The stability, characteristics and magnetic properties of iron carbide from an oolitic hematite†

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Iron carbide (Fe₃C) is a magnetic material but it is not stable when it is prepared. Iron carbide was first prepared from high phosphorus oolitic hematite pellets using hydrogen reduction and was subsequently carburized with methane (CH₄). The products were then cooled down to room temperature using three different cooling conditions: fast cooling, furnace cooling, and heat preservation for 2 h with subsequent fast cooling. The results showed that the optimal reaction conditions for the preparation of Fe₃C from hydrogen reduced high phosphorus oolitic hematite are carburizing with CH₄ for 15 min at 1023 K, then fast cooling with argon gas, thus a degree of carburization of 95.12% can be obtained. The heat preservation promotes the self-decomposition of Fe₃C and the occurrence of multi-walled nano carbon fibers. The carbon nanotubes (CNTs) are in the form of rings, chains or nets, which can be attributed to the collisions between the CNTs. The nonuniform nano iron grains restrict the growth of CNTs deposited on the surface of metallic iron, and cause the bending of each CNT, which generates the chain or the net shape. Under fast cooling, the saturation magnetization declines from 82.59 to 61.97 emu g⁻¹ as the time of carburization with CH₄ increases from 10 to 30 min. The addition of the heat preservation and the control of the time of carburization with CH₄ can give the desired saturation magnetization. The Fe₃C prepared from high phosphorus oolitic hematite has relatively high magnetic properties.

Introduction

The high phosphorus oolitic hematite reserves (3–4 billion tons) in China cannot be utilized because of the difficulty of impurity removal.¹ Recently, several technologies for the deporphos- horization and also a functional material, can be widely used in many industrial fields.¹⁰–¹⁴ Nanosized Fe₃C has been mainly considered as a side product during the synthesis of carbon (C) structures, where the metallic iron is used as a catalyst, for example, in chemical vapor deposition,¹⁵ ultrasonic spray pyrolysis,¹⁶ laser ablation in organic solvent,¹⁷ and pyrolysis processes¹⁸ during the synthesis of carbon nanotubes (CNTs). Papers in the literature have presented countless procedures for the production of a plethora of nanoparticles and nanostructures (ranging from physical to chemical approaches, in water or solventless, by using a hard template or soft matter), thus, it is surprising that a synthetic pathway to produce Fe₃C nanoparticles in a reproducible, simple, and fast manner is still missing. This should contribute to the stability of an interstitial compound of Fe₃C, which can also be affected by the experimental conditions. Iron carbide in macro-scale was mainly prepared from high grade of iron ore such as Australian ore, Brazilian ore and other raw iron ores with little gangue, by reacting it with carburization gas, using a carburization process at low temperature (about 773–1123 K).¹⁹–²¹ The Fe₃C can be prepared from the iron ores in the atmosphere of hydrogen/methane (H₂/CH₄) and the previous work by the Authors’ research group has confirmed that 1023 K is the optimum temperature for the preparation of Fe₃C from high phosphorus oolitic hematite by H₂ reduction and subsequent carburization with CH₄, and also verified that the phosphorus in apatite was not reduced during the preparation of Fe₃C using...
thermodynamic calculations and related verification experiments. The results compensate well for the problem where phosphorus reduced from apatite can dissolve into the iron phase by the carbon-based reduction of high phosphorus hematite at relatively high temperature. Until now, the formation of Fe₃C mainly consists of three steps: reduction of iron oxide, CH₄ decomposition on the surface of the metallic iron, and the carburization of the metallic iron. Wang et al. has noted that the content of free carbon in the product prepared from high phosphorus oolitic hematite is relatively high because of the excessive time of CH₄ decomposition and improper cooling treatment of samples after carburizing, which greatly decreased the quality and purity of the carburized specimens. Thus, if the key factor to make full use of the reaction time of CH₄ decomposition and the cooling condition is well controlled, it may be promising in popularizing the use and the technology of Fe₃C preparation in H₂/CH₄.

In order to study the stability and the properties of Fe₃C from high phosphorus oolitic hematite at 1023 K, the high phosphorus oolitic hematite was first reduced in a H₂ atmosphere for the stepwise reduction of hematite to metallic iron, and subsequently with CH₄ for various reaction times (carburization of iron to Fe₃C), before finally applying three different cooling conditions: (i) fast cooling with a large flow rate of argon (Ar); (ii) with furnace cooling under Ar protection; and, (iii) heat preservation for 2 h with subsequent fast cooling by Ar. X-ray diffraction (XRD) and Mössbauer spectroscopy were used for the qualitative and quantitative analysis of iron and Fe₃C content in the products. Transmission electron microscopy (TEM) and vibrating sample magnetometry (VSM) were used for the characterization, and the magnetic property of the products from different cooling conditions were also investigated.

**Experimental**

**Raw materials**

A high phosphorus oolitic hematite from western Hubei province, China, was used as the raw material. Firstly, the iron ore was crushed into small pieces and then screened with a 100–120 particle size mesh. The fine ore powders were pelletized with some distilled water after an adequate mixing process in an agate mortar. The raw pellets were subsequently dried for 2 h at 373 K and then sintered for 15 min at 1173 K. The composition of the high phosphorus oolitic hematite is shown in Table 1.

**Experimental methods**

**Two-step process for the preparation of iron carbide.** The preparation of Fe₃C from high phosphorus oolitic hematite at 10⁷ Pa was carried out using a two-step process, which was H₂ reduction and subsequent carburization with CH₄. Firstly, the basket with a pellet, which was hung on a thermobalance (METTLER TOLEDO, MS105), was placed in a quartz reaction tube. The specific apparatus has been described in previous papers. Then, the tube was placed into a vertical resistance furnace at the predetermined temperature of 1023 K, and the Ar gas was introduced to flush air in to the reaction tube for 5 min. Afterwards the gas was switched to H₂ for the reduction of the pellet, with a flow rate of 150 cm³ min⁻¹. When the pellet was reduced to a certain degree of 0.9 at 1023 K, the H₂ gas was switched to CH₄ gas with a flow rate of 300 cm³ min⁻¹. When the carburization process finished, the sample was cooled down to room temperature with different cooling conditions. Then the sample obtained was ground in an agate mortar and vacuum packaged ready for the subsequent analysis. Additionally, these powder samples obtained from different cooling conditions and corresponding reaction times with CH₄ were marked with specific numbers, as shown in Tables 2 and 3.

**Analytical apparatus.** The powder samples obtained were analyzed using XRD (PANalytical X'Pert PRO MPD) with Cu Kα X-ray radiation. The Mössbauer spectra were recorded at room temperature with a constant acceleration and conventional standard transmission spectrometer (WissEl GmbH) with a ⁵⁷Co(Pd) source. The drive velocity scale was calibrated with a 25 µm thick pure α-Fe foil. All isomer shifts were quoted relative to the center of the α-Fe calibration spectrum. The experimental ⁵⁷Fe Mössbauer spectra were fitted using a least-squares regression fitting procedure. The TEM images were obtained using a 200 kV TEM (Tecnai G2 F20). Isothermal magnetization measurements at room temperature were performed in a commercial superconducting quantum interference device magnetometer (SQUID-VSM: Quantum Design).

**Results and discussion**

**Methane reaction time and cooling conditions on Fe₃C preparation – the weight change during the Fe₃C preparation**

Fig. 1 illustrates the weight change of the pellet sample at 1023 K in the Fe₃C preparation process. It shows that the weight of the pellet sample decreased with increase of time during the first 50 min of the H₂ reduction. After that, the reduced metallic iron was reacted with CH₄. In the first two minutes of reaction with CH₄, the weight decreased at a relatively slow rate. Then the weight increased gradually with the CH₄ reaction time from 52 to 80 min. It is interesting to find the phenomenon of the

| No. condition | A | B | C | D | E | F |
|---------------|---|---|---|---|---|---|
| Fast cooling  | A-1| B-1| C-1| D-1| E-1| F-1|
| Cooling in furnace | A-2| B-2| C-2| D-2| — | — |
| 1023 K holding and fast cooling | A-3| B-3| C-3| D-3| E-3| F-3|

| Composition of high phosphorus oolitic hematite (100–120 mesh) (mass%) |
|--------------------------|-----------|-----------|-----------|-----------|-----------|-----------|
| Fe | SiO₂ | Al₂O₃ | MgO | CaO | P |
| 50.23 | 14.6 | 5.08 | 0.62 | 2.34 | 0.77 |

| Time (min) | A | B | C | D | E | F |
|------------|---|---|---|---|---|---|
| 5          | 7 | 9 | 11 | 13 | 15 | 17 |
| 10         | 8 | 10 | 12 | 14 | 16 | 18 |
| 15         | 7 | 9 | 11 | 13 | 15 | 17 |
| 30         | 6 | 8 | 10 | 12 | 14 | 16 |
| 60         | 5 | 7 | 9 | 11 | 13 | 15 |
| 120        | 4 | 6 | 8 | 10 | 12 | 14 |

- **Table 1** The composition of high phosphorus oolitic hematite (100–120 mesh) (mass%)
- **Table 2** The specimen numbers and the reaction times with CH₄
- **Table 3** The specimen numbers and the cooling condition
continuous decrease of the weight of the pellet sample during the first 2 min of the reaction with CH4 gas. The result indicates that the reduction of iron(II) oxide (FeO) by the CH4 gas had occurred. In the previous work of our research group, a similar result was also achieved, and it is proposed that there is a mechanism of Fe3C formation which, with the additional process of the reduction of FeO to Fe by reaction with the CH4 gas, means that the Fe3C will not be generated.\textsuperscript{24} In order to evaluate the priority of process of FeO reacting with CH4, and the carburization of metallic Fe to Fe3C, the thermodynamics of reactions (1) to (3) has been calculated, and the results are shown in Table 4.\textsuperscript{25}

The three possible reactions that may occur are shown in Table 4. The temperatures at a Gibbs Free energy change (\(\Delta G^0\)) at 0 K show that the beginning temperature for reaction (3) is the lowest of 865.8 K and as the temperature is fixed at 1023 K, the Gibbs Free energy is \(-10 212.4, -6296.2\) and \(-21 806.1\) J mol\(^{-1}\), respectively, for reactions (1)-(3), which reveals that reaction (3) occurs most easily. To further evaluate the thermodynamics of each reaction, the volume percentage of CH4 for reactions (1) to (3) in equilibrium with FeO and Fe or Fe3C were plotted against temperature in Fig. 2. Fig. 2 shows that reaction (3) requires the minimum CH4 volume percentage at temperatures less than 975 K, which further illustrates that the formation of Fe3C by the carburization of metallic Fe with CH4 gas (reaction (3)) will occur after the reduction of the remaining FeO. Compared with reaction (3), reactions (1) and (2) require less volume percentage of CH4 in the temperature range from 975 to 1300 K. The results indicate that reactions (1) and (2) are more susceptible to occurring, which means that the carburization of metallic Fe to Fe3C is begun after the completion of the formation of metallic Fe from FeO. In addition, the required CH4 partial pressure for reaction (1) is lower than that of reaction (2), so the carbon monoxide (CO) gas is easier to generate than the CO2 gas during the deep reduction of FeO with CH4.

Table 4: The possible reactions between FeO and CH4

| No. | Reaction | \(\Delta G^0\) (J mol\(^{-1}\)) | \(T\) at \(\Delta G^0 = 0\) (K) |
|-----|----------|---------------------------|-----------------------------|
| 1   | 3FeO + CH4 = 3Fe+2H2O + CO | 274 181.6–278.427 | 984.8 |
| 2   | 4FeO + CH4 = 4Fe+2H2O + CO2 | 257 494.6–257.867 | 998.6 |
| 3   | 3Fe + CH4 = Fe3C+2H2 | 120 084–138.77 | 865.78 |

XRD qualitative analysis

Fig. S1 (ESI\textsuperscript{†}) shows the XRD patterns of sample groups A-1 to F-3 (referring to ASTM cards as shown in Table S1\textsuperscript{†}). The relative intensity of each characteristic peak qualitatively reflects the Fe3C content and metallic Fe content after reacting with CH4 for different times and using different cooling conditions. Fig. S1(a) (ESI\textsuperscript{†}) shows the XRD patterns of the specimens after reacting with CH4 for 5 min and using different cooling conditions. Two main phases: iron and quartz were in the three specimens. The analysis of the A- groups did not detect the phase of Fe3C, which means that Fe3C was not formed or not detected accurately in the measurements. In addition, the results also confirm that the deep reduction of reaction (2) exists, which is consistent with the results of previous research.\textsuperscript{24} After the unreacted FeO was totally converted into metallic Fe, reaction (3): 3Fe (s) + CH4 (g) = Fe3C (s) + 2H2 (g), occurred.

The XRD patterns for samples B-1 to D-3 are shown in Fig. S1(b) to S1(d).\textsuperscript{†} Research by Narkiewicz \textit{et al.}\textsuperscript{26} has shown that the \(2\theta = 37.63^\circ\) belongs to the (1 2 1) reflection of Fe3C (PDF No. 00-035-0772), and the relative intensity of the characteristic peak reflects well the relative content of the Fe3C. The \(2\theta = 82.34^\circ\) is assigned to the (2 1 1) reflection for the metallic Fe phase, and the intensity can qualitatively distinguish the relative content of Fe. From Fig. S1(b)–S1(d),\textsuperscript{†} the XRD patterns show that with the increase of reaction time with CH4, the intensity of the characteristic peak for the Fe3C phase increases, whereas a relative lower intensity of Fe3C exists after the sample is furnace cooled or if the cooling condition is heat preservation for 2 h and subsequent fast cooling. In addition, based on the intensity of characteristic peak at \(2\theta = 82.34^\circ\), the relative content of metallic Fe of the samples from furnace cooling or heat preservation at 1023 K for 2 h plus fast cooling, is higher than that from the fast cooling. According to the intensity of the characteristic peaks for Fe3C in different samples, it was preliminarily deduced that the optimal cooling conditions for the Fe3C preparation from high phosphorus oolite hematite should be the fast cooling. Meanwhile, the holding the temperature for 2 h plus the fast cooling could not improve the utilization of deposited carbon from the CH4 decomposition, but will lead to the self-decomposition of Fe3C instead.

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XRD analysis of samples from E-1 to F-3 is shown in Fig. S1(e) (ESI†). The difference of relative intensity of the characteristic peak for Fe₃C caused by the different cooling conditions in E-1 and E-3 or F-1 and F-3 was lower than that of B-1 to D-3, but the relative intensity of the Fe₃C peak was still decreased if the holding time at 1023 K was extended for 2 h. Additionally, compared with the relative intensity of the characteristic peak for graphite in B-1 to D-3, the graphite peak was relatively wide and the intensity was high. It can be deduced that with the extension of the reaction time with CH₄, the deposited carbon on the surface of the newly reduced metallic was increased, the self-decomposition of Fe₃C was accelerated, and a relatively fine grained graphite with a low degree of crystallization was also easily generated.

Quantitative Mössbauer analysis

The Mössbauer analysis is an accurate detection method for the distinction of different Fe phases, especially with the relatively low content iron-bearing phases. Fig. 3 shows the results of the quantitative analysis of the content of Fe-containing phases under different cooling conditions using Mössbauer spectroscopy. The Fe-containing phases include metallic Fe, Fe₃C and chlorite. The results in Fig. 3 reflect that the Fe₃C cannot be formed in the initial 5 min of the reaction with CH₄, which is consistent with the results from the XRD analysis shown in Fig. S1(a) (ESI†). Based on the results in Fig. 3, the Mössbauer analysis of the samples from A-1 to D-3, shows that when the cooling condition is fast cooling, the content of Fe₃C increases with the increase of the reaction time with CH₄. But the content of Fe₃C in the E-1 and F-1 samples shows a small decline, which suggests that the extension of the CH₄ reaction time from 30 min to 1 h and 2 h is not of benefit for the increase of the content of Fe₃C. This phenomenon is because the formation rate of Fe₃C is relatively slow when the CH₄ reaction time is too long, and the Fe₃C is a type of metastable interstitial compound. Chlorite also contains a small amount of iron, but the chlorite is chemically stable, so that the iron does not participate in the Fe₃C formation. The effect of the cooling conditions and the reaction time with CH₄ on the degree of carburization and the degree of decomposition of Fe₃C can be deduced from Fig. 3 and 4. Fig. 4 shows that the degree of carburization during the first 10 min of the reaction with CH₄ is up to 85.61%, which indicates that the carburization reaction is rapid after the reduction of FeO to metallic Fe completed. When the reaction time with CH₄ was increased to 15 min with subsequent fast cooling, the degree of carburization was 95.12%. The degree of carburization was 98.32% after 30 min reaction with CH₄ and then fast cooling. Compared with the fast cooling, the degree of self-decomposition of the samples obtained from the 10 min and 30 min reaction with CH₄ and the subsequent heat preservation at 1023 K for 2 h plus the fast cooling was 39.20% and 44.47%, respectively. The degree of self-decomposition of the other samples from the other reaction times with CH₄ and the subsequent heat preservation was approximately 28%.

The results of the Mössbauer analysis confirmed that fast cooling is the optimal cooling condition for the preparation of Fe₃C, and that the heat preservation at 1023 K for 2 h plus the fast cooling will promote the decomposition of Fe₃C. At 1023 K, the product is a mixture of ferrite iron and cementite (Fe₃C), and the heat preservation promotes the conversion of cementite to ferrite, resulting in the self-decomposition of Fe₃C: Fe₃C(s) = 3Fe(s) + C(s). Thus, the content of Fe₃C decreases. Thus, the optimum conditions for the preparation of Fe₃C from high phosphorus oolitic hematite are a temperature of 1023 K, reaction time of 15 min with CH₄ and then fast cooling to room temperature. Under these conditions, the corresponding degree of carburization is 95.12%, and the atom ratio of carbon to iron is 0.51.

The characterization and magnetic properties of the iron carbide

TEM analysis. Fig. 5(a) and (c) show the TEM images of specimens C-1 and C-3. Based on the dark area, which mainly consists of Fe₃C, the relative content of Fe₃C in C-1 is higher than that in C-3. The light area is mainly graphite. The number of multi-walled carbon nanotubes (MWCNTs) from CH₄ decomposition in C-3 is obviously more than that in C-1, which is consistent with the results of the XRD analysis. However, the ring, and chain shaped MWCNTs in C-3 are different to that found by Helminiak et al., because they used nanocrystal iron as the catalyst [aluminium oxide (Al₂O₃), silicon dioxide (SiO₂), calcium oxide (CaO) and other oxides as the precursors] for the
CH₄ decomposition. In this study, the microstructure of the fine dissemination of gangue minerals and hematite in high phosphorus oolitic hematite was not damaged during the H₂ reduction, and the Fe was mostly embedded in the oolide. Although the reduced metallic Fe was similar to the nano-iron grains, the iron was not evenly distributed. The non-uniform growth of nano iron grains contributed to the collisions of each CNT, and restricted the growth of CNTs deposited on the surface of metallic Fe, which caused the bending, and the rolling of the MWCNTs, and formed a chain or a ring shape during the aggregation of the nanotubes. From Fig. 5(b) and (d), the element in C-1 and C-3 samples is basically the same – they are both composed of Fe, C, silicon (Si), aluminium (Al) and oxygen (O), whereas the copper (Cu) element found comes from the standard Cu grid used in the TEM determination. Results from energy dispersive X-ray spectroscopy (EDS) analysis indicated that the Fe₃C and graphite are the main phases in the C-1 and C-3 samples, and the Si, Al and O were the element sources of the gangue minerals in the high phosphorus oolitic hematite. The coexistence of the elements of Fe and C with Si, Al also revealed that the phase of Fe₃C is still embedded closely with the gangue minerals, which explains why it is difficult to dephosphorize the Fe₃C products using magnetic separation, which has been reported before by Wang et al.28 The morphology of the graphite of C-3 grows much better than that of C-1, and the nano carbon fibers are much clearer in C-3 than C-1. These results are in agreement with those from Helminiak et al.,27 Kameya et al.,28 Narkiewicz et al.,29 and Luo et al.,31 and they revealed that the carbon from the decomposition of Fe₃C always presented as nano carbon fibers. Fig. 5(e) to 6(h) show the morphology of the samples that reacted with CH₄ for 60 min and 120 min, respectively, under different cooling conditions (E-1 to F-3 samples). Comparing the TEM images of E-1, F-1 with C-1, the groups of E and F show a higher degree of graphitization, and the crystallinity of the samples is better. Based on the TEM images of E-3 and F-3, the product samples which were treated with heat preservation for 2 h plus the fast cooling, could form the MWCNTs and also some nanotubes coated around the Fe₃C particles and the MWCNTs are onion-like. Because of self-decomposition of Fe₃C, the newly generated carbon fibers were fine with a relatively low degree of crystallization and have many defects.

Fig. 6(a) and (c) represent the TEM images of the product samples that were cooled using fast cooling and heat preservation for 2 h plus the fast cooling, respectively. The image for
C-3 shows that the boundary in the samples which underwent fast cooling is more ambiguous, and the number of the carbon fibers is more than that in C-1. Fig. 6(b) and (d) are the corresponding selected-area polycrystalline electron diffraction (SAPED) images corresponding to Fig. 6(a) and (c). The Fe₃C [(200) reflection, orthorhombic Fe₃C, JCPDS No. 00-035-0772], SiO₂ [(144) plane, hexagonal SiO₂, JCPDS No. 00-086-1628] and graphite [(100) reflection, hexagonal C, JCPDS No. 00-001-0640] are the main phases in the C-1 sample. While the Fe₃C [(200) and (212), orthorhombic], SiO₂ [(115) plane, hexagonal], graphite [(100) reflection, hexagonal] and metallic iron [(200) plane, cubic, JCPDS No. 01-089-7194] exist in the C-3 sample. The index results of the presence of metallic Fe in C-3 further reveal that the self-decomposition of Fe₃C is caused by the heat preservation, which is in good agreement with the previously mentioned XRD and Mössbauer analysis of the corresponding samples. Meanwhile, the dark area in Fig. 6(a) is larger than that in Fig. 6(c), which can also reflect the self-decomposition of Fe₃C by heat preservation.

**VSM magnetic analysis.** The curves of the magnetization versus magnetic field and the magnetization versus temperature of the product samples under different cooling conditions are shown in Fig. 7 and 8, and the specific magnetic property is shown in Table 5. The results in Table 5 indicate that the saturation magnetization \(M_s\) decreases from 82.59 to 61.97 emu g⁻¹ with the extension of the reaction time with CH₄ from 10 min to 30 min (from B-1 to D-1). Compared with F-1, the \(M_s\) of F-3 increases from 68.08 to 78.64 emu g⁻¹. The \(M_s\) of E-1 and F-1 is obviously less than that of B-1 and D-1, which may be attributed to the fact that much carbon was deposited from the CH₄ decomposition wrapping the iron or the Fe₃C particles (Fig. 5) when reacted with CH₄ for too long. The coercive force \((H_c)\) is approximately 20 Oe except for the result of 9.98 Oe obtained for B-1. The magnetic remanence \((M_r)\) increases a little when the CH₄ reaction lasts for more than 15 min, and the \(M_r\) is maintained at 0.2–0.3 emu g⁻¹ if the reaction time with CH₄ is longer than 30 min. If the reaction time with CH₄ is too long, it contributes to the decrease of the magnetic property of the samples. The possible reason for this is that with the increase of reaction time with CH₄, the carbon content of the sample will be increased, however, the \(M_s\) of metallic iron is higher than that of Fe₃C. Fig. 8 represents the magnetization versus temperature of the different specimens (C-1 and C-3) and shows that the two branches: zero field cooling (ZFC) and field cooling (FC) of each curve, do not merge at any point below room temperature. The result proves that the specimens are ferromagnetic and not superparamagnetic, because superparamagnetism is a phenomenon in which hysteresis does not occur at temperatures in the range of room temperature and above.³² Magnetization curves (C-1) and (C-3) represent the Fe₃C and the Fe₃C with some metallic Fe, respectively. As the amount of the metallic Fe phase increases (summarized in Fig. 4), the \(M_s\) value of C-3 is larger than that of C-1, which is because of the self-decomposition of Fe₃C by the different cooling conditions.

The magnetic property of the Fe₃C prepared from high phosphorus oolitic hematite was also assessed in a previous study.³⁴ The heat preservation for 2 h plus fast cooling caused the decline of the Fe₃C content while the content of metallic iron increased, thus the \(M_s\) will increase, but the carbon from the self-decomposition of Fe₃C will increase the degree of carbon coated on the surface of Fe₃C, which will hinder the magnetic property of the sample. Table 5 shows that the increase of the content of metallic Fe is more beneficial than that of MWNT coating on the surface of Fe₃C. The magnetic property of Fe₃C powder from the high phosphorus oolitic hematite is generally excellent.

**Table 5** The specific parameters of saturation magnetization \(M_s\), coercivity \((H_c)\) and remanent magnetization \(M_r\).

|     | \(M_s\) (emu g⁻¹) | \(H_c\) (Oe) | \(M_r\) (emu g⁻¹) |
|-----|------------------|--------------|-------------------|
| B-1 | 82.59            | 9.98         | 0.39              |
| C-1 | 68.08            | 19.80        | 0.56              |
| C-3 | 78.64            | 20.08        | 0.72              |
| D-1 | 61.97            | 19.91        | 0.25              |
| E-1 | 42.43            | 19.89        | 0.29              |
| F-1 | 32.26            | 19.91        | 0.21              |
Conclusions

(1) The optimal reaction conditions for the preparation of Fe₃C from a high phosphorus oolitic hematite are a temperature of 1023 K, a CH₄ reaction time of 15 min, and the cooling condition of fast cooling using Ar gas, and a degree of carburization of 95.12% can be obtained under these conditions.

(2) The heat preservation promotes the self-decomposition of Fe₃C and the occurrence of nano carbon fibers. With the increase of the reaction time of CH₄, the degree of graphitization increases, and the MWCNTs are in a ring, chain or net formation. The morphology is attributed to the fine dissemination of gangue minerals and the structure of hematite in the high phosphorus oolitic hematite was not damaged during the H₂ reduction, which means that the Fe was mostly embedded in the oolite. The nonuniform growth of nano iron grains contributes to the collisions of each carbon nanotube, and restricts the growth of CNTs deposited on the surface of metallic Fe, resulting in the bending, and generates the chain or the net shape.

(3) With the increase of the reaction time with CH₄, the saturation magnetization (Mₛ) of the Fe₃C powder containing gangue minerals decreases. Under fast cooling, the saturation magnetization declines from 82.59 to 61.97 emu g⁻¹ as the CH₄ reaction time increases from 10 to 30 min. Whereas the addition of the heat preservation can promote the increase of the Mₛ. The results are because of the free carbon content increase with the increase of reaction time with CH₄, and also the Mₛ of the metallic Fe is higher than that of Fe₃C. The Fe₃C prepared from high phosphorus oolitic hematite is a type of soft magnetic material with a relatively high magnetic property.

Conflicts of interest

There are no conflicts to declare.

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