Research Article

The Electrical Performance of Polyamide 66/Poly(vinylidene fluoride) with Vinyl Acetate-Maleic Anhydride Copolymer

Rui Li, 1 Zhiliang Shi, 1 Dongliang Kuang, 2 and Jianzhong Pei 1

1Highway School, Chang’an University, Xi’an 710064, China
2School of Materials Science and Engineering, Chang’an University, Xi’an 710064, China

Correspondence should be addressed to Jianzhong Pei; peijianzhong@126.com

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The electric performance of the PA66/PVDF blends filled with various amount of copolymer synthesized from vinyl acetate–maleic anhydride (VAMA) was investigated. PA66/VAMA/PVDF blends show high dielectric constants, low dielectric loss, and excellent breakdown strength, which were important indexes in the actual application of dielectric material. The VAMA copolymer improves the dielectric and piezoelectric performance of the PA66/PVDF blends. Meanwhile, the addition of VAMA obviously decreases the dielectric loss and breakdown strength of the blends. PA66/PVDF blends filled with 3 wt% VAMA exhibited the best electric ability. The stable dielectric constants of the all-polymeric blends can be tuned by adjusting the content of the VAMA. The created all-polymeric blends represent a novel dielectric material that is technologically simple and easy to process forward application for flexible electronics.

1. Introduction

With the continuous development of science and technology, the need of high performance material urges people to open up the composites, which display their advantages outweighing their components, and also it is hard for single polymer to meet requirements [1–3]. Polivinylidene fluoride (PVDF) and Polyamides (PA) are two famous polar polymers. In particular, PVDF is known as polymer composites materials characterized by excellent dielectric and piezoelectric properties [4, 5]. However, in the actual application, the high cost of PA11 and the high melting point (220–240°C) of PA66 limited the applications in many fields. Also there are still few studies about PA66/PVDF blends. In addition, the PA66/PVDF blends are an interesting system because of some specific interactions [6–8]. As a new material, a blend of PA66 and PVDF offers a strong potential for the application in the field of capacitor [9]. According to the property index requirements of energy storage material, the polymers selected should have two features: (1) the polymer itself should be partly crystallized and (2) the molecular chain of the polymer should have polar groups. This makes the polymer have a more sophisticated orientation under the action of electric field. While improving the dielectric constant of polymer, it can be used as energy storage material with the inherent low breakdown strength [10]. Therefore, VAMA offers a strong potential in the application of the all-polymer blends [11].

The purpose of this work is to investigate the dielectric and piezoelectric performance of the PA66/PVDF blends filled with various amount of VAMA. Firstly, the copolymer VAMA from acetic acid vinyl ester and maleic anhydride was synthesized. Through grafting reaction can lower the melting point of PA66 and the PA66/PVDF blends obtained from PA66/VAMA/PVDF by melt blending through torque rheometer. It is expected that this research can be useful in promoting the applications of the PA66/PVDF blends under oil-free lubrication conditions.

2. Experimental

2.1. Materials. PVDF is from Shanghai 3F Material Co., Ltd., \( T_m = 175^\circ C \), and \( d = 1.75 \text{ g/cm}^3 \). PA66 is from Sinopec Baling Branch Co., Ltd., \( T_m = 225^\circ C \), and \( d = 1.15 \text{ g/cm}^3 \). The related products are commercially available.
2.2. Synthesis of Graft Copolymer of Vinyl Acetate-Maleic Anhydride. The methylbenzene was added to a three-mouth flask, which was equipped with stirrer, thermometer, and reflux condensing tube, and then the maleic anhydride was added and dissolved into methylbenzene slowly after stirring and heating. After the solution turns to be transparent, acetic acid vinyl ester is added. Also, 1.4% initiator azodisobutyronitrile was added. The temperature is controlled at 90°C. The copolymer is subsided for 3 h, filtered, and washed with dichloroethane for 5 times, removing the unreacted maleic anhydride. After vacuum drying under 60°C, white powdered graft copolymer can be obtained [12]. Figure 1 is the reaction scheme of VAc and MAH copolymerization.

2.3. Preparation of PA66/VAMA/PVDF. PA66, VAMA, and PVDF were mixed for 45 min based on different volume ratios in torque rheometer, and then press vulcanizer was used to press the evenly pressed PA66/VAMA/PVDF blends into thin plate. In addition, microtablet machine is used to press plate into around 1 mm. Finally, the samples were polished and silvered by silver paint.

2.4. Instrumental Analysis. The macrostructures of blends were characterized by scanning electron microscopy (SEM) and Fourier transform infrared spectroscopy (FTIR). The electric properties of the samples were performed by HIOKI3532-50 LCR; the frequency is from $10^2$ to $10^6$ Hz at room temperature.

3. Results and Discussions

3.1. Structure Analysis. The photographs of the films were shown in Figure 2(a). It can be seen from Figure 2(b) that PA66/PVDF blends present a unique structure of ordered arrangement which is evenly distributed. The infrared spectrum of PA66/VAMA/PVDF was tested by Fourier transformation infrared spectrometric analyzer. It can be seen from Figure 3 that the absorption peak where the wavenumber of 1780 cm$^{-1}$ was the stretching vibration of carboxylic acid C=O and 1374 cm$^{-1}$ was the flexural vibration of methyl.
The strength of the absorption band of 1314 cm\(^{-1}\) was greater than that of 1433 cm\(^{-1}\), which is the significant characteristic of vinyl acetate structure, which proves the reaction of copolymerization between acetic acid vinyl ester and maleic anhydride. Also, it can be found that the two acid anhydride peaks disappear at 1780 cm\(^{-1}\) and 1740 cm\(^{-1}\). When the infrared spectral curve of PA66/VAMA/PVDF was compared with that of copolymer VAMA, it can be found that PA66/VAMA/PVDF characteristic peak of cyclic amide disappears at 1414 cm\(^{-1}\), which shows that the anhydride of VAMA molecule has chemical reaction with -NH of PA66.

3.2. Dielectric Properties. At room temperature, when the frequency is \(10^3\) Hz, the dielectric properties of PA66/VAMA/PVDF blends are obtained for analysis after adding different mass fractions of VAMA (1 wt%, 2 wt%, 3 wt%, and 5 wt%) as shown in Figure 4. The quantities of VAMA added are those by which it can improve not only the dielectric constant of PA66/PVDF blends that have been improved but also the dielectric loss of blends that have been down. It proved that VAMA can make polar dipoles easier to be oriented within PA66/PVDF blends, thus lowering the dielectric loss of blends. The relation curve of the dielectric constants between the PA66/VAMA/PVDF blends and the content of VAMA was shown in Figure 4. Obviously, the influence of copolymer VAMA on the dielectric constant of PA66/VAMA/PVDF blends is 3 wt% > 2 wt% > 5 wt% > 1 wt%. When PA66 and PVDF were melted and blended, adding VAMA can improve the molecules flexibility of PA and PVDF, and also the dielectric constant of PA66/VAMA/PVDF can be improved. However, excessive VAMA was prone to the grafting ratio of PA66, which is not good for the collection and delivery of the system charge. Therefore, the results indicate that, after grafting, a distinct layer structure is formed in PA66/VAMA/PVDF blends. The optimum content of the VAMA was 3%.

3.3. Piezoelectric Properties. The PA66/PVDF blends with 0 wt%, 1 wt%, 2 wt%, 3 wt%, and 5 wt% VAMA were chosen for studying piezoelectric coefficient \(d_{33}\) owing to the highest dielectric constant and good interfacial compatibility. Figure 5 shows the relationship between the piezoelectric constant \(d_{33}\) and the different VAMA and PA66 content. Although the samples do not exhibit super piezoelectric properties, PA66/VAMA/PVDF blends have a better enhancement in the piezoelectric constant \(d_{33}\) than that of the PA66/PVDF at room temperature. The results show
that the polarization degrees of PA66/VAMA/PVDF were enhanced by VAMA so as to improve the piezoelectric properties.

3.4. Breakdown Strength. Breakdown strength is an important index in the actual application of dielectric material. Regarding the material structure, there are many factors which influence the breakdown strength, which are distribution uniformity of crystal particle, crystal size, and the material defects such as internal air holes. In addition, to varying degrees, the shape of samples, thickness, and the shape of electrode can also influence the final breakdown strength. In order to reduce the influence of material structure on the breakdown strength, samples which were 1 mm thick were selected in this experiment.

The breakdown strength of PA66/VAMA/PVDF blends with PA66 volume fraction increase was shown in Figure 6. When the volume of PA66 was 40%, PA66/VAMA/PVDF blends reach the maximum breakdown strength \( (E_b = 15.3 \text{ kV/mm}) \). And when the volume content of PA66 was 60 vol.%, the breakdown strength of the material has the minimum \( (E_b = 9.9 \text{ kV/mm}) \). This is because the two polymer phases (PA66 and PVDF) within PA66/VAMA/PVDF samples were not bonded well (PA66 : PVDF = 60 : 40), which cause a low breakdown strength.

Combining the dielectric property test result of PA66/VAMA/PVDF blends, the sample with PA66 : PVDF = 20 : 80 has the best dielectric properties. In order to study the influence of thickness on the breakdown strength of samples, PA66/VAMA/PVDF blends were applied to prepare different thicknesses of samples to have breakdown experiment in the volume ratio of PA66 : PVDF = 20 : 80. Figure 7 is the relational curve between breakdown strength and blends thickness. It can be found from it that, with the increase of thickness, the breakdown strength declines sharply. When the thicknesses were increased to 2.0 mm from 0.5 mm, the breakdown strength declined from 11.9 kV/mm continuously. Obviously, the thickness of samples has a significant influence on the breakdown strength. The reason is because the defects in thicker blends than in thin blends. Thereby, the breakdown strength of blends can be decreased greatly. On the other hand, due to the fringe effect of the electric field intensity, the increase of thickness strengthens the electric field for few times on the fringe of electrode. Consequently, the breakdown strength of PA66/VAMA/PVDF blends declined.

The result shows that it is easier for these defects to cause local electric field distortion under the same electric field and testing conditions. Thus, the probability to cause breakdown to dielectric medium is increased. Therefore, with the increase of testing thickness, the breakdown strength will be decreased sharply. Meanwhile, Kuang et al. found that the electric field distortion of plate electrode was significant by applying electric field simulation to different electrode structures, and the electric field being enhanced on the fringe of electrode makes it easier for the air holes on the surface of medium around electrode fringe to cause gas discharge, which caused breakdown to the entire medium \([12, 13]\).
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