



Supporting Information

for

Synthesis and investigation on optical and electrochemical properties of 2,4-diaryl-9-chloro-5,6,7,8-tetrahydroacridines

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Synthesis and analytical data of starting compound 2 and copies of spectra for the synthesized compounds

1. Procedure for preparing the starting material 2,4-dibromo-9-chloro-5,6,7,8-tetrahydroacridine (2)

3,5-Dibromoanthranilic acid (2.92 g, 10 mmol, 1 equiv) and cyclohexanone (1.07 mL, 11 mmol, 1.1 equiv) were stirred in an ice bath. Then, 15 mL of POCl_3 were carefully added and the mixture was heated under reflux for 4 h. The mixture was cooled to room temperature and concentrated to give a slurry. The residue was diluted with dichloromethane, neutralized with aqueous NaHCO_3 , and washed with brine. The organic layer was dried over anhydrous K_2CO_3 and concentrated to afford a yellow solid. It was recrystallized from acetone to give **2** as pale yellow solid (3.24 g, 87%).

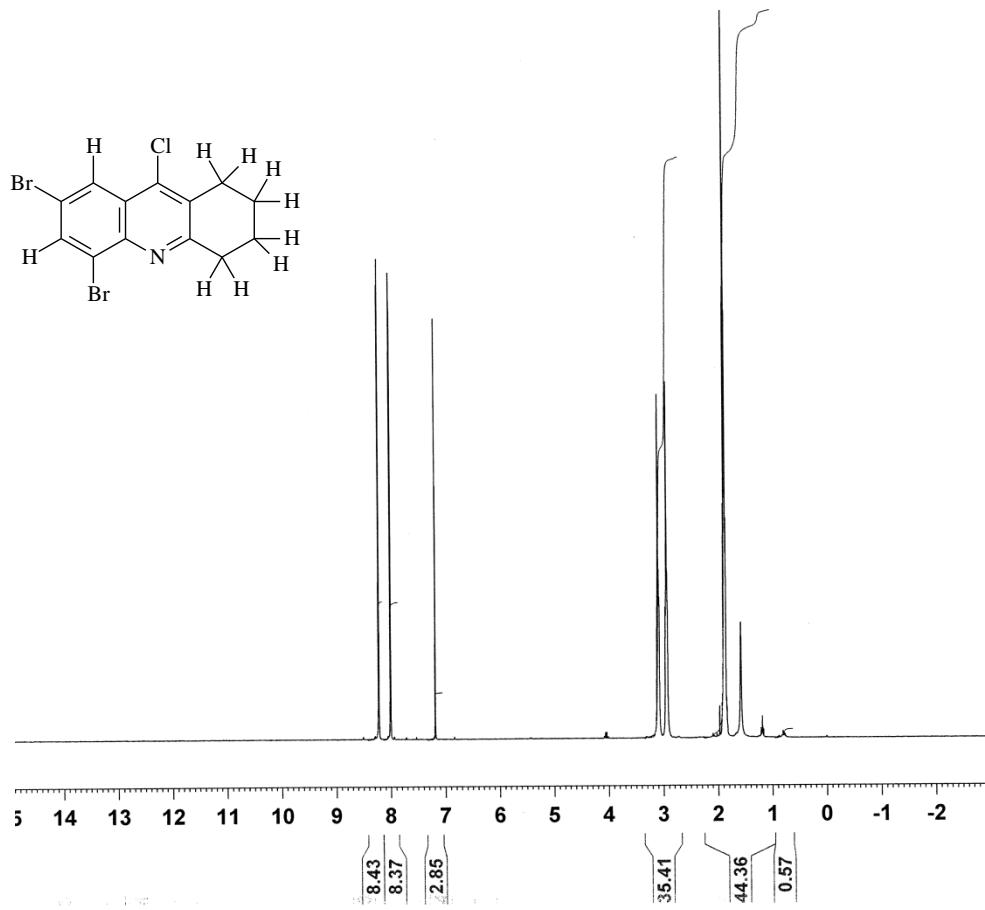


Figure S1: ^1H NMR (300 MHz, CDCl_3) of 2,4-dibromo-9-chloro-5,6,7,8-tetrahydroacridine (**2**).

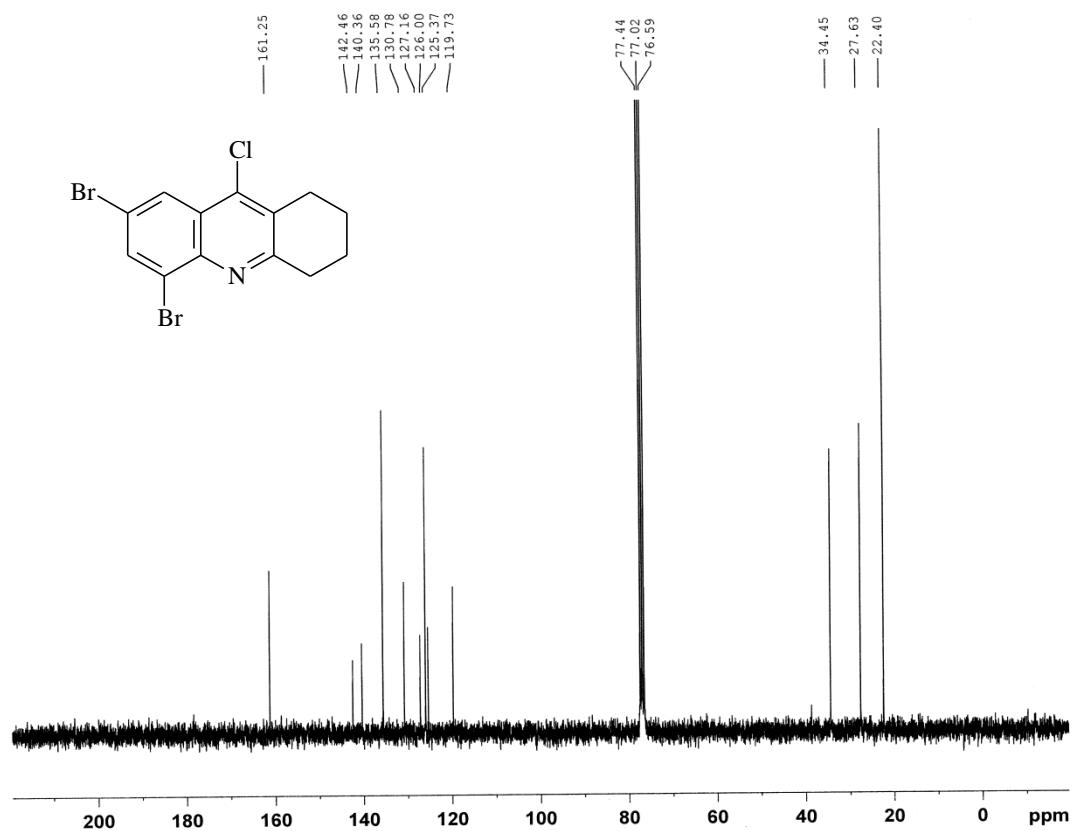


Figure S2: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-dibromo-9-chloro-5,6,7,8-tetrahydroacridine (**2**).

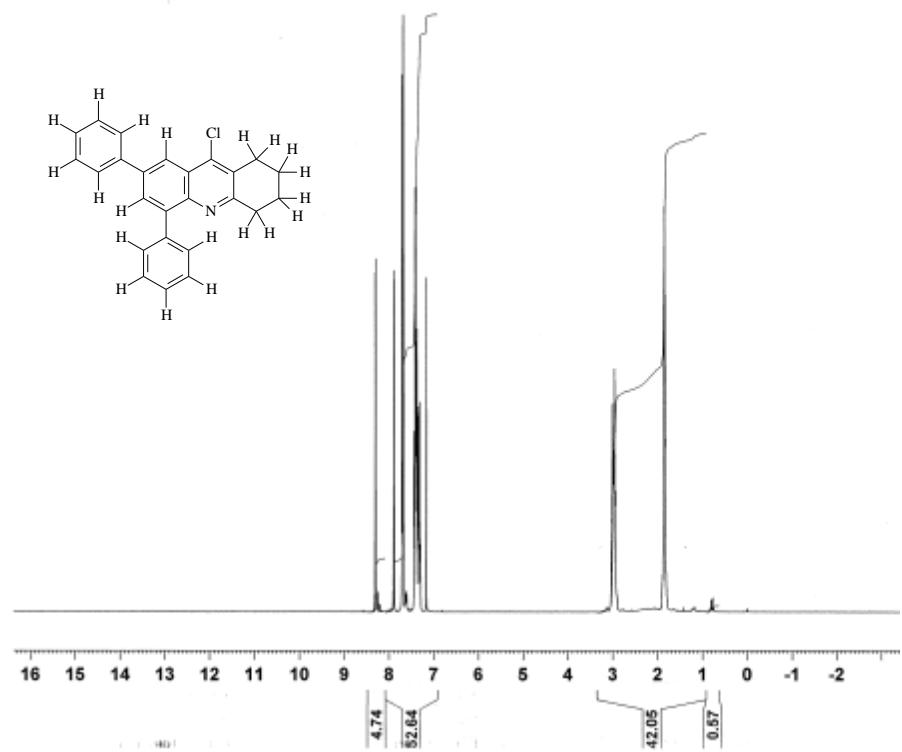


Figure S3: ¹H NMR (300 MHz, CDCl₃) of 2,4-diphenyl-9-chloro-5,6,7,8-tetrahydroacridine (**4a**)

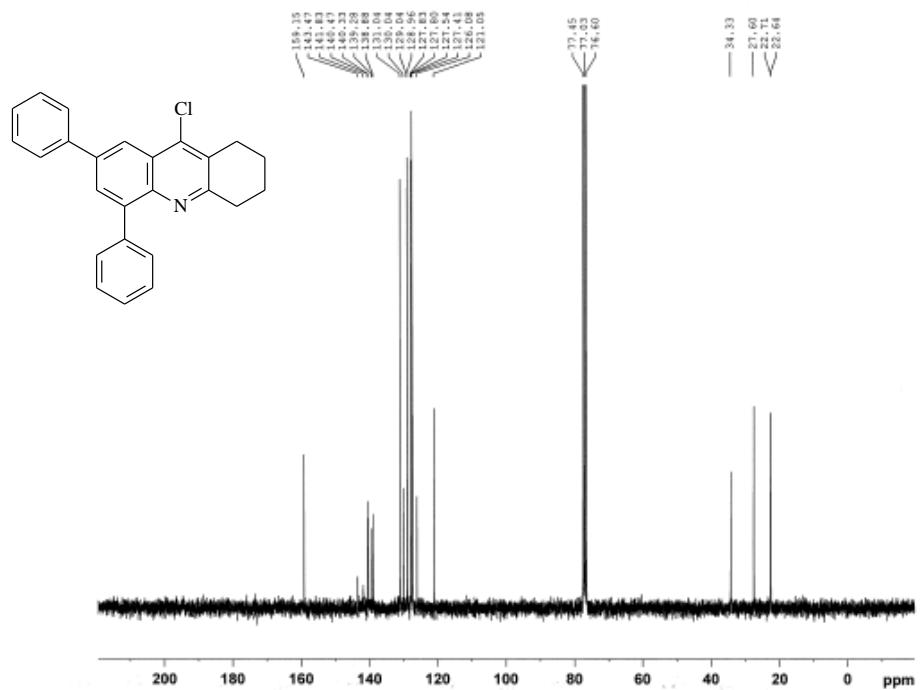


Figure S4: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-diphenyl-9-chloro-5,6,7,8-tetrahydroacridine (**4a**)

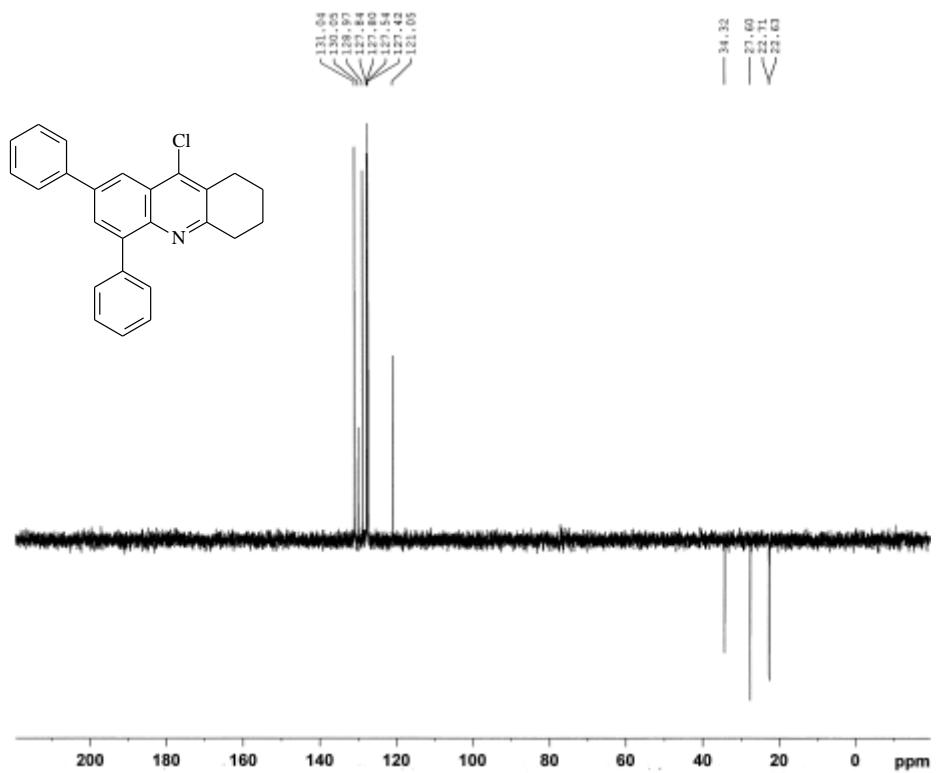
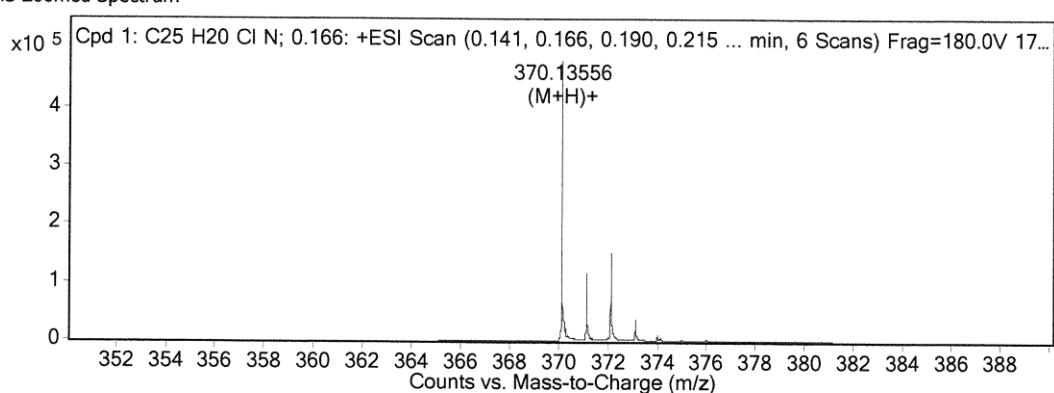


Figure S5: ^{13}C NMR DEPT 135 spectrum of 2,4-diphenyl-9-chloro-5,6,7,8-tetrahydroacridine (**4a**)

MS Zoomed Spectrum



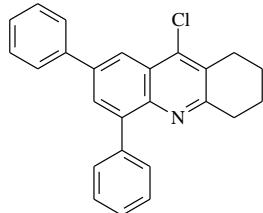
MS Spectrum Peak List

Ion	Abund	Formula	Calculated Mass	Measured Mass	Difference	Diff (ppm)
(M+H)+	485856.59	C25H20ClN	370.1357	370.13556	0.14	0.38

--- End Of Report ---

Figure S6: MS spectra of of 2,4-diphenyl-9-chloro-5,6,7,8-tetrahydroacridine (**4a**)

calculated mass: 369.1357



measured mass: 369.1355

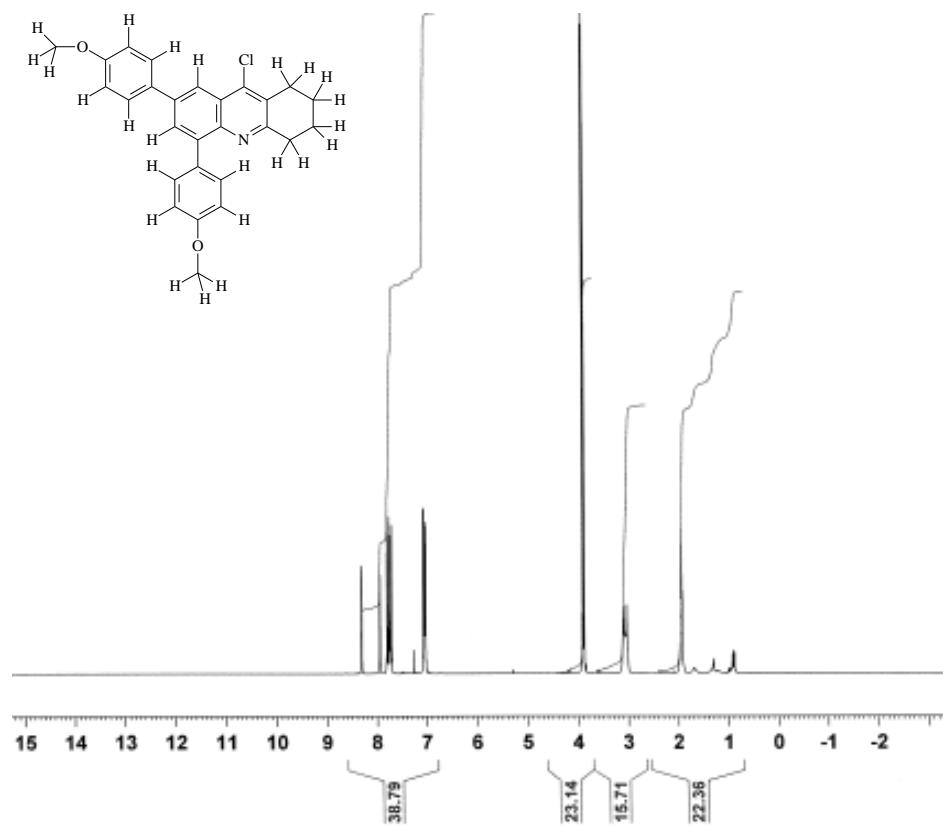


Figure S7: ¹H NMR (300 MHz, CDCl₃) of 2,4-bis(4-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4b**)

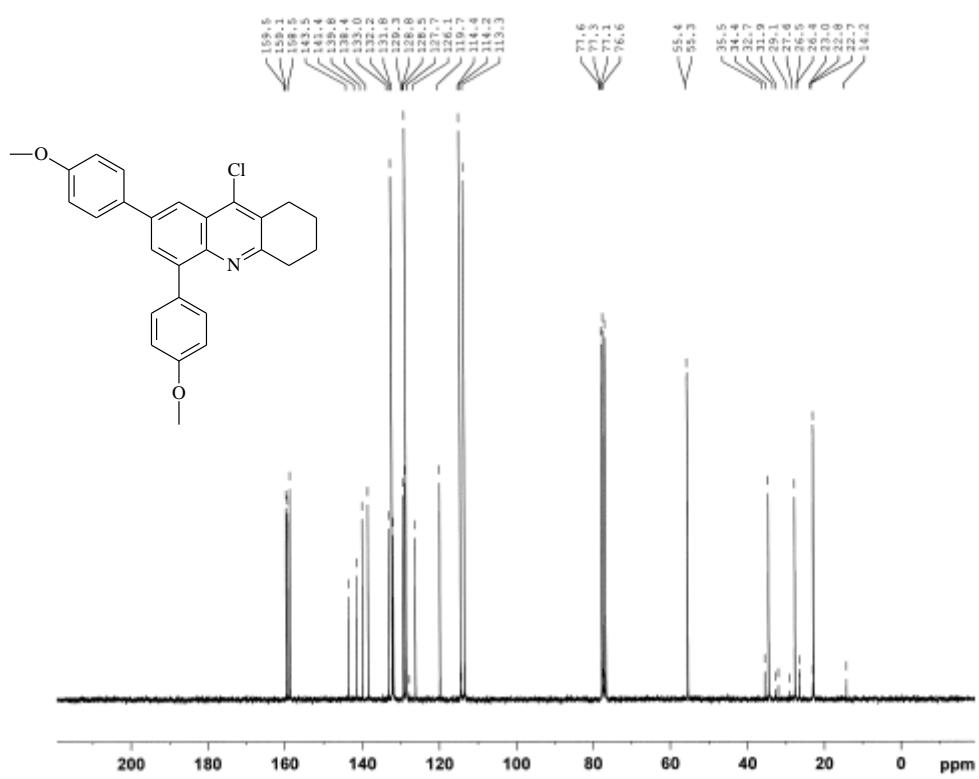


Figure S8: ¹³C NMR (75 MHz, CDCl₃) of 2,4-bis(4-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4b**)

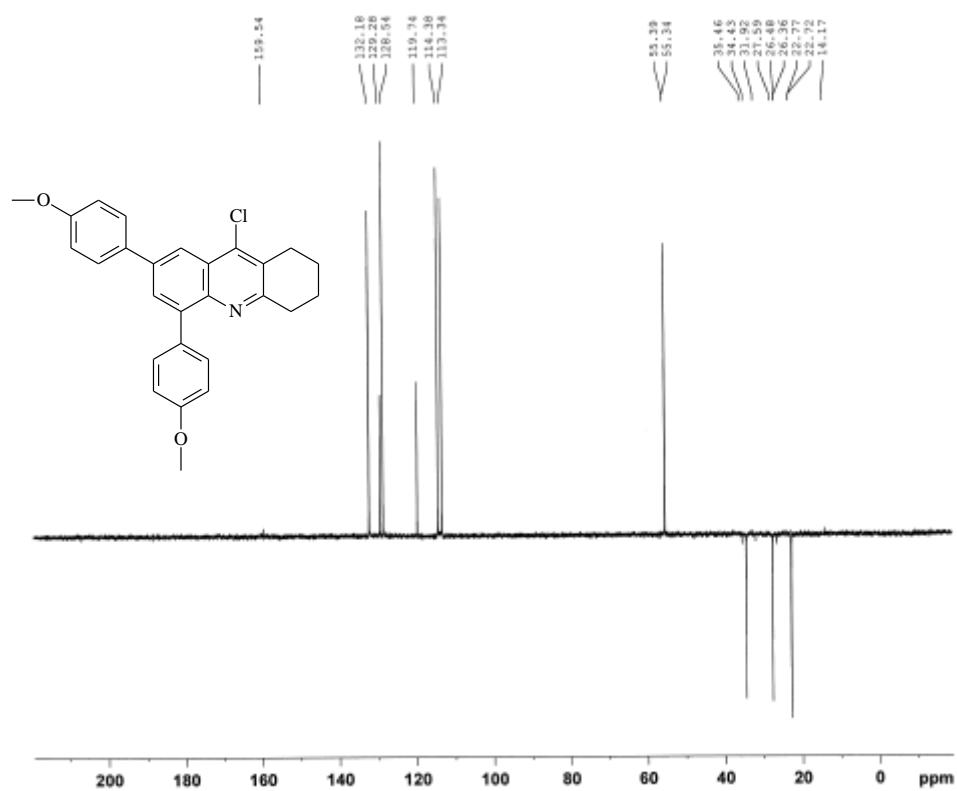


Figure S9: ^{13}C NMR DEPT 135 spectrum of 2,4-bis(4-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4b**)

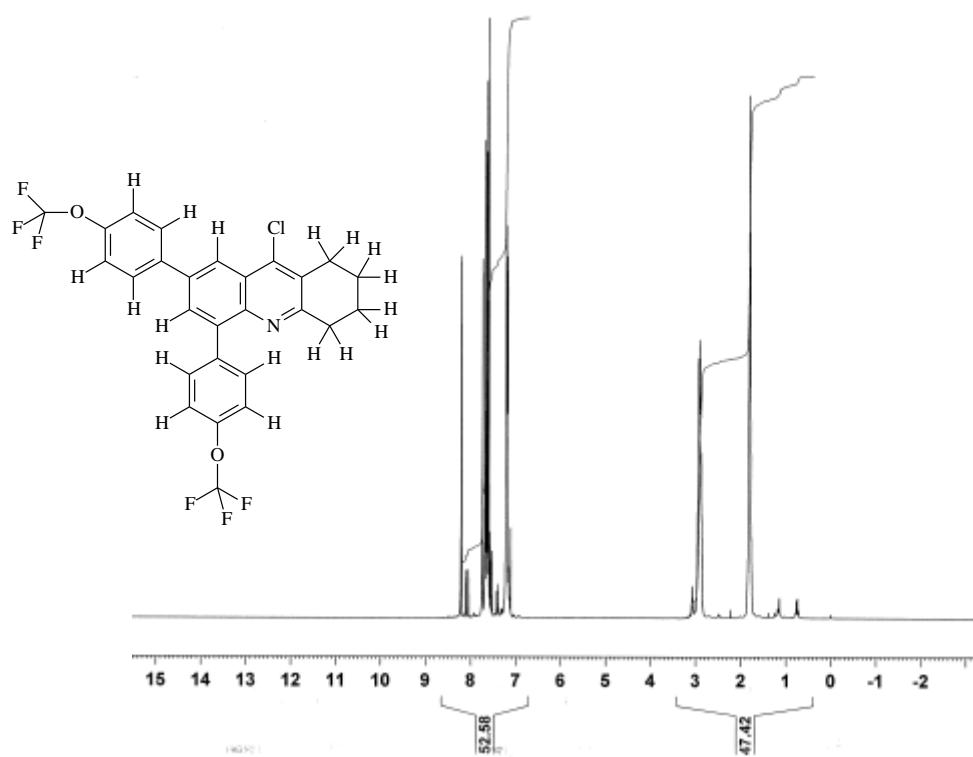


Figure S10: ^1H NMR (300 MHz, CDCl_3) of 2,4-bis(4-trifluoromethoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4c**)

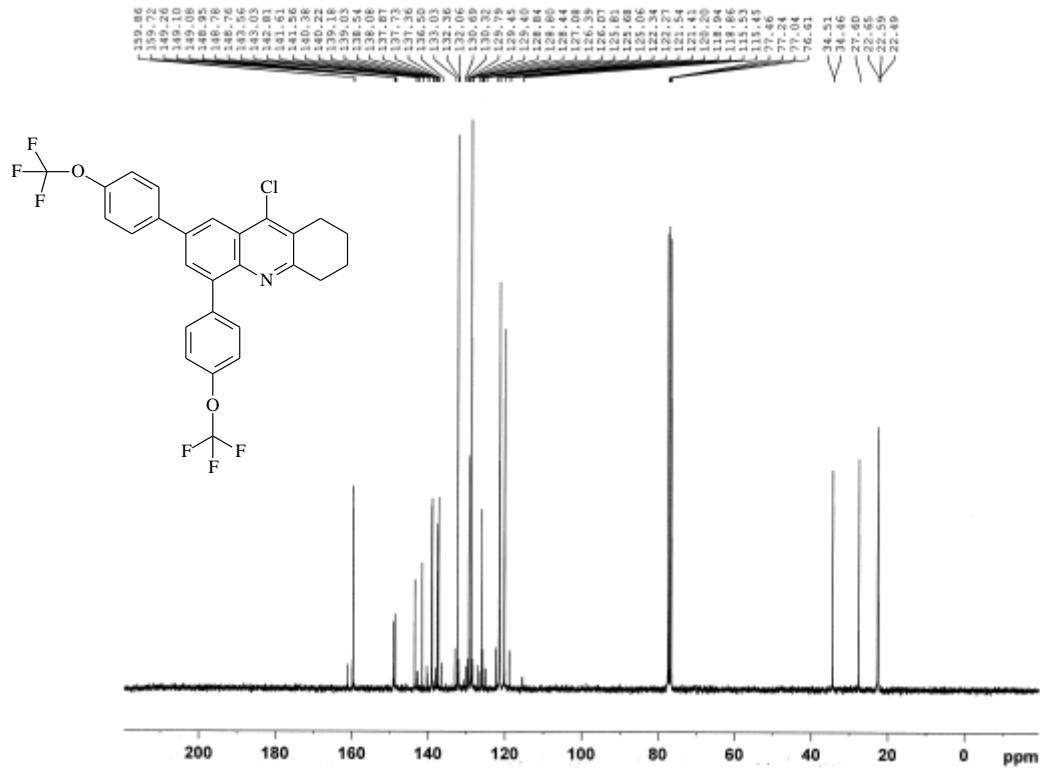


Figure S11: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-bis(4-trifluoromethoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4c**)

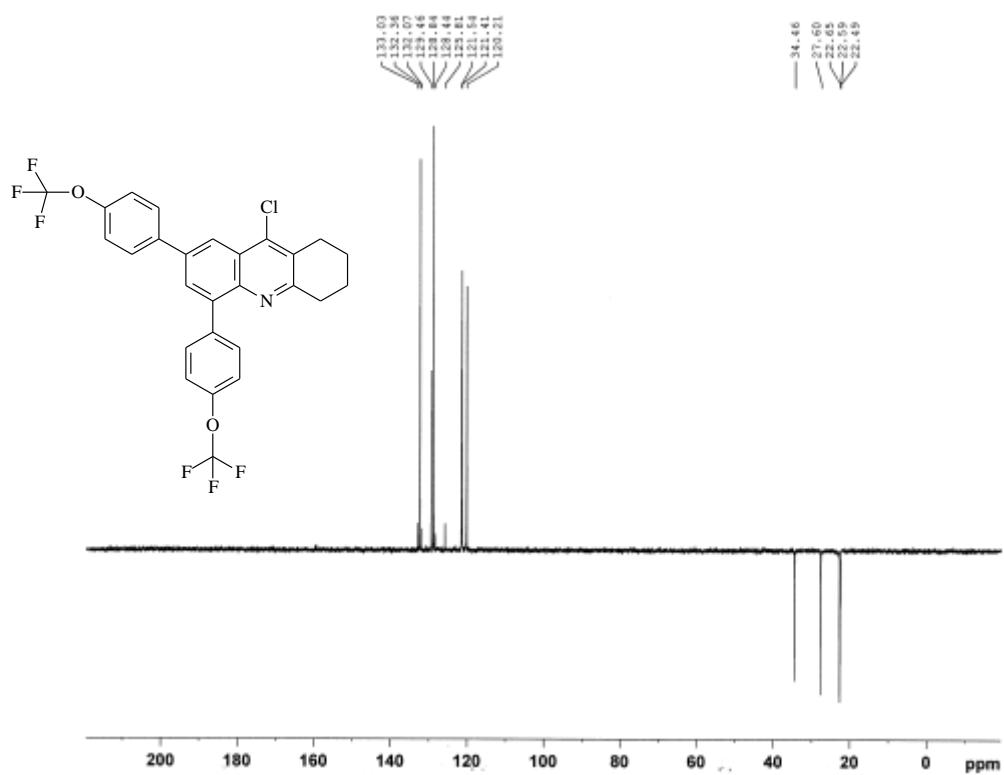


Figure S12: ^{13}C NMR DEPT 135 spectrum of 2,4-bis(4-trifluoromethoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4c**)

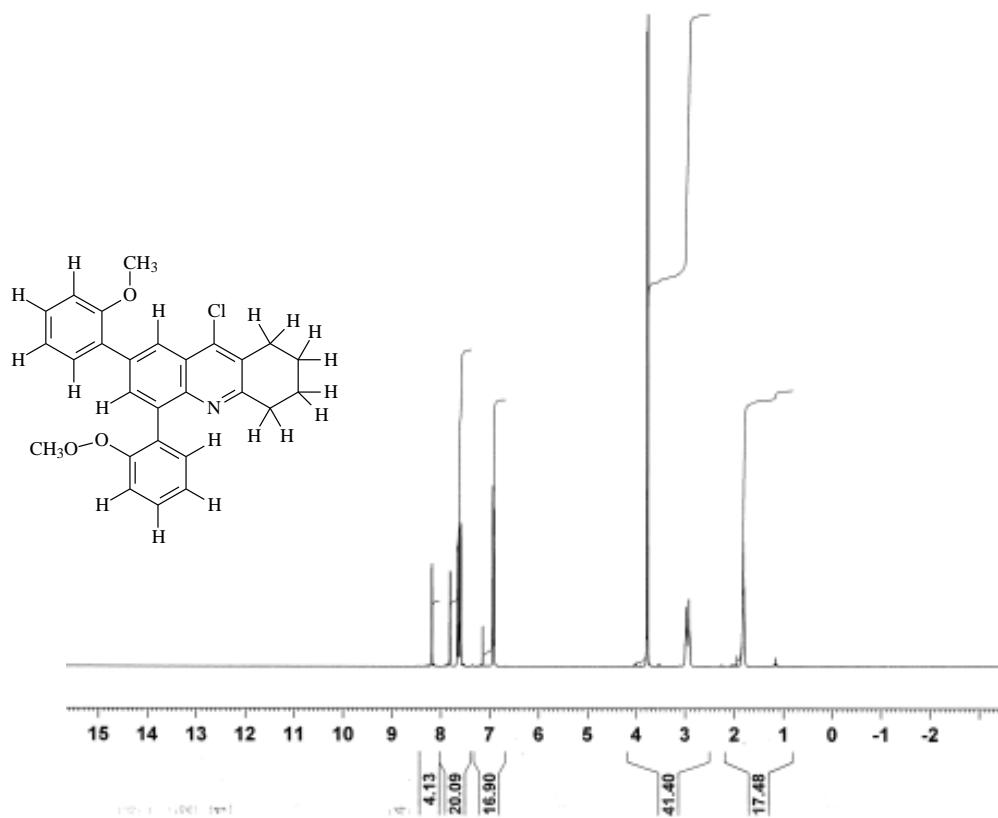


Figure S13: ^1H NMR (300 MHz, CDCl_3) of 2,4-bis(2-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4d**)

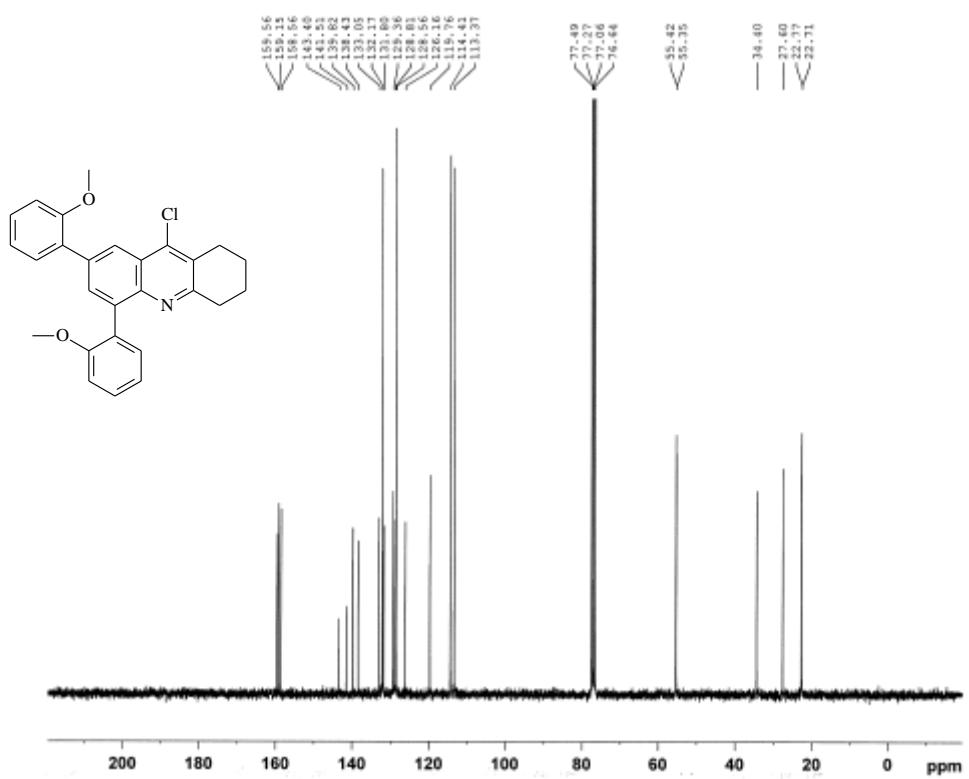


Figure S14: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-bis(2-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4d**)

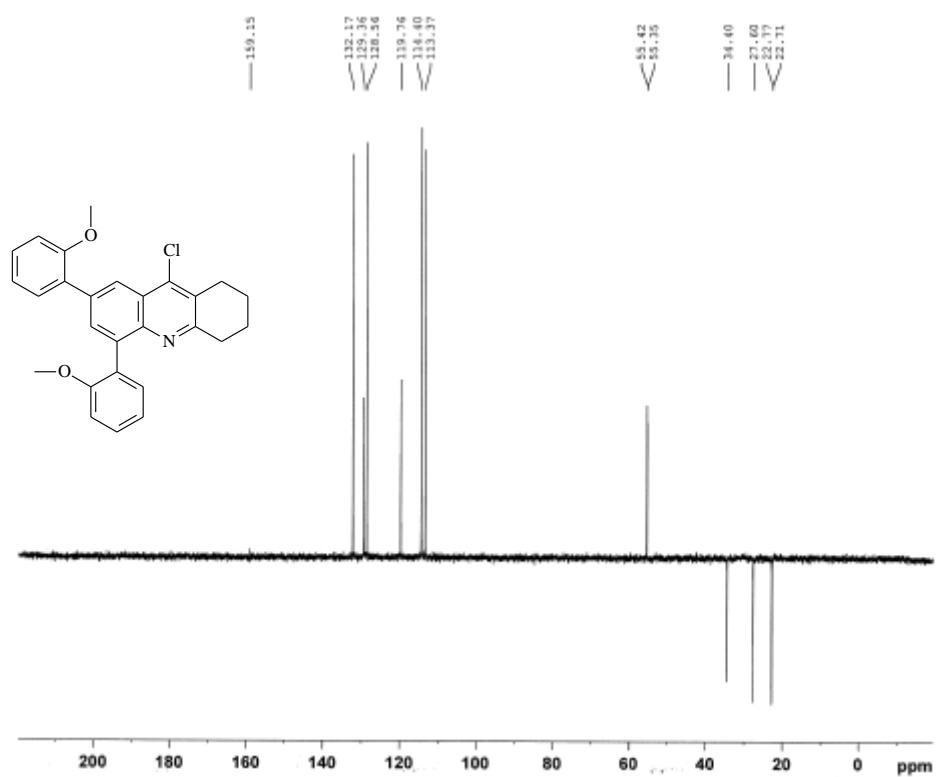


Figure S15: ¹³C NMR DEPT 135 of 2,4-bis(2-methoxyphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (4d)

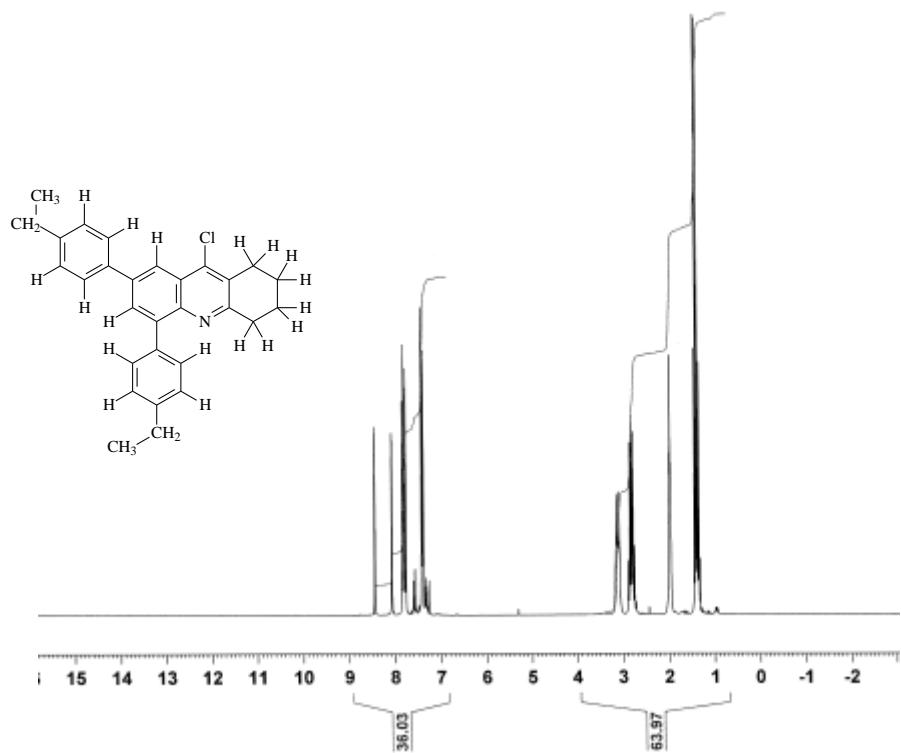


Figure S16: ^1H NMR (300 MHz, CDCl_3) of 2,4-bis(4-ethylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4e**)

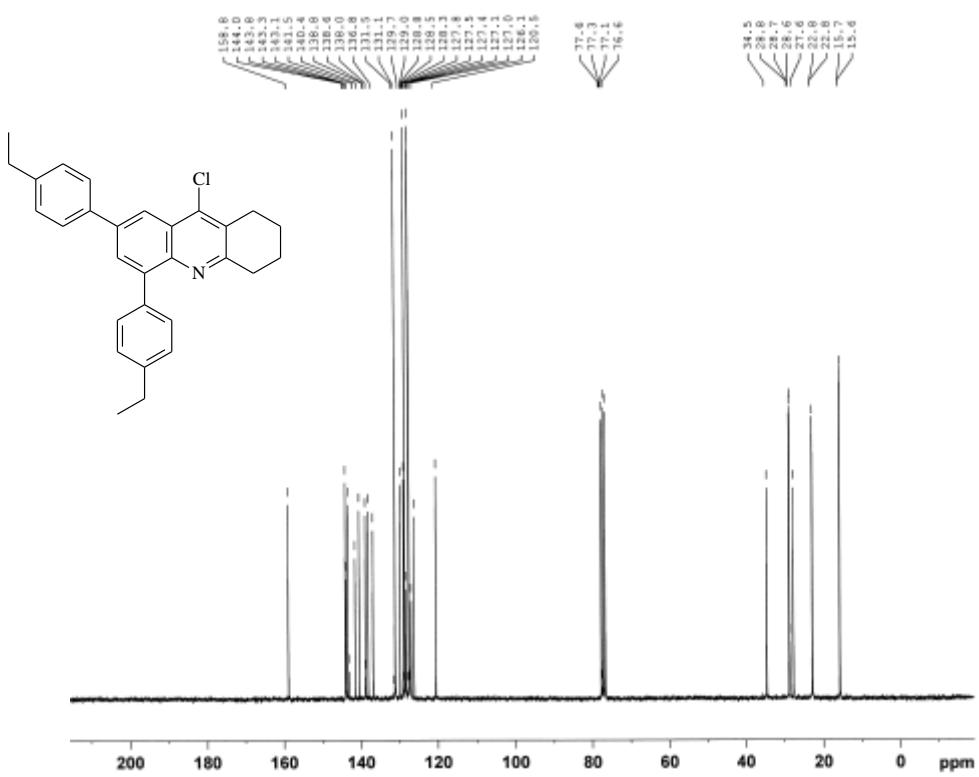


Figure S17: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-bis(4-ethylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (4e)

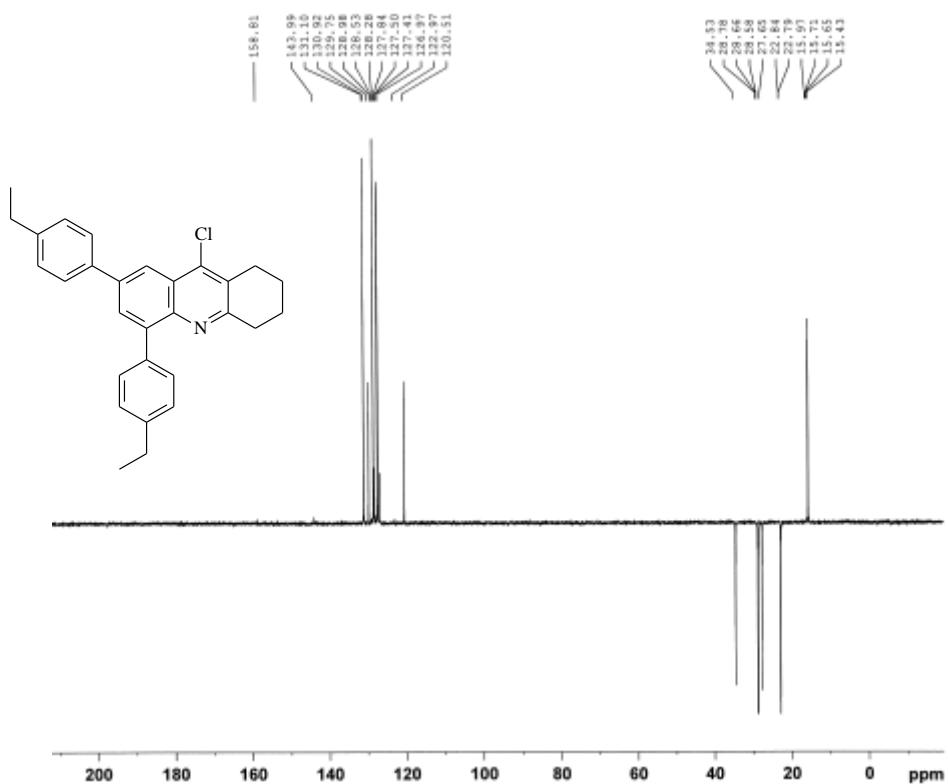


Figure S18: ^{13}C NMR DEPT 135 spectrum of 2,4-bis(4-ethylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4e**)

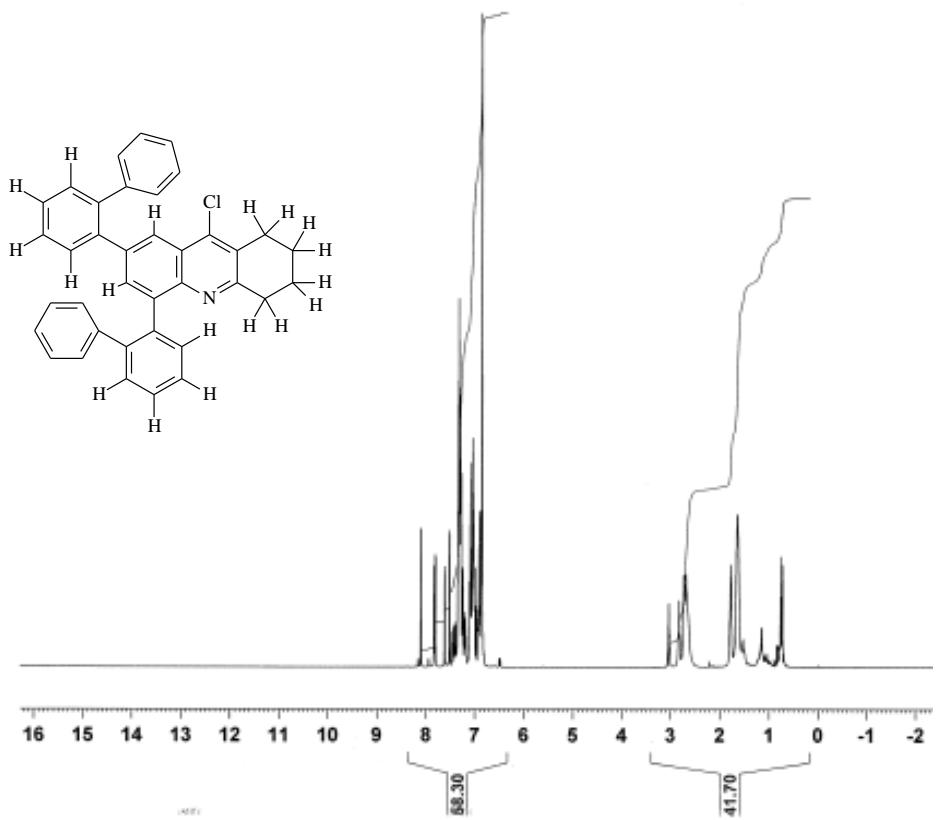


Figure S19: ^1H NMR (300 MHz, CDCl_3) of 2,4-bis(2-phenylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4f**)

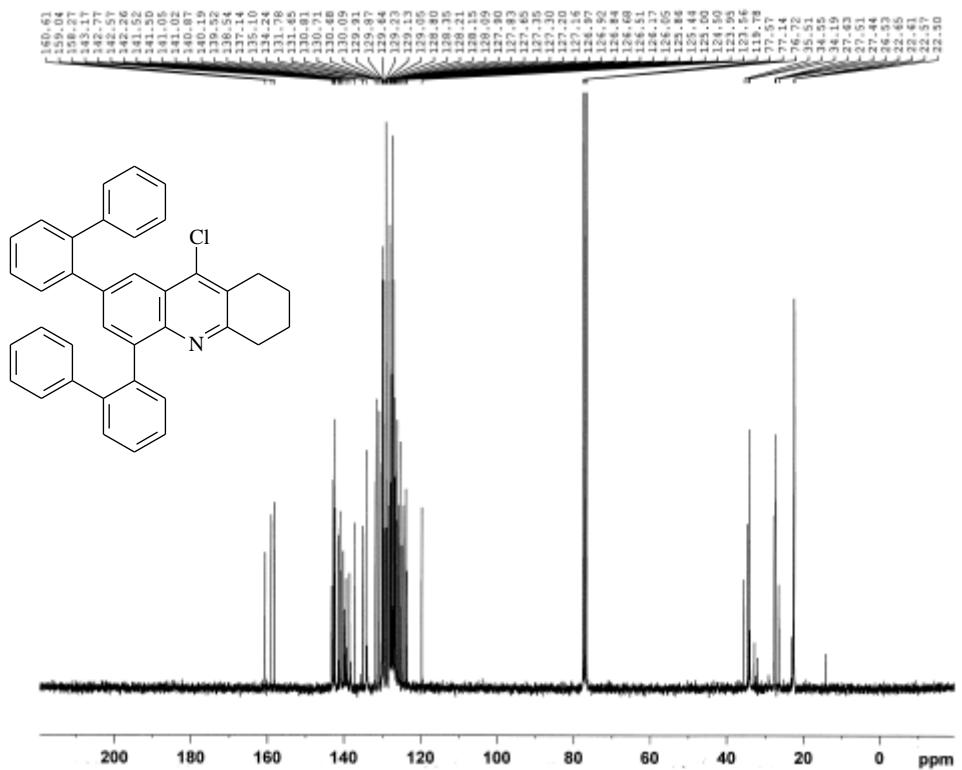


Figure S20: ^{13}C NMR (75 MHz, CDCl_3) of 2,4-bis(2-phenylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4f**)

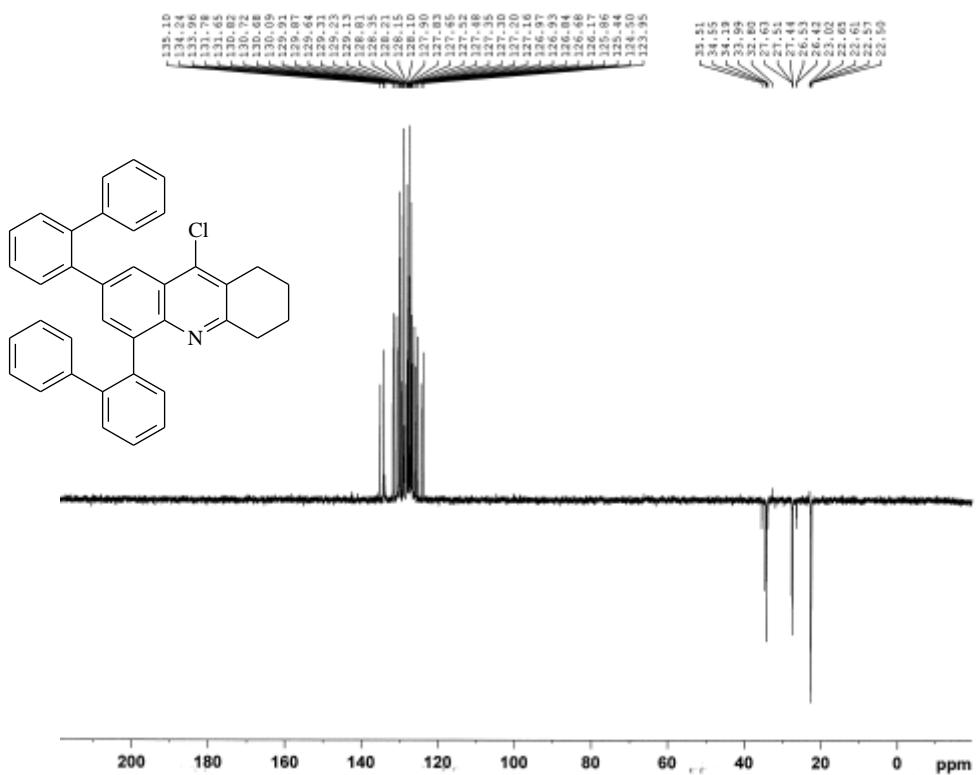


Figure S21: ^{13}C NMR DEPT 135 spectrum of 2,4-bis(2-phenylphenyl)-9-chloro-5,6,7,8-tetrahydroacridine (**4f**)