

Preparation of Amphiphilic Block Copolymers using Canola Oil Fatty Acids to Incorporate Carbamazepine Drug

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Introduction

Micelles are an important area of study due to their use in pharmaceuticals and their advantages over conventional therapeutics, such as targeted drug-delivery. Micelles have diameters generally between 10 – 200nm¹.

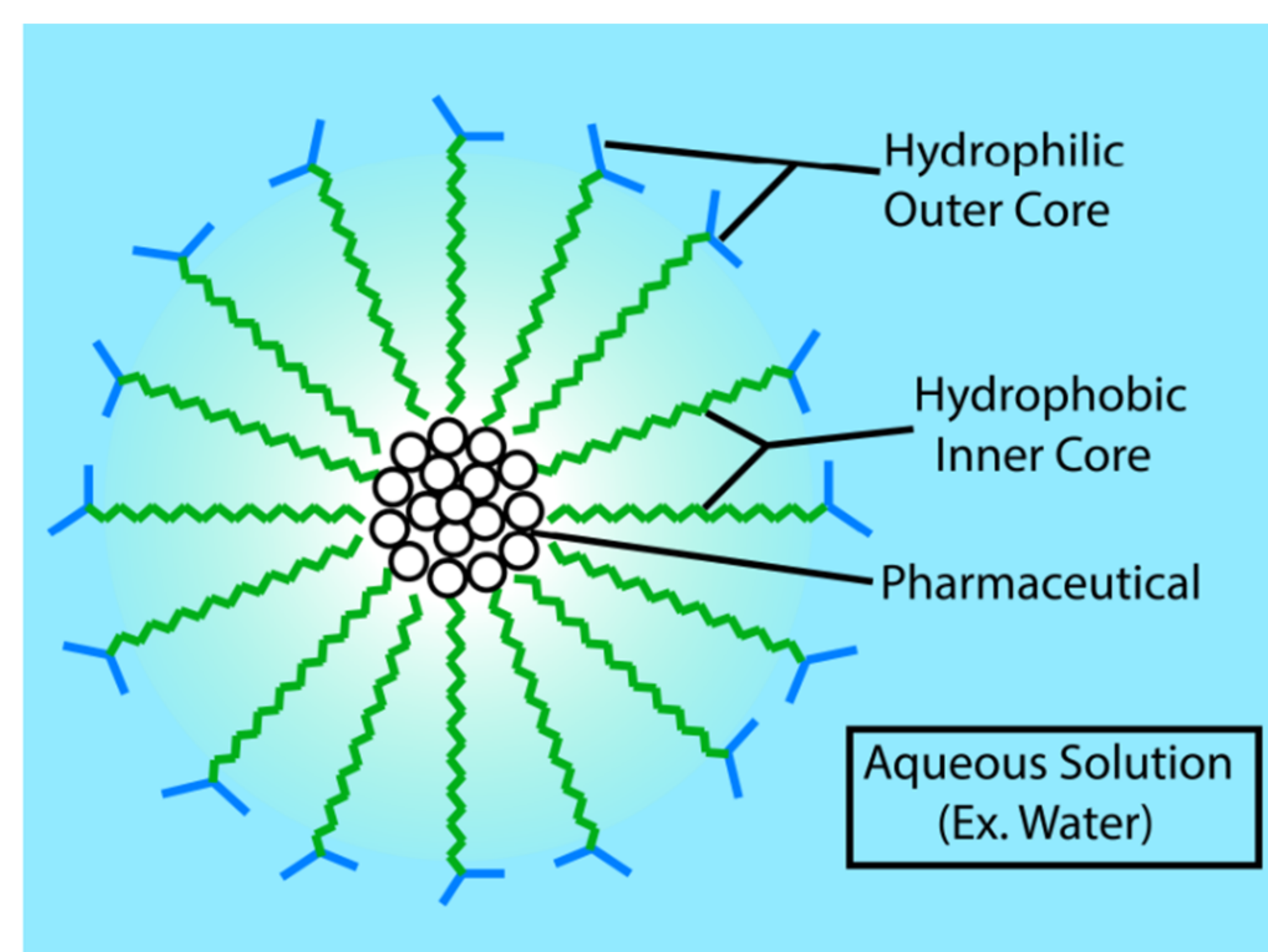


Fig. 1: Diagram of a micelle encapsulating a pharmaceutical within an aqueous medium.

Purpose

- ~ Produce a drug-carrying micelle
- ~ Replace the synthetic hydrophobic component within existing micelle designs with Canola Fatty Esters (CFEs)
- ~ Use of CFEs as a hydrophobic component as lipids are :
 - Biocompatible
 - Biodegradable
 - Non-toxic
 - Non-allergenic
 - Inexpensive

Methods

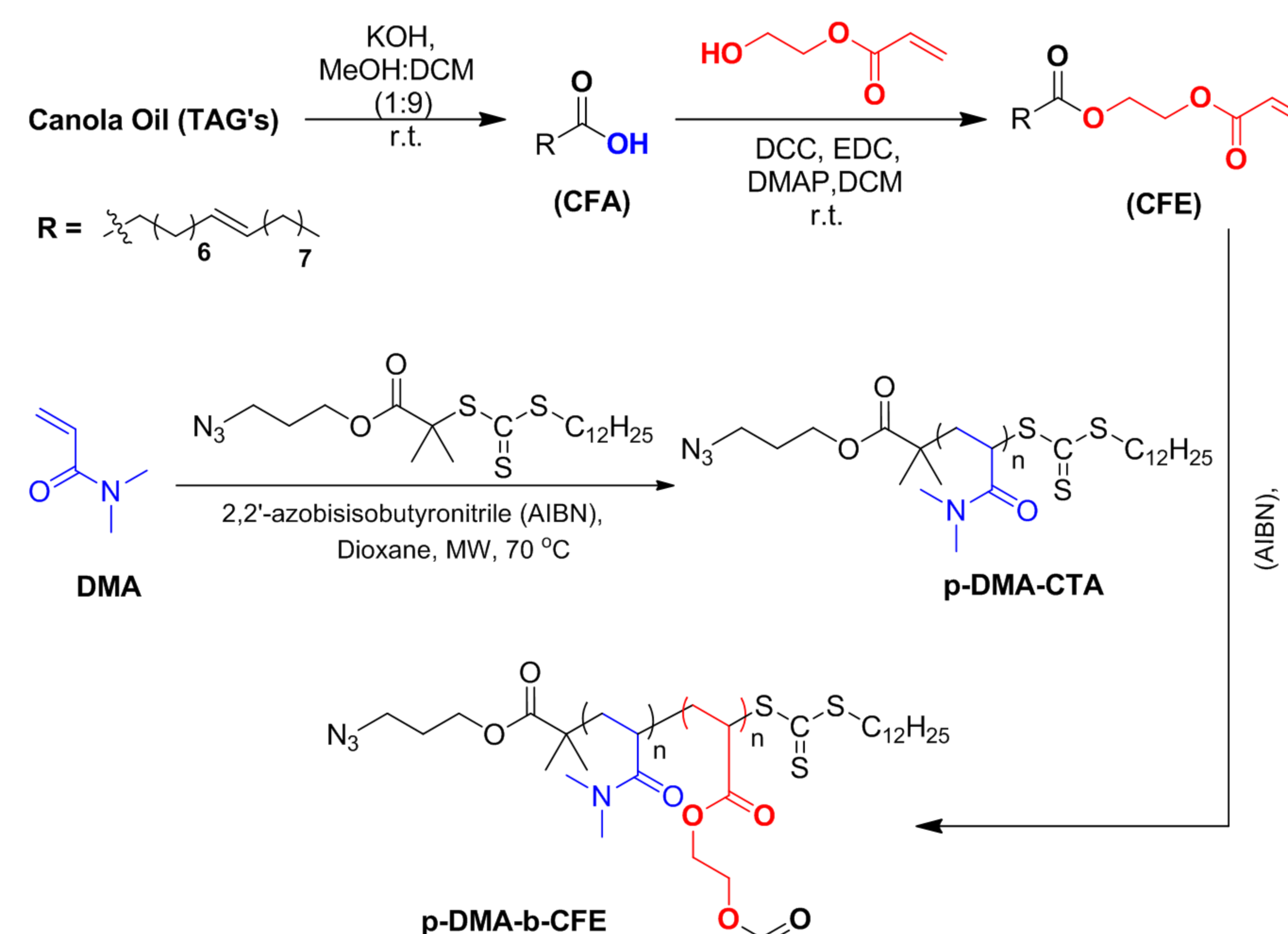
The micelle in question will be made of two main components:

1. poly(*N, N'*-Dimethylacrylamide) (**p-DMA**)
2. CFEs refined from canola triacylglycerols (**TAGs**)

~ The hydrophobic component was obtained from canola oil by the hydrolysis of TAGs followed by the esterification of carboxylic acid with 2-hydroxyethylacrylate.

~ The hydrophilic component was obtained via RAFT (Reversible Addition–Fragmentation chain Transfer) polymerization using DMA.

Both parts were copolymerized by RAFT method into an amphiphilic bio-conjugate.



Scheme, 1: Preparation of amphiphilic block copolymer **p-DMA-b-CFE** by RAFT polymerization

Results

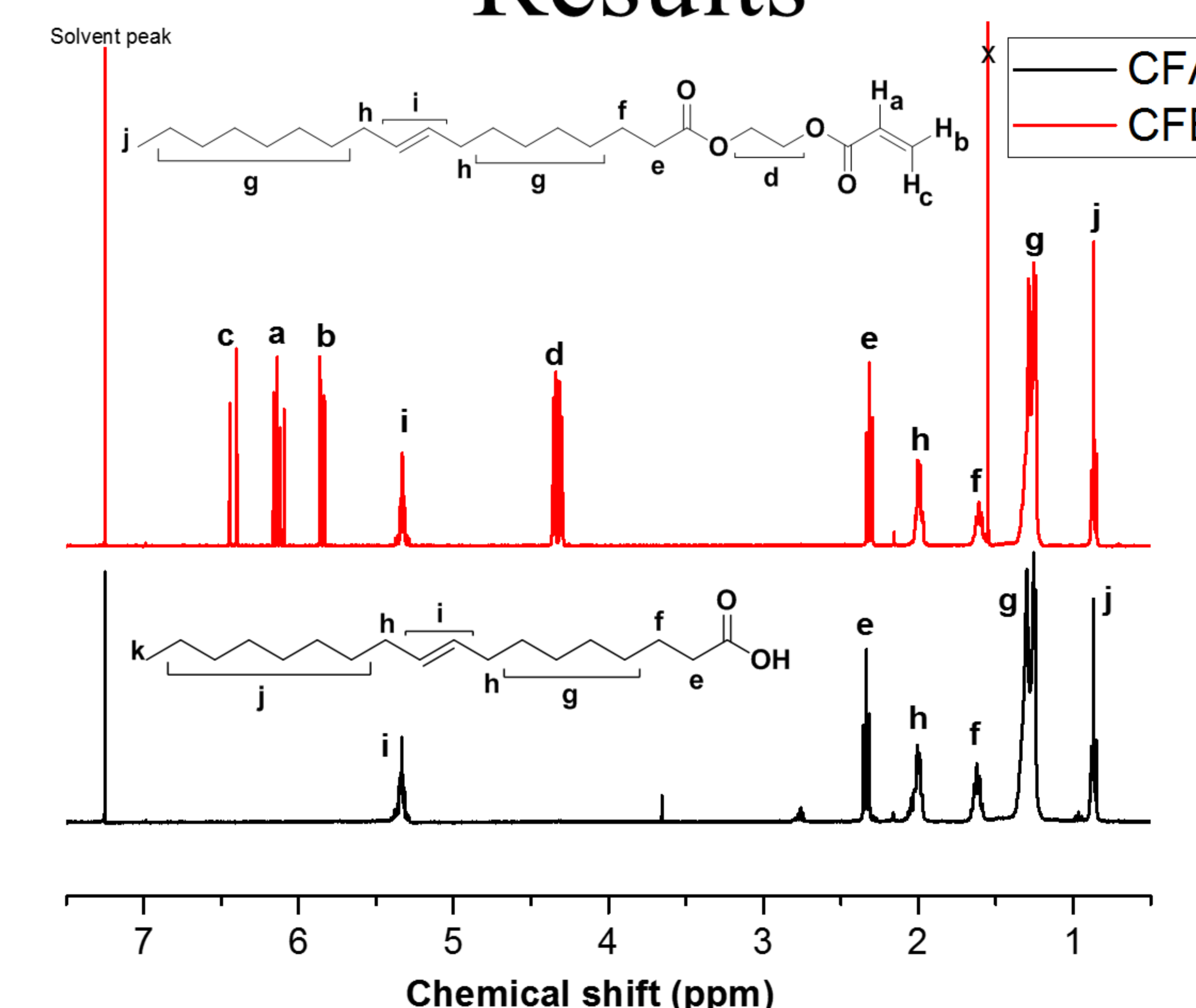


Fig. 2: Proton NMR spectra of canola fatty acid (CFA) and canola fatty esters (CFE) in deuterated chloroform

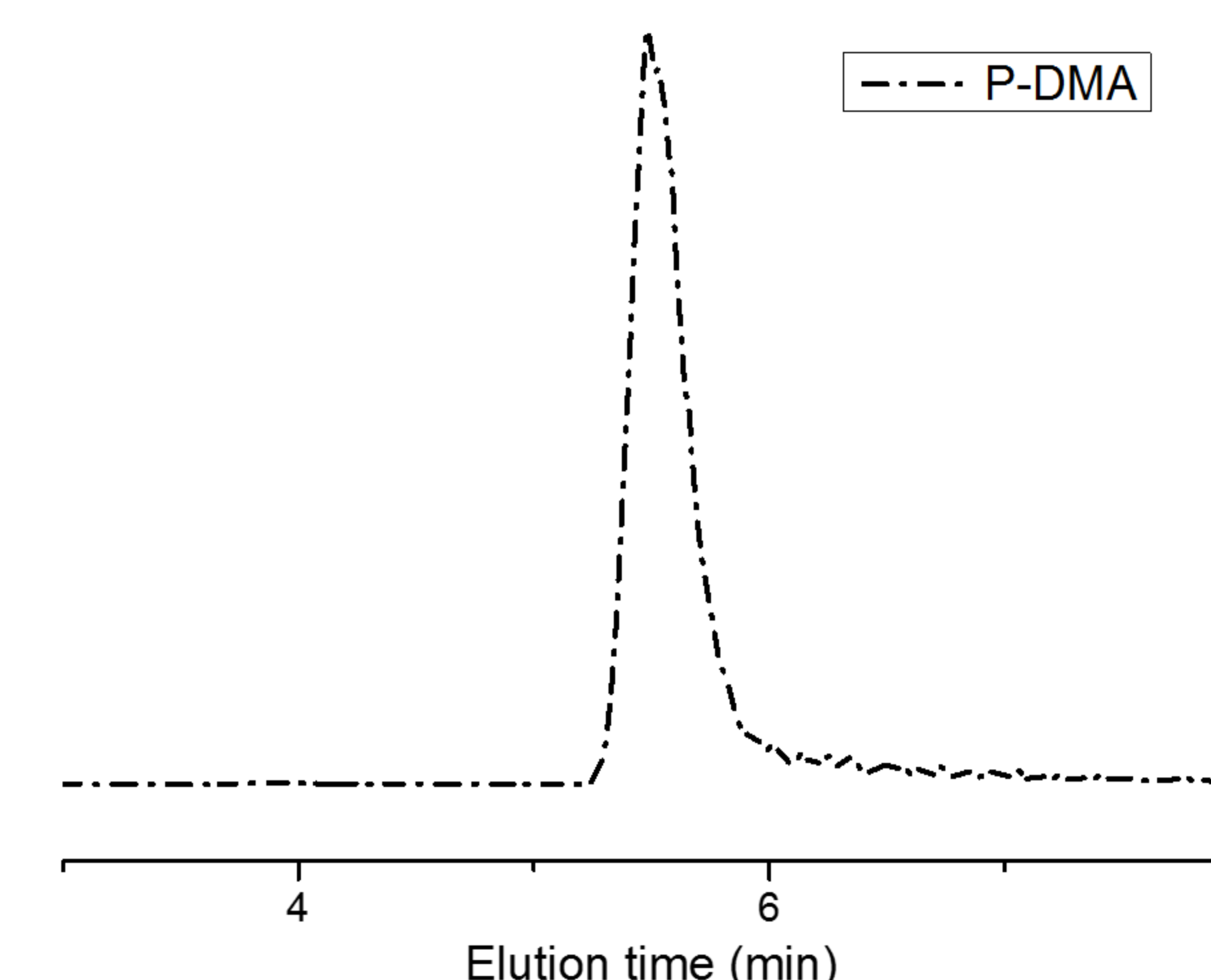


Fig. 3: Gel permeation chromatography (GPC) traces of **P-DMA** (Mw = 6200 g/mol, PDI = 1.03)

Conclusion

- ~ Obtained the monomer CFE, confirmed with HNMR
- ~ Obtained the DMA polymer, confirmed with GPC
- ~ Synthesized the bio-conjugate copolymer, HNMR tests required to confirm product has successfully reacted
- ~ Future research into drug encapsulation and release trials are needed after acquiring the block copolymer