

SYNTHESIS AND CHARACTERIZATION OF A FLUORESCENT DERIVATIVE OF PMVEMA WITH PYRENE

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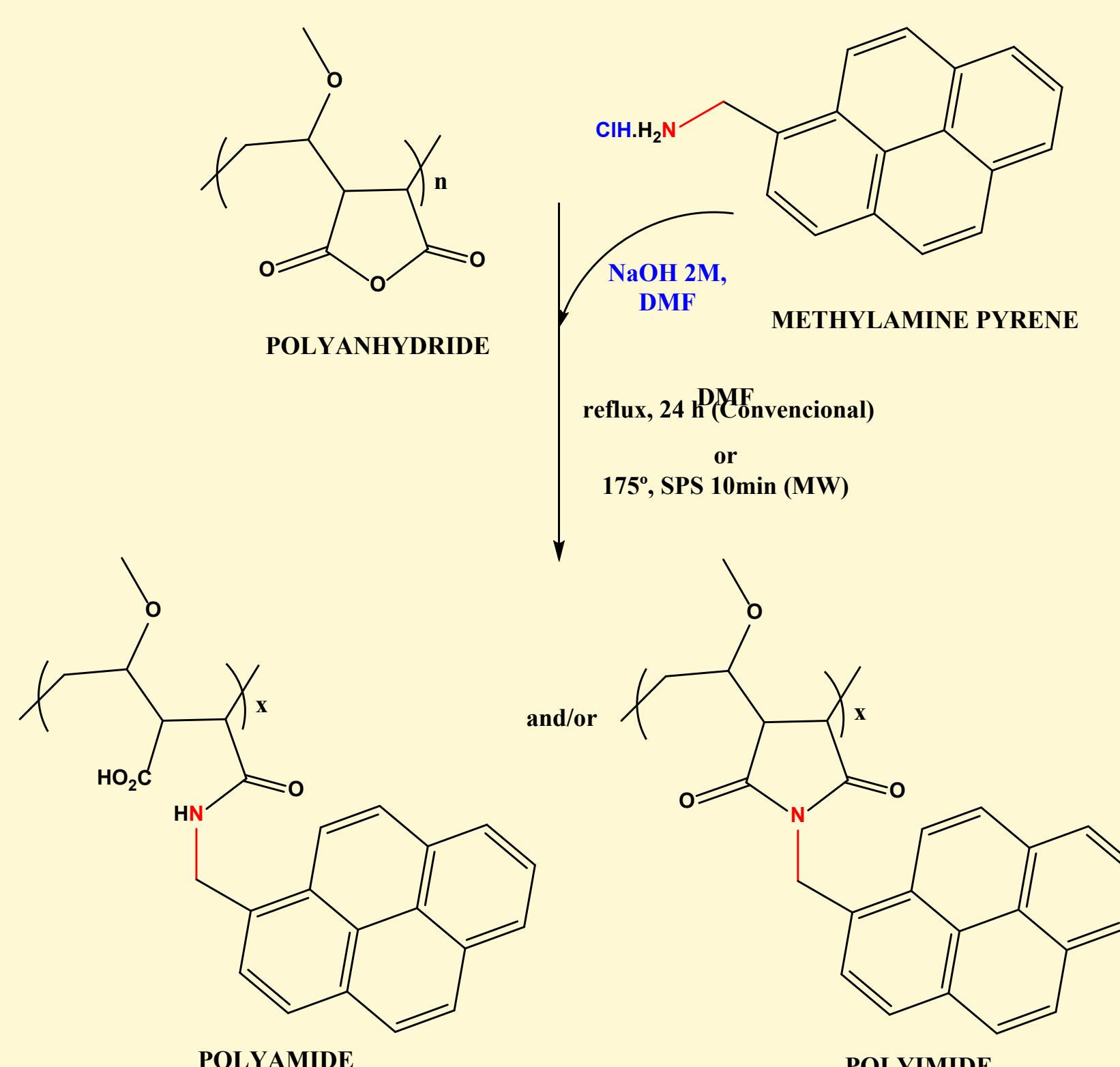
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INTRODUCTION

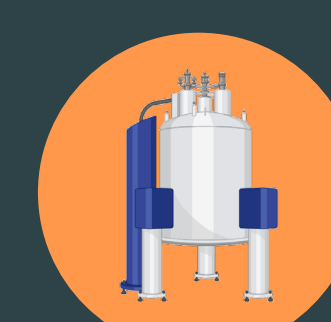
Poly(methyl vinyl ether-alt-maleic anhydride) (PMVEMA) is a versatile polymer known for its properties such as biodegradability and biocompatibility, making it suitable for biomedical applications. Furthermore, its use as a base polymer for the synthesis of nanofibers via electrospinning has already been described. Its derivatization with fluorescent moieties, such as pyrenes, opens new fields of application. In this work, a derivative of the PMVEMA polymer with pyrene (PMVEMA-Pyr) was synthesized, characterized and used for nanofiber synthesis through electrospinning.

SYNTHETIC SCHEME

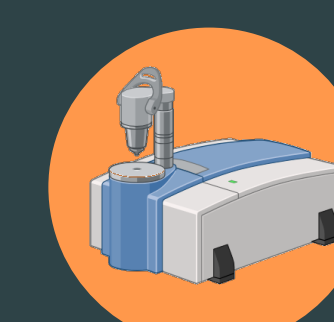


METHODOLOGY

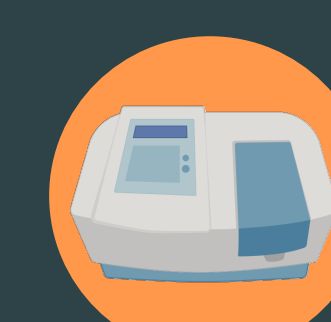
Pyrene 1-pyrenemethylamine was incorporated into the PMVEMA backbone via post-polymerization modification reaction through nucleophilic attack. The molar ratio of PMVEMA to pyrene in the reaction was 2:1. The chemical structure, thermal property, and photophysical performance were analyzed. Nanofibers were synthesized via electrospinning using a blend of PMVEMA-ES 50% w/w in ethanol and PMVEMA-Pyr 5% w/w in DMSO.



Nuclear Magnetic Resonance (NMR)



Fourier Transform Infrared Spectroscopy (FTIR)



Absorbance and emission spectra



Thermogravimetric analysis (TGA)

NMR

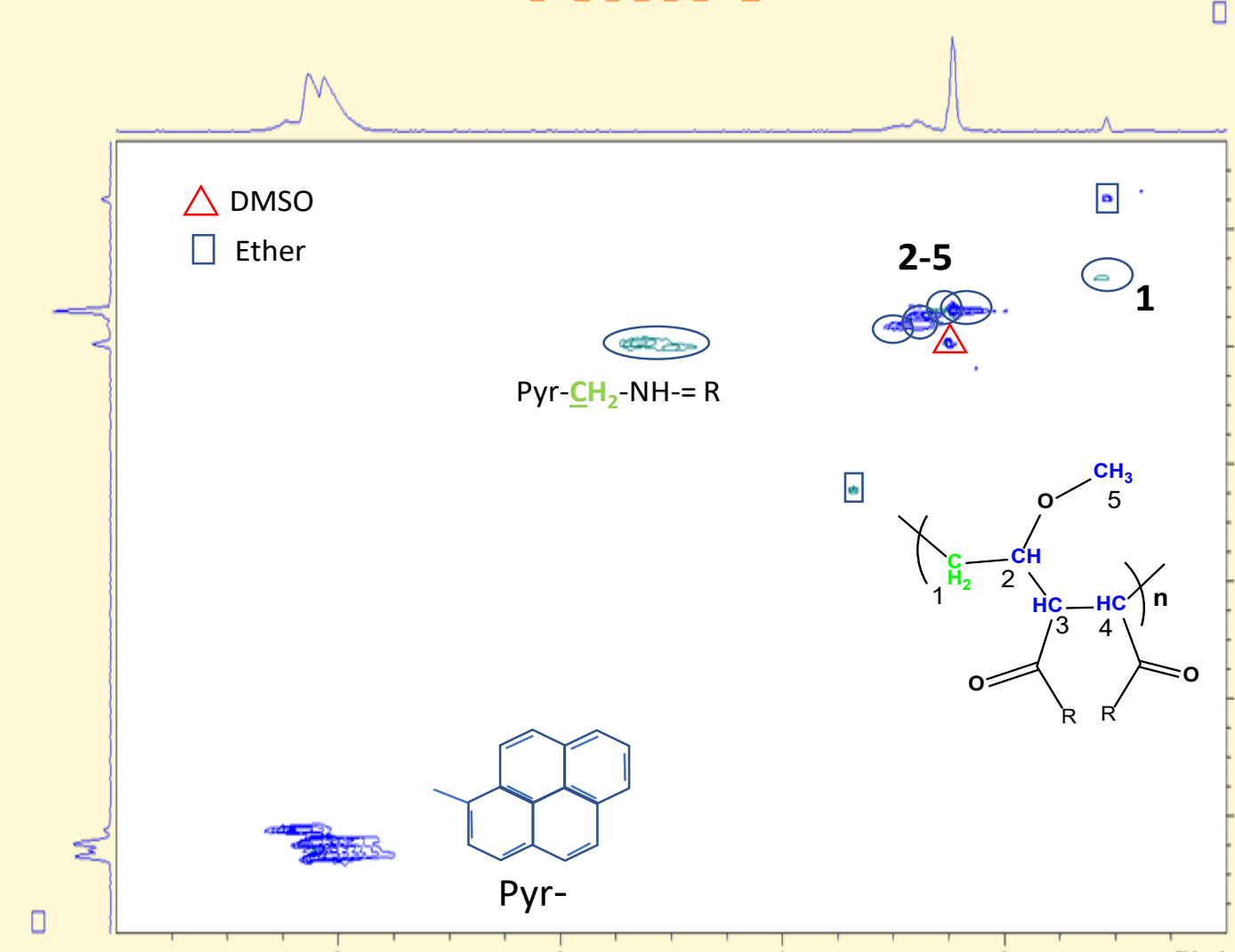


Figure 1. ^1H - ^{13}C HSQC spectrum of PMVEMA-Pyr in DMSO-d_6 (as internal reference). Solvent signals were showed (square and triangle).

RESULTS

UV-vis

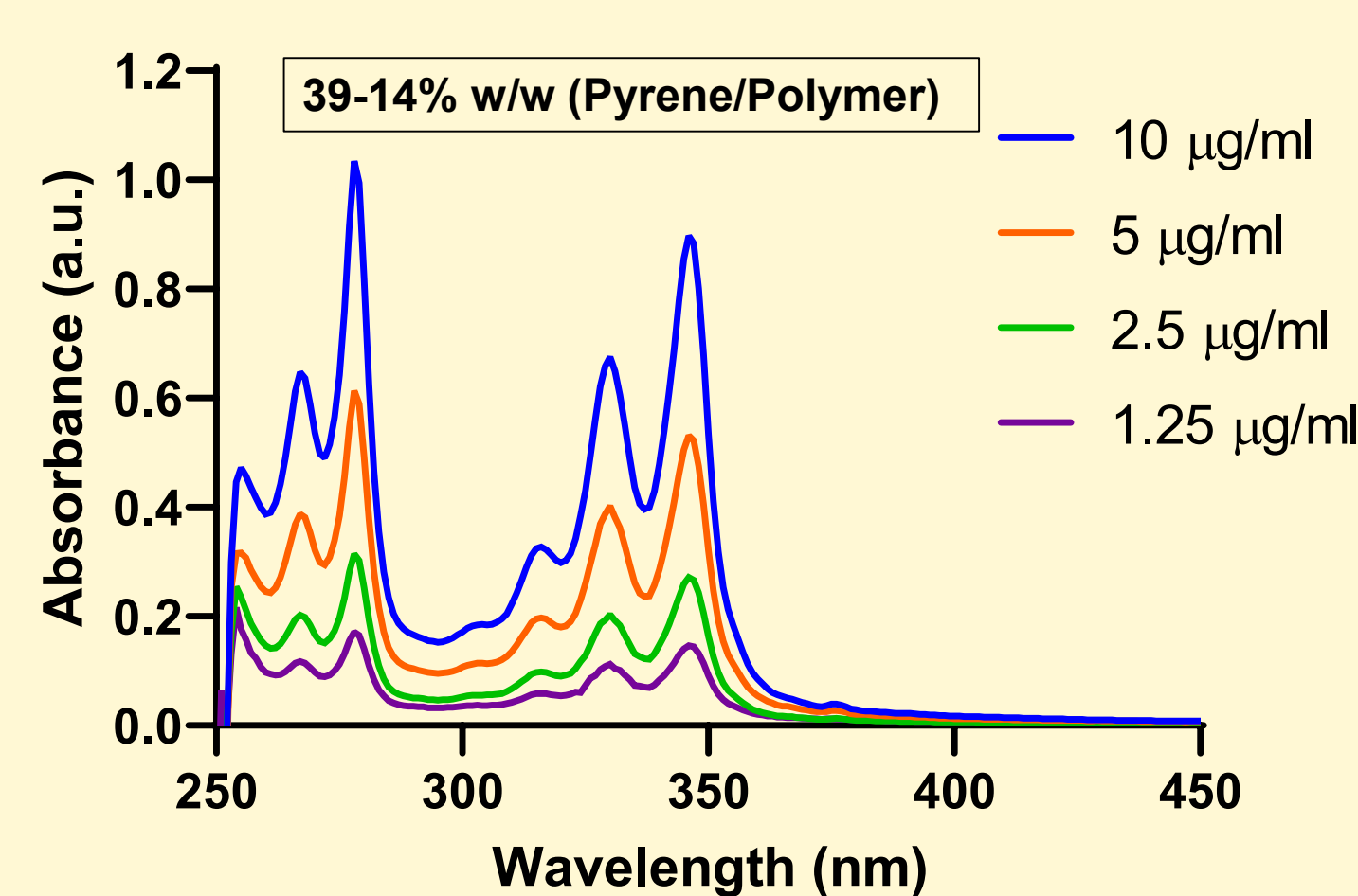


Figure 2. UV-vis absorption spectra of PMVEMA-Pyr in DMSO at different concentrations. The main absorption peaks are found at 267, 278, 330 and 346 nm and correspond to the characteristic absorption bands of pyrene.

Excitation and emission

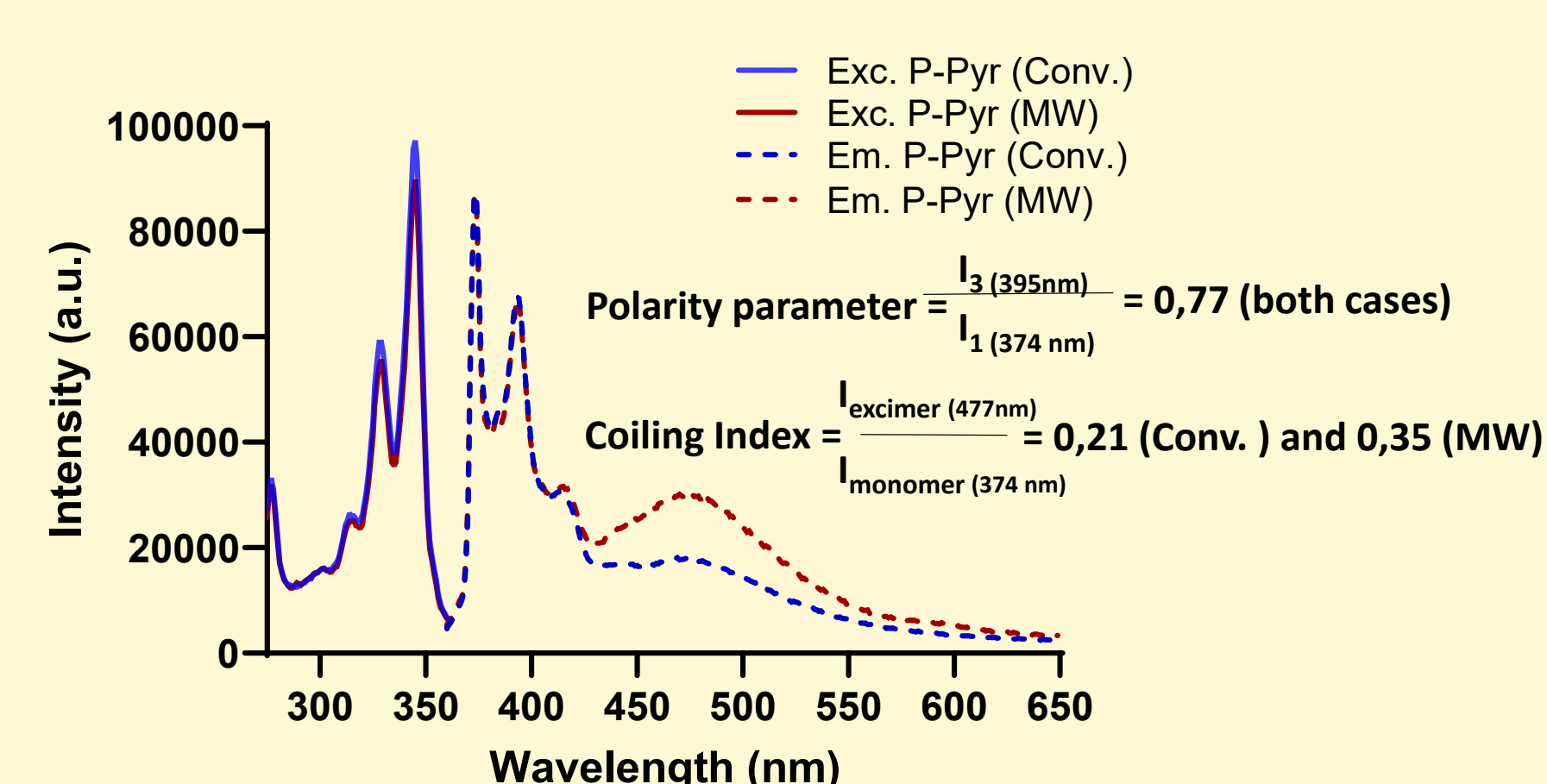


Figure 3. Excitation and emission spectra of PMVEMA-Pyr obtained via conventional and microwave assisted synthesis. Two main peaks are observed at 377 and 398 nm, and a broad excimer emission band around 480 nm.

FTIR

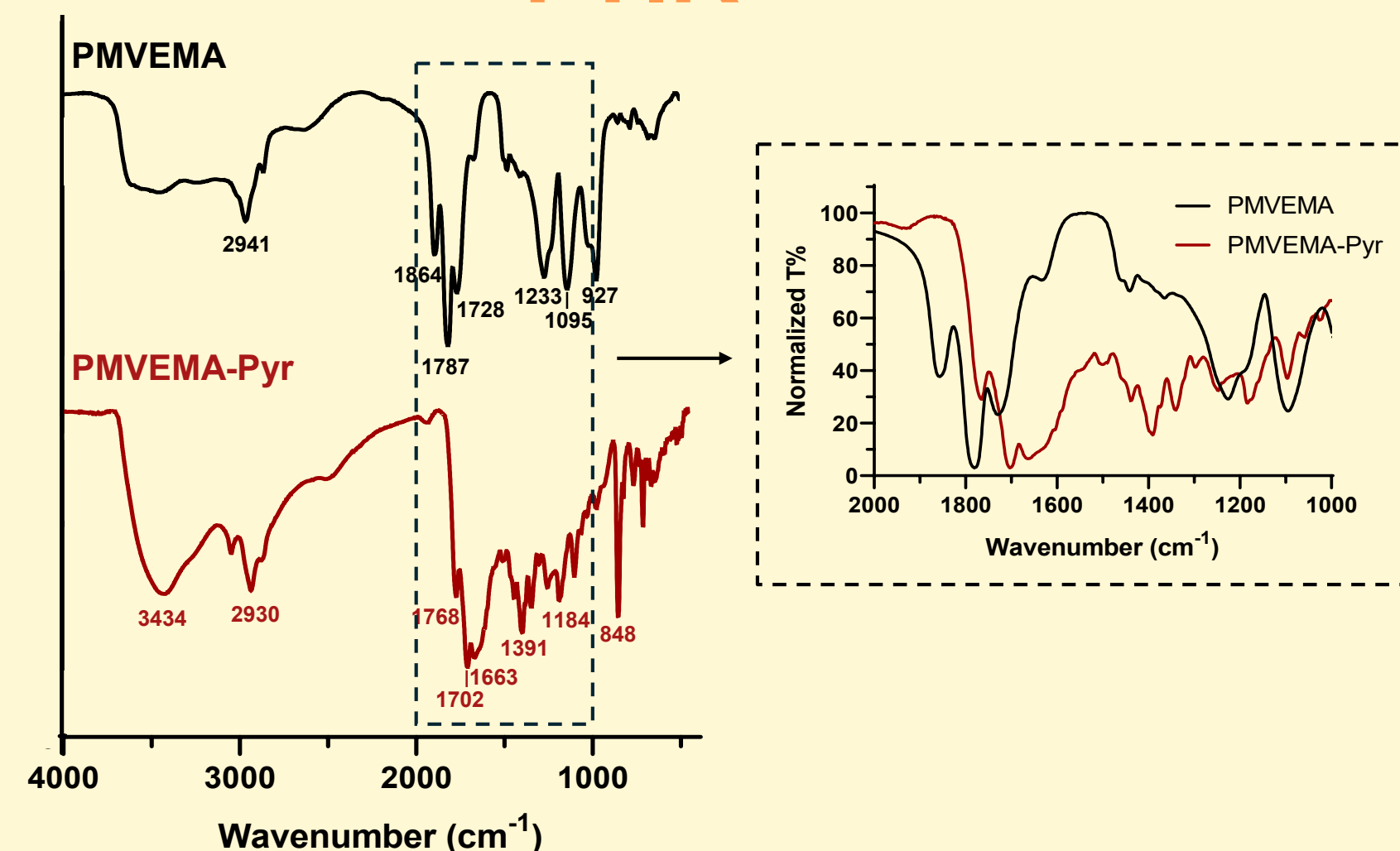


Figure 2. FTIR spectra of PMVEMA-Pyr. The peaks at 1900-1600 cm^{-1} in PMVEMA can be attributed to the asymmetric vibrations of the carbonyl $\text{C}=\text{O}$ groups. However, these peaks are shifted to the right in the PMVEMA-Pyr spectrum, which may be due to the transformation into amide groups.

TGA

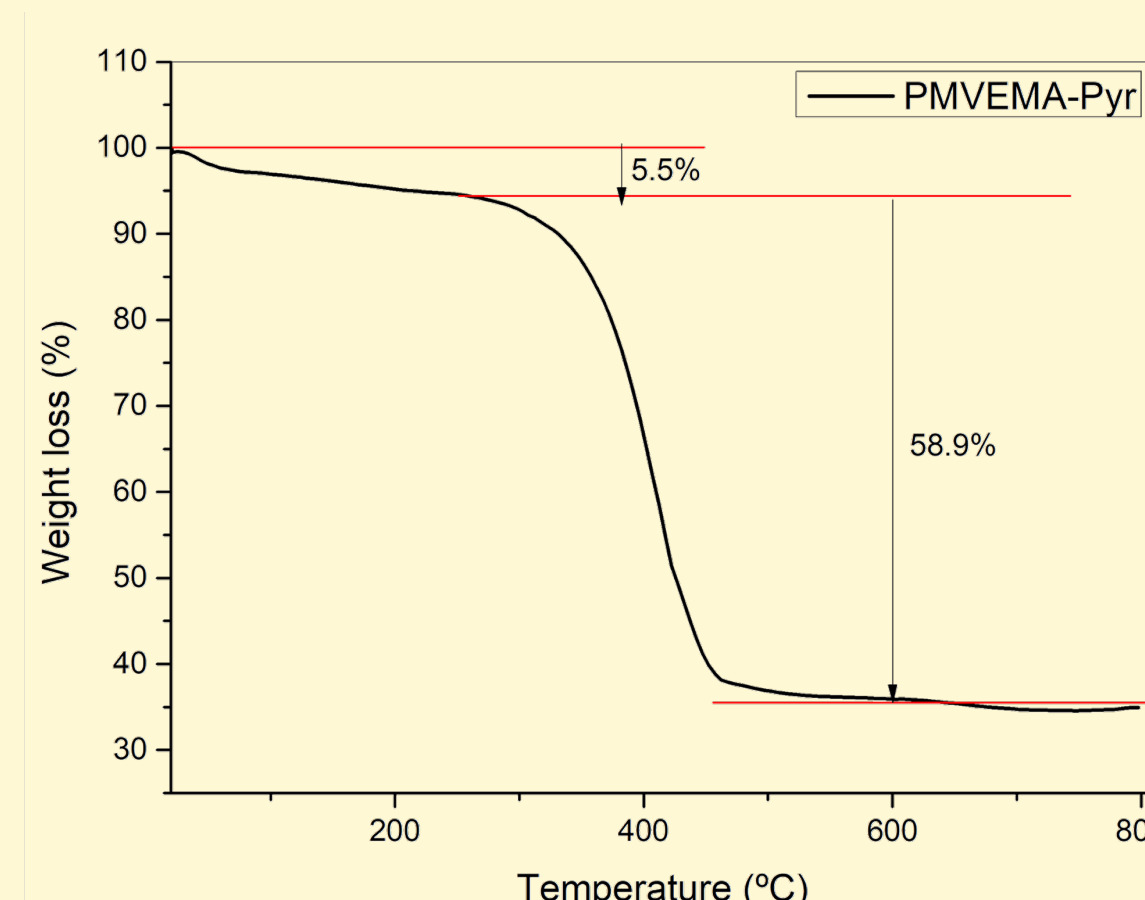


Figure 5. TGA-DGT curve of PMVEMA-Pyr. The main decomposition stage occurs at around 270 and 550 $^\circ\text{C}$ with a mass loss of 58.9%.

Nanofibers

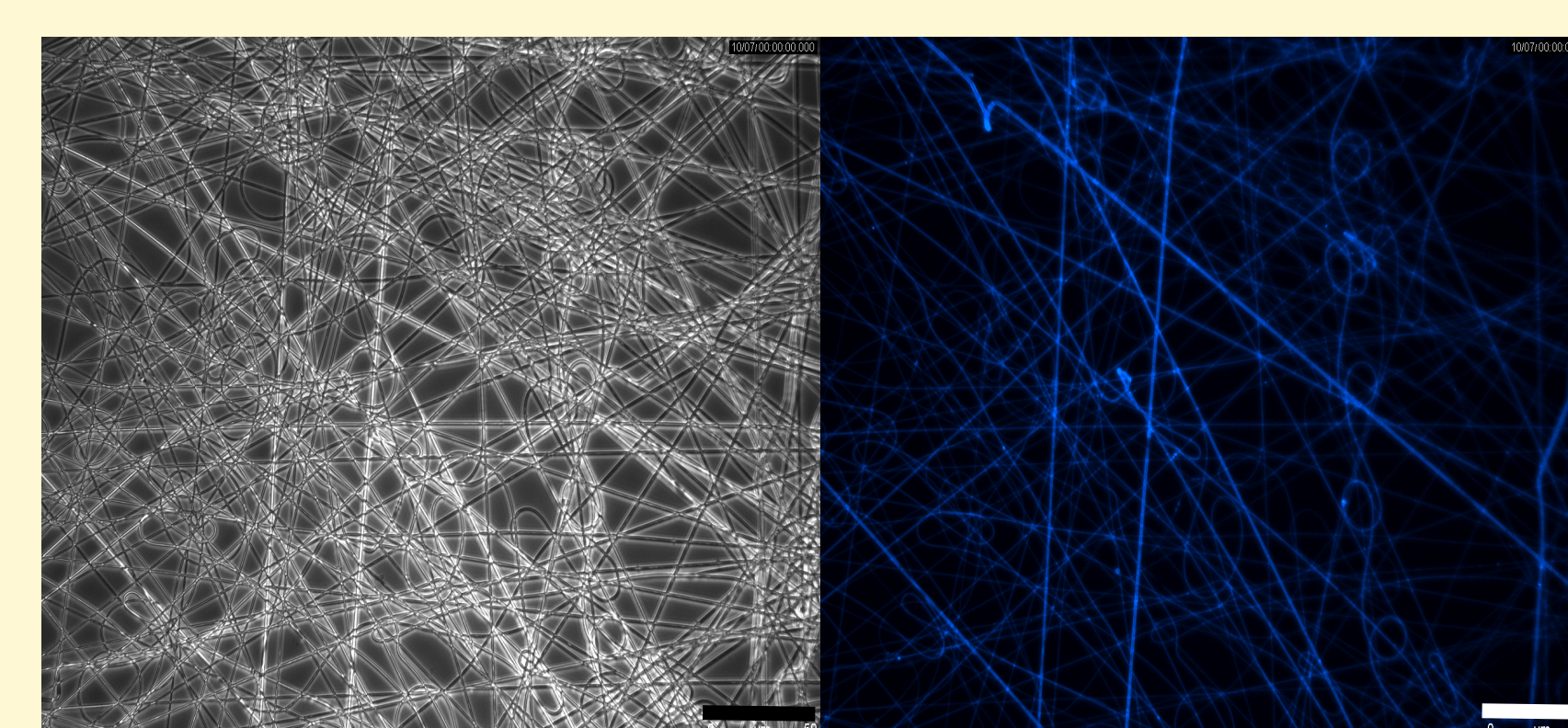


Figure 6. Fluorescent microscopy images of the PMVEMA-ES/PMVEMA-Pyr nanofibers. Scale bar: 50 μm .

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Jeon, H.K., Feist, B.J., Koh, S.B., Chang, K., Macosko, C.W., & Dion, R.P. (2004). Reactively formed block and graft copolymers as compatibilizers for polyamide 66/PS blends. *Polymer*, 45, 197-206.

CONCLUSIONS

- PMVEMA derivative with pyrene (PMVEMA-Pyr) was achieved via conventional synthesis and MW assisted synthesis.
- The different characterization methods confirmed the formation of PMVEMA-Pyr.
- No differences were found between the products obtained using the two types of synthesis, however MW synthesis allows for faster product acquisition.
- Fluorescent nanofibers were synthesized via electrospinning using a blend of PMVEMA-ES and PMVEMA-Pyr.
- Further studies will involve the determination of the biological effects of PMVEMA-Pyr and PMVEMA-Pyr nanofibers.

ACKNOWLEDGEMENTS

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