Open Archive TOULOUSE Archive Ouverte (OATAO)

OATAO is an open access repository that collects the work of Toulouse researchers and makes it freely available over the web where possible.

This is an author-deposited version published in: http://oatao.univ-toulouse.fr/
Eprints ID: 16765

To link to this article: DOI:10.1149/2.0101608jes
URL: http://dx.doi.org/10.1016/j.carbon.2016.05.010


Any correspondence concerning this service should be sent to the repository administrator: staff-oatao@listes-diff.inp-toulouse.fr
Non-Equilibrium Ionic Liquid-Electrode Interface at Elevated Temperature and Its Influence on Co$^{2+}$ Reduction Process

M. Tułodziecki, a J.-M. Tarascon, b,c P.-L. Taberna, d and C. Guéry a,z

a Laboratoire de Réactivité et Chimie des Solides and ALISTORE European Research Institute, UMR CNRS 7314, UFR des Sciences, 60039 Amiens cedex, France
b Chimie du Solide et Energie, UMR 8260, Collège de France, Berthelot, 75005 Paris, France
c Réseau sur le Stockage Electrochimique de l’Energie (RS2E), FR CNRS 3459, France
d CIRIMAT/ICMIE and ALISTORE European Research Institute, UMR CNRS 5085, 31062 Toulouse cedex 9, France

Electrodeposition in Ionic Liquid (IL) media is still at its infancy stage. Recently much attention is devoted to exploration of the IL-electrode interface for better understanding of the electrochemical processes at the electrified surfaces. Herein, we provide a new perspective of the temperature-driven IL-substrate interface structuring. We observed an increase in Co$^{2+}$ reduction kinetics induced by applying a high temperature gradient. In contrast, keeping the cell at a high constant temperature (100°C) leads to slower Co$^{2+}$ reduction kinetics, it results in a shift of the reduction onset and evolution of the cyclic voltammogram shape with time. We could ascribe these phenomena to the disturbance of the electrochemical double layer (induced by heating) and its further slow reorganization at constant temperature. We showed that such non-equilibrium behavior can be beneficial for the synthesis of metallic thin films from ILs. Electrochemical quartz crystal microgravimetry, impedance spectroscopy, cyclic voltammetry and chronoamperometry were employed in this work.

Ionic Liquids (ILs) are superior media for electrodeposition process as they overcome the main drawbacks of common aqueous, organic and molten salts electrolytes. They offer high thermal and electrochemical stability, are not flammable, have negligible vapor pressure and high ionic conductivity.1–2 Thanks to these attractive properties, ILs are considered as alternative media to molecular solvents for electrolytes in electrochemical devices (batteries, supercapacitors, transistors)3–8 or reacting media for synthesis of organic and inorganic compounds (metals, alloys, sulfides, oxides).9–12 Moreover, ILs can be tailor designed to meet specific properties, hence broadening their area of application.13

A primary issue, regarding electrochemical devices, is the structure of the electrochemical double layer (EDL), which depends on the solvent and the electrode’s substrate. It has been proven both theoretically14–17 and experimentally18–21 that the IL’s ions are highly structured and form layered EDL on either metal19 or metal oxide20 based electrodes. Depending on surface polarization, the arrangement and conformation of the anions and cations may change to screen the excess surface charge more efficiently.14–16,18,21,24 When polarized, the biggest changes within EDL are observed in the first adjacent layer; the strength of ion-electrode interactions,18 the compacity,23,25 the conformational changes of ions26 vary greatly within this particular layer. In case of further layers mineral effects were observed as thickening of the ordered layer.26 Moreover, different kinds of superstructures and surface reconstructions have been observed, especially at single crystal electrodes.18,21,26–28

The kinetics of the interfacial processes can vary from microseconds to even few days.29–31 In this context, Anderson et al. 32 found that 2h conditioning of a bismuth single crystal electrode in pyridinium based IL leads to a thicker EDL and the creation of superstructures. Endres and Atkin et al. 30,34 reported that nearly one day is required for an IL-gold electrode to reach equilibrium after its disturbance by polarization. Lastly, Drüschler et al. 37 reported a slow EDL charging process with a time scale of few seconds as probed by electrochemical impedance spectroscopy.

Such complex structures have already shown to have a tremendous impact on the electrochemical processes. Katayama et al.35,36 found that the strong ordering of the EDL causes the slow reduction kinetics of metal cations (i.e. need for high overpotentials). Endres and coworkers37,38 reported that the electrochemical response of the system as well as physical properties of metal deposits were very different, depending on the IL cation, and they ascribed it to the EDL structure. Additionally, the strong electric field induced at the Ionic Liquids-oxides interface was found to activate the oxygen vacancy formation which is of great importance in transistors field.6–8

However, despite intensive investigation of the anomalous EDL structure in ILs (reconstructions, slow relaxations, conformational changes), our understanding of the influence of different parameters (temperature, bath composition) on the EDL and consequently on the real electrochemical process is still slim. Any change in experimental conditions that can affect the EDL can also affect the rates of the heterogeneous electron transfer processes. In our recent studies, we have shown that the changes in EDL structure upon polarization had a tremendous impact on the reduction kinetics of metal cations depending on the applied negative potential.39 Herein, we report the influence of the heating rate (rapid or slow) on the Co$^{2+}$ reduction process as well as evolution of the system at constant high temperature. We show that both the way of heating and keeping the cell at elevated temperatures affects the Co$^{2+}$ reduction kinetics in great extent. We also show the beneficial effect of disturbing the interfacial structure for growing metallic thin films of controlled morphology by electro-deposition in ILs.

**Experimental**

Various electrochemical baths were made by dissolving M(TFSI)$_2$ (M = Co, Zn, Ni, Mn) salts in three selected ILs EMImTFSI, BMPTFSI, and BMImTfO, all purchased from Solvionic, Table I. These 99.9% pure (H$_2$O < 0.005%) ILs were purified under vacuum at 75°C for 2 days prior to being used, so that the final water content was ~6–20 ppm as deduced by Karl Fischer measurement. All chemicals were handled and stored in an argon-filled glove box (O$_2$ and H$_2$O < 0.1 ppm).

A three-electrode set-up was used for electrochemical measurements (chronoamperometry, cyclic voltammetry, impedance spectroscopy). Three different substrates, planar platinum electrode ($P_{Pt}$)
S = 0.020 cm², platinum wires (Pt₆) and stainless steel wires (SS₆), were used as working electrodes (WE). The surface area of the wire-type electrodes was estimated according to the length of the submerged fraction of the electrode (surface ranging from 0.017 to 0.033 cm²). A platinum grid with high surface area (S ≈ 2 cm²) and an Ag wire covered with Ag₂O were used as counter electrode (CE) and pseudo reference electrode (RE), respectively. The stability of RE was checked by reproducibility of the cyclic voltammetry (CV) curves of Co²⁺/Co and ferrocenium/ferrocene redox processes between multiple experiments and with time in single experiment. The CVs were highly reproducible even after few days of immersion in the cell. The WE was calibrated versus the ferrocenium/ferrocene redox couple. The mid-point potentials (Eₑ₀/₂ = Eₓ’anodic| cathodic) at different conditions are given in Table II. The obtained values were reproducible with maximum deviation of ±13 mV. All potentials are given versus the pseudo Ag|Ag₂O reference electrode. Prior to being used, the platinum WE wires were polished with 0.1 µm abrasive paper and electrochemically activated in 0.5 M H₂SO₄ by CV with a scanning rate of 500 mV/s (the CV profile was comparable with standards representative of an impunity free Pt surface). Next, the electrodes were rinsed with distilled water, ethanol, acetone and deionized water. The counter electrode was cleaned in 0.5 M nitric acid in an ultrasonic bath for 1h, and then intensively rinsed with distilled water, ethanal, acetone and deionized water. All the components of the cell were dried in an oven at T = 50°C for at least 12 hours prior to the experiment. Preparation of the solution, assembly and sealing of the cell were done in a glove box. The sealed cell (with controllable atmosphere) was then transferred out of the glove box for further manipulation. The seal of the cell was checked by monitoring the water content evolution over time. After 6 hours outside the glove box, water uptake of only 1–2 ppm was detected meaning that the cell was well sealed. Two heating methods were studied i) fast heating that induces high temperature gradients and ii) slow heating resulting in low temperature gradients. In case of fast heating, the electrochemical cell was immersed directly in the hot bath (either 60°C or 100°C). The temperature equilibration time was 15 min – as measured by EQCM. For slow heating method, the cell was heated progressively in few steps with heating rate of 1°C per minute. After each step of ΔT = 20°C, the cell was isothermally conditioned for 1 h. All heating processes of the cell were done with the WE electrode being at open circuit potential (OCP) if not otherwise specified. The temperature of the oil bath was controlled within ±2°C accuracy at 100°C.

The electrochemical measurements were carried out with an Autolab PGSTAT30 potentiostat (Eco Chemie BV). The electrochemical impedance spectroscopy (EIS) measurements were done on the Pt₆ electrode at a fixed potential that is the OCP value determined prior to starting each EIS measurement. It means that during the measurement the potential is fixed while in between the measurements the cell is left at OCP. The fixed potential is superimposed with 10 mV amplitude AC signal. The impedance data were measured within the frequency range from 1 MHz to 100 mHz.

Electrochemical quartz crystal microbalance (EQCM) measurements were conducted using a commercial SEIKO microbalance (SEIKO QCA 922) with AT-cut 9 MHz quartz covered with platinum on both sides. The electro-active geometric surface area was S = 0.196 cm². Simultaneous measurements of the quartz frequency and motional resistance were performed in order to follow both the change in weight of electrode and the variation in Co²⁺ concentration in the vicinity of the electrode (by following the change of viscosity and density product of the solution). The EQCM was calibrated with a CV measurement in 0.05 M solution of AgTFSI in EMImTFSI, at 100°C, with 2 mV·s⁻¹ scanning rate. The proportionality constant (C = Δm/Δf) was determined between the change of quartz frequency (Δf) and change of deposited mass (Δm). A constant value C = 1.056 · 10⁻⁹ g·Hz⁻¹ close to the theoretical one C₄ = 1.068 · 10⁻⁹ g·Hz⁻¹ was found, suggesting that issues regarding viscosity of the bath or roughness of the deposit can be neglected. Three different experiments were performed: 1) identifying the frequency and motional resistance changes with time as the bath is being fast heated up and then kept at 100°C for 8 h 2) checking the effect of long time conditioning at 100°C on chronoamperometric deposition of Co at −0.9 V and 3) CV in anodic direction after 8 h of heating.

The obtained deposits on quartz crystal were soaked in chloroform to remove adsorbed IL from the surface. Their morphology and elemental analysis was further checked using a Philips XL 30 field emission gun FEG microscope.

All experiments were performed at least twice to check the repeatability.

### Results

#### Cyclic voltammetry in pure EMImTFSI—Before studying the reduction process of Co²⁺, the blank CVs were conducted in pure EMImTFSI on a Pt₆ electrode at RT and 100°C. Two half cycles (cathodic and anodic, each on a freshly prepared electrode) were recorded starting from OCP (Fig. 1). During the cathodic half cycle (red curves), a small reduction wave – (R₁) was observed at both temperatures. Most likely, the R₁ signal originates from the reconstruction of an interfacial layer, as reported in literature (specific adsorption of ions, rearrangement of ions). On the anodic half cycle, an oxidation signal – (O₁) and corresponding reduction signal – (R₂, clearly visible in CV at RT) were observed. The nature of O₁ signal is not clear both the EDL reconstruction and/or beginning of ILs decomposition could be responsible for it.

**Figure 1.** Half cycles in pure EMImTFSI under Ar atmosphere a) at RT and b) at 100°C. The red curves represent the cathodic half cycle and black anodic ones. WE – Pt₆, CE – Pt, RE – Ag|Ag₂O, scan rate 5 mV·s⁻¹. A freshly prepared electrode was used for each measurement.

| Table II. Calibration of the Ag|Ag₂O reference electrode at different conditions. |
|--------------------------|-------------------|-------------------|
| Solution | Temperature | Atmosphere | Eₑ₀/₂ (V vs FeFc/FeFc⁺) |
| Pure EMImTFSI | RT | Ar | 0.244 ± 0.008 |
| 0.017 | 100°C | Ar | 0.225 ± 0.008 |
| 0.12 M Co(TFSI)₂ | RT | Ar | 0.102 ± 0.005 |
| In EMImTFSI | 100°C | Ar | 0.103 ± 0.007 |

### Table II. Calibration of the Ag|Ag₂O reference electrode at different conditions.
Both, the cathodic and the anodic sweeps end with a progressive increase in current density due to the decomposition of the IL. The cathodic ($E_c$) and anodic ($E_a$) limits were determined from Tafel plots (Supporting Information Fig. 1+). Values of $-1.85$ and $-1.70$ V for $E_c$ at RT and $100\, ^\circ$C were obtained, respectively. In the case of the anodic limit, the potential of $1.7$ V was estimated for both RT and $100\, ^\circ$C. Within these potential ranges, electrochemical processes should not be affected by the decomposition of EMImTFSI, hence it was selected for our experiments.

**CV of 0.12 M Co(TFSI)$_2$ solution in EMImTFSI.** Heating method dependence.—Next, the electrochemical response of 0.12 M Co(TFSI)$_2$ solution at $P_{eq}$ electrode was studied at different temperatures. Firstly, we compared the influence of the heating step on the reduction process of the Co$^{2+}$ cations. Two experiments were performed i) fast heating by dipping the cell in pre-heated bath and ii) slow heating by a step-wise increase of temperature (see experimental). For fast heating, the CVs were recorded with $30$ min delay after immersion of the cell in the hot bath, while for slow heating, the CVs were recorded $15$ min after the bath reached the desired temperature. The difference in delay accounts for the time needed for the bath to reach desired temperature (see experimental).

For both heating methods, the reduction sweeps at RT, $60\, ^\circ$C and $100\, ^\circ$C, are shown in Figure 2. In all the cases, two reduction peaks (I, II) separated by a drop in current density were observed. As studied in our previous work, the two reduction waves are due to Co plating, while the intermittent drop in current is caused by double layer reconstruction that hinders the electrochemical process. Nevertheless, the shape and reduction onset of the first peak changes greatly with both the temperature and heating method. In case of fast heating (solid lines), the intensity of the first reduction peak increased greatly with temperature. Moreover, the onset of the reduction process shifted to higher potentials (RT $-0.94$ V; $60\, ^\circ$C $-0.85$ V; $100\, ^\circ$C $-0.70$ V). For slow heating (dashed lines), the intensity of the first reduction signal increased as well, however only small changes of the reduction onset were observed (RT $-0.94$ V; $60\, ^\circ$C $-0.94$ V; $100\, ^\circ$C $-0.92$ V). Lastly, comparing the CVs for the two heating methods, we can clearly see that the onsets of the reduction processes are shifted to more positive values for the fast heating method. Considering that the reduction overpotential is related to EDL structure, the difference in onset potential is most probably due to the disturbance of the EDL caused by a high temperature gradient. This effect is discussed more in detail in the discussion part.

The CV dependence on temperature in case of the fast heating is in agreement with studies by Katayama et al. where they also observed a shift of the reduction signal to positive values with increasing temperature (Co$^{2+}$ and Ni$^{2+}$ reduction processes). However, lack of information on the method of heating and huge differences in the shape of the reduction curve limits direct comparison with our results.

**Evolution of the system with time at high constant temperature.** Further, we checked the evolution of the reduction process with time when the cell is being kept at a constant temperature (RT, $60\, ^\circ$C and $100\, ^\circ$C), which was reached with either a slow or fast heating method. At room temperature (Fig. 3a), the reduction sweeps did not reveal any significant changes with time; the CVs were reproducible even after $64\, h$.

In the case of the fast heated system to $60\, ^\circ$C (Fig. 3b), a small shift of the Co$^{2+}$ reduction onset was observed from $-0.85$ V to $-0.96$ V after $11\, h$. For $100\, ^\circ$C, more drastic changes were witnessed (Fig. 3c). After $2\, h$, the onset of the first reduction peak rapidly shifted to lower potentials (from $E = -0.7$ V to $E = -0.92$ V) and its shape changed. For longer times, the reduction onset stabilized at around $-1$ V (Fig. 3b), while the intensity of the first reduction peak strongly diminished. Finally, after $8\, h$, we obtained a curve that was similar in shape to the one found at RT, with the exception of the lower voltage part where the decomposition of the ionic liquid takes place ($< -1.7$ V). Such changes are accompanied by a slight variation of the OCP value ($\Delta E_{OCPP} = 70$ mV) that stabilizes for times longer than $8\, h$(SI, Fig. 2+).

In the case of a slow heated system to $100\, ^\circ$C (Fig. 4c), an evolution in the intensity of the first peak was also observed. After $5\, h$ (black line), we obtained a curve like the one observed for fast heated system after $8\, h$ (Fig. 4a - black solid line). However, only minor changes in the reduction onset were observed (from $E_{onset} = -0.92$ V to $E_{onset} = -0.95$ V after $5\, h$).

From the above constant temperature experiments, the presence of two coupled effects can be distinguished. First, the reduction onset...
b) Reduction onset shift, fast initial heating

In order to distinguish between these two and/or ii) a passivation layer due to chemical decomposition – solid electrolyte interface (SEI). In addition, the intensity of the first reduction wave decreased with time (SI, Fig. 3∗) in a similar manner to previous experiments at 100 °C. This indicates that the change in the curve shape at 100 °C is due to interfacial phenomena rather than diffusion of cobalt species. There are at least two possibilities to account for this scenario, which include formation of i) a strongly blocking double layer and/or ii) a passivation layer due to chemical decomposition – solid electrolyte interface (SEI). In order to distinguish between those two paths, the evolution of the interface at 100 °C was investigated by an EQCM and impedance spectroscopy experiments.

EQCM measurements.—The applied EQCM experiment protocol consisted of i) setting the potential at a predetermined OCP value at RT, ii) fast heating to 100 °C (dipping the cell in the pre-heated bath) and maintaining it at this temperature for 8 hours while monitoring the frequency and motional resistance of the quartz. Similar experiments were also conducted in pure IL for reference.

Figures 5a, 5b, 5c shows the EQCM results, in 0.12 M Co(TFSI)2 solution in EMIM-TFSI, of the heating (domain I) and constant temperature (domain II) steps together. During the initial heating (I), a sharp increase in frequency and decrease in motional resistance was observed. We ascribed it mainly to changes in viscosity/density parameters of the electrolyte, which are temperature dependent.49

During the process of keeping the cell at 100 °C (II), we observed a stabilization of both frequency and resistance. The curves are not smooth but rather noisy due to the presence of small intensity oscillations (30–80 Hz, 4–13 Ω). We found that small variations in temperature (±2 °C) are at the origin of these oscillations, as both the frequency and the temperature oscillation periods were very similar (8–9 min). The maximum/minimum of the frequency corresponds very well with the minimum/maximum of the motional resistance, even for the highest variation, SI Fig. 4∗. Thus, we ascribed them to changes in both viscosity/density of the IL and properties of the quartz crystal.50,51 The linear trend plot of the frequency and motional resistance (domain II), resulted in almost parallel lines to the x axis (Fig. 5 green lines), indicating no mass changes at the electrode surface.

For the sake of completeness, a similar EQCM measurement was carried-out on pure IL, with the only difference being that the cell

Potential shift to lower values (only for the fast heated system for both 60°C and 100°C). Second, the diminishment of the first reduction wave with time that occurs in both systems (fast and slow heated).

To gain further insight into the CV shape change at 100 °C, we studied its variation as function of the stirring conditions and cobalt concentration. These two parameters did not affect the trend in reduction curve change: Strong decrease at the beginning of the heating process and its further stabilization at E ~ −1 V was again observed (Fig. 4b). In addition, the intensity of the first reduction wave decreased with time (SI, Fig. 3∗) in a similar manner to previous experiments at 100 °C. This indicates that the change in the curve shape at 100 °C is due to interfacial phenomena rather than diffusion of cobalt species. There are at least two possibilities to account for this scenario, which include formation of i) a strongly blocking double layer and/or ii) a passivation layer due to chemical decomposition – solid electrolyte interface (SEI). In order to distinguish between those two paths, the evolution of the interface at 100 °C was investigated by an EQCM and impedance spectroscopy experiments.
was kept at OCP during all the experiment, SI, Fig. 5. The curve trends were similar to the one obtained in 0.12 M Co(TFSI)₂ solution. Namely, we observed a large increase in frequency during initial heating and constancy of the frequency and motional resistance when the temperature is kept constant at 100°C for 8 h.

The constancy of the frequency and motional resistance, in both experiments, suggests that no formation of an extra organic/inorganic layer took place at 100°C.

Electrochemical impedance spectroscopy (EIS).—Next, the evolution of IL-electrode interface at 100°C was probed by EIS measurements at fixed potential (determined OCP prior to each EIS measurement); the results are reported in Figures 6a and 6b. The initial heating, for all the EIS experiments, was done by dipping the cell in a pre-heated bath (fast heating). The Nyquist curves (Fig. 6a) obtained at different times look almost identical except in the very low frequency region where a continuous shift to higher values of the imaginary part of the impedance is observed (the dashed arrow in Fig. 6a). Interestingly, Nyquist curves deviated at low frequencies from the linear behavior characteristic of the blocking electrode; this is commonly observed in systems where either strong adsorption of ions at the electrode takes place or/and due to the roughness of the electrode surface. Parasitic side reactions could also lead to a curvature of the linear law; however bearing in mind the high purity of ILs, the faradaic processes are unlikely (The IL’s stability window is more than 3 V). This conclusion can be further confirmed by the cole-cole capacitance plots (Fig. 6b) showing clear evidence for two capacitive phenomena. First, the high frequency semi-circle, which does not change with time, is ascribed to double layer charging. In contrast, the second arc at low frequencies (rectangular zone) is evolving with time to lower values of the imaginary part of the capacitance (Fig. 6b inset). This low frequency arc can be ascribed to either the specific adsorption of ions at the electrode surface and/or increased ordering of the EDL as suggested by Merlet et al. 56

Different equivalent circuits, presented in Figure 7, were tested to fit the EIS spectra, however only the last two (Figs. 7d, 7e) gave a good fit to experimental values (SI Fig. 6b). Circuit d), consists of an adsorption branch, a resistor (charge transfer) and a capacitor (double layer charging) in parallel. Circuit d) has already been used to describe the double layer phenomena in ILs by Siinor et al. 31 On the other hand, circuit e) is a combination of two parallel adsorption branches 31 in parallel with a capacitor (double layer charging). This circuit was proposed by Gnahm et al. 31 and Muller et al. 56 for the interface between imidazolium-based ILs and a single crystal Au electrode. Nevertheless, both circuits are empirical and the physical meaning of the circuit elements is not well understood. Thus, we intend to use the fitting process mostly for the visualization of the system trends. In the case of circuit e), the elements R₁ and C₁ were omitted, as they resulted in very high fittings errors (see also Ref. 58). Figure 8 represents the evolution of the high frequency capacitance C₁ and the series capacitance C₂ (low frequency). Both circuits give

**Figure 6.** Impedance spectroscopy a) Nyquist plots and b) complex capacitance curves (cole-cole plots) of 0.12M Co(TFSI)₂ solution at 100°C at E = OCP at different times; c) Nyquist plots and d) complex capacitance curves (cole-cole plots) of pure EMImTFSI at 100°C at E = OCP at different times. 30 min - black solid squares, 1 h - green solid triangles, 2 h - blue turned triangles, 4 h - purple open squares, 6 h - brown circles, 8 h - orange open triangles. The inset in part b) represents zoom of the curves at very low frequencies inside the black dashed square. WE – Pt, CE –Pt; RE –Ag|AgCl. 100°C was reached with fast heating method.

**Figure 7.** The equivalent circuits used to fit the EIS data. a) perfect capacitor, b) Frankin-Melik-Gaikazyan circuit, c) standard circuit describing adsorption phenomena Ref. 57 d) circuit used in Ref. 53 e) circuit used in Ref. 31,58.
very comparable results in terms of magnitude and trends. For the high frequency capacitance $C_1$ only a small decrease was observed with time, which accounts for 8% of the total capacitance. The value of $C_1$ ($9 \, \mu$F cm$^{-2}$) is comparable to the one found in literature $^{31,35,38}$, which is ascribed to the standard double layer charging phenomena. In contrast to $C_1$, the $C_2$ value drastically drops with time (35% change after 8 h). Additionally, the shape of the curve (fast drop at the beginning and stabilization after 6 h) is very similar to the change of the reduction onset shift discussed before (Fig. 4b). Both, the trend and magnitude of drop indicate that the low frequency phenomenon is related to the reduction onset shift at constant temperature. The value of $C_2$ ($60-30 \, \mu$F cm$^{-2}$) corresponds well to the one found by Gnahn et al. $^{31}$ and Szinor et al. $^{53}$, where they assigned the low capacitive contrast to $C_2$.

Lastly, it is worth mentioning that we experienced a similar feature of the EIS spectra in pure IL solution (Figs. 6c, 6d), meaning that the changes of the system at 100 °C with time is an internal property of IL itself and not of the [Co(TFSI)$_2$]$^-$ complex.

Effect of the interface evolution on the chronoamperometric Co deposition process. — After establishing the scientific platform of electrochemical behaviors in IL at high temperature, we report here a practical example of system evolution at 100 °C influence on the deposition process and deposit quality. The deposition was done at fixed potential of −0.9 V using EQCM. The limiting parameter was the quartz crystal frequency change of 40 kHz. Two experiments were performed, the first after 30 min (Figs. 9a, 9b, 9c) and the second after 8 h (Figs. 9d, 9e, 9f) of heating at 100 °C. In both cases, the initial heating was done by dipping the cell in pre-heated bath (fast heating).

In case of the experiment recorded after 30 min, a classical chronoamperogram of a plating process was observed (Fig. 9a, black curve). The corresponding changes in resistance (drop) and frequency (drop) indicate consumption of Co$^{2+}$ species and formation of deposit over the electrode surface (Figs. 9a, 9b). The calculated equivalent mass - M / Hz value, from the slope of frequency versus charge, was 27.5 g - mol$^{-1}$ (SI Fig. 7a’) that is close to the theoretical value of a two electron Co$^{2+}$ reduction process (29.5 g - mol$^{-1}$). This indicates the high efficiency of the cobalt plating process. The obtained Co thin films were visually uniform, shiny, and well attached. They were made of ~50 nm grains; additionally, random outgrowths of ~1 μm spherical agglomerates were spotted on the top of the deposit (see insert in Figure 9c). The deposit was composed of Co with traces of F, S and O coming from residual IL, as studied by energy dispersive spectroscopy (EDS).

The chronoamperometric deposition process after 8 hours (Figs. 9d, 9e, 9f) is more complex, as conveyed by the presence of the growth domains (I, II, III) separated by blue dashed lines on the curves. In the first region (up to 400 s), very low current densities together with tiny changes in frequency and motional resistance were observed, indicating high hindrance of the Co$^{2+}$ reduction process. According to the CV (Fig. 4a), the reduction of Co$^{2+}$ should not take place ($E_{\text{red}} < −0.9 \, \text{V after 8 h of heating}$); however the chronoamperometric experiment was done by direct switching of potential from OCP to −0.9 V. This method is very different from the one used for CV experiments (slow decrease in potential). This change in the conditions might lead to a different state of the interface as it was briefly addressed in the recent paper by Carrie and others. $^{59}$ Nevertheless, the reduction of Co$^{2+}$ was greatly hindered indicating the presence of a very strong barrier. Moving toward the end of region I (300–400 s), a significant increase in current density was observed. The increased reduction kinetics with time could be due to perforation of the blocking layer by slow deposition of Co particles. Despite the higher current rates, the frequency changes remained small, meaning that most of the produced Co was poorly adhesive. This is in agreement with our visual observation as we observed black particles floating in the bath. Reaching the second region (400–500 s), a continuous increase in current density accompanied by a progressive decrease in quartz frequency was observed. This indicates an enhancement of the adhesive plating process. For times greater than 500 s (region III), the drop in frequency was significantly increased, with the stabilized current density being nearly the same as for the short heating deposition process. The continuous increase in motional resistance in region III (Fig. 9e) is most likely due to the rough morphology of the deposit as deduced from scanning electron
microscopy (SEM) images (Fig. 9f). The obtained deposit was well attached to the electrode but not uniform and not shiny compared to the previous one. It was composed of a very thin compact layer with large ball-like agglomerates on the surface (Fig. 9f). The EDS elemental analysis showed that deposit was made of Co with traces of Fe, S and O coming from residual IL. Lastly, the calculated M/z value in region III was equal to 30.8 g·mol⁻¹, (SI, Fig. 7b). Despite high increase of resistance (that may potentially affect the frequency change), the obtained M/z value is close to the theoretical one for the Co plating reaction (M/z = 29.5 g·mol⁻¹). For regions I and II, the M/z values lower than 10 g·mol⁻¹ were solely found due to deposit detachments.

**Expanding observations to other systems.—** At this stage, we have unambiguously established the existence of a temperature-driven evolution of Co²⁺ reduction process that has a tremendous impact on the properties of the deposited cobalt thin films. Thus, the question remains whether these phenomena are specific to the nature of the cation, IL formulation, and substrate used in the above experiments. For all the experiments in this part fast heating process was done.

Firstly, the IL formulation has been investigated by changing either the cation to BMP²⁺ or the anion to TfO⁻ (Figs. 10a and 10b, respectively). The CV curves revealed similar features in both cases, with 2 reduction waves separated by a shallow drop in current density, which is less pronounced for the BMP²⁺ based IL. Moreover, the shift in reduction onsets and evolution of the shape of the curve at 100°C (fast heated cell) is similar to the one observed for EMImTFSI.

Turning to the importance of the electroactive species, we investigated three different cations (Zn²⁺, Ni²⁺, Mn²⁺) together with molecular oxygen (Figs. 11a, 11b, 11c and 11d, respectively). All baths containing metal cations show shifts of the reduction potential onsets and change of the curve shape. However, the magnitude of the shift varied depending on the cation and is the largest for Ni²⁺ as compared to both Mn²⁺ and Zn²⁺ (SI, Fig. 8a). In contrast, measurements of the oxygen reduction revealed no change of the current profile shape and only small shift (<30 mV) of the reduction potential onset. The drastically different behavior of molecular oxygen reduction, as compared to metal cations might be related to the nature of the O₂ molecule. We previously found⁵⁰ that the oxygen reduction process is overruling metal plating when both O₂ and metal cations are present in the electrochemical bath even if the later one starts to reduce at higher potentials. Such behavior was attributed to the small size and neutrality of oxygen molecules, which can easily penetrate through the blocking layers at the interface, in contrast to large metal cation complexes.

Finally, the nature of the substrate was investigated using a stainless steel wire as the working electrode (Fig. 12a). The shift in reduction onset and the curve shape change were observed with however a milder evolution as compared to a Pt substrate, especially at the beginning of heating process (Fig. 12b).

The above experiments clearly show that the temperature driven changes in electrochemical reaction are not specific to Co²⁺ reduction in EMImTFSI. The observed phenomena can be extended to other...
systems considering change in IL, electroactive species and electrode nature.

**Reversing the evolution of the system.**—The hindrance of the reduction signal at 100 °C is rather undesirable since the reason of increasing the temperature is to improve the reduction process rate. Nevertheless, we found that an enhancement in the deposition rate, after long time conditioning at 100 °C, can be partially restored by applying an oxidative pretreatment. Fig. 13 shows the CV curve after 8 h of heating initiated in anodic direction (the black curve). After the anodic sweep, the reduction profile has changed greatly; the first reduction wave followed by a drop in current density appear again on the CV curve in contrast to untreated electrode (green dashed curve). Thus, the system state was partially reset with the CV response corresponding to 2–3 h of heating according to Figures 4a and 4b. Interestingly, the shape of the CV and reduction onset potential (~0.95 V) are very similar to the one found for slow heating method, Figure 2 brown dashed line. This indicates, that the oxidative pretreatment suppress the system changes caused only by high temperature T = 100 °C (disappearance of the first reduction peak) rather than the one coming from the high temperature ramp (shift of the reduction onset).

From chronocoulometric and CV experiments carried out at various oxidative potentials (SI Fig. 9^a and 10^b), we could deduce that the process responsible for the partial system reset is already triggered at 1.3 V. This potential corresponds to the onset of the small oxidation wave (green arrow, inset Fig. 13). The same oxidation signal was observed in pure IL at both RT and 100 °C (Fig. 1 – O_2) independently on the heating process.

To characterize the nature of the oxidation process the EQCM measurement was done in 0.12 M Co(TFSI)_2 solution by means of CV initiating in anodic direction. The collected data including first CV curve and the corresponding changes in frequency and resistance, are shown in Figure 14. For the sake of clarity and conciseness, the characteristic color domains, which are repeated in the three plots, were used. During the oxidation process ranging from 1 to +1.5 V (blue/violet domain Fig. 14a), we observed a decrease in frequency of ~50 Hz (Fig. 14b) which corresponds to a constant motional resistance (Fig. 14c). This implies that the frequency change can be attributed to a mass increase at the electrode surface of about 270 ng/cm^2. Such small mass change could be due to ion exchange phenomena as studied by Tsai et al. ^4b We further analyzed the oxidation signal by performing a second subsequent CV scan. Interestingly, the oxidation signal did not appear (SI Fig. 11^a) despite the very slow scan rate (2 mV/s) that allows for diffusion of the species toward the electrode. This suggests that the nature of the signal is not due to some electroactive species in the bulk, as those would give at least a small signal on the subsequent scan. Consequently, the second scan can be treated as a background that can be subtracted from the first CV scan; as a
result (Fig. 14a - inset) a well-shaped symmetrical oxidative peak is obtained confirming the lack of diffusion controlled phenomena.

Additional CV experiments with extended reduction reverse potential were conducted on a Pt electrode (Figs. 15a, 15b). Once the reduction sweep is reversed at ~0.5 V, the oxidation signal appears again on the subsequent oxidation sweep, with however slightly shifted potential to more negative values. The existence of a corresponding reduction process indicates the reversible nature of the phenomenon. The above characteristics suggest a surface phenomenon without a diffusion-limiting step to be at the origin of the oxidative process at 1.5 V.

For the sake of completeness, similar EQCM measurement was carried-out on pure IL. The CV result, after 8 h of heating (SI, Figure 12*), has shown a small increase in weight during the oxidation at +1.5 V, which was equal to the weight decrease during the corresponding reduction sweep. This indicates again high reversibility of this phenomenon independently of the presence of Co\(^{2+}\) ions. Surprisingly in pure IL the corresponding reduction process takes place at much higher potentials that could be due to differences in the EDL structure when Co\(^{2+}\) is not present in the solution.

Similarly to the reset of the CV curve shape by oxidative pretreatment we should expect same changes on EIS spectra. Indeed, the evolution of EIS spectra is also reset by performing an oxidative pretreatment (SI, Fig. 13*). The imaginary part of impedance shifts to higher values at low frequency after oxidative pretreatment, while only slight change was observed for the high frequency process. This confirms the correlation of slow frequency phenomena with the evolution of the system at constant temperature.

**Discussion**

Through our study, we have reported that the reduction process of metal cations in IL media at 100 °C is highly dependent on the heating method and is changing with time at constant high temperature. We have shown that improved kinetics of Co\(^{2+}\) reduction can be assessed when the cell is exposed to high temperature gradients. However, during the long conditioning at constant temperature a shift in the reduction onset toward lower potentials and significant changes in the CV curve shape were observed, leading to a hindrance of the reduction process. This indicates formation of a blocking layer at the electrode surface. Our EQCM results have shown no change in the electrode mass with time at T = 100 °C. This indicates that the phenomena responsible for the system evolution are rooted in the existence of physical interactions taking place at the electrode-IL interface rather than the formation of an extra organic/inorganic layer. This was further confirmed by the EIS results that have shown the presence of a slow capacitive phenomenon due to specific ion adsorption to be at the origin of the system evolution. It suggest that the temperature-driven shift of the reduction process is of a physical nature, but the remaining burning question regards the mechanism of this phenomenon.

It has been previously reported that temperature gradients can disturb the ordered structure of the EDL in aqueous solutions followed by reorganization when the temperature gradient is absent. There is no reason why such a situation would not hold for ILs. Figure 16 represents a schematic of possible mechanism of the EDL changes at elevated temperature at negatively charged surface. At RT, when increasing the electrode’s negative charge, the EDL tends to strongly order forming ion layers with the IL’s cations facing directly the electrode surface.\(^{18–22}\) It is worth noting that even at OCP a layered ordering is present.\(^{62}\) During fast heating to 100 °C, the system is exposed to high temperature gradients with a concomitant disturbance of the EDL structure present at RT. This is in agreement with the time/temperature driven variation of OCP value (SI, Fig. 2*). We believe that applying negative polarization just after heating leads to lower ordering (Fig. 16I) as compare to RT (Fig. 16d). Such a disturbance leads to easier accessibility of the electrode surface, thus a much smaller overpotential is needed for reduction of the large \([\text{Co(TFSI)}]^-\) complexes. This is in accordance with our CV experiments performed with different heating rates. For small gradients of temperature, the EDL disturbance is not expected and consequently it will result in smaller changes of reduction onset, as shown in Figure 2.

At constant temperature T = 100 °C, the EDL tends to reorganize with time (formation of ordered structures, Fig. 16II), as observed in the case of aqueous solutions by Garcia-Araez et al.\(^{46,47,61}\) However, the time of this process in our case might be significantly longer owing to larger thermal movements of the ions at elevated temperature, as well as a complex structure of the ions.\(^{18,33,34}\) The increased ordering

---

**Figure 15.** Successive CV curves of 0.12 M Co(TFSI)\(_2\) solution in EMImTFSI at 100 °C a) 1\(^{st}\) and 2\(^{nd}\) scans b) 3\(^{rd}\) and 4\(^{th}\) scans, showing the corresponding reduction process. WE – Pt, CE – Pt, RE – Ag/Ag\(_2\)O, scan rate 5 mV·s\(^{-1}\). Successive scans were performed on the same electrode.

**Figure 16.** Schematic representation of EDL structure at negatively charged electrode at different steps: RT state, after fast heating, long term conditioning at 100 °C. We stress that this schematic is just a visualization of our hypothesis.
leads to greater overpotentials for the Co^{2+} reduction process and the evolution of the CV curve shape to a state somewhat similar to the one found at RT, where the EDL is highly ordered. Nevertheless, some differences in EDL are present as suggested from the observed CV at RT (SI Fig. 14+) of heat-treated and pristine electrodes, with the former one being more blocking. The exact nature of the interfacial rearrangement is not well understood; an increase in temperature could enable additional stable conformations of ions or induce strong phys/chemisorption.\(^{13}\) Certainly further in-situ studies are needed to gain deeper molecular inside into the interfacial processes. In partic-
ular, AFM and STM techniques on single crystal electrodes would be crucial to access this information.

Going deeper into the +1.5 V oxidation process, we believe that applying positive polarization triggers a major reconstruction of the EDL with the cations being repelled from the electrode surface, while the anions are attracted (Fig. 16V). This exchange of ions could explain both i) the gain in mass observed by the EQCM experiment during oxidation at +1.5 V and ii) the loss in mass during the corresponding reduction process, as the TFSI\(^-\) ions have almost three times higher molar mass than EMIm\(^+\) ions. On the successive reduc-
tion sweep, at ~0.5 V, the reverse reconstruction takes place, namely the anions are repelled from the surface while the cations are attracted, thus forming an ordered EDL with adjacent layer reach in-
cations (Fig. 16V). Nevertheless, the properties of this EDL (V) are different as compared to the one obtained during long term isothermal heating (III). The new ordered EDL is less blocking leading to en-
hanced kinetics of Co^{2+} reduction (as was shown on Figure 13). This suggestion is further supported by the shift of the oxidation signal at 1.5 V to more negative potentials during successive CV cycling (Fig. 15). During the first oxidation scan (after 8 h of heating), the EDL reconstruction might be harder to take place due to the spe-
cific adsorption of ions; thus a higher oxidative potentials have to be applied to trigger the reconstruction. Moreover, the electrochem-
ically induced reconstruction does not lead to a disordered EDL as the one found after applying high temperature gradient (II). Therefore, no significant shift of the reduction onset will take place. Summing up this part, the oxidative pretreatment resets the specific adsorption phenomena responsible for the hindrance of the first reduction peak, but it does not make disordered EDL structure as the one from high temperature gradient. This reasoning is in agreement with our CV curves; the CV obtained after oxidative pretreatment was similar (Fig-
ure 13) to the one recorder after 15 min of heating for the slow initial heating method (small temperature gradient, Figure 2 brown dashed curve).

With respect to the electrochemical growth of materials as thin films, a strong evolution of the EDL layers during the heating process can lead to non-reproducible data if monitoring of the time domain is omitted. However, mastering of temperature effects can lead to the formation of high quality (uniform, compact, adherent) metal coatings together with an enhancement of the deposition rates. This is a serious asset when using ILs that usually show sluggish kinetics. We believe that our results could be of great help in solving today’s challenges encountered when depositing metals from ILs. Disturbing interfacial structures with the methods shown in this report can improve these processes.

**Conclusions**

We have reported the influence of fast and slow heating process and long term conditioning at high temperatures on the reduction of Co^{2+} cations in ionic liquid media, based on the electrochemical quartz crystal microgravimetry, cyclic voltammetry, chronoumperom-
etry, impedance spectroscopy and scanning electron microscopy ex-
periments. We observed improved kinetics of Co^{2+} reduction when the system is exposed to a high temperature gradient. However, at 100 °C, the Co^{2+} reduction kinetics slowed down with time. The reduction onset potential shifted to more negative values while the intensity of the first reduction wave greatly diminished. The EQCM and EIS measurements suggest physical interactions of ions and the electrode surface cause these phenomena. We deduced that the system evolu-
tion is related to the ordering process of the EDL that was disturbed by applying a temperature gradient during heating from RT to 100 °C (dynamic state of the EDL). This process is not specific to the reduc-
tion of Co^{2+} in EMImTFSI since we have showed herein that it also occurred in BMImTFSI and BMImTIO and for different cations (Zn\(^{2+}\), Ni\(^{2+}\), Mn\(^{2+}\)). Finally, the paper shows the importance of controlling the heating process and its influence on the deposit quality. We found that the de-
position of cobalt in non-equilibrated systems leads to better quality deposits (uniformity, morphology, adhesion). It also gave an ad-
nitional parameter for controlling the deposit morphology (thin compact layer and 3D agglomerates made of ball-like grains were synthesized). We hope that our findings will help in mastering the deposition of both metal and oxide thin films from ILs at elevated temperature.

**Acknowledgment**

The authors thank to ALISTORE European Research Institute for the financial support to conduct the research.

**References**

4. N. Madria, T. A. Arunkumar, N. G. Nair, A. Vadugali, Y.-W. Huang, S. C. Jones, and V. P. Reddy, “Ionic liquid electrolytes for lithium batteries: Synthesis, electro-
19. T. Carstens, R. Hayes, S. Zein El Abedin, B. C. G. Webber, N. Borisenko, R. Atkin, and F. Endres, “In situ STM, AFM and DTS study of the interface 1-hexyl-3-
methylimidazolium tris(pentafluoroethyl)trifluorophosphate/Au(111) interface: potential dependent sol-


