## **CHAPTER 5 CONCLUSIONS AND SUGGESTIONS FOR FUTURE WORKS**

## **5.1 Conclusions**

Three Schiff bases, H<sub>2</sub>L1, H<sub>2</sub>L2 and H<sub>2</sub>L3 were successfully prepared from the reactions between 2-hydroxybenzaldehyde and 1,8-diaminooctane, 1,10-diaminodecane and 1,3-diaminopentane, respectively. These Schiff bases reacted with  $[M(CH_3(CH_2)_{14}COO)_2]$  (M = Cu, Ni, Co, Fe; R = to form 16 complexes showing differences in structures and properties (magnetic, thermal, and mesomorphic).

 $H_2L1$  formed mononuclear square planar or distorted square complexes with Cu(II), Ni(II) and Co(II) (general formula, [M(L1)]), and formed dinuclear octahedral complex with Fe(II), [Fe<sub>2</sub>(CH<sub>3</sub>(CH<sub>2</sub>)<sub>14</sub>COO)<sub>2</sub>(L1)(H<sub>2</sub>O)<sub>4</sub>].2.5H<sub>2</sub>O. All complexes, except Ni(II), were paramagnetic. Their decomposition temperatures ranged from 198°C to 313°C. However, these complexes did not exhibit liquid crystal properties.

H<sub>2</sub>L2 formed mononuclear Cu(II) complexes (general formula [M(L2)] by both one-pot and step-wise methods. The complexes were square planar, magnetic and thermally stable. For Ni(II), the step-wise method formed a mononuclear complex, [Ni(L2)].CH<sub>3</sub>CH<sub>2</sub>OH, while the one-pot method formed a dinuclear Ni(II) complex,  $[Ni_2(L2)(HL2)_2]$ .H<sub>2</sub>O. Both complexes square planar (hence diamagnetic). The mononuclear complex has a higher thermal stability than the dinuclear complex.For Co(II), method formed dinuclear Co(II) the step-wise a complex, [Co<sub>2</sub>(L2)(HL2)<sub>2</sub>(EtOH)<sub>4</sub>], while one-pot method formed a mononuclear Co(III) complex, [Co(L2)(HL2)]. The geometry of Co(II) and Co(III) was octahedral. The thermal stability of both complexes were almost similar ( $T_{dec} = 205^{\circ}C$ ). The dinuclear complex was paramagnetic while the mononuclear complex was diamagnetic.For Fe(II), both methods formed dinuclear mixed-valence complexes in good yields. The difference in the two complexes was the presence of coordinated and non-coordinated solvent. The magnetic susceptibility and thermal stability for the complex formed from the one-pot method were slightly higher than from the step-wise method. All of these complexes did not exhibit liquid crystal properties

H<sub>2</sub>L3 formeda dinuclearCu(II) complex,  $[Cu_2(CH_3(CH_2)_{14}COO)(L3)(HL_3)]$ , and mononuclear complexes with Ni(III), Co(II) and Fe(III) complexes. These complexes were paramagnetic, with thermal decomposition temperatures in the range of 142 °C to 209 °C. However, these complexes also did not have liquid crystal properties.

## **5.2 Suggestions for Future Works**

All hexadecanoato-Schiff base complexes were not mesomorphic. However, they may be made to be mesogenicby reacting them with neutral N-donor amide ligands, such as shown in **Figure 5.1**.

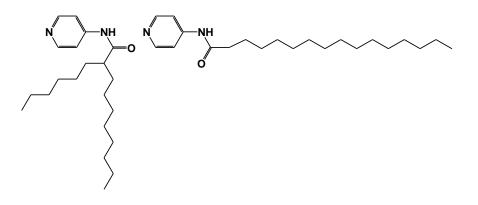


Figure 5.1Structural formulae of neutral N-donor ligands

The thermal SCO behavior of cobalt(II) and iron(II) complexes should be further probed using SQUID magnetometer on both heating and cooling cycles. It would also be interesting to know the thermoelectric behavior of Co(III) and mixed-valence Fe complexes by measuring their electronic conductivities at different temperatures.Lastly, the potential uses of these complexes, especially Co(II), Fe(II), as catalysts and dyesensitised cell materials, should be explored.