surface after conducting experiments by CLSM.

| Sample          | Fe-Cr alloy |
|-----------------|-------------|
| Temperature (°C)| 1350 1550 1400 100 |
| Slope (°C/min)  | 270 25 -60 -30 -2600 |
| Hold (sec)      | 0 300 0 0 |
| Frame count     | 1 5 15 5 |
3 Results and Discussion

3.1 Microstructure of the prepared sample

Figure 2 shows the internal structure of the Fe-C-Cr alloy produced using an arc furnace. CLSM was used for observation. Figure 2(a) shows the internal structure of the Fe-C-Cr alloy. Hexagonal crystals are crystallized throughout the sample. From the crystal shape, it is thought that M_7C_3 carbide is crystallized. Figure 2(b) is an enlargement of the position shown by the square in Figure 2(a). The size of M_7C_3 carbide was about 50 µm.

3.1.1 Analysis by XRD

Figure 3 shows the results of Fe-Cr-C alloy by XRD. Figure 4 shows the peak tendency of Cr carbides Cr_7C_3 and Cr_23C_6 in the database inside the XRD. The highest peak appears in the vicinity of 45° along the horizontal axis, and the peak is generally observed at similar positions. However, in Figure 3, a peak appears at 0° to 20°. This seems to be the peak due to the sample fixing material fixed when analyzing the Fe-Cr-C alloy. Hence, Cr carbides Cr_7C_3 and Cr_23C_6 exist in the prepared Fe-Cr-C alloy, and they mix with Fe to form M_7C_3 and M_23C_6.

3.1.2 Analysis by EPMA

From qualitative analysis and quantitative analysis by EPMA, it was confirmed that only three types of Fe, C, and Cr were present in the internal structure of the Fe-Cr-C alloy and contained no impurities. Figure 5 shows the results of area analysis by EPMA using Fe-Cr-C alloy. M_7C_3 carbides are observed, and the cross section is hexagonal, with a hollow at the center. Figure 5(a)–(c) shows the color maps of Fe, C, and Cr, respectively. Most of M_7C_3 carbides are occupied by Cr, and most of matrix structure and around M_7C_3 carbides are Fe. C was included evenly.
3.2 Solidification process observation of Fe-Cr-C alloy

3.2.1 Solidification process with a cooling rate of 60°C/min

Figure 6 shows the solidification process at the cooling rate of 60°C/min. In Figure 6(a), the corner portion of the M₇C₃ carbide is crystallized. As cooling is continued as shown in Figure 6(b)–(c), M₇C₃ carbides form a hexagon so as to connect the corners. However, not all M₇C₃ carbides form a perfect hexagon. In Figure 6(d), a gap is present between the M₇C₃ carbides at the positions shown by squares at 1466.5°C. This is considered to be the same phenomenon as the previous work by Liu et al. (2016). It was found by observation that while M₇C₃ carbides form hexagons, the corners are immediately connected, but it takes time to fill the gaps existing between the carbides. The size of the M₇C₃ carbides was about 50 µm, similar to those seen in Figure 2 and Figure 5. Figure 7 shows the solidification process in another observation plane. Rectangular crystals are stretched. This crystal is considered to be M₇C₃ carbide because it is the same as the temperature zone where M₇C₃ carbide crystallizes. Since M₇C₃ carbides preferentially grow along the c-axis, they become hexagonal columnar crystals. Therefore, this elongated shape is considered to be due to growth along the preferred orientation. From these results, the growth process of M₇C₃ carbide is predicted as shown in Figure 8. First, corner portions of M₇C₃ carbides grow in the preferential growth direction (Figure 6(a)), and hexagons are formed so as to connect the subsequent corners. M₇C₃ carbides have a preferred growth orientation in the c-axis, so they become elongated and become hexagonal columnar crystals.
3.2.2 Solidification process with a cooling rate of 30℃/min

Figure 9 shows the solidification process at a cooling rate of 30℃/min. M₇C₃ carbide crystallizes out as M₇C₃ is surrounded from the outside in Figure 9(a). Since this is a behavior not seen in the solidification process at a cooling rate of 60℃/min, it is considered that there is another crystallization behavior in the solidification process at a cooling rate of 30℃/min. When the cooling is continued as shown in Figure 9(b)–(d), it appears to form a hexagon, but as in Figure 6(d), a gap is present without forming a complete hexagon. The size of M₇C₃ carbide was about 50 µm. Now, Figure 10 shows the solidification process in another observation plane. The arrows indicate the direction in which the crystals were crystallized. From the place where it crystallized out in Figure 10(a), it can be seen that the crystals are growing in a hexagonal shape gradually. Therefore, it is considered that crystallization as in the growth model of Figure 8 is not performed. Thus, it is considered that there is another pattern in the crystallization behavior of the M₇C₃ carbide. An expected growth process of another pattern of M₇C₃ carbide is shown in Figure 11. M₇C₃ carbides grow diagonally to the observation plane. First, only a few areas can be observed, and most tissues are hidden in the fluid (Figure 10(a)). After that, the tissue in the liquid starts to grow, and eventually gets a hexagonal shape.

3.3 Structure observation after melting

Figure 12 shows how the sample surface after the melting experiment was observed using OM. The M₇C₃ carbide was found to have more crystallization in the transverse direction than in the longitudinal direction. Figure 13 shows how the sample surface after the experiment is observed by SEM.
shown in Figure 13(a), it can be seen that the entire sample is covered with M7C3 carbide in the lateral direction. The M7C3 carbide from the longitudinal direction is seen at the position enclosed by the square. The M7C3 carbides shown in Figure 13(b) appears to have a difference in height between the inside and the outside of the carbides. From the change of the height and the result of the melting experiment, it is thought that the formation of M7C3 carbide is formed preferentially in the outside and the formation of the inside is delayed.

Figure 12. OM image of Fe-C-Cr alloy

Figure 13. SEM image of Fe-C-Cr alloy (a) Sample Overall picture, and (b) Unevenness of M7C3

4 Conclusions

- As Fe-C-Cr alloy observed by the CLSM, the crystallization of M7C3 carbide was confirmed during the solidification process.
- Corners of the Hexagonal structure grew in the preferred orientation direction of M7C3 carbide at 60°C/min. After that, the M7C3 carbide formed to connect between the corners.
- At a cooling rate of 30°C/min, M7C3 carbides grew in oblique direction to the observation area. At the beginning, the structures were hidden in the fluid. But the structures started to grow, the shape of the hexagon became eventually.
- Since hexagonal structure arose a difference in height between the outside and the inside observed by OM and SEM. M7C3 carbides grew from the outside in the solidification process.

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