

Supporting Information

Preparation of catalytic deep eutectic solvents-based silica microreactors using a non-aqueous sol-gel route

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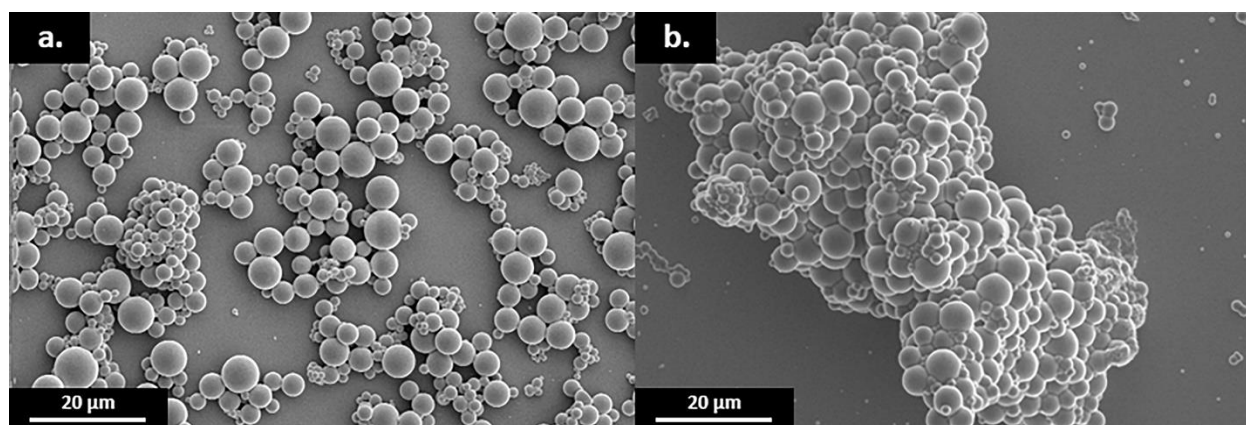


Figure S1. SEM images of microcapsules prepared from 20% core instead of 36% a. CHCl:TA@SiO₂ using xylene as the solvent in the dispersing phase and b. CHCl:S@SiO₂ using cyclohexane as the solvent in the dispersing phase.

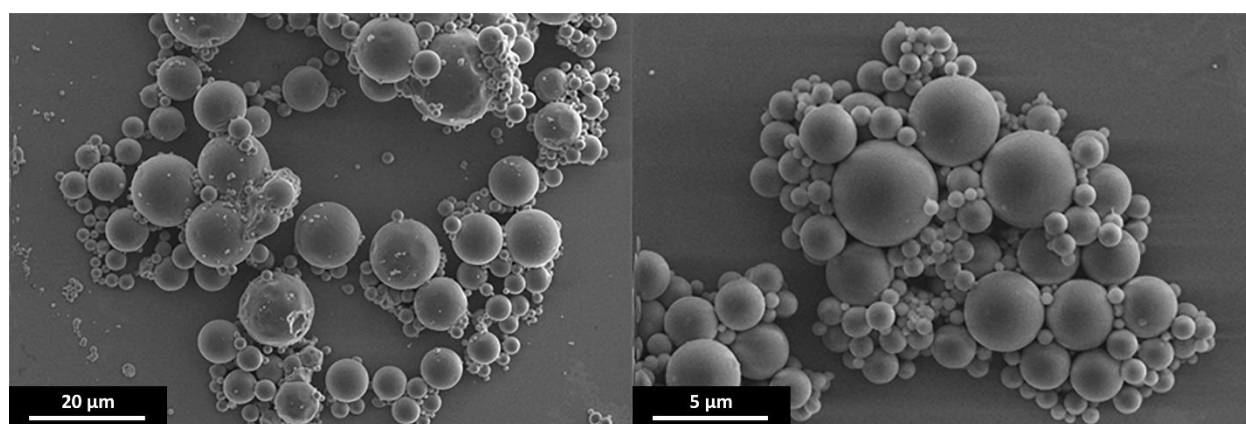


Figure S2. SEM images of CHCl:TSA@SiO₂ prepared using L-lactic acid instead of formic acid for the non-aqueous sol-gel process.

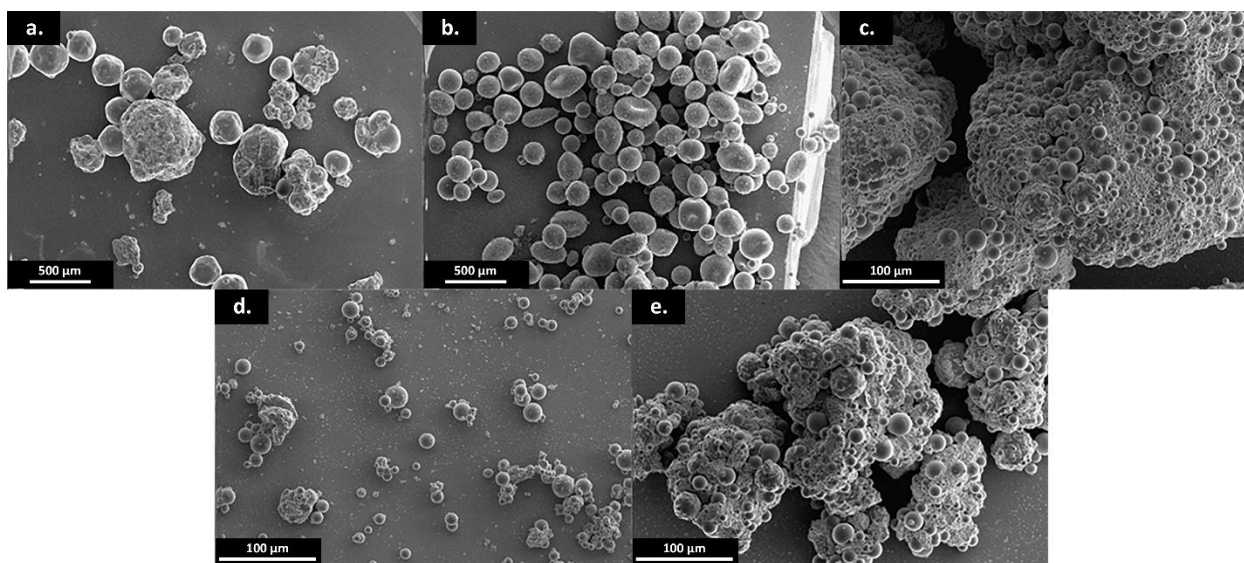


Figure S3. SEM images of CHCl:TA@SiO₂ microcapsules prepared using a. 0.05 g, b. 0.1 g, c. 0.2 g, d. 0.4 g, and e. 0.8 g of ABIL EM 90 and using xylene as the solvent in the dispersing phase.

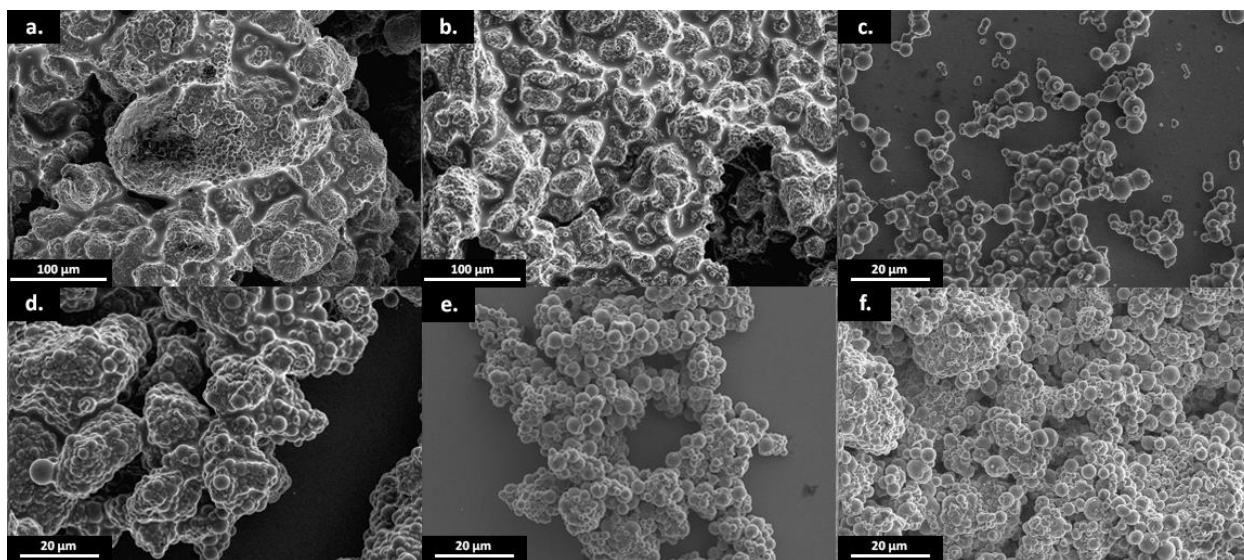


Figure S4. SEM images of CHCl:S@SiO₂ microcapsules prepared using a. 0.05 g, b. 0.2 g, c. 0.4 g, d. 0.8 g, e. 1 g, and f. 1.25 g of ABIL EM 90 and using heptane as the solvent in the dispersing phase.

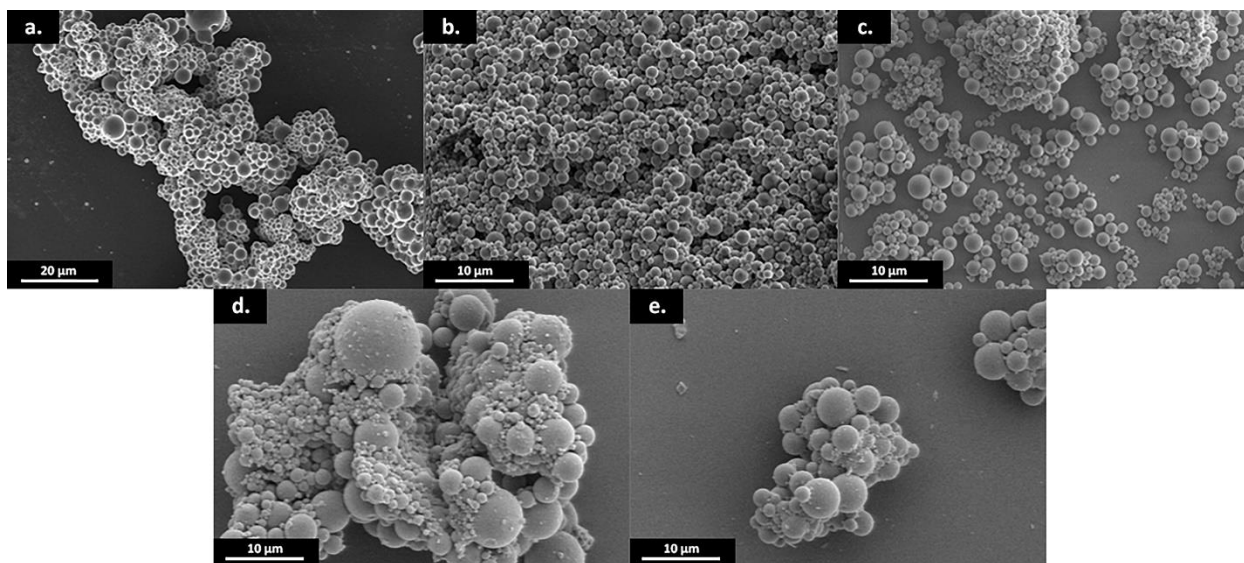


Figure S5. SEM images of a. CHCl:S@SiO₂, b. CHCl:U@SiO₂, c. CHCl:LA@SiO₂, d. CHCl:TSA@SiO₂, and e. CHCl:CSA@SiO₂ microcapsules prepared according to the general procedure in the experimental section but with PDMS as the silane monomer.

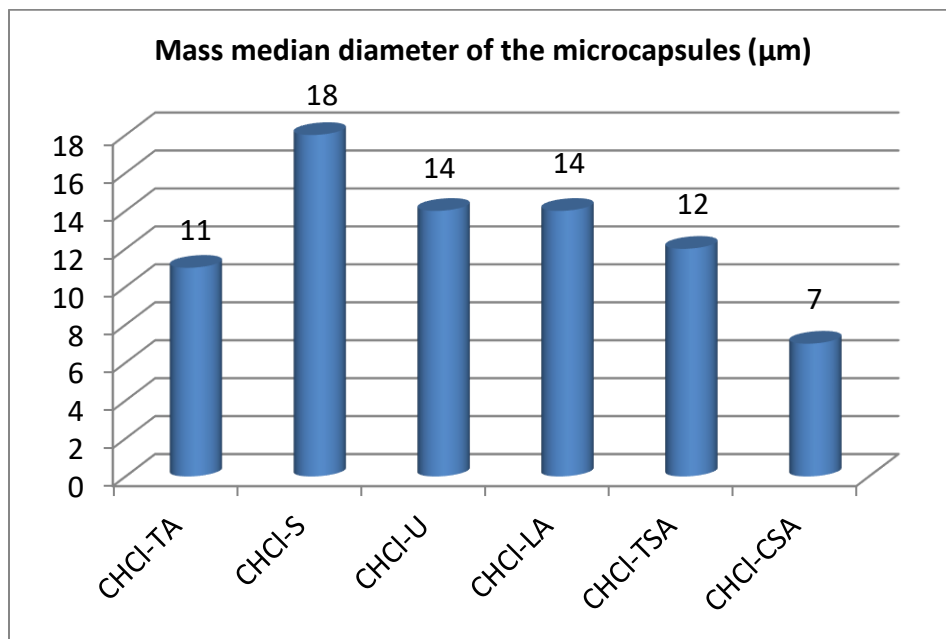


Figure S6. Mass median diameter in microns of the microencapsulated DESs according to Mastersier analysis.

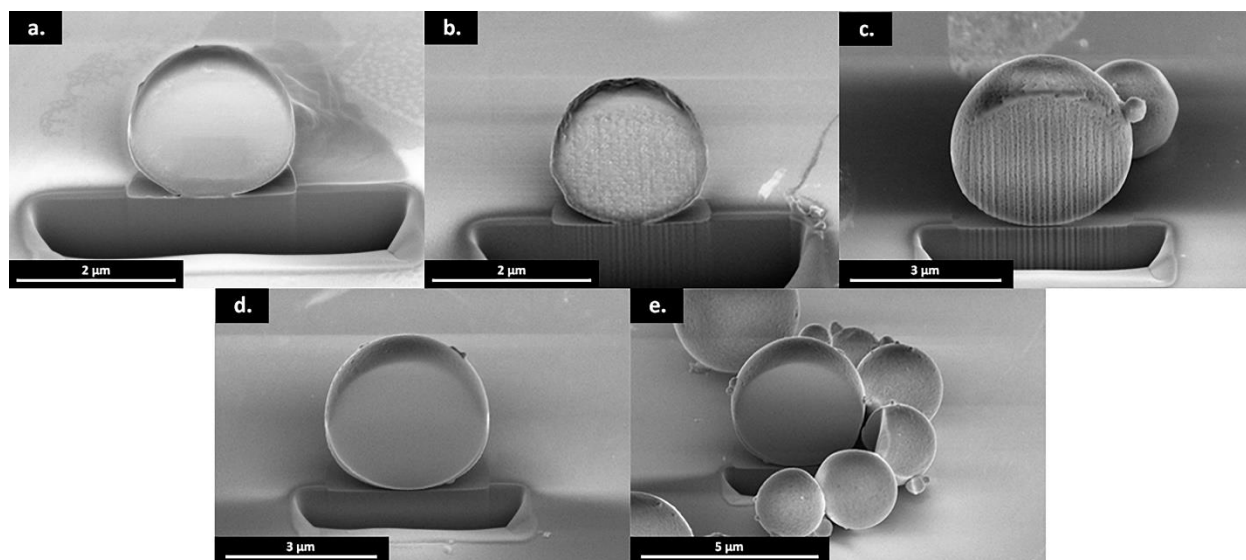


Figure S7. FIB images of a. CHCl:S@SiO₂, b. CHCl:U@SiO₂, c. CHCl:LA@SiO₂, d. CHCl:TSA@SiO₂, and e. CHCl:CSA@SiO₂ microcapsules prepared according to the general procedure in the experimental section but with PDMS as the silane monomer.

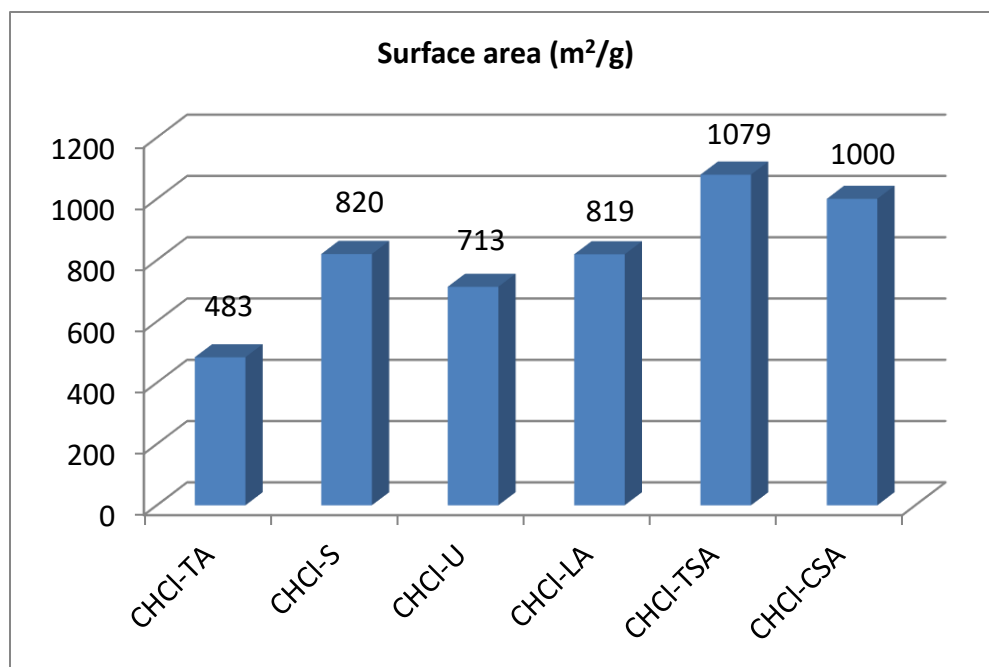


Figure S8. Surface area (m²/g) of the microencapsulated DESs according to BET analysis.

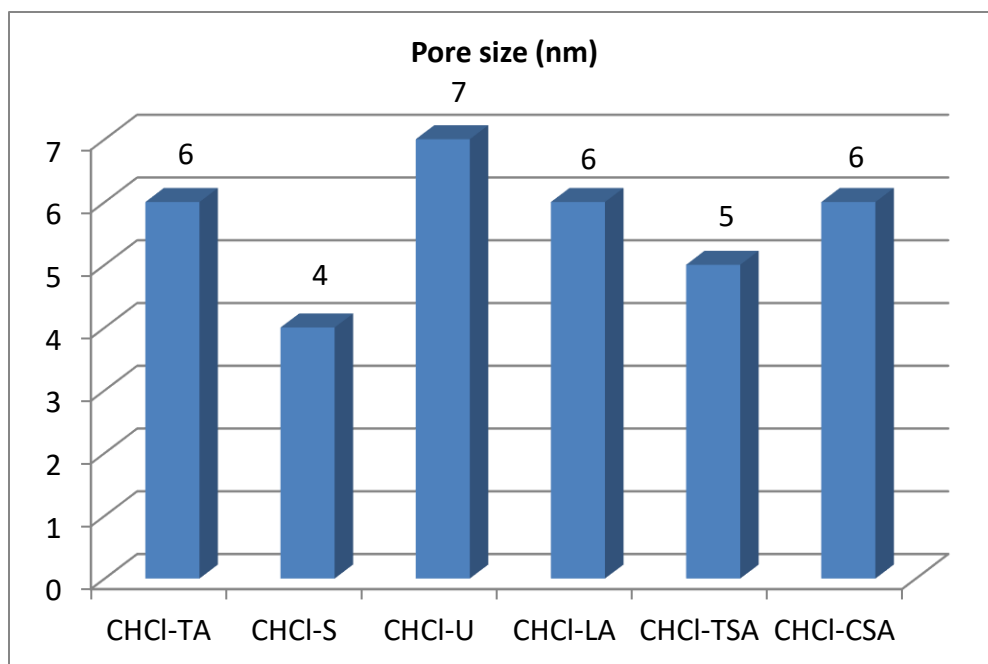


Figure S9. Pore size (nm) distribution of the microencapsulated DESs according to BET analysis.

¹³C-CPMAS
065-030-1

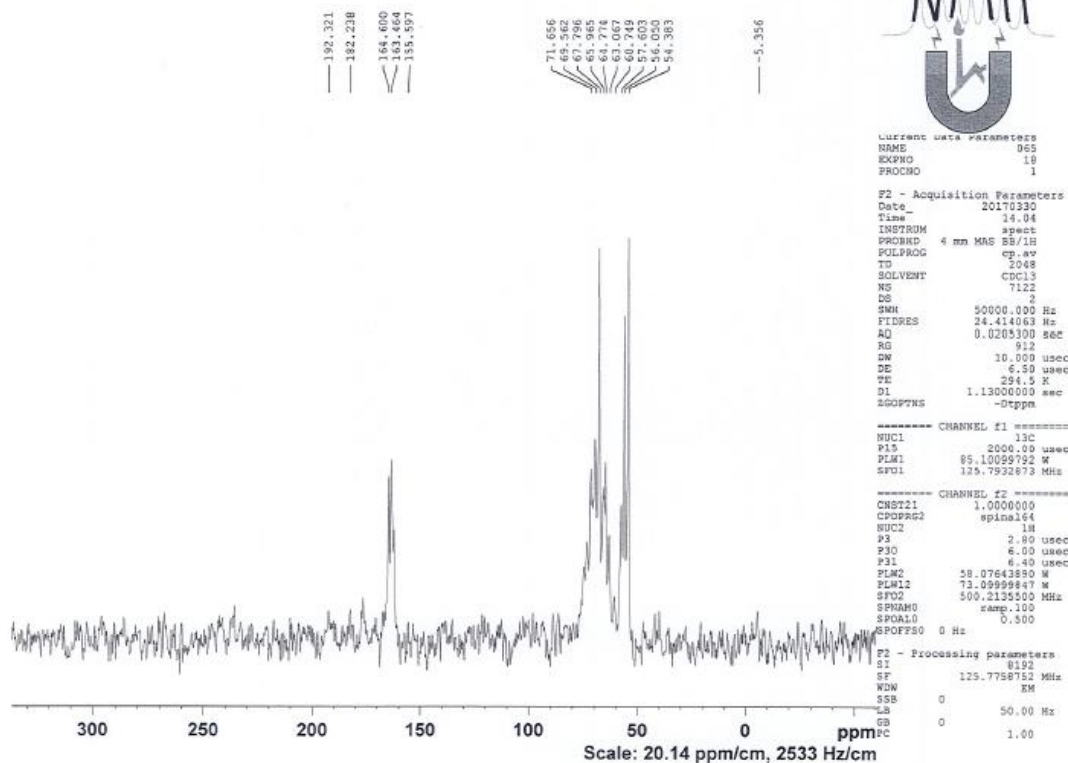


Figure S10. ¹³C CP-MAS NMR of CHCl:S@SiO₂ microcapsules.

²⁹Si-CPMAS
065-030-1

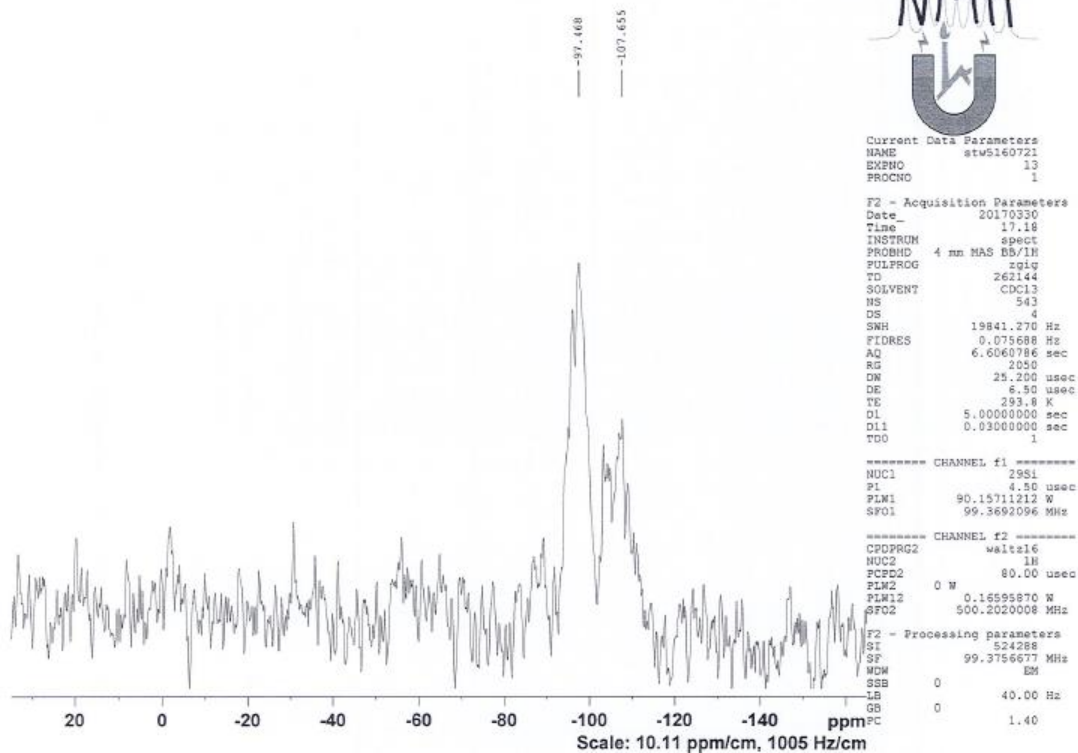


Figure S11. ²⁹Si CP-MAS NMR of CHCl:S@SiO₂ microcapsules

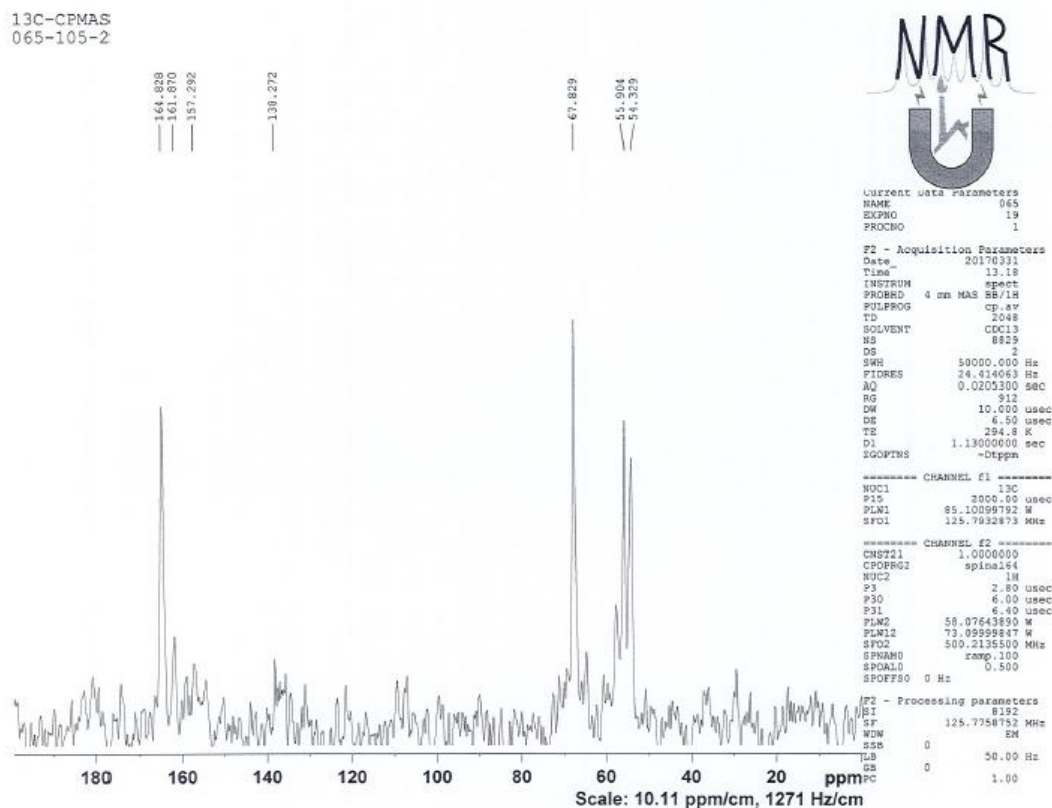


Figure S12. ^{13}C CP-MAS NMR of CHCl:U@SiO_2 microcapsules.

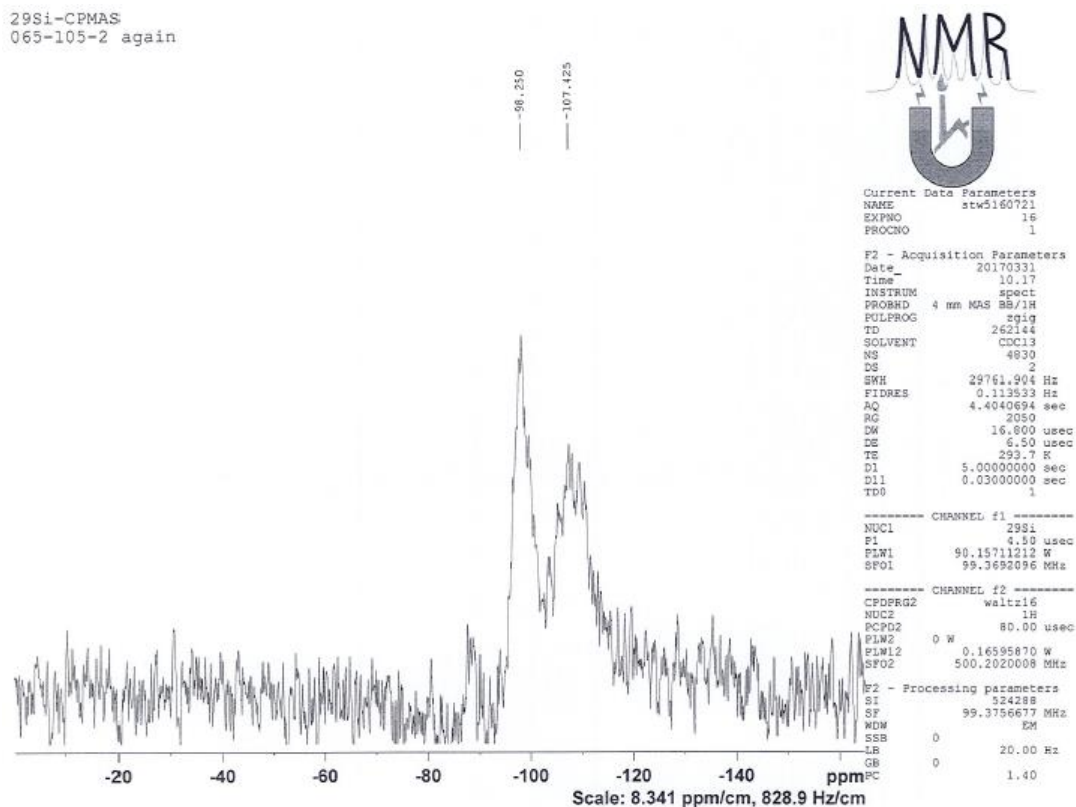


Figure S13. ^{29}Si CP-MAS NMR of CHCl:U@SiO_2 microcapsules.

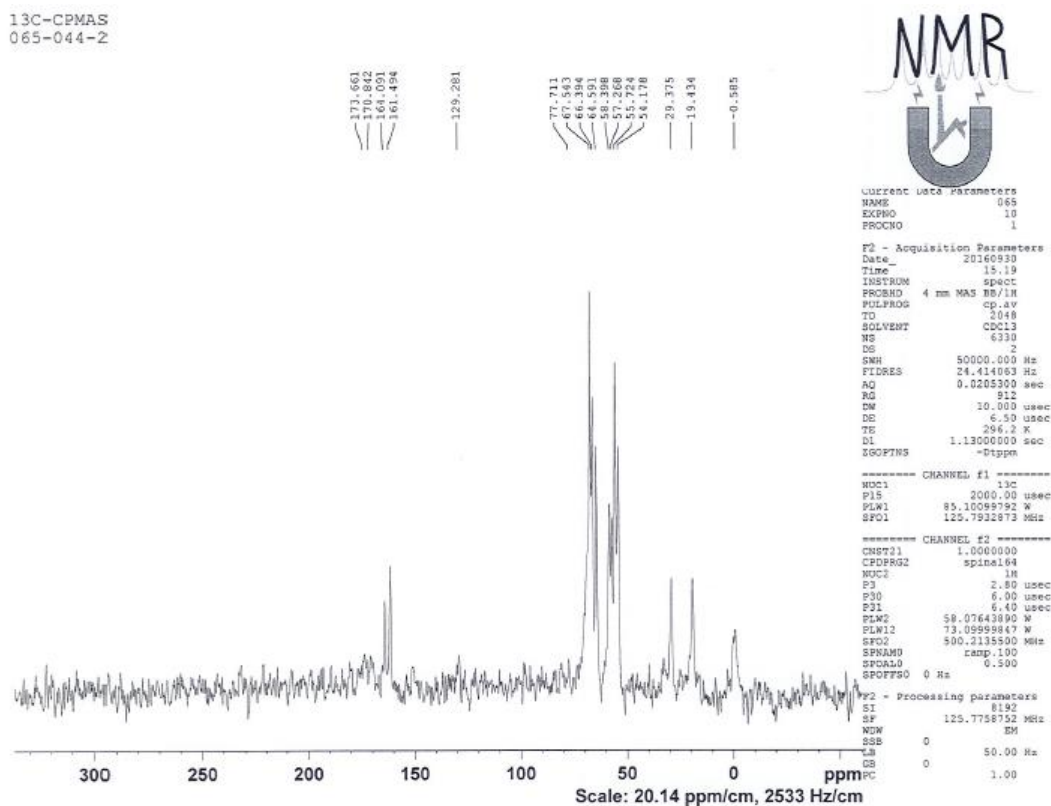


Figure S14. ^{13}C CP-MAS NMR of CHCl:LA@SiO_2 microcapsules

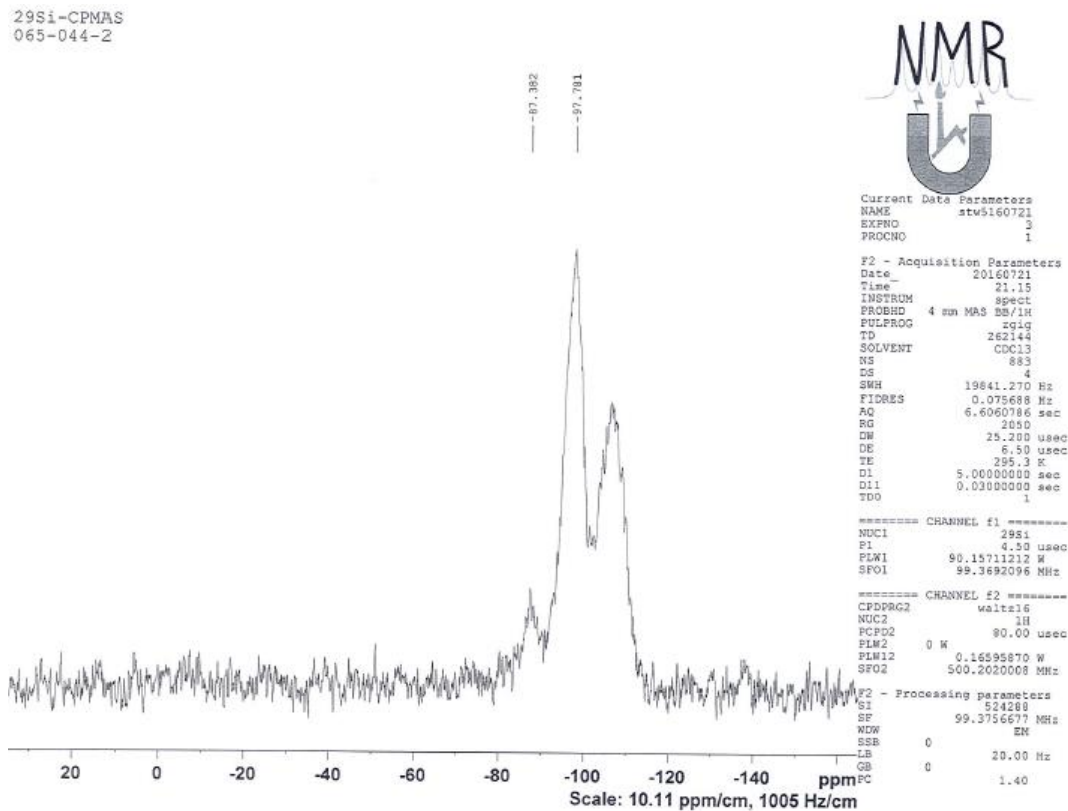


Figure S15. ^{29}Si CP-MAS NMR of CHCl:LA@SiO_2 microcapsules.

13C-CPMAS
065-044-3

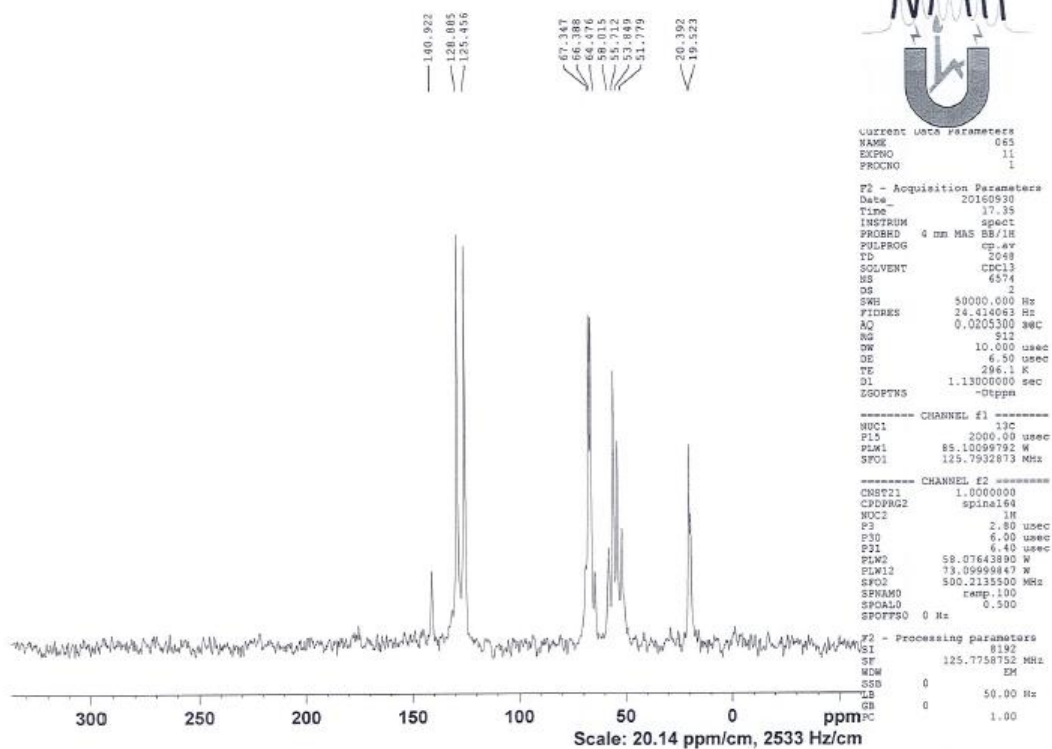


Figure S16. ¹³C CP-MAS NMR of CHCl₃:TSA@SiO₂ microcapsules.

29Si-CPMAS
065-044-3

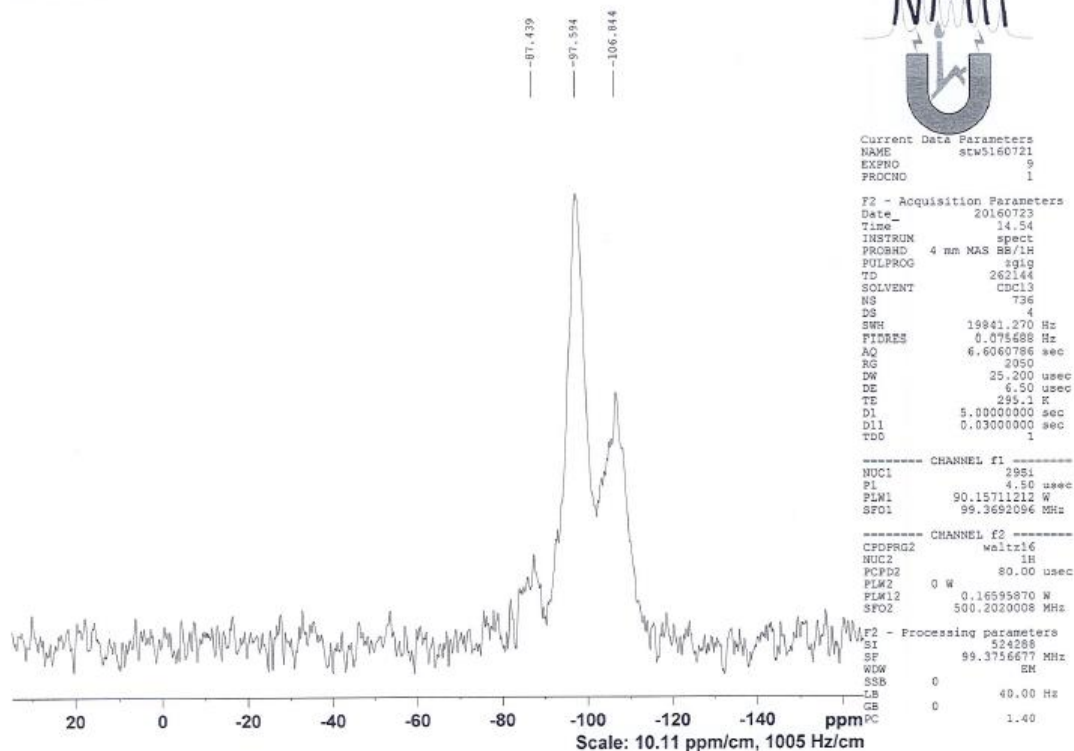


Figure S17. ²⁹Si CP-MAS NMR of CHCl₃:TSA@SiO₂ microcapsules.

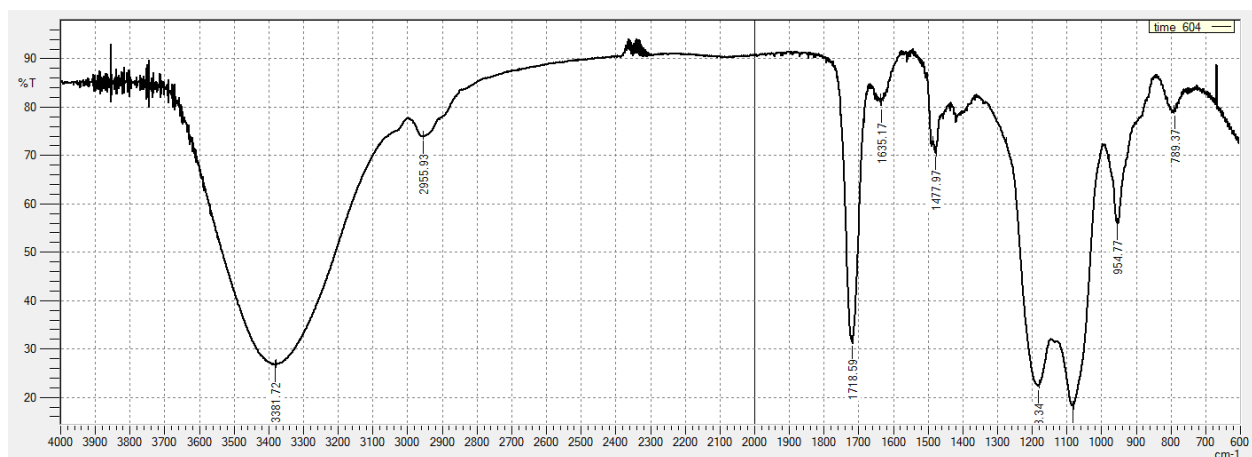


Figure S20. FTIR analysis of CHCl:S@SiO₂ microcapsules.

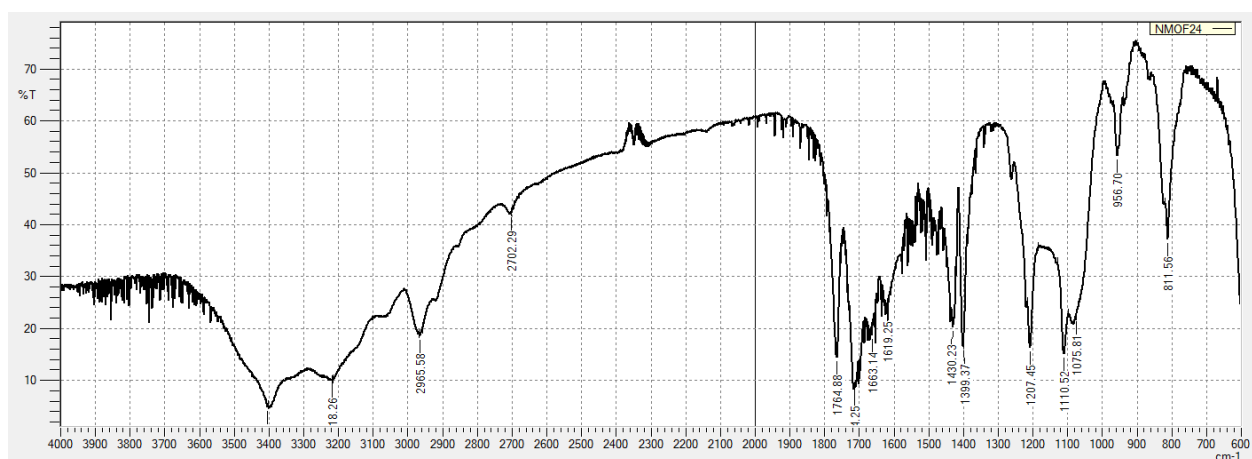


Figure S21. FTIR analysis of CHCl:U@SiO₂ microcapsules.

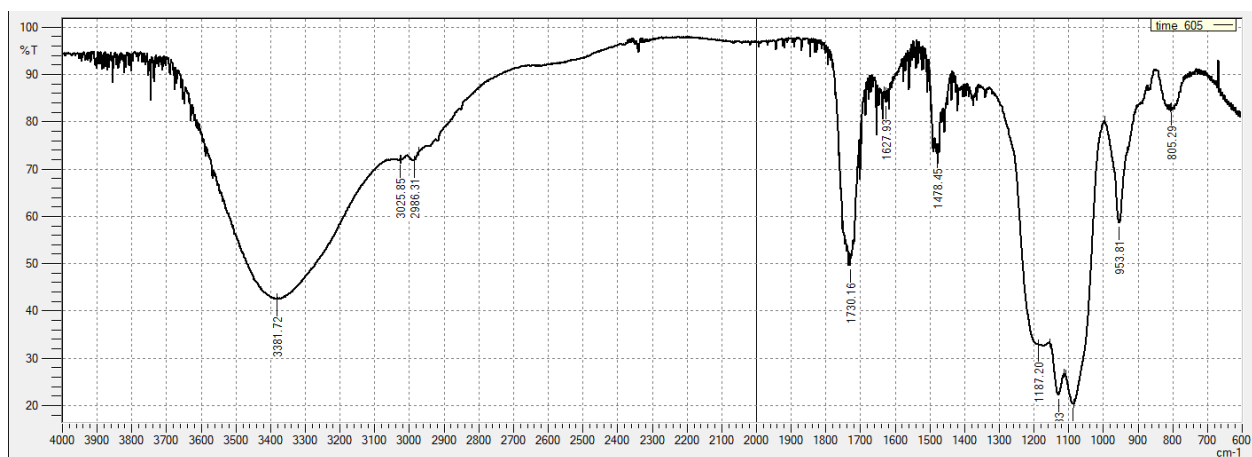


Figure S22. FTIR analysis of CHCl:LA@SiO₂ microcapsules.

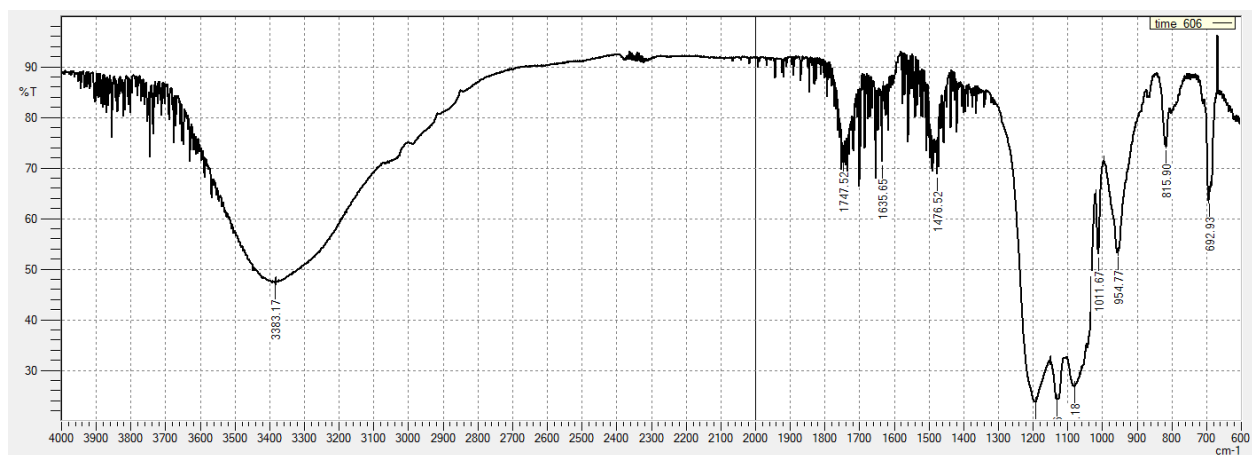


Figure S23. FTIR analysis of CHCl:TSA@SiO₂ microcapsules.

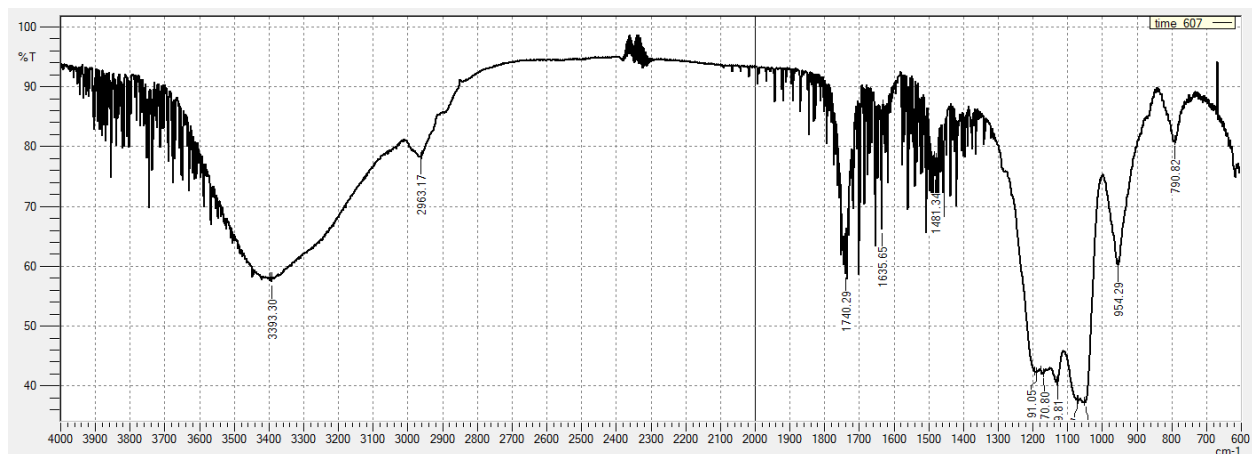


Figure S24. FTIR analysis of CHCl:CPA@SiO₂ microcapsules.

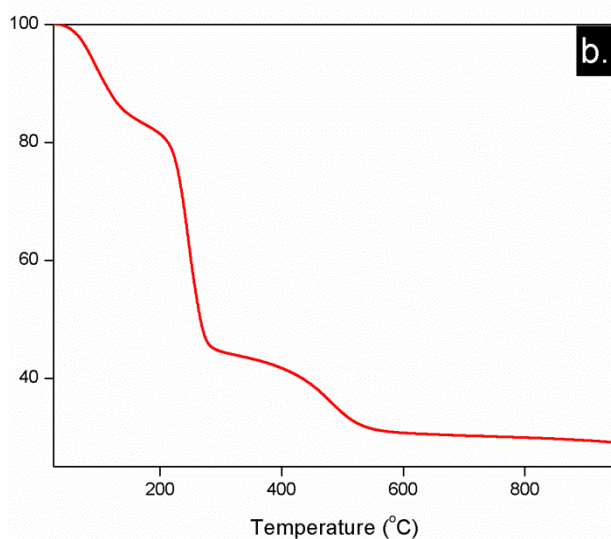
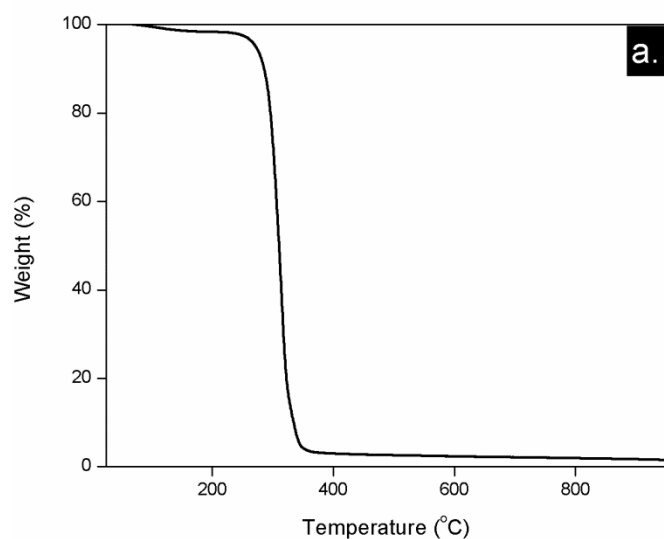


Figure S25. TGA analysis of a. CHCl:S, and b. CHCl:S@SiO₂ microcapsules under N₂ atmosphere.

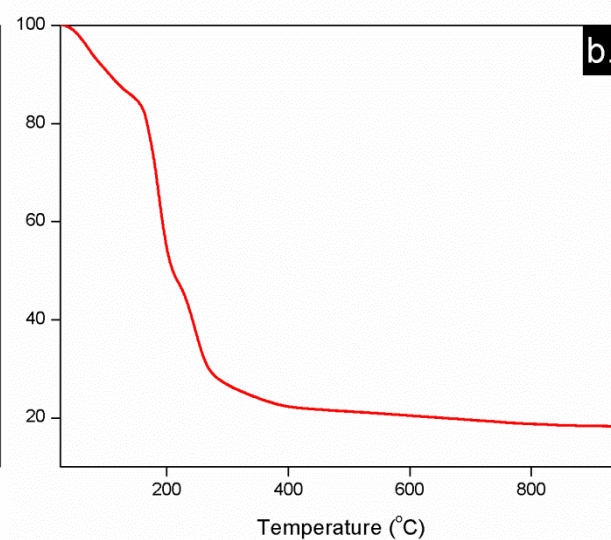
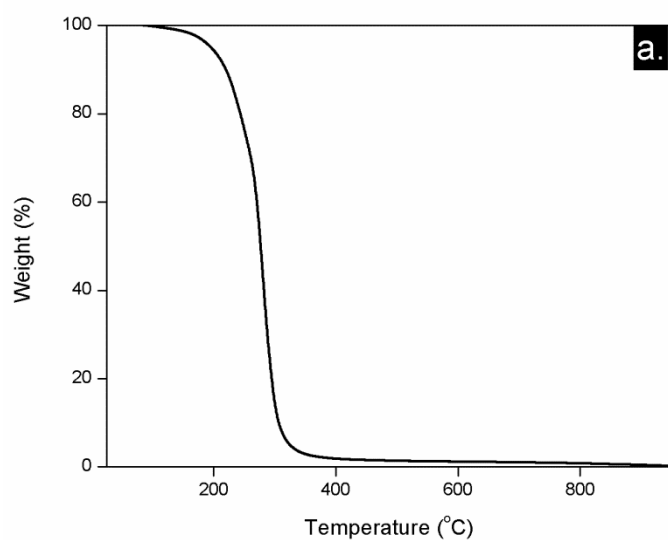


Figure S26. TGA analysis of a. CHCl:U, and b. CHCl:U@SiO₂ microcapsules under N₂ atmosphere.

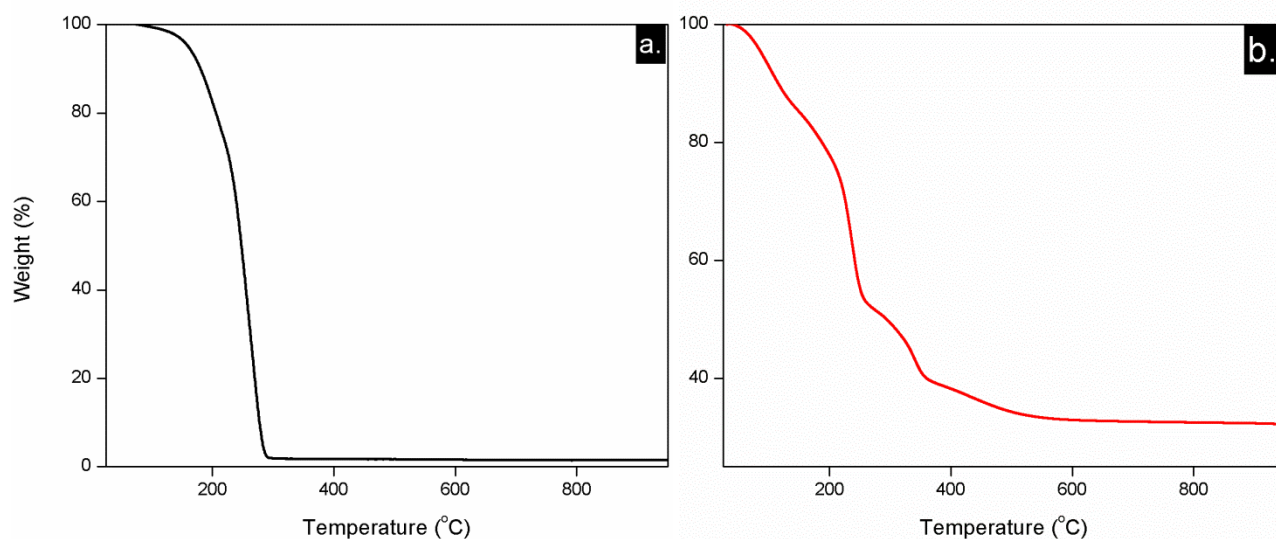


Figure S27. TGA analysis of a. CHCl:LA, and b. CHCl:LA@SiO₂ microcapsules under N₂ atmosphere.

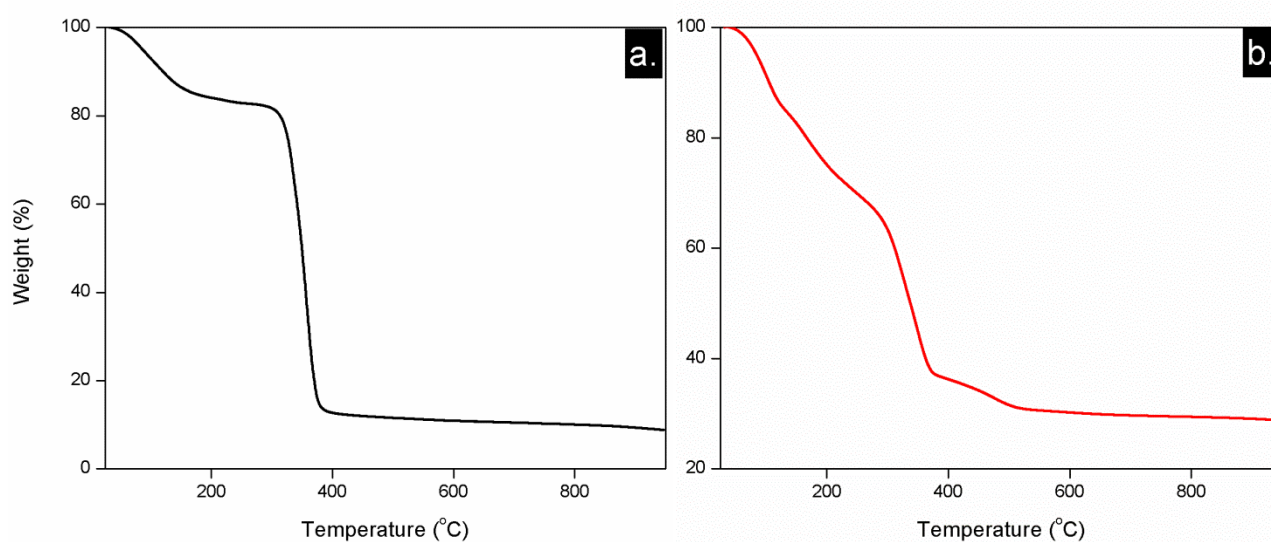


Figure S28. TGA analysis of a. CHCl:TSA, and b. CHCl:TSA@SiO₂ microcapsules under N₂ atmosphere.

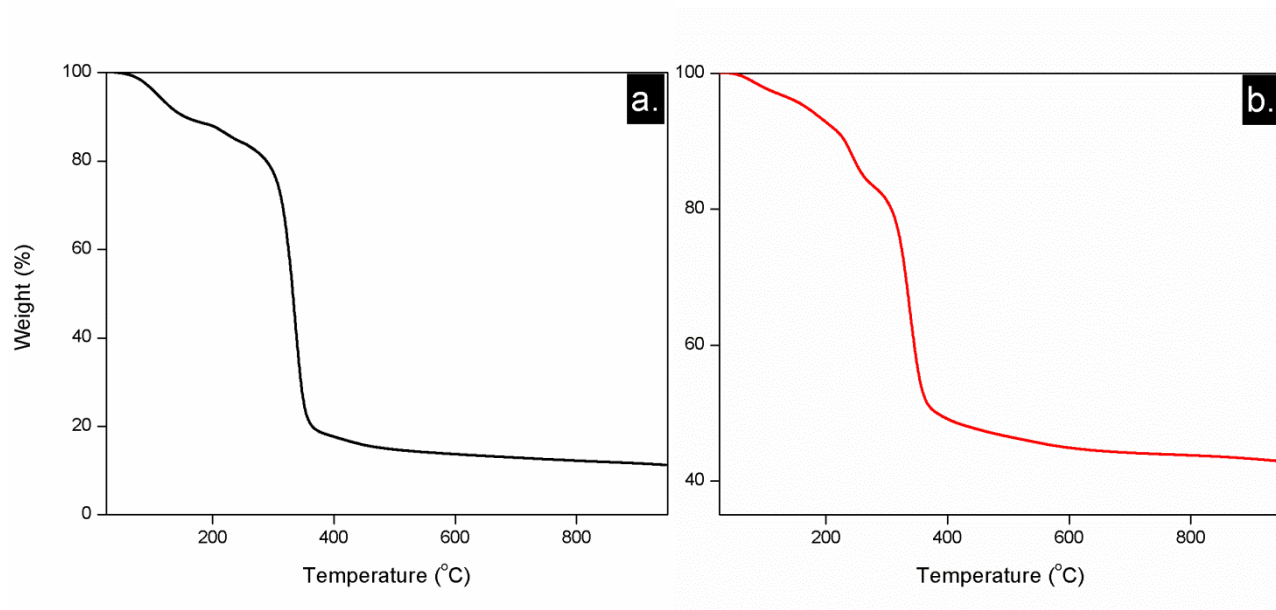


Figure S29. TGA analysis of a. CHCl:CSA, and b. CHCl:CSA@SiO₂ microcapsules under N₂ atmosphere.

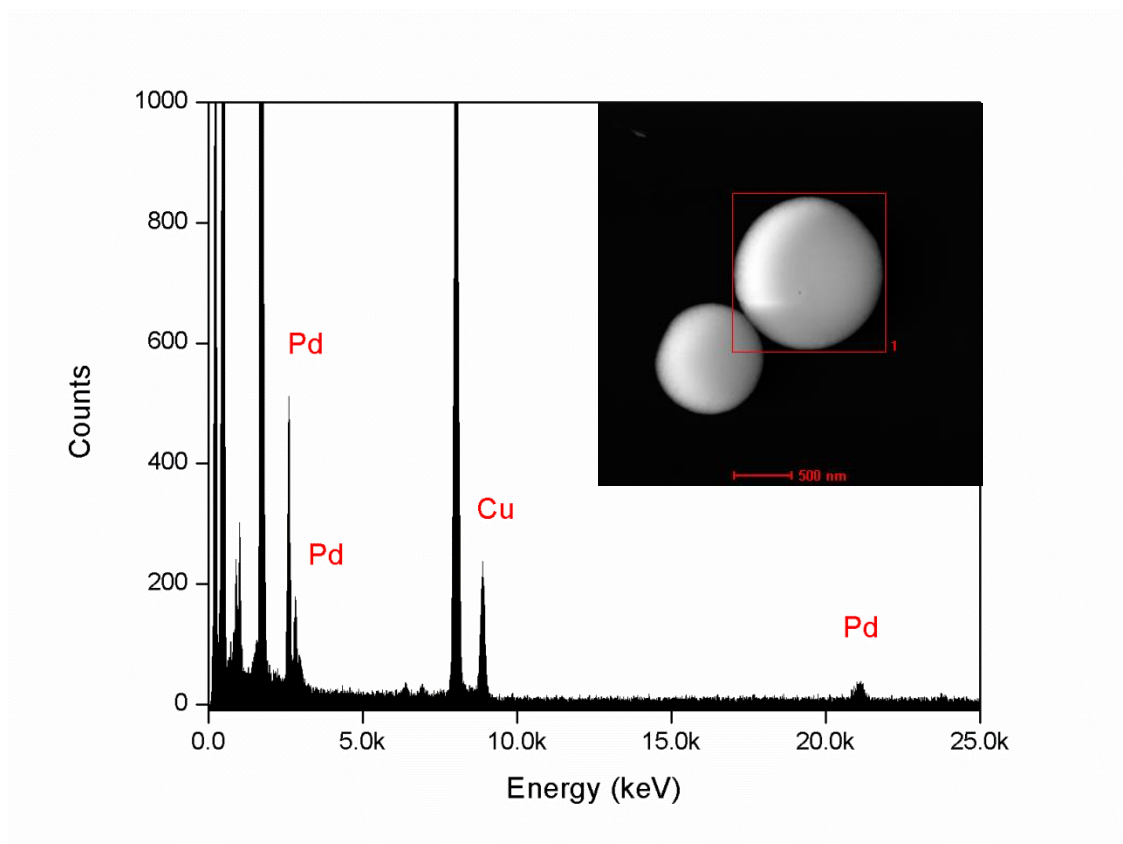


Figure S30. STEM image and EDS analysis of Na₂PdCl₄ entrapped inside CHCl:TA@SiO₂ microcapsules.

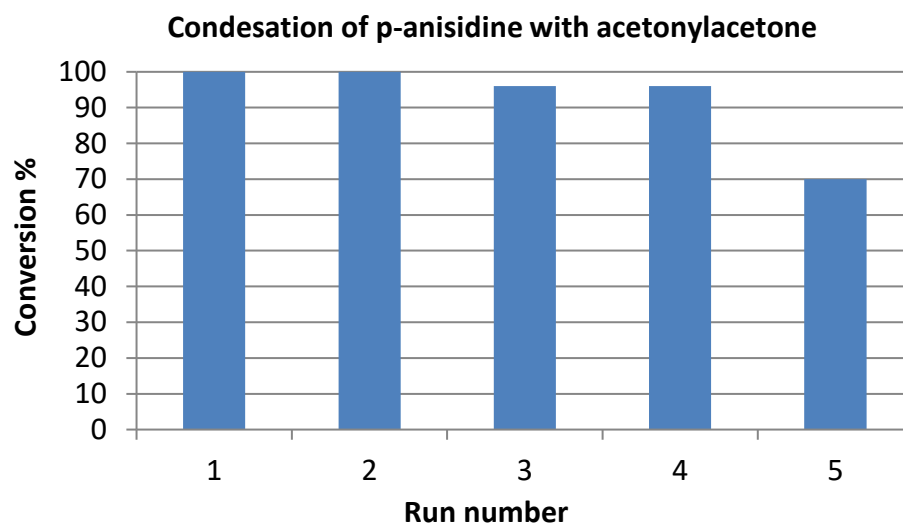


Figure S31. Recycling of CHCl:TA@SiO₂ microcapsules following the condensation of p-anisidine with acetonylacetone.

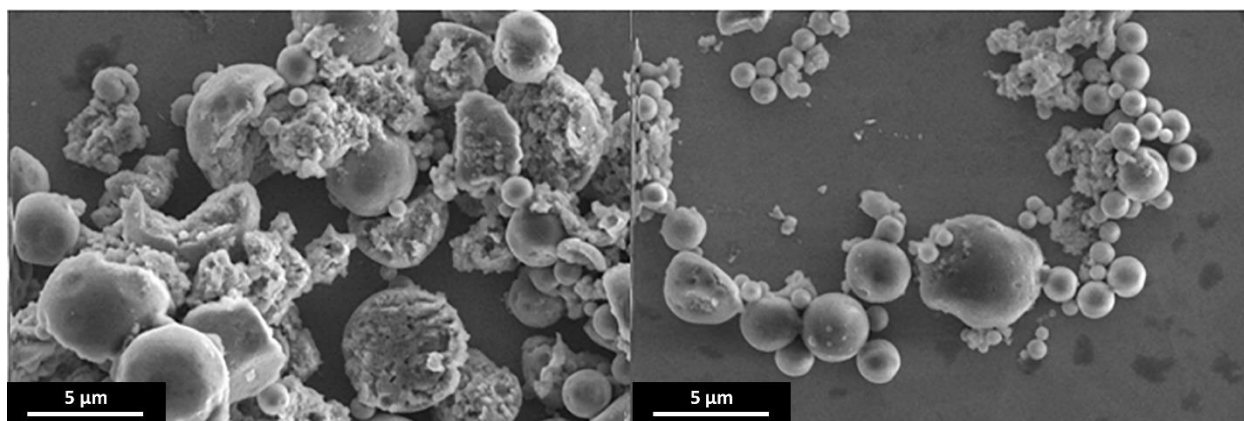


Figure S32. SEM images of CHCl:TA@SiO₂ microcapsules following the condensation of p-anisidine with acetonylacetone after the 4th run.