

Supplementary Information

Synthesis of N-cinnamoyl dipeptide esters and investigation of their self-assembly leading to nanorods formation

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Spectral Details of N-Cinnamoyl-Gly-Gly-OEt

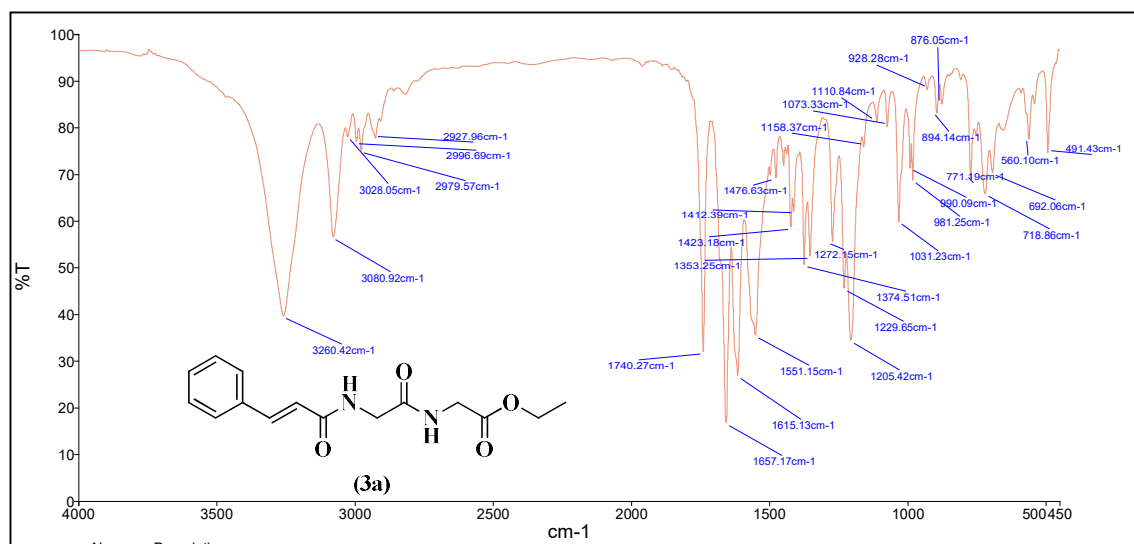


Figure 1: IR Spectrum of N-Cinnamoyl-Gly-Gly-OEt (3a)

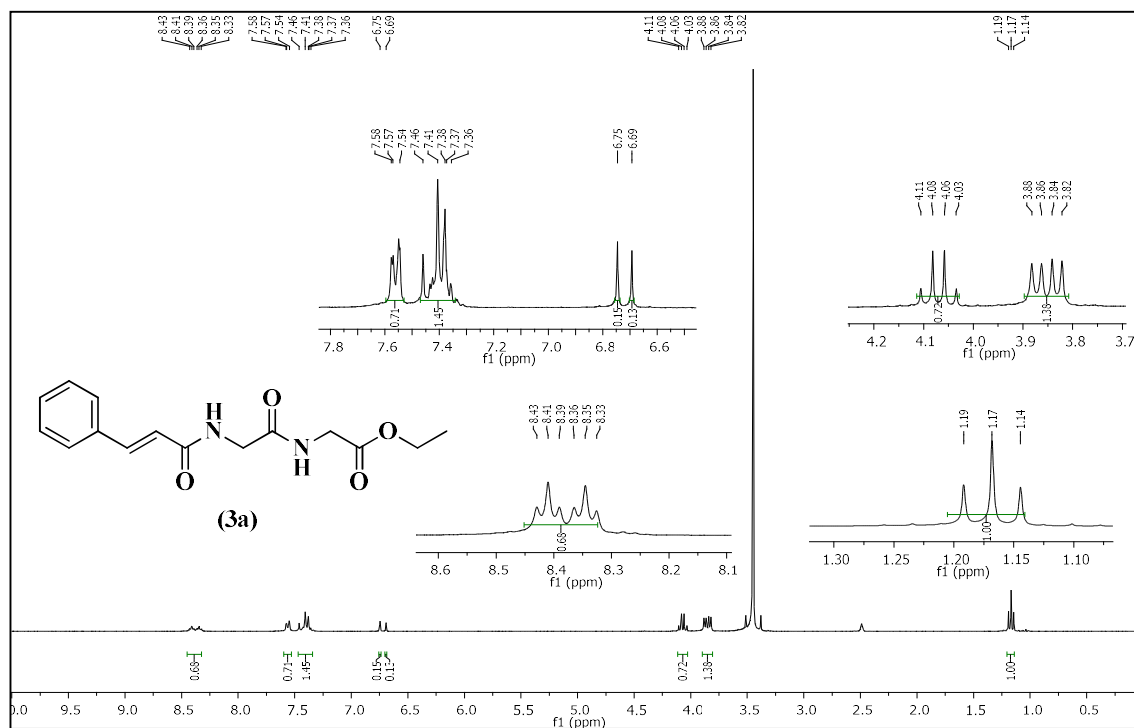


Figure 2: ¹H NMR Spectrum of N-Cinnamoyl-Gly-Gly-OEt (3a)

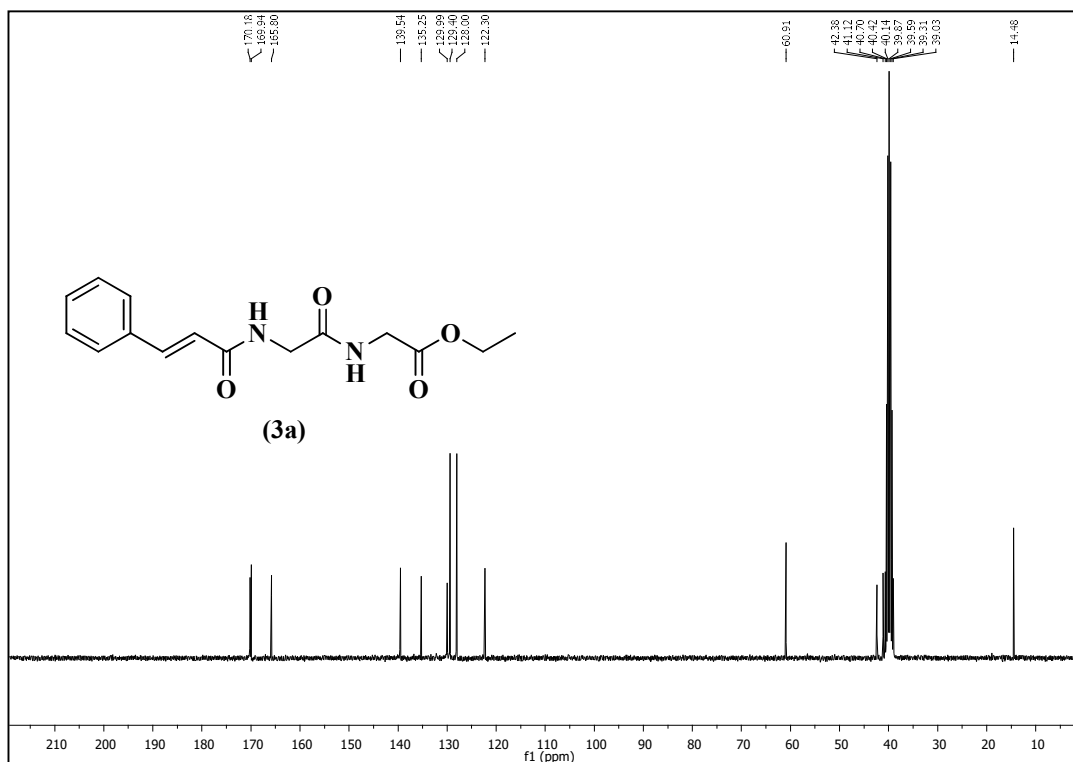


Figure 3: ¹³C NMR Spectrum of N-Cinnamoyl-Gly-Gly-OEt (**3a**)

Spectral Details of N-Cinnamoyl-Ala-Gly-OEt

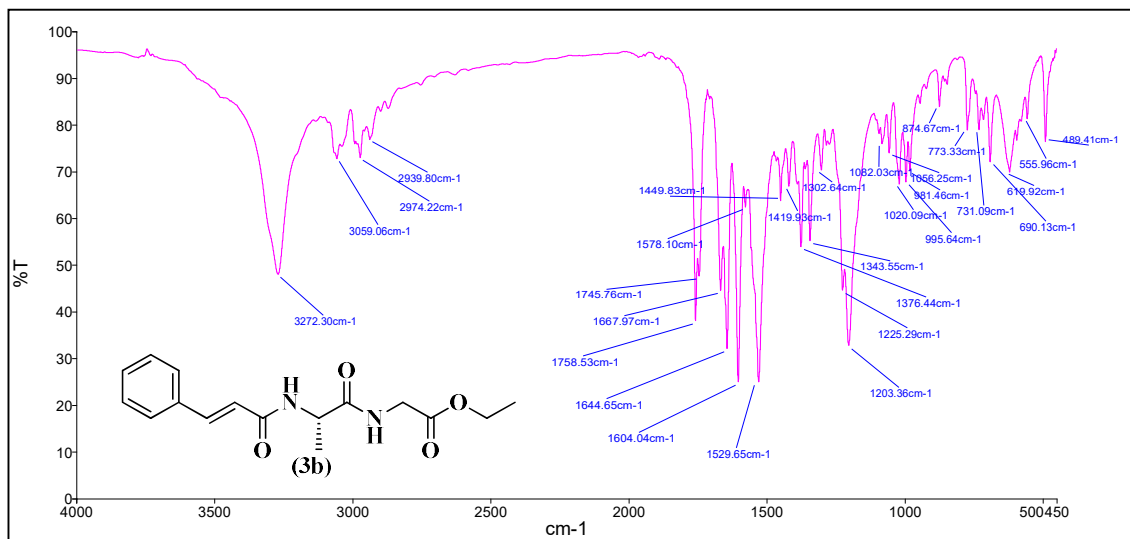


Figure 4: IR Spectrum of N-Cinnamoyl-Ala-Gly-OEt (**3b**)

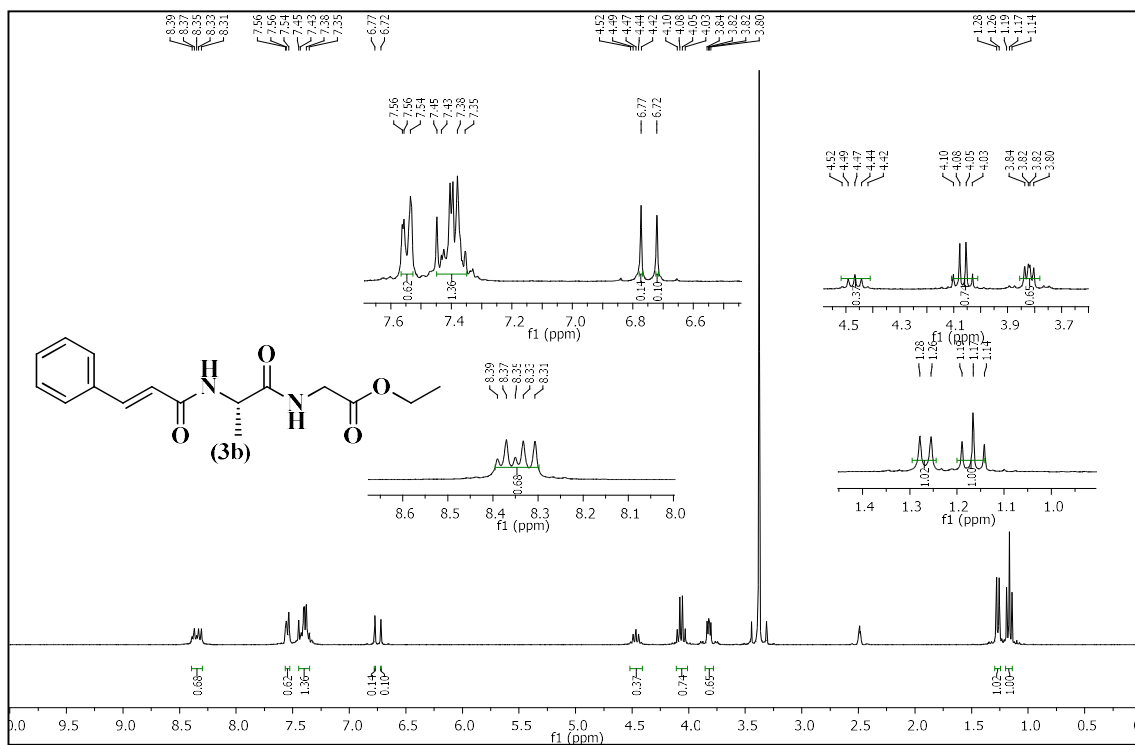


Figure 5: ¹H NMR Spectrum of N-Cinnamoyl-Ala-Gly-OEt (3b)

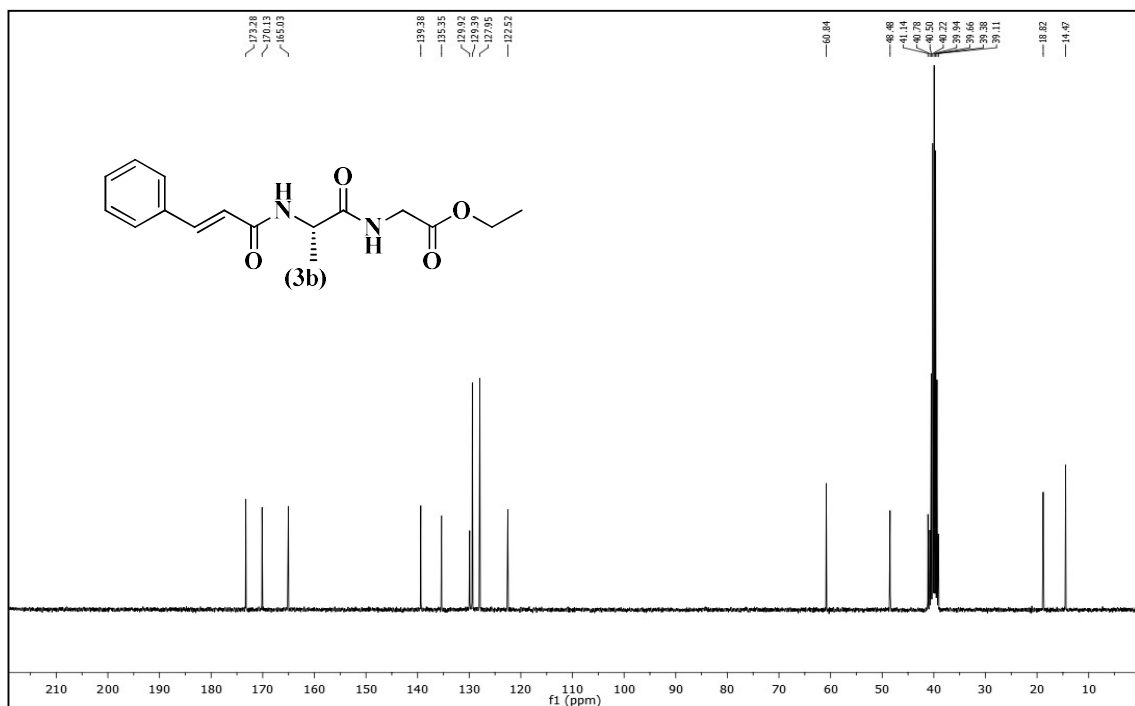


Figure 6: ¹³C NMR Spectrum of N-Cinnamoyl-Ala-Gly-OEt (3b)

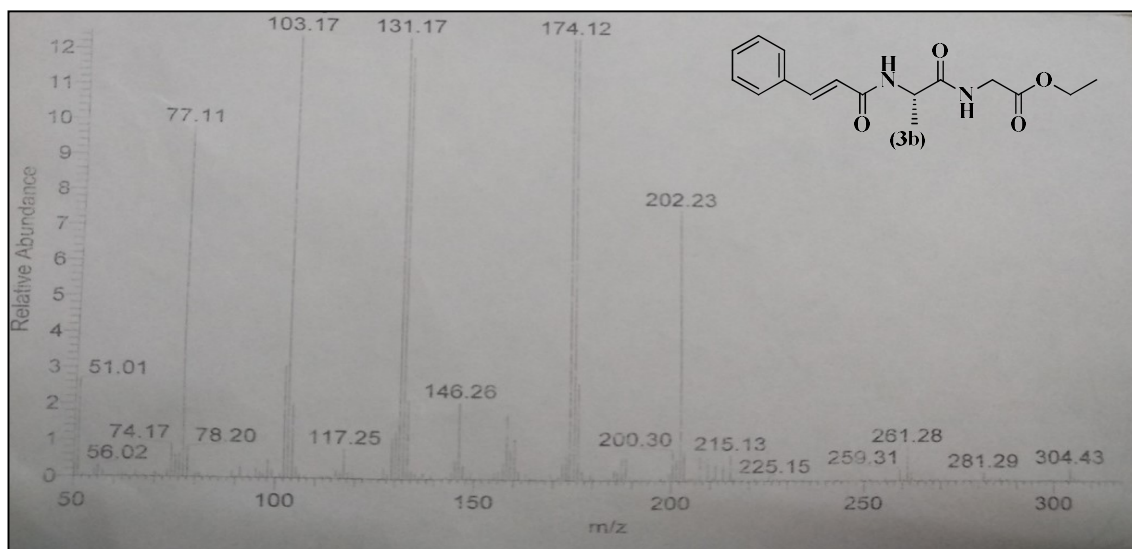


Figure 7: Mass Spectrum of N-Cinnamoyl-Ala-Gly-OEt (**3b**)

Spectral Details of N-Cinnamoyl-Phe-Gly-OEt

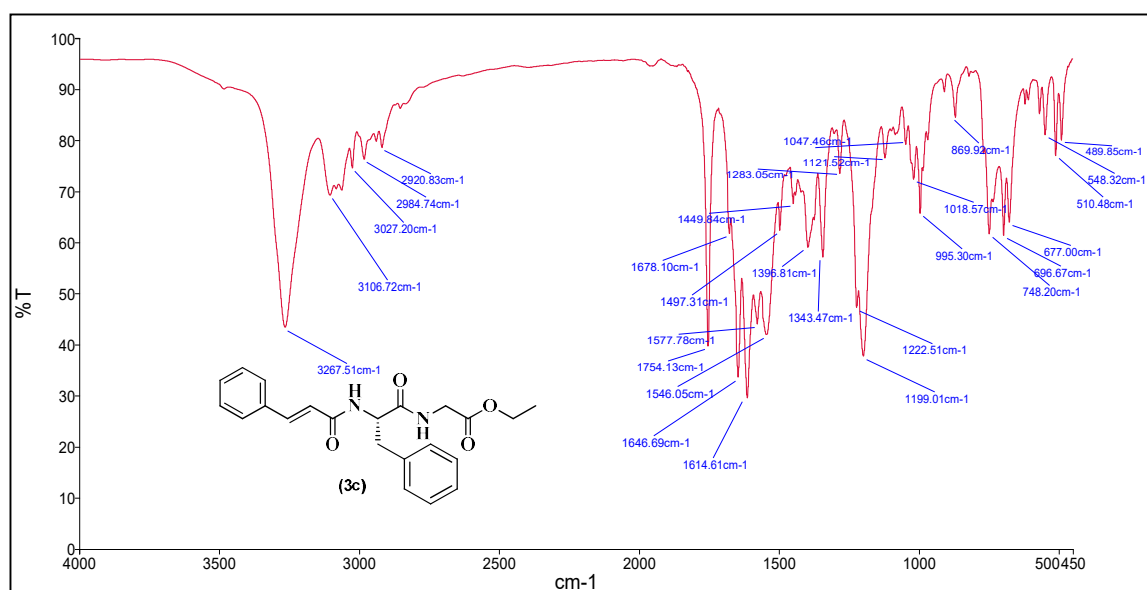


Figure 8: IR spectrum of N-Cinnamoyl-Phe-Gly-OEt (**3c**)

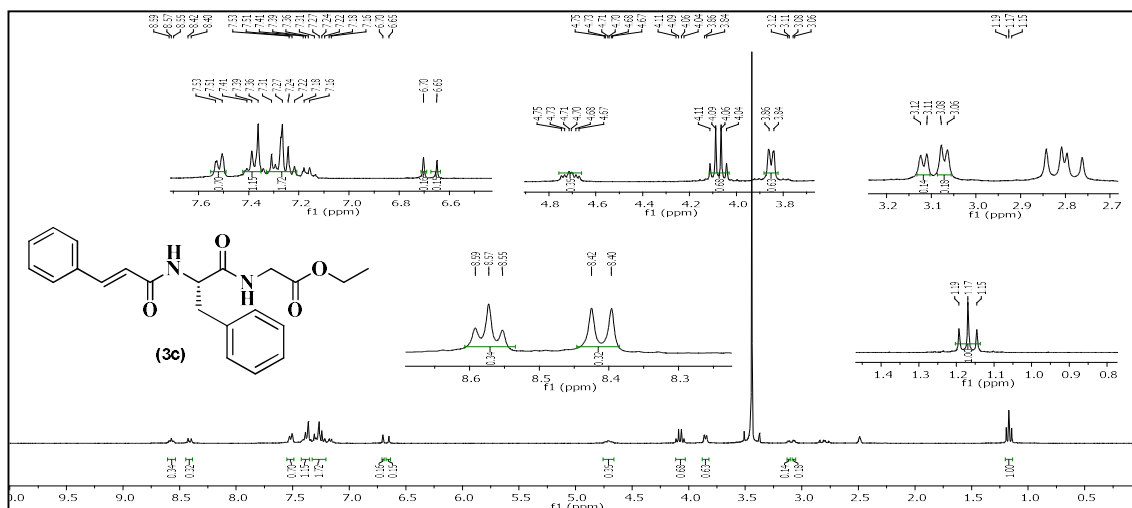


Figure 9: ^1H NMR spectrum of N-Cinnamoyl-Phe-Gly-OEt (3c)

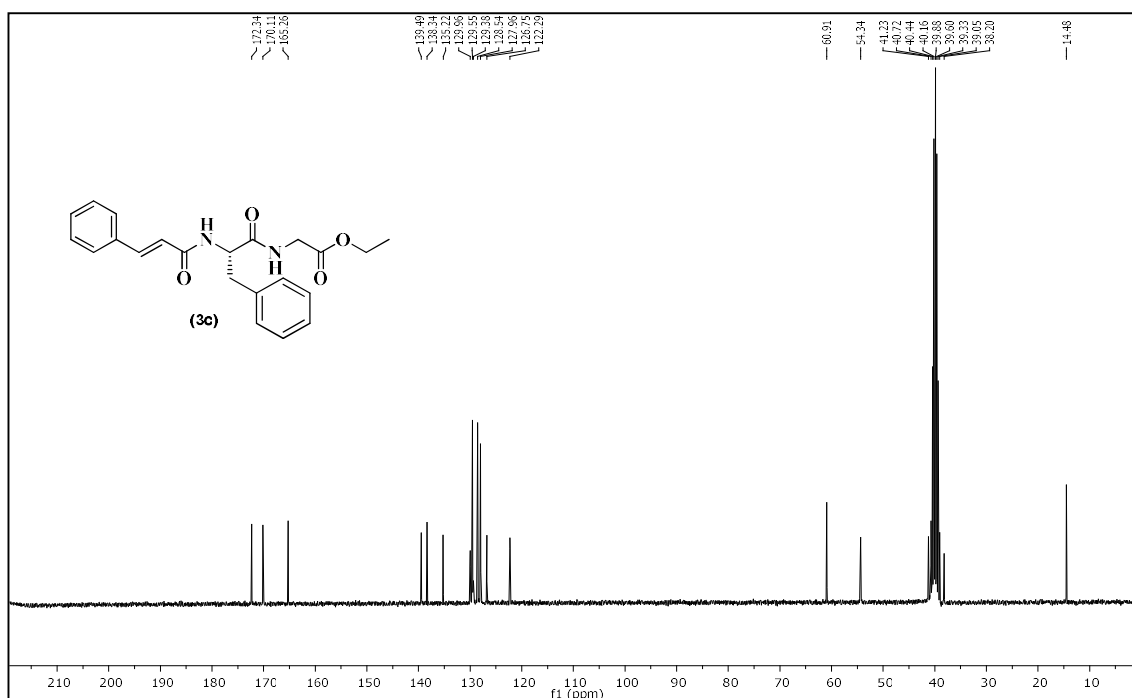


Figure 10: ^{13}C NMR Spectrum of N-Cinnamoyl-Phe-Gly-OEt (3c)

Spectral Details of N-Cinnamoyl-Gly-Phe-OEt

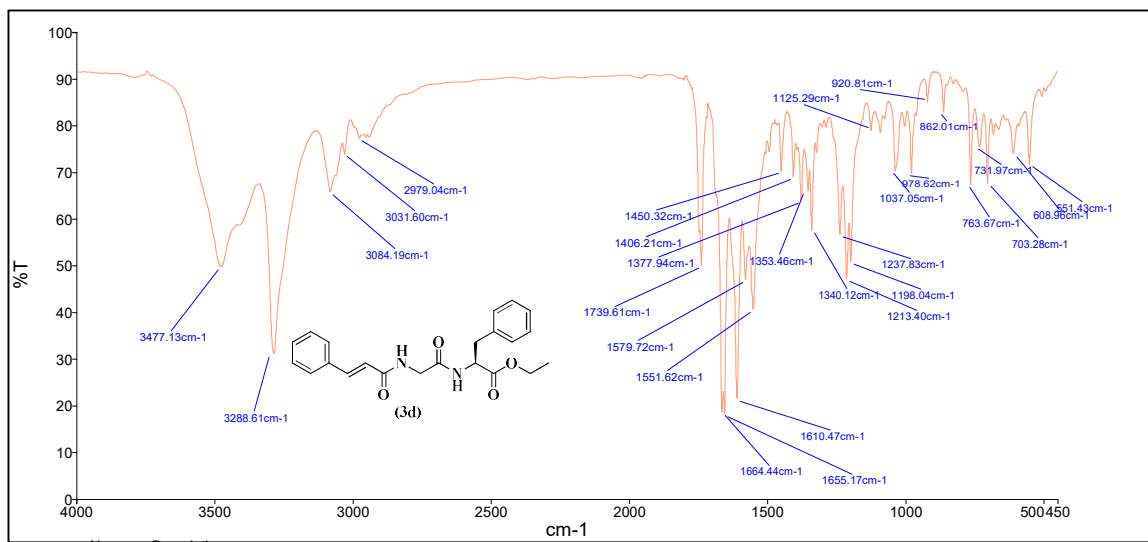


Figure 11: IR Spectrum of N-Cinnamoyl-Gly-Phe-OEt (3d)

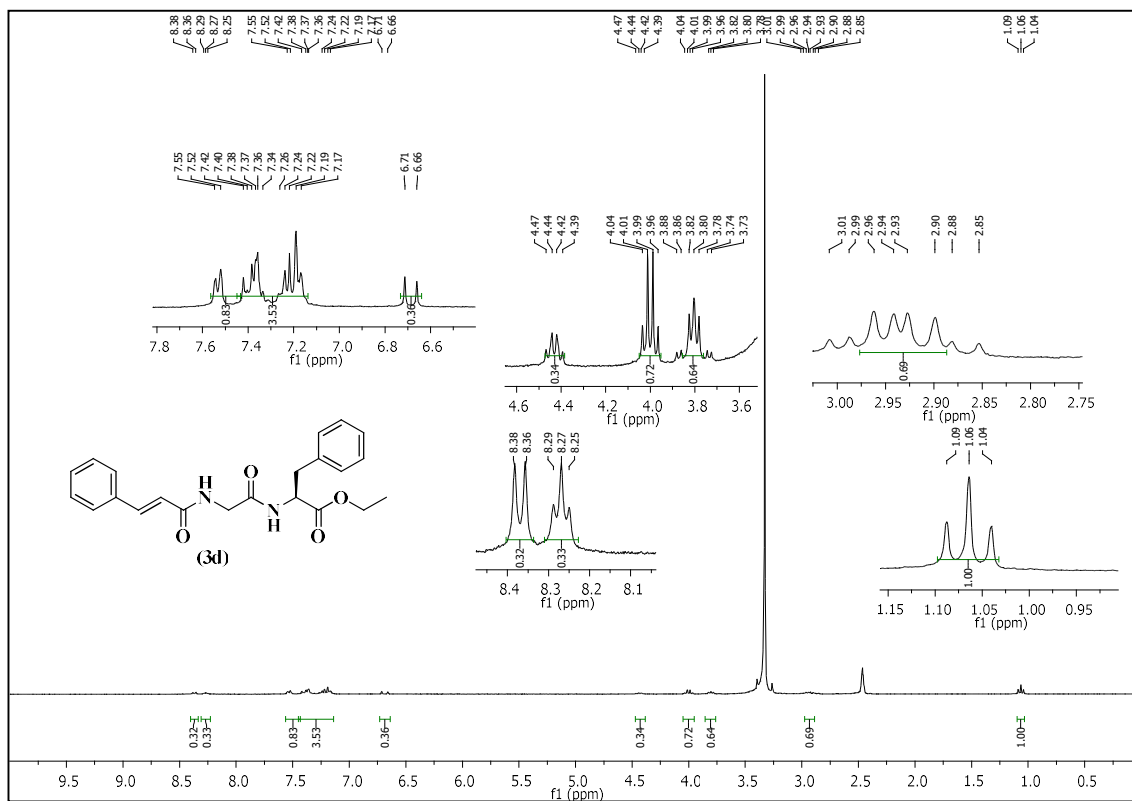


Figure 12: ¹H NMR Spectrum of N-Cinnamoyl-Gly-Phe-OEt (3d)

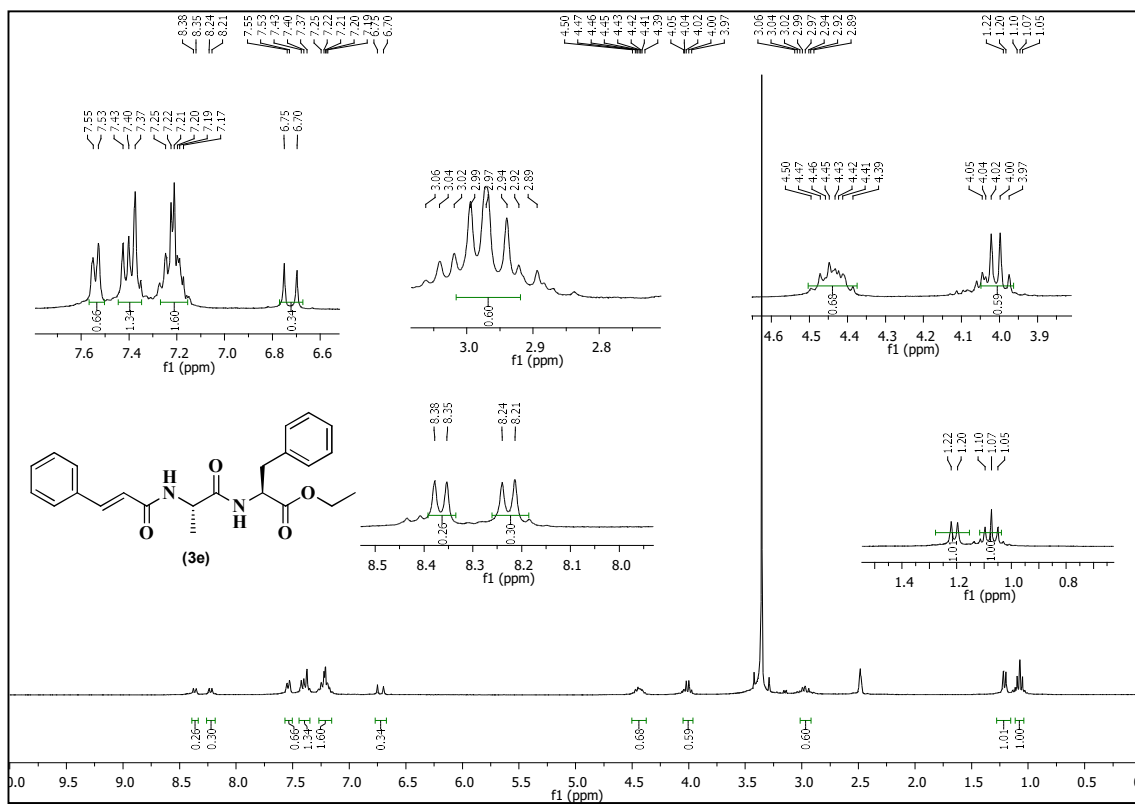


Figure 15: ¹H NMR Spectrum of N-Cinnamoyl-Ala-Phe-OEt (3e)

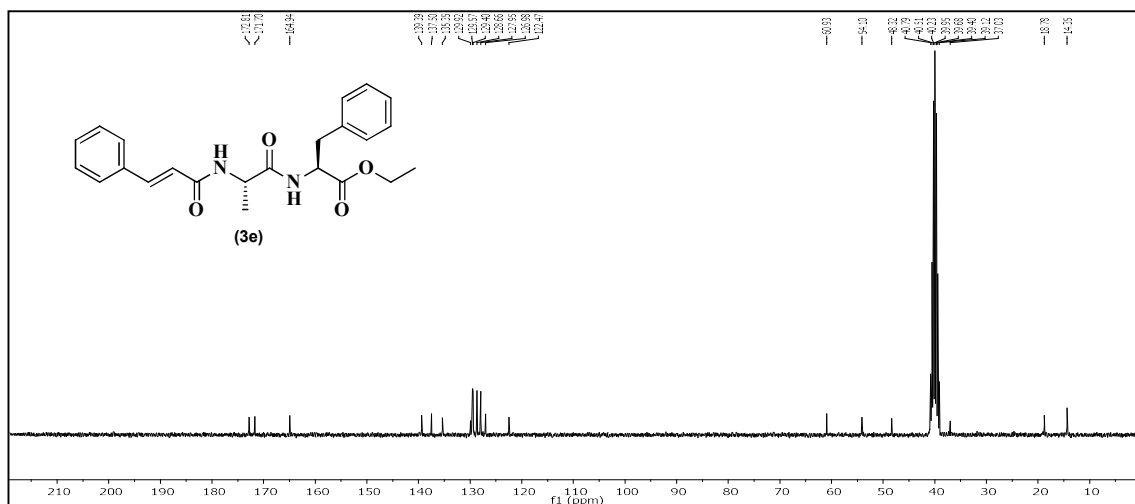


Figure 16: ¹³C NMR Spectrum of N-Cinnamoyl-Ala-Phe-OEt (3e)

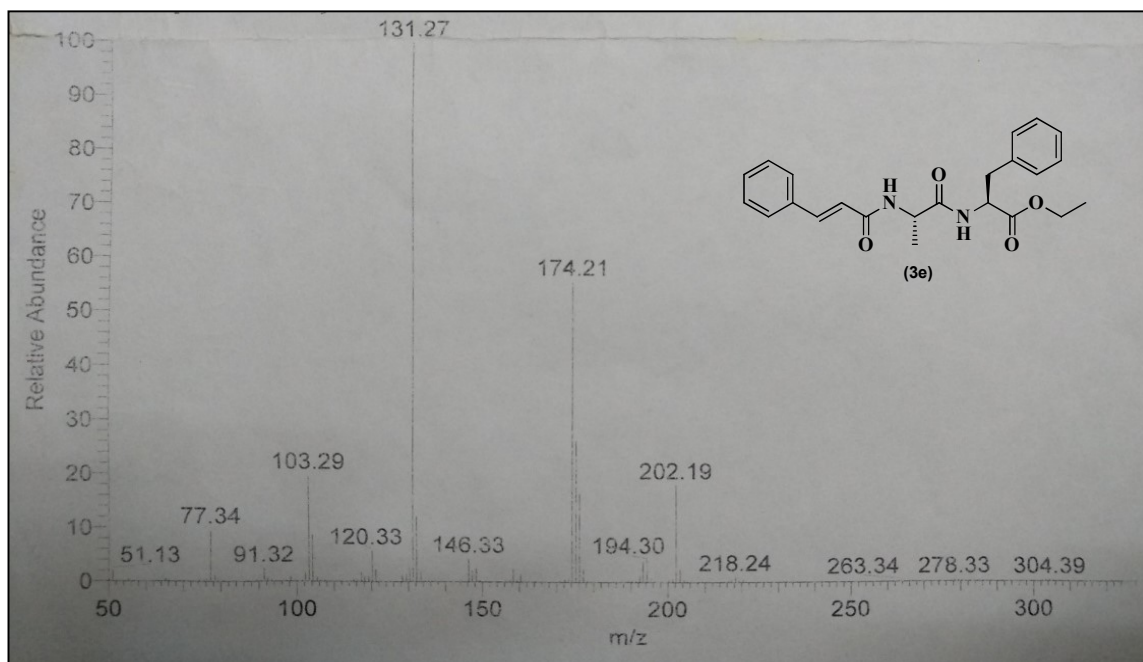


Figure 17: Mass spectrum of N-Cinnamoyl-Ala-Phe-OEt (**3e**)

Sample preparation procedure for the Morphological Study of N-Cinnamoyl-dipeptide ester (3):

1. Preliminary step for SEM sample preparation of N-Cinnamoyl-dipeptide ester (3)

A fresh stock of solution was prepared by dissolving N-Cinnamoyl dipeptide ester in 1,1,1,3,3,3-Hexafluoro-2-propanol (HFIP) at a concentration of 100 mg/mL. 10mg (100 μ L) of compound was then taken out from the stock solution and diluted into final concentration of 2 mg/mL in double distilled water. After dilution with water, spontaneous precipitation was observed. The precipitated sample was further analysed using SEM.

2. SEM Analysis of N-Cinnamoyl-dipeptide ester (3) : 10 uL of the solution prepared in step 1 of N-Cinnamoyl-dipeptide ester of concentration 2 mg/mL was spotted on a aluminium plate. The plate was allowed to dry in vacuum desiccator and was subjected to SEM analysis.

3. TEM Analysis of N-Cinnamoyl-dipeptide ester (3) 10 uL of the solution prepared in step 1 of N-Cinnamoyl-dipeptide ester of concentration 2 mg/mL was spotted on a copper grid, coated by carbon stabilized Formvar film. The grid was dried in a vacuum desiccator. 10 uL of 2 % uranyl acetate solution in water was then spotted on the same grid and then it was dried in vacuum. The grid was then studied using TEM.