

Synthesis and Characterization of Porous Polymer-based Adsorbents for CO₂ Capture

Kehinde A. Fayemiwo^{*a}, Seyed Ali Nabavi^{a,b}, Goran T. Vladislavljević^a, Vasilije Manović^b, Brahim Benyahia^a



^aLoughborough University, Chemical Engineering Department, LE11 3TU, Loughborough, UK

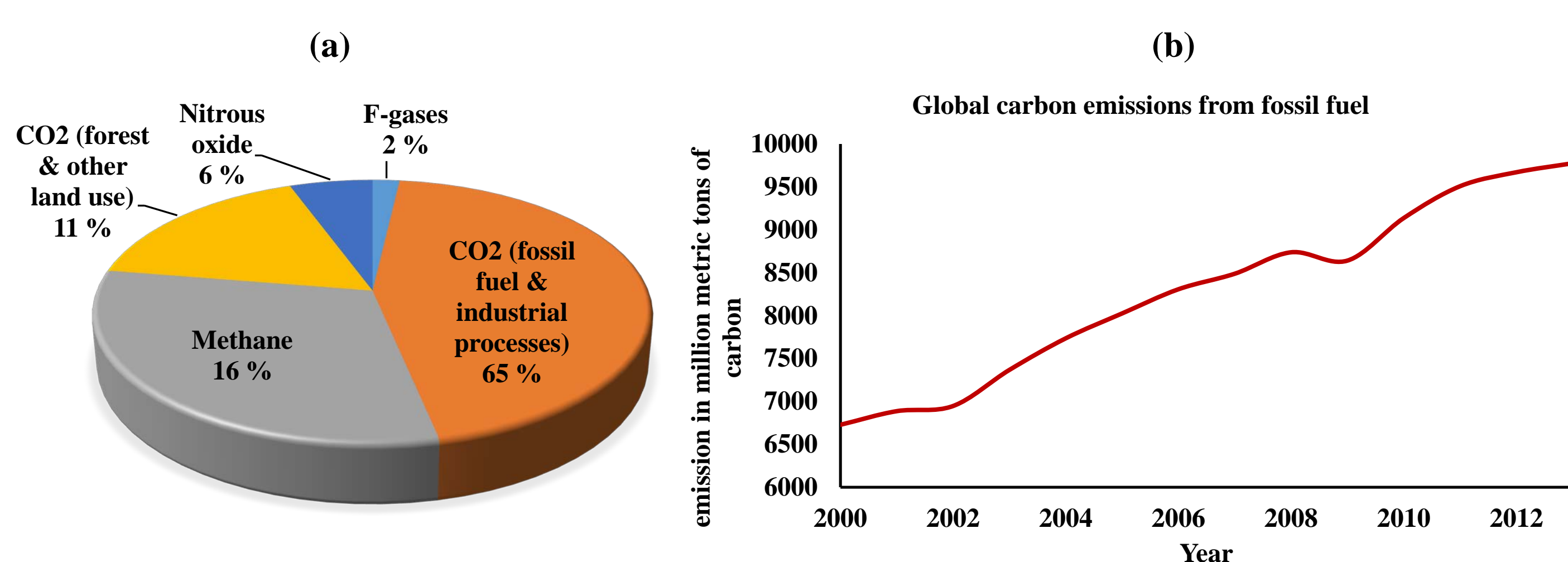
^bCranfield University, Centre for Combustion, Carbon Capture & Storage, MK43 0AL, Cranfield, UK

*corresponding author. Email: k.a.fayemiwo@lboro.ac.uk



Introduction

Combustion of fossil fuels for energy and transport is highly responsible for CO₂ emission, a major greenhouse gas contributing to an increasing global warming. Carbon capture and storage (CCS) has been regarded as the best approach to reduce CO₂ released into the atmosphere. Various CCS technologies include: physical absorption, chemical absorption, adsorption, membrane separation; however, each of these technologies has its own inherent limitations such as high equipment corrosion rate, high energy requirement, poor selectivity, operational limitation, toxicity and environmental unfriendly. In this work, a Porous Polymeric Material (PPM) with CO₂-philic NH₂ groups from non-toxic, inexpensive and readily available materials was synthesized and its CO₂ adsorption capacity was investigated.



(a) Global greenhouse gas emissions in 2010 (b) Global carbon emissions from fossil fuel

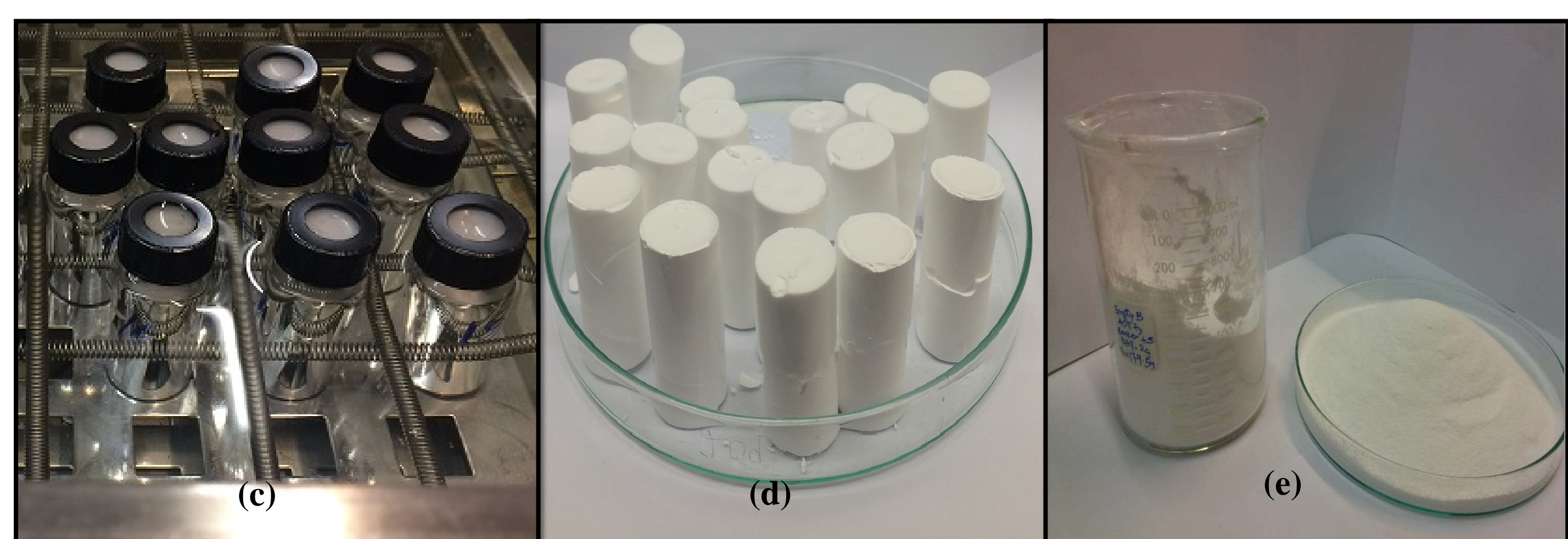
Methodology

Materials

- Methacrylamide (MAAM) – Functional monomer
- Ethylene glycol dimethacrylate (EGDMA) - crosslinker
- Azobisisobutyronitrile (AIBN) – initiator
- Acetonitrile – porogent
- Methanol - solvent

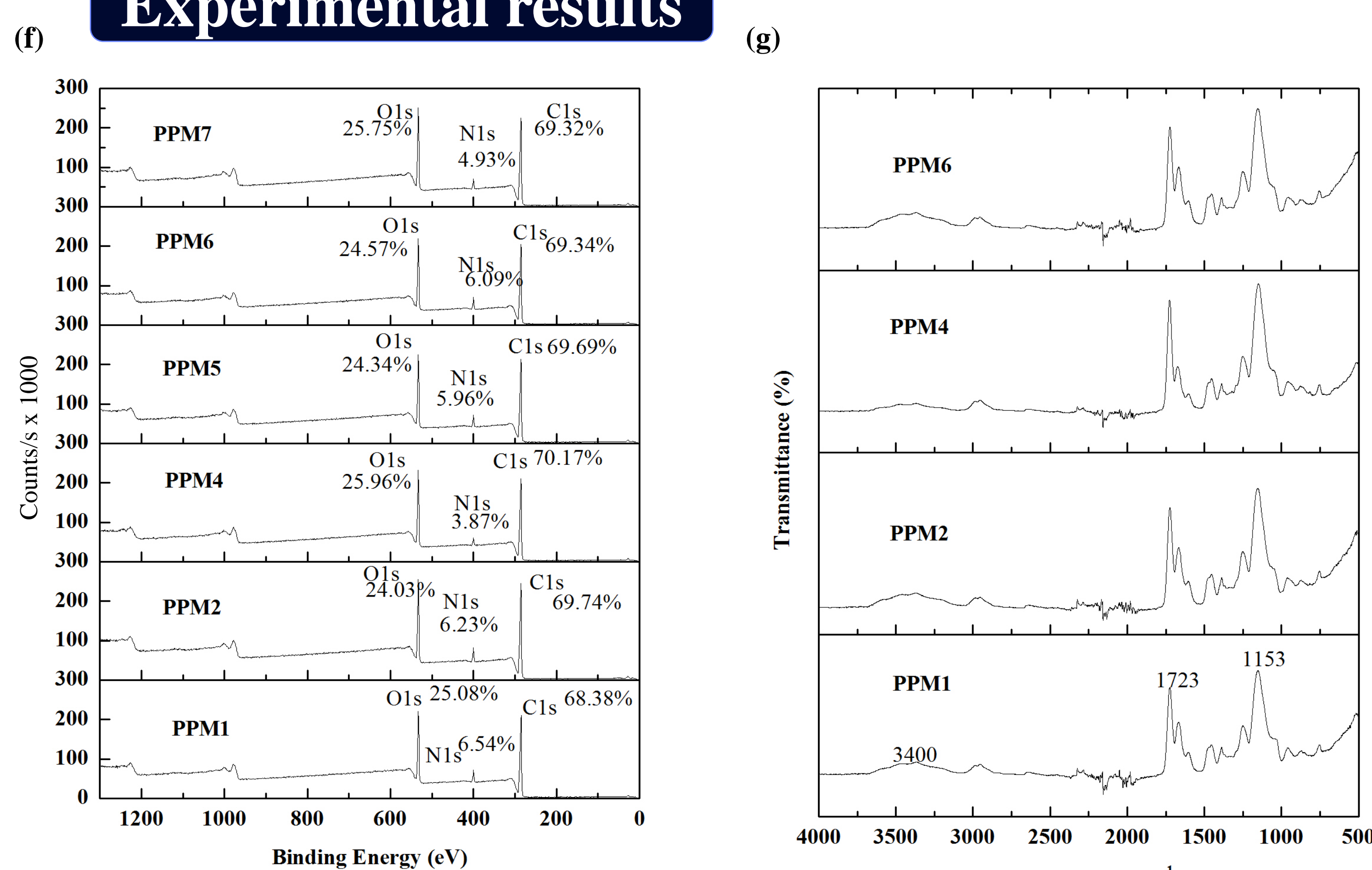
Experimental

MAAM was dissolved in AN followed by adding EGDMA and AIBN. The mixture was degassed and purged with N₂, then sealed up and placed in closed water bath (60 °C) for 24 h. The resultant bulk polymer particles were ground and screened to 90-212 μm, washed with methanol, and dried overnight in a vacuum oven (60 °C).



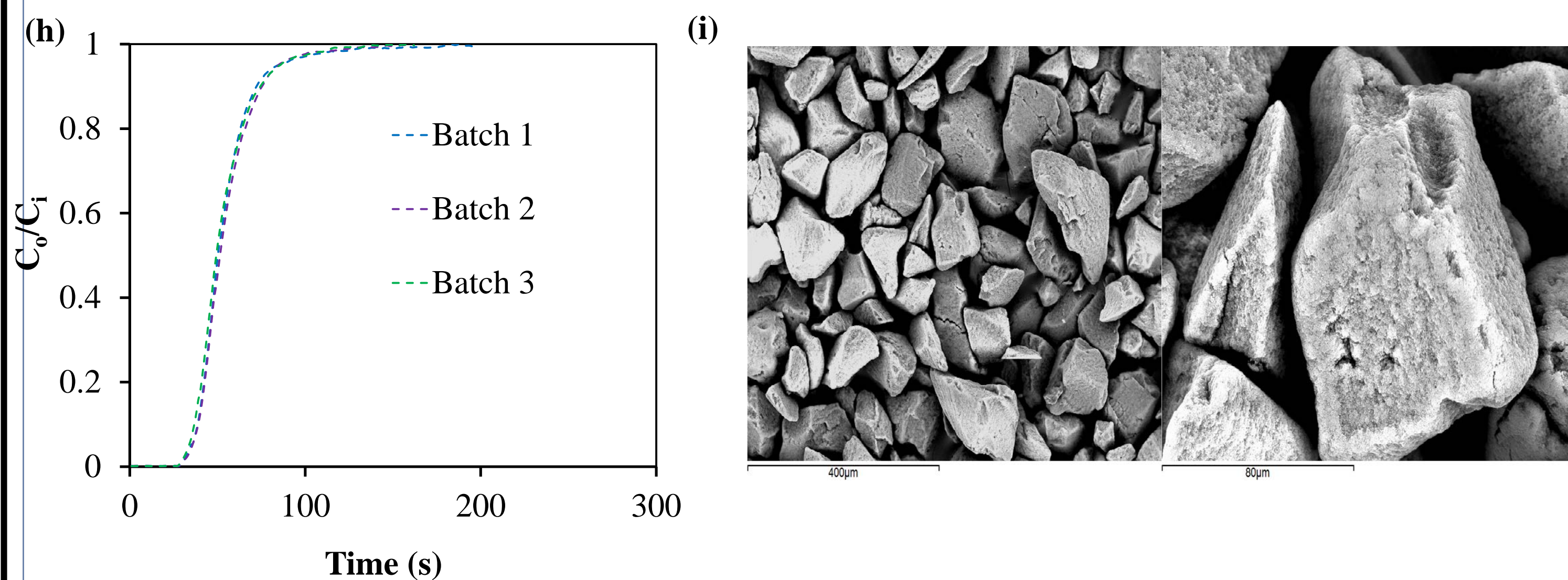
(c) Prepared mixture placed in water bath (d) the bulk sample after polymerization (e) the final product after grinding, sieving and drying

Experimental results



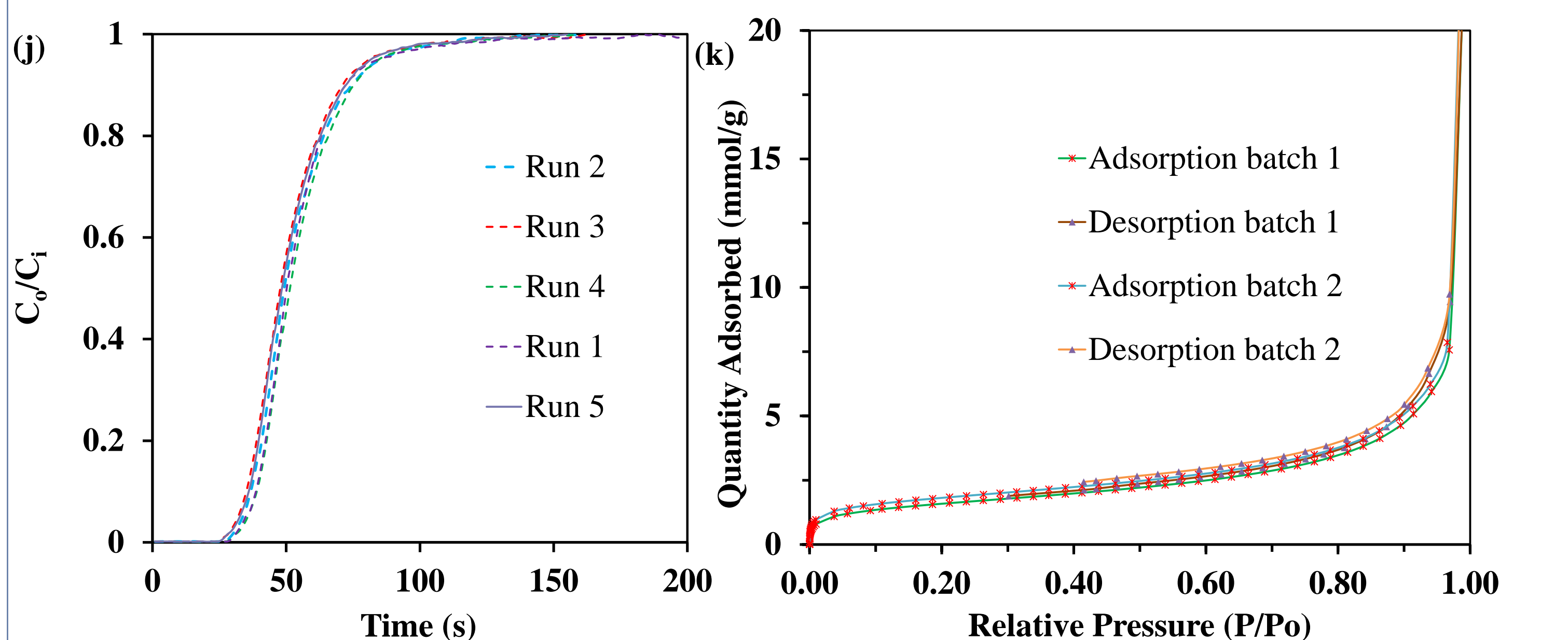
(f) XPS survey Al K photoelectron spectra of PPM.

(g) FTIR spectra of PPMs.



(h) Reproducibility test based on of PPMs.

(i) SEM images of polymer PPM particles



(j) Re-usability test based on of PPMs.

(k) Nitrogen adsorption/desorption isotherms measured at 77 K.

Conclusions

- A series of simple, inexpensive, non-toxic and environmental friendly PPM was developed for CO₂ adsorption with a promising CO₂ capture capacity.
- The adsorbents retained its -NH₂ functional group of the based monomer and also, the C=C of the monomer, MAAM and cross-linker, EGDMA were completely broken as confirmed in the XPS and FTIR analysis.
- All the adsorption isotherms of PPMs as shown exhibit a typical shape of type II featuring a non-uniform distribution of pore size.
- The PPM exhibited CO₂ uptake capacity up to 0.64 mmol/g at 313 K and 0.15 bar CO₂ partial pressure and consistent in both reusability and reproducibility test run.

References

- [1] S.-Y. Lee and S.-J. Park, "A review on solid adsorbents for carbon dioxide capture," *J. Ind. Eng. Chem.*, 2015, (23) pp. 1–11.
- [2] J. Gibbins and H. Chalmers, "Carbon capture and storage," *Energy Policy*, 2008, pp. 4317–22.
- [3] Nabavi, S. A.; Vladislavljević, G. T.; Wicaksono, A.; Georgiadou, S.; Manović, V. Production of Molecularly Imprinted Polymer Particles with Amide-Decorated Cavities for CO₂ Capture Using Membrane Emulsification/suspension Polymerization. *Colloids Surfaces A Physicochem. Eng. Asp.* 2016.
- [4] Nabavi, S. A.; Vladislavljević, G. T.; Eguagie, E. M.; Li, B.; Georgiadou, S.; Manović, V. Production of Spherical Mesoporous Molecularly Imprinted Polymer Particles Containing Tunable Amine Decorated Nanocavities with CO₂ Molecule Recognition Properties. *Chem. Eng. J.* 2016, 306, 214–225.