

## Dielectric Properties of Magnetite-Containing Core–Satellite-Type Structures

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### Abstract

BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> core–satellite particles were synthesized and incorporated into a PVDF matrix to obtain hybrid nanocomposite films. XRD analysis revealed reflections at 31.53°, 35.59°, 56.12°, and 62.73°, confirming the coexistence of tetragonal BaTiO<sub>3</sub> and cubic Fe<sub>3</sub>O<sub>4</sub> phases in the hybrid structure. Dielectric spectroscopy in the range 25 Hz to 1 MHz showed a frequency-dependent decrease in dielectric permittivity for all compositions, consistent with the reduced contribution of interfacial polarization at higher frequencies. The dielectric constant of neat PVDF was about 8 at 1 kHz, while the addition of hybrid filler increased this value. For the sample containing 7% BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub>, the dielectric permittivity reached values close to 60. These findings confirm that the introduction of BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> hybrid particles substantially enhances the polarization capability of the polymer matrix and improves the dielectric characteristics of the resulting nanocomposites

**Keywords:** Nanocomposite, magnetite, dielectric properties, dielectric polarization

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### 1. Introduction

In recent decades, the intensive development of nanotechnology has led to the creation of a new class of functional materials, namely nanocomposites with con-

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trollable structure and properties[1]. Of particular interest are magnetic nanostructures based on iron oxides, especially  $\text{Fe}_3\text{O}_4$ , due to their unique combination of magnetic, catalytic, and electrochemical characteristics[2-3]. Such materials have found wide application in catalysis, biomedicine, sensing, energy, and environmental technologies [4].

One of the most promising directions is the development of nanocomposites with core-shell and core-satellite architectures, which make it possible to purposefully combine the properties of different components within a single system[5-8]. In core-shell structures, the inner core provides the basic functional properties, such as magnetic or conductive behavior, while the shell performs a protective, stabilizing, or functionalizing role [9]. In turn, core-satellite structures are characterized by the presence of nanoparticles distributed over the surface of a magnetic core, which increases the active surface area and enhances interfacial interactions [10].

Such hybrid structures exhibit a pronounced synergistic effect [11]. For example, the presence of metallic nanoparticles promotes faster electron transfer and higher catalytic activity, while the magnetic core enables efficient separation of the material from the reaction medium by means of an external magnetic field. The relevance of research in this field is also determined by the need to address environmental and energy-related challenges. In particular, the development of efficient nanocatalysts makes it possible to reduce toxic organic compounds into less hazardous products and to create highly sensitive sensor systems for detecting environmental pollutants. Thus, the development and study of  $\text{Fe}_3\text{O}_4$ -based nanocomposites with controlled core-shell and core-satellite morphology represent an important scientific task. Investigation of their structure, properties, and interaction mechanisms opens broad prospects for the creation of new functional materials with tailored characteristics for application in various fields of science and technology.

The aim of this work is to synthesize  $\text{BaTiO}_3$ - $\text{Fe}_3\text{O}_4$  core-satellite structures, to fabricate nanocomposites based on the thermoplastic polymer PVDF with uniform distribution of these structures in the polymer matrix, and to investigate the structure and dielectric properties of the obtained nanocomposites.

## **2. Experimental methodology**

### **2.1. Synthesis of $\text{BaTiO}_3$ - $\text{Fe}_3\text{O}_4$ core-satellite structures**

0.002 g of polyvinylpyrrolidone was first dissolved in 50 mL of distilled water. Subsequently, 1 g of  $\text{BaTiO}_3$  was introduced into the solution, and the mixture was stirred at 70°C for 1 h. In the next step,  $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$  and  $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$  were added in amounts of 0.008 and 0.015 mol, respectively. Ammonium hydroxide ( $\text{NH}_3 \cdot \text{H}_2\text{O}$ ) was then added until the pH of the reaction medium reached 11–12. After comple-

tion of the reaction, the mixture was cooled to room temperature, and the resulting product was washed six times with ethanol and dried under vacuum at 45°C. This procedure yielded BaTiO<sub>3</sub> particles whose surfaces were decorated with Fe<sub>3</sub>O<sub>4</sub> nanoparticles.

## **2.2. Preparation PVDF- BaTiO<sub>3</sub>-Fe<sub>3</sub>O<sub>4</sub> nanocomposites**

For the preparation of nanocomposite films, the synthesized BaTiO<sub>3</sub>-Fe<sub>3</sub>O<sub>4</sub> particles were introduced into a thermoplastic PVDF matrix. Thin films were fabricated by solution mixing. First, PVDF was completely dissolved in N,N-dimethylformamide, after which the required amount of BaTiO<sub>3</sub>-Fe<sub>3</sub>O<sub>4</sub> particles was added to the solution. The resulting mixture was stirred for 3 h to ensure uniform filler dispersion, then cast into a glass vessel and dried. As a result, homogeneous PVDF-based nanocomposite structures containing the hybrid BaTiO<sub>3</sub>-Fe<sub>3</sub>O<sub>4</sub> filler were obtained.

## **2.3. Characterization of nanocomposites**

### **2.3.1. X-ray analysis**

Rigaku Mini Flex 600 XRD diffractometer (CuK $\alpha$  radiation from Cu X-ray tube, run at 15 mA and 30 kV) was used for X-ray diffraction analysis at ambient temperature. The nanocomposite films were scanned in the 2 $\theta$  angle range from 10 to 100°.

### **2.3.2. Transition electron microscopy (SEM) analysis**

TEM analysis of nanocomposites was performed on a JEOL JEM-1400 (Japan), mode at a working voltage of 80-120kV.

### **2.3.3. Dielectric measurement**

Dielectric permittivity of the nanocomposites was determined using a broadband immittance analyzer (MNIPI E7-20). Frequency-dependent measurements were conducted in the range of 25 Hz to 1 MHz at room temperature.

## **3. Experimental results and their discussion**

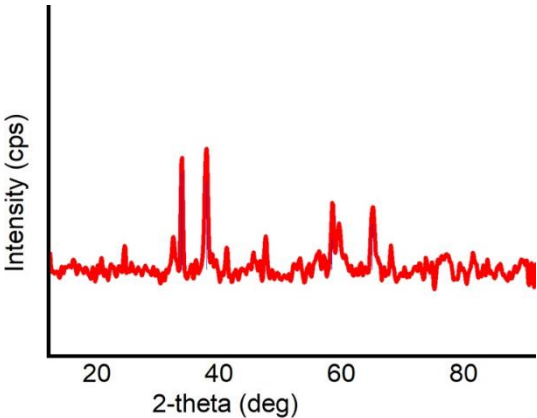
The structure of BaTiO<sub>3</sub> nanoparticles coated with Fe<sub>3</sub>O<sub>4</sub> nanoparticles on their surface was investigated by the X-ray diffraction method. The table 1. presents various parameters of the diffraction lines corresponding to the observed peaks. Based on the data presented in the table, it can be concluded that the sample contains both a tetragonal BaTiO<sub>3</sub> core and a cubic Fe<sub>3</sub>O<sub>4</sub> shell, while PVP, being amorphous,

does not produce diffraction peaks [12]. In the 56–58° range, the BaTiO<sub>3</sub> (211) and Fe<sub>3</sub>O<sub>4</sub> (511) reflections overlap.

**Table 1.** XRD data of BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> particles

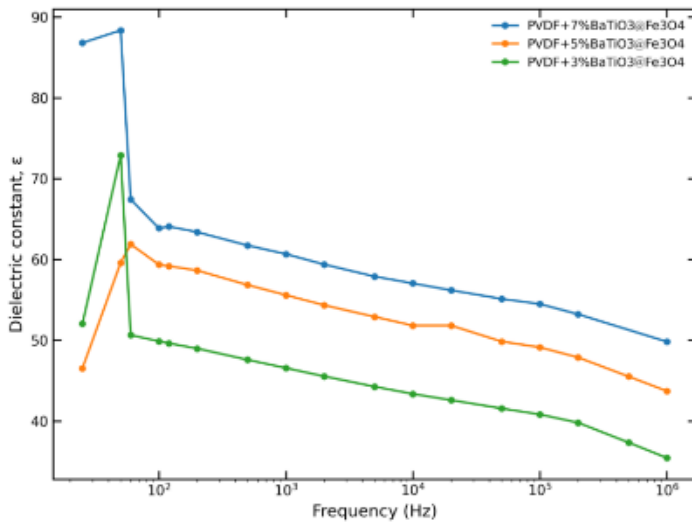
| 2θ(deg) | (hkl) index                                                                | Characteristic phase                  |
|---------|----------------------------------------------------------------------------|---------------------------------------|
| 31.53   | BaTiO <sub>3</sub> (110)/(101)                                             | a tetragonal BaTiO <sub>3</sub> core- |
| 35.59   | Fe <sub>3</sub> O <sub>4</sub> (311)                                       | magnetite                             |
| 56.12   | Fe <sub>3</sub> O <sub>4</sub> (511)+BaTiO <sub>3</sub> (211) (overlapped) | Both phase                            |
| 62.73   | Fe <sub>3</sub> O <sub>4</sub> (440)                                       | Magnetite                             |

Figure 2 presents the variation of dielectric constant ( $\epsilon$ ) with frequency for PVDF nanocomposites containing BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> hybrid fillers. In all cases,  $\epsilon$  decreases with increasing frequency, which is a characteristic feature of polymer-based dielectric composites. This behavior is generally associated with interfacial polarization effects. At lower frequencies, space charges can migrate and accumulate at the interfaces between phases, resulting in large dielectric constant values. With increasing frequency, the ability of these charges to respond to the alternating field becomes limited, and the contribution of interfacial polarization correspondingly diminishes, causing a reduction in  $\epsilon$ .



**Figure. 1.** X-ray diffraction (XRD) patterns of the BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> particles

The dielectric permittivity of neat PVDF is about 8 at 1 kHz. In contrast, the incorporation of BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> hybrid nanoparticles leads to a significant increase in dielectric constant at the same frequency. According to Ma et al. (2017), particles with a grain size of 122 nm possess a dielectric permittivity of approximately 2000



**Figure 2.** Dielectric constant as a function of frequency for PVDF and BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub>- based nanocomposite

at 1 kHz [13]. However, in the present nanocomposites, the measured dielectric constant is considerably lower, and even for the sample filled with 7% BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub>, the  $\epsilon$  value remains below 60. Such a difference may be related to the magnetite coating on the surface of BaTiO<sub>3</sub> particles, which can alter their dielectric contribution, as well as to the relatively low amount of hybrid filler dispersed in the PVDF matrix. Despite this, the increase in dielectric permittivity from about 7–8 for neat PVDF to values approaching 60 clearly demonstrates that the hybrid filler effectively enhances the overall polarization of the system.

## Conclusion

In conclusion, novel core–satellite structures consisting of BaTiO<sub>3</sub> particles decorated with Fe<sub>3</sub>O<sub>4</sub> nanoparticles on their surface were successfully synthesized, and on this basis PVDF-matrix nanocomposites were prepared. The X-ray diffraction results confirm the successful formation of BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> hybrid nanoparticles composed of a tetragonal BaTiO<sub>3</sub> core and a cubic Fe<sub>3</sub>O<sub>4</sub> shell, while the absence of diffraction peaks from PVP is consistent with its amorphous nature. This structural organization plays a key role in determining the dielectric behavior of the PVDF-based nanocomposites. The incorporation of these hybrid fillers leads to a pronounced increase in dielectric constant compared with neat PVDF, confirming that the hybrid particles enhance the overall polarization of the system. At the same time, the observed decrease in dielectric constant with increasing frequency indicates that interfacial po-

larization is the dominant mechanism governing the dielectric response. Although the dielectric permittivity of the nanocomposites remains much lower than that of pure BaTiO<sub>3</sub>, this can be explained by the Fe<sub>3</sub>O<sub>4</sub> coating on the BaTiO<sub>3</sub> surface and the relatively low filler content in the PVDF matrix. Nevertheless, the increase in dielectric permittivity from about 8 for neat PVDF to values approaching 60 demonstrates that even a limited amount of BaTiO<sub>3</sub>–Fe<sub>3</sub>O<sub>4</sub> hybrid filler is sufficient to significantly improve the dielectric performance of the polymer matrix.

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