



## Molecular Crystals and Liquid Crystals Science and Technology. Section A. Molecular Crystals and Liquid Crystals

Publication details, including instructions for authors and subscription information:

<http://www.tandfonline.com/loi/gmcl19>

## Synthesis and Physical Properties of a Novel Difluoroterphenyl Ferroelectric Liquid Crystal with a Fluorine at the Chiral Centre

N. Gough<sup>a</sup>, D. D. Parghi<sup>a</sup>, R. A. Lewis<sup>a</sup> & M. Hird<sup>a</sup>

<sup>a</sup> Department of Chemistry, University of Hull, Cottingham Road, Hull, HU6 7RX, UK

Version of record first published: 24 Sep 2006

To cite this article: N. Gough, D. D. Parghi, R. A. Lewis & M. Hird (1999): Synthesis and Physical Properties of a Novel Difluoroterphenyl Ferroelectric Liquid Crystal with a Fluorine at the Chiral Centre, *Molecular Crystals and Liquid Crystals Science and Technology. Section A. Molecular Crystals and Liquid Crystals*, 332:1, 347-354

To link to this article: <http://dx.doi.org/10.1080/10587259908023778>

PLEASE SCROLL DOWN FOR ARTICLE

Full terms and conditions of use: <http://www.tandfonline.com/page/terms-and-conditions>

This article may be used for research, teaching, and private study purposes. Any substantial or systematic reproduction, redistribution, reselling, loan, sub-licensing, systematic supply, or distribution in any form to anyone is expressly forbidden.

The publisher does not give any warranty express or implied or make any representation that the contents will be complete or accurate or up to date. The accuracy of any instructions, formulae, and drug doses should be independently verified with primary sources. The publisher shall not be liable for any loss, actions, claims, proceedings, demand, or costs or damages whatsoever or howsoever caused arising directly or indirectly in connection with or arising out of the use of this material.

# Synthesis and Physical Properties of a Novel Difluoroterphenyl Ferroelectric Liquid Crystal with a Fluorine at the Chiral Centre

N. GOUGH, D. D. PARGHI, R. A. LEWIS and M. HIRD

*Department of Chemistry, University of Hull, Cottingham Road,  
Hull HU6 7RX, UK*

This paper details the synthesis and mesomorphic properties of a novel SmC\*-forming liquid crystalline compound with potential applications in fast-switching multi-component ferroelectric mixtures. The material (NG158) possesses a difluoro-substituted *p*-terphenyl core and a fluorine at the chiral centre. The spontaneous polarisation at frequencies of 20Hz and 50Hz, tilt angles and switching speeds, are also reported.

**Keywords:** Ferroelectric; mixtures; fluorine

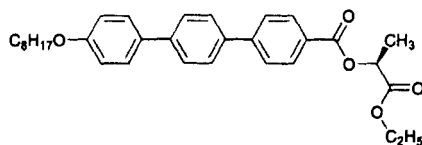
## INTRODUCTION

Liquid crystals possessing the ferroelectric SmC\* phase are being increasingly accepted as the materials of choice for the next generation of fast electro-optic devices<sup>[1]</sup>. These materials possess a number of attractive properties that make them particularly suitable for full-colour high-resolution display devices (fast response times, bistability, good contrast and excellent viewing angles)<sup>[2]</sup>.

By far the most practical technique for obtaining materials with the most suitable properties, for use in any twisted nematic-based device, is the combining of individual liquid crystalline components to form mixtures<sup>[3]</sup>. In order to exploit the ferroelectric SmC\* phase in a similar manner<sup>[4]</sup>, individual materials with wide SmC\* phases, good switching characteristics, low viscosities and a reasonably large spontaneous polarization are required.

Tailoring these mixtures to further enhance these properties, and moreover lower their viscosities, can be achieved by replacing individual components in present mixtures with those that have the more preferred properties.

Derivatives based on the *p*-terphenyl core, such as the material shown in Figure 1, have been widely used in both multi-component chiral and chiral-dopant mixtures<sup>[5]</sup>.



K 127 SmC\* 158.5 SmA\* 180.2 I (°C)

FIGURE 1 An example of a *p*-terphenyl-based material used in ferroelectric mixtures

The aim of this work was to synthesize a novel liquid crystal NG158 (Figure 2), with a difluoro-substituted *p*-terphenyl core and a fluorine at the chiral centre, and to determine its suitability as a component for ferroelectric mixtures for use in electro-optic devices. The material is structurally tailored as a co-dopant for use with the low viscosity difluoroterphenyl hosts favoured for  $\tau_{Vmin}$  FLC mixtures<sup>[6]</sup>. It was postulated that the fluorine at the chiral centre would give the material a reasonably high spontaneous polarization. Tilt angles and switching speeds were also measured.

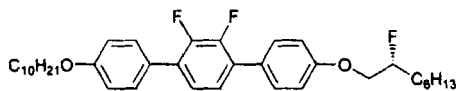
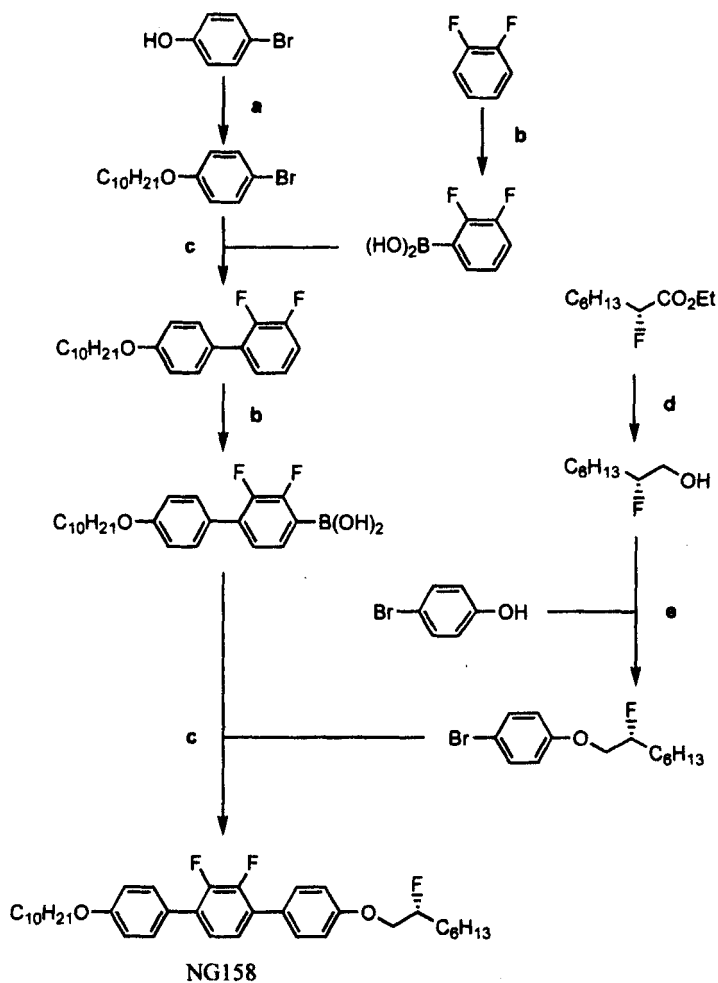


FIGURE 2 NG158

## RESULTS AND DISCUSSION

### Synthesis

The compound was prepared according to the synthesis outlined in Scheme 1.



### Reactions and reagents used in the preparation of NG158:

a... $\text{CH}_3(\text{CH}_2)_9\text{Br}$ ,  $\text{Na}_2\text{CO}_3$ , acetone.

b...i)  $n\text{BuLi}$ , THF,  $-76^\circ\text{C}$ , ii)  $\text{B}(\text{OMe})_3$ ,  $-76^\circ\text{C}$ , iii) 10% HCl.

c... $\text{Pd}(\text{PPh}_3)_4$ , 2M  $\text{Na}_2\text{CO}_3$ , MTBE.

d... $\text{LiAlH}_4$ , THF.

e...DEAD,  $\text{PPh}_3$ ,  $\text{Et}_2\text{O}$ .

SCHEME 1

### **Mesomorphic Properties**

The transition temperatures of the material were determined by optical microscopy and verified by DSC. The phase sequence of the material is as follows:-

K 88.6 SmC\* 144.1 SmA\* 154.9 N\* 157.3 BPI 157.8 I (°C)

The material possesses the greater part of preferred phase sequence on cooling for alignment in a display device (N\* - SmA\* - SmC\*). No ordered phases (SmI\*, SmF\* etc.) were observed in the material; this was more-than-likely due to the inhibiting steric effect of the lateral fluorines on the central phenyl ring.

The photographs below (Figures 3 and 4) show two of the textures observed on cooling from the isotropic liquid;

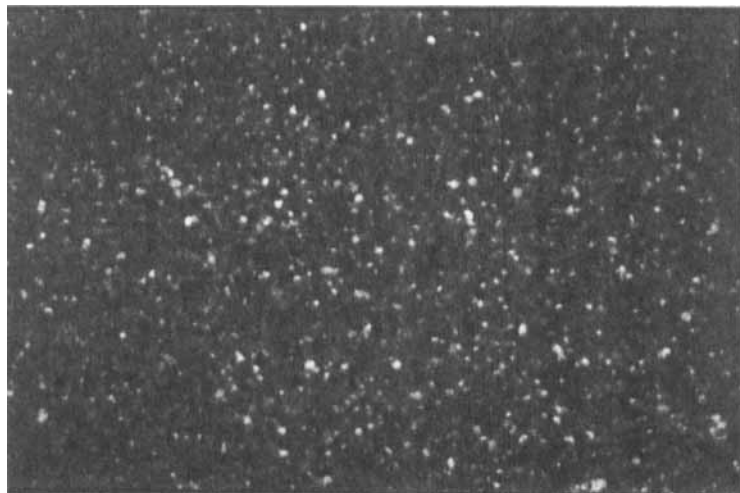


FIGURE 3 Blue Phase I (157.5°C)

(See Color Plate X at the back of this issue)

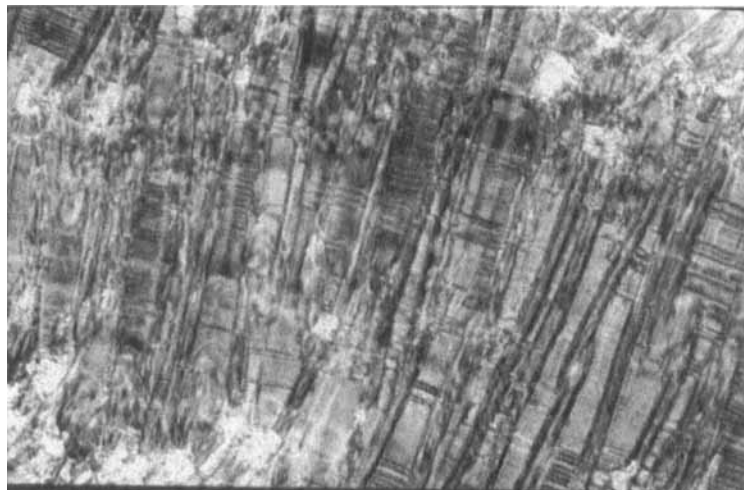


FIGURE 4 Smectic C\* (120°C)

(See Color Plate XI at the back of this issue)

### **Physical Measurements**

The spontaneous polarization ( $P_s$ ) of the compound was measured at switching frequencies of 20Hz and 50Hz under an applied triangular-wave AC field of  $\pm 50V$  ( $\pm 10V\mu m^{-1}$ ). The cells used were standard Linkam cells ( $5\mu m$  thick,  $0.81cm^2$  area, ITO-coated, antiparallel-aligned). The change in observed tilt angles of the material, from the temperature at which the  $P_s$  is detected (on cooling from the  $SmA^*$  phase), is also shown in Figure 5.

The material showed excellent alignment on cooling from the  $N^*$  phase to  $SmA^*$  phase under an applied square-wave AC field of  $\pm 90V$  ( $\pm 18V\mu m^{-1}$ ) at switching frequency of 100Hz.

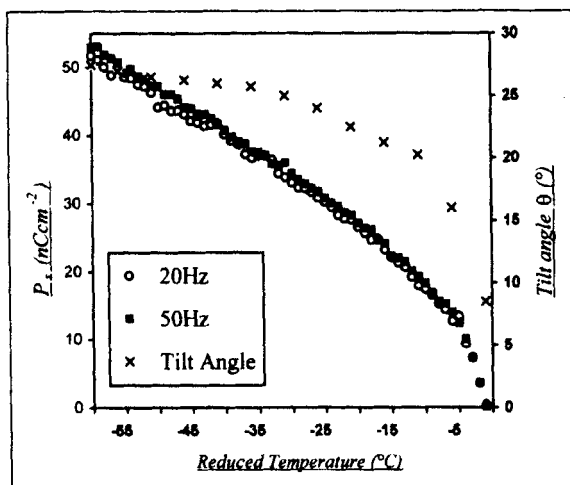


FIGURE 5  $P_s$  (at 20Hz and 50Hz) and observed tilt angle of NG158

The material has a maximum  $P_s$  of approximately  $50 \text{ nC cm}^{-2}$ , and a maximum tilt angle of  $28.5^\circ$ . The value of the  $P_s$  increases with observed tilt angle although the material showed a tilt angle of approximately  $10^\circ$  before the  $P_s$  was detected. This was due to electroclinic (soft-mode) switching in the  $\text{SmA}^*$  phase. The  $P_s$  measured at 50Hz is slightly higher than that measured at 20Hz; this may be due to increased saturated switching taking place at 50Hz in comparison to 20Hz.

Figure 6 shows the triangular-wave response of the material at  $90^\circ\text{C}$ , at a switching frequency of 20Hz. The first plot is the input triangular-wave, the second is the detected electrical response from the cell; ferroelectric switching is clearly visible.



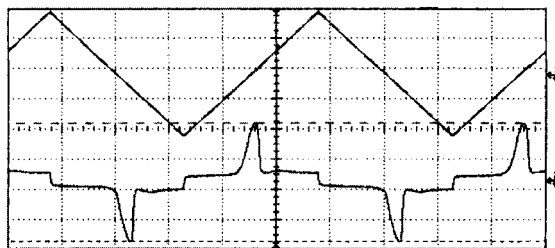


FIGURE 6 Triangular-wave response at 90° C ( $\pm 50$ V at 20Hz)

The energy-minimised molecular model below (Figure 7) shows the fluorine at the chiral centre forms an angle of approximately 90° with the lateral fluorines on the central phenyl ring. The combination of the molecular dipole at the chiral centre due to the fluorine, with the lateral dipole associated with the di-fluorinated core may account for a reasonable value for the  $P_s$  of the material, given that the material does not possess any ester linkages.

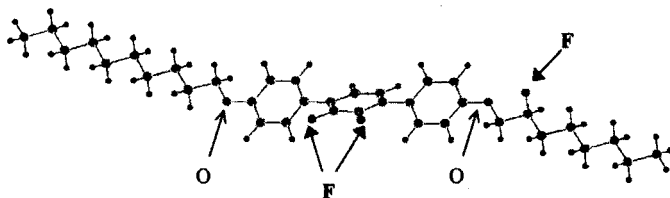


FIGURE 7 Energy-minimised molecular model of NG158 (COSMIC force field)

Table 1 shows the switching speeds of the material, obtained by shining a red laser (734nm) at a photodetector through a cell of the liquid crystal, switched under an applied square wave of  $\pm 50$ V at 50Hz. The material showed very fast response times, this indicates that the material possesses a low viscosity, also due to the absence of any ester functional groups. This suggests that the difluoroterphenyl core, in combination with a fluorine at the chiral centre and ether linkages between the core and the alkyl chains, are potentially useful in the preparation of suitable ferroelectric SmC\* components.

Temperature (°C)	Switching Speed
140	6.8 $\mu$ s
130	8.0 $\mu$ s
120	8.4 $\mu$ s
110	9.2 $\mu$ s
100	11.6 $\mu$ s
90	13.2 $\mu$ s

TABLE 1 Switching speeds of NG158 at different temperatures.

## CONCLUSIONS

A novel difluoroterphenyl ferroelectric SmC\*-forming liquid crystal, with a fluorine at the chiral centre, has been synthesized and its physical properties investigated. A postulated trade-off between  $P_s$  and viscosity, as reflected by the moderate  $P_s$  but very fast switching times, is achieved by using a material with a difluoroterphenyl core and a fluorine at the chiral centre. The viscosity is reduced by the absence of any ester groups in the material. The material may then be suitable as a component for tailoring fast-switching ferroelectric mixtures for display devices. The rotational viscosity and mixture studies of the material will be reported in a future paper.

## Acknowledgements

The authors wish to thank DERA and EPSRC for funding this work.

## References

- [1] N. A. Clark, S. T. Lagerwall, *Appl. Phys. Lett.*, **36**, 899 (1980).
- [2] T. Harada, M. Taguchi, K. Iwasa, M. Kai, *SID Digest*, **16**, 131 (1985).
- [3] N. Miyauro, T. Yanagawi, A. Suzuki, *Synth. Comm.*, **11**, 513 (1981).
- [4] T. Geelhaar, *Ferroelectrics*, **85**, 329 (1988); G. W. Gray, M. Hird, D. Lacey, K. J. Toyne, *J. Chem. Soc. Perkin Trans.*, **II**, 2041 (1989).
- [5] M. F. Bone, M. J. Bradshaw, L. K. M. Chan, D. Coates, J. Constant, P. A. Gemmell, G. W. Gray, K. J. Toyne, *Mol. Cryst. Liq. Cryst.*, **164**, 117 (1988); L. K. M. Chan, G. W. Gray, D. Lacey, *Mol. Cryst. Liq. Cryst.*, **123**, 185 (1985).
- [6] H. Orihara, K. Nakamura, Y. Ishibashi, Y. Yamada, N. Yamamoto, M. Yamakawi, *Jpn. J. Appl. Phys.*, **25**, L839 (1986).