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Catalytic dissolution of metals from printed circuit board using a calcium chloridebased deep eutectic solvent

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Supplementary information



Figure S1: Gold-coated printed circuit board with block terminals of circular (in red circles) and square (in green squares)-shaped geometries.



Figure S2: Cyclic voltammograms of 0.01 mol dm⁻³ solutions of $K_3Fe(CN)_6$ in CaCl₂·6H₂O: EG (1:1 ratio). Scans were recorded at a 1 mm diameter Pt disc working electrode vs. Ag/AgCl in 3 M KCl reference at room temperature.



Figure S3: Cyclic voltammograms of the solvents obtained after copper powder dissolution (5 mg) in the presence of 0.1 mol dm⁻³ of (a) FeCl₃ and (b) CuCl₂ as oxidising agents in CaCl₂· $6H_2O$: EG. The legend describes the CaCl₂· $6H_2O$: EG (DES)-to-water ratio. The experiments were performed in 10 mL of CaCl₂· $6H_2O$: EG.



Figure S4: UV-vis spectra of the solvents obtained after copper powder dissolution (5 mg) in the presence of 0.1 mol dm⁻³ of (a) FeCl₃ and (b) CuCl₂ as oxidising agents in CaCl₂·6H₂O: EG at 50 °C. The legend describes the CaCl₂·6H₂O: EG (DES)-to-water ratio. The experiments were performed in 10 mL of CaCl₂·6H₂O: EG.



Figure S5: Etch depth of copper over time from the cross section of a square-shaped block terminal with two different concentrations of (a) FeCl₃ and (b) CuCl₂ as oxidising agents in the CaCl₂·6H₂O: EG eutectic system at 25 °C. The corresponding trend lines are also shown.



Figure S6: 3D reflected light image of the cross section of a printed circuit board block terminal with (a) 0.02 mol dm⁻³ CuCl₂ and (b) 0.01 mol dm⁻³ FeCl₃ in CaCl₂· $6H_2O$:EG (1:1 ratio), for 1 hour at 50 °C. The reddishbrown area describes the copper layer and, the white-greyish area describes the nickel layer. The dark area corresponds to the resin holding the printed circuit board sample.