Superconducting properties of carbonaceous chemical doped MgB2

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Abstract
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Superconducting Properties of Carbonaceous Chemical Doped MgB$_2$

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1. Introduction

The discovery of superconductivity in magnesium diboride (MgB$_2$: 39 K, in January 2001) (Nagamatsu et al., 2001) has generated enormous interest and excitement in the superconductivity community and the world in general, but especially among researchers into superconductivity in non-oxide and boron related compounds. MgB$_2$ possesses an AlB$_2$-type hexagonal structure (space group $P6/mmm$) with alternating boron honeycomb planes and magnesium triangular planes, as shown in Fig. 1. Each Mg atom is located at the center of a hexagon formed by boron, and it donates its electrons to the boron planes; hence, the B-B bonding is strongly anisotropic. The unit cell parameters are $a = 0.3086$ nm and $c = 0.3524$ nm at room temperature. These values of lattice parameters for MgB$_2$ are in the middle of the values of lattice parameters of AlB$_2$-type compounds. Owing to the simple hexagonal structure with space group $P6/mmm$, four optical modes at the $\Gamma$ point of the Brillouin zone are predicted for MgB$_2$ (An & Pickett, 2001): a silent $B_{1g}$ mode (at 87.1 meV, ~700 cm$^{-1}$), the $E_{2g}$ Raman mode (at 74.5 meV, ~600 cm$^{-1}$), and the infrared active $E_{1u}$ (at 40.7 meV, ~330 cm$^{-1}$) and $A_{2u}$ (at 49.8 meV, ~400 cm$^{-1}$) modes. The $E_{2g}$ mode is responsible for the high transition temperature, $T_c$, in MgB$_2$.

Fig. 1. Hexagonal structure of MgB$_2$ with space group $P6/mmm$ (Nagamatsu et al., 2001)

Further studies based on a number of experimental techniques, such as angle-resolved photoemission spectroscopy (ARPES), the de Haas-van Alphen effect, and Hall resistivity
measurements, have found that MgB₂ exhibits a rich multiple-band structure. These results are in agreement with band structure calculations and reveal strongly two-dimensional \( sp, p_y(\sigma) \) bands, as well as three-dimensional \( p_z(\pi) \) bands. The identification of MgB₂ as a two gap superconductor has resulted in much research associated with the spectroscopy of this material. It has become generally accepted that the larger gap is associated with the 2D \( \sigma \) bands arising from the boron planes, which has the value of \( \Delta_\sigma \approx 7.069 \) meV, while the 3D \( \pi \) bands have a gap of \( \Delta_\pi \approx 2.70 \) meV (Bouquet et al., 2001; Kortus et al., 2001).

MgB₂ has been fabricated in bulks, single crystals, thin films, tapes, and wires for different applications (Eisterer & Weber, 2009). In addition to the relatively high critical transition temperature, \( T_c \), and the simple crystal structure, MgB₂ possesses a large coherence length, high critical current density, and transparency of grain boundaries to current flow. The \textit{in-situ} route seems to be the most promising method to improve the upper critical field, \( H_{c2} \), and the critical current density, \( J_c \), performance of MgB₂. MgB₂ is a promising superconductor for high-magnetic field applications because of its already high \( J_c \). The grain boundaries in MgB₂ do not significantly degrade \( J_c \) and even serve as pinning centers, which is different from the weak-link effects in high-\( T_c \) superconductors.

For single-gap dirty limit superconductors, the upper critical field \( H_{c2}(0) = 0.69T_c (dH_{c2}/dT)_{T_c} \) and \( (dH_{c2}/dT)_{T_c} \propto \rho_n \), where \( \rho_n \) is the normal state resistivity (Werthamer et al., 1966); therefore, \( H_{c2} \) increases with \( \rho_n \), which can be achieved by adding impurities and defects into the superconductor. Gurevich pointed out that the two-band superconductor MgB₂ can be understood as a weakly-coupled bilayer in which two thin films corresponding to the \( \sigma \) and \( \pi \) bands are in contact through Josephson coupling (Gurevich, 2007). Using the dirty-limit, weak-coupling, multiband Bardeen Cooper Schrieffer (BCS) model and taking into account both interband and intraband scattering by nonmagnetic impurities, Gurevich showed that the temperature dependence of \( H_{c2}(T) \) is influenced by whether the \( \sigma \) bands or \( \pi \) bands are dirtier, making it very different from the temperature dependence in the one-band theory (Gurevich, 2003). The global \( H_{c2}(T) \) of the bilayer is dominated by the layer with the higher \( H_{c2} \). If the \( \pi \) layer is dirtier, it will have higher \( H_{c2} \) at low temperature, even though its \( T_c \) is much lower. As a result, an upturn in the global \( H_{c2}(T) \) occurs at low temperature. \( H_{c2}(0) \) of MgB₂ can exceed 0.69\( T_c (dH_{c2}/dT)_{T_c} \) considerably because of the existence of the two bands. Considering the electron–phonon coupling effect, Gurevich argued that the strong coupling paramagnetic limit in MgB₂ can be as high as 130 T; thus, there is still room for further enhancement of \( H_{c2} \) by engineering the \( \sigma \)- and \( \pi \)-band scattering (Gurevich, 2007). The high \( H_{c2} \) in MgB₂ is very attractive for high-magnetic-field applications. The \( H_{c2} \) behavior described by Gurevich has been observed in experimental results. For example, Braccini et al. observed different types of temperature dependence of \( H_{c2} \), including the anomalous upturn at low temperature, reflecting different multiband scattering in thin film samples from various groups, with disorder introduced in different ways. The value of \( H_{c2} \) in carbon-doped thin films has reached over 60 T at low temperature, approaching the BCS paramagnetic limit of 65 T (Braccini et al., 2005).

The depairing current density, \( J_d \), is \( \approx 8.7 \times 10^8 \) A/cm² for pure MgB₂, as estimated from the Ginzburg-Landau (GL) formula:

\[
J_d = \Phi_0 / \left[ 3 / \left( \sqrt{3} \right) \mu_0 \lambda^2(T) \xi(T) \right]
\]

where \( \Phi_0 \) is the flux quantum, \( \mu_0 \) the permeability of vacuum, \( \lambda \) the penetration depth, and \( \xi \) the coherence length. The self-field critical current density, \( J_c(0) \), in the best connected
samples indicates the ultimate current-carrying potential in the superconductor, which has been reported as $3.5 \times 10^7$ A/cm$^2$ at 4.2 K and $1.6 \times 10^8$ A/cm$^2$ at 2 K in clean films made by hybrid physical-chemical vapour deposition (HPCVD). These values are about 4% and 20% of the $J_d$ values. Compared with these values, the $J_c(0)$ values in polycrystalline MgB$_2$ bulks and wires are very low and have great potential to be improved.

It was pointed out soon after the discovery of MgB$_2$ that clean grain boundaries are, in principle, no obstacles for supercurrents (Finnemore et al., 2001; Kawano et al., 2001). Such obstacles are known as weak links in the high temperature superconductors. Nevertheless, the connections between the grains remain delicate, since dirty grain boundaries potentially reduce the critical current. Insulating phases have been found at the grain boundaries, consisting of MgO, boron oxides, or boron carbide. Cracks, porosity, or normal conducting phases can further reduce the cross-section over which supercurrents effectively flow. The density of in-situ prepared MgB$_2$ is typically only about half (or less) of its theoretical value, which leads to high porosity.

The in-situ route seems to be the most promising method to improve the $H_{c2}$ and $J_c$ performance of MgB$_2$. Magnesium or MgH$_2$ reacts with boron after mixing and compacting of these precursor powders. MgB$_2$ samples with small grains of poor crystallinity can be obtained at low processing temperatures, resulting in strong pinning and high $H_{c2}$. The stoichiometry can be modified to yield samples with magnesium deficiency, which induces lattice strain, decreases $T_c$, and increases $H_{c2}$. An excess of magnesium in the starting powders may compensate the loss of magnesium due to evaporation or due to a reaction with other elements (e.g. with oxygen or with the sheath material). The precursor powders are very important for the properties of the final samples (Yamada et al., 2004). They should be clean to ensure good grain connectivity. The grain size is strongly influenced by the grain sizes of the precursor powders, especially of the boron powders. Ball milling or mechanical alloying of the precursor powders reduces the grain size and improves the critical current.

Chemical or compound doping changes the reaction kinetics and therefore influences the grain growth, the formation of secondary phases, the density, and the stoichiometry. Carbon doping can be easily performed by the addition of B$_4$C, carbon, carbon nanotubes, nanodiamonds, NbC, SiC, or organic compounds. SiC is by far the most popular dopant, because carbon can be doped into MgB$_2$ at low temperatures (600 °C), according to the dual reaction model (Dou et al., 2007). Higher processing temperatures are necessary for most of the other carbon sources, leading to more grain growth and worse pinning. However, comparable results have also been obtained with nanoscale carbon powder, stearic acid, and carbon nanotubes. It should be noted that the electromagnetic properties of MgB$_2$ are greatly dependent on the starting materials, shielding metals, processing techniques, and measurements. That is why the irreversibility field ($H_{irr}$), $H_{c2}$, and $J_c$ values are different from one batch to another, even for pristine samples, as shown in the figures in this text. All the $J_c$ values are based on the transport measurements reported in this chapter.

2. Nanosized carbon doping effects

The carbon atom has one more electron than boron, and the two-gap feature of MgB$_2$ can be modified if the extra electron is interposed properly into the system. Fortunately, the carbon atoms show strong substitution effects on the boron sites, both theoretically and experimentally, ranging from 1.225% to 30%. As a result, the enhancement of $H_{irr}$, $H_{c2}$, and $J_c$ can be achieved by a controlled carbon doping. The $T_c$ decreases monotonically with
increasing carbon content in the full investigated range of substitution. By adjusting the nominal composition, $T_c$ of substituted crystals can be tuned over a wide temperature range between 10 and 39 K. However, carbon solubility and the effects of carbon doping on $T_c$ vary significantly due to differences in the precursor materials, fabrication techniques, and processing conditions used, because polycrystalline carbon substituted samples may contain significant amounts of impurity phases and the nominal content is assumed most often to be equal to the actual one. Avdeev et al. first suggested the relationship between carbon concentration and lattice parameters. The level of C substitution, $x$, in the formula $\text{Mg}(B_{1-x}C_x)_2$ can be estimated as $x = 7.5\Delta(c/a)$, where $\Delta(c/a)$ is the change in $c/a$ compared to a pure sample (Avdeev et al., 2003).

Fig. 2. The effects of sintering temperature on $J_c(H)$ performance of $\text{MgB}_{1.9}C_{0.1}$ (Yeoh et al., 2006b)

The $J_c$ performance is greatly dependent on the sintering temperature, as shown in Fig. 2. High sintering temperature is essential for a strong flux pinning force because of the intensive carbon substitution effects. Under the optimum conditions, transport $J_c$ has been enhanced by a factor of 5.7 at 12 T and 4.2 K as compared to the pure MgB$_2$ wire. The increased $H_{c2}$ shown in Fig. 3 is in agreement with the high carbon substitution effects. $H_{c2}(0)$ of pure MgB$_2$ increased from 16.0 to 32.5 T in a carbon doped MgB$_2$ filament with slight depression of $T_c$ from 39.2 to 36.2 K for 3.8% C substitution, using the chemical vapor deposition (CVD) method to co-deposit B together with carbon (Wilke et al., 2004). The carbon substitution effects on $H_{c2}$ have shown an encouraging enhancement, with a range of enhanced $H_{c2}$ values from 25 to 40 T at temperatures of 4.2 K and below (Masui et al., 2004; Ohmichi et al., 2004; Putti et al., 2004). Furthermore, $H_{c2}$ with a value of 52–55 T has been commonly observed for carbon alloyed thin films at temperatures around 1.5–4.2 K (Ferdeghini et al., 2005; Ferrando et al., 2005). The enhancement of $H_{c2}$ is in agreement with predictions of the model of two-band impurity scattering of charge carriers in MgB$_2$, which indicates increased intraband scattering via shortening of the electron mean free path, $l$ (Gurevich, 2003). The coherence length, $\xi$, will be shortened according to the equation $1/\xi = 1/l + 1/\xi_0$, where $\xi_0$ is the coherence length at 0 K.
3. Carbon nanotube (CNT) doping effects

Compared with other nano-carbon precursors, carbon nanotubes (CNTs) are particularly interesting because their special geometry (high aspect ratio and nanometer diameter) may induce more effective pinning centers. CNTs can form column-like strong pinning centers to enhance $J_c$ in the Bi-based superconductors (Fossheim et al., 1995; Huang et al., 1999). The flux pinning force depends greatly on the geometry of the different CNTs. Furthermore, the CNT doping significantly improves heat transfer and dissipation during materials processing (Dou et al., 2006), due to the high thermal conductivity and stable electric conductivity of CNTs (Kim et al., 2001; Wei et al., 2001). With CNT properties of high axial strength and stiffness, approaching values for an ideal carbon fiber (Treacy et al., 1996), CNT doping can improve the current path and connectivity between the grains in MgB$_2$.

Transmission electron microscope (TEM) images have shown that CNTs are easy to align in the wire processing direction, as shown in the TEM images in Fig. 4.

The doping effects of single-walled carbon nanotubes (SWCNTs) include amazing pinning effects in MgB$_2$ at 4.2 K, as shown in Fig. 4. Similarly to the case of ordinary carbon-doped MgB$_2$, the best performance in $J_c(H)$ was shown by SWCNT doping with sintering at 900 °C, where the high processing temperature encourages better carbon substitution compared to lower processing temperatures. The $J_c(4.2 \text{ K})$ reached the values of $\sim$51,000 and $\sim$3500 A/cm$^2$ at 7 and 12 T, respectively, as shown in Fig. 5.

Multi-walled carbon nanotubes (MWCNT) have also shown positive effects on the $J_c$ of MgB$_2$, however, the results are not as significant as with the SWCNTs. Furthermore, the $J_c$ is dependent on the length of the MWCNTs: short MWCNTs give rise to a stronger flux pinning force than long ones. Yeoh et al. have shown that there is a correlation between the reactivity of the CNTs and the amount of carbon substitution in the MgB$_2$, with the substitution of carbon for boron only occurring after the carbon atoms break free from the CNT (Yeoh et al., 2007a). Longer CNTs tend to entangle and agglomerate, which results in
Fig. 4. TEM images of CNT doped MgB$_2$ show straightened CNTs in the same processing direction in the MgB$_2$ matrix. The inset is a high resolution image of a CNT (Dou et al., 2006)

Fig. 5. Transport critical current at 4.2 K at fields up to 12 T for different CNT doped wires produced at sintering temperatures of 800 and 900 °C (Kim et al., 2006a)

inhomogeneous mixing of the CNTs with the precursor powder, blocking the current transport and suppressing the $J_c$ (Yeoh et al., 2005). Ultrasonication of CNTs has been introduced to improve the homogenous mixing of the CNTs with the MgB$_2$ matrix, resulting in a significant enhancement in the field dependence of the critical current density (Yeoh et
al., 2006a). The $J_c$ performance of different types of CNT doped MgB$_2$ is in agreement with the $H_{c2}$ shown in Fig. 6.

![Fig. 6. The $H_{c2}$ of different CNT doped MgB$_2$ samples sintered at 900 °C. The temperature has been normalized by $T_c$ (Kim et al., 2006a)](image)

4. Nanosized SiC doping effects

Nanosized doping centers are highly effective, as they are comparable with the coherence length of MgB$_2$ (Soltanian et al., 2003). MgB$_2$ has a relatively large coherence length, with $\xi_{ab}(0) = 3.7$–12 nm and $\xi_c(0) = 1.6$–3.6 nm (Buzea & Yamashita, 2001), so a strong pinning force can be introduced by nanoparticles that are comparable in size. Nanoscale SiC has been found to be the right sort of candidate, providing both second phase nanoscale flux pinning centers and an intensive carbon substitution source (Dou et al., 2002a; Dou et al., 2002b; Dou et al., 2003b). 10 wt% nano-SiC doped MgB$_2$ bulk samples showed $H_{irr} \approx 8$ T and $J_c \approx 10^8$ A cm$^{-2}$ under 3 T at 20 K. The $T_c$ reduction is not pronounced, even in heavily doped samples with SiC up to 30% (Dou et al., 2002b).

Fig. 7 compares the $J_c$ values of pure MgB$_2$ and those of MgB$_2$ doped with 10 wt% nanosized SiC at different temperatures. There are crossover fields for the $J_c$ at the same temperature for different samples, due to the different reductions in slope of the flux pinning force when the temperature is lower than 20 K. The carbon substitution effects in the SiC doped sample are very strong, and therefore, the $J_c$ decreases steadily with increasing field. The $J_c$ drops quickly when the temperature approaches $T_c$. An increase in $H_{c2}$ from 20.5 T to more than 33 T and enhancement of $H_{irr}$ from 16 T to a maximum of 28 T for an SiC doped sample were observed at 4.2 K (Bhatia et al., 2005). Matsumoto et al. showed that very high values of $H_{c2}(0)$, exceeding 40 T, can be attained in SiC-doped bulk MgB$_2$ sintered at 600 °C (Matsumoto et al., 2006). This result is considerably higher than for C-doped single crystal (Kazakov et al., 2005), filament (Wilke et al., 2004; Li et al., 2009a), or bulk samples (Senkowicz et al., 2005). Low temperature sintering is beneficial to both the $H_{irr}$ and the $H_{c2}$. 
as shown in Fig. 8, which suggests that significant lattice distortion is introduced by alloying and by reaction at low temperature. This has important consequences for the application of MgB$_2$ wires and tapes in the cable and magnet industries.

Fig. 7. Comparison of $J_c$ of pure MgB$_2$ with that of a nanosized SiC doped sample at different temperatures (Dou et al., 2002b; Shcherbakova et al., 2006)

Fig. 8. The effects of sintering temperature on $H_{c2}$ and $H_{irr}$ of 10 wt%, ~15 nm SiC doped MgB$_2$ (Soltanian et al., 2005). The insets show the resistance as a function of temperature at different magnetic fields for samples sintered at 640 °C (upper right) and 1000 °C (lower left)
Fig. 9 shows the critical current density of MgB$_2$ in comparison with other commercial superconductor materials. It should be noted that the $J_c$ of SiC-doped MgB$_2$ stands out very strongly, even at 20 K in low field, and that it is comparable to the value of $J_c$ for Nb–Ti at 4.2 K, which is very useful for application in magnetic resonance imaging (MRI). At 20 K, the best $J_c$ for the 10 wt% SiC doped sample was almost $10^5$ A/cm$^2$ at 3 T, which is comparable with the $J_c$ of state-of-the-art Ag/Bi-2223 tapes. These results indicate that powder-in-tube-processed MgB$_2$ wire is promising, not only for high-field applications at 4.2 K, but also for applications at 20 K with a convenient cryocooler. Fig. 10 shows TEM and high resolution TEM (HRTEM) images of 10 wt% nanosized SiC doped MgB$_2$. A high density of dislocations and different sizes of nano-inclusions can be observed in the MgB$_2$ matrix. Furthermore, the HRTEM images indicate that the MgB$_2$ crystals display nanodomain structures, which is attributed to lattice collapse caused by the carbon substitution.

However, similar to the doping effects of carbon and CNTs, the connectivity of nanosized SiC doped MgB$_2$ is quite low. To improve the connectivity, additional Mg was added into the precursor mixture (Li et al., 2009a; Li et al., 2009b). To explore the effects on connectivity of Mg excess, microstructures of all the samples were observed by scanning electron microscope (SEM), as shown in Fig. 11. The grains in the stoichiometric MgB$_2$ samples show an independent growth process, which is responsible for their isolated distribution. The grains in Mg$_{1.15}$B$_2$ have clearly melted into big clusters because the additional Mg can extend the liquid reaction time. The grain shapes in MgB$_2$ + 10 wt % SiC are different from those in pure, stoichiometric MgB$_2$ because the former crystals are grown under strain due to the C substitution effects. The strain is also strong in Mg$_{1.15}$B$_2$ + 10 wt % SiC, as long bar-shaped grains can be observed under SEM. The strain is released in the high Mg content samples ($x$ > 1.20), judging from the homogeneous grain sizes and shapes. Compared with MgB$_2$ + 10 wt % SiC, the grain connectivity improved greatly with the increasing Mg addition. The
grains were merged into big particles, and grain boundaries have replaced the gaps between grains. However, more impurities are induced in forms such as residual Mg and MgO.

Fig. 10. TEM images of SiC-doped MgB₂ showing the high density of dislocations (a), inclusions larger than 10 nm (b), inclusions smaller than 10 nm (c), and HRTEM image of the nanodomain structure (d) (Dou et al., 2003a; Li et al., 2003)

The concept of the connectivity, \( A_F \), was introduced to quantify this reduction of the effective cross-section, \( \sigma_{\text{eff}} \), for supercurrents (Rowell, 2003; Rowell et al., 2003): \( A_F = \sigma_{\text{eff}} / \sigma_0 \), where \( \sigma_0 \) is the geometrical cross-section. The connectivity can be estimated from the phonon contribution to the normal state resistivity by

\[
A_F = \frac{\Delta \rho_{\text{ideal}}}{\Delta \rho(300 \text{ K})}
\]

where \( \Delta \rho_{\text{ideal}} = \rho_{\text{ideal}}(300 \text{ K}) - \rho_{\text{ideal}}(T_c) \approx 9 \mu\Omega \cdot \text{cm} \) is the resistivity of fully connected MgB₂ without any disorder, and \( \Delta \rho(300 \text{ K}) = \rho(300 \text{ K}) - \rho(T_c) \). This estimate is based on the assumption that the effective cross-section is reduced equivalently in the normal and superconducting states, which is a severe simplification. The supercurrents are limited by the smallest effective cross-section along the conductor, and the resistivity is given more or less by the average effective cross-section. A single large transverse crack strongly reduces
Fig. 11. SEM images of MgB$_2$ (a), Mg$_{1.15}$B$_2$ (b), MgB$_2$+10 wt % SiC (c), Mg$_{1.15}$B$_2$+10 wt % SiC (d), Mg$_{1.20}$B$_2$+10 wt % SiC (e), Mg$_{1.25}$B$_2$+10 wt % SiC (f), and Mg$_{1.30}$B$_2$+10 wt % SiC (g) (Li et al., 2009a)
Fig. 12. (Color online) Ambient Raman spectra of MgB$_2$, Mg$_{1.15}$B$_2$, and Mg$_x$B$_2$+10 wt % SiC ($x = 1.00, 1.15, 1.20, 1.25,$ and 1.30) fitted with three peaks: $\omega_1$, $\omega_2$, and $\omega_3$. The dashed line indicates the vibration of the $E_{2g}$ mode ($\omega_2$) in different samples (Li et al., 2009a).

$J_c$, but only slightly increases the resistivity of a long sample. Un-reacted magnesium decreases $\Delta\rho(300$ K) (Kim et al., 2002) and the cross-section for supercurrents. Thin insulating layers on the grain boundaries strongly increase $\Delta\rho(300$ K), but might be transparent to supercurrents. Finally, $\Delta\rho_{\text{ideal}}$ within the grains can change due to disorder. Even a negative $\Delta\rho(300$ K) has been reported in highly resistive samples (Sharma et al.,
2002). Despite these objections, $A_F$ is very useful, at least if the resistivity is not too high. A clear correlation between the resistivity and the critical current has been found in thin films (Rowell et al., 2003). Nevertheless, one should be aware of the fact that this procedure is not really reliable, but just a possibility for obtaining an idea about the connectivity. It should be noted that the connectivity is far removed from that found in ideal crystals, as reflected by the low $A_F$ values. Although the $A_F$ values of pure and 10% SiC doped MgB$_2$ are just 0.106 and 0.062, additional Mg can improve them to 0.162 and 0.096 for 15 wt % Mg excess samples, respectively. High $A_F$ values are the reflection of a broad channel of supercurrents, while impurities reduce the connectivity in large $x$ samples. High connectivity improves the supercurrent channels because the currents can easily meander through the well-connected grains. The results show that excess Mg in Mg$_{1.15}$B$_2$ + 10 wt% SiC composite effectively improves the connectivity, as evidenced by its higher $A_F$. Its promising $J_c(H)$ is attributed to both the high connectivity and the improved $H_{irr}$ and $H_{c2}$.

Raman scattering is employed to study the combined influence of connectivity and lattice distortion. Chemical substitution and lattice distortion are expected to modify the phonon spectrum, by changing the phonon frequency and the electron-phonon interaction. The effects of C substitution include an increase in impurity scattering and band filling, which reduces the density of states (DOS) and alters the shape of the Fermi surface. The $E_{2g}$ phonon peak shifts to the higher energy side, and the peak is narrowed with increasing $x$ in Mg(B$_{1-x}$C$_x$)$_2$ (Li et al., 2008). As a carbon source, nano-SiC shows a similar influence, due to its C atoms, on the $J_c$, $H_{irr}$, $H_{c2}$, and even Raman spectra in MgB$_2$. Figure 12 shows the Raman spectra fitted with three peaks: $\omega_1$, $\omega_2$, and $\omega_3$. The $\omega_1$ and $\omega_3$ peaks are understood to arise from sampling of the phonon density of states (PDOS) due to disorder, while $\omega_2$ is associated with the $E_{2g}$ mode, which is the only Raman active mode for MgB$_2$ (Kunc et al., 2001). A reasonable explanation for the appearance of $\omega_1$ and $\omega_3$ is the violation of Raman selection rules induced by disorder. All three peaks are broad, as in previous results, due to the strong electron-phonon coupling. The influence of $\omega_1$ on the superconducting performance is negligible compared with those of $\omega_2$ and $\omega_3$ because of its weak contribution to the Raman spectrum. The frequency and full width at half maximum (FWHM) of $\omega_2$ and $\omega_3$ are shown in Fig. 13. Both $\omega_2$ and $\omega_3$ are hardened with SiC addition. The $\omega_2$ frequency is reduced with further Mg addition, whereas the $\omega_3$ frequency remains almost stable. The frequencies of $\omega_2$ for the $x \geq 1.20$ samples are even lower than for the pure, stoichiometric MgB$_2$. The FWHM of $\omega_2$ decreases with SiC doping, while the Mg excess weakens this trend. On the contrary, the $\omega_3$ FWHM increases with SiC addition and becomes narrow with more addition of Mg. The Raman scattering properties are the direct reflection of the phonon behavior of MgB$_2$. The parameters of Raman spectra vary with the composition of MgB$_2$ crystals and the influence of their surroundings, which depends on both the connectivity and the disorder of the samples. Furthermore, the disorder should be considered as composed of intrinsic and extrinsic parts based on their different sources. The crystallinity and chemical substitution are believed to be responsible for the intrinsic disorder effects, while the grain boundaries and impurities are treated as responsible for the extrinsic disorder effects. The influences of intrinsic disorder on the basic characteristics of Raman spectra are significant because the physical properties of MgB$_2$ depend on the intrinsic disorder. The Raman parameters can also be tuned by the extrinsic disorder. Especially in samples with good connectivity, the influences of grain boundaries and impurities on the Raman spectra need to be taken into account because of their strain effects on the MgB$_2$ crystals (Zeng et al., 2009). The
differences between shifts and FWHMs in the Raman spectra for MgB$_2$, Mg$_{1.15}$B$_2$, MgB$_2$ + 10 wt % SiC, and Mg$_{1.15}$B$_2$ + 10 wt % SiC are mostly attributable to their intrinsic characteristics because of their different chemical compositions. The Raman spectra of Mg$_x$B$_2$ + 10 wt % SiC ($x > 1.20$) can be considered as gradual modifications of that of Mg$_{1.15}$B$_2$ + 10 wt % SiC. The weakened C substitution effects are responsible for the decreased frequencies and slightly increased FWHMs of $\omega_2$ with Mg addition. Accordingly, the FWHMs of $\omega_3$ decrease with increased Mg due to the weakened lattice distortion. Although the $A_F$ values are quite low for Mg$_x$B$_2$ + 10 wt % SiC ($x > 1.20$), the effects of extrinsic disorder on Raman parameters are considerable, through the MgB$_2$-MgB$_2$ and MgB$_2$-impurity interfaces, and the connectivity deteriorates with the increased $x$ values due to the decreased number of MgB$_2$-MgB$_2$ interfaces. A high FWHM value for $\omega_2$ is correlated with high self-field $J_c$ due to high carrier density, while a high FWHM value for $\omega_3$ is correlated with strong high-field $J_c$ because of the strong flux pinning force due to the large disorder. The FWHM behaviors show that high connectivity and strong disorder are best combined in Mg$_{1.15}$B$_2$ + 10 wt % SiC among all the samples.

Fig. 13. Fitted parameters of Raman shifts for $\omega_2$ (a) and $\omega_3$ (b), and FWHMs for $\omega_2$ (c) and $\omega_3$ (d). The sample labels are defined as A for Mg$_{1.15}$B$_2$, B for MgB$_2$, C for MgB$_2$+10 wt % SiC, D for Mg$_{1.15}$B$_2$+10 wt % SiC, E for Mg$_{1.20}$B$_2$+10 wt % SiC, F for Mg$_{1.25}$B$_2$+10 wt % SiC, and G for Mg$_{1.30}$B$_2$+10 wt % SiC (Li et al., 2009a)
5. Organic dopants

Most dopants have been introduced into MgB$_2$ superconductors via solid state reaction using a dry mixing process, which is responsible for the common inhomogeneous distribution of dopants. Therefore, the soluble nature and low melting point of hydrocarbons and carbohydrates give these dopants advantages over the other carbon based dopants. The homogeneous distribution of hydrocarbons and carbohydrates results in high $J_c$ values comparable with those from the best SiC nanoparticles (Kim et al., 2006b; Yamada et al., 2006; Li et al., 2007; Zhou et al., 2007).

Fig. 14 shows the $J_c$ performance of MgB$_2$ doped with malic acid and sintered at different temperatures. Low temperature sintering has significant benefits for the $J_c$. Moreover, the malic acid (C$_4$H$_6$O$_5$) doping technique provides additional benefits to the $J_c(H)$ performance in low fields, that is, $J_c$ at low fields is not degraded at certain doping levels as it is for any other C doping method. A cold, high pressure densification technology was employed for improving $J_c$ and $H_{irr}$ of monofilamentary in-situ MgB$_2$ wires and tapes alloyed with 10 wt% C$_4$H$_6$O$_5$. Tapes densified at 1.48 GPa exhibited an enhancement of $J_c$ after reaction from 2 to $4 \times 10^4$ A cm$^{-2}$ at 4.2 K/10 T and from 0.5 to $4 \times 10^4$ A cm$^{-2}$ at 20 K/5 T, while the $H_{irr}$ was enhanced from 19.3 to 22 T at 4.2 K and from 7.5 to 10.0 T at 20 K (Flukiger et al., 2009; Hossain et al., 2009). Cold densification also caused a strong enhancement of $H(10^4)$, the field at which $J_c$ takes the value $1 \times 10^4$ A cm$^{-2}$. For tapes subjected to 1.48 GPa pressure, $H(10^4)\parallel$ and $H(10^4)\perp$ at 4.2 K were found to increase from 11.8 and 10.5 T to 13.2 and 12.2 T, respectively. Almost isotropic conditions were obtained for rectangular wires with aspect ratio $a/b < 2$ subjected to 2.0 GPa, where $H(10^4)\parallel = 12.7$ T and $H(10^4)\perp = 12.5$ T were obtained. At 20 K, the wires exhibited an almost isotropic behavior, with $H(10^4)\parallel = 5.9$ T and $H(10^4)\perp = 5.75$ T, with $H_{irr}(20 \text{ K})$ being ~10 T. These values are equal to or higher than the highest values reported so far for isotropic in-situ wires with SiC or other carbon based additives. Further improvements are expected in optimizing the cold, high pressure densification process, which has the potential for fabrication of MgB$_2$ wires of industrial lengths.

![Fig. 14. Sintering temperature effects on the $J_c$ performance of MgB$_2$ doped with malic acid (Kim et al., 2008)](image-url)
Fig. 15. Field emission SEM images: (a) pure MgB$_2$, (b) MgB$_2$ + 10 wt% malic acid, and (c) MgB$_2$ + 30 wt% malic acid (Kim et al., 2006b)

Fig. 16. $H_{irr}$ and $H_{c2}$ variations with doping content of malic acid in MgB$_2$ (Kim et al., 2006b)

Highly reactive and fresh carbon on the atomic scale can be introduced into the MgB$_2$ matrix because the organic reagents decompose at temperatures below the formation temperature of MgB$_2$. The carbon substitution is intensive at temperatures as low as the formation temperature of MgB$_2$. Microstructural analysis suggests that $J_c$ enhancement is due to the substitution of carbon for boron in MgB$_2$, liquid homogenous mixing, and highly homogeneous and highly connected MgB$_2$ grains, as shown in Fig. 15. MgB$_2$ with
hydrocarbon-based carbonaceous compounds has also demonstrated great application potential due to the improvements in both $J_c$ and $H_{c2}$ as shown in Fig. 16, while the $T_c$ just decreases slightly. It should be noted that 30 wt% doping with malic acid is still effective for the improvement of $H_{c2}$, which benefits from the high density of flux pinning centers in the MgB$_2$ matrix.

6. Doping effects of other carbon sources

Diamond, Na$_2$CO$_3$, carbon nanohorns, graphite, and carbide compounds have also been employed as dopants to achieve flux pinning in MgB$_2$ (Zhao et al., 2003; Ueda et al., 2004; Xu et al., 2004; Ban et al., 2005; Yamamoto et al., 2006). All show positive effects on $J_c$ performance. B$_4$C appears to be an ideal carbon source to avoid excessive carbonaceous chemical addition. Ueda et al. and Yamamoto et al. showed that C could substitute into the B sites when a mixture of Mg, B, and B$_4$C was sintered at 850 °C for bulk samples (Ueda et al., 2005; Yamamoto et al., 2005a; Yamamoto et al., 2005c). Substantially enhanced $J_c$ properties under high magnetic fields were observed in the B$_4$C doped samples due to the relatively low processing temperature and carbon substitution effects. Lezza et al. successfully obtained a $J_c$ value of $1 \times 10^4$ A cm$^{-2}$ at 4.2 K and 9 T for 10 wt% B$_4$C powders added to MgB$_2$/Fe wires at a reaction temperature of 800 °C (Lezza et al., 2006). Despite the carbon substitution effects, the homogeneous microstructure of the dopants provides the MgB$_2$ composites with good grain connection for the MgB$_2$ phase and a high density of flux pinning centers.

7. Mechanism of doping effects — dual reaction model

Carbon substitution in the boron sites is the dominant factor for the enhancement of $J_c(H)$ and $H_{c2}$ in all carbonaceous chemical doped MgB$_2$ because of the strong disorder effects. Furthermore, the defects, grain sizes, second phases, grain boundaries, and connectivity are also important for the superconducting properties. The study of reaction kinetics for different carbonaceous chemicals during the MgB$_2$ synthesis is a crucial issue for understanding the $H_{ Ir}$, $H_{c2}$, and $J_c$ performance in MgB$_2$. A systematic correlation between the processing temperature, $J_c$, and $H_{c2}$ has been observed in pure, nano-carbon, CNT, SiC, and hydrocarbon doped MgB$_2$ samples (Dou et al., 2007; Yeoh et al., 2007b). The processing temperature is believed to be the most important factor influencing the electromagnetic properties because both the carbon substitution intensity and the microstructure are dependent on it.

Fig. 17 shows the effects of sintering temperature on the $J_c(H)$ for different carbon based dopants. The hydrocarbon and SiC doped MgB$_2$ show significant enhancement in $J_c$ for the samples sintered at lower temperature, whereas the carbon and CNT doped MgB$_2$ need to be sintered at higher temperature for high $J_c$. The low sintering temperature results in small grain size, high concentrations of impurities and defects, and large lattice distortion, which are all responsible for a strong flux pinning force (Soltanian et al., 2005; Yamamoto et al., 2005b). Furthermore, the hydrocarbon and SiC can release fresh and active free carbon at very low temperature, which means that the carbon substitution effects take place simultaneously with the MgB$_2$ formation. A high sintering temperature will perfect the crystallization and decrease the flux pinning centers in the MgB$_2$ matrix. That is the reason why high sintering temperature degrades the $J_c$ performance. Although high sintering
temperature has the same shortcomings in nanosized carbon and CNT doped MgB$_2$, the carbon substitution effects improve their $J_c$ values. The high sintering temperature is necessary for carbon and CNT doped MgB$_2$ because the carbon and CNT are quite stable at low temperature and the substitution effects are absent if the sintering temperature is not high enough.

Fig. 17. The critical current density ($J_c$) at 4.2 K versus magnetic field for wires of pure MgB$_2$ and MgB$_2$ doped with C, SiC, SWCNTs, and malic acid that were sintered at different temperatures (Dou et al., 2002b; Yeoh et al., 2006b; Dou et al., 2007; Kim et al., 2008)

A dual reaction model has been suggested to explain the improvement of the superconducting properties in SiC doped MgB$_2$, based on the $J_c$ dependence on the sintering temperature (Dou et al., 2007). The reaction of SiC with Mg at low temperature will release fresh and active carbon, which is easily incorporated into the lattice of MgB$_2$ at the same temperature. The reaction product Mg$_2$Si and excess carbon are also high quality nanosized flux pinning centers. The low temperature substitution is accompanied by small grain size, high density of grain boundaries, and high density of all kinds of defects, which are all favorable to the high superconducting performance. Another example for the dual reaction model is the high $J_c$ malic acid doped MgB$_2$ shown in Fig. 14. The carbonaceous chemical doping effects on the superconducting performance can be predicted according to the dual reaction model as arising from the combination of defects and carbon substitution effects. Most dopants, such as TiC and NbC, show very small effects towards the enhancement of $J_c$ compared with carbon, SiC, CNTs, and hydrocarbons because the substitution effects are very weak and there are no efficient flux pinning centers either.

8. Conclusions

The experimental results on $H_{c2}$ and $J_c$ strongly suggest that MgB$_2$ doped with carbonaceous sources shows remarkable enhancement of superconducting performance if the carbon substitution effects are intensive. In particular, nanosized SiC and malic acid are the most promising dopants to advance the high field $J_c$ performance for practical application. The
enhancement of $J_c$, $H_{irr}$, and $H_{c2}$ for MgB$_2$ with carbon substituted into boron sites is due to its intrinsic properties arising from the strong two-band impurity scattering effects of charge carriers. The carbonaceous chemical doping effects have been attributed to a dual reaction model, based on the sintering effects on superconducting properties for different kinds of carbonaceous chemicals. The fresh, active, and free carbon atoms are very easy to substitute onto B sites in the MgB$_2$ lattice if the carbonaceous decomposition temperatures are close to the formation temperature of MgB$_2$, ~650 °C. The dual reaction model can explain and predict the doping effects of carbonaceous chemicals on the superconducting properties very well. The high density of defects is another factor that improves the $J_c$, $H_{irr}$, and $H_{c2}$. However, the connectivity of the samples is also responsible for the low field $J_c$ performance, which is free from the flux pinning force and can be attributed to the density of supercurrent carriers. Both microstructure observations and Raman scattering measurements have confirmed the great influence of connectivity on $J_c$ behavior, as shown by the effects of extra Mg addition in nanosized SiC doped MgB$_2$. The proper Mg content will improve the connectivity greatly to improve the density of supercurrent carriers.

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