

Theoretical and Experimental Exploration of the Photochemistry of Resveratrol: Beyond the Simple Double Bond Isomerization

Roi Álvarez, Inma R. Lahoz, Olalla Nieto Faza, María-Magdalena Cid, Carlos Silva*

*Lopez**

Departamento de Química Orgánica, Facultade de Química, Edificio de Ciencias
Experimentais, Campus Lagoas-Marcosende, 36310 Vigo (Spain).

mcid@uvigo.es, carlos.silva@uvigo.es

Figure 1S. Comparison of ^1H -NMR spectra of compound **10**: 1) Obtained from photochemical irradiation of resveratrol **1**; 2) obtained from MOM-**8** after treatment with MeOH/HCl; 3 and 4) same as 2 but after longer reaction times.

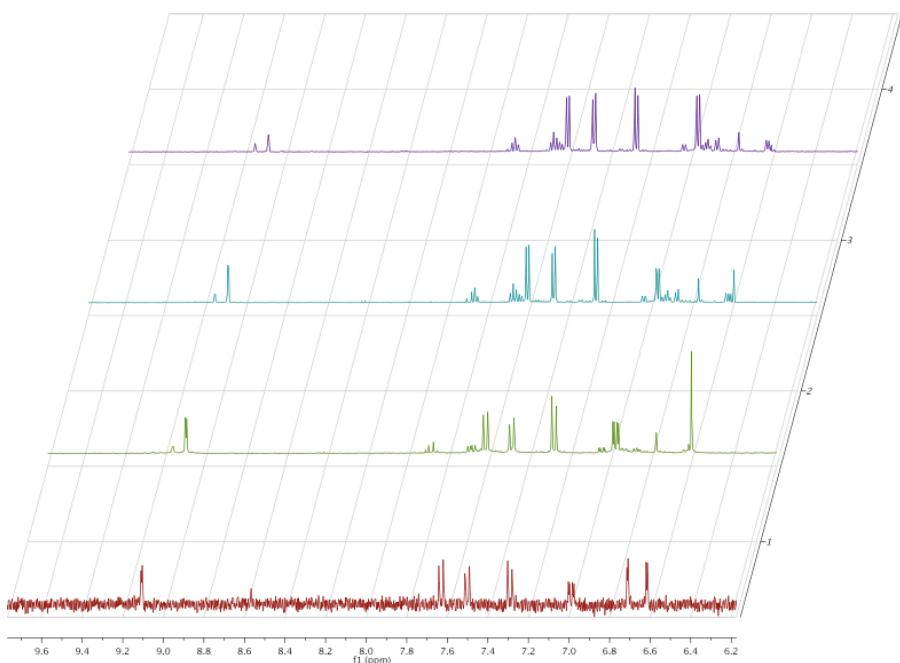
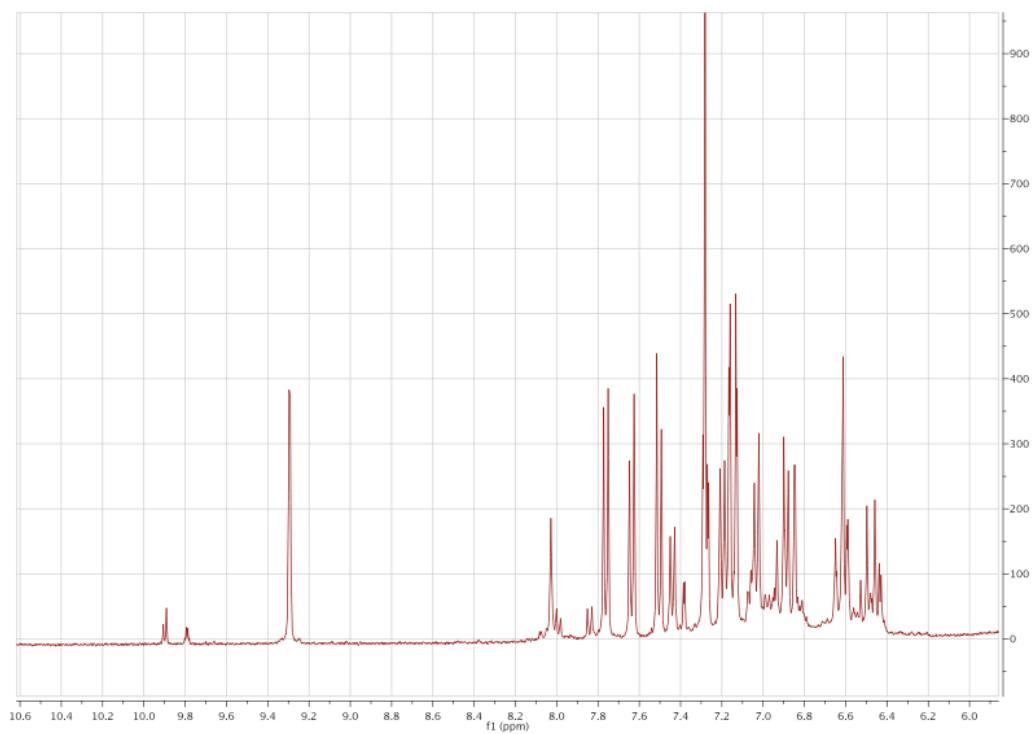
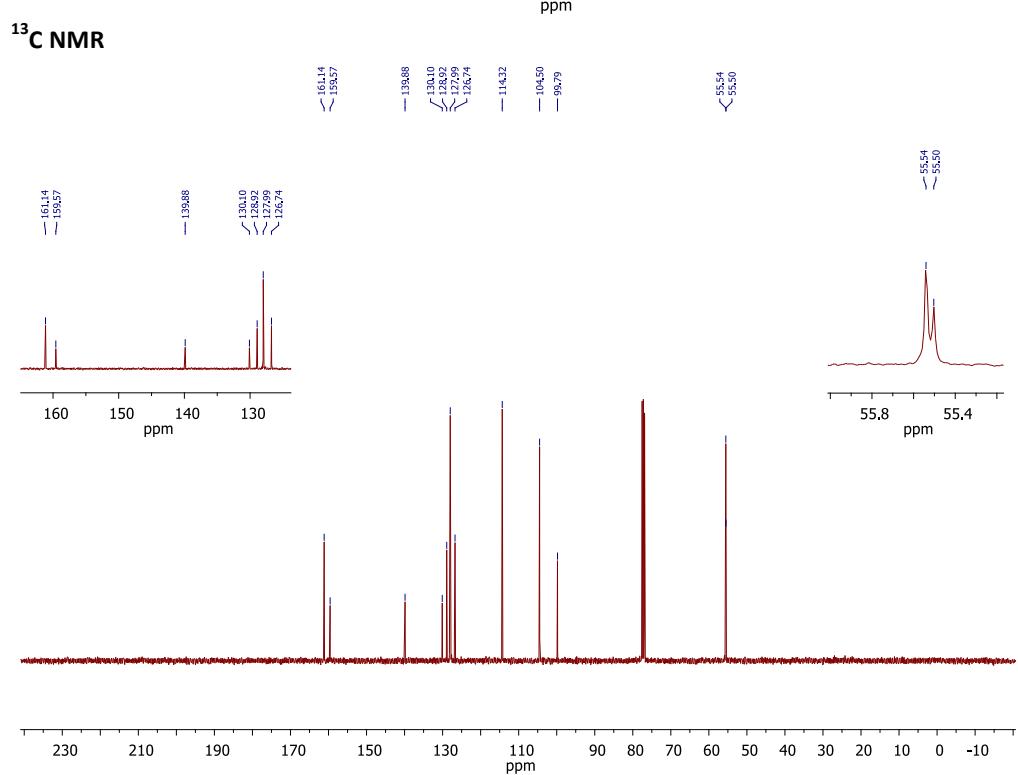
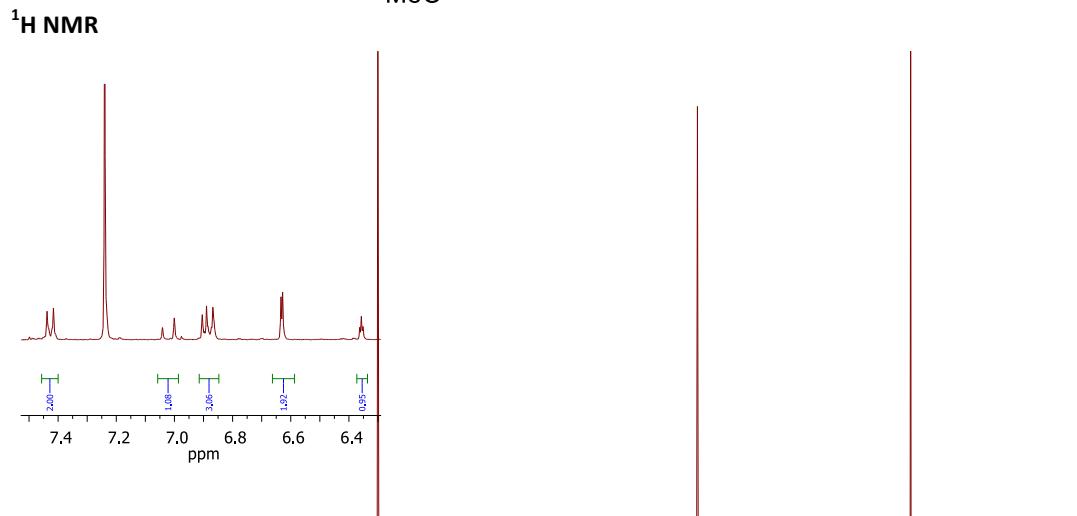
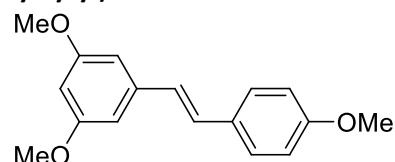


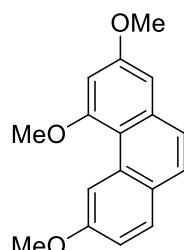
Figure 2S. ^1H NMR of EOM-**8** after 7 h of irradiation



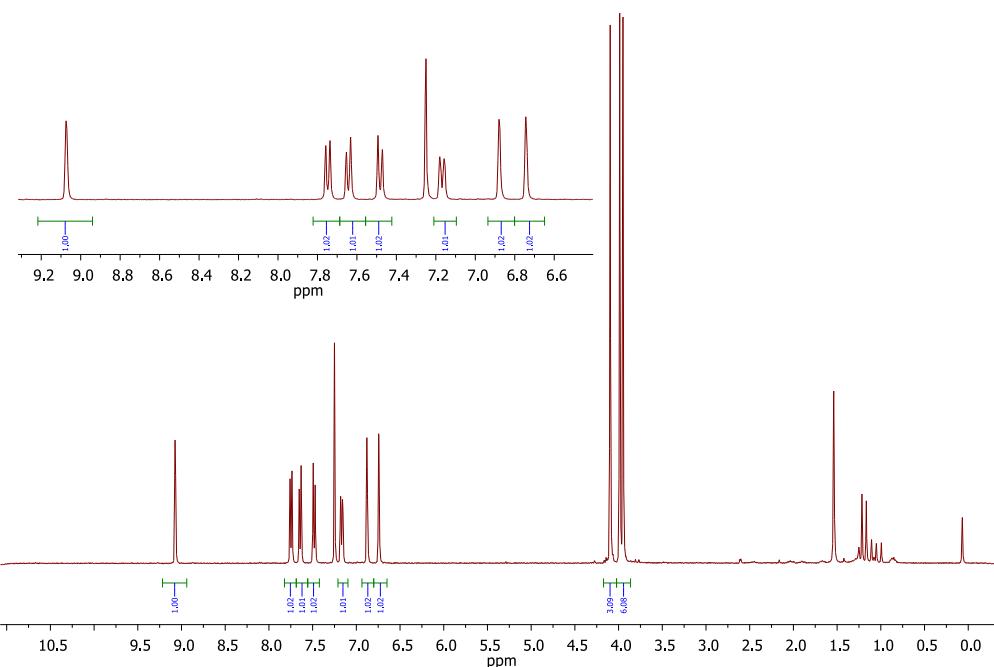
(E)-1,3-dimethoxy-5-(4-methoxystyryl)benzene



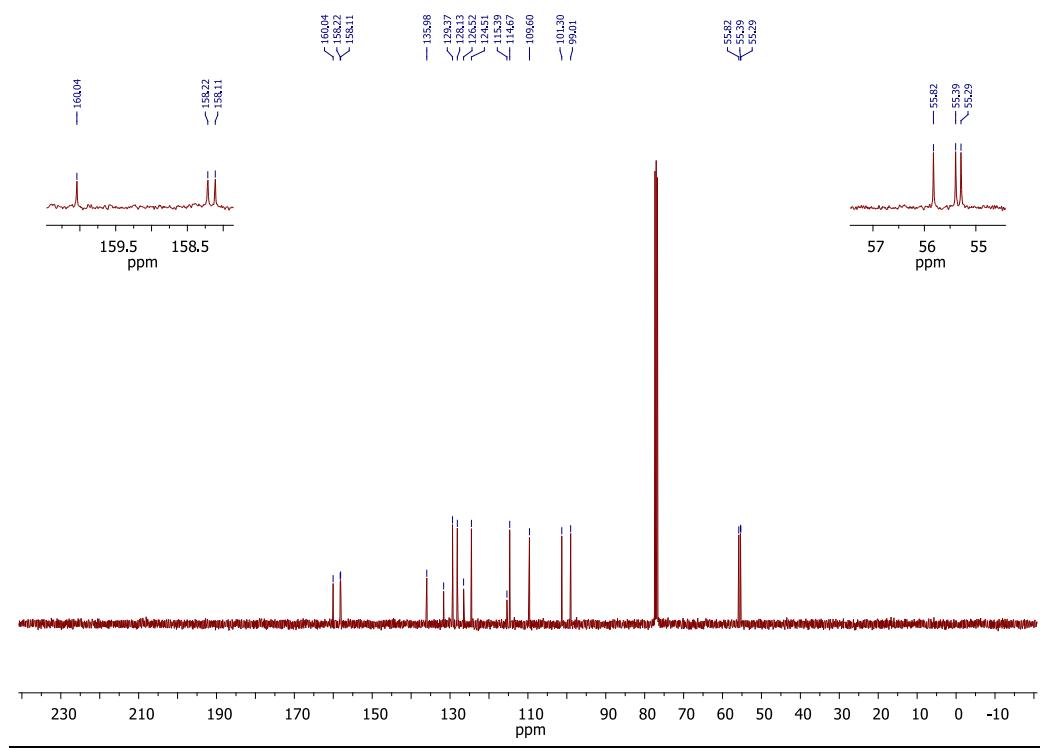
2,4,6-Trimethoxyphenanthrene



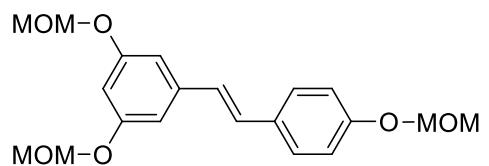
$^1\text{H-NMR}$



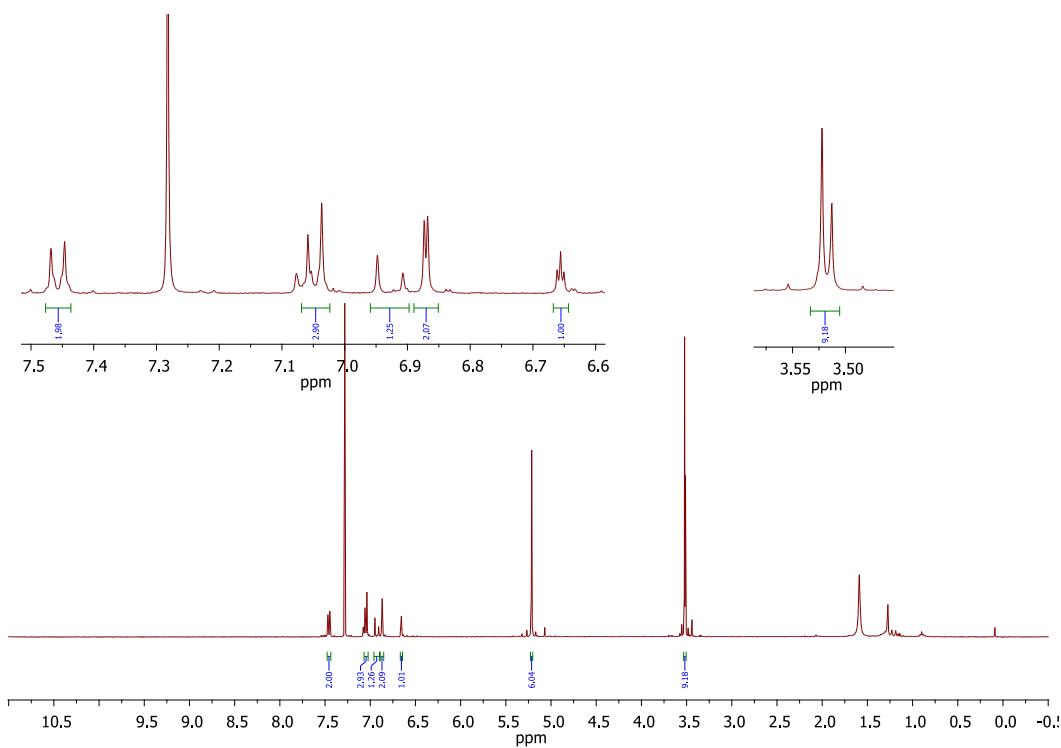
$^{13}\text{C-NMR}$:



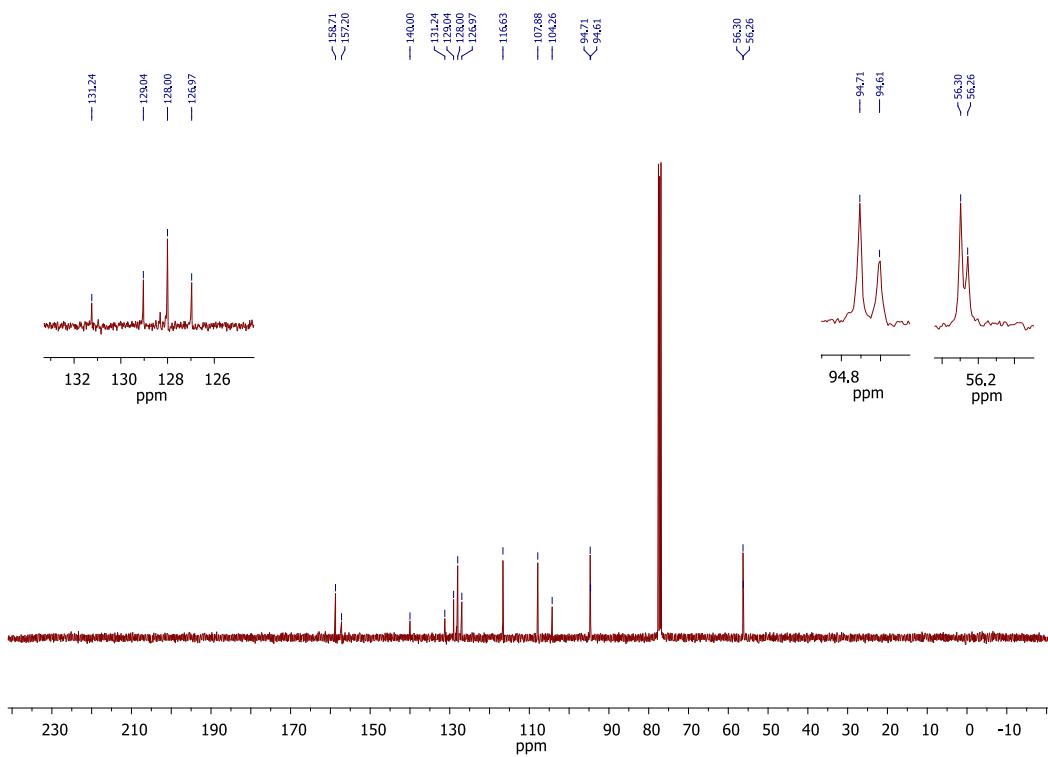
(E)-1,3-bis(Methoxymethoxy)-5-(4-methoxymethoxy)styrylbenzene.



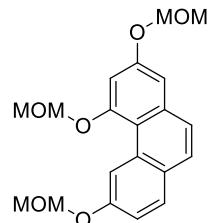
¹H NMR:



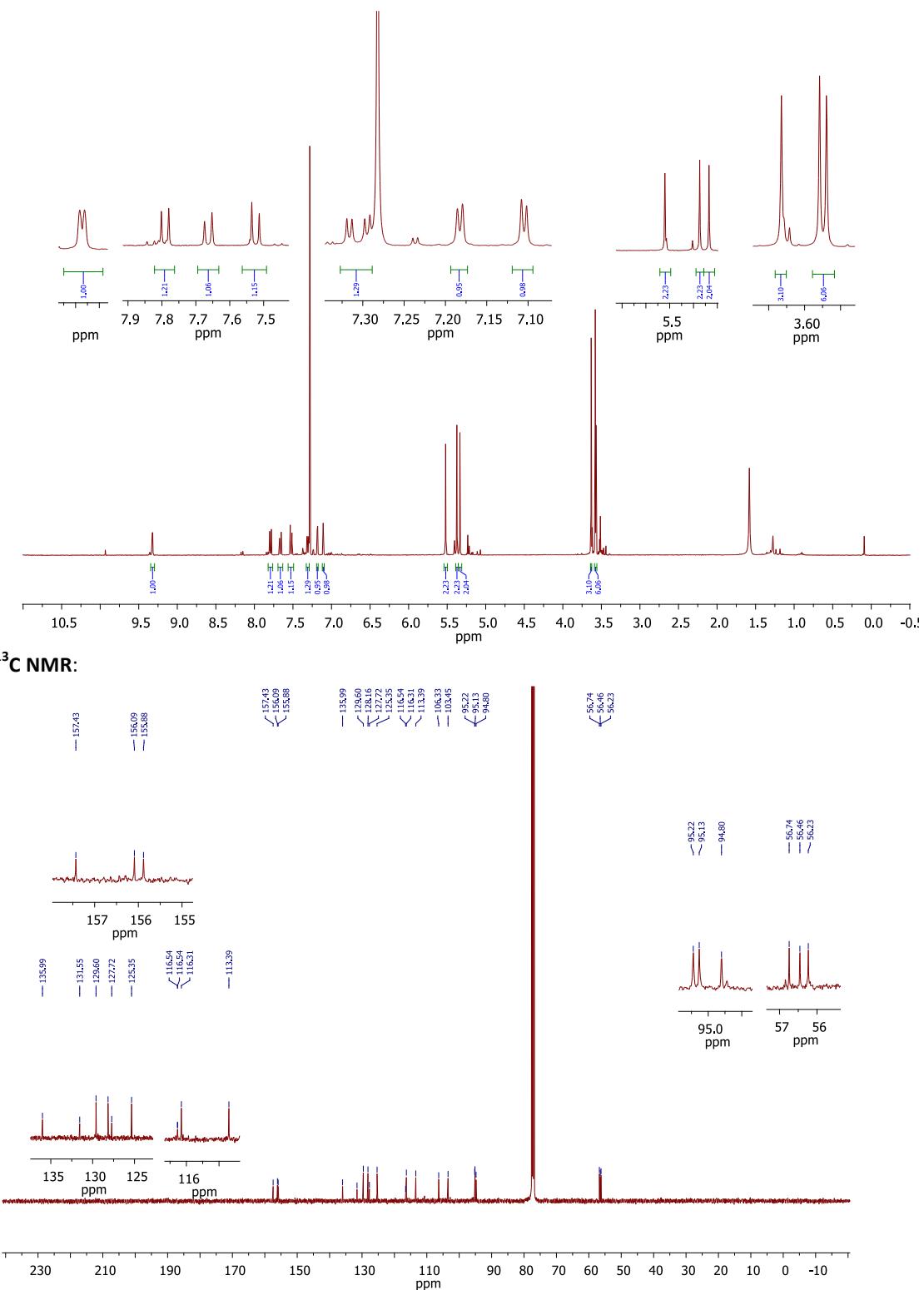
¹³C-NMR:



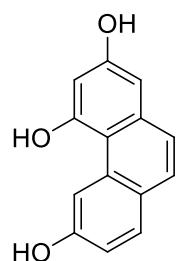
2,4,6-tris(methoxymethoxy)phenanthrene



¹H NMR:



Phenanthrene-2,4,6-triol.



¹H NMR:

