

# Variation in molecular stacking resulting from the different polarity of liquid crystalline molecules: synthesis and study of azo dye compounds

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To investigate the effect of molecular polarity on the packing of liquid crystalline molecules, two liquid crystals, *N,N*-disubstituted aminophenylazo-4-alkylbenzenes were synthesized and studied by single crystal structure determination. A comparison of the resulting molecular stacking with that of *N,N*-disubstituted aminophenylazo-4-butylbenzoate was made.

## 1. Introduction

Supramolecular aggregation by molecular self-assembly is an important issue in the field of structural chemistry [1]. In addition to electrostatic interaction, non-covalent forces also play a significant role in determining the structural stacking and properties of molecular assemblies [2]. The interaction between the functional molecules is found to be critical for molecular packing in crystallization [3]; furthermore the stability and phase behaviour of mesogenic molecules have been reported to arise therefrom [4]. Previously, we studied two liquid crystals, azo dyes **1** and **2**, by single crystal structure determination [5]. During examination of their molecular stacking, we found that the polarity of the liquid crystals affects the molecular arrangement during crystallization. On the basis of crystallographic study, the molecular

stacking of compound **1** is shown in figure 1. Molecules in the same layer are directed head-to-head and tail-to-tail (for example, molecules a1–a3 in the layer A), but molecules in adjacent layers are directed by a head-to-tail arrangement (for example, molecules a1–a3 in layer A and molecules b1–b3 in layer B) [5*a*]. Apparently, the carboxylate and amido groups in compound **1** are both strongly polar moieties. For greatest molecular attraction and minimum dipole moment in the solid state, the carboxylate group of one molecule is directed toward the amido group of another molecule in an adjacent layer. The closest distance between different layers (not including hydrogen atoms) thus appears in the vicinity of these groups, and the distances of the C1–O1 and C2–O2 between A and B layers are 3.48 and 3.25 Å, respectively. For a better understanding of

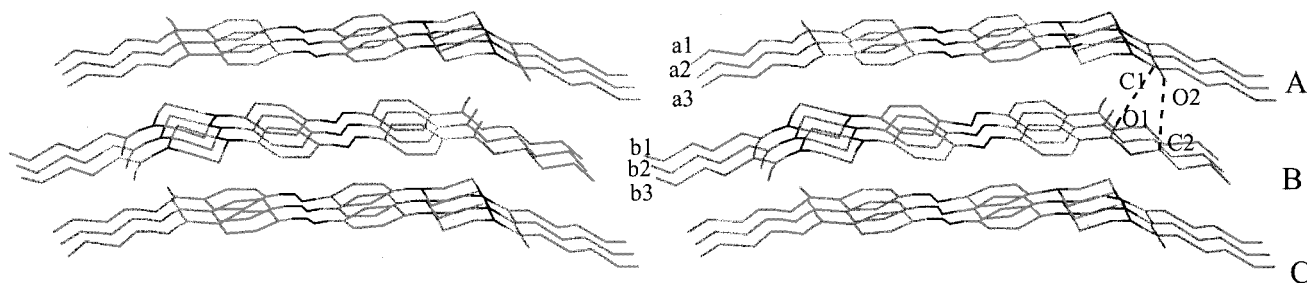
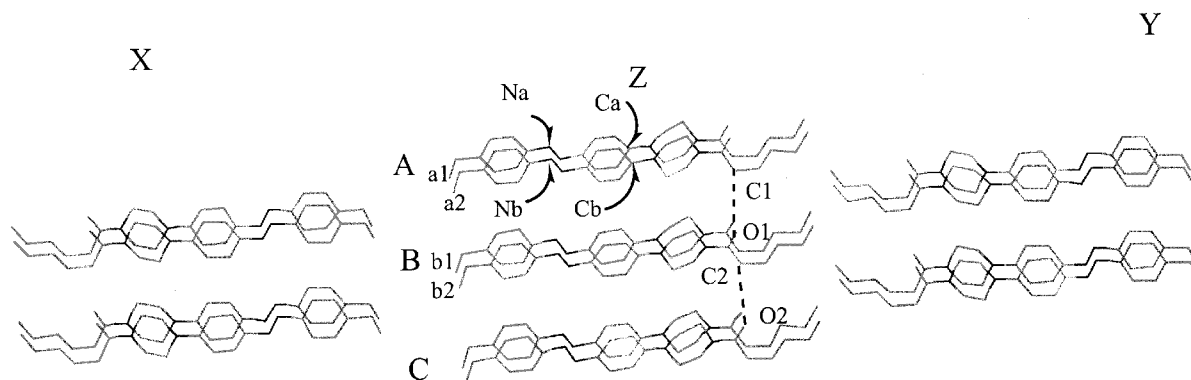
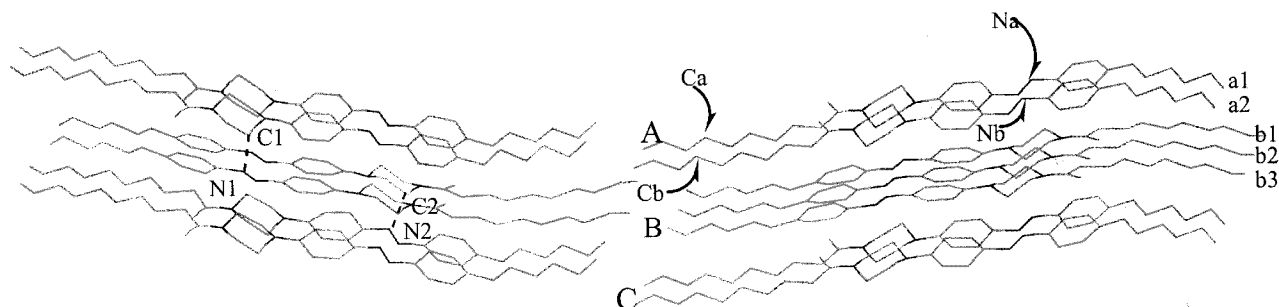


Figure 1. The molecular stacking of azo dye **1**.

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Figure 2. The molecular stacking of azo dye **5a**.Figure 3. The molecular stacking of azo dye **5b**.

the molecular stacking of liquid crystals with different polar moieties, a similar azo dye compound without the carboxylate group has been synthesized and studied by single crystal structure determination. As the driving force for the molecular stacking could be different from the case of azo dye **1** which contains amido and carboxylate groups, compounds **5a** and **5b** containing only an amido group were accordingly prepared and their molecular stackings and mesogenic behaviours investigated. We now wish to report preliminary results.

## 2. Synthesis

Compounds **5a** and **5b** were synthesized according to the scheme. Compound **3** was first obtained by reaction of *N*-phenylpiperazine with alkanoyl chloride in dichloromethane in an almost quantitative yield. The diazonium salt **4** was then prepared according to the literature method [6]; compounds **5a** and **5b** were subsequently obtained in about 15% yields, and characterized to be the correct compounds by NMR (Varian 300) and high resolution mass spectroscopy (VG70-250; EI, 70ev).

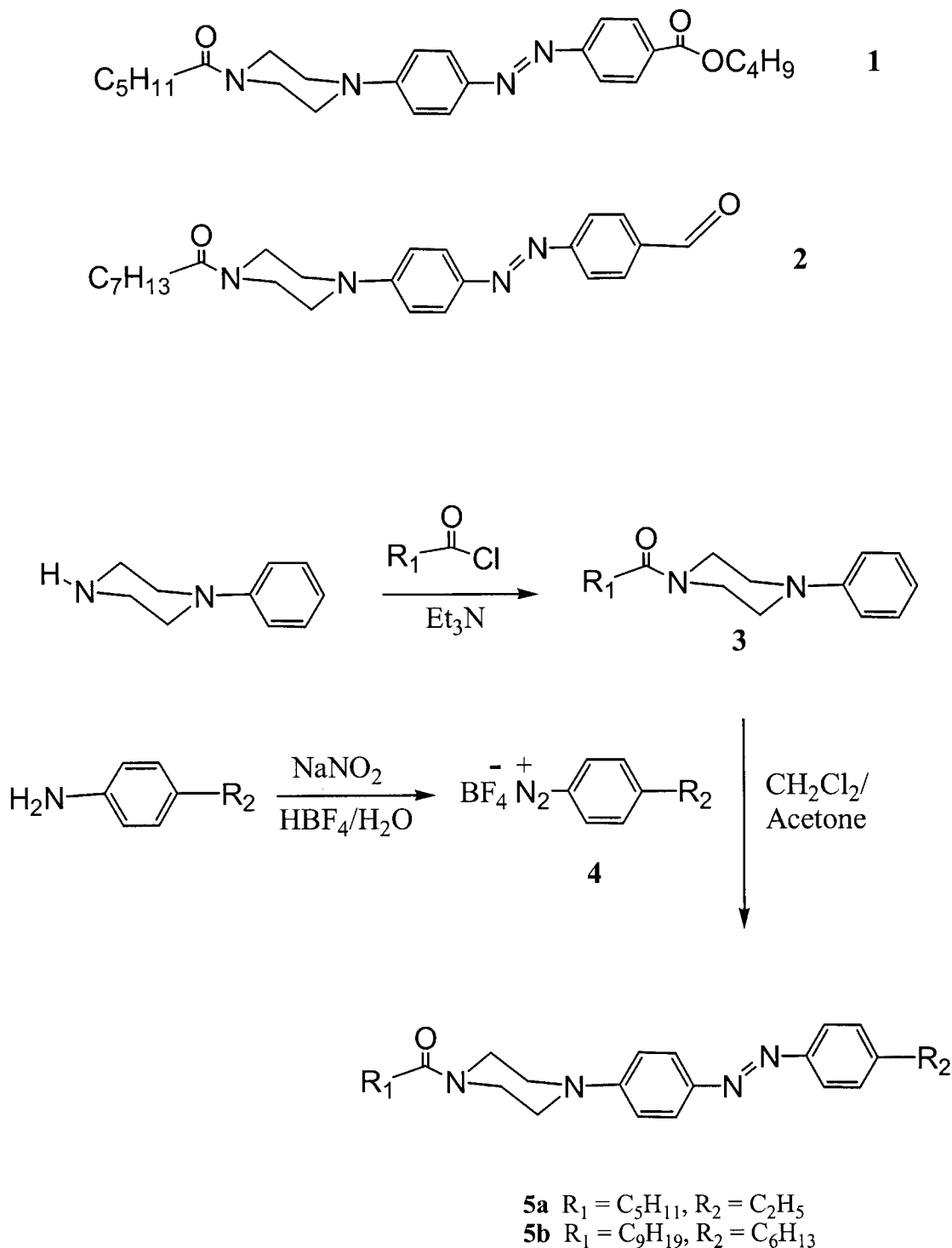
**5a**. 17.1%.  $^1\text{H}$  NMR:  $\delta$  ( $\text{CDCl}_3$ ) 0.94 (t, 3H,  $\text{CH}_3$ ), 1.33 (t, 3H,  $\text{CH}_3$ ), 1.69–1.78 (m, 6H,  $3 \times \text{CH}_2$ ), 2.41 (t, 2H,  $\text{CH}_2$ ), 2.75 (quart, 2H,  $\text{CH}_2$ ), 3.38 (s, b, 4H,  $2\text{CH}_2$ ), 3.68 (t, 2H,  $\text{CH}_2$ ), 3.82 (t, 2H,  $\text{CH}_2$ ), 7.01 (d, 2H,  $J = 8.7$  Hz,  $2 \times \text{Ar-H}$ ), 7.34 (d, 2H,  $J = 8.1$  Hz,  $2 \times \text{Ar-H}$ ), 7.82 (d, 2H,  $J = 8.7$  Hz,  $2 \times \text{Ar-H}$ ), 7.91 (d, 2H,  $J = 8.1$  Hz,  $2 \times \text{Ar-H}$ ). HRMS for  $\text{C}_{24}\text{H}_{32}\text{N}_4\text{O}$ :

392.2576; found 392.2578. **5b**. 16.5%.  $^1\text{H}$  NMR  $\delta$  ( $\text{CDCl}_3$ ) 0.78–0.88 (m, 6H,  $2 \times \text{CH}_3$ ), 1.23–1.69 (m, 22H,  $11 \times \text{CH}_2$ ), 2.35 (t, 2H,  $\text{CH}_2$ ), 2.65 (t, 2H,  $\text{CH}_2$ ), 3.34 (s, b, 4H,  $2\text{CH}_2$ ), 3.67 (s, b, 2H,  $\text{CH}_2$ ), 3.80 (s, b, 2H,  $\text{CH}_2$ ), 7.01 (d, 2H,  $J = 8.7$  Hz,  $2 \times \text{Ar-H}$ ), 7.27 (d, 2H,  $J = 8.1$  Hz,  $2 \times \text{Ar-H}$ ), 7.78 (d, 2H,  $J = 8.7$  Hz,  $2 \times \text{Ar-H}$ ), 7.92 (d, 2H,  $J = 8.1$  Hz,  $2 \times \text{Ar-H}$ ). HRMS for  $\text{C}_{32}\text{H}_{48}\text{N}_4\text{O}$ : 504.3828; found 504.3830.

## 3. Characterization and results

Compounds **5a** and **5b** were studied by single crystal structure determination†, and their corresponding molecular stackings are shown in figures 2 and 3, respectively. In figure 2, molecules of **5a** in the same column are directed regularly head-to-head and tail-to-tail (for example, molecules a1–a2 and b1–b2 in column Z). Furthermore, molecules are found to be

†Crystal data, **5a**:  $\text{C}_{24}\text{H}_{32}\text{N}_4\text{O}$ , crystal dimension  $0.45 \times 0.25 \times 0.10 \text{ mm}^3$ ,  $P2_1$ , monoclinic,  $a = 7.602(2)$ ,  $b = 5.7559(12)$ ,  $c = 49.912(10) \text{ \AA}$ ,  $\beta = 90.19(3)^\circ$ ,  $Z = 4$ ,  $\rho_{\text{calcd}} = 1.194 \text{ g cm}^{-3}$ , reflections measured = 12334, reflections used = 3830 ( $(R_{\text{int}}) = (0.0390)$ ),  $R_1 = 0.0873$ ,  $[I > 2\sigma(I)]$ ,  $R_1 = 0.1083$  (all data), GOF on  $F^2 = 1.158$ . **5b**:  $\text{C}_{32}\text{H}_{48}\text{N}_4\text{O}$ , crystal dimension  $0.35 \times 0.25 \times 0.02 \text{ mm}^3$ ,  $P2_12_12_1$ , orthorhombic,  $a = 5.724(2)$ ,  $b = 7.364(2)$ ,  $c = 67.76(2) \text{ \AA}$ ,  $Z = 4$ ,  $\rho_{\text{calcd}} = 1.174 \text{ g cm}^{-3}$ , reflections measured = 8198, reflections used = 4980 ( $(R_{\text{int}}) = (0.0843)$ ),  $R_1 = 0.0953$ ,  $[I > 2\sigma(I)]$ ,  $R_1 = 0.1760$  (all data), GOF on  $F^2 = 1.003$ .

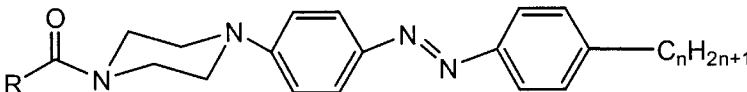


Scheme.

parallel to the adjacent molecules in the same layer (for example, molecules a1–a2 in layer A) and the distances between two corresponding atoms in the same layer (for example, Na–Nb and Ca–Cb) are all about 7.6 Å. The

plane containing diazobenzene moieties of the molecule in layer A is approximately parallel to the corresponding plane in layer B and the distance between two parallel planes is about 3.8 Å, which is half of the distance between

Table. Phase transition temperature ( $^{\circ}\text{C}$ ) and corresponding enthalpies ( $\text{J g}^{-1}$ ), in parentheses, of **5**. The phase transition temperatures and corresponding enthalpies of **5** were determined by 2nd DSC scans using a Perkin Elmer DSC-6 at heating and cooling rates of  $10^{\circ}\text{C min}^{-1}$  between 50 and  $220^{\circ}\text{C}$ . Abbreviations: Cr = crystalline, SmX = unidentified smectic phase, SmX' = unidentified smectic phase, SmA = smectic A phase, I = isotropic liquid.



**5**

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<b>5a</b> ,	$n = 2$ $R = \text{C}_5\text{H}_{11}$	Cr1	$\xrightleftharpoons[56.6]{125.7(40.1)}$	SmX	$\xrightleftharpoons[63.0^b]{138.6}$	SmX'	$\xrightleftharpoons[137.6(3.6)]{141.1^a}$	SmA	$\xrightleftharpoons[180.0(12.8)]{187.4(11.6)}$	I
<b>5b</b> ,	$n = 6$ $R = \text{C}_9\text{H}_{19}$	Cr1	$\xrightleftharpoons[56.6]{104.1(16.0)}$	SmX	$\xrightleftharpoons[153.1(5.4)]{154.1(6.8)}$	SmA	$\xrightleftharpoons[192.7(14.5)]{195.1(15.5)}$	I		

<sup>a</sup> The peaks are overlapped and their total enthalpy is  $4.5 \text{ J g}^{-1}$ .

<sup>b</sup> The peaks are overlapped and their total enthalpy is  $26.5 \text{ J g}^{-1}$ .

corresponding atoms in the same layer. The closest distance between two adjacent layers (not including hydrogens) is  $3.88 \text{ \AA}$ , found, for example, at the C1–O1 between layers A and B as well as at C2–O2 between layers B and C. The distances O1–H (at C1) and O2–H (at C2) are both  $3.01 \text{ \AA}$ , which are in the range of a normal hydrogen bond [7*a*]. After careful examination of molecular stacking, we do not find that the interatomic distance in different molecules is closer than this value. We thus believe that the driving force for the stabilization of molecular stacking of azo dye **5a** mostly arises from intermolecular hydrogen bonding.

In figure 3, molecules of **5b** are directed regularly head-to-head and tail-to-tail in the same layers (for example, molecule a1–a2 in layer A). However, molecules in adjacent layers are arranged head-to-tail (for example, molecules a1–a2 in layer A and molecules b1–b3 in layer B). In the same layer, the molecules are found to be parallel to adjacent molecules and the distances between two corresponding atoms in the same layer (for example, Na–Nb and Ca–Cb) are all about  $5.72 \text{ \AA}$ . Molecules in different layers are not parallel to each other. The intercept angle between the plane containing diazobenzene moieties of the molecules in layer A and the corresponding plane in layer B is estimated at about  $40^{\circ}$  on the basis of crystallographic data. The closest distance between two adjacent layers (not including hydrogens) is  $3.37 \text{ \AA}$ , which is, for example, found at the C1–N1 between A and B layers, as well as C2–N2 between B and C layers. The distances of N1–H (at C1) and N2–H (at C2) are both  $2.82 \text{ \AA}$  which is comparable to the sum of the van der Waals radii of H and N (Bondi radius: H  $1.20$ , N  $1.55$ ) [7*b*]. The H-bond interaction arising therefrom should be influential in

the crystallizing process and is believed to be the major contribution to stabilization of the molecular stacking of compound **5b**.

Based on the previous results, we may reasonably regard that for a bi-polar compound such as azo dye **1**, the polarity of functional groups determines the orientation of the molecular arrangement in the crystal structure. However, for mono-polar compounds such as azo dyes **5a** and **5b**, the intermolecular hydrogen bonding interaction is most influential for the molecular stacking. Additionally, it is interesting to note that molecule **5a** is in a linear arrangement, but there is a slight bending between the piperazine and decanoyl moieties in structure **5b**. The hexyl tail at the other end of molecule **5b** makes this bending more pronounced and thus the whole molecule is arranged in a slightly curved shape. Compounds **5a** and **5b** both show a SmA phase in thermal cycling, characterized by a focal-conic texture under polarizing optical microscopy (see the table), which is different from the mesogenic behaviour of azo dye **1** which shows a SmC phase [5*a*].

Based on the crystallographic study, the extended molecular lengths of azo dyes **5a** and **5b** are calculated to be  $24.08$  and  $32.83 \text{ \AA}$ , respectively, which are quite close to the corresponding *d*-spacing distances of azo dyes **5a** and **5b** in the SmA phase, obtained from powder X-ray diffraction (XRD)<sup>‡</sup>. Although the amido moiety

<sup>‡</sup> The *d*-spacings of compound **5a** are  $24.95$ ,  $25.09$ ,  $25.09$ ,  $25.11$ ,  $25.24 \text{ \AA}$  at  $180$ ,  $170$ ,  $160$ ,  $150$ ,  $140^{\circ}\text{C}$ , respectively, during cooling; the *d*-spacings of compound **5b** are  $32.88$ ,  $32.86$ ,  $32.70$ ,  $2.45$ ,  $32.41 \text{ \AA}$  at  $188$ ,  $185$ ,  $175$ ,  $165$ ,  $155^{\circ}\text{C}$ , respectively, during cooling. Powder XRD patterns were obtained from a Siemens D-5000 X-ray Diffractometer equipped with a TTK 450 temperature controller.

of compounds **5a** and **5b** may vibrate up and down during the thermal process, the extended length of the molecules may not change much as a result. According to the literatures, the *N*-substituent of the piperazine favours equatorial conformation, and the energy barrier between the equatorial and axial conformations is only about 1.9–4 kcal mol<sup>-1</sup> [8]. As we have shown in the case of azo dye **1** [5a], the *N*-substituent in either equatorial or axial conformation does not significantly change the extended molecular length. Thus the *d*-spacing of the molecules will not vary much during thermal processing; this is consistent with the observations obtained from powder XRD.

#### 4. Conclusion

While exploring crystal stacking with varying molecular polarity, compounds **5a** and **5b** were investigated by crystallography and the study of mesogenic behaviour. As mentioned previously, for the bi-polar azo dye **1**, the mutual attraction from amido and carboxylate groups of different molecules affects the molecular stacking in the crystal structure. For the mono-polar azo dyes **5a** and **5b**, intermolecular hydrogen bonding is the major contribution to stabilization of the molecular stacking. These results provide not only a better understanding of the effect of the functional group and hydrogen bonding on the stacking of liquid crystals in the solid state, but also provide a useful guide for the future design of desired materials.

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#### References

- [1] (a) LEHN, J.-M., 1995, *Supramolecular Chemistry* (Weinheim: VCH); (b) HAMILTON, A. D., 1996, *Perspectives in Supramolecular Chemistry* (John Wiley).
- [2] (a) HUNTER, C. A., 1993, *Angew. Chem. int. Ed. Engl.*, **32**, 1584; (b) HUNTER, C. A., 1994, *Chem. Soc. Rev.*, **23**, 101.
- [3] (a) DESIRAJU, G. R., 1989, *Crystal Engineering: The Design of Organic Solids* (New York: Elsevier); (b) DESIRAJU, G. R., 1997, *Chem. Commun.*, 1475.
- [4] (a) LEE, K. M., LEE, C. K., and LIN, I. J. B., 1997, *Angew. Chem. int. Ed. Engl.*, **36**, 1850; (b) CHANG, J., and MOORE, J. S., 1994, *J. Am. chem. Soc.*, **116**, 2655.
- [5] (a) LAI, L. L., LEE, L. J., LEE, G. H., WANG, Y., LU, K. L., and LEE, S. J., 2001, *Liq. Cryst.*, **28**, 1513; (b) LAI, L. L., LIN, Y. J., HO, C. H., WANG, E., LIU, Y. H., WANG, Y., LIN, Y. C., and CHENG, K. L., 2001, *Helv. Chim. Acta* (in the press).
- [6] McNAB, H., and MONAHAN, L. C., 1989, *J. chem. Soc. Perkin Trans. 1*, 419.
- [7] ALLEN, F. H., LOMMERSE, J. M. P., HOY, V. J., HOWARD, J. A. K., and DESIRAJU, G. R., 1996, *Acta Cryst.*, **B52**, 734; (b) BONDI, A., 1964, *J. phys. Chem.*, **68**, 441.
- [8] (a) BAKER, V. J., FERGUSON, I. J., KATRITZKY, A. R., and PATEL, R., 1976, *Tetrahedron. Lett.*, 4735; (b) NIEMEYER, H. M., 1979, *J. mol. Struct.*, **57**, 241.

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