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## Synthesis and Characterization of Phosphorous Containing NLO Polymers

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### Abstract

Two diol monomers 2,6-Bis (4-hydroxybenzylidene)cyclohexanone (BHBC) and 2,6-Bis (4-hydroxy-3-methoxy benzylidene) cyclohexanone (BHMBC) were used as monomer for preparing phosphorus NLO containing polyesters. The monomers BHBC and BHMBC were prepared respectively by refluxing 4-hydroxybenzaldehyde and 3-methoxy-4-hydroxy benzaldehyde with cyclohexanone. The polyesters were synthesized by interfacial polycondensation of photosensitive diols with Phenyl dichloro phosphate using hexadecyltrimethyl ammonium bromide (HDTMAB) as phase transfer catalyst. Copolymers were also prepared by incorporating Bisphenol-A in the polymer backbone. The synthesized monomers and polymers were characterized by UV-vis, FT-IR,  $^1\text{H}$ ,  $^{13}\text{C}$ ,  $^{31}\text{P}$ -NMR spectroscopic techniques. Thermal properties of the polymers were studied by thermogravimetric analysis and differential scanning Calorimetry under nitrogen atmosphere. The polymers show good SHG efficiency and thus it may be considered as a promising material for NLO application.

**Keywords:** Interfacial polycondensation; NLO; Phase transfer catalyst; Thermogravimetric analysis.

### 1. INTRODUCTION

Photosensitive polymers have gained remarkable interest and a great deal of attention in recent times because of their wide variety of applications such as printing plates, photoresists, stereolithography, photocurable coatings, photorecorders, energy exchange materials (Murugavel *et al.* 1997; Borden *et al.* 1978) and microlithography. They induce many changes in photochemical properties such as solubility, optical transparency, dielectric constant and refractive index (Devia *et al.* 1998). Hence, for these photosensitive polymers their inherent inflammability is one of the serious drawbacks. Simple halogenated or phosphorus containing compounds are commonly

used to repress flammability. But these flame retardant additives are susceptible for migration leading to inhomogeneity of their distribution besides getting exuded out of the system. Incorporation of halogen and phosphorus moieties into the backbone of these photosensitive polymers may be sought as a promising approach to overcome these problems. The present work, deals with the synthesis of photosensitive bisbenzylidene diol with  $\alpha,\beta$ -unsaturated ketone moiety in the main chain and polymerized using phenyl dichloro phosphate by interfacial polycondensation technique. Two copolymers containing bisphenol-A were also synthesized for comparing the thermal properties. Spectral, thermal, Non Linear Optical property studies were also carried out in this investigation.

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## 2. EXPERIMENTAL DETAILS

### 2.1 Materials

Vanillin (sdfine-India), p-hydroxybenzaldehyde (sdfine-India), Boron trifluoride diethyl etherate (Fluka), Cyclohexanone (E.Merck, India), Ethanol (Changshu Yangquan-China), Chloroform (E.Merck, India), Hexa decyl trimethyl ammonium bromide (HDTMAB) (Fluka), Phenyl dichloro phosphate (sigma Aldrich) and Bisphenol-A (sdfine-India) were used as received. The other reagents and solvents were purified by adopting the standard procedure (Moran *et al.* 1965).

### 2.3 Synthesis of monomers

#### 2.3.1 2,6 - Bis (4 - hydroxy - 3 - methoxy benzylidene) cyclohexanone (BHMBC)

A mixture of cyclohexanone (1 mmol) and 4-hydroxy-3-methoxy benzaldehyde (vanillin, 2 mmol) was dissolved in 25 ml of absolute ethanol and the solution refluxed for 4 h with three drops of boron trifluoride ethyl etherate. The reaction mixture was then cooled to 0 °C and the crystallized product filtered and washed with cold ethanol. Recrystallization from ethanol gave more than 75% yield of fine yellowish green crystals of the title compound. Melting point 179 °C [lit.179 °C] (Balaji *et al.* 2011). This monomer BHBC was prepared in a similar manner to BHMBC. Recrystallization from a 3:1 mixture of ethanol/water gave fine yellow crystals of the title monomer with 82% yield, m.p.278 °C.

### 2.4 Synthesis of Polymers

All the polymers were prepared by the interfacial polycondensation method using HDTMAB as a phase transfer catalyst (Kurtz *et al.* 1968). A typical procedure for the synthesis of polymer 1 is as follows: in a solution of BHMBC (1 mmol) in 20 ml of aqueous NaOH (1 N), HDTMAB (2 wt.% of the diol) was dissolved, then 20 ml chloroform solution of phenyl

dichloro phosphate (1 mmol), was added to this mixture with vigorous stirring at 20 °C. After 30 min of stirring the solid polymer obtained was filtered and washed with water and acetone. It was then dried to a constant weight *invacuo* at 50 °C, the yield was 80%. The other polymers and copolymers II-V were also prepared in similar manner. These polymers were soluble in chloroform, DMSO, THF, DMF, DMAC and insoluble in benzene, toluene and pentane.

### 2.5 Characterization

Infrared spectra from 4000 to 600 cm<sup>-1</sup> of solid samples of the synthesized monomers and polymers were obtained by KBr method using a PERKIN ELMER spectrophotometer. <sup>1</sup>H-NMR spectra was run on JEOLGSX 400 MHz NMR spectrophotometer at room temperature (RT) in d4-dimethylsulphoxide (DMSO) or CDCl<sub>3</sub> using TMS as an internal reference. The UV-visible spectra of the polymers were scanned on a SYSTRONICS 2202 spectrophotometer in DMSO or CHCl<sub>3</sub>. Thermo gravimetric analyses (TGA) of the polymers were examined in nitrogen atmosphere using a thermal analyzer (TA) PERKIN ELMER at a heating rate of 10 °C/min. Differential Scanning Calorimetry (DSC) was examined in nitrogen atmosphere using a PERKIN ELMER (thermal analyzer). Second harmonic generation test for the polymers was performed by the powder technique of Kurtz and Perry (Kurtz *et al.* 1968) using a pulsed Nd:YAG laser (Model: YG501C, λ = 1064 nm). Pulse energy of 4 mJ/pulse, pulse width of 10 ns and repetition rate of 10 Hz were used.

### 2.6 Non Linear Optical studies

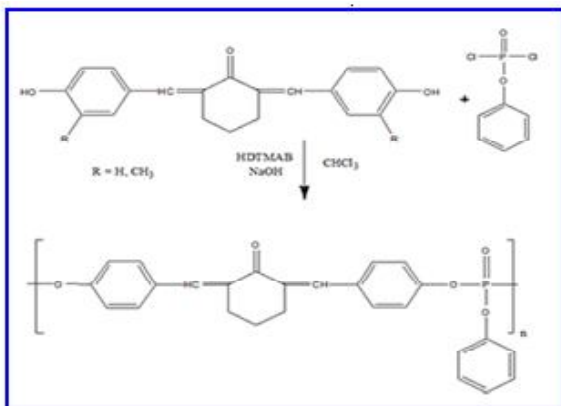
The presence of SHG in the synthesized polymer samples was confirmed by using the modified powder SHG test of Kurtz and Perry. The polymer samples were ground into powder particles of uniform size and packed in a glass container. A Q switched Nd:YAG laser is used as source for IR radiation of fundamental wavelength 1064 nm, 10 ns pulse width, 10 Hz pulse rate and 2.5 mJ energy per pulse is made

to fall on the sample uniformly. The SHG is confirmed by the emission of green radiation of wavelength 532 nm. The SHG efficiencies of the polymer samples were measured by keeping KDP and Urea as the reference material.

### 3. RESULTS & DISCUSSION

All the polymers were obtained by interfacial polycondensation method using a phase transfer catalyst at 20 °C with >75% yield (Scheme I). All the polymers were soluble in chloroform, tetrahydrofuran, dimethylformamide, dimethylsulfoxide and dimethylacetamide and insoluble in benzene, toluene, and n-hexane. The good solubility in polar organic solvents may be attributed to the introduction of phosphorous segments in the polymer backbone (Delaviz *et al.* 1993). The results are summarized in Table 1.

The infrared spectrum of polymer I showed the absorption band near 3063  $\text{cm}^{-1}$  corresponds to aromatic -CH stretching, 1665 and 1600 corresponds to cyclohexanone and exocyclic double bond stretching's respectively. All the polymers show absorption around 1300 and 960  $\text{cm}^{-1}$  corresponds to PP=O and P-O-C (aromatic) stretching's. These results support the formation of polyphosphate esters.



Scheme I

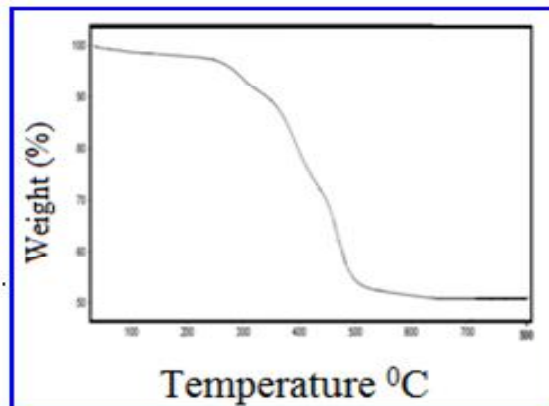


Fig. 1: TGA traces of polymer II

$^1\text{H-NMR}$  spectrum of polymer I showed the aromatic protons of the main chain and pendent phenyl ring as a broad multiplet in the region of 6.4 to 7.1  $\delta$ . The signals corresponding to Olefinic protons appear at 7.5  $\delta$ . The  $\beta$  and  $\gamma$  methylene protons of cyclohexanone appear at around 2.87  $\delta$  and 1.8  $\delta$  respectively. The  $-\text{CH}_3$  protons of bisphenol-A moiety in homo and copolymers appear in the range of 1.58-2.19  $\delta$ . These data strongly confirmed the formation of the polymers.

Table 1. Yield and solubility data of polymer II

Polymer	Yield %	Solvents							
		$\text{CHCl}_3$	DMSO	THF	DMF	DMAC	$\text{C}_6\text{H}_6$	Toluene	n-hexane
I	80	+	+	+	+	+	.	.	.
II	78+	+	+	+	+	.	.	.	.
III	76+	+	+	+	+	.	.	.	.
IV	82+	+	+	+	+	.	.	.	.
V	79	+	+	+	+	+	.	.	.

The proton decoupled spectra  $^{13}\text{C-NMR}$  spectrum of polymer I shown in Fig.2 the aromatic carbons of the main and side chains are in the region 115-150  $\delta$ . The methoxy carbon resonates at 56  $\delta$ . The thermal stability of the polymers was ascertained by



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