

New carbon capture materials: Novel Approaches to Post-Combustion CO₂ Capture

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Introduction

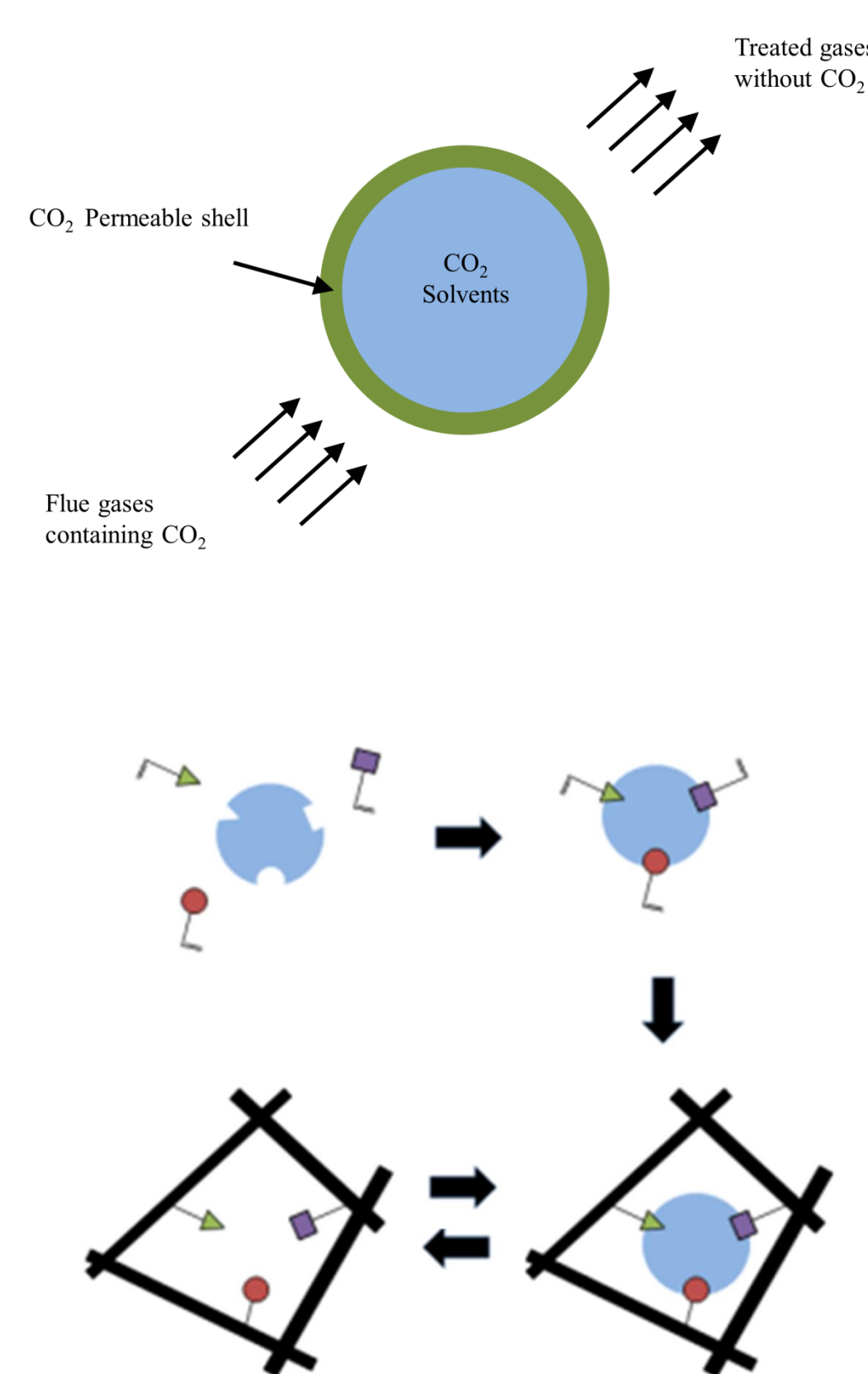
The most commercially viable capture method in carbon capture and storage (CCS) has been attributed to post-combustion carbon capture using chemical solvents. Although the conventional chemical solvents such as MEA solutions have high selectivity and capture capacity, they are highly corrosive and required high regeneration energy. In addition, volatilisation of MEA at elevated temperature and its release to the atmosphere can lead to major human and environment concerns. In this study two alternative carbon capture materials have been investigated.

- **CO₂ solvent microcapsules** where the CO₂ solvents are encapsulated within a CO₂ permeable polymer shells.

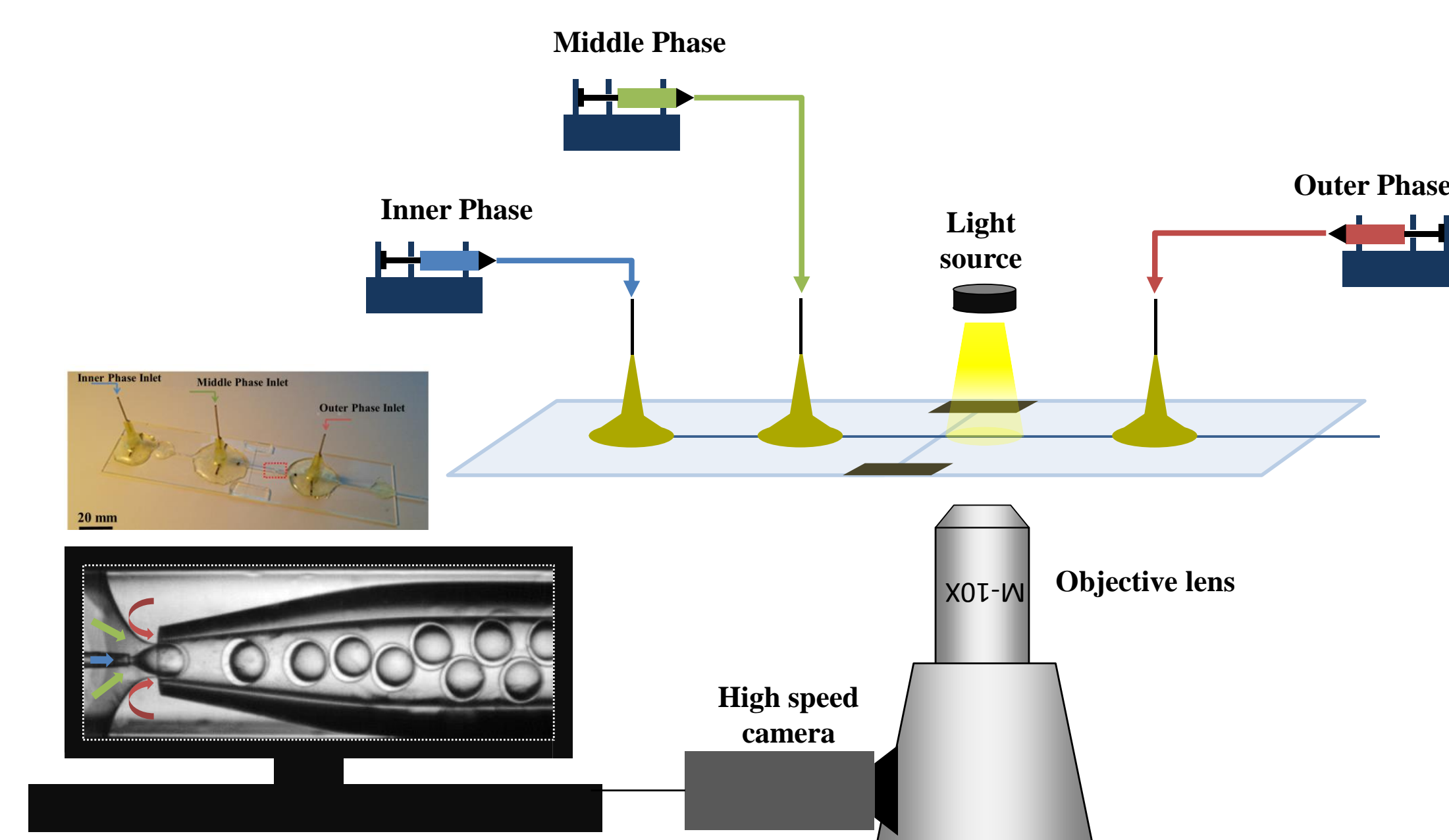
- Adv. 1.** Prevents direct contact of solvents with system.
- Adv. 2.** Reduction in solvent volatilisation.
- Adv. 3.** Provides much larger surface area and consequently increases the capture rate.

- **CO₂ based imprinted polymers (CO₂-MIPs)** where recognition cavities are formed within the polymer, based on the target molecules (template)

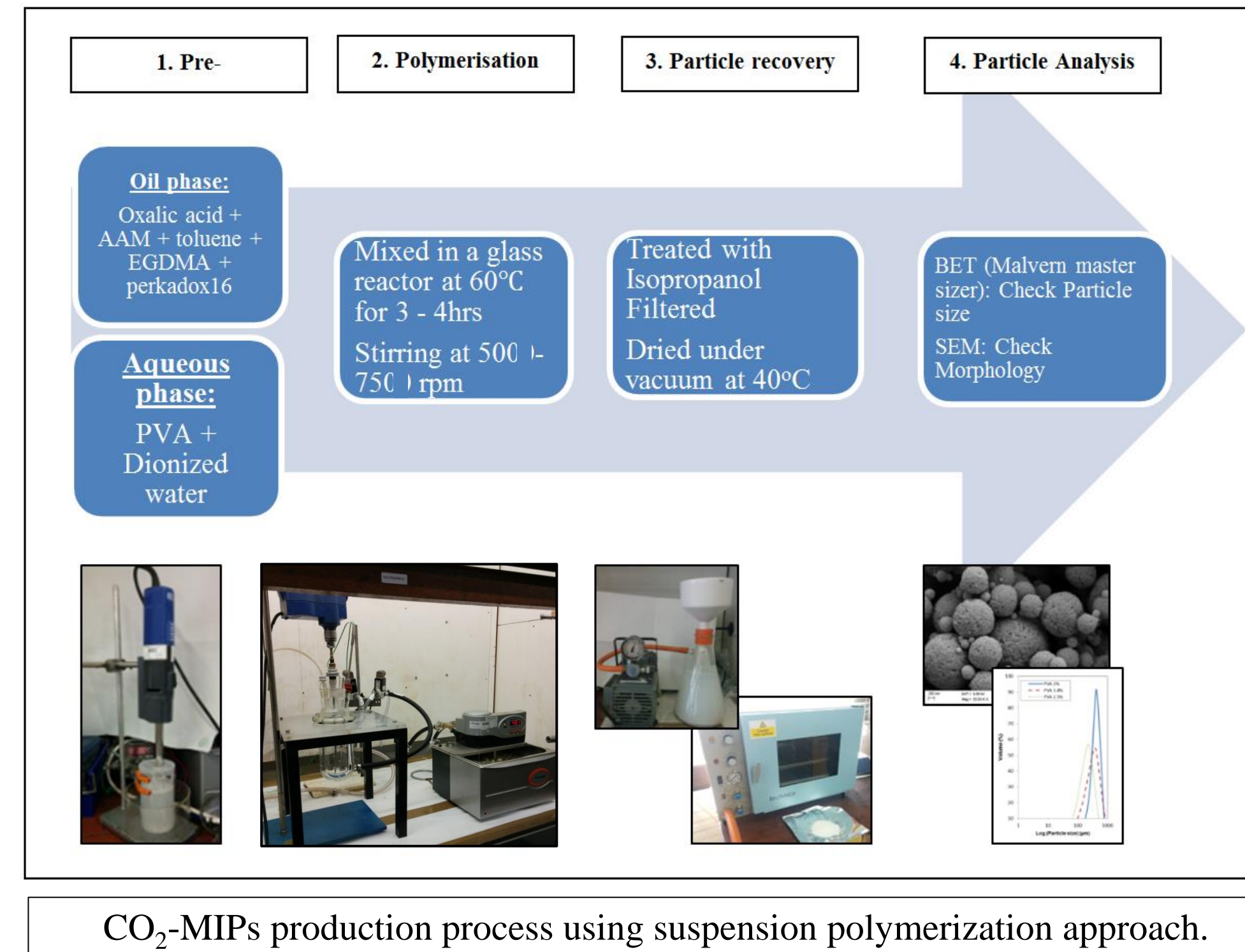
- Adv. 1.** High selectivity.
- Adv. 2.** Stable capture efficiency in present of impurities.
- Adv. 3.** Stable capture efficiency in repetitive cycles.
- Adv. 4.** Lower required regeneration energy.



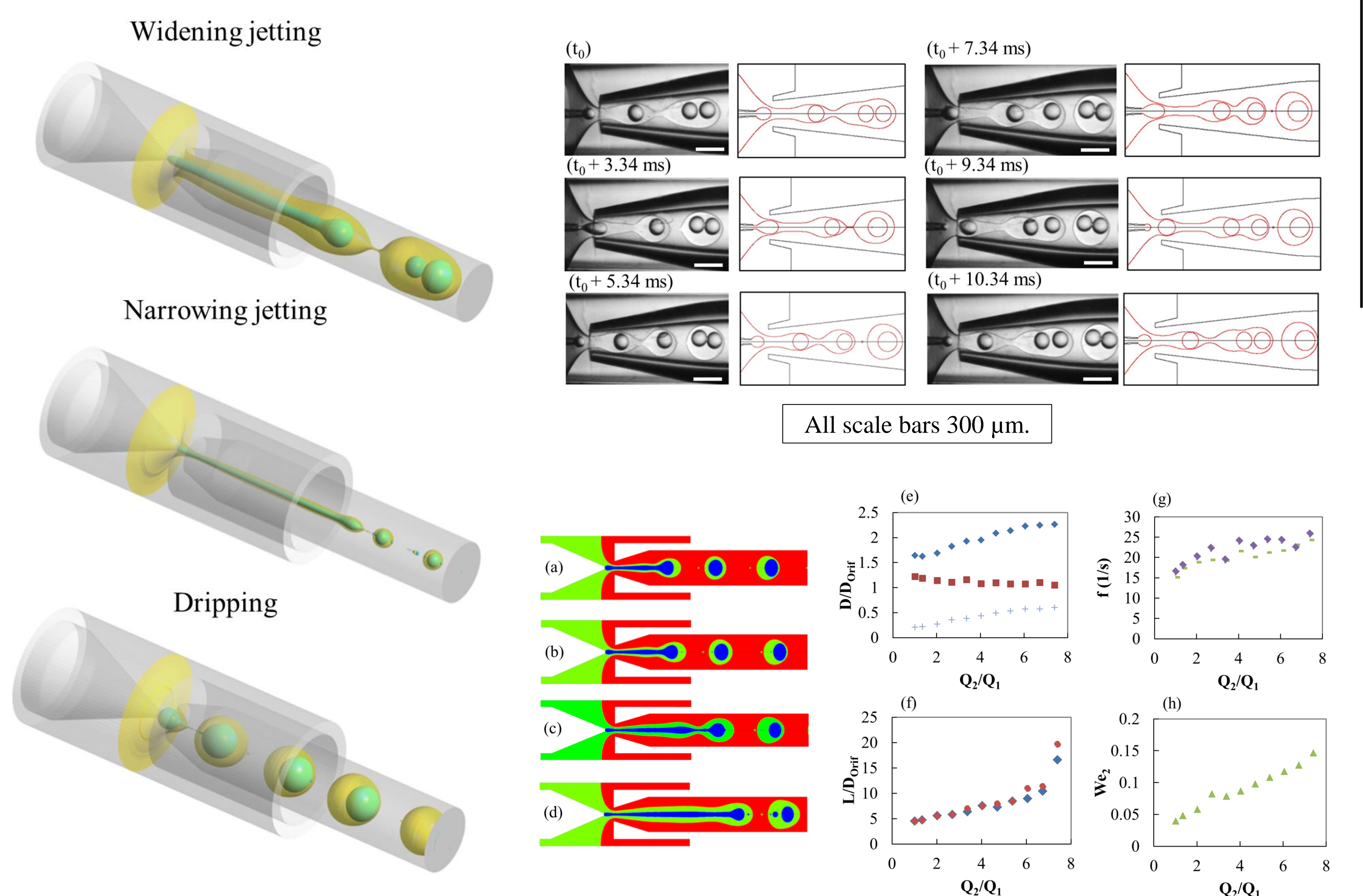
Experimental Setup



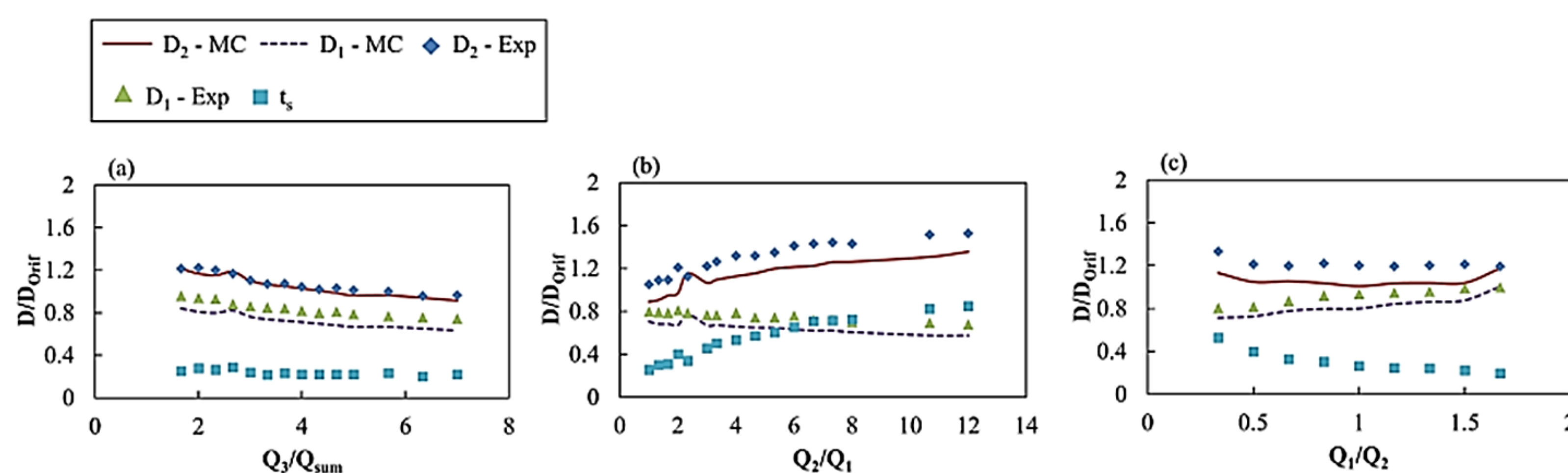
A fabricated three-phase glass capillary device for production of CO₂ solvent microcapsules. The direction of inner, middle and outer fluid is shown by the blue, green and red arrow, respectively.



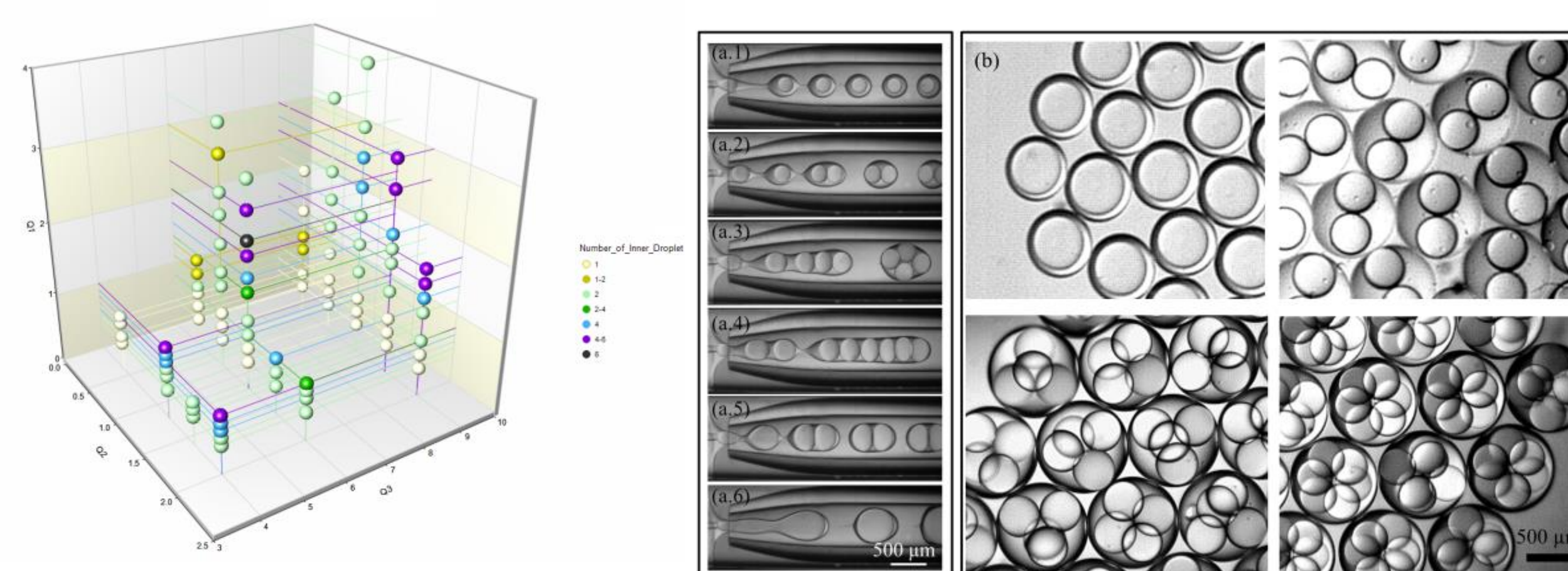
Numerical and Experimental Results



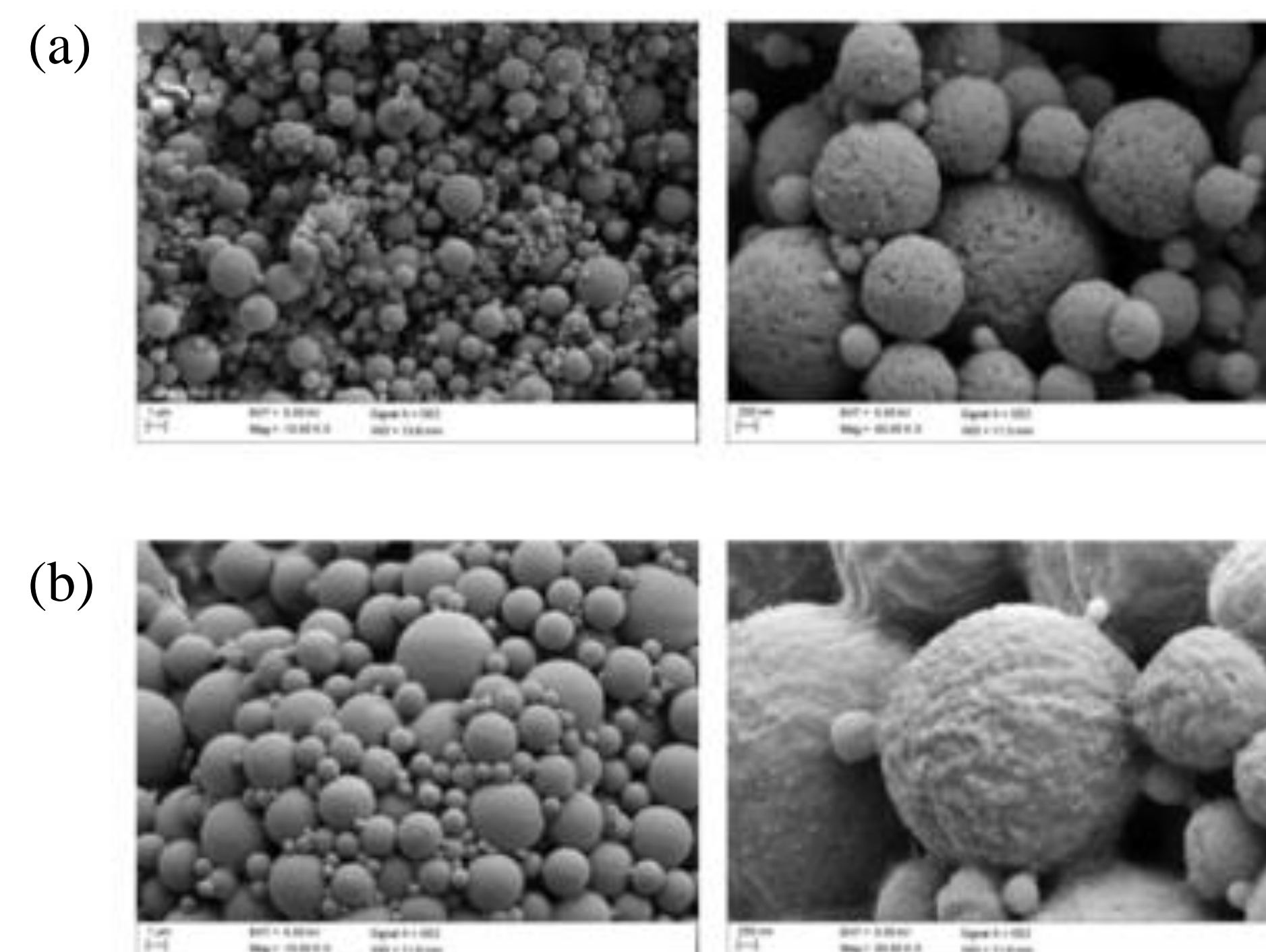
A VOF/CSF Numerical model has been developed and validated with experiments and analytical solutions.



Experimental data: The size of inner and outer droplets, and the shell thickness as a function of phase flow rates.



The smaller inner or outer droplets, the higher reacting area with flue gases. Therefore, beside controlling the size of inner and outer droplet, the number of encapsulated droplet based on flow rates has been controlled.



SEM images of CO₂-MIPs morphology as a function of initiators: (a) Perkadox 16 (case 3), (b) AIBN; (c) the particle size distributions.

Conclusions

Production of two promising carbon capture material was investigated. Regarding the CO₂ solvent microcapsules both experiments and numerical modelling were used to study the effect of flow rates, fluid properties and microfluidic geometry to achieve an active control on the microcapsule size, shell thickness and the number of encapsulated inner droplets. The microcapsules with size over the range of 50-600 μm were produced. Concerning CO₂-MIPs, suspension polymerisation method was used to achieve spherical particles with controllable size over the range of 1-100 μm. The effect of operative parameters on particle morphology has been investigated. Both materials due to wide range of particle size can be used for industrial and domestic applications.