Preparation and Characterization of Modified
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Abstract

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acrylonitrile-butadiene-styrene (ABS), acrylonitrile-styrene-acrylate (
butadiene-styrene (SBS) with different mass ratios. Mecha But yomene Butaniche stythe (ADS), act yiomene stythe act yiate (ADS), or stythe
butadiene-styrene (SBS) with different mass ratios. Mechanical tests indicated t
the appropriate blending could result in good compatibility, integrity and homogeneity, this side effectratio between different compositions was we
Key words: Polycarbonate, ABS, ASA, SBS, m
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1. Introduction

In the past decades, polycarbonate has become a widely used engineering plastic^[1]. Associated with advantages of mechanical strength, toughness, optical transparency, acid resistance, the polycarbonate has been widely used in many fields^[2]. For the purpose of improving its overall performances and overcoming the inherent shortages, polycarbonate was often blended with other resins^[3-7]. Many researches proved that styrene-based thermoplastic is one of the best mate for the blending among various candidates^[8-10]. It is also well established that the chemical compositions of the polymer blend would directly affect on its compatibility and overall performance in real applications, therefore choosing the appropriate components is the critical step in the blending.

In this work, three type of styrene thermoplastics, acrylonitrile-butadiene-styrene (ABS), acrylonitrile-styrene-acrylate (ASA), and styrene-butadiene-styrene (SBS) were selected to blend with polycarbonate with different mass ratios. The blends were prepared through extrusion and injection molding. The products were characterized of their mechanical and thermal properties for the performance evaluations.

2. Experiment

2.1 Granulation of polycarbonate blends

The polycarbonate (PC, Qimei Corp, Taiwan) was granulated acrylonitrile-butadiene-styrene (ABS, LG Corp, South Korea), acrylonitrile-styrene-acrylate (ASA, LG Corp), or styrene-butadiene-styrene (SBS, LG Corp) via extrusion molding by using the the twin screw extruder (SHJ-20, provided by Nanjing Giant Machinery Co.,Ltd., China). The mixture formula were shown in Table 1. The temperatures were well controlled and screened as : zone 1: 225°C, zone 2: 230°C, zone 3: 235°C, zone4: 240°C, extruder head: 240°C, polymer melt: 210°C.

Samples	Formula			
Pure polycarbonate	PC 100g			
PC/ABS 9010	PC: 90g, ABS: 10g			
PC/ABS 8020	PC: 80g, ABS: 20g			
PC/ABS 7030	PC: 70g, ABS: 30g			
PC/ABS 6040	PC: 60g, ABS: 40g			
PC/ASA 9010	PC: 90g, ASA: 10g			
PC/ASA 8020	PC: 80g, ASA: 20g			
PC/ASA 7030	PC: 70g, ASA: 30g			
PC/ASA 6040	PC: 60g, ASA: 40g			
PC/SBS 9010	PC: 90g, SBS: 10g			
PC/SBS 8020	PC: 80g, SBS: 20g			
PC/SBS 7030	PC: 70g, SBS: 30g			
PC/SBS 6040	PC: 60g, SBS: 40g			

Table 1. Formulation of polycarbonate blend

2.2 Preparation of polycarbonate blends standard test samples

The PC blends standard test samples were obtained via injection molding (TTI-130F2, Welltec Machinery Ltd, China)^[11]. The temperatures of different zones were set as a range of:

injection head: 250-255°C, zone 1: 230-235°C, zone 2: 235-240°C, zone 3: 240-245°C, zone 4: 245-250°C, mold: 85°C. The injection pressure was set as 10MPa, the holding pressure was set as 11.5MPa and the pressure was kept for 10 seconds.

2.3 Characterization of polycarbonate blends

Morphology investigation: a JEOL JSM-6700F scanning electron microscope was used to observe the surfaces of polycarbonate blends.

Mechanical properties tests: all polycarbonate blends were characterized of mechanical properties including tensile strength (GB/T1040.1-2006), Shore D hardness (GB/T2411-2008), flexural strength (GB/T9341-2008), notched impact strength (GB/T1043.1-2008, 7.5J of impact energy).

Thermal properties characterization: the differential scanning calorimetry (DSC, NETZSCH DSC204, Germany) was used to study the thermal properties of polycarbonate blends, the heating and cooling rate was precisely controlled as 20°C/min.

Thermal stability characterization: the thermogravimetric analysis were performed on polycarbonate blends to evaluate their thermal stabilities. In addition, the blends were measured of the Vicat (ZWK1302-B, MTS Systems Corporation) softening temperature (GB/T1633-2000, heating rate: 150°C/h).

3. Result and discussion

3.1 Morphology

The Fig.1 showed the SEM images of polycarbonate based samples. The polycarbonate blends, though were not as smooth as pure polycarbonate, displayed an homogeneous and integrated surface morphology without presenting component aggregation and phase separation.

Fig.1. SEM images of (a).Pure polycarbonate; (b). PC/ABS; (c). PC/ASA; (d). PC/SBS.

3.2 Mechanical properties

The mechanical properties of all polycarbonate blends were displayed in Table 2. For these blends, the mechanical properties were critically determined by the types and application amounts of styrene thermoplastics. As shown in the table, PC/ABS blends possessed a good overall mechanical properties, PC/ASA blends possessed the betterimpact strengths, and PC/SBS blends possessed the better tensile strengths and hardness. The differences of mechanical properties could be attributed the chemical structures of styrene thermoplastics. The ABS has the butadiene components that contains the C=C double bonds, when blending with PC, these double bonds could increase the flexibilities of the blends and thus improving the impact strength^[12]. The ASA has the acrylonitrile components instead of butadiane, the polar nitrile groups on acrylonitrile provide the blends with rigidities so the tensile strengths increased. For the SBS, besides flexible butadiene it also has high content of bulk styrene which result in high tensile strength and hardness. The mass ratios between each components were also significant, overdose of styrene thermoplastics would cause the decrease of mechanical properties of the blends. The reason was that the homogeneity of PC blend would be damaged if the amount of styrene thermoplastics was too high.

	Flexural	Tensile	Shore D	Notched impact
	strength/MPa	strength/MPa	hardness	strength/kJ*m ⁻²
Pure PC	62.20	61.30	75	26
PC/ABS 9010	60.60	53.90	74	38
PC/ABS 8020	54.50	56.70	73	41
PC/ABS 7030	55.60	54.80	73	45
PC/ABS 6040	53.70	56.20	72	40
PC/ASA 9010	60.40	53.10	74	29
PC/ASA 8020	53.70	52.90	73	32
PC/ASA 7030	53.00	53.10	71	33
PC/ASA 6040	50.70	49.90	70	32
PC/SBS 9010	64.00	61.90	75	29
PC/SBS 8020	66.30	62.70	77	34
PC/SBS 7030	67.40	63.50	78	38
PC/SBS 6040	66.90	63.50	78	36

Table 2. Mechanical properties of polycarbonate blends

3.3 Thermal properties

The DSC figure was displayed in Fig.2, the blends displayed two endothermic peaks, indicating two glass transition points of the blends, the higher one belonged to the PC components. The two glass transitions indicated that in blends, the two separated phases have been formed. In addition, the glass transition point of polycarbonate was found to be affected by the existence of styrene thermoplastics, that might be caused by the intermolecular forces between two components that affect on the polycarbonate segments movements.

Fig.2 DSC curves of samples

Fig.3 was the TGA analysis of samples. Compared to the pure polycarbonate, the thermal stabilities of PC blends decreased, the Vicat test (Table 3) also indicate that the PC blends became less thermal resistant. This was due to the damage of homogeneities. In addition, the PC/ABS and PC/SBS were less thermal stable than PC/ASA, that was because the PC/ABS and PC/SBS contained the butadiene, whose diene segments sensitive to thermal oxidation, under high temperature, the butadiene components were vulnerable and decomposed.

Fig.3 TGA curves of samples

	Vicat softening point/°C	
Polycarbonate	152	
PC/ABS 7030	140	
PC/ASA 7030	146	
PC/SBS 7030	142	

Table 3. Vicat softening temperatures

4. Conclusion
In this study, three types of poly
and PC/ASA) were prepared and chara
blends were significantly determined **nclusion**
In this study, three types of polycarbonate/styrene thermoplastics blends (PC/ABS, PC/SBS
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