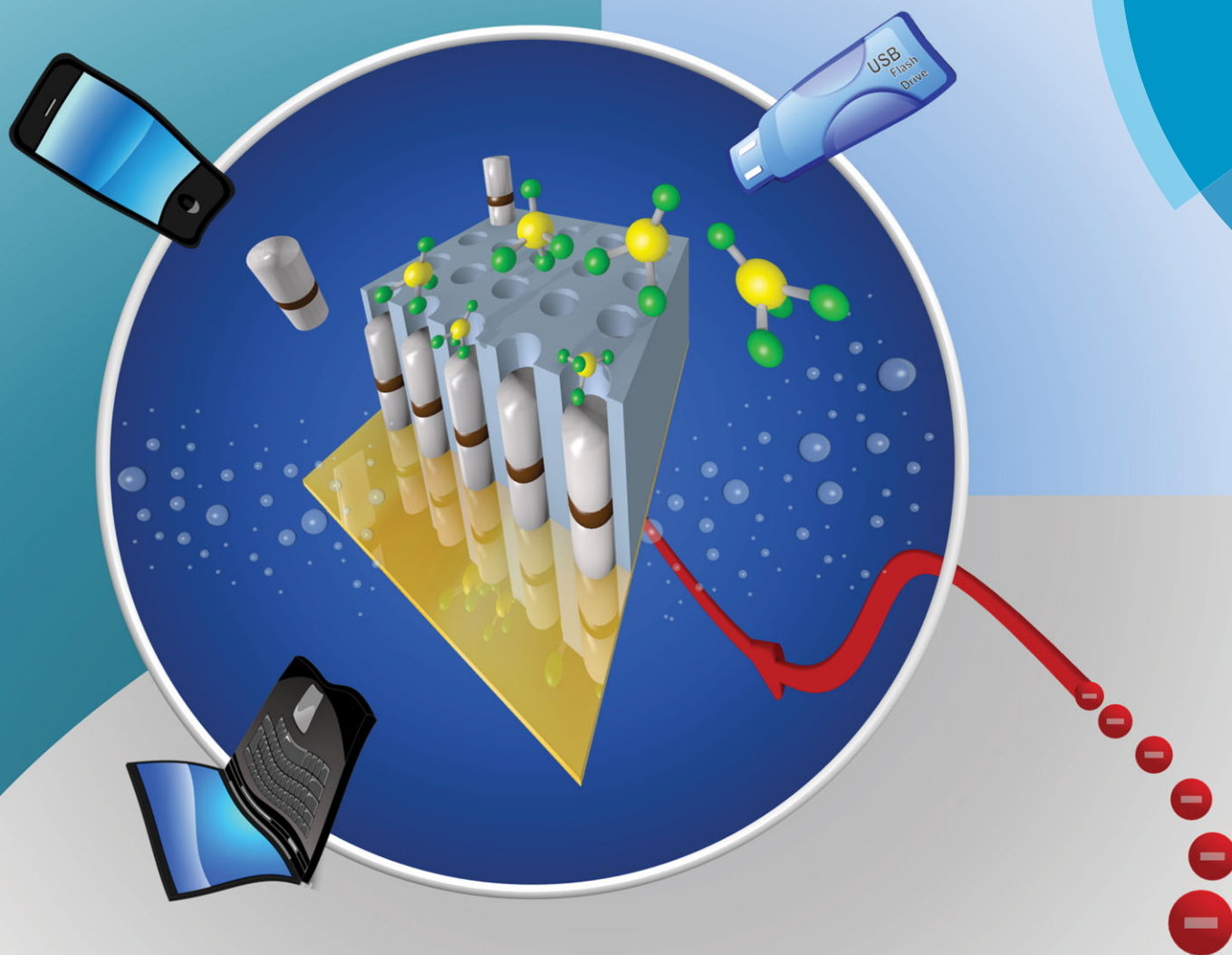


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PERSPECTIVE
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Recent studies have shown that it is possible to electrodeposit a range of materials, such as Cu, Ag and Ge, from various supercritical fluids, including hydrofluorocarbons and mixtures of CO₂ with suitable co-solvents. In this perspective we discuss the relatively new field of electrodeposition from supercritical fluids. The perspective focuses on some of the underlying physical chemistry and covers both practical and scientific aspects of electrodeposition from supercritical fluids. We also discuss possible applications for supercritical fluid electrodeposition and suggest some key developments that are required to take the field to the next stage.

1. Introduction

The field of supercritical fluid electrodeposition continues a long history of development of materials production and deposition methods which underpin nearly all modern technology. The importance and sophistication of materials deposition techniques can be seen in the cost of silicon fabrication facilities used to produce integrated circuits which are measured in billions of dollars. There now exists a wide range of deposition technologies. Vacuum techniques such as thermal evaporation, sputtering, chemical vapour deposition and molecular beam epitaxy are particularly suited to the deposition of high quality thin films. Such techniques have enabled the engineering of the fundamental quantum mechanical states of electrons in semiconductor structures leading to quantum well lasers in DVD players, quantum cascade lasers for chemical sensing, and entirely new fundamental physics, *e.g.* the fractional quantum Hall effect. Spin-coating and printing are two deposition techniques which have been used for a very long time, but which have been exploited in new ways recently. Much of this renewed interest has come about because of the discovery and application of organic semiconductors for applications such as flexible electronics. Whilst materials deposition is ubiquitous in technology, specific techniques can be found in particular applications in which their unique characteristics can out perform other techniques. For instance, as supercritical fluids have no surface tension and improved mass transport over liquids they have advantages for thermal deposition into high aspect ratio trenches and pores.¹ Supercritical fluids are also particularly good at swelling polymers in a controllable manner to allow the deposition of nanoparticles into existing polymers structures.²

In this perspective we will focus on electrodeposition from supercritical fluids and in particular on the underlying physical chemistry of this new field. Since its introduction in the early 1800s, electrodeposition has become widely exploited because it has a combination of capabilities not found together in other deposition techniques.^{3,4} These include: conformal deposition onto and into large complex objects; spatially defined growth allowing deposition onto complex structures only where required and allowing contiguous axial deposition through pores without blocking; direct control over the driving force for deposition through the applied potential; *in operando* measurement of electrical properties of the deposit during growth allowing active feedback; highly efficient use of reagents.

This perspective focuses on a relatively new development in the field of electrodeposition, supercritical fluid electrodeposition (SCFED). Supercritical fluids have a number of key advantages for electrodeposition. These include (i) a lack of surface tension which enables electrodeposition onto fragile substrates, (ii) good pore penetration and mass transport allowing deposition into high aspect ratio nanopores¹ and small diameter (< 4 nm) nanopores,⁵ (iii) a range of relatively easily accessible supercritical fluids with high chemical stability and thus large deposition windows, which allow the deposition of reactive materials such as germanium,⁶ and (iv) the possibility to increase the deposition temperature with a specific fluid to above the critical temperature. Whilst the technique is still in its infancy, it is possible to envisage technological areas in which SCFED may find unique applications. A likely example is in the production of few nanometre diameter nanowires and nanowire based devices, *e.g.* Fig. 1.

There are many, varied reasons for the interest in nanowires for applications. These include at the simplest level the possibility of further miniaturisation of existing electronic⁷ and data storage⁸ components. The proximity of the surface to the whole of the bulk of the nanowire makes nanowire chemical and biochemical sensors capable of single molecule sensitivity.⁹ In the field of thermoelectric

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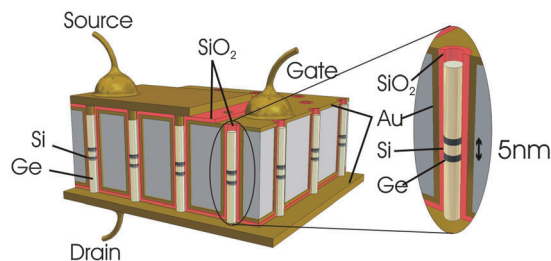


Fig. 1 Concept of a device which supercritical fluid electrodeposition may enable in the future. The concept consists of a multiple resonant tunnel diode memory with element separation of 5 nm. Electrodeposition is particularly suited to the production of axial heterostructures like the Si/Ge quantum dot structure shown here. Using a supercritical fluid as the solvent would have advantages for pore penetration, deposition of reactive materials and high temperature deposition for materials quality.

materials nanowires show increased figure of merit over bulk materials because of decreased phononic thermal conductivity.¹⁰ Other routes by which nanowires could contribute to energy efficiency and generation include photovoltaic nanowire devices¹¹ and piezoelectric nanowires.¹² The fact that nanowires are less susceptible to defects means that it may be possible to produce ultra-strength materials from them.¹³ Most of the near to market applications use nanowires with diameters in excess of 10 nm, however initial investigations of ultrathin nanowires¹⁴ suggest that these may be even more exciting in the medium term.

Supercritical fluids may bring a number of advantages for electrodeposition and for the electrodeposition of nanowires and nanostructures. However, there are significant challenges associated with trying to utilise SCFs for electrochemistry and electrodeposition. These include dissolution of a sufficient concentration of ionic species to produce a suitably conductive fluid, identifying sufficiently soluble and stable electrochemical precursors to produce useful deposition rates, ensuring the formation of a homogeneous solution and the development of safe and practical high pressure apparatus. Below we discuss these issues.

2. Properties of SCFs and measurement of solubility and conductivity

A supercritical fluid is defined as a substance above its critical temperature (T_c) and critical pressure (p_c), and as such they are unique solvents that combine the properties of liquids and gases, Table 1. SCFs have recently been exploited as green

alternatives to conventional organic solvents in a variety of applications including: (i) separation processes such as extraction, sorption processes, chromatography, and drying; (ii) mechanical processes such as extrusion, homogenisation, emulsification, micronisation, crystallisation, impregnation and encapsulation, and (iii) chemical and biochemical reactions.^{15–21} Many processes utilising sub- and supercritical fluids have been investigated for a wide range of industries including agriculture and food, cosmetics, pharmaceuticals, medicine, coatings, textiles, electronics and semiconductors, and waste treatment.^{22–27} The properties of SCFs have received considerable attention, and have been extensively reviewed.^{28–33}

Unlike conventional solvents, the physical properties of SCFs can be easily tuned by changing either temperature or pressure, or both. This advantage has been widely exploited in supercritical fluid extraction for separating products and in supercritical fluid reactions for achieving high yield and selectivity.^{28–33} The tuneable properties of SCFs also have applications in processing nano-materials, *e.g.* supercritical drying of highly porous β -chitin structures,³⁵ and preparation of free-standing arrays of CdS nanowires.³⁶

In order to successfully electrodeposit from a supercritical fluid we must consider their phase behaviour. SCFs are often illustrated with a p - T phase diagram of a single-component system. One such diagram is presented in Fig. 2 using trifluoromethane (CHF_3) as an example, for which T_c and p_c are 299.1 K and 4.82 MPa, respectively. The supercritical state is represented by the shaded area in Fig. 2. Density is one of the most important physical properties of SCFs; density can be shown as isochores in the p - T phase diagram, *e.g.* 7 isochores of CHF_3 are depicted in Fig. 2, with the density between 0.25 and 1.45 g cm⁻³. It is worth pointing out that only a certain combination of temperature and pressure within the shaded area can provide the desired properties for use of SCFs as solvents. For example, consider CHF_3 under the conditions of 400 K and 15 MPa, which is ~ 100 K above its T_c , and ~ 10 MPa above its p_c . At 400 K and 15 MPa CHF_3 is in the supercritical state; however its density is only 6.4 mol dm⁻³ and compared with liquid H_2O , having a density of 55.6 mol dm⁻³ at 298 K, or liquid CHF_3 , having a density of 20 mol dm⁻³ at its boiling point, the density of CHF_3 at 400 K and 15 MPa is too low for it to be a good solvent for ionic species, and therefore too low for it to be a good solvent for electrodeposition.

The dissolution of ionic species in SCFs is one of the key challenges to carry out supercritical fluid electrodeposition because most of the SCFs with easily accessible critical temperatures are non-polar or low polarity molecules and therefore are

Table 1 Density, viscosity, and diffusion coefficient of SCFs, gases and liquids^{28,34}

State	Gas	Supercritical fluid	Liquid
Temperature/K	298	T_c	298
Pressure/MPa	0.1	p_c	0.1
Density/g cm ⁻³	$0.6\text{--}2 \times 10^{-3}$	0.2–0.5	0.6–2
Diffusion coefficient/cm ² s ⁻¹	0.1–0.4	$0.5\text{--}4 \times 10^{-3}$	$0.2\text{--}2 \times 10^{-5}$
Viscosity/MPa s	0.01–0.03	0.01–0.03	0.2–20



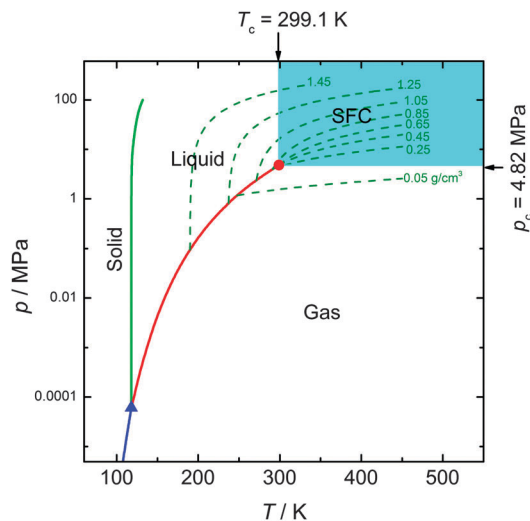


Fig. 2 Phase diagram of pure CHF_3 . ●, the critical point of CHF_3 ; ▲, the triple point of CHF_3 ; SCF, supercritical fluid, shaded in cyan. The red, green and blue curves are the gas–liquid, liquid–solid, and gas–solid phase boundary, respectively. The dashed curves represent the isochores corresponding to the density between 0.05 and 1.45 g cm^{-3} . The critical temperature (T_c) and critical pressure (p_c) are 299.1 K and 4.82 MPa, respectively, indicated by the arrows in the figure.

not good media for ion solvation. Electrostatic interactions contribute more than 80% of the total ion–solvent interactions.³⁷ Based on the Born equation, the electrostatic solvation energy (ΔG_{ES}), defined as the difference between the electrostatic free energy of an ion *in vacuo* and that in a solution, can be calculated from the ionic charge, ionic radius, and dielectric constant (ϵ_r). Fig. 3a shows the calculated ΔG_{ES} for a univalent ion with a radius of 0.2 nm in a medium with a dielectric constant ranging from 1 to 40. It can be seen that ΔG_{ES} decreases very rapidly with the increase of ϵ_r in the low dielectric constant region ($\epsilon_r < 7$), but decreases rather slowly in the high dielectric constant region ($\epsilon_r > 20$). In order for the electrolyte to dissolve and dissociate in the solution ϵ_r needs to be large enough so that the Gibbs free energy for solvation, ΔG_{ES} , is sufficiently negative to overcome the lattice energy. On the other hand, for those solvents with dielectric constant greater than 20, the difference in ΔG_{ES} is rather small so that the effect of dielectric constant on the solubility is not significant when only the electrostatic solvation energy is taken into account. The ϵ_r region between 7 and 20 is highlighted in Fig. 3. This is why most non-aqueous electrochemistry is carried out in solvents with a dielectric constant above 10 such as acetonitrile ($\epsilon_r = 36.6$), *N,N'*-dimethylformamide ($\epsilon_r = 38.3$), dimethylsulfoxide ($\epsilon_r = 47.2$) or propylene carbonate ($\epsilon_r = 66.1$).

Fig. 3b plots the dielectric constant of three fluorinated hydrocarbons (HFCs) as a function of pressure at temperatures above their corresponding T_c . It is clear that the dielectric constant increases with increasing pressure for all three HFCs, suggesting that the solubility of ionic species is higher at high pressures. In addition, the solubility of ionic species can be substantially higher in CH_2F_2 than in the other two HFCs (*i.e.* CHF_3 and $\text{CF}_3\text{CH}_2\text{F}$) because the dielectric constant of

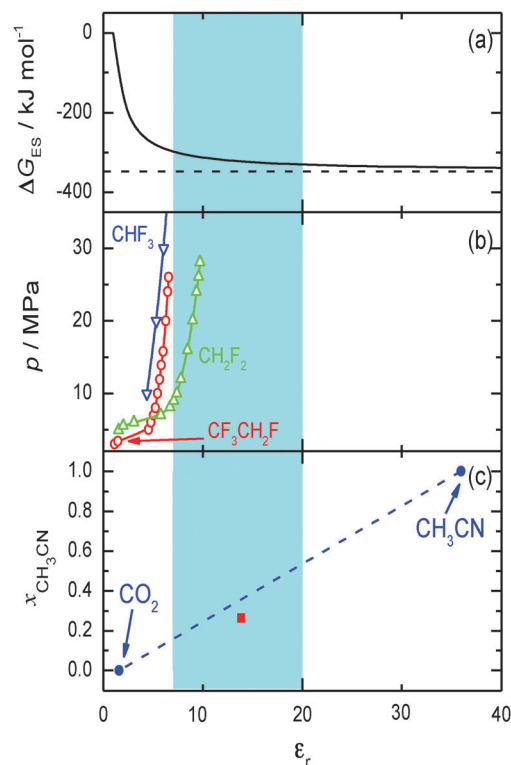


Fig. 3 (a) Plot of the electrostatic solvation energy (ΔG_{ES}) of a univalent ion against dielectric constant (ϵ_r), calculated by the Born equation with the ionic radius of 0.2 nm.³⁷ (b) The effect of pressure (p) on ϵ_r for three fluorinated hydrocarbons at $T > T_c$ of each solvent: ▽, CHF_3 at 335 K; ○, $\text{CF}_3\text{CH}_2\text{F}$ at 383 K; △, CH_2F_2 at 363 K.^{38–40} (c) ϵ_r of the binary system of $\text{CO}_2 + \text{CH}_3\text{CN}$. The dashed line represents the linear interpolation of ϵ_r as a function of the mole fraction of CH_3CN ($x_{\text{CH}_3\text{CN}}$).^{41,42} ■, ϵ_r of a mixture of $\text{CO}_2 + \text{CH}_3\text{CN}$ measured at 25 °C and 5.0 MPa.⁴³ The composition of the mixture is interpolated from the vapour–liquid phase equilibrium data reported by Reighard *et al.*⁴⁴

CH_2F_2 reaches 7 when the pressure is over 8 MPa. Hence, the selection of SCFs with high dielectric constant is the key to increasing the solubility of ionic species.

An alternative method to increase the dielectric constant is the use of mixed solvents, *e.g.* adding polar co-solvents into non-polar SCFs. This has been a widely accepted concept in all sorts of applications of SCFs.^{45–49} In particular, the enhancement of the solubility of polar substances in CO_2 has been extensively studied by using acetone, methanol, ethanol, acetonitrile, *etc.* as co-solvents. Fig. 3c shows the effect of the mole fraction of CH_3CN ($\epsilon_r = 36$) on the dielectric constant of the binary system of $\text{CO}_2 + \text{CH}_3\text{CN}$. One can see that it is crucial to add 10–18% (mole fraction) of CH_3CN into CO_2 to achieve a dielectric constant of 7. A much higher concentration of CH_3CN (*e.g.* $x_{\text{CH}_3\text{CN}} > 0.5$) may not be necessary because it will not significantly decrease ΔG_{ES} , but makes the solution more ‘liquid-like’, and hence slows down the mass transport.

High-pressure electrochemistry cells are the key equipment for electrodeposition in SCFs. The general design considerations include maximum operating temperature and pressure, electrode insulation and installation, material comparability,



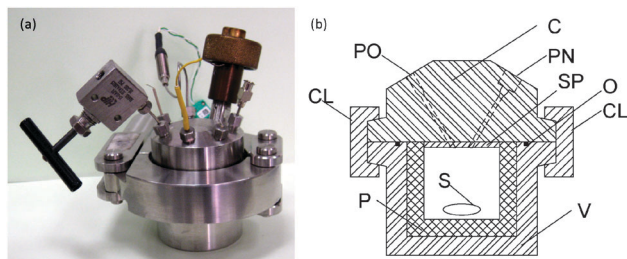


Fig. 4 High-pressure electrochemistry cell: (a) the electrochemistry cell with the electrodes installed; (b) schematic diagram, C, cap; CL, clamp; O, O-ring; P, PEEK pot; PO, SSI type port; PN, port for the needle valve; S, magnetic stirrer; SP, PTFE spacer; V, stainless steel vessel. More details can be found from ref. 6 and 34.

and high-pressure safety. (**Safety note:** the illustrations shown here are purely schematic and should not be regarded as engineering drawings for the construction of safe high-pressure cells. Anyone wishing to build such equipment for themselves must carry out their own safety evaluation.) There are also a number of more subjective requirements, such as stirring the fluid in the cell, dismantling and reassembling the cell frequently and quickly, and convenience of transferring reagents. One cell design is given in Fig. 4. Briefly, the cell consists of a 10 cm³ cylindrical vessel and a cap at its top, constructed from stainless steel. The cap has a fitted needle valve and 6 ports for SSI (Scientific Systems Inc.) type electrodes, thermocouples and fluid inlet/outlet. The vessel and the cap are assembled using a failsafe clamp system, incorporating a disposable elastomeric O-ring.⁵⁰ Recently, a pot made from PEEK (polyether ether ketone) was added to the vessel, together with a PTFE spacer fixed on the cap to reduce contamination from the stainless-steel inner surface.⁶

Electrodeposition requires working solutions with low ohmic resistance and high concentrations of supporting electrolytes and reagents. Since such data on working solutions are not readily available in the literature for supercritical fluid systems, it is essential to acquire a fundamental understanding on the electrical conductivity and phase behaviour of working solutions at high temperature and high pressure. For this purpose, specialised high pressure equipment has been utilised for studying the conductivity and phase behaviour of multi-component systems; two examples of such equipment are shown in Fig. 5. The detailed description of the equipment can be found from the respective publications.^{34,51,52}

3. Electrochemistry in supercritical fluids

It is appropriate to begin this section with a quotation from Darr and Poliakoff which is useful to keep in the forefront of one's mind "Working with supercritical fluids involves high pressures and sometimes high temperatures. Nearly always, it will always be easier to carry out an experiment under conventional conditions than supercritically".⁵³ This, in our experience, is definitely true in the case of electrochemistry where, when working at elevated temperatures and pressures it is no longer a trivial matter to polish the electrode between measurements,

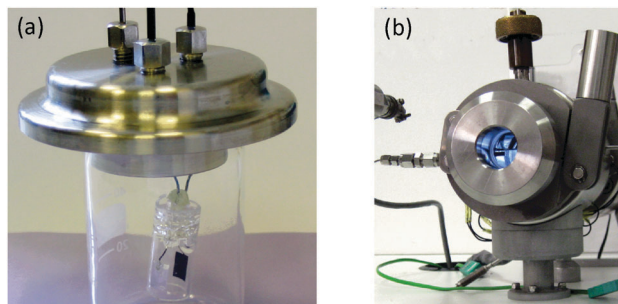


Fig. 5 (a) Conductivity cell with two platinum electrodes. The electrode is soldered to a platinum wire, which is insulated with PEEK tubing and sealed into a stainless steel holder for installing in a high-pressure autoclave. (b) Variable-volume view cell for studying the phase behaviour of multi-component systems. The fluid mixture with a given concentration of supporting electrolyte and reagents is introduced into the cell and sealed. The conditions (temperature and pressure) corresponding to a homogeneous, single-phase state can be determined visually by changing the volume of the cell at a given temperature.

to find well-behaved reference electrodes or to take for granted the function of the counter electrode. On the other hand supercritical fluids have unique properties and sometimes the additional complexity allows results which cannot be achieved another way.

The earliest studies in supercritical fluids other than water appear to have been by Williams and Naiditch who reported the electrodeposition of Ag from "dense gaseous solutions" of AgNO₃ in ammonia.⁵⁴ These experiments were carried out in sealed glass vessels using two Pt electrodes. Dendritic growth of Ag was observed on the cathode. The deposition was carried out slightly (7 K) above the critical temperature for NH₃ so it is not certain that the fluid was in the supercritical state.

Subsequently in 1981 Silvestri *et al.* reported results for a preliminary study of supercritical CO₂, bromotrifluoromethane, hydrogen chloride and ammonia as solvents for electrochemistry.⁵⁵ Their focus was on the possibility of using these supercritical fluids as solvent for electrosynthesis. They report that "a solution of tetrabutylammonium iodide in CO₂ was a poor conductor in the liquid, as well as in the supercritical state. Bromotrifluoromethane, in which the electrolyte was practically insoluble, also proved to be a very poor conductor". They had greater success with the more polar, and therefore higher dielectric, solvents scHCl and scNH₃ and they report long-term electrolyses (4000 C charge passed) for Ag and Fe electrodes in scNH₃. scHCl was found to be difficult to work with because it is highly corrosive. Nevertheless they were able to produce elemental iodine by electrolysis of KI in scHCl.

Between 1984 and 1997 Bard's group published a series of studies of electrochemistry in supercritical fluids, NH₃,^{56–58} water,^{59–62} acetonitrile^{63,64} and SO₂.⁶⁵ In these studies they developed the techniques for working with supercritical fluids, and particularly corrosive supercritical fluids such as water, and studied the electrochemistry of both inorganic and organic redox systems, including Cu deposition,⁶² halide oxidation,⁶⁰ oxygen reduction,⁶⁰ and hydroquinone oxidation⁶⁰ all from scH₂O, and solvated electrons and organic redox couples in scNH₃.^{56–58}

The systems investigated by Bard's group were all quite polar fluids. These, in general, have the disadvantage of high



critical temperatures and pressures. Although less polar supercritical fluids such as scCO_2 are more benign to work with, they present the electrochemist with a significant problem in that they have very low dielectric constants and it is therefore difficult to achieve sufficient dissolution and dissociation of the electrolyte to achieve reasonable solution conductivity. Thus, for example, Abbott and Harper⁶⁶ briefly studied the electrochemistry of $[\text{Ndodecyl}_4][\text{Ni}(\text{mnt})_2]$ (mnt = maleonitrile) in scCO_2 using hydrophobic electrolytes (tetradecylammonium tetraphenylborate), they found some conductivity ($\sim 10^{-6} \text{ S cm}^{-1}$) and obtained some, poorly resolved, voltammetry.

One approach to overcome this problem is to use co-solvents such as methanol or acetonitrile. These mixtures still form single phase supercritical systems, although it is of course essential to characterise the phase behaviour to establish the appropriate conditions in terms of T and p , but the presence of the co-solvents increases the solubility of ionic species and dissociation of the electrolyte, and hence the conductivity of the fluid. To increase the conductivity of scCO_2 ($\epsilon_r = 1.5$ at 308 K and 10 MPa⁴¹), CH_3OH and CH_3CN have been previously explored as co-solvents.⁶⁷ A review of electrochemistry in scCO_2 and scCO_2 with various co-solvent, covering the literature up to 1998, has been provided by Grinberg and Mazin.⁶⁸

An alternative approach is to use hydrofluorocarbons (HFCs) as supercritical solvents as these are polar and give higher dielectric constant fluids whilst retaining reasonable critical temperatures and pressures. Early work in this area was carried out by Olsen and Tallman⁶⁹ who studied the electrochemistry of ferrocene and the cobalticenium cation at microdisc electrodes in scCHClF_2 with millimolar concentrations of $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ electrolyte. In addition they also investigated the use of scCHF_3 and found good voltammetry for ferrocene and the cobalticenium cation with microdisc electrodes (Fig. 6). Subsequently, in a series of papers, Goldfarb and Corti^{70–72} studied electrochemistry of decamethylferrocene in scCHF_3 containing $[\text{N}^n\text{Bu}_4][\text{PF}_6]$ electrolyte.

At around the same time Abbott's group also published a series of papers on electrochemistry in supercritical hydrofluorocarbons. Thus they investigated the use of supercritical 1,1,1,2-tetrafluoroethane ($\text{CF}_3\text{CH}_2\text{F}$) and scCH_2F_2 and demonstrated a wide potential window for these solvents with $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ or $[\text{N}^n\text{Bu}_4][\text{ClO}_4]$ electrolytes allowing oxidation of $[\text{Cs}(18\text{-crown-6})]^+$ and Xe in liquid $\text{CF}_3\text{CH}_2\text{F}$.⁷³ They also used supercritical 1,1,1,2-tetrafluoroethane with $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ electrolyte as a solvent for the electrochemical reduction of CO_2 at Pt and Pb electrodes.⁷⁴

In addition to these studies of electrode reactions there have been a limited number of studies of more fundamental issues such as diffusion, effects of solvation and the structure of the double layer. For example Goldfarb and Corti⁷² have published an in depth study of the diffusion coefficient for decamethylferrocene in scCHF_3 with $[\text{N}^n\text{Bu}_4][\text{PF}_6]$ electrolyte as a function of the temperature and pressure, taking account of the effects of ion pairing and correlating the results on the basis of the Stokes–Einstein model. Abbott *et al.*⁷⁵ investigated the effects of solute on the viscosity of scCH_2F_2 using quartz crystal microbalance measurements and showed that the viscosity could be surprisingly high (8–32 fold increase) at pressures

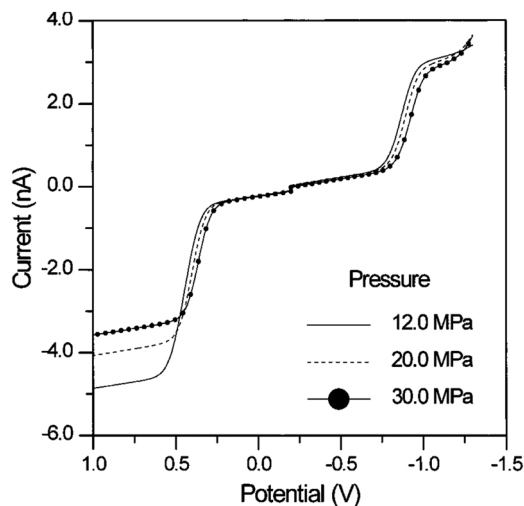


Fig. 6 Comparison of voltammetry of ferrocene and the cobalticenium cation as a function of pressure in supercritical chlorodifluoromethane. Conditions: 90 μM Fc, 90 μM Cc^+ , 10.0 mM $[\text{N}^n\text{Bu}_4][\text{BF}_4]$; 388 K; 25 μm -diameter Pt disk electrode; scan rate, 20 mV s^{-1} . For clarity, only the forward sweep of each voltammogram is displayed, with the ferrocene wave recorded first and the cobalticenium cation wave immediately after, each wave originating from -0.20 V . Reprinted with permission from S. A. Olsen and D. E. Tallman, *Anal. Chem.*, 1996, **68**, 2054–2061. Copyright 1996 American Chemical Society.

close to the critical value. They also investigated the effects of ion-pairing on the redox potential of ferrocene carboxylic acid in scCH_2F_2 .⁷⁶ Abbott's group are also the only ones to investigate the structure of the double layer in supercritical fluid electrolytes; scCH_2F_2 containing $[\text{NR}_4][\text{BF}_4]$ ⁷⁷ and scCO_2 containing long chain quaternary ammonium electrolytes.⁷⁸

4. New electrolytes

One of the enduring challenges of electrochemistry in supercritical fluids is the choice of electrolyte in order to achieve high solution conductivity and thus reduce iR drop due to the uncompensated solution resistance. This is particularly important for applications in electrosynthesis or electrodeposition where it is necessary to pass larger currents. A number of studies have been reported in the literature on the conductivity of supercritical fluid electrolytes.

Olsen and Tallman⁷⁹ reported values for the equivalent conductivity for $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ in scCHClF_2 at 388 K between 10 and 24 MPa for concentrations between 6 and 13 mM, Fig. 7. They found a non-linear increase in equivalent conductivity with the square root of electrolyte concentration; behaviour indicative of the formation of triple ions $(+)(-)(+)$ or $(-)(+)(-)$ as predicted by the Fuoss–Kraus equation.⁸⁰ Abbott and Eardley⁸¹ investigated the conductivity of $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ in scCH_2F_2 and found similar behaviour with evidence for a contribution from triple ions and Goldfarb and Corti,⁷¹ in an extensive study, found similar effects for $[\text{N}^n\text{Bu}_4][\text{ClO}_4]$ in scCHF_3 . Jun and Fedkiw reported conductivities in the range 10^{-5} to $10^{-4} \text{ S cm}^{-1}$ for alkali-metal salts of trifluoroacetates in $\text{scCO}_2\text{--CH}_3\text{OH}$, with



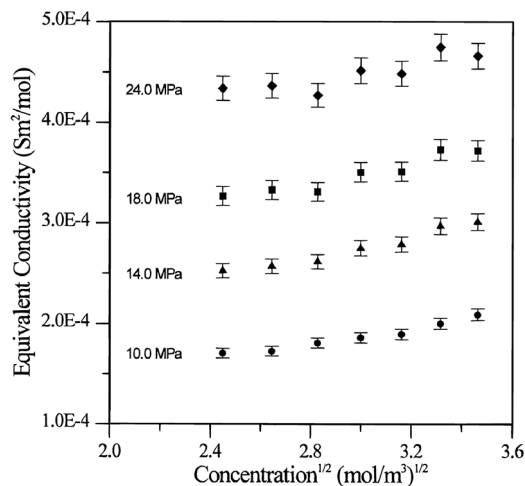


Fig. 7 Equivalent conductivity of supercritical CHClF_2 at 388 K as a function of the square root of $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ concentration. The points (error bars) are the averages (standard deviations) of 5–6 measurements. Data at four densities are shown corresponding to fluid pressures of 10.0 (●), 14.0 (▲), 18.0 (■), and 24.0 MPa (◆). Reprinted with permission from S. A. Olsen and D. E. Tallman, *Anal. Chem.*, 1996, **68**, 2054–2061. Copyright 1996 American Chemical Society.

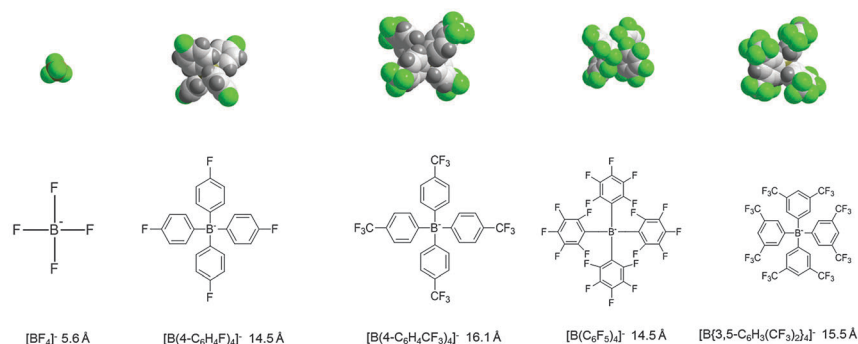
the highest values in the single phase region obtained for the lithium salt.⁸² In all cases these commercially available electrolytes give solutions with reasonable conductivity but, for electro-synthesis and electrodeposition applications, it would be desirable to find electrolytes with higher conductivity.

The successful development of electrodeposition from supercritical fluids requires careful choice of both electrolytes and reagents. The electrolytes must (i) have good solubility in the SCF, (ii) be dissociated into the constituent ions in the

solution, (iii) have a wide electrochemical window (*i.e.* redox inactive in the potential range to be used), and (iv) be chemically inert with respect to the reagents added. The low polarity of the SCFs used in this work makes (i) and (ii) in particular, key issues to be addressed. Weakly coordinating anions have long been used in coordination and organometallic chemistry with aims including stabilisation of reactive cations, allowing the coordination of weakly bound ligands to metal centres, and enhancing catalysis rates and activities. Typical examples include $[\text{BF}_4]^-$, $[\text{ClO}_4]^-$, $[\text{PF}_6]^-$, $[\text{SO}_3\text{F}]^-$, $[\text{BPh}_4]^-$ usually in association with alkali metal or large tetra-alkylammonium cations.⁸³ The search for even more weakly coordinating anions has resulted in development of the chemistries of fluorinated analogues of $[\text{BPh}_4]^-$, $[\text{M}\{\text{OTeF}_5\}_m]^-$ ($\text{M} = \text{B}, \text{As etc.}$), $[\text{Al}\{\text{OC}(\text{CF}_3)_3\}_4]^-$, and various carboranes.⁸⁴ It is important to remember that weakly coordinating anions are not necessarily chemically inert, many examples of fluoride abstraction from $[\text{BF}_4]^-$ or $[\text{PF}_6]^-$, and B–C bond cleavage in $[\text{BPh}_4]^-$ are known.⁸³ For SCFED, electrolytes based upon tetraalkylammonium cations and weakly coordinating anions seem to be a promising approach. Conductivity studies of scCO_2 , CH_2F_2 and CHF_2CHF_2 by Abbott *et al.*^{73,81} used $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ or $[\text{N}^t\text{Bu}_4][\text{ClO}_4]$ as electrolytes. We have used fluorinated tetra-aryl borates ($[\text{B}(\text{Ar}^F)_4]^-$) which were initially developed by organometallic chemists to address both the tendency of $[\text{BPh}_4]^-$ to π -aryl coordination to metal centres and B–C bond fission; they are also much more resistant to fluoride abstraction by electrophilic metals than $[\text{BF}_4]^-$.^{83,84}

The $[\text{B}(\text{Ar}^F)_4]^-$ anions with $\text{Ar}^F = \text{C}_6\text{H}_4\text{F}$, C_6F_5 , $4\text{-CF}_3\text{C}_6\text{H}_4$ and $3,5\text{-(CF}_3)_2\text{C}_6\text{H}_3$, (Fig. 8) are readily made as sodium salts by reaction of $\text{Na}[\text{BF}_4]$ with the appropriate Grignard reagent $\text{BrMg}(\text{Ar}^F)$,^{85–87} and can be converted to $[\text{NR}_4]^+$ salts by metathesis with $[\text{NR}_4]\text{I}$ in CH_2Cl_2 .⁸⁸ Various $[\text{B}(\text{Ar}^F)_4]^-$ salts are commercially

Fluorinated aryl borate anions



Tetraalkylammonium cations

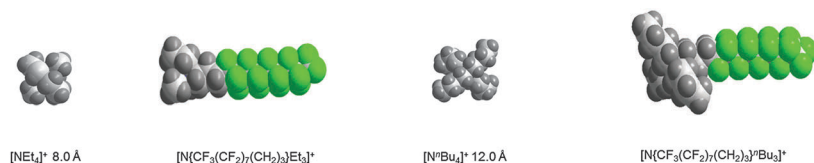


Fig. 8 Space filling representations of the cations and anions used as supporting electrolytes (green = F), illustrating their approximate sizes. The dimensions correspond to the estimated diameter across the respective ions based upon van der Waals radii.



available, although very expensive, but we have found that, depending on the metal reagents used, it is often possible to recover the electrolyte after deposition, re-purify it and hence recycle it.

We also explored the effects of using alkylammonium cations incorporating fluororous "ponytails" to increase solubility. $[\text{CF}_3(\text{CF}_2)_7(\text{CH}_2)_3\text{N}^+\text{Bu}_3]\text{I}$ was made by quaternisation of $\text{N}^+\text{Bu}_3\text{N}$ with $\text{CF}_3(\text{CF}_2)_7(\text{CH}_2)_3\text{I}$, and $[\text{CF}_3(\text{CF}_2)_7(\text{CH}_2)_3\text{NMe}_3]\text{I}$ from $\text{CF}_3(\text{CF}_2)_7(\text{CH}_2)_3\text{NH}_2$ and CH_3I , and metathesis with $\text{Na}[\text{B}(\text{Ar}^{\text{F}})_4]$ in CH_2Cl_2 gave the appropriate $[\text{CF}_3(\text{CF}_2)_7(\text{CH}_2)_3\text{NR}_3][\text{B}(\text{Ar}^{\text{F}})_4]$.⁸⁸ Thermogravimetric analysis studies showed that the cations were thermally stable up to $\sim 180^\circ\text{C}$, although subsequent conductivity measurements showed the cations were not significantly better than $[\text{N}^+\text{Bu}_4]^+$.⁸⁸

As stressed above, it is important to fully characterise the phase behaviour of the system since addition of significant concentrations ($>10\text{ mM}$) of electrolyte is likely to alter the phase behaviour. This can be done using a view cell of the type described above. These studies should then be complemented by measurements of the conductivity to characterise the behaviour of the different electrolytes. In our work⁸⁸ we started by comparing the behaviour for mixtures of CO_2 with $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ and either methanol (CH_3OH) or acetonitrile (CH_3CN) as the co-solvent. These experiments showed that at similar temperatures and pressures, the solubility of $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ was ~ 5 times higher in $\text{CH}_3\text{CN} + \text{CO}_2$ than in $\text{CH}_3\text{OH} + \text{CO}_2$.

Fig. 9 shows the phase behaviour of a ternary CO_2 – $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ – CH_3CN system and illustrates the potential complexity. For the mixture shown in Fig. 9, the three-phase region is a surface with 2 degrees of freedom as required by the phase rule for a ternary system with three phases ($F = 3 - 3 + 2 = 2$). At high T and p there is a single supercritical phase region which gives way to two and three phase regions as the pressure is decreased. The phase transition is a normal bubble-point transition ($\text{L}_1 \rightarrow \text{L}_1 + \text{V}$) when the temperature is below T^* . However, when T is above T^* , the sequence of the phase transition on decreasing the pressure isothermally is: $\text{L}_1 \rightarrow \text{L}_1 + \text{L}_2 \rightarrow \text{L}_1 + \text{L}_2 + \text{V} \rightarrow \text{L}_1 + \text{V}$.

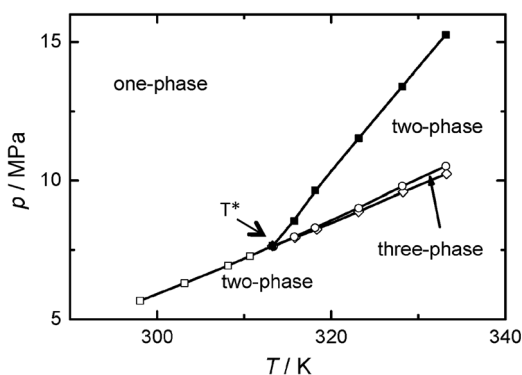


Fig. 9 p – T phase diagram of the mixture of CO_2 (1) + CH_3CN (2) + $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ (3) with $x_1 = 0.871$, $x_2 = 0.129$, and $x_3 = 8.3 \times 10^{-4}$. (■) and (□), the phase boundary between one-, and two-phase region; (○) and (◇) mark the region where three phases are in co-existence ($\text{L}_1\text{L}_2\text{V}$). The pressure difference between the two three-phase boundaries is less than 0.3 MPa at 333 K.³⁴

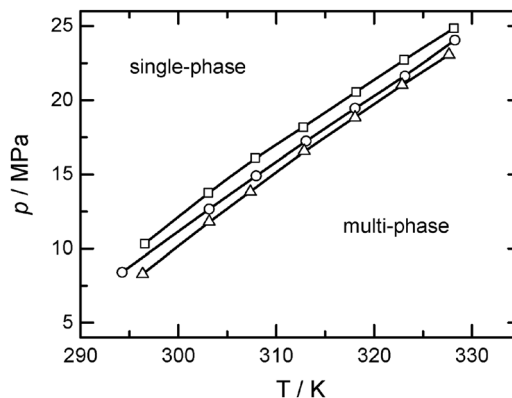


Fig. 10 p – T phase diagram of the mixture CO_2 (1) + CH_3CN (2) + $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ (3) with and without metal salts (4). $x_1 = 0.89$, $x_2 = 0.11$ and $x_3 = 1.34 \times 10^{-3}$. (○) no metal salt; (△) with $[\text{Ag}(\text{hfac})(\text{cod})]$, $x_4 = 3.4 \times 10^{-4}$, and (□) with $[\text{Cu}(\text{hfac})_2]$, $x_4 = 3.6 \times 10^{-4}$.³⁴

From these phase behaviour studies for different compositions of the ternary mixture we find that increasing the concentration of $[\text{N}^+\text{Bu}_4][\text{BF}_4]$ increases the pressure required to form a homogeneous solution at a given temperature, and furthermore that the more CO_2 there is in the system, the higher the pressure that is required because CO_2 is a poor solvent for the ionic species (in this case $[\text{N}^+\text{Bu}_4][\text{BF}_4]$). Again this illustrates the potential complexity of the phase behaviour of these systems and why it is essential to experimentally establish the conditions required to form a single supercritical phase.

In principle, the addition of the electrochemical reagents used in the electrodeposition will also alter the phase behaviour. In practice for this system, as shown in Fig. 10, the effects are much smaller, not least because the concentrations of the dissolved reagents are much smaller – the addition of $\sim 3.5 \times 10^{-4}$ mole fraction of $[\text{Ag}(\text{hfac})(\text{cod})]$ or $[\text{Cu}(\text{hfac})_2]$ (cod = cycloocta-1,5-diene, $\text{hfacH} = \text{CF}_3\text{C}(\text{O})\text{CH}_2\text{C}(\text{O})\text{CF}_3$) produces only slight shifts in the supercritical phase boundary ($<2\text{ MPa}$ at fixed temperature).

In order to improve upon the commercial electrolytes we have synthesised a range of anions and cations selected to achieve greater solubility in scCO_2 – CH_3CN by increasing the ion size and by increasing the degree of fluorination. In our conductivity studies we found that the effects of adding a fluororous ponytail (replacing $[\text{N}^+\text{Bu}_4]^+$ by $[\text{NR}^{\text{F}}\text{Me}_3]^+$ or $[\text{NR}^{\text{F}}\text{Bu}_4]^+$) to the cation were small and did not justify the synthetic effort. This is presumably because the fluororous ponytail does not alter the separation in the ion pair because it is flexible. A similar effect was found by Abbott and Schiffrin⁸⁹ for tetrafluoroborate salts of long chain alkylammonium cations in low dielectric solvents, where the interionic distance in the ion pairs is largely independent of the alkyl chain length due to interpenetration.

Table 2 summarises the data, the structures of the ions and their relative sizes are shown in Fig. 8. In terms of the effect of the anions on the molar conductivity, the general trend is $[\text{BF}_4]^- \sim [\text{B}(4\text{-C}_6\text{H}_4\text{F})_4]^- < [\text{B}(4\text{-C}_6\text{H}_4\text{CF}_3)_4]^- \sim [\text{B}(\text{C}_6\text{F}_5)_4]^- < [\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]^-$ with the highest molar conductivity, $22\text{--}26\text{ S cm}^2\text{ mol}^{-1}$, achieved using the $[\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]^-$.



Table 2 Molar conductivity of the fluorinated supporting electrolytes in a mixed fluid of CO₂ (1) + CH₃CN (2) with a molar ratio of $x_2/x_1 = \sim 0.12$ at 328.15 K and 20 MPa³⁴

	[N ⁿ Bu ₄] ⁺	[NR ^f Me ₃] ⁺	[NR ^f Bu ₃] ⁺
[BF ₄] [−]	2.3 ^{a,b} (2.4) ^c	—	—
[B(4-C ₆ H ₄ F) ₄] [−]	—	3.1 (2.2)	3.6 (2.4)
[B(4-C ₆ H ₄ CF ₃) ₄] [−]	—	8.9 (0.9)	10.6 (1.3)
[B(CF ₃) ₄] [−]	10.6 (2.4)	8.8 (1.0)	6.8 (0.7)
[B{3,5-C ₆ H ₃ (CF ₃) ₂ }] [−]	22.7 (2.4)	21.5 (2.4)	26.2 (2.5)

^a The supporting electrolyte represented here consists of a cation and an anion which can be found from the column name and row name respectively. ^b Molar conductivity in units of S cm² mol^{−1}. ^c The numbers in brackets represent the concentration of electrolytes in mmol dm^{−3}.

This is about an order of magnitude improvement over [NⁿBu₄][BF₄]. For a solution of 0.07 M [NⁿBu₄][B{3,5-C₆H₃(CF₃)₂}] dissolved in scCO₂–CH₃CN (molar ratio CH₃CN:CO₂ ~ 0.12) at 20 MPa and 328.15 K we obtain a maximum measured conductivity of ~ 3 mS cm^{−1}.

Studies of the conductivity as a function of concentration, Fig. 12, show an increase in molar conductivity with concentration for both [NⁿBu₄][B{3,5-C₆H₃(CF₃)₂}] and [NR^fBu₃][B{3,5-C₆H₃(CF₃)₂}]. At low electrolyte concentrations, Kohlrausch's law predicts that the molar conductivity should decrease with increasing electrolyte concentration.

$$\Lambda = \Lambda_0 - A\sqrt{c} \quad (1)$$

where Λ is the molar conductivity, Λ_0 is the sum of the limiting molar conductivities of the single ions (A⁺ and B[−]), A is a constant and c the concentration of the electrolyte.

However, for low dielectric constant solutions, such as supercritical fluids, the association of ions to form ion pairs and ion triples becomes significant. According to the Fuoss–Kraus model⁸⁰ the conductivity under these conditions is given by

$$\Lambda = \frac{\Lambda_0\sqrt{K}}{\sqrt{c}} + \frac{\lambda_0\sqrt{Kc}}{k} \quad (2)$$

where K is the equilibrium constant describing formation of the AB ion pairs, λ_0 is the sum of the limiting molar conductances of the two kinds of triple ions (ABA⁺ and BAB[−]) and k is the equilibrium constant describing the formation of the triple ions (assumed to be the same for both types of triple ion for simplicity). As we can see from Fig. 12 there is reasonable agreement between the data and the Fuoss–Kraus model at the lower electrolyte concentrations. The simple Fuoss–Kraus model of eqn (2) neglects the effects of the ion interactions on the conductivity (the electrophoretic and relaxation effects) and assumes that the activity coefficient is unity. More sophisticated treatments have been described in the literature that take these effects into account (see for example the work of Goldfarb and Corti⁷¹ or Abbott and Schiffrin⁸⁹ for applications of these) but these require values for the dielectric constant, viscosity or limiting molar conductivity which we do not have for this supercritical system. The results in Fig. 11 and 12 also show that the fluorinated ponytail does not have a significant influence

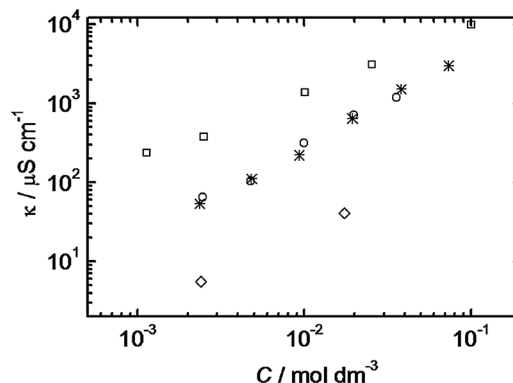


Fig. 11 Conductivity of different supercritical fluid electrolyte solutions. Electrical conductivity of supporting electrolytes in a supercritical fluid mixture of CO₂ + CH₃CN ($x_{\text{CO}_2}:x_{\text{CH}_3\text{CN}} = 0.89:0.11$) at 328.15 K and 20.0 MPa. $\diamond$, [NⁿBu₄][BF₄]; $*$, [NⁿBu₄][B{3,5-C₆H₃(CF₃)₂}]; $\circ$, [NR^fBu₃][B{3,5-C₆H₃(CF₃)₂}]; $\square$, [NⁿBu₄][BF₄] in pure CH₃CN at ambient condition.⁵

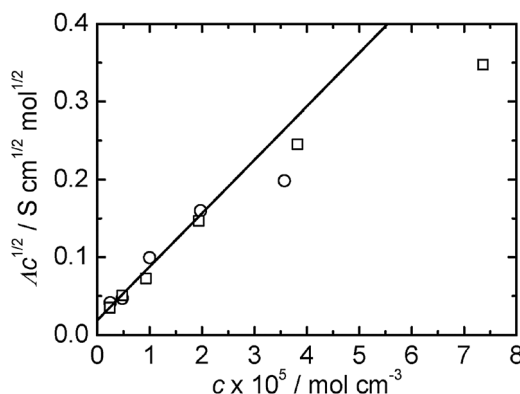


Fig. 12 Plot of $\Lambda\sqrt{c}$ against c for the data shown in Fig. 11: ($\square$) [NⁿBu₄][B{3,5-C₆H₃(CF₃)₂}] and ($\circ$) [NR^fBu₃][B{3,5-C₆H₃(CF₃)₂}]. The line is the best fit to the low concentration (below 3×10^{-5} mol cm^{−3}) data.³⁴

on the extent of ion pairing. From these experiments it is clear that triple ions make a significant contribution to conduction in scCO₂–CH₃CN.

We have carried out similar phase behaviour and conductivity studies for supercritical CHF₃, CH₂F₂, and CH₂FCF₃ containing [NⁿBu₄][BF₄], [NⁿBu₄][B{3,5-C₆H₃(CF₃)₂}] and Na[B{3,5-C₆H₃(CF₃)₂}]⁵². As we can see from Table 3 these three hydrofluorocarbons have reasonably accessible critical temperatures and pressures and all have dipole moments greater than 1.6 D. Compared to the CO₂ + CH₃CN system ($x_{\text{CO}_2}/x_{\text{CH}_3\text{CN}} = 0.14$) discussed above, the pressures required to form a homogeneous solution for [NⁿBu₄][BF₄] electrolyte are much lower (Fig. 13).

Table 3 Physical properties of some hydrofluorocarbons

Physical property	CHF ₃	CH ₂ F ₂	CH ₂ FCF ₃
T_c/K	299.29	351.26	374.21
p_c/MPa	4.832	5.782	4.049
$\rho_c/\text{kg m}^{-3}$	526.5	424.00	511.90
μ/D	1.65	1.98 ± 0.02	1.80 ± 0.22



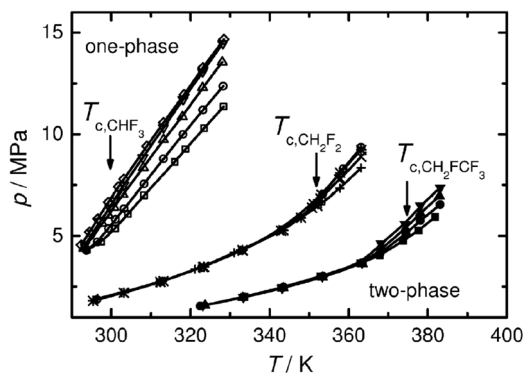


Fig. 13 p - T phase diagram of the mixture of $[N^nBu_4][BF_4]$ (1) + HFC (2). HFC = CHF_3 : $\square$, $x_1 = 0.82 \times 10^{-3}$; $\triangle$, $x_1 = 1.59 \times 10^{-3}$; ∇ , $x_1 = 4.83 \times 10^{-3}$; $\diamond$, $x_1 = 6.90 \times 10^{-3}$. HFC = CH_2F_2 : $+$, $x_1 = 1.67 \times 10^{-3}$; $\times$, $x_1 = 2.55 \times 10^{-3}$; $*$, $x_1 = 3.29 \times 10^{-3}$; $\oplus$, $x_1 = 4.17 \times 10^{-3}$; HFC = CH_2FCF_3 : $\blacksquare$, $x_1 = 1.33 \times 10^{-3}$; $\bullet$, $x_1 = 2.62 \times 10^{-3}$; $\blacktriangle$, $x_1 = 3.91 \times 10^{-3}$; $\blacktriangledown$, $x_1 = 5.14 \times 10^{-3}$. The arrows indicate the critical temperature of the HFCs.⁵²

For all three HFCs we find that when the temperature is below the corresponding T_c (marked by the arrows in Fig. 13) the p - T boundaries are overlapped for the mixtures with different composition of $[N^nBu_4][BF_4]$ but at higher temperature the solubility of $[N^nBu_4][BF_4]$ in the HFC fluid phase increases with increasing pressure. Although all three HFCs are good solvents for $[N^nBu_4][BF_4]$, CH_2F_2 gives the lowest reduced pressure, p_r (where $p_r = p/p_{c,HFC}$ the ratio of the pressure on the phase to the critical pressure of the corresponding HFC), for dissolving the same amount of $[N^nBu_4][BF_4]$. This may be explained by the fact that CH_2F_2 has the highest dipole moment of the three (Table 3).

Fig. 14 shows p - T phase diagrams for three binary mixtures of CH_2F_2 and three different electrolytes, $[N^nBu_4][BF_4]$, $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$, and $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$, at similar mole fraction ($x_{\text{electrolyte}} \sim 1.65 \times 10^{-3}$). The p - T phase boundaries for all of the mixtures are almost superimposed with only a slight difference in the phase separation pressure observed when $T > 353$ K. It is not surprising that two binary

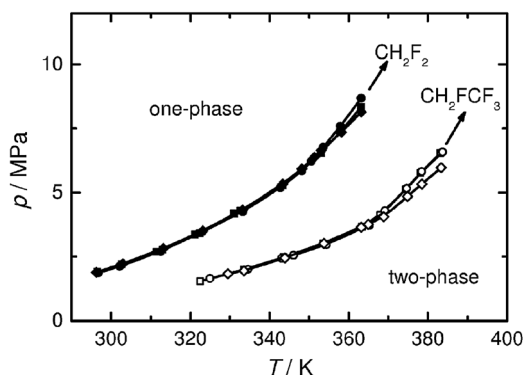


Fig. 14 p - T phase diagram of the mixtures of electrolyte (1) + CH_2F_2 (2): $\blacksquare$, $[N^nBu_4][BF_4]$, $x_1 = 1.67 \times 10^{-3}$; $\bullet$, $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$, $x_1 = 1.65 \times 10^{-3}$; $\blacklozenge$, $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$, $x_1 = 1.62 \times 10^{-3}$; and of electrolyte (1) + CH_2FCF_3 (2): $\square$, $[N^nBu_4][BF_4]$, $x_1 = 2.62 \times 10^{-3}$; $\circ$, $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$, $x_1 = 2.58 \times 10^{-3}$; $\diamond$, $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$, $x_1 = 2.59 \times 10^{-3}$.⁵²

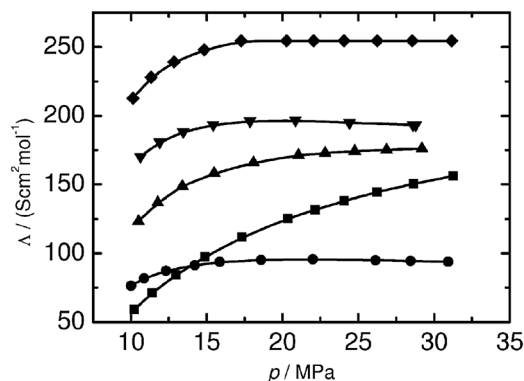


Fig. 15 Molar conductivity of the supporting electrolytes in CH_2F_2 at 363.15 K. The molar concentration of the electrolytes is $\sim 9.0 \text{ mol m}^{-3}$. $\blacksquare$, $[N^nBu_4][BF_4]$; $\bullet$, $[N^fBu_3][B\{3,5-C_6H_3(CF_3)_2\}_4]$; $\blacktriangle$, $[N^nBu_4][B(C_6F_5)_4]$; $\blacktriangledown$, $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$; $\blacklozenge$, $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$.⁵²

mixtures with $[N^nBu_4][BF_4]$ and $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$ show similar behaviour because CH_2F_2 is such a good solvent for tetrabutylammonium salts and, consequently, the effect of replacing $[BF_4]^-$ by the $[B\{3,5-C_6H_3(CF_3)_2\}_4]^-$ anion is not very significant. However, it is surprising that $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$ shows such similar behaviour, given that sodium salts often have low solubility in organic solvents. Here it would appear that the $[B\{3,5-C_6H_3(CF_3)_2\}_4]^-$ anion is a key component to enhance the solubility of $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$ in CH_2F_2 , and the large size difference between Na^+ and $[B\{3,5-C_6H_3(CF_3)_2\}_4]^-$ is probably important. Similar results are found for the same three supporting electrolytes in CH_2FCF_3 at a fixed mole fraction of $\sim 2.60 \times 10^{-3}$, also shown in Fig. 14.

Studies of the conductivity of $[N^nBu_4][B(C_6F_5)_4]$, $[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$, $[N^fBu_3][B\{3,5-C_6H_3(CF_3)_2\}_4]$ ($R^f = (CH_2)_3C_7F_{15}$), and $Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$ in $scCH_2F_2$, Fig. 15 and Table 4, show that these salts are more conducting than $[N^nBu_4][BF_4]$ under the same conditions although the increase is much less significant than that for supercritical $CO_2 + CH_3CN$. Consequently, either $[N^nBu_4][BF_4]$ or the corresponding $[B(R^f)_4]^-$ salts are good choices for supporting electrolytes for electrodeposition from $scCH_2F_2$.

Typical conductivities for 1 M electrolyte solutions in acetonitrile⁹⁰ are in the range $20\text{--}55 \text{ S cm}^2 \text{ mol}^{-1}$ so the values we obtain for the supercritical fluids are quite respectable for a

Table 4 Molar conductivity of the supporting electrolytes in CH_2F_2 at 363 K and 20 MPa. The concentration of all the electrolytes is $\sim 9.0 \text{ mol m}^{-3}$. Piecewise cubic Hermite polynomials were used to interpolate the data shown in Fig. 15 to the fixed pressure (20 MPa)

Electrolyte	$\Delta / \text{S cm}^2 \text{ mol}^{-1}$
$[N^nBu_4][BF_4]$	124
$[N^fBu_3][B\{3,5-C_6H_3(CF_3)_2\}_4]^a$	95
$[N^nBu_4][B(C_6F_5)_4]$	170
$[N^nBu_4][B\{3,5-C_6H_3(CF_3)_2\}_4]$	197
$Na[B\{3,5-C_6H_3(CF_3)_2\}_4]$	255

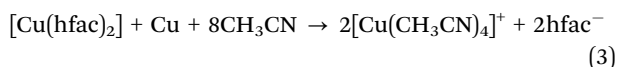
^a $R^f = CF_3(CF_2)_7(CH_2)_3$.⁵²



non-aqueous electrolyte and sufficient to carry out electrochemical studies.

5. Reagents

The reagents used in SCFED also require a number of key properties. In addition to solubility in the SCF, they need to be both stable to the electrolyte and at the working temperature, and ideally readily synthesised in high purity. Unwanted electrochemical side reactions which could result in fouling of the electrode or counter-electrode by the reagents and reactive liberated ligands must also be avoided. For example, $[\text{Ag}(\text{cod})(\text{hfac})]\text{BF}_4$ (cod = cyclo-octadiene; hfac = hexafluoroacetylacetonyl) is unsuitable for Ag deposition, probably because the cod fouls the counter electrode, whilst if $[\text{Cu}(\text{hfac})_2]$ is used for copper deposition in $\text{scCO}_2\text{-CH}_3\text{CN}$ the deposited copper can redissolve due to the reaction⁹¹



For deposition of transition metals from $\text{scCO}_2\text{-CH}_3\text{CN}$ the nitrile complexes $[\text{M}(\text{CH}_3\text{CN})_x][\text{BF}_4]_n$ ($\text{M} = \text{Fe}, \text{Co}, \text{Cu}, \text{Ag}, \text{etc.}$) are attractive reagents since the anion can be present in the electrolyte and the deposition liberates CH_3CN which is present as co-solvent, hence no chemical incompatibilities are presented.^{5,91} However in scCH_2F_2 , the complexes are unstable in the absence of added CH_3CN , which limits their usefulness, whilst $[\text{Cu}(\text{hfac})_2]$ is stable in scCH_2F_2 .⁵² The nitrile complexes are readily made from the metal powders and $\text{NO}[\text{BF}_4]$ in CH_3CN .^{92,93} In the case of $\text{Cu}(\text{I})$, $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ is obtainable from Cu_2O and aqueous HBF_4 in CH_3CN , or by refluxing $\text{Cu}[\text{BF}_4]_2$ with copper in CH_3CN .⁹⁴ The nitrile complexes can also be converted to $[\text{B}(\text{Ar}^F)_4]^-$ salts to increase solubility in the SCF if required.^{92,93} For silver electrodeposition several reagents were considered for use in $\text{scCO}_2\text{-CH}_3\text{CN}$ including $[\text{Ag}(\text{hfac})(\text{cod})]$, $[\text{Ag}(\text{CH}_3\text{CN})_4][\text{BF}_4]$, and $[\text{CF}_3(\text{CF}_2)_6\text{CO}_2\text{Ag}(\text{PPh}_3)_2]$ and $[\text{Ag}(\text{PPh}_3)_4][\text{BF}_4]$, in which the arylphosphine groups and the perfluorocarboxylate ligand were introduced to increase solubility in hydrofluorocarbons. Of these, $[\text{Ag}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ was found to be the most suitable for silver deposition from $\text{scCO}_2\text{-CH}_3\text{CN}$ (see below).⁹⁵

The choice of reagents to electrodeposit germanium is less obvious than for copper or silver. Germanium forms compounds in two oxidation states, $\text{Ge}(\text{II})$ and $\text{Ge}(\text{IV})$, and the chemical stability of its compounds varies widely with the ligands present.^{6,96} Many $\text{Ge}(\text{IV})$ complexes are moisture sensitive, whilst the thermal stability of $\text{Ge}(\text{II})$ complexes is often poor. Among the $\text{Ge}(\text{II})$ complexes explored were the commercially available GeBr_2 , a complex of the functionalised diimine $[\text{GeCl}_2(4,4'\text{-didecyl-2,2'-bipyridyl})]$,⁶ the trichlorogermanate(II) salt $[\text{N}^t\text{Bu}_4][\text{GeCl}_3]$, made by reduction of GeCl_4 in aqueous hydrochloric acid with H_3PO_2 , followed by addition of $[\text{N}^t\text{Bu}_4]\text{Cl}$,⁶ and $[\text{Ge}(12\text{-crown-4})_2][\text{CF}_3\text{SO}_3]_2$, made from $[\text{GeCl}_2(\text{dioxane})]$ and 12-crown-4. The first two proved to be too poorly soluble in scCH_2F_2 , and whilst it was soluble, $[\text{Ge}(12\text{-crown-4})_2][\text{CF}_3\text{SO}_3]_2$ degraded in the same solvent.⁶ $[\text{N}^t\text{Bu}_4][\text{GeCl}_3]$ was both soluble and stable in scCH_2F_2 and its electrochemistry is described below. Among the $\text{Ge}(\text{IV})$

reagents assessed were $[\text{GeF}_4(\text{CH}_3\text{CN})_2]$, made by passing GeF_4 gas into dry CH_3CN , and $[\text{GeF}_4(\text{Me}_2\text{NCH}_2\text{CH}_2\text{NMe}_2)]$, prepared from $[\text{GeF}_4(\text{CH}_3\text{CN})_2]$ and $\text{Me}_2\text{NCH}_2\text{CH}_2\text{NMe}_2$ in CH_2Cl_2 .⁹⁷ These two reagents were chosen as they are thermally relatively stable, and since they are coordinatively saturated, they are moisture stable solids and relatively resistant to hydrolysis in solution. Unfortunately, they proved to be insufficiently soluble in the SCFs. As a non-polar liquid, GeCl_4 (B.P. 365 K) was expected to be readily miscible with SCFs, but it is moisture sensitive and readily hydrolysed. It is commercially available and easily dried and purified by distillation from a mixture of Na_2CO_3 and CaH_2 .

6. Electrodeposition

There is very little published work on electrodeposition from supercritical fluids. As described above, the earliest work is probably that of Williams and Naiditch⁵⁴ in 1976 and the work of Silvestri *et al.*⁵⁵ five years later. Finally, MacDonald *et al.*⁶² in 1986 reported deposition of Cu from CuCl_2 under supercritical, or close to supercritical, H_2O containing KCl at 573 K and 8.2 MPa. Apart from the three papers on electrodeposition of metals there are also a few papers which describe the electrodeposition of conducting polymers from supercritical fluids. This is slightly different because in this case the neutral solution species (the heterocyclic monomer) is oxidised to produce a charged product (the oxidised polymer) which deposits with the counter ion on the electrode surface. The first paper claiming to electrodeposit poly(pyrrole) and poly(aniline) from a supercritical fluid appeared in 2002.⁹⁸ This group used CO_2 containing 13.1% acetonitrile, 0.16 M pyrrole and 0.16 M $[\text{N}^t\text{Bu}_4][\text{PF}_6]$ at 323 K and 9.65 MPa. Subsequent work published in 2005 by Yan *et al.*⁹⁹ using a view cell showed that under these conditions the system was biphasic and not in the supercritical state. Yan *et al.* were, however, able to deposit poly(pyrrole) films from supercritical fluid using CO_2 containing 13.1% acetonitrile, 0.16 M pyrrole and 0.04 M $[\text{N}^t\text{Bu}_4][\text{PF}_6]$ at 323 K and 10 MPa. This illustrates one of the potential pitfalls of working with, or trying to work with, supercritical fluids. It is not sufficient to rely on the critical temperature and critical pressure of the pure fluid, but rather it is essential to properly characterise the phase behaviour of the whole system. The honour of reporting the first successful electropolymerisation of a conducting polymer from a supercritical fluid consequently falls to Atobe *et al.*¹⁰⁰ who, in 2004, reported the electropolymerisation of poly(pyrrole) and poly(thiophene) from scCHF_3 containing 10 mM pyrrole or thiophene with 40 mM $[\text{N}^t\text{Bu}_4][\text{PF}_6]$ at 323 K and 15 MPa.

Before moving on to discuss our recent work, it is appropriate to mention the work of Sone's group on electrodeposition from dense emulsions of CO_2 . The group has published extensively, starting in 2002,¹⁰¹ on the electrodeposition of Ni from a dense emulsion of an aqueous Ni plating (Watts) bath formed in scCO_2 using surfactants (typical conditions 323 K and 15 MPa and 60 vol% of the aqueous plating bath stabilised with 1 vol% surfactant).¹⁰² This is not a single phase supercritical fluid. The advantages of the system



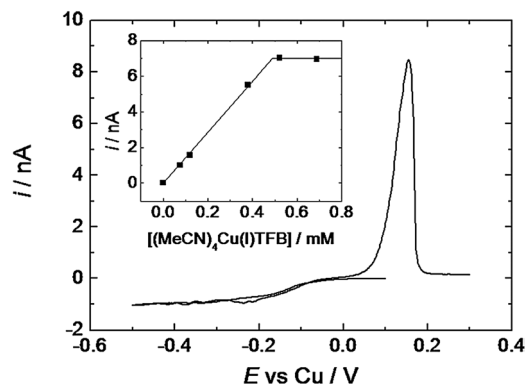


Fig. 16 Copper voltammetry performed in scCO_2 with 12.1 wt% CH_3CN and $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ (20 mM); at 310 K and 17.24 MPa. Electrodes were: 25 μm diameter platinum disc working electrode, 0.5 mm diameter platinum disc pseudo reference, and 0.5 mm diameter platinum wire counter electrode. The sweep rate was 20 mV s^{-1} . The insert shows the limiting current as a function of the concentration of the copper complex.⁵

are that it produces high quality Ni films with higher uniformity, smaller (sub 100 nm) grain size and significantly higher Vickers hardness than conventional electroplating from aqueous solution.¹⁰³ This is attributed, at least in part, to the enhanced desorption of hydrogen gas bubbles during deposition.¹⁰⁴ The same group has also applied this approach to the deposition of poly(pyrrole)¹⁰⁵ and copper.¹⁰⁶

In our work we have studied the electrodeposition of copper from supercritical fluids.^{5,91} Fig. 16 shows cyclic voltammetry for $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ in scCO_2 - CH_3CN with $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ electrolyte. The voltammetry shows the characteristic features expected for electrodeposition with mass transport limited reduction of $\text{Cu}(\text{I})$ to $\text{Cu}(0)$ at negative potentials, accompanied by a stripping peak for Cu on the return scan. The stripping peak is sharp and undistorted, showing that iR drop is insignificant in these experiments. The inset in Fig. 16 shows a plot of the mass transport limited currents recorded at a microdisc electrode for different concentrations of $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ added to the solution. This is a very sensitive way to determine the solubility of the redox active species since the mass transport limiting current at the microdisc is given by

$$I_L = 4nFaDc \quad (4)$$

where n is the number of electrons transferred and a is the radius of the microdisc electrode. Thus, assuming the diffusion coefficient, D , does not change with concentration, the limiting current should be proportional to the concentration of species dissolved in the solution. From the inset in Fig. 16 we can see that at low concentrations the current increases linearly with the concentration, as expected, but then abruptly reaches a plateau value, indicating that the solution is saturated. From the intersection of the two lines we obtain an estimate of the solubility of $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ under these conditions of 0.49 mM. From the slope of the initial part of the curve, using the microdisc equation (eqn (4)), we obtain a diffusion coefficient of $3.5 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$. This is approximately 1.6 times larger than the value measured for the same complex in acetonitrile at 311 K ($2.2 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$).

By changing the counter anion from $[\text{BF}_4]^-$ to $[\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]^-$ it is possible to significantly increase (>30 times) the solubility of $[\text{Cu}(\text{CH}_3\text{CN})_4]^+$ in scCO_2 - CH_3CN . This illustrates a significant advantage of careful choice and design of the electrolyte to enhance electrodeposition from supercritical fluids, the higher solubility of the reagent allows a greater rate of metal plating. For $[\text{Cu}(\text{CH}_3\text{CN})_4]^+$ in supercritical CO_2 with 12.1 wt% CH_3CN and 20 mM $[\text{N}^n\text{Bu}_4][\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]$ electrolyte at 310–311 K the diffusion coefficient is $3.3 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$. This is ~ 1.5 times larger than the corresponding value in scCO_2 - CH_3CN with 20 mM $[\text{N}^n\text{Bu}_4][\text{BF}_4]$, suggesting that there may be less ion pairing between the copper and the large $[\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]^-$ ion, leading to higher ion mobility. It is worth noting that these diffusion coefficients are relatively low compared to values for complexes dissolved in supercritical CO_2 .¹⁰⁷ This is probably due to the addition of 12–13 wt% acetonitrile as a co-solvent which significantly increases the viscosity of the solution.^{108,109}

Using $[\text{Cu}(\text{CH}_3\text{CN})_4]^+$ in scCO_2 - CH_3CN it is possible to electroplate smooth, reflective copper films onto 1 cm^2 electrodes. Fig. 17 shows SEM images of films deposited at different potentials. The morphology of the deposits varies with the applied potential as expected. At high overpotentials, when the deposition is mass transport limited, the films are rough and dendritic, whereas at low overpotentials the films are smooth, shiny and adherent. EDX, SIMS, and Auger studies all show that the films are high purity copper. Four-point probe measurements gave film resistivities for the best films of $4 \times 10^{-6} \Omega \text{ cm}$, which is comparable to values reported for copper electrodeposition from aqueous solution (1.75 to $2 \times 10^{-6} \Omega \text{ cm}$ depending on plating bath additives¹¹⁰) or by chemical deposition in supercritical CO_2 as reported by Watkins *et al.*¹¹¹ ($2 \times 10^{-6} \Omega \text{ cm}$). Thus the quality of the electrodeposited films is high and already close to that required for device applications.

An important feature of the copper plating bath described above is that it uses the $\text{Cu}(\text{I})$ complex. The relative stability of the $\text{Cu}(\text{I})$ and $\text{Cu}(\text{II})$ redox states strongly depends on the choice

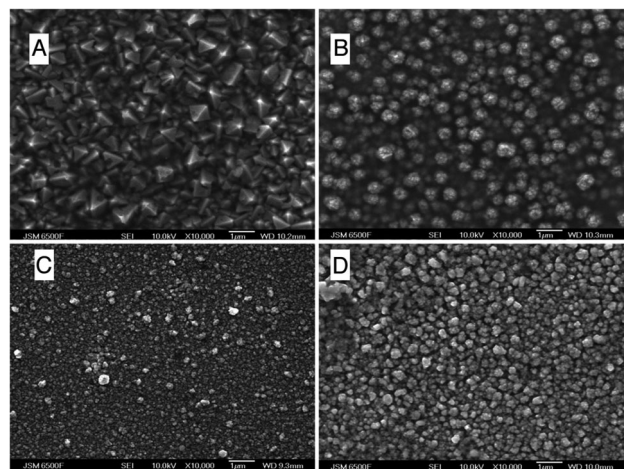


Fig. 17 Electroplated copper SEM images of copper films deposited from a saturated solution of $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ in 87.4 wt% CO_2 , 12 wt% acetonitrile, 0.6 wt% $[\text{N}^n\text{Bu}_4][\text{BF}_4]$ 309–311 K and 13.8 MPa at (A) -0.6 V , (B) -0.9 V , (C) -1.5 V , and (D) -1.7 V vs. Pt.⁵



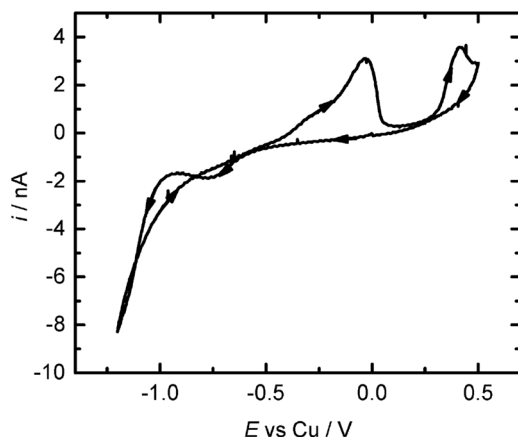


Fig. 18 Cyclic voltammetry for 1.2 mM $[\text{Cu}(\text{hfac})_2]$ in supercritical CO_2 with 12.1 wt% CH_3CN containing 20 mM $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ electrolyte. $T = 310\text{ K}$, $p = 17.2\text{ MPa}$. The working electrode was a 25 μm diameter platinum disc, the counter electrode was a platinum grid and the reference a 0.5 mm polished copper disc. Sweep rate 20 mV s^{-1} .⁵⁰

of ligands. For $[\text{Cu}(\text{CH}_3\text{CN})_4]^+$ the redox potential for oxidation to $[\text{Cu}(\text{CH}_3\text{CN})_4]^{2+}$ is high ($>1.25\text{ V vs. Cu in CH}_3\text{CN}$). It is possible to electroplate copper from the $[\text{Cu}(\text{hfac})_2]$ from $\text{scCO}_2\text{-CH}_3\text{CN}$, Fig. 18. Now however the comproportionation reaction, eqn (3), complicates the process so that this is not a good choice of reagent in this case. EDX analyses of these films shows contamination with fluorine and carbon.

We have also electrodeposited copper from supercritical hydrofluorocarbons. Studies of the phase behaviour of CH_2F_2 containing $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ and $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ showed that the copper complex was unstable in the absence of added CH_3CN .⁵² Consequently, a small amount of CH_3CN was added to the CH_2F_2 not only to increase the solubility of $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$, but also to stabilise it. Phase behaviour studies show that addition of $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ to the $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ system in scCH_2F_2 drastically shifts the phase boundary to high pressures at $T > T_{\text{c,CH}_2\text{F}_2}$.⁵² We believe that this is caused by selective solvation of the $[\text{N}^t\text{Bu}_4]^+$ and $[\text{BF}_4]^-$ ions by CH_3CN , depleting the CH_3CN molecules in the bulk phase, so that the density, and hence the pressure of the system, needs to be increased considerably for the quaternary mixture to form a homogeneous phase. Again this illustrates the importance of carefully characterising the phase behaviour of the electrolyte system.

$[\text{Cu}(\text{hfac})_2]$ is more soluble and more stable than $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ in scCHF_3 and it only increases the phase-separation pressure by a moderate amount when added to the CHF_3 solution with $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ as the supporting electrolyte.⁹¹ Fig. 19 shows a cyclic voltammogram of $[\text{Cu}(\text{hfac})_2]$ in scCHF_3 with free hfacH ligand added to the solution. The voltammetry is similar to that in CH_3CN and in $\text{scCO}_2\text{-CH}_3\text{CN}$, with the $[\text{Cu}(\text{hfac})_2]$ reduced to metallic copper in one step. In this case there is a significant cathodic background current beyond -0.7 V . Deposition onto large area ($\sim 1\text{ cm}^2$) electrodes was found to give good copper films with resistivities as low as $5.8 \times 10^{-6}\ \Omega\text{ cm}$. The best films were obtained with stirring and in the presence of the free ligand (hfacH).

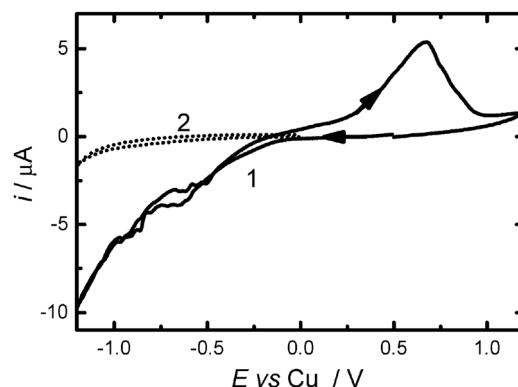


Fig. 19 Scan 1, cyclic voltammetry for 5.3 mM $[\text{Cu}(\text{hfac})_2]$ in scCHF_3 with 20 mM $[\text{N}^t\text{Bu}_4][\text{BF}_4]$ and 0.1 M hfacH at 311 K and 17 MPa. Scan 2 shows the background current in the absence of the Cu complex. The working electrode was a 0.37 mm diameter platinum disc, and the counter and reference electrodes were 0.5 mm diameter copper wires. Sweep rate 100 mV s^{-1} .⁵⁰

We have also studied the electrodeposition of silver from $\text{scCO}_2\text{-CH}_3\text{CN}$.⁹⁵ Five silver reagents were investigated. Of these the most suitable was found to be $[\text{Ag}(\text{CH}_3\text{CN})_4][\text{BF}_4]$ which gives good quality metallic deposits without any complicating electrochemical reactions. Of the other complexes $[\text{Ag}(\text{hfac})(\text{cod})]$ was found to give silver deposits, but polymerisation of the cod ligand on the counter electrode caused problems, and the silver deposition from $[\text{CF}_3(\text{CF}_2)_6\text{CO}_2\text{Ag}(\text{PPh}_3)_2]$, $[\text{Ag}(\text{hfac})(\text{PPh}_3)]$ and $[\text{Ag}(\text{PPh}_3)_4][\text{BF}_4]$ were all accompanied by electrochemical reduction of the PPh_3 ligand leading to a complex set of reactions with the electrolyte.

To exploit supercritical fluid electrodeposition to make devices it will be necessary to electrodeposit not only simple metals but also functional materials such as semiconductors. Germanium is an extremely important material in a wide range of technologies and single crystalline SiGe layers are widely used in integrated “silicon” electronics.¹¹² Electrodeposition of nanostructured germanium would offer a number of advantages over other deposition techniques if sufficient material quality could be achieved. In addition, electrodeposition of germanium is a stepping stone to the electrodeposition of silicon-germanium and silicon layers.¹¹³ We have investigated several reagents for Ge deposition from $\text{scCO}_2\text{-CH}_3\text{CN}$ and scCH_2F_2 .⁶ For Ge(II) reagents the most promising results were obtained using $[\text{N}^t\text{Bu}_4][\text{GeCl}_4]$. Better results were obtained using GeCl_4 . Fig. 20 shows the phase behaviour for binary mixtures of CH_2F_2 with GeCl_4 and $[\text{N}^t\text{Bu}_4]\text{Cl}$. At 357 K and 15 MPa the system forms a single phase, supercritical system. Fig. 21 shows typical cyclic voltammograms of GeCl_4 in scCH_2F_2 under these conditions. At the 250 μm Au disc electrode reduction starts at $\sim 0.7\text{ V vs. Pt}$ followed by a less well resolved increase in current cathodic of -1.5 V , presumably corresponding to the two-step reduction of GeCl_4 . In the limiting current region there are large, random fluctuations in the reduction current. These fluctuations arise from unsteady convective flow in the high-pressure electrochemistry cell because of the low viscosity of the fluid when the operating conditions are close to the critical



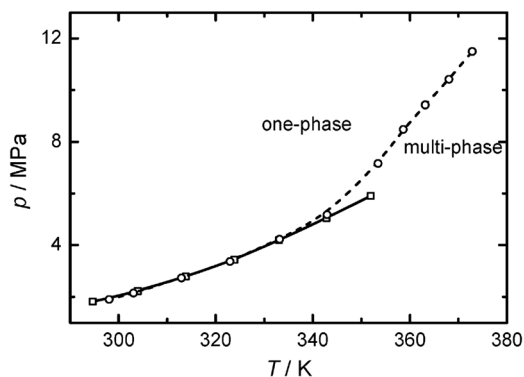


Fig. 20 p - T phase diagrams of the binary mixtures of CH_2F_2 + A: A = GeCl_4 ($\square$), $x_A = 4.4 \times 10^{-3}$; and A = $[\text{N}^+\text{Bu}_4]\text{Cl}$ ($\circ$), $x_A = 2.95 \times 10^{-3}$.⁶

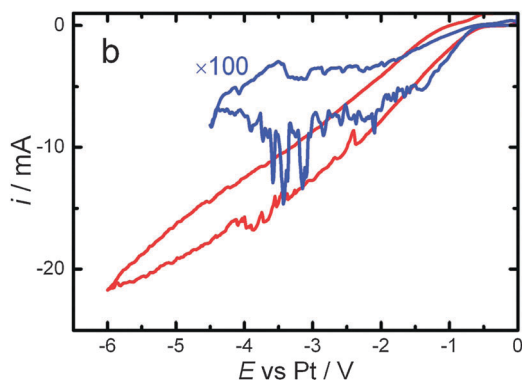


Fig. 21 Cyclic voltammogram of $0.06 \text{ mol dm}^{-3} \text{ GeCl}_4$ in scCH_2F_2 with $0.06 \text{ mol dm}^{-3} [\text{N}^+\text{Bu}_4]\text{Cl}$ as the supporting electrolyte. The working electrodes were a $250 \mu\text{m}$ Au microdisc (blue line, scan rate 0.1 V s^{-1}), and a 0.36 cm^2 gold-coated glass slide (red line, scan rate 0.1 V s^{-1}). The measurements were carried out at 357 K and 15 MPa .⁶

point of CH_2F_2 . We can also see that there is significant iR drop in this case. This is because the conductivity of the $[\text{N}^+\text{Bu}_4]\text{Cl}$ supporting electrolyte is low in scCH_2F_2 . These conditions can be used for bulk deposition of Ge, the ‘as deposited’ material was amorphous, but crystallised under high intensity laser illumination.⁶

7. Templates

Electrodeposition into small and high aspect ratio pore structures is a specific advantage of SCFED, and provides opportunities to grow high aspect ratio nanowires or to construct multi-component devices in electrically directed locations. Aligned pore structures will need to be used in future to deliver the promise of linking properties to designed nanostructures.

Electrochemical reaction of aluminium in an acidic environment is well known to produce porous anodic aluminium oxide (AAO) films and well-ordered hexagonal AAO films (Fig. 22a) are achievable if the anodisation conditions are well optimised.^{114–116} These materials can be produced with pore sizes from around 5 nm ¹¹⁷ to 500 nm ,¹¹⁸ and the resulting templates are electrically

and thermally insulating, optically transparent and robust. Typically they are grown on the surface of a solid aluminium foil electrodes, but can also be produced in sputtered aluminium films on other surfaces such as ITO-glass,¹¹⁹ suggesting opportunities to make porous structures on various substrates. Several authors have published electrodeposition of metals and other materials into these structures from more conventional electrolytes.^{120,121} Many of these porous films are now commercially produced, where they are cleaved from the aluminium surface to produce self-supporting membranes. The chemistry to produce similar structures in other oxides, especially titanium dioxide, is also now well developed.¹²²

Surfactant- and block copolymer-templated porous films are the commonest porous materials with small pore sizes, with pores ranging from below 3 nm to above 10 nm . It is possible to produce a variety of 1-, 2- or 3-dimensional pore structures controlled by a combination of surfactant type, concentration and temperature. Evaporation induced self-assembly (EISA) is a process in which the concentration increase that results from solvent evaporation during coating drives both the formation of micelles from the surfactant and condensation of the silica species in the sol. However, the EISA process is difficult to control. The linear pores that are present in hexagonal phase silica are particularly attractive hosts for electrodeposition and are the most heavily studied morphology. The alignment of the pores is controlled by the relative strength of the interaction between the micelle surfaces and the substrate compared with micelle-micelle interactions. In most cases the pores are either randomly oriented (referred to as ‘worm-like’) or are parallel to the substrate.¹²⁸ Parallel pores can be brought into long range alignment (Fig. 22b) by the addition of a low molar mass homopolymer to a block copolymer template film and then infiltrating a tetraethylorthosilicate (TEOS) precursor as a supercritical CO_2 solution.¹²³ Filling pores in this orientation is very interesting from an applications viewpoint due to the anisotropy that is present in the plane of the film, but pore access is more limited than in films with pores perpendicular to the film surface.

The tendency for pores to align with the substrate surface has been used to good effect in formation of mesoporous silica inside AAO membranes.¹²⁹ These have well-formed larger pores that are perpendicular to the face of the membrane. When a silica sol containing a surfactant at an appropriate concentration is infiltrated into one of these membranes the mesopores can be oriented along the length of the host pore (columnar growth) or can form toroidal or helical channels depending on deposition conditions. High humidity is frequently the key parameter in obtaining columnar growth, which is the most accessible morphology for electrodeposition (although electrodeposition into the other structure types has also been demonstrated). Copper, silver and tellurium nanowires have already been deposited in larger (P123-templated) columnar pores synthesised in this way (Fig. 22c).¹²⁴

Thin hexagonal mesoporous films with pores perpendicular to the plane of the film have been grown at an oil-water interface using CTAB (cetyltrimethylammonium bromide) surfactant.¹³⁰ These films were very thin and not useful as electrodeposition hosts. However, Nagarajan *et al.* have shown that supercritical



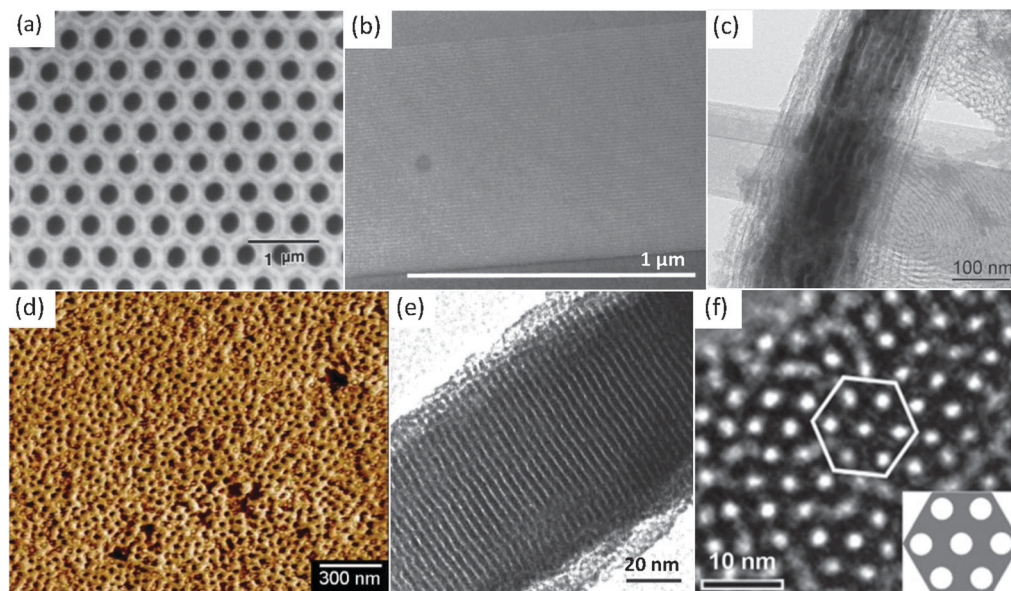


Fig. 22 Images of ordered porous structures. (a) SEM of anodically oxidised alumina with a hexagonal array of large ordered pores¹¹⁸ (Copyright 1998 The Japan Society of Applied Physics). (b) Cross-sectional TEM of long range ordered pores parallel to substrate surface produced from a block copolymer-homopolymer blend¹²³ (1 μm scale bar; reprinted (adapted) with permission from V. R. Tirumala *et al.*, *Chem. Mater.*, 2007, **19**, 5868. Copyright 2007, American Chemical Society). (c) Cu nanowires in columnar mesoporous silica originally grown in AAO¹²⁴ (reprinted (adapted) with permission from A. Keilbach *et al.*, *Chem. Mater.*, 2010, **22**, 5430. Copyright 2007, American Chemical Society). (d) Phase scanning force micrograph of a block copolymer film with cylindrical domains perpendicular to the surface infused with silica¹²⁵ (© WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim). (e) A cross-sectional TEM of an EASA-derived film with well-aligned perpendicular pores¹²⁶ (reprinted by permission from Macmillan Publishers Ltd.: A. Walcarius *et al.*, *Nat. Mater.*, **6**, 602, copyright 2007). (f) TEM of a flake of Stöber silica film showing the hexagonal pore structure¹²⁷ (20 nm scale bar; © 2012 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim).

CO_2 infiltration of TEOS into spin-coated films of the amphiphilic block copolymer, poly(α -methyl styrene- β -hydroxy styrene), which exhibits orientation of the cylindrical microdomains perpendicular to the film surface, results in perpendicular pore structures with pore sizes of 20 nm or larger (Fig. 22d).¹²⁵ These could be grown on conductive surfaces and hence be useful electrodeposition hosts. Recently three approaches have used more conventional surfactant-silica sol mixtures to make well-ordered hexagonal pore structures. Tolbert used a (111)-oriented cubic porous titania film as the substrate for growth of hexagonal porous silica, with the surfactant chosen to give a lattice match such that the perpendicular pore structure is inherently favoured from the base of the pores.¹³¹ Walcarius has used an “Electro-assisted self-assembly” (EASA) method to control pore orientation.^{126,132} The application of an electric field has two roles in this method, it aids the formation of micelles from the cationic surfactant (CTAB) on the electrode at the same time as generating hydroxyl ions *via* water oxidation which aids local condensation of the sol particles at the electrode surface. This method produces well-aligned pores with ~ 3 nm diameter (Fig. 22e) in films with thickness up to ~ 100 nm. A key advantage is that a conductive substrate, typically indium tin oxide (ITO), is used in growth and so is inherently available as the electrode for electrodeposition. Thicker films with similar pore size can be grown by a related route developed by Zhao and carried out with a dilute Stöber silica solution containing CTAB.¹²⁷ The method relies on the formation of CTAB micelles at the substrate surface under weakly basic conditions in which TEOS undergoes slow hydrolysis and again produces a hexagonal array of pores

perpendicular to the substrate surface (Fig. 22f). Whilst these films do not need to be grown on conductive surfaces the authors report the best ordered films on ITO.

Fig. 23 shows results for the deposition of copper into nanoscale (<10 nm) pores. The templates were prepared by

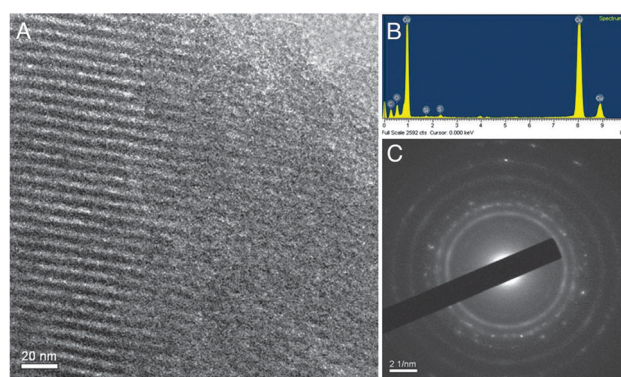


Fig. 23 Copper nanorods in mesoporous silica. (A) TEM of copper nanorods electrodeposited into silica mesopores from a solution of scCO_2 with 12.1 wt% CH_3CN , 20 mM $[\text{N}^+\text{Bu}_4][\text{B}(\text{3,5-}\text{C}_6\text{H}_3(\text{CF}_3)_2)_4]$, and 2 mM $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{B}(\text{3,5-}\text{C}_6\text{H}_3(\text{CF}_3)_2)_4]$ at 311 K and 17.24 MPa. The working electrode was a 0.8×0.5 cm ITO on glass slide modified with approximately 250 nm thick film of mesoporous silica; the counter electrode was a large, coiled, copper wire and the reference electrode was a 0.5 mm diameter copper disc. (B) EDX spectrum recorded from the same TEM sample. (C) Selected area electron diffraction (SAED) pattern recorded on 1 of the copper rods.⁵



dip coating ITO slides in a solution of Brij56, TMOS, 0.5 M HCl and MeOH (1 : 1.8 : 1 : 3.8 by weight). After calcining, this produces an adherent mesoporous silica film roughly 200 nm thick containing a regular hexagonal array of approximately 3 nm diameter cylindrical pores approximately 6 nm apart but randomly aligned with respect to the substrate. Electrochemical deposition of copper was carried out from $[\text{Cu}(\text{CH}_3\text{CN})_4][\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]$ in $\text{scCO}_2\text{-CH}_3\text{CN}$ containing 20 mM $[\text{N}^+\text{Bu}_4][\text{B}\{3,5\text{-C}_6\text{H}_3(\text{CF}_3)_2\}_4]$. The TEM image in Fig. 23A shows the regular array of cylindrical pores clearly visible in cross section on the left hand side of the image. Evidence that the pores are filled with copper is provided by the EDX analysis, which shows strong copper as well as the expected silicon and oxygen signals, Fig. 23B, and by the selected area electron diffraction results, Fig. 23C, which show a combination of diffuse rings attributed to silica together with rings that are consistent with the expected (111), (200), (220), and (311) diffraction peaks for copper. These results demonstrate the proof-of-principle, but much remains to be done to improve alignment and produce nanowire devices.

8. Conclusions and future prospects

The barriers to successful electrodeposition from a supercritical fluid are significant, but the field has now progressed beyond the stage of preliminary studies to the point where generally applicable methods for supercritical fluid electrodeposition have been developed. An important contribution has been to establish a range of supporting electrolytes capable of achieving high ionic conductivities in the low dielectric constant fluids. Supercritical fluid electrodeposition has been used to deposit a range of different materials, *e.g.* Ag, Cu, and Ge, from complex ions with different overall charges, different ligand types and with different oxidation states. It has been demonstrated that a range of supercritical fluids, including both those with moderate dielectric constants and those with low dielectric constant when combined with a suitable co-solvent, can be used to electrodeposit films. Whilst not every combination of supercritical fluid and target material will be possible, the existing work clearly indicates that the range of materials that can be deposited could be expanded significantly, for example to a wider range of metals, semiconductors, binary and ternary compounds and alloys. The work so far has demonstrated that, by suitable design and choice of the combination of anion and cation, it is possible to achieve supercritical fluid electrolytes with surprisingly large conductivities that will allow deposition at sufficiently high rates to be useful. The field of supercritical fluid electrodeposition is still at an early stage but progress has been good and the prospects are very encouraging. In taking this work forward our increasing knowledge of the underpinning physical chemistry, in terms of the phase behaviour of the supercritical fluid mixtures, the solubility, stability and conductivity of electrolytes and complex reagents in supercritical fluids, the phase behaviour of supercritical fluids in confined spaces, the structure of the double layer at the supercritical fluid/electrode interface, the kinetics

of ligand exchange and electron transfer reactions in supercritical fluids, *etc.*, will be very important.

Considerable advances have been achieved in the field of supercritical fluid electrodeposition, but it is important not to lose sight of the fact that electrodeposition from a supercritical fluid will always involve significant practical challenges due to the high pressures involved. This means that supercritical fluid electrodeposition will only become technologically important if there are very clear advantages and novel applications which only supercritical fluid electrodeposition can fulfil. The particular properties of supercritical fluids strongly suggest that just such novel applications will exist, most probably in the area of complex and extreme nanostructures. However, as yet definitive proof of this has not been demonstrated and this has to be one of the key priorities for the field. What are the most likely candidates for these novel applications? One way of viewing supercritical fluid electrodeposition is as an extension of non-aqueous electrodeposition to higher temperature. Whilst lower deposition temperatures are often advantageous economically, better material quality, *e.g.* crystallinity, often requires an elevated deposition temperature. For any specific solvent the critical temperature is a fundamental limit above which electrodeposition can only be performed using the supercritical fluid electrodeposition technique. One area in which supercritical fluids are already showing promise is their ability to penetrate into very small pores. In particular supercritical fluid electrochemistry has been used to electrodeposit into 3 nm pores within mesoporous silicas.

It will take some time, and more work, to discover whether supercritical fluid electrodeposition can open up novel areas in nanomaterials and structures; however there are a number of shorter term goals which will enable the field to develop into new and exciting areas. These include, from a physical chemistry perspective, a better understanding of the structure of the double layer and of electrolyte solvation in supercritical fluids, studies of electron transfer kinetics at the electrode supercritical fluid interface, studies of speciation in supercritical fluids, and a greater understanding of the phase behaviour of supercritical fluids and supercritical fluid electrolytes in confined spaces. In addition there is much scope for practical advances in the design of new cells, in solvent purification and for the application of supercritical fluids in electrosynthesis.

In conclusion, supercritical fluid electrochemistry has come a long way in a relatively short time and has achieved a number of goals which many people thought were not possible. It is an exciting field with significant scientific challenges and we hope that this perspective will stimulate others to consider how electrochemistry in supercritical fluids might be applied not only in materials deposition, but also in other areas.

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