



## Preparation and Aggregation-Induced Emission of New 1,3,5-Triazine-2,4,6-tricarboxamide with Liquid Crystal Properties

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### ABSTRACT

The combination of aggregation-induced emission (AIE) and liquid crystal properties generates solid-state efficient luminescent liquid crystal materials. Here in, we reported the synthesis of 1,3,5-triazine-2,4,6-tricarboxamide and utilized it as a supramolecular organic motif for the AIE-active liquid crystal material. The compound exhibits high-intensity emission maxima at 417 and 468 nm in the solid state with excitation at 254 nm, whereas it shows weak emission in the solution phase. Also, this compound behaves as liquid crystalline material and shows columnar hexagonal mesophase with endothermic peaks at 73.4°C, 185.6°C, and exothermic peaks were observed at 181.9°C and 66.1°C with focal conic fan shape texture. The thermal data showed that the compound is stable up to 200°C.

**Keywords:** Luminescence, 1,3,5-Triazine-2,4,6-Tricarboxamide, Aggregation-induced emission, liquid crystal.

### INTRODUCTION

Organic solid luminescent materials have been attracting particular attention due to their vast potential applications, in displays, optoelectronics, organic solar cells, organic light emitting diode (OLED) as well as in the field of sensing and imaging<sup>1,2</sup>. Conventional solid organic molecules are known to exhibit strong  $\pi$ - $\pi$  stacking interactions due to their extended  $\pi$ -conjugation and planar structure.

This interaction promotes loss of excitation energy. Consequently, the molecules are poorly or non emissive in the solid state, even though they emit efficiently in solution. The phenomenon is called aggregation-caused quenching (ACQ). Although many attempts have been made to overcome the ACQ effect, such as introductions of large cyclic species and dendritic wedges to Fluorophores, these approaches often led not only to complicated syntheses, but also undesired photophysical



properties<sup>5,6</sup>. In 2001, Tang *et al.*, reported an unusual phenomenon with silacyclopentadine (silole) derivatives that are non emissive in solution but become highly emissive upon aggregation. The phenomenon is termed aggregation-induced emission (AIE)<sup>7</sup>. Restriction of intramolecular rotation was identified as the mechanism for AIE effect in aggregate state<sup>8</sup>.

Recently, many attention has been focused on luminescent liquid crystals (LCs) organic materials<sup>9,10</sup>, due to their combination of light emission, supramolecular organization and broad technological applications such as organic light-emitting diodes, polarized organic lasers<sup>11</sup> and sensors<sup>12</sup>. In spite of the unique properties of luminescent liquid crystal molecules, their syntheses remain challenging due to difficulty incorporating emissive functional group and liquid crystal properties. Also, many conjugated molecular emitters suffer aggregation-cause quenching effect leading to the quenching of fluorescence<sup>12</sup>. Therefore, the utilization of aggregation-induced emission could circumvent the ACQ effect since AIE molecules usually show strong emission in the solid or aggregate state. Although AIE-active LCs show promising prospects, the research is still rarely reported since the requirements for the LC and AIE characteristic are hard to fulfil at the same time in one single molecule. Here in, we report the first example of 1,3,5-triazine-2,4,6-tricarboxamide as an AIE-active liquid crystal (AIE-LC) bearing hydrophobic alkyl side chain with simple aromatic 1,3,5-triazine AIE core. It has been known that attaching either long alkyl/alkoxy groups to the AIE core creates light-emitting liquid crystal materials<sup>13</sup>.

## MATERIALS AND METHODS

### General

All chemicals were obtained from Aldrich and used as received without further purification unless otherwise stated. <sup>1</sup>H-NMR spectra were recorded on a Bruker Ultra-Shield 400 MHz spectrometer. Samples were dissolved in CDCl<sub>3</sub> or D<sub>2</sub>O as solvent and chemical shifts are reported in ppm. FT-IR spectra were recorded on a Perkin Elmer spectrometer in the range of 4000–400 cm<sup>-1</sup> using KBr disk. Mass spectra were obtained using a Kratos MALDI-TOF mass spectrophotometer system. UV-Vis

absorption spectrum was recorded on a Shimadzu UV-2600 spectrophotometer. Photoluminescence spectrum was recorded on JASCO FR 8500 spectrophotometer. TGA was performed a Mettler Toledo tga/851e thermal analyser under nitrogen atmosphere at a heating rate of 10°C min<sup>-1</sup>. The liquid crystal properties were determined using differential scanning calorimetry (DSC) on a Mettler Toledo calorimeter model DSC822e at range of temperature of 25 to 200°C. Throughout the DSC analysis, the sample was heated and cooled at the rate of 10°C min<sup>-1</sup>. Polarized optical microscopy (POM) was performed on a Leica DM2700 P model optical polarizing microscope equipped with a Linkam LTS420 hot-stage.

### Synthesis of 1,3,5-Triazine-2,4,6-Tricarboxylic acid (2)

Triethyl-1,3,5-triazine-2,4,6-tricarboxylate **1** (1.0 g, 3.36 mmol) was transferred into a 250 mL round bottom flask fitted with a Liebig condenser. Separately, NaOH (0.672 g, 16.8 mmol) was dissolved in 50 mL mixture of water-ethanol (1:1) to form a homogenous solution. The solution was then added dropwise to the triethyl-1,3,5-triazine-2,4,6-tricarboxylate, and the resultant mixture was heated to reflux for 10 h while stirring. After cooling to room temperature, the pH of the resulting solution was adjusted to pH 2 with 0.1 M HCl after which a white precipitate was obtained. It was then filtered, recrystallized in ice-cold water and dried to give a white powder solid. Yield: 0.7 g, 3.28 mmol, 97%. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub> δ/ppm): 10.68 (s, 3H, OH). FTIR (KBr): 3447 (m), 1744 (s), 1651 (w), 1416 (m) 1248 (s) 852 cm<sup>-1</sup> (m).

### Synthesis of 1,3,5-triazine-2,4,6-tricarbonyl trichloride (3)

1,3,5-triazine-tricarboxylic acid **2** (0.5 g, 2.35 mmol) was transferred into a 250 mL two-necked round-bottom flask and was treated under vacuum for 10 min. before being flowed with nitrogen gas after which 40 mL DMF was added and stirred at 0°C for 1 hour. Then, oxalyl chloride (1 mL, 7.75 mmol) was added dropwise, and the resulting solution was stirred at room temperature overnight. The mixture was then heated for 2 h at 70°C to remove any excess oxalyl chloride before allowing it to cool to room temperature. The solvent was removed by heating at 90°C under reduced

pressure and vacuum dried overnight to give a pale pink solid product. The obtained solid was used for further synthesis without purification.

### Synthesis of 1,3,5-triazine-2,4,6-tricarboxamide (5)

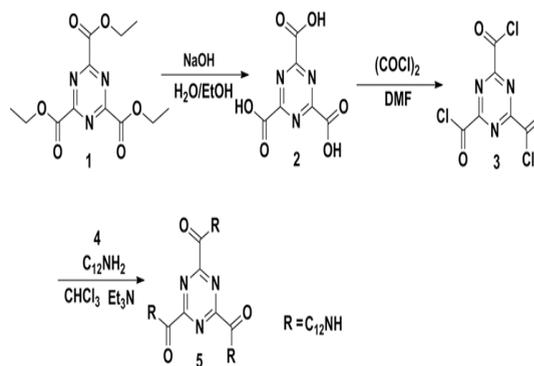
1,3,4-triazinetriacarbonyl trichloride **3** (0.631 g, 2.35 mmol) was placed into a two-neck round bottom flask and was treated under vacuum for 10 min. before being flowed with nitrogen gas. Then 1-aminododecane (1.44 g, 7.76 mmol) was added dropwise, and the mixture was added with 50 mL dried chloroform, followed by the addition of triethylamine (1.09 mL, 7.76 mmol). After the addition was completed, the reaction mixture was stirred overnight at room temperature, and the resulting mixture was filtered. The targeted product was successfully isolated using column chromatography with chloroform/methanol 50:1 as an eluent and subsequently evaporated and dried under inert condition by using vacuum pump under reduced pressure at room temperature to afford a white solid powder product. Yield: 0.87g, 1.22 mmol, 52 %. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub> δ/ppm): 0.83-0.87 (t, J = 7.2 Hz, 9H, -CH<sub>3</sub>), 1.22- 1.35 (overlapped, 54H, -CH<sub>2</sub>(CH<sub>2</sub>)<sub>10</sub>) 1.70-1.79 (m, 6H -CH<sub>2</sub>) 2.93-2.97 (t, J = 8.0 Hz, 6H -NHCH<sub>2</sub>), 8.20(s, 3H, NH). FTIR (KBr): 3430 (M), 1630 (s), 1585 (m), 1516 (w) 1475 (m), 1371 (m), 1143, (s) 835 cm<sup>-1</sup> (m) MS (MALDI-TOF): m/z 753.5757 [M + K<sup>+</sup>] observed, Calcd for C<sub>42</sub>H<sub>78</sub>N<sub>6</sub>O<sub>3</sub>: 753.5772 [M+K<sup>+</sup>].

## RESULTS AND DISCUSSION

Scheme 1 illustrates the structures and the synthetic route to 1,3,5-triazine-2,4,6-tricarboxamide (**5**). The target compound was synthesized by Schotten-Baumann amidation reaction<sup>14</sup>, starting with the base hydrolysis of triethyl-1,3,5-triazine-2,4,6-tricarboxylate **1** to afford 1,3,5-triazine-2,4,6-tricarboxylic acid **2** in good yield. Subsequently, compound **2** was subjected to acylation reaction with oxalyl chloride, (COCl)<sub>2</sub>, in the presence of DMF resulting in the formation of 1,3,4-triazinetriacarbonyl trichloride **3** which was then treated with 1-aminododecane in the presence of CHCl<sub>3</sub>/Et<sub>3</sub>N to obtain **5** as white powder in 52% yield.

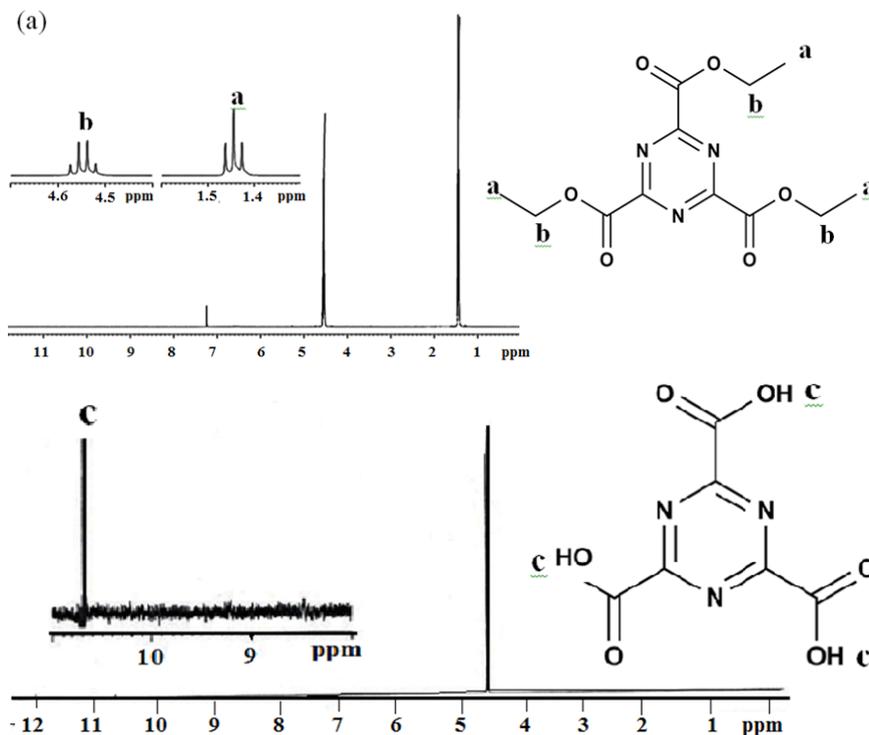
Based on the H-NMR spectra in Fig. 1 summarized in Table 1, the spectrum of precursor 1

dissolved in CDCl<sub>3</sub> showed two signals corresponding to the methylene and methyl protons while the product **2** dissolved in D<sub>2</sub>O showed one signal belong to the hydroxyl proton. The chemical shift at 1.43-1.47 ppm (triplet, 9H, = Ha -CH<sub>3</sub>) and 4.52-4.57 ppm (quartet, 6H, Hb = -CH<sub>2</sub>) of compound **1** disappeared in the spectrum of **2** due to cleavage of the ethyl group. Moreover, the appearance of peak at 10.68 ppm (singlet, 3H, Hc = OH) indicate the successful formation of compound **2**. It was further confirmed with FTIR as shown in Fig. 2a, by the disappearance of C-H vibrations in the region 3000–2883 cm<sup>-1</sup> indicating the cleavage of OCH<sub>2</sub>CH<sub>3</sub> bond in compound **2** and the appearance of broad peak at 3447 cm<sup>-1</sup> assigned to O-H stretching vibration of **2** in Fig. 2b<sup>15</sup>.



Scheme 1. Synthesis route for 1,3,5-Triazine-2,4,6-Tricarboxamide

Figure 3 showed <sup>1</sup>H NMR spectra of the reactants 1-aminododecane **4** and product 1,3,5-triazine-2,4,6-tricarboxamide **5** as summarized in Table 2. The chemical shift of 1-aminododecane at 2.60-2.65 ppm (t, J = 8.0 Hz, 2H -NHCH<sub>2</sub>) in Fig. 3a was shifted to 2.93-2.97 (t, J = 8.0 Hz, 6H -NHCH<sub>2</sub>) in Fig. 3b due to the successful transformation of acyl chloride group of **3** to form carboxamide group. Also the appearances of a broad peak at chemical shift 8.20 ppm (s, 3H, NH) in Fig. 3b indicates the attachment of alkyl side chains to triazine ring. Moreover, the formation of **5** was further confirmed by FTIR spectra in Fig. 2c by the appearance of vibrational peaks at 3436 cm<sup>-1</sup> attributed to N-H stretching and 1558 cm<sup>-1</sup> N-H bending as well as 2849-2950 cm<sup>-1</sup> assigned to C-H stretching and 1475 cm<sup>-1</sup> C-H bending, which show that the alkyl side chains was successfully bonded to the triazine ring. It was Further supported by MALDI-TOF-MS molecular weight m/z 753.5757 Da [M+K<sup>+</sup>] observed, 753.5772 Da [M+K<sup>+</sup>] Calculated for chemical formula C<sub>42</sub>H<sub>78</sub>N<sub>6</sub>O<sub>3</sub>.

Fig. 1 <sup>1</sup>H-NMR spectra of (a) 1 and (b) 2Table 1: <sup>1</sup>H-NMR data for 1 and 2

Proton (H)	Chemical shift (ppm)	
	1	2
a	1.43-1.47 (9H)	–
b	4.52-4.55 (6H)	–
c	–	10.68 (3H)

### Photoluminescent behaviour

1,3,5-triazine-2,4,6-tricarboxamide exhibits fluorescent emission maxima centered at 417 nm and 468 nm on excitation at 254 nm. The luminescent response was investigated in solid and solution. It shows weak fluorescent emission intensity in chloroform solution, in contrast, it gave strong emission intensity in the solid state as shown in Fig. 4a. Upon excitation at 365 nm with hand-held UV lamp, the solid powder gave blue emission, but when the solution of **5** was illuminated at 365 nm, practically no emission was observed Fig. 4b. Furthermore, the luminescent behaviour was investigated using UV-Vis absorption in solid state and solution as shown in Fig. 4c. In solid state **5** exhibit intense broadened absorption bands from 250-800 nm red shifted with respect to absorption in

solution, suggesting intermolecular charge transfer (CT) interactions due to formation of supramolecular J-aggregate. However, in chloroform solution, the broad intense absorption band collapsed suggesting dissolved molecular species and the decreased in intermolecular charge transfer (CT) interactions, which is evident by a blue shift observed and a sharp absorption band at 231 nm suggesting molecular isolated species.

The intense emission of **5** in solid could be due to aggregation-induced emission (AIE). And its AIE mechanism could be accounted for similar to the previous reports<sup>9,16</sup>. The plausible AIE mechanism is shown in Fig. 5. In contrast to the ACQ effect as a result of free intramolecular rotations along C-C bonds, molecules with AIE properties exhibit strong fluorescent in the aggregate state, attributed to the restriction of intramolecular rotations. It was suggested that multiple non covalent interactions such as hydrogen bonding would stiffen the structure of the molecules upon aggregation hindering free intramolecular rotations<sup>17,18</sup>.

It is proposed that in the solution state, **5** exist as isolated molecular species which undergo free

intramolecular rotations along the amide units upon excitation, which may result in the loss of excitons energy leads to the quenching of the fluorescent emission intensity. However, in the aggregate or

solid state, the free intramolecular rotations are restricted due to multiple intermolecular interactions which blocked the non radiative relaxation pathway resulting in the stronger fluorescent emission.

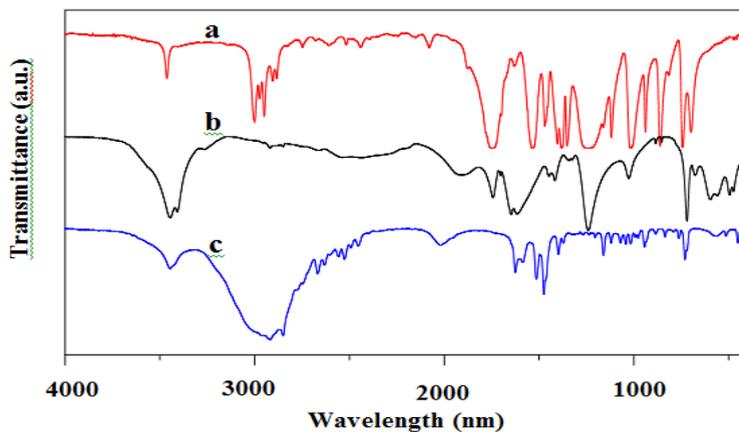


Fig. 2. FTIR spectra of compounds (a) 1 (b) 2 and (c) 5

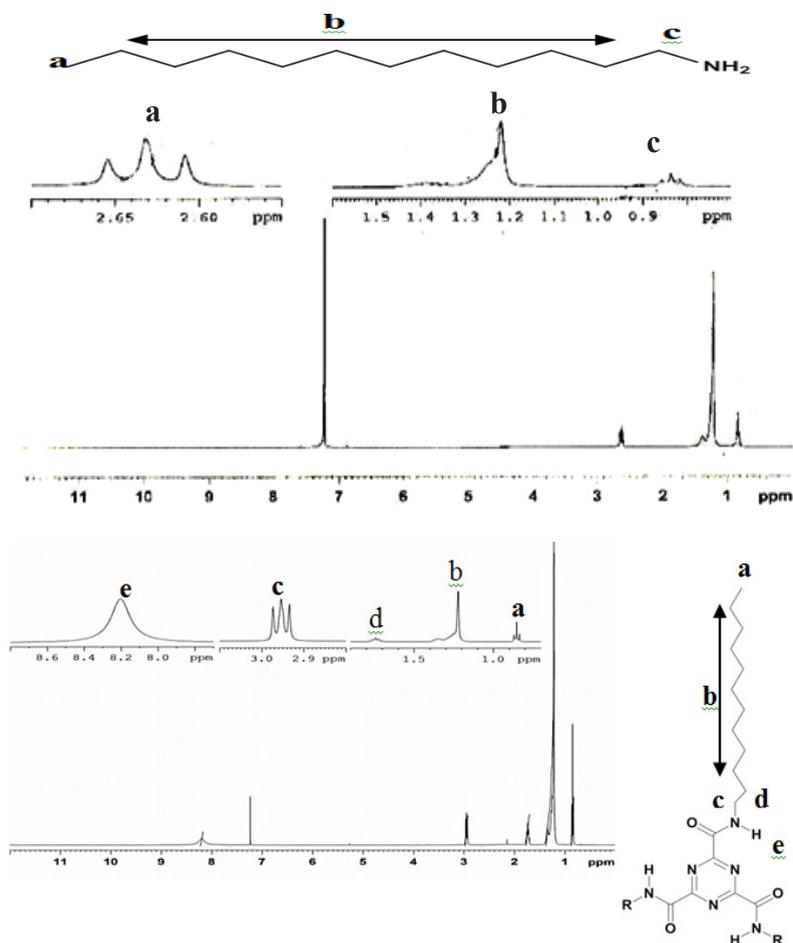


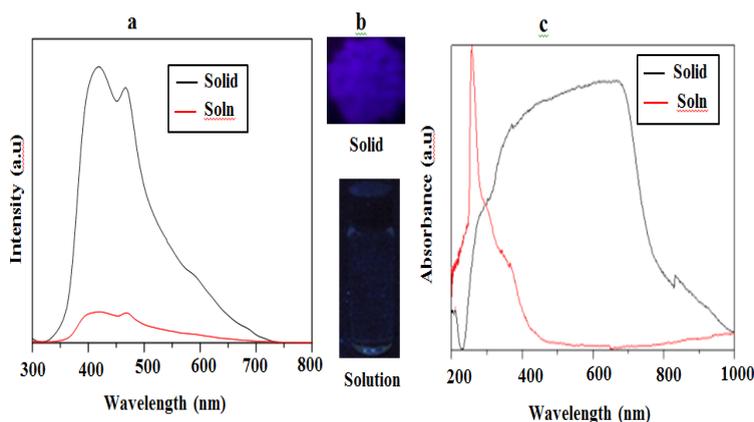
Fig. 3.  $^1\text{H-NMR}$  spectra of (a) 4 and (b) 5

**Table 2: <sup>1</sup>H-NMR data for 4 and 5**

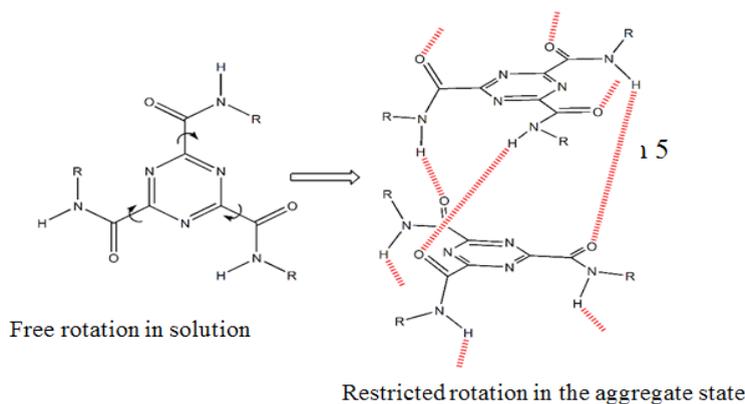
Proton (H)	Chemical shift (ppm)	
	4	5
a	0.77-0.86 (3H)	0.83-0.87 (9)
b	1.29-1.41 (20H)	1.22-1.35 (54H)
c	2.60-2.65 (2H)	2.93-2.97 (6H)
d	–	1.70-1.79 (6H)
e	–	8.20 (3H)

**Liquid crystal properties**

The liquid crystal behaviour was investigated using polarized optical microscopy (POM) and differential scanning calorimetry (DSC). The heating and cooling rate were 10°C min<sup>-1</sup>. Upon cooling from its isotropic phase at 190°C to anisotropic texture the POM showed the appearance of focal conic fan-like texture a typical characteristic of hexagonal columnar mesophase (Colh)<sup>19</sup>, as shown



**Fig. 4.** (a) Emission spectra of 5 in solid and solution (b) Photograph of its emission in solid and solution under illumination of UV lamp (c) Absorption spectra in solid and solution



**Fig. 5.** Plausible AIE mechanisms in 5

in Fig. 6a and b. The DSC thermogram displays two endothermic transitions at 73.4°C and 185.6°C upon heating from crystalline (Cr) phase to isotropic (Iso) phase and two exothermic peaks at 181.9°C and 66.1°C when cooled from the isotropic to crystalline phase as shown in Fig. 6c. In addition, two crystalline modifications peaks were observed upon cooling (Cr and Cr1) related to the crystallization process<sup>20</sup>.

We believed that the transition from the isotropic phase to crystal phase on cooling is accompanied by a reorganization of the molecule in which crystal structural modification is likely to occur.

**Thermal behaviour**

The thermogravimetric analysis (TGA) property of 5 has two identifiable weight loss stages as shown in Fig. 7. It starts in a temperature range

of 200°C with a 1% weight loss corresponding to the organic structural collapse of the triazine ring and the alkyl chain. Above 200°C, the compound showed

gradual decrease in weight up to 95% and finally decomposed at 330°C. Thus this thermal analysis data suggested that the compound is stable up to 200°C.

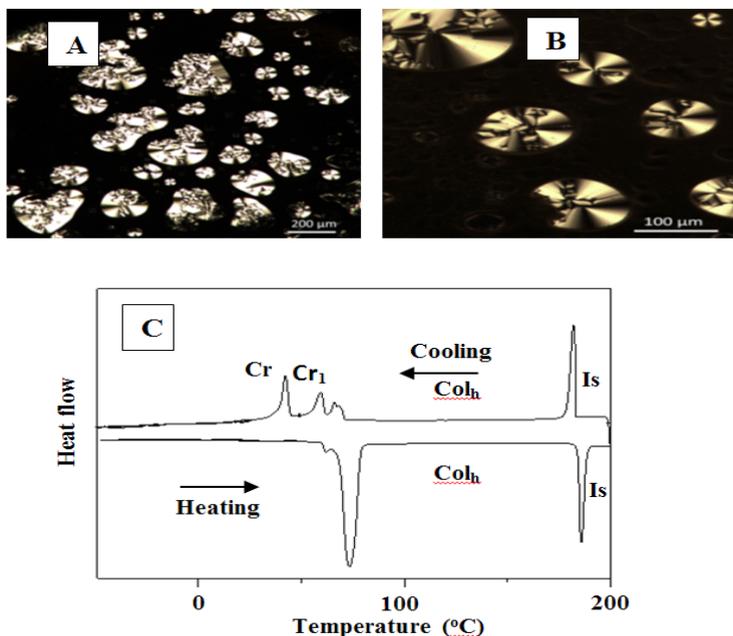


Fig. 6. POM images of 5 upon cooling from isotropic state (a) at 200  $\mu\text{m}$  (b) at 100  $\mu\text{m}$  and (c) DSC thermogram of 5 upon heating and cooling cycle

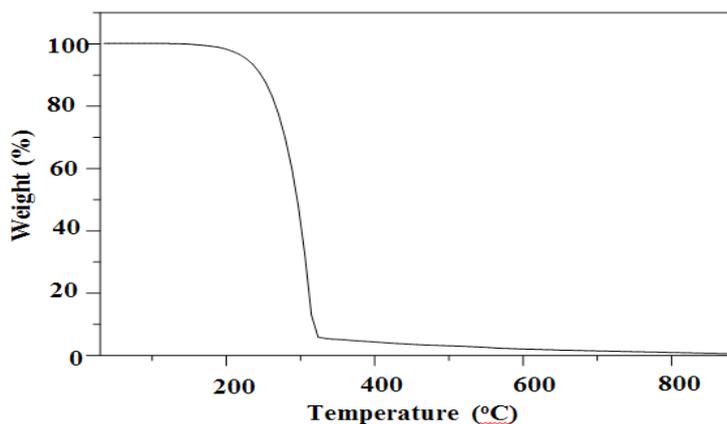


Fig. 7. TGA plot for 5

## CONCLUSION

In summary, we successfully prepared and characterized AIE-active liquid crystal material based on 1,3,5-triazine-2,4,6-tricarboxamide. On the basis of the results obtained, the material was weakly emissive in solution but emit efficiently in the aggregate state, and formed columnar hexagonal LC phase with focal conic-fan shape texture in addition the molecule exhibit high thermal stability.

By combining the advantages of aggregation induce emission and liquid crystal properties such as long-range self-assembly and charge carrier mobility, make this material good candidate for optoelectronic applications.

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