

CHAPTER V

POLY(VINYLDENE FLUORIDE-CO-HEXAFLUOROPROPYLENE) /BACTERIAL CELLULOSE (PVDF-HFP/BC) COMPOSITE FILM

5.1 Abstract

Poly(vinylidene fluoride-co-Hexafluoropropylene) (PVDF-HFP) composites were fabricated via twin screw extruder to achieve the high dielectric and piezoelectric properties and also utilized as a touch sensor. This chapter, extracted bacterial cellulose (BC) from Nata de coco in 1-5 wt.% were provided as an appropriate filler to improve the dipole alignment of PVDF-HFP matrix. The cast film extruder was used to produce the transparent composite films with unique properties. The increment in β -phase crystalline presented with higher amount of BC. The dielectric constant corresponded to piezoelectric coefficient was enhanced from 2.00 of neat PVDF-HFP to 3.25 with 5 wt.% BC loading. Besides, the presence of BC in the composite films led to an improvement in thermal properties, and mechanical properties in terms of Young's modulus and tensile strength with no dimensional changes at 110 °C due to the excellent thermal, and mechanical properties of BC structure.

5.2 Introduction

Films of some polymers, such as polyvinylidene fluoride (PVDF) and its copolymers, opportunely conditioned, exhibit piezoelectric and pyroelectric properties [1–7] with excellent sensitivity and a satisfactory dynamic response. Moreover, they can be reduced to tiny compliant strips for embedding in a continuous elastic layer. Used as isolated sensors/actuators or assembled into more complex multimodal devices, they offer challenging solutions in many application fields including robotics. For these reasons they can be regarded as interesting candidates for touch sensors or touch screen panel [8]. PVDF is synthesized by addition polymerization of the CH₂CF₂ monomer. When produced as the homopolymer (i.e. from 100% CH₂CF₂ monomer), the majority of the PVDF chains have a regular structure of alternating CH₂ and CF₂ groups. The crystalline phase of interest for PVDF ferroelectricity is

the polar β -phase. This structure can be obtained either by using the PVDF trifluoroethylene copolymer [P(VDF-TrFE)] which has the tendency to crystallize directly in the polar β -phase or by mechanical stretching of the α -phase [9-12], from melt under specific conditions such as high pressure [13-14], external electric field [15] and ultra-fast cooling [16]

However, there is a need to develop a touch sensor with high mechanical properties for their long term operation and tolerance to the applied force during electronic device lifetime. Up to the present time, in order to fulfill flexible characteristic and maintain the excellence on dimensional stability, one of the materials found to be promising was transparent cellulose composite. The use of cellulose also generally provides numerous advantages including biodegradability, light weight, transparency, including high strength. Cellulose is typically separated from lignocellulosic plants, such as wood and agricultural crops, using mechanical treatment. Usually, lignin is also removed from the plant cell wall prior to fibrillation using chemical treatment. Moreover, some of researchers have proposed the novel method of cellulose preparation from bacterial species which called bacterial cellulose (BC). The uniformity of cellulose prepared from bacterial is easier to control by means of bacterial digestion mechanism than the plant-based cellulose [17]. So in this chapter, we focus on the BC as a filler.

Due to the fact that cellulose is hydrophilic and forms irreversible aggregates when it dries, the most successful methods used to prepare the cellulose composites have been solution casting of dilute slurry of matrix and cellulose. However, for larger scale production these methods are slow and expensive. Extrusion compounding is one of the most promising methods for industrial processing due to easy scale-up and the possibility of further molding of the materials. It also tends to reduce environmental hazards. There are only few published studies on extrusion processing of cellulose composites which is why more research is needed.

The objective of this work is to develop the transparent and flexible touch sensor based on piezoelectric material for touch screen applications. We used bacterial cellulose (BC), extracted from *Nata de coco*, as a filler to improve the dipole alignment of PVDF-HFP matrix by using melt mixing method followed by cast film process. The dielectric constant was measured as the electrical and piezoelectric

properties. In addition, the morphological, thermal and mechanical properties were analyzed and discussed based on basis properties for touch screen applications.

5.3 Experimental

5.3.1 Materials

Poly(vinylidene fluoride-co-Hexafluoropropylene) or PVDF-HFP with a melt flow rate of 7 g/10 min (5 kg at 230°C). Solef[®] 1010/1001 from Solvay was purchased from Prostar Chemicals Co., Ltd., Thailand.; Nata de coco was purchased from local food market; Sodium hydroxide (NaOH) (AR grade, 98%) were purchased from Merck Ltd.

5.3.2 Methods

5.3.2.1 *Bacterial cellulose (BC) preparation [17]*

BC was extracted from the Nata de coco product. The Nata de coco was rinsed with distilled water to remove some excess sugar and blended using a laboratory blender to reduce the size and then was treated in 0.1M sodium hydroxide (NaOH) at 80°C for an hour to remove any remaining microorganisms, medium component and soluble polysaccharides. The purified bacterial cellulose (BC) was then thoroughly washed with distilled water until reaching neutral pH. After the extraction procedure, the BC was dried at 60 °C for 2 days. Then, the dry BC was sieved to produce uniform powder with size less than 74 µm.

5.3.2.2 *PVDF-HFP/BC composite film preparation*

All materials were generously pre-compounded and pelletized using a co-rotating twin-screw extruder (LTE-20-40 Lab Tech). The PVDF-HFP pellets and BC powders were dried at 60°C for 48 hours prior to remove adsorbing water. The amount of blends prepared was 1 kg per each blend ratio, depending on main polymer. Pre-mixtures of PVDF-HFP with BC with the dry weight ratio between PVDF-HFP:BC varied at 1000:0, 99:1, 97:3, and 95:5 % by weight were blended in a laboratory blender to form uniform powder. The amount of materials prepared of each blend ratio was reported in Table 5.1.

Table 5.1 Amount of materials prepared for each blend ratio

Formula	PVDF-HFP (g)	BC (g)
PVDF-HFP	1000	-
PVDF-HFP/BC (1 wt.%)	990	10
PVDF-HFP/BC (3 wt.%)	970	30
PVDF-HFP/BC (5 wt.%)	950	50

PVDF-HFP/BC composites were melted mixing in a twin screw extruder. The processing temperature, residence time and the screw speed were maintained at 150-160-165-170-175°C, 10 min and 30 rpm, respectively. The extruder barrel was equipped with two atmospheric vents and vacuum ventilation in purpose to remove the vaporized water from the material.

After pelletizing, the PVDF-HFP and its composite pellets were dried at 60°C for 48 hours before film casting process to prepare the thin film. The temperatures of the extruder were set at 170-180°C. The screw speed for the samples was 65 rpm. The die temperature was fixed at 180°C. The chill roll was placed in 2 cm from the die and its temperature was kept at 20°C by temperature controller. The film was quenched by a chill roll and was transported to a pulling station using a nip roll. The speed of chill and nip rolls were controlled separately from the extruder using a dial and digital display. As reference, the neat PVDF-HFP film was also prepared in similar manner.

5.4 Results and Discussions

5.4.1 Bacterial Cellulose (BC) Characterization

5.4.1.1 *Morphological Properties*

Figure 5.1 shows the visual appearance and SEM images of BC. The BC displays irregular shape and apparent size dispersion. The particles size varied widely in the range of 30-70 μm .

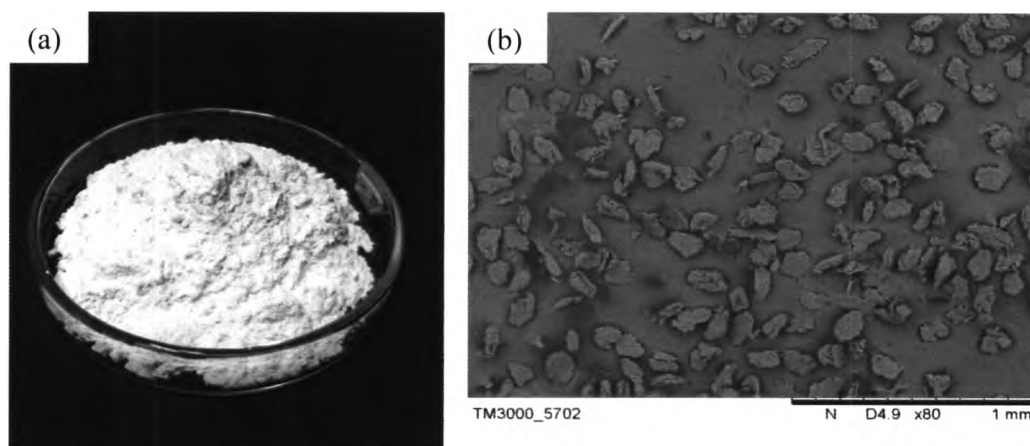


Figure 5.1 (a) The visual appearance and (b) SEM image of BC.

5.4.1.2 Crystallinity

XRD diffractogram of BC is shown in Figure 5.2. The result shows three main diffraction peaks at 2θ of 15.5° , 17.0° , 22.5° , and 35.0° which refer to the 110, 110, 020, and 004 diffraction planes, respectively. These diffractions correspond to the typical profile of the cellulose I allomorph [18-19]. The crystallinity of BC is 70% which was calculated based on crystalline peak.

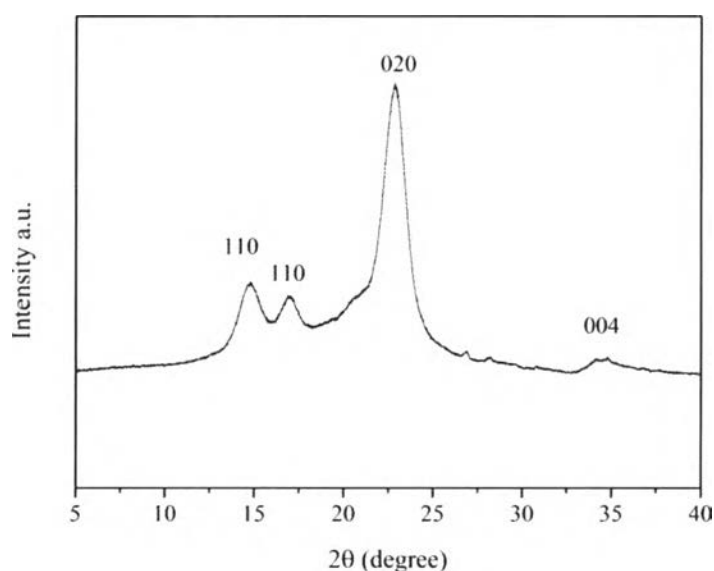


Figure 5.2 XRD diffractogram of BC.

5.4.1.3 Chemical Properties

The FT-IR spectra of BC is shown in Figure 5.3. The structure of BC is linear homopolymer composed of D-glucopuranose units linked by β -1,4-glycosidic linkage at C₁ and C₄ carbon position [20]. The peak at 3332 cm⁻¹ referred to O-H stretching vibration, while the peak at 2893 cm⁻¹ and 1429 cm⁻¹ referred to C-H stretching and symmetric blending of -CH₂, respectively. The peak at 897 cm⁻¹ indicated to C-O stretching [21]. The peaks at 1429, 1110, and 897 cm⁻¹ were thought to be the typical bonds of cellulose I (I β) [22], which was consistent with XRD diffractogram.

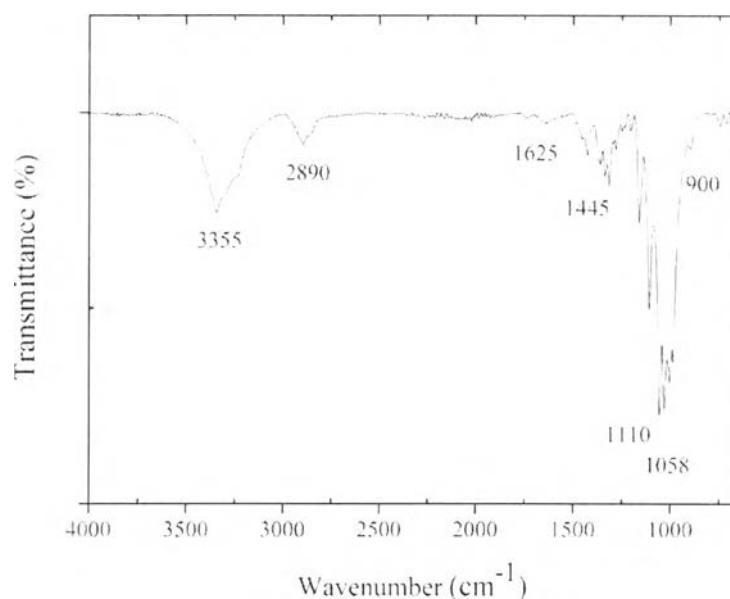


Figure 5.3 FT-IR Spectra of BC.

5.4.1.4 Thermal Properties

TGA thermogram of the BC (a) and its derivative peak (b) are shown in Figure 4.4. There are two thermal degradation steps corresponding to the vaporization of adsorbed water in the BC (start from 40°C) and the degradation process such as depolymerization, dehydration and carbonization of BC chain (start from 300 °C). The degradation temperature (T_d) of BC was 343°C [23]. When temperature reaches 400 °C the formation of charred residue is occurred and its remaining weight was evaluated at approximately 8% referring to char residue.

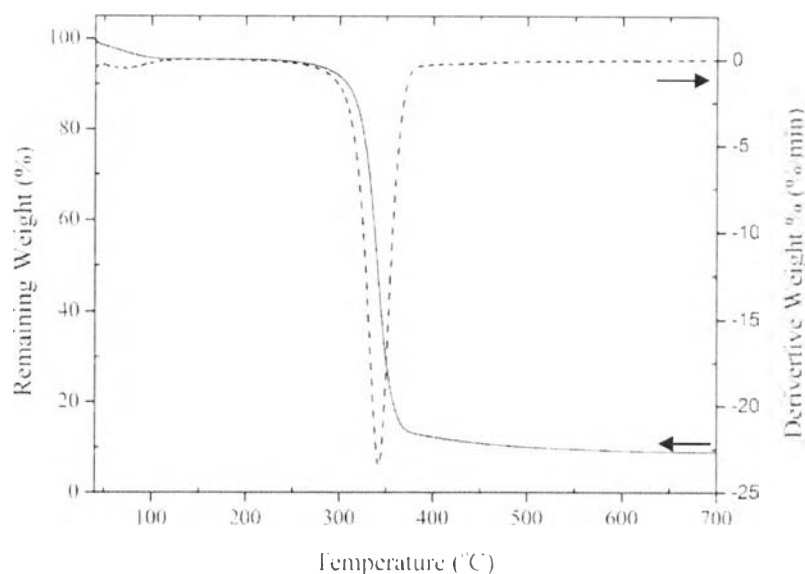


Figure 5.4 TGA thermogram of BC (—) and its derivative (---).

5.4.2 PVDF-HFP and PVDF-HFP/BC Films Characterization

5.4.2.1 Morphological Properties

Extruded pellet and casted films with a thickness ranging between 0.1 and 0.2 mm are shown in Figure 5.5. Both of neat PVDF-HFP pellet and casted film were transparency. After mixing, the composite pellets and films were more opaque than neat PVDF-HFP in agreements with observations reported previously by Fortunati (2010) [24-25]. This optical result can be confirmed by UV/Vis spectra (Figure 5.6). Neat PVDF-HFP film has 50-70% transmittance in the visible light region (400-750 nm), but when adding BC, the transmittance tends to decrease, which means BC exhibits the opacity property. It is due to the formation of a network structure in the BC. It can be suggested that with high amount of BC may segregate from the blend to form a micro-sized aggregation which can be observed in the visible light region. Similar to Auras's work, cellulose show a blocking effect on the ultra violet C region (280–100 nm), usually produced by artificial light sources [26].

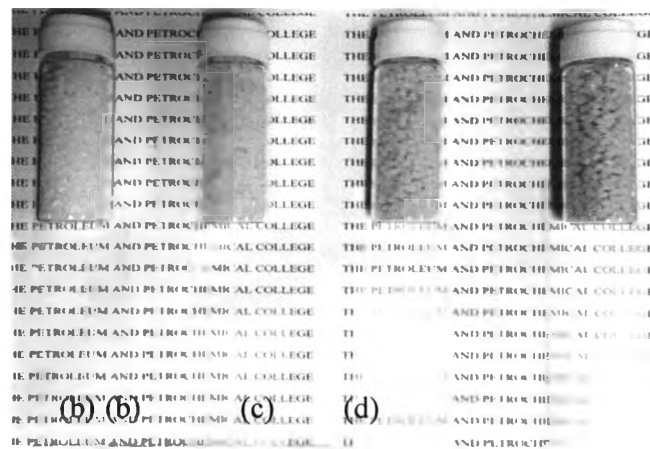


Figure 5.5 The visual appearance of extruded pellet and casted film of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

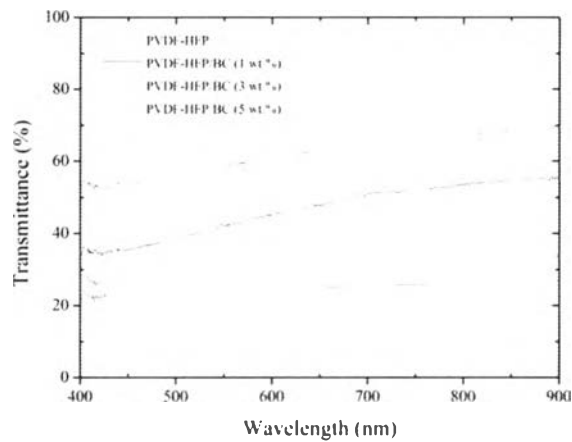


Figure 5.6 The UV/Vis spectra of neat PVDF-HFP and PVDF-HFP/BC composite films.

The dispersion of BC particles in the PVDF-HFP matrix were also observed by an optical microscope. Figure 5.8 shows the Optical microscope images. At low content of 1-3 wt.% (Figure 5.8b-c), most BC particles dispersed uniformly in the PVDF-HFP matrix without agglomeration and obtain a good orientation in machine direction. While the aggregation of BC particles increased with an increasing BC content (Figure 5.8d).

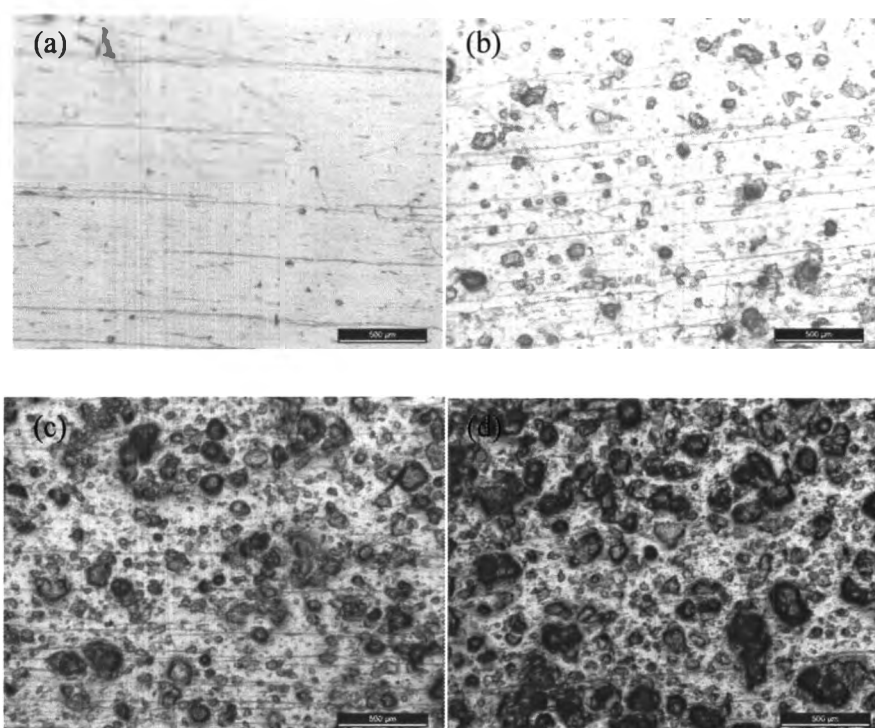


Figure 5.7 The Optical microscope images of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

5.4.2.2 Crystallinity

PVDF, the main constituent of the PVDF-HFP copolymer used in this work as polymer host, can present several different crystalline structures. In general, the functional group of PVDF ($-\text{CH}_2-\text{CF}_2-$) exist 5 molecular conforma-

mations (α , β , δ , γ , and ϵ). The α and β phases are commonly studied and also the most important polymorphs for PVDF and its copolymers.

The absorption bands characteristic of neat PVDF-HFP and PVDF-HFP/BC composite films was shown in Figure 5.8. The basic characteristic absorption peaks of PVDF-HFP, *i.e.* CH₂ wagging, antisymmetric CF₂ stretch, and CF₃ out-of-plane deformation absorb IR radiation at 1402, 1180 and 1065 cm⁻¹ [26-27], respectively. Moreover, the absorption band of α -phase, which is a non-polar trans-gauche-trans- gauche' (TGTG') conformation, were observed at 763 cm⁻¹ (CF₂ bending), and 795 cm⁻¹ (CF₂ rocking). While the presence of β -phase, all-trans (TTTT) or planar zigzag conformations observed by the specific absorption bands at 840 and 1285 cm⁻¹ (CH₂ rocking and bending, respectively) [26-29]. While the amorphous region, which was observed at 873 cm⁻¹, come from the HFP phase of PVDF-HFP.

According to Equation 3.2, the fraction of β -phase, $F(\beta)$, was calculated from FTIR spectra and presented in Table 5.2, assuming that there is no absorbance contribution from the BC. With the higher amount of BC incorporated, β phase tends to be increased because crystallization was obstructed by the network of BC. The β -phase content for neat PVDF-HFP film is 47% and increase with adding BC. This result indicated that the introduction of BC probably acting as a nucleation site led to a change of the crystalline phase from α to β phase crystals.

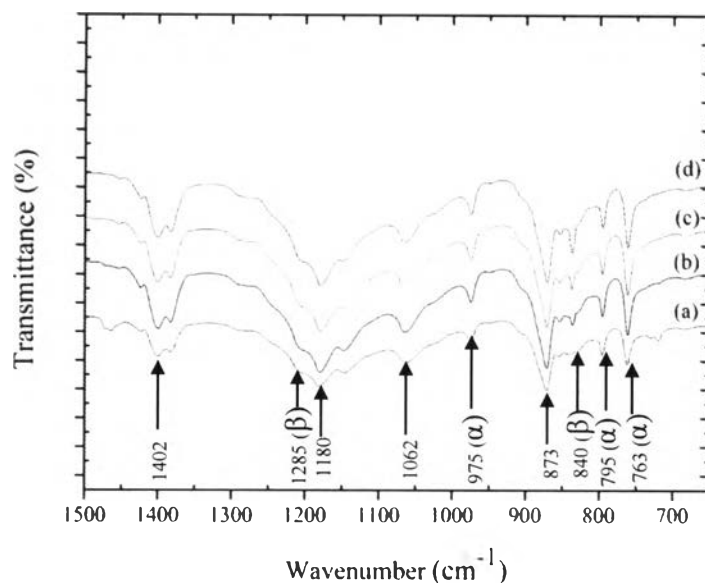


Figure 5.8 FT-IR spectra of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

Table 5.2 β -phase fraction; $F(\beta)$, of neat PVDF-HFP and its composite films

Formula	$F(\beta)$
PVDF-HFP	0.47
PVDF-HFP/BC (1 wt.%)	0.55
PVDF-HFP/BC (3 wt.%)	0.56
PVDF-HFP/BC (5 wt.%)	0.56

To confirm the presence of β -phase in PVDF-HFP/BC composites, XRD analysis was also used to investigate. Figure 5.9 shows XRD patterns of crystalline phase of PVDF-HFP and its composite films with 1, 3, and 5wt.% of BC loading. Similar to Chang's work [31], there were two characteristic peaks which refers to α and β -phases. In our study, the characteristic peaks at 17.5° , 17.9° , 20.0° , and 27.0° corresponding to monoclinic α phase of $\alpha(0\ 2\ 0)$, $\alpha(1\ 0\ 0)$, $\alpha(1\ 1\ 0)$, and $\alpha(0\ 2\ 1)$, respectively [32]. In the case of PVDF-HFP/BC composites, a new reflection peak at $2\theta = 20.4^\circ$ corresponding to β phase, which has orthorhombic unit cell,

appeared, and the peak intensity increased with increasing BC content which was corresponded to the FT-IR result. This result indicated that the introduction of BC probably acting as a nucleation site led to a change of the crystalline phase from α to β phase crystals. As well documented in literature [33-34], α -form crystal is nonpolar phase (trans-gauche-trans-gauche' (TGTG')), while β -form crystal shows polar properties. Therefore, it can be inferred that the polarity of the materials increased as the content of BC increasing.

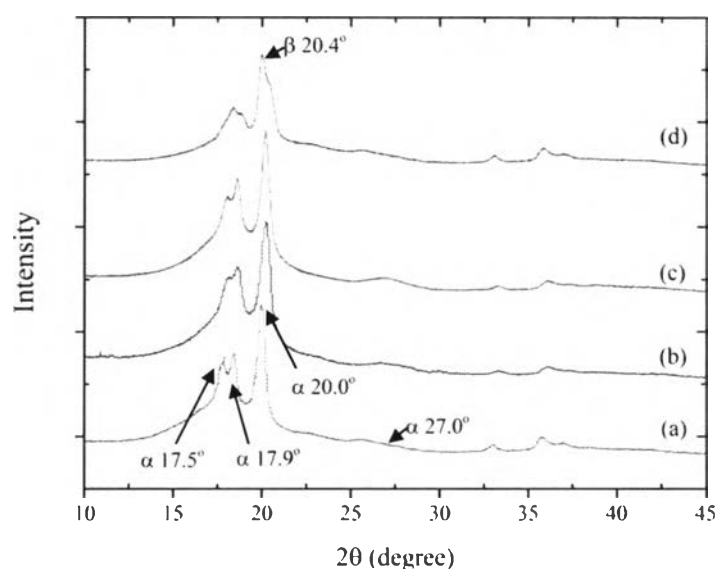


Figure 5.9 XRD patterns of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

5.4.2.3 Thermal Properties

In order to determine the thermal behavior of the neat PVDF-HFP and PVDF-HFP/BC composite film, DSC studies were carried out. The DSC parameters are shown in Table 5.3. From the result, there are transition temperature peaks observed at about 160 °C and 120 °C which corresponding to T_m and T_c of PVDF-HFP, respectively.

Figure 5.10 shows the second-heating curves of neat PVDF-HFP and its composite films. The inclusion of BC in polymer matrix shows slightly effects on T_m of PVDF-HFP matrix. The increase in the T_m value was due to the

higher thermal stability of BC and the interaction between oxygen atoms of BC with hydrogen atoms of PVDF-HFP, while fluorine atoms in PVDF-HFP may also be attracted to hydrogen atoms in BC [35].

Exothermic peaks of crystallinity temperature are showed in Figure 5.11 indicate that higher amount of BC shift T_c to higher temperature because the composite films need higher energy to crystallize with the presence of BC.

Furthermore, the degree of crystallinity (X_c) will be measured as the ratio between ΔH_m and ΔH_m^0 , as equation 3.1. As shown in Table 5.3, the increasing of BC causes slight decrement of X_c from 55.5 to 53.0. It can be described that the PVDF-HFP crystallization process may be disturbed by the micro structure of BC. Conclusively, although the BC induced the crystal transformation from α to β form, it reduced the crystallinity of PVDF-HFP [36-38].

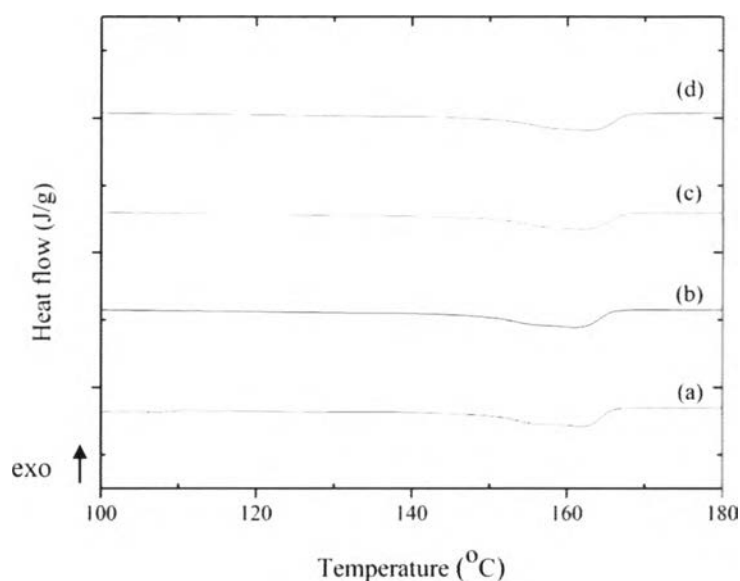


Figure 5.10 DSC second-heating curves of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

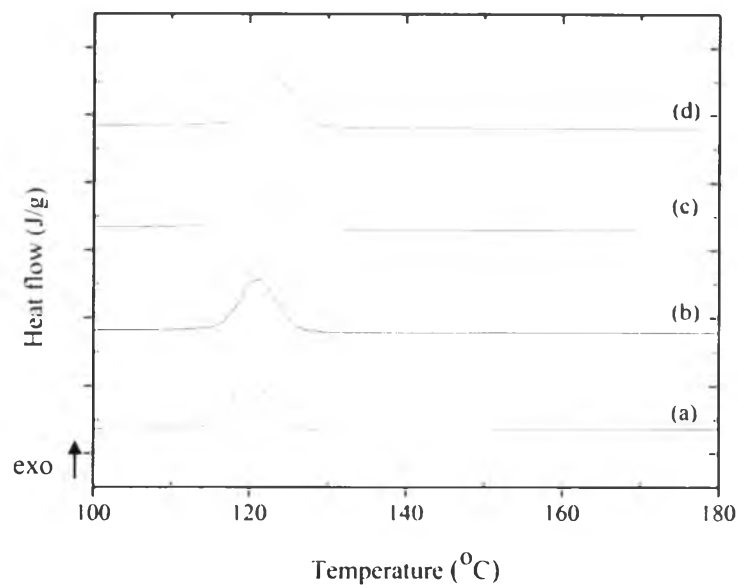


Figure 5.11 DSC first-cooling curves of (a) PVDF-HFP, (b) PVDF-HFP/BC (1 wt.%), (c) PVDF-HFP/BC (3 wt.%), and (d) PVDF-HFP/BC (5 wt.%).

Table 5.3 DSC parameters of neat PVDF-HFP and its composite films

Formula	T_m^a (°C)	T_c^b (°C)	ΔH_m^c (J/g)	X_c^d (%)
PVDF-HFP	161.6	120.0	57.5	55.5
PVDF-HFP/BC (1 wt.%)	159.3	121.0	59.1	57.0
PVDF-HFP/BC (3 wt.%)	162.0	122.0	56.8	54.8
PVDF-HFP/BC (5 wt.%)	162.6	122.3	55.0	53.0

a is melting peak temperature, *b* is cooling peak temperature, *c* is melting enthalpy, and *d* is Crystallinity of PVDF-HFP

Note: Crystallinity of PVDF-HFP was calculated by using the data on heating process.

Thermal degradation behavior was observed by using TGA. Figure 5.11 showed TGA thermograms of BC, neat PVDF-HFP, and its composite films. It was found that the thermal degradation behavior of the PVDF-HFP/BC

composite films was divided into two steps. The first step started at 300 °C, which corresponded to the degradation temperature of BC (see Figure 5.4) [31]. Where, the degradation of second step started at 420 °C, which corresponded to the degradation temperature of PVDF-HFP matrix [32]. The degradation temperature of PVDF-HFP matrix was not affected by the dispersion of BC. When temperature reaches 550 °C, the formation of char residue is presented. The degradation temperature and char residue are shown in table 5.4.

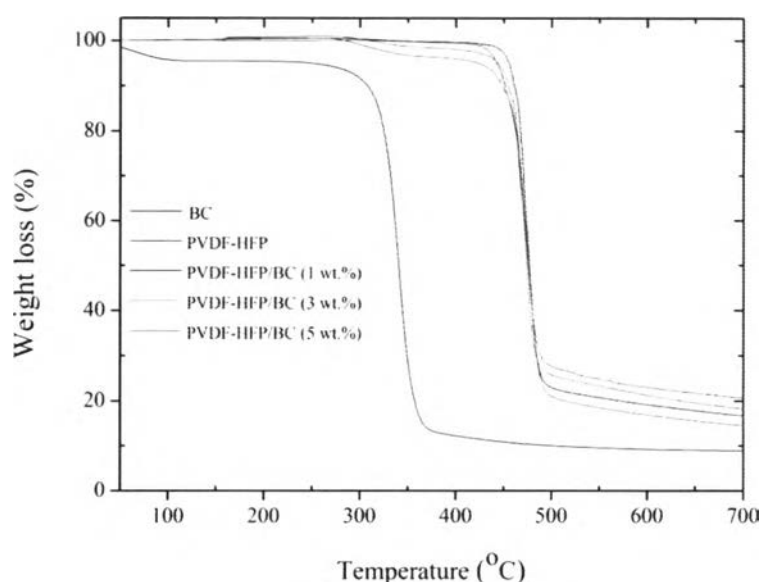


Figure 5.12 TGA thermograms of neat PVDF-HFP and its composite films.

Table 5.4 The degradation temperature (T_d) and char residue of PVDF-HFP and its composite films

Formula	T_{d1} (°C)		T_{d2} (°C)	
	onset	peak	onset	peak
PVDF-HFP	-	-	421	475
PVDF-HFP/BC (1 wt.%)	310		421	471
PVDF-HFP/BC (3 wt.%)	310	340	422	473
PVDF-HFP/BC (5 wt.%)	300	341	420	476

5.4.2.4 Dielectric Properties

The frequency dependence at 20°C of the dielectric constant at low and high frequency range of the films were shown in Figure 5.13(a) and 5.14(a), respectively. The decrease in dielectric constant with increasing frequency could be explained on the basis of dipole relaxation phenomenon. At low frequency, the dipoles in polymeric chain had sufficient time to align with the electrical field direction causing the higher dielectric constant and the inability of the electric dipoles to be in pace with the frequency of applied electric field at high frequency. It can be concluded that the dielectric behaviors of the PVDF-HFP and its composite film strongly depend on the variation of frequency. Dielectric constant of the composites slightly increases with increasing BC content. The highest dielectric constant value of PVDF-HFP composite was observed at 1 kHz when the content of BC was added 5 wt.%, the upper limit of the BC content in this study.

Figure 5.13(b) and 5.14(b) showed the frequency dependence of the dissipation factor at low and high frequency regions of all samples. It was observed that the dielectric loss increased as frequency increased for both neat PVDF-HFP and PVDF-HFP/BC composite films at low frequency (from 1 kHz to 1 MHz), but decreased as frequency increased at high frequency up to 1 GHz, the upper limit of the frequency range in this study. The maximum value of dielectric loss is reached at the frequency of 1 MHz.

Figure 5.15 exhibits the temperature dependence of the dielectric constant and dissipation factor was at 2-5 and 0.02-0.20, respectively at wide range of temperature and frequency range from 10 MHz to 1 GHz. At constant frequency with high temperature, the dielectric constant tended to be high due to polymer chains can be free to vibrate, move and reorient which allowing them to keep up with the changing electric field. When the successfully orientation was obtained, the dielectric constant became constant. With lower temperature, the segmental motion of the chain is practically frozen when the temperature was much lower than dynamic T_g which causing the lower dielectric constant [33]. Moreover, the result showed that dielectric constant was increase as increasing BC content which correspond to the result from previous paragraph. The dielectric relaxation mechanism of neat PVDF-HFP and composite films can be observed in Figure 5.15 (right), it was found

that there's only one relaxation mechanism appeared which was referred to the dynamic glass transition temperature (T_g) or the β relaxation that can be obtained by the dielectric response at very low frequency [34]. With the increasing of frequency (10 MHz to 100 MHz), this relaxation peak shifted to higher temperature (20 °C to 60 °C) because of polymeric chains in amorphous region has not enough energy to relax due to the very short time to store the energy, so the higher energy was needed for the chains to vibrate and reorient along the electrical field. For the PVDF-HFP/BC composite films, there's only one relaxation mechanism, which means that the incorporation of BC didn't have the effect on the relaxation behavior of the blend films.

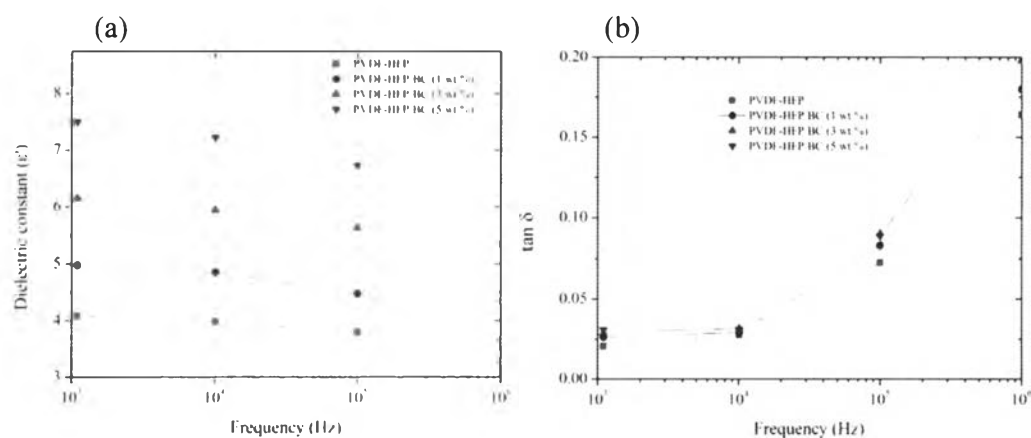


Figure 5.13 The frequency dependence of dielectric constant (a) and dissipation factor (b) of PVDF-HFP and its composite films at low frequency (1kHz-1MHz) ($T = 20$ °C)

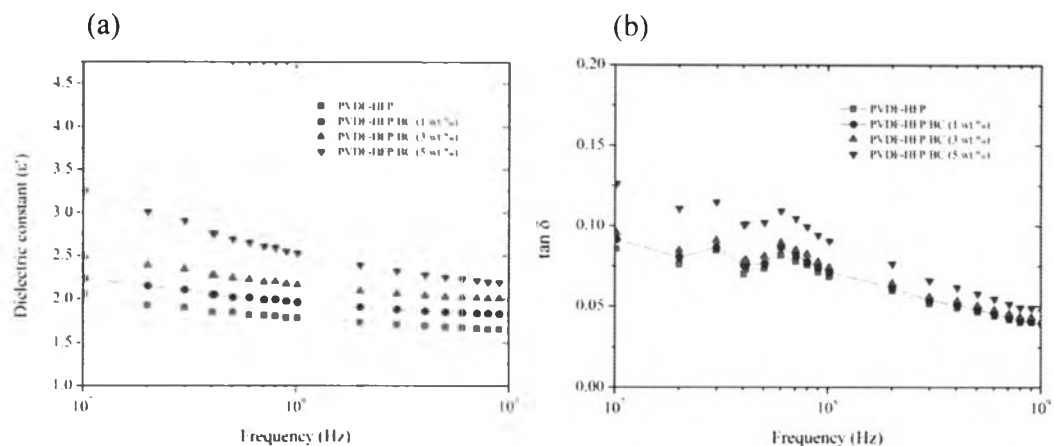
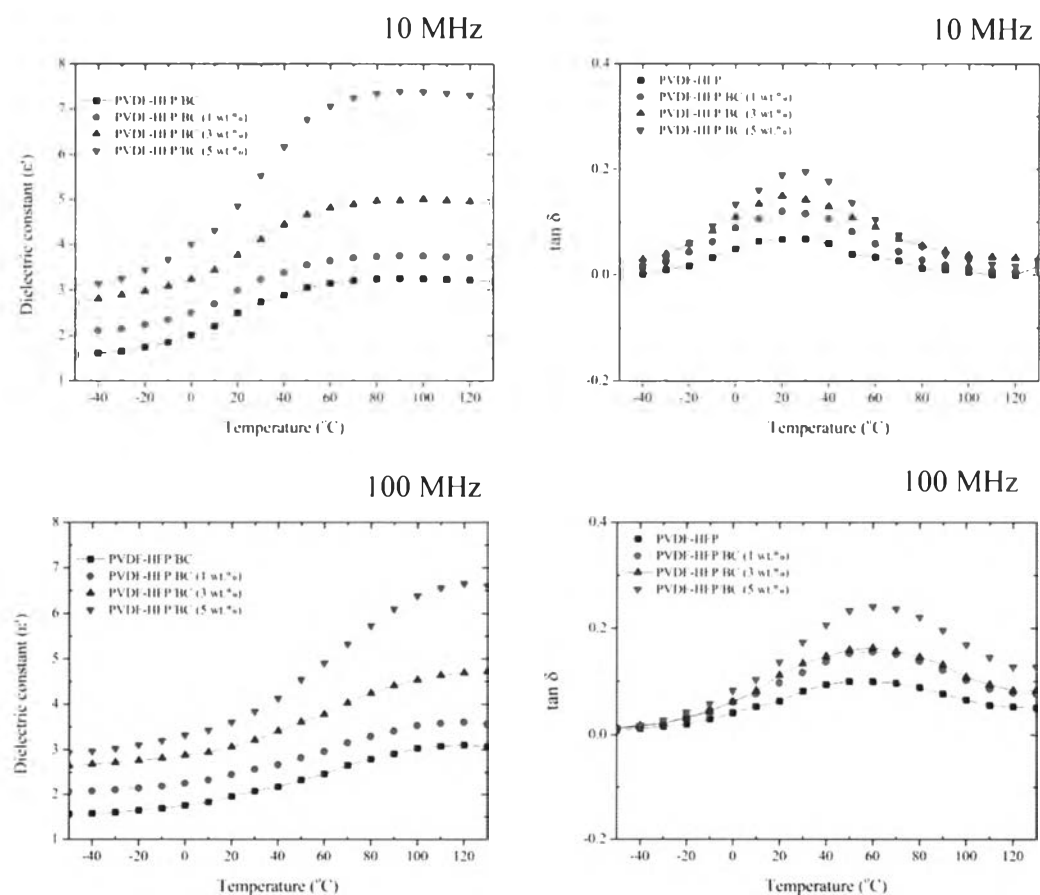


Figure 5.14 The frequency dependence of dielectric constant (a) and dissipation factor (b) of PVDF-HFP and its composite films at high frequency (10MHz-1GHz) ($T = 20^\circ\text{C}$).



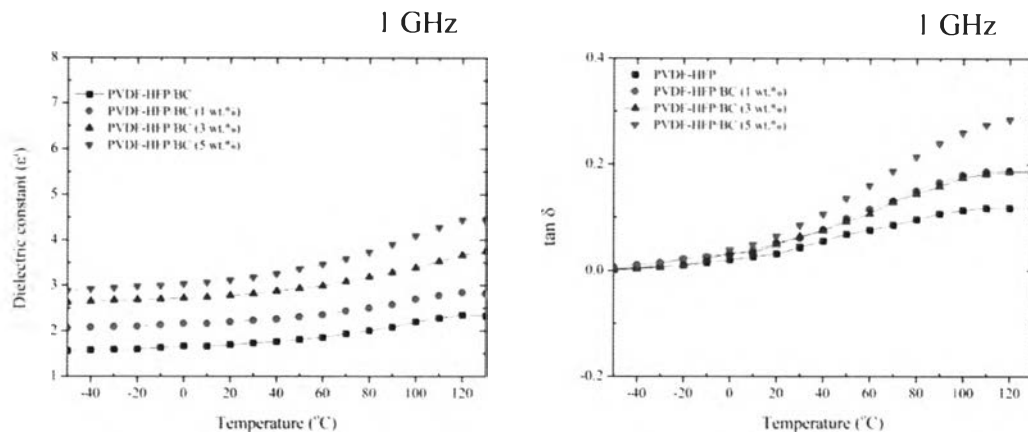


Figure 5.15 The temperature dependence of dielectric constant (left) and dissipation factor (right) of PVDF-HFP and its composite films (-50 $^{\circ}\text{C}$ - 130 $^{\circ}\text{C}$) at frequency 10 MHz, 100 MHz, and 1 GHz.

Figure 5.16, shown the dielectric constant and dissipation factor at 10 MHz as a function of BC content, it can be seen that incorporation with higher amount of BC exhibited significantly higher dielectric constant and dissipation factor. This increasing occurred due to the interfacial polarization between the hydroxyl group in the BC chain and fluorine atom in PVDF-HFP chain. Moreover, as the particle size is in micrometer order, the number of particles per unit volume is large, hence the dipole moment per unit volume increases, and also the dielectric constant increases [35]. However, higher amounts of BC, more free space occurred from repulsive force between the hydroxyl groups, and it consequently decreased the polarization, resulted in increasing on the dissipation factor. Moreover, a higher content of BC in the composite film caused higher water content, which increased the loss of the composite films due to lower stability of the PVDF-HFP structure and a higher dissipation factor from the water [30].

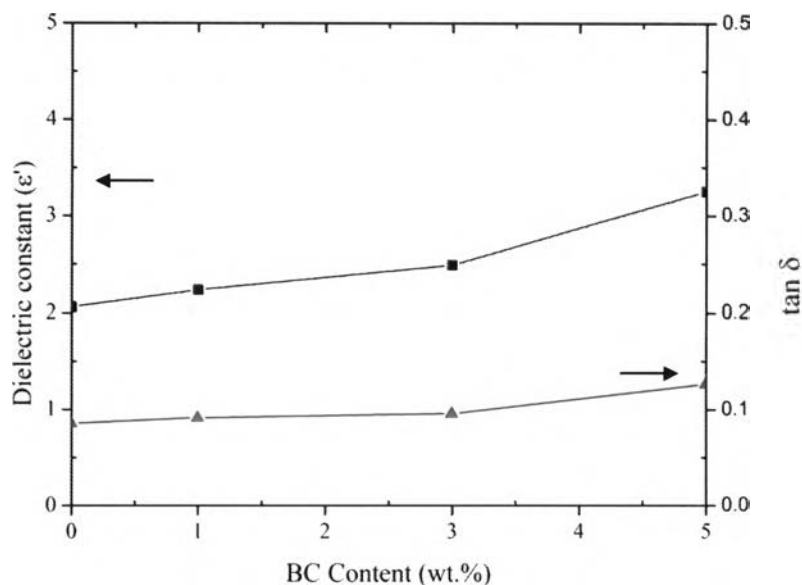


Figure 5.16 Dielectric constant and dissipation factor of PVDF-HFP and its composite films as a function of BC content measured at 10 MHz and 20 °C.

5.4.2.5 Mixture of dielectrics

Figure 5.17 shows the typical experimental and theoretical dielectric permittivities of the PVDF-HFP/BC composite films as a function of BC content; Maxwell (equation 3.6), and log (equation 3.7) mixing rule. It can be seen here that dielectric constant measurement was always greater than theories. From experimental, the dielectric constant increase with increasing BC content up to 5 wt.%. It was also observed that the results of the Maxwell and log mixing rule calculations were very different from the experimental results in this frequency range. These theories are based on a spherical morphology of the filler particles. Due to the non-spherical morphology of the BC (see Figure 5.1), the theoretical calculations of the dielectric constant were not completely in accord with the experimental results. In this case, the results showed that the Maxwell and log mixing rule were not suitable theory for calculating the dielectric constant of the PVDF-HFP/BC composite films which prepared by melt process.

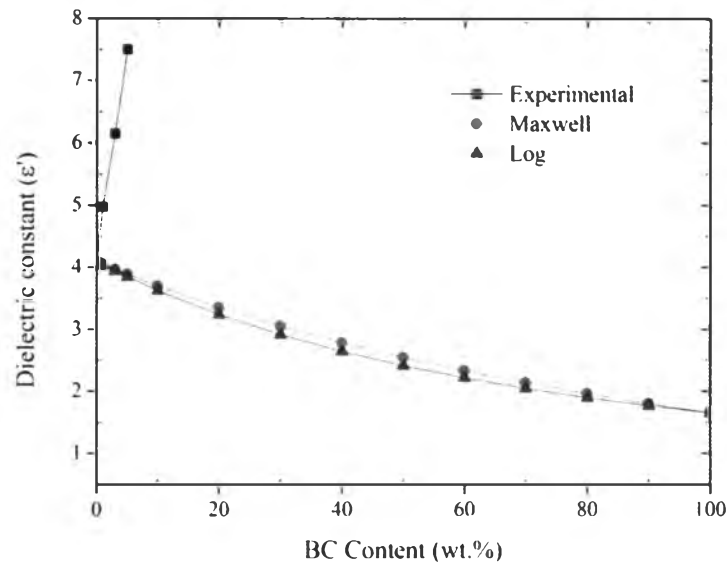


Figure 5.17 Dielectric constant of PVDF-HFP/BC composite obtained from experimental test and theoretical calculation (Maxwell, and Log mixing rule) at 10 MHz and 20°C.

5.4.2.6 Mechanical Properties

Touch sensor for used as piezoelectric touch screen require high mechanical properties for their long term operation and tolerance to the applied force during operation. Here, Young's modulus, tensile strength, and elongation at break were measured with universal testing and results are given in Figure 5.18, 5.19, and 5.20, respectively. Loading of BC was effective for improving the Young's modulus and Tensile strength of PVDF-HFP. The Young's modulus which measures the stiffness of the film was dramatically improved from initial 649 to 1151 MPa in transverse direction and from 696 to 877 MPa in machine direction for 5 wt.% BC content. This outstanding property may be attributed to the formation of a networked structure above percolation threshold which was produced by BC interactions through hydrogen bonding and attributed to the stiff nature of the BC dispersed [38]. Furthermore, the tensile strength of the composites also increases. The maximum strength was 39 MPa in transverse direction and 56 MPa in machine direction with 5 wt.% loading of BC. These results further confirm the presence of strong interfacial

of PVDF-HFP matrix is reduced when BC is added and the elongation drops from 285% to 209% in transverse direction when the BC content is 5 wt.%. On the other hand, there is no obvious decrease in the elongation at break in machine direction with increasing BC content because BC are good orientation in machine direction as observed in an optical microscope image.

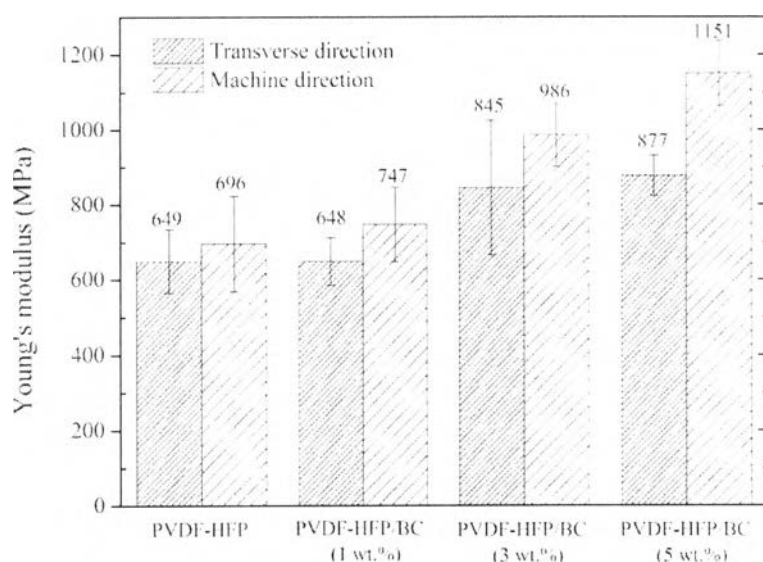


Figure 5.18 Young's modulus of neat PVDF-HFP and composite films.

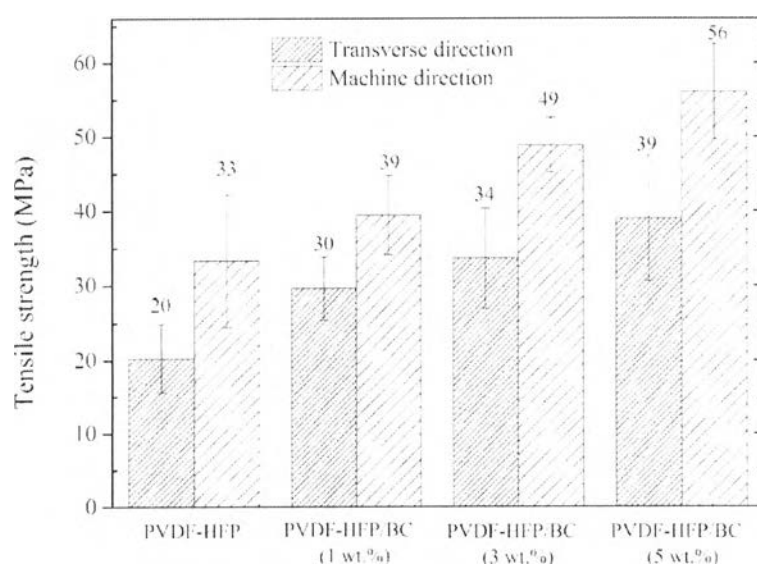


Figure 5.19 Tensile strength of neat PVDF-HFP and composite films.

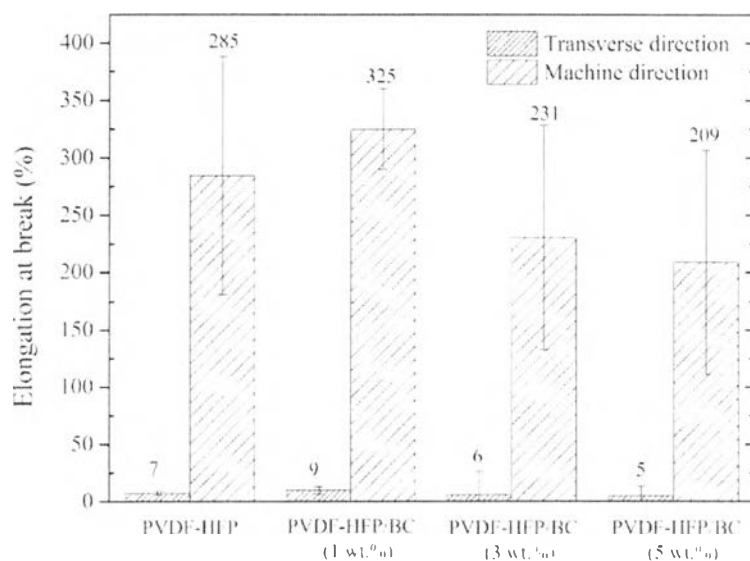


Figure 5.20 Elongation at break of neat PVDF-HFP and composite films.

5.4.2.7 Thermal shrinkage

Thermal shrinkage at high temperature causes the internal short circuit, measuring the thermal shrinkage is very important to evaluate the dimensional thermal stability of the touch sensor. The neat PVDF-HFP and PVDF-HFP/BC composite film were annealed at 130 °C for 2 hours to further compare their thermal stability at high temperatures. The dimensional changes of neat PVDF-HFP film in the machine direction is 5% and no shrinkage happens in the transverse direction. For high BC content (1-5 %wt.), no dimensional changes can be observed at 130 °C, indicating excellent dimensional thermal stability because the excellent dimensional stability of BC at high temperatures makes it promising for the application in high temperature.

5.5 Conclusion

The PVDF-HFP reinforced with bacterial cellulose (BC) at different weight percent, ranging from 1 to 5 wt.%, were successfully prepared by using melt mixing method follow by cast film process. The structure and morphology of the films were

studied using SEM. The existences of BC in the composite films were confirmed by the analysis of FTIR, TGA, DSC, and XRD. The PVDF-HFP/BC composite film had appealing feature of optical transparency, high thermal properties, and dielectric properties in term of dielectric constant, which corresponded to piezoelectric coefficient was improve from 2.0 to 3.25 with the addition of 5%wt, the upper BC content in this study. of BC. In addition, their Young's modulus and also tensile strength were dramatically increased comparing with neat PVDF-HFP to the property of BC. These features satisfied the criteria for the touch sensor application.

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5.7 Reference

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