Synthesis, Characterization and Study of Thermally Stable Fluorescent Polyesters

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Abstract. Numerous polyesters containing heterocyclic ring have been synthesized by the polycondensation method. Ten Polyesters were synthesised having different aliphatic-aromatic diols in the chain having s-triazine ring as main moiety. The polyesters were synthesized by polycondensation of 6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carbonyl Chloride)-1,3,5-triazine [PCTC] with aliphatic and aromatic diols. Dark brown, light brown, golden and maroon colour showed by novel synthesized polyesters. The solubility of synthesized polyesters was observed in different solvents. The viscosity was studied for all the polyesters. This polyester showed inherent viscosity ranging from 0.402- 0.709 dl/g in DMF at 25°C. The synthesized polyesters were characterized by ¹H-NMR, FT-IR, fluorescence spectra. Fluorescent polymers are macromolecules with enormous important applications. The measured data indicates the synthesized aliphatic-aromatic fluorescent polyesters, focussing on synthetic methods, properties and applications. Due to their unique mechanical property and workability, their emerging applications involve the fields of fluorescent probe, smart polymer machines, chemo sensors, biological imaging, pH/temperature sensor, drug delivery and in many fields. Moreover high quality fluorescent polymers give widespread application in advanced bio imaging, ultrasensitive molecular diagnosis, and novel light emitting nano devices.

1. Introduction

The polymer chemistry is a multidisciplinary science that deals with the chemical synthesis and chemical properties of polymers. It is emerging as a necessary part of human life and it impacted on the system in which world exists. Amazing variety in the properties of polymeric materials exists [1,2]. Polymers are used in a broad variety of applications [3]. A great opportunity is there to become aware of these materials and expand their properties as polymeric material. Growing demands for extremely specialized materials for utilization with specific properties are there in today’s scenario. There is massive request for fluorescent polymers for past several decades. Organic fluorescent polymers have been extensively studied [4]. Research on fluorescent material has attracted great attentions due to wide range of applications such as clean energy technologies [5], explosive sensing, pH/temperature sensor, biological imaging [6], life science, material science, ultrasensitive molecular diagnosis and novel light-emitting nano devices [7], and biological detection schemes, smart polymer machines, fluorescent chemo sensor, fluorescent molecular thermometers, fluorescent imaging, drug delivery carriers, and so on [8,9]. Different methods have been used developed to synthesize fluorescent polymers by many research groups, with new architecture which contain various amounts of fluorophores and show bright fluorescence are synthesized for various applications [10,11]. Research on fluorophore-enriched polymers has become an interesting field because of their multipurpose applications as luminescent probes in various areas[12-15]. In general, there are two ways to synthesize such fluorophore-enriched polymers. Fluorescent polymers can be synthesized by polymerization of fluorescent functional monomers, using fluorescent compounds as initiator, fluorescent compounds as chain transfer agents, chemical bonding between fluorescent groups and polymers, and polymerization of non-fluorescent functional monomer[16-20], the second is the chemical variation of polymers by
fluorescent molecules or fluorescent oligomers [21,22]. The aim of present work was to synthesize polyesters with high fluorescence intensity for the coming use in different field.

2. Experiment work

2.1 Material and Methods

Freshly prepared double distilled water was used for the preparation of solutions. Chemicals such as Sodium hydroxide pallets, Sodium bicarbonate, Thionyl chloride, Cetramide were used as received. The solvents carbon tetrachloride, chloroform, methanol, ethanol, acetone, Benzene, n-butanol, ethylacetate, amyl acetate, isoamyl acetate, dimethyl formamide(DMF), dimethyl sulfoxide(DMSO) were received from Merck. Cyanuric chloridewas purified by repeated crystallization from pure benzene(m.p146°C). pipridine(BDH), Phenolphthalein (BDH LR),Catechol, Diethylene glycol, Ethylene glycol, Hydroquinone, Resorcinol, 1,4-dihydroxy Anthraquinone, 1,5-dihydroxy Anthraquinone, 1,8-dihydroxy Anthraquinone (Cibatul,Atul) were received and purified by recrystallization from rectified spirit.

2.2 Synthesis of Monomer PCT

2.2.1 Synthesis of 2, 4-dichloro-6-(Piperidinyl)-1, 3, 5-triazine [PT]

A solution of cyanuric Chloride (18.44g, 0.1mol) in a 60ml Acetone was added with stirring to a cooled solution (0-5°C) of sodium bicarbonate (10.6g) in 100ml of distilled water, in a three-necked flask (250ml) equipped with a mechanical stirrer. This resulted in the formation of slurry of cyanuric chloride. A solution of piperidine (10.2ml, 0.1mol) in 10ml of acetone was added to the cold slurry of cyanuric chloride. The mixture was stirred for 2 hour at 0-5°C. The white colored product was filtered, and recrystallized from ethanol and dried in vacuum desiccators. The yield was 82%. M. P. 120°C.

2.2.2 Synthesis of 6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carboxylic acid)-1,3,5-triazine [PCT]

A solution of 2,4-dichloro-6-(Piperidinyl)-1,3,5-triazine [PT] (23.3g, 0.1mol) in 40ml of acetone was mixed with solution of NaOH (16g, 0.4mol) and 7-Hydroxy-Coumarin-3-carboxylic acid (41.2g, 0.2mol) in 80ml double distilled water. Reaction was stirred for 2 hr at room temperature and 2 hr at 80°C. Separated solid was collected and washed with hot water. Product was dried in vacuum at 100°C. The yield was 86%. The product was recrystallized from acetone. M. P.153°C.

2.2.3 Synthesis of 6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carbonyl Chloride) 1,3,5-triazine [PCTC]

Thionyl chloride (11.9ml, 0.1mol) was added into 6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carboxylic acid)-1,3,5-triazine [PCT] (5.68g, 01mol) in a dry round bottom flask. The reaction mixture was refluxed at 78°C for 2 hr. At the end of the reaction, excess thionyl chloride was distilled and dry product was collected. The yield was about 78%. The product was recrystallized from dimethyl formamide. M. P. 218°C.

2.3 Synthesis of Polyesters

6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carbonyl Chloride)-1,3,5-triazine [PCTC] (0.01mol) in the minimum quantity of DMF (approx. 10ml) was stirred into round bottom flask, then initiator citrimide (0.25g) was added and heated up to 150°C. Then 1, 4 -Anthraquinone (0.02mol) was added. The reaction mixture was raised to 160-180°C and heated for 8 hrs. The reaction mixture was cooled and poured with constant stirring in 250ml of ice-cooled water. Solid was filtered, washed with hot water and dried. The yield was about 79%. The other polyesters from 6-(N-Piperidinyl)-2,4-bis-(7-Hydroxy-Coumarin-3-carbonyl Chloride)-1,3,5-triazine [PCTC] and various diols were synthesized by the similar method. PHE-1 to PHE-10 polymers are synthesized by using catechol, diethylene glycol, ethylene glycol, hydroquinone, phenolphthalein, resorcinol, bisphenol-A, 1,4antraquinone, 1,5antraquinone, 1,8 antraquinone respectively.
2.3.1 Properties of Polyesters
Polyesters prepared from coumarin and a variety of diols have exclusive colours. Coumarin and piperidine based polyesters show different colours. Most of the polyesters are brown in colour. PHE-1, PHE-3 were dark brown, PHE-2, PHE-6, PHE-7 were light brown, but PHE-4, PHE-5, PHE-8, PHE-9, and PHE-10 were pink, coffee, maroon, golden and black respectively.

2.3.2 Solubility
Relative solubility of various polyesters obtained from coumarin and piperidine in different solvents were carried out at room temperature and 50°C. The solubility of the polyesters in different solvents reveals that polyesters were insoluble in aliphatic chlorinated solvents like chloroform, carbon tetrachloride. It was also found that polyesters were insoluble in halogenated and non-halogenated aromatic compounds like chlorobenzene, benzene and ether. Polyesters were soluble in dimethyl formamide, dimethyl sulfoxide, tetra hydro furan, ether and ethyl acetate. Polyesters were soluble at higher temperature which is described in table-1. Polyesters were soluble in halogenated aromatic compounds like chlorobenzene, benzene and ether. Polyesters were soluble in dimethyl formamide, dimethyl sulfoxide, tetra hydro furan, ether and ethyl acetate. Polyesters were partly soluble at room temperature in methanol, ethanol, n-butanol, and isopropyl alcohol but some of polyesters are soluble at higher temperature which is described in table-1.

Table 1. Relative solubility of polyesters

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<th>Solvent</th>
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<th>PHE-3</th>
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<th>PHE-7</th>
<th>PHE-8</th>
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(+) = Soluble, (-) = Insoluble, (±) = Partly soluble

2.3.3 Viscosity
Intrinsic, reduced and inherent viscosities for all the polyesters at various concentrations were determined at 25 ± 0.1°C. The relative (n rel) and specific (n sp) viscosities, reduced and inherent viscosities were calculated from experimental data. Typical Huggins and Kraemer plots were used to obtain intrinsic viscosity for some of polyesters. Intrinsic, reduced and inherent viscosities along with Huggin’s and Kraemer’s constants for 1% solution are shown in figure 1.

The observations of intrinsic viscosities of Polyesters PHE 1-10 reveals that PHE-7 which contain Bisphenol –A as diol has the highest solution viscosity and which can be due to high molecular weight compound, whereas PHE-3 has the lowest intrinsic viscosity which contain aliphatic Ethylene Glycol. The range of viscosity in this series was 0.455 -0.615 g/dl. The polymers which contain aromatic diols have high viscosity and the aliphatic diols have low viscosity such as PHE-3 and PHE-2 which has diethylene glycol and ethylene glycol as diols. The viscosity of PHE-5 is less than PHE-7 but higher than the rest of the polyesters. The intrinsic viscosity of the polyesters obtained follows the sequence given here which is described in table 2.
2.3.4 Characterization

All the synthesised polyesters were characterised by visible absorption spectra, some representative IR spectra and NMR spectra. The visible absorption spectra of all synthesised polyesters have been recorded in DMF and observed in range of 550 to 380nm. The colour observed for each of the polyester was due to oscillation of electrons and the presence of substituents.

2.3.5 IR Spectra

IR spectra of polyester PHE-3 showed O-H stretching vibration of –OH group at 3445 cm\(^{-1}\), N-H stretching vibration of secondary amine at 3424 cm\(^{-1}\), C-H stretching vibration of -OCH\(_3\) group at 2960 cm\(^{-1}\), C-H stretching vibration of -CH\(_2\) group at 2842 cm\(^{-1}\), C=C stretching vibration of aromatic ring at 1607 cm\(^{-1}\).

IR spectra of polyester PHE-6 showed O-H stretching vibration of –OH group at 3438 cm\(^{-1}\), N-H stretching vibration of secondary amine at 3424 cm\(^{-1}\), C-H stretching vibration of -OCH\(_3\) group
at 2940 cm⁻¹, C-H stretching vibration of -CH₂- group at 2843 cm⁻¹, C=O stretching vibration of – COOR group at 1708 cm⁻¹.

IR spectra of polyester PHE-7 showed O-H stretching vibration of –OH group at 3291 cm⁻¹, N-H stretching vibration of secondary amine at 3424 cm⁻¹, C-H stretching vibration of -OCH₃ group at 2960 cm⁻¹, C-H stretching vibration of -CH₂- group at 2845 cm⁻¹, C=C stretching vibration of aromatic ring at 1608 cm⁻¹, C-N stretching vibration of tertiary amine group at 1533 cm⁻¹.

IR spectra of polyester PHE-9 showed O-H stretching vibration of –OH group at 3620 cm⁻¹, N-H stretching vibration of secondary amine at 3544 cm⁻¹, C-H stretching vibration of -OCH₃ group at 2440 cm⁻¹, C-H stretching vibration of -CH₂- group at 2843 cm⁻¹, C=O stretching vibration of – COOR group at 1708 cm⁻¹.

Examination of IR absorption frequencies reveals that the polyesters exhibit strong or moderately strong absorption at a frequency of 803cm⁻¹ and 1464-1511cm⁻¹ which may be attributed to out of plane and in-plane vibrations of s-triazine ring respectively, 1737cm⁻¹ and 1612cm⁻¹ which may be attributed to vibrations involving lactone and ester groups respectively.
2.3.6 1HNMR Spectra

In figure (6) and figure 7, 1H-NMR spectrum of polyesters PHE-3 and PHE-6 singlet appear at 3.94δ were of -NH-protons, the presence of –CH₂ at 3.67 and 3.69δ. The protons at 4.90δ, and 6.78-7.80 were of -OH proton and aromatic protons. The multiple signals of aromatic protons appear at 7.05-7.63δ, 6.66-7.08δ, 6.63-7.28δ, and 7.89-7.96δ, in PHE-3, PHE-6, PHE-7, and PHE-9 respectively. The shift at 0.87δ indicates the presence of –CH₂- group of Diethylene Glycol in figure (6). Chemical shifts at 3.63-3.69δ prove the presence of –CH₂- group in piperidine. A singlet of 1.55δ gives the conformation of –CH₃ group of Bisphenol-A in figure (6). In the same way, in figure (7), the presence of aromatic protons appears at 6.66-7.08 δ and the presence of –CH₂ at 3.7δ-3.8 δ in PHE-6. The protons at 6.8 δ, 6.7-6.9 δ proves the presence of aromatic protons in PHE-7 and PHE-9 in figure (8) and figure (9). Chemical shifts at 2.2-2.6δ, 2.5δ prove the presence of ester linkage in PHE-7 and PHE-9.
Figure 6. $^1$H-NMR spectrum of PHE-3

Figure 7. $^1$H-NMR spectrum of PHE-6
2.3.7 Thermogravimetric analysis:

The TGA data were used to determine the initial degradation temperature, degradation rate, residual weight at the end of degradation, and degradation kinetic parameters. The weight loss of polyesters during thermal degradation is a function of degradation rate and degradation time. At higher heating rate, although the degradation rate was higher, the time to reach a given temperature became shorter and the residual weight at a given temperature during thermal degradation may be higher. The remaining solid residue after complete pyrolysis was char and ash. TGA was used to determine the thermal degradation characteristics and kinetic parameters. An attempt is made to assess the thermal stability of some of the polyesters qualitatively based on visual obtained thermograms. \( T_0 \) and \( T_{10} \) are some of the main criteria of the thermal stability of the polyesters. The higher the \( T_0 \) and \( T_{10} \), the higher will be the heat stability of polyesters. TGA curve obtains at a scan rate of 10°C/min for polyesters.
Figure 10. TGA OF PHE-3

Figure 11. TGA OF PHE-6
The thermal properties of the PHE-3, PHE-6, PHE-7, and PHE-9 were evaluated by TGA at heating rate of 10°C/min under a nitrogen atmosphere. The 10% weight loss temperature of the aromatic polyesters in nitrogen was recorded at 300°C for PHE-3, PHE-6, PHE-7, and PHE-9. TGA thermogram of PHE-6 polyesters in figure (11) exhibited high thermal stability. The amount of residue of PHE-6 polyesters in a nitrogen atmosphere was more than 44% at 550°C. The polyesters PHE-3, PHE-7 and PHE-9 were degraded at 530°C, 510°C, 540°C respectively. The high char yields of these polyesters could be ascribed due to their high aromatic content. The series of polyesters showed higher thermal stability compared to the corresponding counterpart without coumarin moiety.
2.3.8 *Fluorescence Spectra:*

The fluorescence spectrums of synthesized polyesters PHE-3, PHE-6, PHE-7, PHE-9 and PHE-10 were coumarin based piperidine monomer with the Ethylene glycol, resorcinol, hydroquinone and 1,8 Anthraquinone as diols. Polyesters were dissolved in DMSO-$D_6$ and then excited at 190nm. Emission spectra were recorded between 110 and 800 nm. Upon excitation at 110 nm, the fluorescence spectrum of PHE-3, PHE-6, PHE-7, PHE-9 and PHE-10, indicates broaden emission range between 100-600nm and emission peak in between 290nm-300nm. The highest intensity peak was observed in figure (18) and the lowest intensity peak was observed in figure (14) due to aliphatic diol. The decrease of fluorescence intensity does not follow the first-order kinetics model. One can suggest that at least two processes are responsible for the decrease of fluorescence: (1) concentration of fluorescent decreases due to photo cyclization reaction, and (2) fluorescence photo quenching operates in this system via intermolecular mechanism of energy transfer in photo stationary state.

![Fluorescence spectrum of PHE-3](image)

*Figure 14. Fluorescence spectrum of PHE-3*
Figure 15. Fluorescence spectrum of PHE-6

Figure 16. Fluorescence spectrum of PHE-7
Figure 17. Fluorescence spectrum of PHE-9

Figure 18. Fluorescence spectrum of PHE-10
3. Result and Discussions

The resulting novel polyesters were obtained in good yields (80-95%), with variety of colours. PHE-1 and PHE-3 showed light brown colour, PHE-2 and PHE-6 showed pink colour, PHE-7 and PHE-8 were maroon PHE-9 and PHE-10 showed light yellow colour. All the novel polyesters were insoluble in benzene, chloroform, carbon tetrachloride and chlorobenzene but completely soluble in Ethyl acetate, DMF and DMSO and partly soluble in methanol, tetrahydrofuran, n-butanol, isopropyl alcohol and ether. The inherent viscosities of novel polyesters showed ranging between 0.402- 0.709 dLg⁻¹. The intrinsic viscosity of polyesters were given this sequence PHE-3<PHE-2<PHE-1<PHE-8<PHE-6<PHE-4<PHE-9<PHE-10<PHE-5<PHE-7. The highest intrinsic viscosity was of PHE-7 due to molecule structure of polyester in which bisphenol-A was used as diol. The thermal properties of different polyesters were evaluated by TGA at heating rate of 10°C/min under a nitrogen atmosphere. TGA thermogram in figure (11) polyesters exhibited high thermal stability and lowest thermal stability was observed in figure (12). The high char yields of these polyesters could be ascribed to their high aromatic content. The series of polyesters showed higher thermal stability compared to the corresponding counterpart without coumarin moiety. Hence, the presence of heterocyclic compounds in novel polyesters proves the higher thermal stability of synthesized polyesters. The fluorescence spectrums of synthesized polyesters were observed. The presence of heterocycles and aromatic diols showed the highest intensity peak in figure (18) (PHE-10) and due to aliphatic diol the lowest intensity peak was observed in figure (14) (PHE-3).

4. Conclusions

The data indicates the creation of fluorescence and thermally stable polyesters by polycondensation of coumarin probe carboxylic acid and aliphatic-aromatic diols. The resulting polyesters show outstanding colours, solubility in organic solvents and good thermal stability. All the synthesized polyesters show excellent fluorescence which can be used in fluorescent probe, advanced bio imaging, ultrasensitive molecular diagnosis, novel light emitting nano devices, chemosensors, biological imaging and drug delivery.

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References


