Synthesis and Characterization of Cholesteric Thermotropic Liquid Crystalline Polyesters Based on Isosorbide

Nayaku N. Chavan

Polymer Science and Engineering Division, National Chemical Laboratory, Pune, India.
Email: nn.chavan@ncl.res.in

Received July 23rd, 2011; revised August 20th, 2011; accepted September 27th, 2011.

ABSTRACT

Background: Generally main chain cholesteric thermotropic liquid crystalline polymers are prepared from chiral diacid or diol monomer. But these monomers are costly. Isosorbide is chiral cycloaliphatic diol accessible from renewable resources in the form of pure enantiomers. Thus it is used to synthesize main chain cholesteric thermotropic liquid crystalline polymers. Incorporation of phenyl hydroquinone into the backbone of the main chain frustrates chain packing, thus lowering the crystallinity and depressing the melting point below the degradation temperature, also improves the solubility due to disruption of packing and maintains the mechanical and thermal performance.

Results: Optical microscopy study reveals that more than 50% of isosorbide content with phenyl hydroquinone and terephthalic acid showed "yellow iridescent oily streaks" with a background of mosaic/marble texture. These are the typical textures of cholesteric liquid crystalline phase. Copolyesters based on phenyl hydroquinone, isosorbide and terephthalic acid are soluble in aprotic solvents like N,N-dimethylacetamide (DMAC), dimethyl sulfoxide (DMSO), N,N-dimethylformamide (DMF), and N-methyl-2-pyrrolidone (NMP). Solubility increases with the content of isosorbide percent. Thermal stability of all copolyesters was more than 300°C on the basis of 10 wt% loss.

Conclusions: It was found that main chain cholesteric thermotropic liquid crystalline polymers can be prepared from chiral cycloaliphatic diol such as isosorbide. Main chain cholesteric thermotropic liquid crystalline polyesters are prepared from phenyl hydroquinone, isosorbide and terephthalic acid showed thermal stability more than 300°C. Main chain cholesteric thermotropic liquid crystalline polymers are soluble in aprotic solvents like DMAC, DMSO, DMF and NMP.

Keywords: Liquid Crystalline Polymers, Thermotropic, Cholesteric, Differential Scanning Calorimetry, Optical Microscopy, Thermo Gravimetric Analysis, Wide Angle X-Ray Diffraction, Crystallinity, Inherent Viscosity, Solubility and Solution Polycondensation

1. Introduction

Cholesteric liquid crystalline polymers (ChLCPs) have ability to produce supramolecular structures characterized by a helical organization with certain identity period. This special organization is responsible for unique optical properties of cholesteric polymer liquid crystals such as selective reflection of light at different wavelengths, very high optical activity, high sensitivity of selective reflection of light to temperature variations and to the action of external electrical and magnetic fields [1-5]. The requirement of principal condition to develop a cholesteric phase is associated with the presence of an chiral (asymmetric) center in monomer which produces a helical supramolecular structure [6]. Cholesteric phase is characterized by circular dichroism: the light selective reflected by cholesteric liquid crystal is circularly polarized. The direction of circular polarization of the reflected beam coinsides with the direction of the twisting of cholesteric helix. The principal feature of cholesteric structure is associated with a strong rotation of polarization plane, which may be as high as 10⁴ deg·mm⁻¹. In conventional organic liquids this parameter is provided by only chirality of molecules and it does not exceed 10² deg·mm⁻¹. The helical ordering is a result of the frustration between molecular chirality and layer ordering and is unusual in the fact that the twist is manifested by a regular array of screw dislocations. High sensitivity of
the pitch of the helix changes with respect to temperature as well as the action of electrical and magnetic fields offers substantial advantages for practical applications of such ChLCPs are as thermometers and thermoindicators for techniques and medicine. To prepare ChLCPs different polymerization methods are applied: 1) homopolymerization of chiral mesogenic monomers; 2) copolymerization of nematogenic monomers with the chiral mesogenic monomers; 3) copolymerization of chiral mesogenic monomers with non-mesogenic monomers. The most advantageous procedure for the development of cholesteric mesophase is copolymerization of mesogenic and chiral monomers. This approach offers a convenient means for the preparation of the cholesteric copolymers and allows one to control phase and optical properties of cholesteric polymer by varying their composition [7-9]. Addition of flexible aliphatic spacer increases solubility but simultaneously it reduces the thermal and mechanical performance of polymers. According to structure property relationship study terephthalic acid is highly rigid diacid based monomer and it enhances performance of polymers due to para-para orientation. Hydroquinone is also highly rigid diol based monomer and it improves performance of polymers due to para-para orientation. Homopolymer of hydroquinone and terephthalic acid is insoluble due to highly integrated para-para orientation. Incorporation of aromatic unit such as phenyl group on the backbone of the main chain frustrates chain packing, thus lowering the crystallinity and depressing the melting point below the degradation temperature, also improves the solubility due to disruption of packing and maintains the mechanical and thermal performance [10].

Prime aim of the study was to synthesize high performance main chain cholesteric thermotropic liquid crystalline polyesters using an isosorbide as chiral cycloaliphatic diol. Isosorbide is an attractive building block because it is easily accessible from renewable resources in the form of pure enantiomers [11-15]. It is well known that aliphatic spacer reduces thermal and mechanical performance of polymers. Hence it is not advisable to use aliphatic spacer even though it improves solubility. Isosorbide structure contains two cycloaliphatic rings and ether bridges. It improves the rigidity as well as performance of polymers due to cycloaliphatic group. In our earlier communication isosorbide based main chain cholesteric thermotropic liquid crystalline polyesters were reported [5].

In this communication, we attempted to use an isosorbide as chiral building block to prepare main chain cholesteric thermotropic liquid crystalline polyesters. Thermal, spectral, mesomorphic and physical properties of polyesters were characterized by thermogravimetric analysis (TGA), $^1$H nuclear magnetic resonance ($^1$H NMR), polarizing optical microscopy (POM), wide angle X-ray diffraction (XRD), solubility and inherent viscosity. The influence of the concentration of the chiral monomer on cholesteric thermotropic liquid crystalline phase behavior of the polyesters is discussed.

2. Experimental

2.1. Materials

Phenyl hydroquinone purchased from Aldrich and was recrystallized from chloroform. Terephthalic acid received from Sisco and was used as received. Thionyl chloride procured from Loba and was distilled prior to use. Isosorbide received from Aldrich and was used as such. Benzyl triethyl ammonium chloride (BTEAC) purchased from Spectrochem and was used without purification. Analytical reagent grade sodium hydroxide received from Merck and was used as received. Dichloromethane procured from Merck and was distilled over calcium hydride. Dimethylacetamide (DMAC) purchased from Aldrich and was distilled under reduced pressure on calcium hydride.

2.2. Measurements

Inherent viscosities of copolyesters were measured at a concentration of 0.5 dL/g in dimethyl acetamide using an Ubbelhode viscometer at 30°C. $^1$H NMR spectra were recorded on a Bruker AC300 spectrometer at room temperature in deuterated solvents. Thermo gravimetric analyses (TGA) were carried out on Perkin Elmer TGA-7 model under N$_2$ atmosphere using heating rate of 10 °C/min. The optical characterization was performed on a Mettler polarizing optical microscope equipped with a FP2 Mettler hot stage, at magnification of 50X. Wide angle x-ray scattering (WAXS) curves were obtained by a Philips PW X-ray diffractometer equipped with 1830 generator and 1710 adjustment with CuKα radiation 1.54 Å.

2.3. Polymerization (Solution Method)

Into three necked 100 mL round bottom flask, equipped with high speed mechanical stirrer, ice bath etc., phenyl hydroquinone (1.0 g, 5.3 mmol) and sodium hydroxide solution (14 mL, 10.6 mmol) were added. The solution was stirred for 15 min at room temperature. BTEAC (20 mg) added and the solution was cooled to 5°C. To this solution 10 mL dichloromethane containing terephthaloyl chloride (1.076 g, 5.3 mmol) was added at once. The solution vigorously stirred at 2000 rpm for 30 min and polymeric solution was poured into hot water. Filtered the solution and washed with methanol for several times. Polymer obtained was dried under reduced pressure at
3. Results and Discussion

Chiral polymers have been attracting much more interest because of their chiroptical and stereospecific properties. The synthesis of cholesteric liquid crystalline polyesters was performed with sugar diol such as isosorbide an attractive chiral building block. Several isotropic polyesters of this cycloaliphatic diol were synthesized by other research groups [11-15].

Synthesis of cholesteric thermotropic liquid crystalline polyesters from phenyl hydroquinone, isosorbide and terephthalic acid is depicted in Scheme 1 and compositions used for the syntheses of copolyesters are listed in Table 1. Polycondensation was performed by interfacial polymerization method using BTEAC as a phase transfer catalyst at 5°C.

3.1. Solubility

Solubility of cholesteric liquid crystalline polymers based on phenyl-hydroquinone, isosorbide and terephthalic acid are reported in Table 2. All these cholesteric thermotropic liquid crystalline polyesters were insoluble in common organic solvent like chloroform due to high rigidity.

\[ \text{Scheme 1. Synthesis of cholesteric thermotropic liquid crystalline polyesters based on phenyl hydroquinone, isosorbide and terephthaloyl chloride.} \]

| Table 1. Compositions of copolyesters based on phenyl-hydroquinone, isosorbide and terephthaloyl chloride. |
|---------------------------------------------------------------|
| Polymer code | ph-Hq:Iso:TPCl (Compositions) | ph-Hq (mmol) | Isosorbide (mmol) | TPCl (mmol) | NaOH (mmol) | BTEAC (mg) |
|---------------|--------------------------------|-------------|------------------|-------------|-------------|------------|
| NNC-1         | 100:00:100                     | 5.300       | -                | 5.300       | 10.600      | 20         |
| NNC-2         | 85:15:100                      | 4.505       | 0.795            | 5.300       | 10.600      | 20         |
| NNC-3         | 70:30:100                      | 3.710       | 1.590            | 5.300       | 10.600      | 20         |
| NNC-4         | 50:50:100                      | 2.650       | 2.650            | 5.300       | 10.600      | 20         |
| NNC-5         | 35:65:100                      | 1.855       | 3.445            | 5.300       | 10.600      | 20         |
| NNC-6         | 20:80:100                      | 1.060       | 4.240            | 5.300       | 10.600      | 20         |
| NNC-7         | 00:100:100                     | -           | 5.300            | 5.300       | 10.600      | 20         |

ph-Hq = phenyl-Hydroquinone; Iso = Isosorbide; TPCl = Terephthaloyl chloride; BTEAC = Benzyl triethyl ammonium chloride; NaOH = Sodium hydroxide.

| Table 2. Solubility of liquid crystalline homo/copolymers based on phenyl-hydroquinone, isosorbide and terephthaloyl chloride. |
|---------------------------------------------------------------|
| Polymer code | CHCl₃ | DMAc | DMSO | DMF | NMP |
|---------------|-------|------|------|-----|-----|
| NNC-1         | -     | -    | -    | -   | -   |
| NNC-2         | -     | -    | -    | +   | +   |
| NNC-3         | -     | +    | -    | +   | +   |
| NNC-4         | -     | +    | +    | +   | +   |
| NNC-5         | -     | +    | +    | +   | +   |
| NNC-6         | -     | +    | +    | +   | +   |
| NNC-7         | -     | +    | +    | +   | +   |

+ = soluble at room temperature; - = insoluble at room temperature.
Synthesis and Characterization of Cholesteric Thermotropic Liquid Crystalline Polyesters Based on Isosorbide

Solubility of cholesteric thermotropic liquid crystalline copolyesters was tested in aprotic solvents like dimethyl acetamide (DMAC), dimethyl sulphoxide (DMSO), dimethyl formamide (DMF) and N-methyl-2-pyrrolidone (NMP). Solubility increases with isosorbide content [5, 16]. Homopolymer based on phenyl hydroquinone and terephthalic acid is insoluble in chloroform as well in aprotic solvents like DMAC, DMSO, DMF and NMP. It clearly indicates that introduction of phenyl ring on hydroquinone does not improve solubility in aprotic solvents also. Homopolymer of isosorbide and terephthalic acid is also insoluble in chloroform. It dictates that isosorbide contain cycloaliphatic ring which acts as rigid moiety. Incorporation of terephthalic acid increases rigidity due to para-para orientation and reduces solubility of homopolyester also.

3.2. Thermal Properties

The thermo gravimetric analyses (TGA) of cholesteric liquid crystalline copolymers based on phenyl-hydroquinone, isosorbide and terephthaloyl chloride are summarized in Table 3. Thermo gravimetric analyses (TGA) of cholesteric liquid crystalline copolyesters were determined in nitrogen atmosphere using 10°C/min heating rate. Higher the molar fraction of phenyl hydroquinone stabilizes cholesteric liquid crystalline phase and rises the isotropization temperature above 300°C. TGA thermogram of NNC-1 showed 5 wt% loss at 350°C and 10 wt% loss at 395°C. DTG curve of the same polyester showed no decomposition up to 375°C and maximum decomposition was at 509°C. TGA thermogram of NNC-2 showed maximum wt. loss at 536°C, but DTG peak was broad. The T<sub>max</sub> temperature was on higher side due to the broadness of DTG peak. All copolyesters showed two stage degradation. It clearly indicates that liquid crystalline phases are formed. Homopolyester based on isosorbide and terephthalic acid showed first stage degradation above 300°C due to degradation of ether linkages of isosorbide and second stage degradation above 400°C due to degradation of polymeric unit.

Thermal stability decreases with isosorbide content increase. It is due to two ether linkages are present in isosorbide unit. Bridging group like ether degrade faster than aromatic ring due to bond energy of C=C (145 kcal/mol) is higher than C-O linkage (78 kcal/mol). The thermal stability of all copolyesters was more than 300°C on the basis of 10% wt. loss. Even though 10 wt%. loss of homopolyester based on isosorbide and terephthalic acid (NNC-7) was at 299°C. It clearly indicates that homopolyester based on isosorbide and terephthalic acid is also highly thermally stable. Maximum decomposition temperature (T<sub>max</sub>) decreases with isosorbide content increase.

3.3. Inherent Viscosity

Inherent viscosities of liquid crystalline copolyesters were determined in dimethyl acetamide (DMAC) at 30°C using 0.5% concentration in Ubbelhode viscometer. Polyester NNC-1 and NNC-2 are insoluble in DMAC. Inherent viscosities of liquid crystalline copolyesters were in the range of 0.11 to 0.15 dL/g. It indicates that molecular weights of copolyesters were not high. It may be due to interfacial polycondensation method is not suitable for the polymerization of isosorbide with acid chloride in the presence of strong base like sodium hydroxide. During interfacial polycondensation, isosorbide containing ether linkages may not be stable and lead to imbalance the stoichiometry of diols to acid chloride ratio. Polycondensation of diols with acid chlorides may be carried out with weak base like pyridine, and solution polycondensation may be carried out at high temperature using toluene or o-dichlorobenzene as solvent [5].

| Polymer code | T<sub>5</sub> (°C) | T<sub>10</sub> (°C) | T<sub>max</sub> (°C) | η<sub>inh</sub> (dL/g) | Crystallinity (%) |
|--------------|-----------------|-----------------|----------------|----------------|-----------------|
| NNC-1        | 350             | 395             | 509            | b              | 31.17           |
| NNC-2        | 301             | 330             | 536 (broad peak) | b              | 37.81           |
| NNC-3        | 286             | 362             | 514            | 0.15           | 28.37           |
| NNC-4        | 303             | 326             | 545 (broad peak) | 0.12           | 38.75           |
| NNC-5        | 299             | 321             | 512 (broad peak) | 0.13           | 33.71           |
| NNC-6        | 344             | 369             | 464            | 0.12           | 33.44           |
| NNC-7        | 272             | 299             | 418            | 0.11           | 37.62           |

T<sub>5</sub> = 5 wt% loss, T<sub>10</sub> = 10% weight loss; T<sub>max</sub> = maximum decomposition; a = inherent viscosity determined in DMAC at 30°C; b = insoluble in DMAC.
3.4. $^1$H NMR Spectrum

$^1$H NMR spectrum of isosorbide was carried out in CDCl$_3$ and is presented in Figure 1. The protons of isosorbide are observed in 3.50 - 4.75 δ ppm range and hydroxyl protons are observed at 2.45 and 5.58 δ ppm. $^1$H NMR spectrum of cholesteric thermotropic liquid crystalline copolyester NNC-4 in deuterated DMSO based on phenyl hydroquinone:isosorbide:terephthaloyl chloride (50:50:100) is presented in Figure 2. It showed that the incorporation of isosorbide protons which are observed at 4.0 - 5.5 δ ppm range. The position of hydroxyl groups also shifted. It clearly indicates that the copolymer formation. The intensity of isosorbide peaks was low because inherent viscosities of these copolyesters were low.

3.5. Optical Microscopy

Higher concentrations of phenyl hydroquinone like 85% or 100% did not melt up to 350°C on polarizing microscope, hence photomicrographs were not performed. The most interesting textures of cholesteric phase observed for copolymer NNC-5 and NNC-6 and are presented in Figures 3-10. It clearly showed “yellow iridescent oily streaks” having a background of mosaic texture which is a special characteristic of cholesteric liquid crystalline phase.

Optical textures of copolyester NNC-5 at 245, 273, and 297°C were depicted in Figures 3-5 respectively. It showed “yellow iridescent streak” with background of mosaic texture, which is a typical texture of cholesteric phase. Optical texture of polymer NNC-5 at 297°C depicted in Figure 5. It showed “yellow broad and bright oily streak” with background of mosaic texture, which is also a typical texture of cholesteric phase. Copolyesters NNC-5 showed broad range of cholesteric liquid crystalline phase i.e. 245°C - 297°C. Optical texture of copolymer NNC-5 at 315°C was illustrated in Figure 6 and it showed “yellow broad blackish oily streak” with background of mosaic texture. It clearly indicates the initiation of degradation of cholesteric phase.

Optical textures of copolyester NNC-6 at 174°C, 184°C, 198°C, and 231°C are depicted in Figures 7-10. Optical texture of copolyester NNC-6 at 174°C showed “small yellow oily streak” having a background of marble texture indicates the initiation of formation of cholesteric phase. Optical textures of copolyester NNC-6 at

![Figure 1. $^1$H NMR spectrum of Isosorbide.](image-url)
184°C and 198°C showed “broad yellow oily streaks” with background of mosaic texture. It indicated that the broad range of cholesteric liquid crystalline phase i.e. 174°C - 231°C. Optical texture of polymer NNC-6 at 231°C is depicted in Figure 10. It showed blackish “yellow iridescent oily streaks” with background of marble texture, which is a typical character of initiation of degradation. Homopolyester of isosorbide with terephthalic acid did not show cholesteric liquid crystalline texture due to depression of the liquid crystalline orientation order. All the optical photomicrographs were performed on heating cycle of optical microscope using magnification 50×.

3.6. Crystallinity

Crystallinity percentage of cholesteric liquid crystalline
Synthesis and Characterization of Cholesteric Thermotropic Liquid Crystalline Polyesters Based on Isosorbide

Figure 5. Optical texture of copolyester NNC-5 at 297°C.

Figure 6. Optical texture of copolyester NNC-5 at 315°C.

Figure 7. Optical texture of copolyester NNC-6 at 174°C.

Figure 8. Optical texture of copolyester NNC-6 at 184°C.

Figure 9. Optical texture of copolyester NNC-6 at 198°C.

Figure 10. Optical texture of copolyester NNC-6 at 231°C.

copolyesters was in the range was 31.17 to 38.75. It shows that copolyesters are semicrystalline. The WAXD patterns did not show definite trend.

4. Conclusions

The present study allows the conclusion that the chole-
teric thermotropic liquid crystalline polyesters can be synthesized from isosorbide, phenyl hydroquinone and terephthalic acid. Isosorbide is an optically active and inexpensive cycloaliphatic diol. Due to its cycloaliphatic nature it helps to increase the performance thermotropic liquid crystalline polyesters. Homopolyesters of isosorbide and terephthalic acid is thermally stable up to 300°C. Copolyesters based on isosorbide, phenyl hydroquinone and terephthalic acid showed thermal stability more than 300°C on the basis of 10 wt% loss. Optical microscopy study reveals that more than 50% of isosorbide content with phenyl hydroquinone and terephthalic acid showed “yellow iridescent oily streaks” with a background of mosaic/marble texture. These are the typical textures of cholesteric liquid crystalline phase.

Interfacial polymerization method is not suitable to get high molecular weight polyesters, because isosorbide containing two ether linkages may not be stable with strong inorganic base like sodium hydroxide lead to imbalance the stoichiometry of diols to acid chloride ratio. High performance cholesteric liquid crystalline polyesters with high molecular weight will be synthesized from same monomers by solution polycondensation method using weak organic base.

REFERENCES

[1] V. Shibaev, A. Bobrovsky and N. Boiko, “Photoactive Liquid Crystalline Polymer Systems with Light-Controlle

[lable Structure and Optical Properties,” Progress in Polymer Science, Vol. 28, No. 5, 2003, pp. 729-836. do

io:10.1016/S0079-6700(02)00086-2

[2] S. Nakanishi and M. Ueda, “Synthesis of Novel Glass-Forming Liquid Crystals Containing Acrylic Acid Trimer Core Unit and Mesogenic Moiety, and Their Use in Cholesteric Reflection Films,” Polymer Journal, Vol. 39, No. 3, 2007, pp. 252-258. doi:10.1295/polymj.PJ2006115

[3] B. Zhang, J. Hu, B. Wang, D. Yao and H. Li, “Synthesis and Characterization of Side-Chain Cholesteric Elastomers Derived from an Isosorbide Cross Linking Agent,” Colloid & Polymer Science, Vol. 285, No. 15, 2007, pp. 1683-1690. doi:10.1007/s00396-007-1742-5

[4] J. Lub, W. P. M. Nijssen, R. T. Wegh, J. P. A. Vogels and A. Ferrer, “Synthesis and Properties of Photoisomerizable Derivatives of Isosorbide and Their Use in Cholesteric Filters,” Advanced Functional Materials, Vol. 15, No. 12, 2005, pp. 1961-1972. doi:10.1002/adfm.200500127

[5] S. A. Jadhav, R. R. Chougule, Y. A. Shinde and N. N. Chavan, “Synthesis and Characterization of Cholesteric Thermotropic Liquid Crystalline Polyesters,” Journal of Applied Polymer Science, Vol. 103, No. 2, 2007, pp. 1232-1237. doi:10.1002/app.24610

[6] H. Finklemann, H. Ringsdorf, W. Siol and J. H. Wendtorf, “Synthesis of Cholesteric Liquid Crystalline Polyesters,” Macromolecular Chemistry, Vol. 179, No. 3, 1978, pp. 829-832. doi:10.1002/macp.1978.021790326

[7] J. Asrar, J. Toriumi, J. Watanabe, W. R. Krigbaum, A. Ciferri and J. Preston, “Thermotropic Homopolymers. I. The Preparation and Properties of Polymers Based on 4,4’-Dihydroxybiphenyl,” Journal of Polymer Science Part B: Physics Edition, Vol. 21, No. 7, 1983, pp. 1119-1131. doi:10.1002/pol.1983.180210712

[8] K. Fujishiro and R. W. Lenz, “Main-Chain Cholesteric Liquid Crystalline Polyesters with Chiral Pendant Groups. I. Model Compounds and Polyesters Containing a Chiral Substituted Hydroquinone,” Macromolecules, Vol. 25, No. 1, 1992, pp. 81-87. doi:10.1021/ma00027a014

[9] H. Park, J. Jin and R. W. Lenz, “Liquid Crystal Polymers: 19. Cholesteric Main Chain Polyesters with Triad Aromatic Mesogenic Units and Chiral Polyalkylane Spacers,” Polymer, Vol. 26, No. 9, 1985, pp. 1301-1306. doi:10.1016/0032-3861(85)90303-9

[10] N. Khan, Z. Bashir and D. M. Price, “Liquid Crystalline Aromatic Polymers Formed with Terephthalic Acid, Phenyl Hydroquinone, and Naphthalene or Anathracene Diols,” Journal of Applied Polymer Science, Vol. 58, No. 9, 1995, pp. 1509-1516. doi:10.1002/app.1995.070580913

[11] H. R. Kircheldorf and N. Probst, “Liquid-Crystalline Polyamides. 16. Chiral Thermotropic Copoly(Ester-Imide)s Based on Isosorbide and N-(4-Carboxyphenyl) trimellitimide,” Macromolecular Rapid Communications, Vol. 16, No. 4, 1995, pp. 231-237. doi:10.1002/marc.1995.030160401

[12] G. Schwarz and H. R. Kircheldorf, “New Polymer Synthesis. LXXXIII. Synthesis of Chiral and Cholesteric Polyesters from Silylated ‘Sugar Diols’,” Journal of Polymer Science Part A: Polymer Chemistry Edition, Vol. 34, No. 4, 1996, pp. 603-611. doi:10.1002/(SICI)1099-0518(199603)34:4<603::AID-POLA6>3.0.CO;2-R

[13] H. R. Kircheldorf, S. Sun, A. Gerken and T. Chang, “Polymers of Carbonic Acid. 22. Cholesteric Polycarbonates Derived from (S)-(2-Methylbutyl)hydroquinone or Isosorbide,” Macromolecules, Vol. 29, No. 25, 1996, pp. 8077-8082. doi:10.1021/ma960494d

[14] Q. Lin, J. Pasatta and T. E. Long, “Synthesis and Characterization of Chiral Liquid-Crystalline Polyesters Containing Sugar-Based Diols via Melt Polymerization,” Journal of Polymer Science Part A: Polymer Chemistry Edition, Vol. 41, No. 16, 2003, pp. 2512-2520. doi:10.1002/pola.10787

[15] B. Zhang, Y. Zheng and H. Lu, “Synthesis and Characterization of Side Chain Cholesteric Liquid Crystalline Polyesters Containing Isosorbide as a Chiral Center,” Liquid Crystals, Vol. 32, No. 3, 2005, pp. 357-364. doi:10.1080/0267829050034065

[16] M. Flugel and W. Heitz, “Synthesis and Properties of Polyamides Based on (+)-Nonbornane-2-Endo, 3 Exo-Di-carboxylic Acid,” Macromolecular Rapid Communications, Vol. 18, No. 6, 1997, pp. 523-528. doi:10.1002/marc.1997.030180611