Superconducting Properties of \((\text{MnFe}_2\text{O}_4)_x/\text{CuTl-1223}\) Composites

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Abstract

The effects of nanometer size \(\text{MnFe}_2\text{O}_4\) particles addition on different properties of \((\text{Cu}_{0.5}\text{Tl}_{0.5})\text{Ba}_2\text{Ca}_n\text{Cu}_{2n+4-\delta}\) (CuTl-1223) superconducting phase was studied. \(\text{MnFe}_2\text{O}_4\) nanoparticles were synthesized by sol-gel method and were added to CuTl-1223 superconducting phase synthesized by solid-state reaction to get \((\text{MnFe}_2\text{O}_4)_x/\text{CuTl-1223}\); \(x = 0 \sim 2.0\) wt. % composites. The effects of these magnetic nanoparticles on structural, morphological and superconducting transport properties of host CuTl-1223 phase were investigated via X-ray diffraction (XRD), scanning electron microscopy (SEM), energy dispersive X-rays (EDX) spectroscopy, Fourier transform infrared (FTIR) spectroscopy and Four-Probe Resistivity (\(\rho\)) measurements. Inclusion of these nanoparticles exhibited no change in crystal structure of host CuTl-1223 superconducting phase. Suppression of superconducting properties was probably due to enhanced effective scattering of carriers across magnetic natured \(\text{MnFe}_2\text{O}_4\) nanoparticles residing at the grain-boundaries of CuTl-1223 superconductor.

Keywords: \((\text{MnFe}_2\text{O}_4)_x/\text{CuTl-1223}\) composites; Crystal structure; Morphology; Superconducting properties.

PACS codes: 74.25.-q, 74.25. F-, 74.72.-h, 74.81.Bd

Introduction

Selection of any superconducting material for practical applications highly depends upon different critical parameters like critical temperature (\(T_c\)), critical current density (\(J_c\)) and critical magnetic field (\(H_c\)). Different phases of \((\text{Cu}_{0.5}\text{Tl}_{0.5})\text{Ba}_2\text{Ca}_n\text{Cu}_{2n+4-\delta}\) high temperature superconductor (HTSCs) family can be synthesized at higher pressure of (~5 GPa) as well as at ambient pressure\(^{[1]}\). HTSCs samples synthesized at high pressure exhibited relatively higher superconducting properties due to reduction in bulk porosity and higher value of superconducting volume fraction. But large scale production of HTSCs at higher pressure for industrial based applications is a biggest challenge in present time. Therefore, synthesis of HTSCs at ambient pressure is an appropriate way for large production. But presence of voids and pores in higher density is the main disadvantage of this synthesis route, which reduces overall superconducting volume fraction. Another reason for reduction in carrier’s density from an optimal level is mostly caused by the existence of various defects in the form of oxygen deficiencies in ambient pressure synthesized HTSCs samples. So, other than higher pressure synthesis route, numbers of efforts have been made to overcome this problem in which main goal was to improve the superconducting properties of different compounds of HTSCs families by different techniques\(^{[2-4]}\). This issue is most effectively resolved by increasing the connectivity between the grains and reduction in density of voids and pores by inclusion of various nanoparticles, which is the easiest and efficient ways in this regards. But control on the size, concentration and distribution of these nanoparticles at grain-boundaries of host bulk HTSCs are the real challenges\(^{[5-8]}\). This uniform distribution of nanoparticles can equally facilitate the transportation of carriers across the grains, reducing the energy required for transport and increasing the supercurrent density.
charge carriers across the grain-boundaries but up to a certain optimal level of nanoparticles inclusion after which prominent reduction in superconducting volume fraction takes place. It was found that nature of nanostructures materials added in the host material plays a very crucial role in deciding the resultant effects on superconducting properties. A prominent improvement in superconducting properties of Bi-based superconductor was observed after inclusion of MgO, ZrO and Al₂O₃ nanoparticles. Increase in superconducting volume fraction and sharpness in transition width of Bi-2212 superconducting phase was observed after inclusion of MgO nanoparticles. The critical current density, volume pinning force density, activation energy and onset temperature in presence of applied magnetic field were improved after alumina nanoparticles inclusion in (Bi, Pb)-2223 superconductor. The inclusion of various nanoparticles of different materials with variant sizes like Fe₃O₄ with required amount was used to generate the artificial flux pinning centers, which can increase the various in-field superconducting parameters. The crystal structure and Jc value of Y123 superconducting thin film improved after the addition of Ag nanoparticles. Addition of Ag nanoparticles improved the superconducting and mechanical properties of (Bi-Pb)-Sr-Ca-Cu-O superconducting phase, which was attributed to the cementing effects of these metallic nanoparticles at grain-boundaries. An improvement was observed in critical parameters and superconducting volume fraction of CuTi-1223 matrix after addition of Ag nanoparticles up to x = 1.5 wt.%. Addition of ZnFe₂O₄ nanoparticles in Gd-Ba₂Cu₃O₇, enhanced superconducting parameters up to a certain optimal concentration level. CoFe₂O₄ nanoparticles addition in YBa₂Cu₃O₇ (Y-123) and Y Ba₂Cu₃O₁₈ (Y-358) suppressed the superconducting parameters as well as grain size. The critical current density Jc of Bi-2223 superconductor was increased after inclusion NiFe₂O₄ nanoparticles.

In this article, we have analyzed and compared the effects of ferri-magnetic MnFe₂O₄ nanoparticles inclusion on structural, morphological and electrical transport properties of CuTi-1223 phase in detail. In literature there is no evidence of investigation on the effects of these nanoparticles inclusion in HTSCs materials.

### Experimental details

Solid-state reaction method was used to prepare bulk CuₓBa₃Ca₂Cu₃O₁₀₋δ precursor. Initially Cu₂(CN)₂.H₂O (99%, BDH), Ba(NO₃)₂ (99.50%, UNI-Chem), and Ca(NO₃)₂.4H₂O (99%, Sigma-Aldrich) were mixed in suitable ratios of 5.149, 7.802 and 7.049 grams to prepared 20 grams initial mixtures. These mixed materials were ground for 1 hour and then pellets were made by using of hydraulic press. Each pellet was enclosed in a gold capsule and sintered in pre-heated chamber furnace at 860°C for 10 min and quenched to room temperature to get required (MnFe₂O₄)/CuTi-1223 nanoparticles-superconductor composites.

### Results and Discussion

Crystal structures of nanoparticles and superconducting composites were determined by XRD technique. XRD pattern of MnFe₂O₄ nanoparticles is shown in Figure.1. Debye-Scherrer’s formula was used to determine the average particle size and was found 20 nm. All the major well indexed diffraction peaks demonstrate the spinal structure of MnFe₂O₄ nanoparticles. MnFe₂O₄ showed cubic structure following space group Fd-3m with calculated lattice parameter a = 4.89 Å. XRD spectra of (MnFe₂O₄)/CuTi-1223 composites are shown in Figure.2. Majority of XRD peaks of these composites are indexed according tetragonal structure of CuTi-1223 phase following P4/mmm space group. XRD of all these samples have also shown the presence of very small amount of impurities and other superconducting phases. Percentage volume fraction of different phases was calculated by using the following relations:

\[
\text{CuTi-1223 \%} = \frac{\sum \text{I(1223)}}{\sum (\text{I(1223)} + \text{I(1234)} + \text{I(1212)} + \text{I(Unknown Impurities)})} \times 100
\]

\[
\text{CuTi-1234 \%} = \frac{\sum \text{I(1234)}}{\sum (\text{I(1223)} + \text{I(1234)} + \text{I(1212)} + \text{I(Unknown Impurities)})} \times 100
\]

\[
\text{CuTi-1212 \%} = \frac{\sum \text{I(1212)}}{\sum (\text{I(1223)} + \text{I(1234)} + \text{I(1212)} + \text{I(Unknown Impurities)})} \times 100
\]

Unknown Impurities \% = \frac{\sum \text{I(Unknown Impurities)}}{\sum (\text{I(1223)} + \text{I(1234)} + \text{I(1212)} + \text{I(Unknown Impurities)})} \times 100

Where, I (1223), I (1234), I (1212) and I (Impurities) are intensities of respective XRD peaks for CuTi-1223, CuTi-1234, CuTi-1212 phases and unknown impurities, respectively. The calculated relative proportions of these phases present in (MnFe₂O₄)/CuTi-1223 composites are listed in Table 1. We observed a clear decrease in % of volume fraction of CuTi-1223 phase with the addition of MnFe₂O₄ nanoparticles. The intensities of diffraction peaks related to nanoparticles crystal structures are very small, which are suppressed under high intensities diffraction peaks of host CuTi-1223 superconductor. But all the major diffraction peaks are well indexed according to CuTi-1223 phase after addition of MnFe₂O₄ nanoparticles, which indicates unaltered crystal structure of host CuTi-1223 matrix.
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Figure 1. XRD spectrum of (MnFe$_2$O$_4$) nanoparticles.

Figure 2. XRD spectra of (MnFe$_2$O$_4$)$_x$/CuTl-1223; $x = 0$, 1.0 and 2.0 wt.% composites.

Table 1. % volume fraction of CuTl-1223, CuTl-1234, CuTl-1212 and unknown impurities present in (MnFe$_2$O$_4$)$_x$/CuTl-1223 nanoparticles-superconductor composites $x = 0$, 1.0 and 2.0 wt. %.

<table>
<thead>
<tr>
<th>Nanoparticles contents (x %)</th>
<th>% of CuTl-1223 Phase</th>
<th>% of CuTl-1234 Phase</th>
<th>% of CuTl-1212 Phase</th>
<th>% of Unknown Impurities</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>94.49</td>
<td>4.9</td>
<td>0.55</td>
<td>---</td>
</tr>
<tr>
<td>1.0</td>
<td>91.48</td>
<td>3.54</td>
<td>3.59</td>
<td>1.31</td>
</tr>
<tr>
<td>2.0</td>
<td>90.50</td>
<td>4.54</td>
<td>3.15</td>
<td>1.78</td>
</tr>
</tbody>
</table>

Morphology of representative (MnFe$_2$O$_4$)$_x$/CuTl-1223; $x = 0$ and 1.0 wt.% composites samples was examined by scanning electron microscope (SEM) as shown in Figure 3. We observed relative improvement in inter-grain weak-links and decrease in density of voids and pores after addition MnFe$_2$O$_4$ nanoparticles in CuTl-1223 superconducting matrix. The presence of different elements in these (MnFe$_2$O$_4$)$_x$/CuTl-1223; $x = 0$ and $x = 1.0$ wt.% composites was determined by energy dispersive X-rays (EDX) spectroscopy as shown in Figure 4. The wt.% of different elements present in (MnFe$_2$O$_4$)$_x$/CuTl-1223; $x = 0$ and $x = 1.0$ wt.% composites are given in Table 2.

Figure 3. SEM images of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites with (a) $x = 0$, (b) $x = 1.0$ wt. %.

Figure 4. EDX spectra of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites with (a) $x = 0$, (b) $x = 1.0$ wt. %.
Table 2. % of different elements present in (MnFe$_2$O$_4$)$_x$/CuTl-1223; $x =$ 0, 1.0 wt.% nanoparticles-superconductor composites.

<table>
<thead>
<tr>
<th>Elements</th>
<th>wt.% of elements in samples with $x =$ 0</th>
<th>wt.% of elements in samples with $x =$ 1.0 Wt.%</th>
</tr>
</thead>
<tbody>
<tr>
<td>O</td>
<td>16.5</td>
<td>18.9</td>
</tr>
<tr>
<td>Ca</td>
<td>9.20</td>
<td>8.9</td>
</tr>
<tr>
<td>Cu</td>
<td>27.7</td>
<td>23.8</td>
</tr>
<tr>
<td>Ba</td>
<td>28.6</td>
<td>28.8</td>
</tr>
<tr>
<td>Tl</td>
<td>18.0</td>
<td>18.0</td>
</tr>
<tr>
<td>Fe</td>
<td>---</td>
<td>1.0</td>
</tr>
<tr>
<td>Mn</td>
<td>---</td>
<td>0.5</td>
</tr>
<tr>
<td>Total %</td>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

The trace amount of impurities and different functional groups can be identified with the help of FTIR spectroscopy. Different vibrational oxygen modes in unit cell of cuprates can also be investigated by FTIR absorption spectroscopy. Different oxygen vibrational phonon modes in our samples has been studied by FTIR absorption spectroscopy as we know optimum oxygen content are very useful for superconductivity in HTSCs.

FTIR absorption spectra of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites in the range from 400 to 700 cm$^{-1}$ are shown in Figure 5. The absorption modes appearing in the range from 400 to 540 cm$^{-1}$ are associated with apical oxygen atoms and 541 - 600 cm$^{-1}$ represents the modes of CuO$_2$ planer oxygen atoms$^{[22]}$. The modes in the range from 600 to 700 cm$^{-1}$ are associated with O$_{\delta}$ oxygen atoms in the charge reservoir layers. Apical oxygen modes of type Ti-O$_6$-Cu(2) and Cu(1)-O$_2$-Cu(2) and planer oxygen modes of type Cu(2)-O$_6$-Cu(2) can be observed around 420 cm$^{-1}$, 459 cm$^{-1}$ and 543 cm$^{-1}$, respectively for un-added pure CuTl-1223 sample. All absorption bands almost remained unchanged after the addition of MnFe$_2$O$_4$ nanoparticles. By comparing the FTIR absorption spectra of pure and nanoparticles added samples, we found that the stoichiometry and structure of host CuTl-1223 matrix remained un-disturbed after the inclusion of these MnFe$_2$O$_4$ nanoparticles. Therefore, FTIR absorption spectroscopy also indicated the occupancy of these nanoparticles at inter-granular sites of host CuTl-1223 matrix.

The resistivity versus temperature measurements of (MnFe$_2$O$_4$)$_x$/CuTl-1223; ($x =$ 0, 0.5, 1.5 and 2.0 wt.% ) nanoparticles-superconducting composites are shown in Figure 6. The variation of $T_c$ (K) with respect to various concentrations of MnFe$_2$O$_4$ nanoparticles in CuTl-1223 superconducting matrix is shown in the inset of Figure 6. Normally $T_c$ (0) depends upon micro-structural characteristics of material and carrier’s density in CuO$_2$ planes$^{[23-27]}$. After the addition of MnFe$_2$O$_4$ nanoparticles, the value of $T_c$ (K) was decreased, which showed the reduction in superconducting volume fraction and cooper pairs formation across transition from normal state to superconducting state. The reduction in cooper pairs formation is most probably due to enhanced scattering cross-section caused by the magnetic nature of these particles. Therefore, these nanoparticles act as affective scattering centers at grain-boundaries due to which overall all superconductivity was suppressed.

**Figure 5.** FTIR absorption spectra of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites with (a) $x =$ 0, (b) $x =$ 0.5 wt.%, (c) $x =$ 1.0 wt.%, (d) $x =$ 1.5 Wt.% and (e) $x =$ 2.0 wt. %.

**Figure 6.** Resistivity versus temperature measurements of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites with $x =$ 0, 0.5, 1.5 and 2.0 wt. % and in the inset is shown the variation in $T_c$ (K) with nanoparticles content ($x$).

**Conclusion**

The effects of magnetic MnFe$_2$O$_4$ nanoparticles addition on micro-structural, morphological and transport characteristics of CuTl-1223 phase were studied and compared. The crystal structure and stoichiometry of CuTl-1223 phase were not influenced by the addition of these nanoparticles at grain-boundaries. The suppression of superconductivity was observed with inclusion these nanoparticles. Superconducting volume fraction and cooper pairs density was suppressed due to enhanced scattering cross-section across these magnetic nanoparticles at grain-boundaries. Most challenging problem in these superconducting composites is to make uniform and homogeneous distribution of MnFe$_2$O$_4$ nanoparticles or any other nanostructures at grain-boundaries of host CuTl-1223 superconductor. Therefore, somehow non-monotonic behavior in variation of normal state resistivity and other superconducting parameters of (MnFe$_2$O$_4$)$_x$/CuTl-1223 composites was observed, which was associated with inhomogeneous distribution of MnFe$_2$O$_4$ nanoparticles at the grain-boundaries of bulk CuTl-1223 superconducting phase.
References


22. Firdous, T., Khan, N.A. Sn doped (Cu$_{0.5}$Tl$_{0.5}$)Ba$_2$Ca$_2$Cu$_{3−x}$O$_{7−δ}$ superconductors: Effect on the diamagnetism and phonon modes. (2009) J Appl Phys 106(8).


