



Cite this: *RSC Adv.*, 2017, 7, 44456

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

H. H. Wang,<sup>ab</sup> G. Q. Li,<sup>ID</sup> \*<sup>ab</sup> J. H. Ma<sup>b</sup> and D. Zhao<sup>a</sup>

Iron carbide (Fe<sub>3</sub>C) 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 (CH<sub>4</sub>). 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 Fe<sub>3</sub>C from hydrogen reduced high phosphorus oolitic hematite are carburizing with CH<sub>4</sub> 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 Fe<sub>3</sub>C 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<sup>-1</sup> as the time of carburization with CH<sub>4</sub> increases from 10 to 30 min. The addition of the heat preservation and the control of the time of carburization with CH<sub>4</sub> can give the desired saturation magnetization. The Fe<sub>3</sub>C prepared from high phosphorus oolitic hematite has relatively high magnetic properties.

Received 18th July 2017  
 Accepted 4th September 2017

DOI: 10.1039/c7ra07886b

[rsc.li/rsc-advances](http://rsc.li/rsc-advances)

## Introduction

The high phosphorus oolitic hematite reserves (3–4 billion tons) in China cannot be utilized because of the difficulty of impurity removal.<sup>1</sup> Recently, several technologies for the dephosphorization have been studied for the utilization of high phosphorus hematite, which were mainly: mineral processing,<sup>2,3</sup> deep reduction<sup>4–8</sup> and smelting technology.<sup>9</sup> Generally, mineral processing has problems of low iron recovery and dephosphorization ratio. The deep reduction leads to the reduction of apatite to phosphorus, which will dissolve into the iron phase, therefore, the decline of phosphorus content in the iron phase needs to be further investigated. The immature technology of smelting for dephosphorization and the energy consumption mean that the technology is unable to be widely applied.

Iron carbide (Fe<sub>3</sub>C), a type of raw material used for steel-making and also a functional material, can be widely used in

many industrial fields.<sup>10–14</sup> Nanosized Fe<sub>3</sub>C 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,<sup>15</sup> ultrasonic spray pyrolysis,<sup>16</sup> laser ablation in organic solvent<sup>17</sup> and pyrolysis processes<sup>18</sup> 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 Fe<sub>3</sub>C nanoparticles in a reproducible, simple, and fast manner is still missing. This should contribute to the stability of an interstitial compound of Fe<sub>3</sub>C, 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).<sup>19–21</sup> The Fe<sub>3</sub>C can be prepared from the iron ores in the atmosphere of hydrogen/methane (H<sub>2</sub>/CH<sub>4</sub>) and the previous work by the Authors' research group has confirmed that 1023 K is the optimum temperature for the preparation of Fe<sub>3</sub>C from high phosphorus oolitic hematite by H<sub>2</sub> reduction and subsequent carburization with CH<sub>4</sub>, and also verified that the phosphorus in apatite was not reduced during the preparation of Fe<sub>3</sub>C using

<sup>a</sup>The State Key Laboratory of Refractories and Metallurgy, Wuhan University of Science and Technology, PO Box 185, 947 Heping Avenue, Qingshan District, Wuhan, Hubei, 430081, PR China. E-mail: [liguangqiang@wust.edu.cn](mailto:liguangqiang@wust.edu.cn); Fax: +86 2768862665; Tel: +86 2768862665

<sup>b</sup>Key Laboratory for Ferrous Metallurgy and Resources Utilization of Ministry of Education, Wuhan University of Science and Technology, Wuhan, Hubei, 430081, PR China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra07886b



thermodynamic calculations and related verification experiments.<sup>22–24</sup> 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 Fe<sub>3</sub>C mainly consists of three steps: reduction of iron oxide, CH<sub>4</sub> decomposition on the surface of the metallic iron, and the carburization of the metallic iron.<sup>20,21</sup> Wang *et al.*<sup>24</sup> 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 CH<sub>4</sub> 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 CH<sub>4</sub> decomposition and the cooling condition is well controlled, it may be promising in popularizing the use and the technology of Fe<sub>3</sub>C preparation in H<sub>2</sub>/CH<sub>4</sub>.

In order to study the stability and the properties of Fe<sub>3</sub>C from high phosphorus oolitic hematite at 1023 K, the high phosphorus oolitic hematite was first reduced in a H<sub>2</sub> atmosphere for the stepwise reduction of hematite to metallic iron, and subsequently with CH<sub>4</sub> for various reaction times (carburization of iron to Fe<sub>3</sub>C), 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 Fe<sub>3</sub>C 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 Fe<sub>3</sub>C from high phosphorus oolitic hematite at 10<sup>5</sup> Pa was carried out using a two-step process, which was H<sub>2</sub> reduction and subsequent carburization with CH<sub>4</sub>. Firstly, the

Table 1 The composition of high phosphorus oolitic hematite (100–120 mesh) (mass%)

Composition	Fe	SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	MgO	CaO	P
Content	50.23	14.6	5.08	0.62	2.34	0.77

Table 2 The specimen numbers and the reaction times with CH<sub>4</sub>

No.	A	B	C	D	E	F
Time (min)	5	10	15	30	60	120

Table 3 The specimen numbers and the cooling condition

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

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.<sup>22–24</sup> 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 H<sub>2</sub> for the reduction of the pellet, with a flow rate of 150 cm<sup>3</sup> min<sup>-1</sup>. When the pellet was reduced to a certain degree of 0.9 at 1023 K, the H<sub>2</sub> gas was switched to CH<sub>4</sub> gas with a flow rate of 300 cm<sup>3</sup> min<sup>-1</sup>. 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 CH<sub>4</sub> 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 K $\alpha$  X-ray radiation. The Mössbauer spectra were recorded at room temperature with a constant acceleration and conventional standard transmission spectrometer (WissEl GmbH) with a <sup>57</sup>Co(Pd) source. The drive velocity scale was calibrated with a 25  $\mu$ m thick pure  $\alpha$ -Fe foil. All isomer shifts were quoted relative to the center of the  $\alpha$ -Fe calibration spectrum. The experimental <sup>57</sup>Fe 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 Fe<sub>3</sub>C preparation – the weight change during the Fe<sub>3</sub>C preparation

Fig. 1 illustrates the weight change of the pellet sample at 1023 K in the Fe<sub>3</sub>C preparation process. It shows that the weight of the pellet sample decreased with increase of time during the first 50 min of the H<sub>2</sub> reduction. After that, the reduced metallic iron was reacted with CH<sub>4</sub>. In the first two minutes of reaction with CH<sub>4</sub>, the weight decreased at a relatively slow rate. Then the weight increased gradually with the CH<sub>4</sub> reaction time from 52 to 80 min. It is interesting to find the phenomenon of the



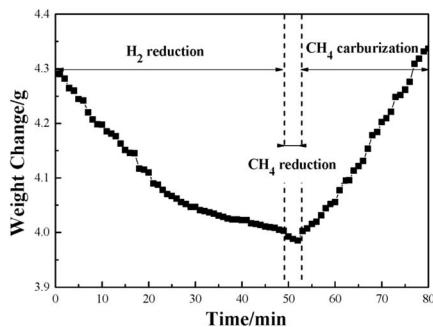


Fig. 1 The curve of the weight change versus reaction time at 1023 K.

continuous decrease of the weight of pellet sample during the first 2 min of the reaction with  $\text{CH}_4$  gas. The result indicates that the reduction of iron(II) oxide ( $\text{FeO}$ ) by the  $\text{CH}_4$  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  $\text{Fe}_3\text{C}$  formation which, with the additional process of the reduction of  $\text{FeO}$  to  $\text{Fe}$  by reaction with the  $\text{CH}_4$  gas, means that the  $\text{Fe}_3\text{C}$  will not be generated.<sup>24</sup> In order to evaluate the priority of process of  $\text{FeO}$  reacting with  $\text{CH}_4$ , and the carburization of metallic  $\text{Fe}$  to  $\text{Fe}_3\text{C}$ , the thermodynamics of reactions (1) to (3) has been calculated, and the results are shown in Table 4.<sup>25</sup>

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  $-21\,806.1$   $\text{J mol}^{-1}$ , 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  $\text{CH}_4$  for reactions (1) to (3) in equilibrium with  $\text{FeO}$  and  $\text{Fe}$  or  $\text{Fe}_3\text{C}$  were plotted against temperature in Fig. 2. Fig. 2 shows that reaction (3) requires the minimum  $\text{CH}_4$  volume percentage at temperatures less than 975 K, which further illustrates that the formation of  $\text{Fe}_3\text{C}$  by the carburization of metallic  $\text{Fe}$  with  $\text{CH}_4$  gas (reaction (3)) will occur after the reduction of the remaining  $\text{FeO}$ . Compared with reaction (3), reactions (1) and (2) require less volume percentage of  $\text{CH}_4$  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  $\text{Fe}$  to  $\text{Fe}_3\text{C}$  is begun after the completion of the formation of metallic  $\text{Fe}$  from  $\text{FeO}$ . In addition, the required  $\text{CH}_4$  partial pressure for reaction (1) is lower than that of reaction (2), so the carbon monoxide ( $\text{CO}$ ) gas is easier to generate than the  $\text{CO}_2$  gas during the deep reduction of  $\text{FeO}$  with  $\text{CH}_4$ .

Table 4 The possible reactions between  $\text{FeO}$  and  $\text{CH}_4$

No.	Reaction	$\Delta G^0$	$T$ at $\Delta G^0 = 0$ (K)
1	$3\text{FeO} + \text{CH}_4 = 3\text{Fe} + 2\text{H}_2\text{O} + \text{CO}$	$274\,181.6 - 278.42T$	984.8
2	$4\text{FeO} + \text{CH}_4 = 4\text{Fe} + 2\text{H}_2\text{O} + \text{CO}_2$	$257\,494.6 - 257.86T$	998.6
3	$3\text{Fe} + \text{CH}_4 = \text{Fe}_3\text{C} + 2\text{H}_2$	$120\,084 - 138.7T$	865.78

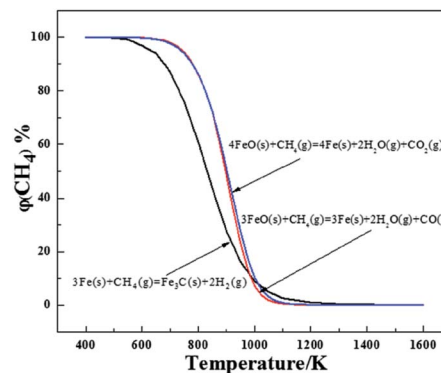


Fig. 2 The equilibrium relationship between the volume percentage of  $\text{CH}_4$  and the temperature of reactions (1) to (3).

### XRD qualitative analysis

Fig. S1 (ESI<sup>†</sup>) shows the XRD patterns of sample groups A-1 to F-3 (referring to ASTM cards as shown in Table S1<sup>†</sup>). The relative intensity of each characteristic peak qualitatively reflects the  $\text{Fe}_3\text{C}$  content and metallic  $\text{Fe}$  content after reacting with  $\text{CH}_4$  for different times and using different cooling conditions. Fig. S1(a) (ESI<sup>†</sup>) shows the XRD patterns of the specimens after reacting with  $\text{CH}_4$  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  $\text{Fe}_3\text{C}$ , which means that  $\text{Fe}_3\text{C}$  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.<sup>24</sup> After the unreacted  $\text{FeO}$  was totally converted into metallic  $\text{Fe}$ , reaction (3):  $3\text{Fe}(\text{s}) + \text{CH}_4(\text{g}) = \text{Fe}_3\text{C}(\text{s}) + 2\text{H}_2(\text{g})$ , occurred.

The XRD patterns for samples B-1 to D-3 are shown in Fig. S1(b) to S1(d).<sup>†</sup> Research by Narkiewicz *et al.*<sup>26</sup> has shown that the  $2\theta = 37.63^\circ$  belongs to the (1 2 1) reflection of  $\text{Fe}_3\text{C}$  (PDF No. 00-035-0772), and the relative intensity of the characteristic peak reflects well the relative content of the  $\text{Fe}_3\text{C}$ . The  $2\theta = 82.34^\circ$  is assigned to the (2 1 1) reflection for the metallic  $\text{Fe}$  phase, and the intensity can qualitatively distinguish the relative content of  $\text{Fe}$ . From Fig. S1(b)–S1(d),<sup>†</sup> the XRD patterns show that with the increase of reaction time with  $\text{CH}_4$ , the intensity of the characteristic peak for the  $\text{Fe}_3\text{C}$  phase increases, whereas a relative lower intensity of  $\text{Fe}_3\text{C}$  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  $\text{Fe}$  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  $\text{Fe}_3\text{C}$  in different samples, it was preliminarily deduced that the optimal cooling conditions for the  $\text{Fe}_3\text{C}$  preparation from high phosphorus oolitic 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  $\text{CH}_4$  decomposition, but will lead to the self-decomposition of  $\text{Fe}_3\text{C}$  instead.



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 Fe<sub>3</sub>C 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 Fe<sub>3</sub>C 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 CH<sub>4</sub>, the deposited carbon on the surface of the newly reduced metallic was increased, the self-decomposition of Fe<sub>3</sub>C 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, Fe<sub>3</sub>C and chlorite. The results in Fig. 3 reflect that the Fe<sub>3</sub>C cannot be formed in the initial 5 min of the reaction with CH<sub>4</sub>, 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 Fe<sub>3</sub>C increases with the increase of the reaction time with CH<sub>4</sub>. But the content of Fe<sub>3</sub>C in the E-1 and F-1 samples shows a small decline, which suggests that the extension of the CH<sub>4</sub> reaction time from 30 min to 1 h and 2 h is not of benefit for the increase of the content of Fe<sub>3</sub>C. This phenomenon is because the formation rate of Fe<sub>3</sub>C is relatively slow when the CH<sub>4</sub> reaction time is too long, and the Fe<sub>3</sub>C 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 Fe<sub>3</sub>C formation. The effect of the cooling conditions and the reaction time with CH<sub>4</sub> on the degree of carburization and the degree of decomposition of Fe<sub>3</sub>C can be deduced from Fig. 3

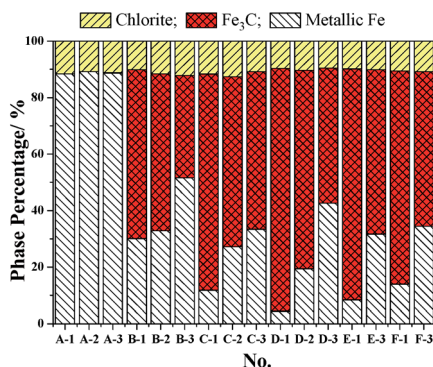


Fig. 3 The relationship between cooling conditions and the content of iron-bearing phases.

and 4. Fig. 4 shows that the degree of carburization during the first 10 min of the reaction with CH<sub>4</sub> 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 CH<sub>4</sub> 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 CH<sub>4</sub> 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 CH<sub>4</sub> 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 CH<sub>4</sub> 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 Fe<sub>3</sub>C, and that the heat preservation at 1023 K for 2 h plus the fast cooling will promote the decomposition of Fe<sub>3</sub>C. At 1023 K, the product is a mixture of ferrite iron and cementite (Fe<sub>3</sub>C), and the heat preservation promotes the conversion of cementite to ferrite, resulting in the self-decomposition of Fe<sub>3</sub>C: Fe<sub>3</sub>C (s) = 3Fe (s) + C (s). Thus, the content of Fe<sub>3</sub>C decreases. Thus, the optimum conditions for the preparation of Fe<sub>3</sub>C from high phosphorus oolitic hematite are a temperature of 1023 K, reaction time of 15 min with CH<sub>4</sub> 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 Fe<sub>3</sub>C, the relative content of Fe<sub>3</sub>C 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 CH<sub>4</sub> 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.*,<sup>27</sup> because they used nanocrystal iron as the catalyst [aluminium oxide (Al<sub>2</sub>O<sub>3</sub>), silicon dioxide (SiO<sub>2</sub>) calcium oxide (CaO) and other oxides as the precursors] for the

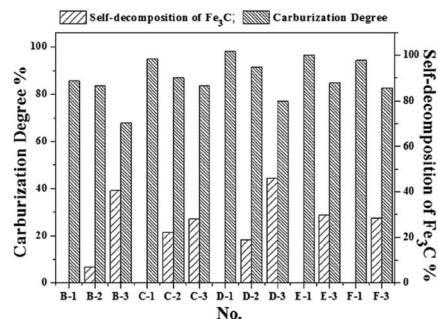


Fig. 4 The relationship between cooling conditions and the content of iron-bearing phases.



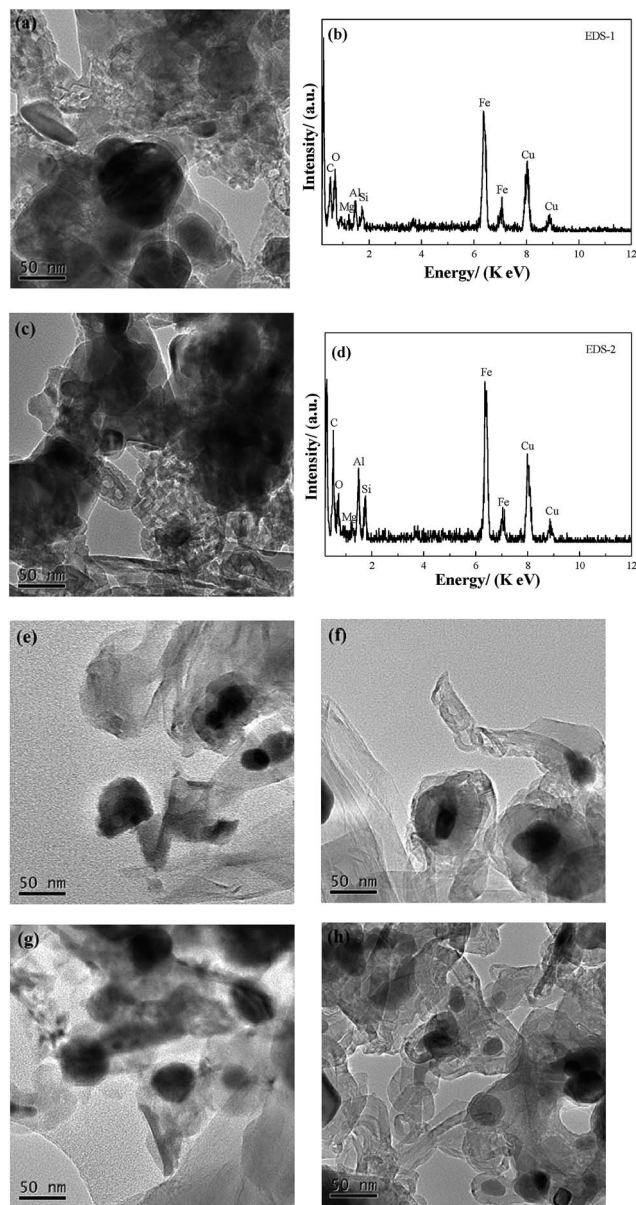


Fig. 5 Results of TEM and EDS analysis of the samples from C-1 to F-3; (a) and (c) are the TEM images of C-1 and C-3; (b) and (d) are the corresponding EDS analysis of (a) and (c); (e) to (h) are the samples of E-1, E-3, F-1 and F-3, respectively.

$\text{CH}_4$  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  $\text{H}_2$  reduction, and the Fe was mostly embedded in the ooid. 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  $\text{Fe}_3\text{C}$  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  $\text{Fe}_3\text{C}$  is still embedded closely with the gangue minerals, which explains why it is difficult to dephosphorize the  $\text{Fe}_3\text{C}$  products using magnetic separation, which has been reported before by Wang *et al.*<sup>28</sup> 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.*,<sup>27</sup> Kameya *et al.*,<sup>29</sup> Narkiewicz *et al.*,<sup>30</sup> and Luo *et al.*,<sup>31</sup> and they revealed that the carbon from the decomposition of  $\text{Fe}_3\text{C}$  always presented as nano carbon fibers. Fig. 5(e) to 6(h) show the morphology of the samples that reacted with  $\text{CH}_4$  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  $\text{Fe}_3\text{C}$  particles and the MWCNTs are onion-like. Because of self-decomposition of  $\text{Fe}_3\text{C}$ , 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

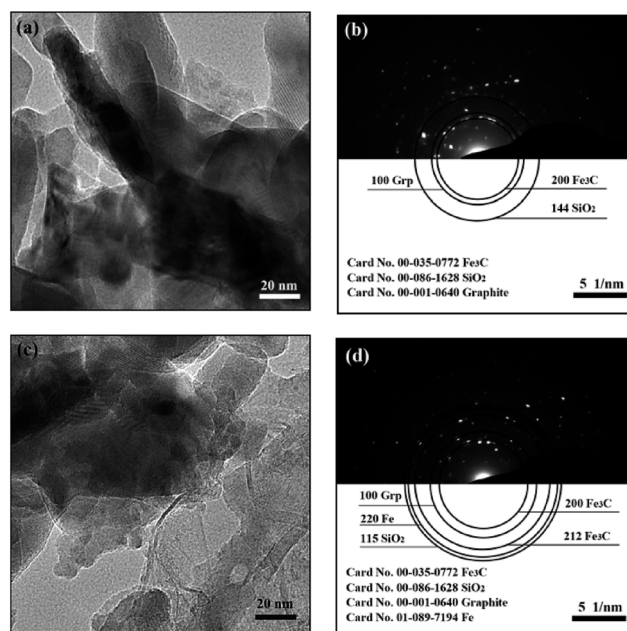


Fig. 6 The TEM images and the corresponding selected-area polycrystalline electron diffraction of C-1 (a, b) and C-3 (c, d) specimens.



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  $\text{Fe}_3\text{C}$  [(200) reflection, orthorhombic  $\text{Fe}_3\text{C}$ , JCPDS No. 00-035-0772],  $\text{SiO}_2$  [(144) plane, hexagonal  $\text{SiO}_2$ , 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  $\text{Fe}_3\text{C}$  [(200) and (212), orthorhombic],  $\text{SiO}_2$  [(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 Fe<sub>3</sub>C 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  $\text{Fe}_3\text{C}$  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  $\text{emu g}^{-1}$  with the extension of the reaction time with  $\text{CH}_4$  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  $\text{emu g}^{-1}$ . 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  $\text{CH}_4$  decomposition wrapping the iron or the  $\text{Fe}_3\text{C}$  particles (Fig. 5) when reacted with  $\text{CH}_4$  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  $\text{CH}_4$  reaction lasts for more than 15 min, and the  $M_r$  is maintained at 0.2–0.3  $\text{emu g}^{-1}$  if the reaction time with  $\text{CH}_4$  is longer than 30 min. If the reaction time with  $\text{CH}_4$  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

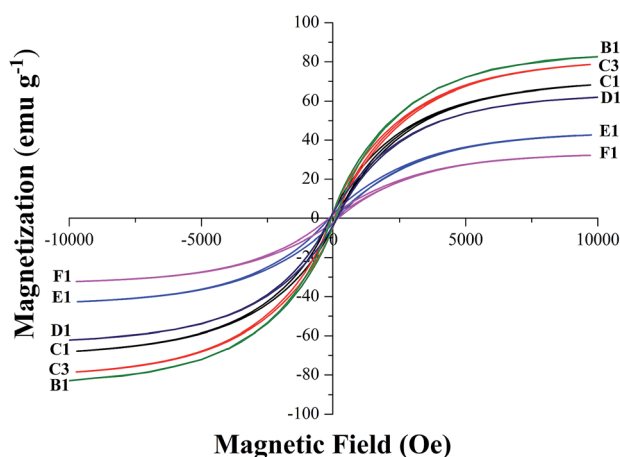


Fig. 7 The magnetic hysteresis loops recorded for the B1 to F1.

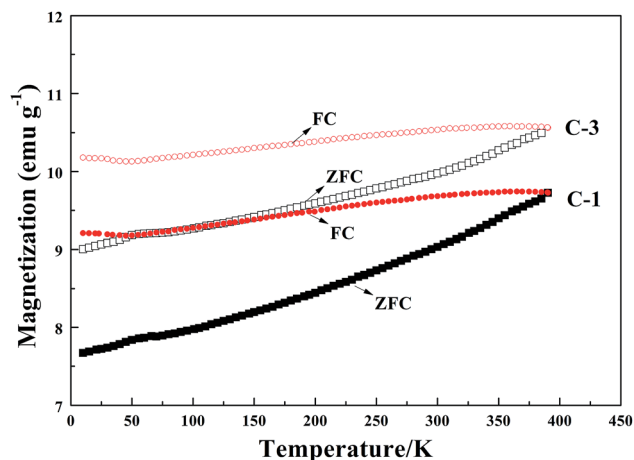


Fig. 8 The magnetization versus temperature of the specimens C-1 and C-3.

reaction time with  $\text{CH}_4$ , the carbon content of the sample will be increased, however, the  $M_s$  of metallic iron is higher than that of  $\text{Fe}_3\text{C}$ . 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.<sup>32</sup> Magnetization curves (C-1) and (C-3) represent the  $\text{Fe}_3\text{C}$  and the  $\text{Fe}_3\text{C}$  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  $\text{Fe}_3\text{C}$  by the different cooling conditions.

The magnetic property of the  $\text{Fe}_3\text{C}$  prepared from high phosphorus oolitic hematite was also assessed in a previous study.<sup>24</sup> The heat preservation for 2 h plus fast cooling caused the decline of the  $\text{Fe}_3\text{C}$  content while the content of metallic iron increased, thus the  $M_s$  will increase, but the carbon from the self-decomposition of  $\text{Fe}_3\text{C}$  will increase the degree of carbon coated on the surface of  $\text{Fe}_3\text{C}$ , 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 MWCNT coating on the surface of  $\text{Fe}_3\text{C}$ . The magnetic property of  $\text{Fe}_3\text{C}$  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 ( $M_r$ )

	$M_s$ ( $\text{emu g}^{-1}$ )	$H_c$ (Oe)	$M_r$ ( $\text{emu g}^{-1}$ )
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 Fe<sub>3</sub>C from a high phosphorus oolitic hematite are a temperature of 1023 K, a CH<sub>4</sub> 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 Fe<sub>3</sub>C and the occurrence of nano carbon fibers. With the increase of the reaction time of CH<sub>4</sub>, 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 H<sub>2</sub> 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 CH<sub>4</sub>, the saturation magnetization ( $M_s$ ) of the Fe<sub>3</sub>C powder containing gangue minerals decreases. Under fast cooling, the saturation magnetization declines from 82.59 to 61.97 emu g<sup>-1</sup> as the CH<sub>4</sub> reaction time increases from 10 to 30 min. Whereas the addition of the heat preservation can promote the increase of the  $M_s$ . The results are because of the free carbon content increase with the increase of reaction time with CH<sub>4</sub>, and also the  $M_s$  of the metallic Fe is higher than that of Fe<sub>3</sub>C. The Fe<sub>3</sub>C 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.

## Acknowledgements

This work is financially supported by the National Natural Science Foundation of China (No. 51374159) and the Open Foundation of State Key Laboratory of Advanced Metallurgy, University of Science and Technology Beijing (No. KF12-06).

## Notes and references

- 1 D. Wei, *Mod. Min.*, 2011, 5, 9, in Chinese.
- 2 Y. B. Dong, M. Qiang, Z. Y. Duan, J. J. Xu and X. Wang, *Metal. Mine*, 2010, 404, 62, in Chinese.
- 3 Y. B. Long and Y. S. Zhang, *Multipurp. Util. Miner. Resour.*, 2011, 1, 3, in Chinese.
- 4 H. Han, D. Duan, P. Yuan and S. Chen, *Ironmaking Steelmaking*, 2015, 42, 542.
- 5 G. F. Li, Y. X. Han, P. Gao and Y. S. Sun, *Ironmaking Steelmaking*, 2016, 43, 163.
- 6 Y. Sun, P. Gao, Y. Han and D. Ren, *Ind. Eng. Chem. Res.*, 2013, 52, 2323.
- 7 Y. Li, T. Sun, J. Kou, Q. Guo and C. Xu, *Miner. Process. Extr. Metall. Rev.*, 2014, 35, 66.
- 8 Z. H. Li, Y. X. Han, Y. S. Sun and P. Gao, *J. Northeast. Univ.*, 2015, 36, 1757, in Chinese.
- 9 J. Wu, Z. J. Wen and M. J. Cen, *Steel Res. Int.*, 2011, 82, 494.
- 10 A. N. Conejo and G. P. Martins, *Ironmaking Steelmaking*, 1999, 26, 111.
- 11 T. Akiyama, A. Miyazaki, H. Nakanishi, M. Hisa and A. Tsutsumi, *Int. J. Hydrogen Energy*, 2004, 29, 721.
- 12 A. N. Conejo and R. S. Estrada, *Steel Res. Int.*, 2007, 78, 3.
- 13 T. Herranz, S. Rojas, F. J. Pérez-Alonso, M. Ojeda, P. Terreros and J. L. G. Fierro, *J. Catal.*, 2006, 8199, 40199.
- 14 M. Bahgat, *J. Mater. Sci. Technol.*, 2006, 22, 423.
- 15 C. P. Deck and K. Vecchio, *Carbon*, 2006, 44, 267.
- 16 J. D. Atkinson, M. E. Fortunato, S. A. Dastgheib, M. Rostam-Abadi, M. J. Rood and K. S. Suslick, *Carbon*, 2011, 49, 587.
- 17 V. Amendola, P. Riello and M. Meneghetti, *J. Phys. Chem. C*, 2011, 115, 5140.
- 18 A. K. Schaper, H. Q. Hou, A. Greiner and F. Phillipp, *J. Catal.*, 2004, 222, 250.
- 19 G. Li, H. Ni, Q. Shen and F. Tsukihashi, *ISIJ Int.*, 2006, 46, 981.
- 20 J. Zhang and O. Ostrovski, *ISIJ Int.*, 2001, 41, 333.
- 21 H. W. Ni, F. S. Zhang, D. Q. Cang and J. P. Jiang, *J. Iron Steel Res. Int.*, 2001, 13, 1, in Chinese.
- 22 G. Q. Li, H. H. Wang, J. Yang and J. H. Ma, *Adv. Mater. Res.*, 2014, 881–883, 98.
- 23 H. H. Wang, G. Q. Li, J. Yang, J. H. Ma and B. S. Khan, *Metall. Mater. Trans. B*, 2016, 47, 2571.
- 24 H. H. Wang, G. Q. Li, J. H. Ma and D. Zhao, *RSC Adv.*, 2017, 7, 3921.
- 25 D. Stull and H. Prophet, *JANAF Thermochemical Tables*, US Government Printing Office, Washington, DC, 2nd edn, 1971.
- 26 U. Narkiewicz, W. Arabczyk, W. Konicki and A. Pattek-Janczyk, *J. Mater. Res.*, 2005, 20, 386.
- 27 A. Helminiak, E. Mijowska and W. Arabczyk, *Mater. Sci.*, 2013, 31, 29.
- 28 H. H. Wang, G. Q. Li, J. H. Ma, Y. X. Hu and D. Zhao, *J. Wuhan Univ. Sci. Technol.*, 2017, 40, 11, in Chinese.
- 29 Y. Kameya and K. Hanamura, *Carbon*, 2012, 50, 3503.
- 30 U. Narkiewicz, W. Arabczyk, I. Pelech, N. Guskos, J. Typek, M. Maryniak, M. J. Woźniak, H. Matysiak and K. J. Kurzydłowski, *Mater. Sci.*, 2006, 24, 1067.
- 31 N. Luo, X. Li, X. Wang, H. Yan, C. Zhang and H. Wang, *Carbon*, 2010, 48, 3858.
- 32 D. L. Huber, *Small*, 2005, 1, 482.

