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Voltammetric behaviour of Cationic Redox Probes at Mesoporous Silica Film Electrodes

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Abstract

The effect of vertically aligned silica nanochannels on the electrochemical response of various cationic redox probes has been investigated. To this end, an oriented mesoporous silica thin film has been generated by electro-assisted self-assembly onto a fluoride-doped tin oxide (FTO) electrode, which was then used to characterize the voltammetric behaviour of redox cations of different nature, size and charge. Methylene Blue (MB⁺), paraquat (PQ²⁺), diquat (DQ²⁺), ruthenium trisbipyridine (Ru(bpy)₃²⁺) and ruthenium hexamine (Ru(NH₃)₆³⁺) were selected for that purpose. Multisweep cyclic voltammetry experiments showed significant accumulation of these cations thanks to favourable electrostatic interactions with the negatively-charge silica surface, but the enhancement of the votammetric signals compared to the bare FTO electrode was strongly dependent on the probe type, according the following trend: MB⁺ > PQ²⁺ > DQ²⁺ > Ru(bpy)₃²⁺ > Ru(NH₃)₆³⁺. The accumulation processes were very fast owing to the vertical orientation of the mesopore channels ensuring fast diffusion from the solution to the electrode surface. Operating at various potential scan rates and different probe concentrations enabled to quantify the relative ratio between surface-confined and diffusion-controlled processes, the adsorption phenomena being clearly more prominent in diluted media and also accentuated at high potential scan rates. The mesoporous silica film electrodes exhibited higher sensitivities at lower concentrations, which can give benefits for sensor applications at trace levels.

Keywords

Mesoporous silica film; modified electrode; redox cations; permeability; diffusion; accumulation.

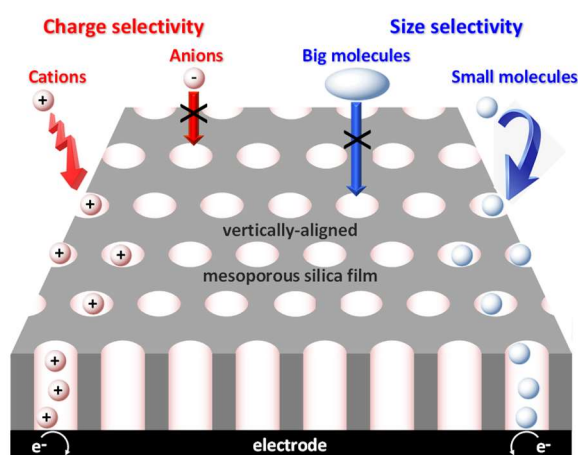
1. Introduction

The transport of ions and molecules in nanochannels and through nanopores has been extensively studied [1–6], notably because nanoporous membranes offer attractive features for applications in separation [7–12], sensors [13–15], catalysis [16,17], or energy storage [17]. Porous membranes made of nanochannels and nanopores are mostly prepared in the form of materials such as anodic aluminum oxide (AAO) [18–20], carbon nanotube (CNT) assemblies [21], polyethylene terephthalate (PET) nanochannels [22], self-organized nanotubes [23], zeolite nanoporous membranes [24–26], or boron nitride nanotube [27]. The nanochannels prepared from such materials are usually characterized by pore diameters above 10 nm, which is basically useful for confinement of large molecules and species such as metal, metal oxide, polymers, and ionic liquids [5]. These are often subject to wettability issues, their mechanical and chemical stability needs to be improved, their pore size and density cannot always be accurately controlled, and vertical alignment of nanochannels may need complex processing [28]. Vertically-aligned nanoporous silica membranes made of a high density of hexagonally-packed mesopore channels with small and monodisperse diameters (in the ~2-3 nm range) [18,29–31] can offer advantageous features, notably for the confinement and transport of small molecules and ions.

A versatile and fast way of preparing vertically-ordered silica nanochannels onto electrode surfaces is the use of the electrochemically assisted self-assembly (EASA) method, as developed in our group some time ago [29,30]. This method is likely to generate uniform thin films (with thicknesses ranging typically between 50 and 150 nm [30], or even more in multilayer configurations [31]) with pore sizes adjustable between 2 and 3 nm [32], and the mesochannels can be further functionalized with a variety of organic moieties [33–35]. Other synthetic approaches include the Stöber-solution growth [36], magnetically-induced orientation [37], epitaxial growth [38], or via an oil-induced co-assembly process [39]. Such oriented nanochannel membranes are very promising for applications, especially in the fields of electrochemical sensing and separation at molecular level [40–43]. Their performance can be affected by various factors such as the mesopore size [32,44], the mesochannel length/thickness [45,46], the solution pH and/or ionic strength [47,48] and the surface charge of the pores [43,49]. They are also attractive hosts for inclusion of redox mediator species or adsorption/confinement of other catalysts, for applications in electrocatalytic or electrochemiluminescence sensing, for instance [50–56].

Vertically ordered silica mesochannel films exhibit molecular sieving properties and permselective ion transport (Scheme 1). The size selectivity of mesoporous silica membranes has been demonstrated using molecular probes with dimensions greater than the pore diameter [28,32,57]. For example, on the basis of silica membranes with distinct pore sizes (i.e., 2.0 and 2.9 nm), it has been shown that a ferrocenyl dendrimer (critical size of about 2.4 nm) can only be detected at the underlying electrode surface by diffusion through the larger nanochannels (2.9 nm) while being totally excluded from the film made of narrower pores (2.0 nm) [32]. Cationic fluorophores of similar geometries but different sizes have been used to evidence possible permeation for the smaller ones (i.e., ruthenium trisbipyridine, $\text{Ru}(\text{bpy})_3^{2+}$, and ruthenium trisphenanthroline, $\text{Ru}(\text{phen})_3^{2+}$, with molecular size of 1.3 nm), whereas bigger species (ruthenium trisdiphenylphenanthroline, $\text{Ru}(\text{dpp})_3^{2+}$, 2.0 nm in diameter) were excluded from the mesoporous silica membrane [28]. The size exclusion of small proteins (such as hemoglobin) from silica mesochannels was demonstrated and applied as protective barrier against biofouling [57,58]. Note that similar molecular sieving effects have been also reported for non-silica mesoporous membranes (permeation of methylene blue, MB^+ (diameter, 1.4nm), and rejection of rhodamine 6G, R6G^+ , (diameter, 1.8nm), at vertically aligned nanopores in a polymer membrane having a pore size of 1.2-1.4 nm) [9]. In addition, the vertically-ordered silica nanochannels are otherwise characterized by charge selectivity issues. At pH greater than 3, the silica surface is expected to be negatively charged owing to its isoelectric point of 2-3 [59,60]. As a consequence, when working in usual media (at moderately acidic or neutral pH values), the mesoporous silica films on electrodes acted as a permselective membrane, enabling an easy transport of positively-charged redox probes (such as ruthenium hexamine, $\text{Ru}(\text{NH}_3)_6^{3+}$, for instance) while rejecting the anionic ones (as ferricyanide, $\text{Fe}(\text{CN})_6^{3-}$), resulting in large electrochemical signals for cations (sometimes larger than at bare electrode due to some accumulation) and low currents for anions (as a result of electrostatic repulsions preventing the negatively-charged redox probes to reach the underlying electrode surface) [44,61]. Such charge selectivity was similarly observed recently with polymer-based nanochannel membranes, exhibiting, e.g. selective permeation of methylene blue, MB^+ , over methyl orange, MO^- [9]. By contrast, small neutral molecules (such as ferrocene dimethanol or catechol) can freely diffuse through the mesoporous silica membranes and be detected onto the underlying electrode surface [47,62], yet with some influence of the mesopore diameter on the transport rates [63]. Attempts to modulate the permselective behaviour, especially to promote the detection of anionic redox species, have been reported in the literature, by means of turning the internal surface of nanochannels from negative to positive (by working in highly acidic media

[48], or by grafting positively-charged ammonium groups to the silica surface [61,64]), by operating in very high supporting electrolyte concentrations [65], or by using neutral redox mediators acting as electron shuttles for carrying charges from through the mesoporous film [62]. The transport of charged species through such regular and narrow channels bearing charges onto their internal surface was found to be highly depending on the ionic strength (defining the thickness of the electrical double layer at the silica surface as an interplay with the mesopore size) [47,48,61]. Hydrophobic effects on mass transport processes have been also reported [66,67]. To date, however, no extensive studies on the factors affecting the accumulation and electrochemical response of redox cations at oriented mesoporous silica film electrode, depending on their characteristics (nature, charge, size), have been made.



Scheme 1: Illustration of charge and size selectivity at vertically aligned mesoporous silica film. The film thickness is generally 80 -150 nm with 2-3 nm pore diameters.

Here, we have thus investigated the voltammetric behaviour of various positively-charged redox species at an fluorine-doped tin oxide (FTO) electrode covered with an oriented mesoporous silica film generated by EASA. The probes have been chosen for their different nature (organic or inorganic), charge (mono-, di- or tri-cationic) and size (yet, all of them being smaller than the nanochannels diameter to avoid any size-exclusion effects). These include paraquat (PQ^{2+} , also called methylviologen), diquat (DQ^{2+}), methylene blue (MB^+), ruthenium hexamine ($Ru(NH_3)_6^{3+}$) and ruthenium tris-bipyridine ($Ru(bpy)_3^{2+}$). Their confinement in the silica nanochannels was mainly monitored by multisweep cyclic voltammetry, in comparison to the bare FTO electrode and at various probe concentrations, as well as after open-circuit

accumulation and subsequent voltammetric analysis. The extent of accumulation was strongly dependent on the probe type, as discussed in this paper.

2. Experimental section

2.1. Chemicals and Reagents

Tetraethoxysilane (TEOS, 98%, Alfa Aesar), cetyltrimethylammonium bromide (C₁₆TAB, 99%, Acros), ethanol (95–96%, Merck), and HCl (1.0 M, Sigma–Aldrich) were used for the electrochemically assisted self-assembly (EASA) of mesoporous silica films on fluorine-doped–tin oxide (FTO) plates (surface resistivity 10–14 Ω cm⁻², Delta Technologies). Potassium ferrocyanide, (K₄Fe(CN)₆, 99.5%, Fluka), Ferrocenemethanol (C₁₁H₁₂FeO, 97%, Alfa Aesar) and hexaammineruthenium(III) chloride (Ru(NH₃)₆Cl₃, 98%, Sigma-Aldrich), Paraquat (1-1'-dimethyl-4-4'-bipyridinium dichloride, 98%, Sigma-Aldrich), Diquat (C₁₂H₁₂Br₂N₂, H₂O, 99%, Riedel-de Haën), Methylene Blue (C₁₆H₁₈ClN₃S xH₂O, Fluka), Ruthenium Trisbipyridine or Tris(2,2'-bipyridyl)dichlororuthenium(II) hexahydrate (C₃₀H₂₄Cl₂N₆Ru·H₂O, 98%, Acros) were used as redox probes in the cyclic voltammetry experiments. Sodium nitrate (NaNO₃, 99%, Fluka) was used as the supporting electrolyte.

2.2. Electrochemical Measurements

Cyclic voltammetry measurements were carried out using either an EMStat2 or a PalmSens potentiostat (PalmSens, Netherlands) in the absence of oxygen (achieved through N₂ purging for 20 min). A three- electrode system using an Ag/AgCl (3M KCl) reference electrode (Metrohm) and a stainless steel rod as the counter electrode, and FTO plates as working electrode, modified with mesoporous silica thin films. The background electrolyte for the cyclic voltammetry experiments was at 70 mM NaNO₃, as it was optimized in a previous study [47].

2.3. Preparation of mesoporous silica thin films

The oriented mesoporous silica thin films were deposited electrochemically into FTO electrodes according to a previously reported procedure [29]. A starting sol solution containing 100 mM TEOS as the silica precursor, 32 mM surfactant as template, and NaNO₃ (0.1 M) as supporting electrolyte was prepared in a water: ethanol (1:1) solution adjusted to a pH of 3 using 0.1 M HCl solution. The solution was hydrolysed for 2.5 h and used for electrochemical deposition of silica by applying -1.5 V for 25 s in FTO as a working electrode with a diameter of 0.8 cm, the reference electrode was a silver wire, and a stainless steel plate as a counter

electrode. The modified FTO electrode surface was rinsed immediately with water and dried overnight at 130°C. The template was removed by solvent extraction, by immersion of the film in 0.1 M HCl / Ethanol solution for a minimum of 15 min. Mesoporous silica films with and without surfactants were further characterized by GISAXS, TEM and electrochemical techniques. The full characterization is described in the supplementary information (Figure S1 and associated text).

Grazing-incidence small-angle scattering (GISAXS) experiments were carried out on "SAXSess mc²" instrument (Anton Paar) with GISAXS equipped "VarioStageXY". This instrument is attached to a ID 3003 laboratory X-ray generator (General Electric) equipped with a sealed X-ray tube (PANalytical, λ Cu, $K\alpha = 0.1542$ nm) operating at 40 kV and 50 mA. Scattering of X-ray beam was registered by a CCD detector (Princeton Instruments, 2084 \times 2084 pixels array with 24 \times 24 μm^2 pixel size) at 299 mm distance from the sample. Using Winspec32 software (Princeton Instruments), the 2D image was integrated into one-dimensional scattering intensities $I(q)$ as a function of the magnitude of the scattering vector $qz = (4\pi/\lambda) \sin(\theta)$, θ is the specular reflectance angle. The level of mesostructural order was evaluated by transmission electron microscopy (TEM) using a PhilipsCM20 microscope at an acceleration voltage of 200 kV. The samples were prepared by removing some pieces of the films, which were supported on a carbon-coated copper grid.

3. Results and Discussion

3.1. Accumulation of different cationic redox probes inside the mesochannels

The effect of oriented mesoporous silica thin film on FTO electrode was investigated by cyclic voltammetry in order to point out possible differences in the accumulation of cationic redox probes with different positive charges (ranging from +1 to +3), sizes, and molecular structure. The redox probes used in this study were paraquat (PQ^{2+}), diquat (DQ^{2+}), methylene blue (MB^+), ruthenium trisbipyridine ($\text{Ru}(\text{bpy})_3^{2+}$), and ruthenium hexamine ($\text{Ru}(\text{NH}_3)_6^{3+}$). Cyclic voltammetry of the various probes was run 50 times to allow reaching steady-state current values (Figure S2). Figure 1 showed the 50th cyclic voltammograms obtained at both bare (dashed curves) and mesoporous silica thin film electrodes (solid lines) for the five redox probes. Since the silica surface has an isoelectric point of 2 [44,68–70], silanolate groups are present on the silica surfaces in pH conditions used here (unbuffered solutions with pH values around 6.5), leading to favourable electrostatic interactions with the cationic redox probes. The

absence of an adsorption prepeak for all redox probes (Figure S2) suggests a weak type of adsorption on mesoporous silica [71], with similar formal potentials for the adsorbed and the non-adsorbed redox couples [72]. For all redox probes investigated, the electrochemical signals were enhanced with mesoporous silica compared to the bare electrode (Figure 1), suggesting that all cationic probes were accumulated in the mesochannels of the silica films. However, the extent of the accumulation varied significantly from one probe to another. The highest current response was observed for MB⁺ with a 13-fold increase of the peak current after accumulation whereas the signal for Ru(NH₃)₆³⁺ was improved only by a factor 1.4. The order for the enhanced electrochemical signal of the cationic redox probes is the following: MB⁺ > PQ²⁺ > DQ²⁺, Ru(bpy)₃²⁺ > Ru(NH₃)₆³⁺. It seems therefore that multicationic species are less efficiently accumulated than the monocation, which can be due to the fact that they required more counter-anions from the electrolyte to maintain charge balance [47] (anion transport is restricted in such silica channels [48]). At same charge, the size of the probe might have some effect (i.e., more effective accumulation of PQ²⁺ having a kinetic diameter of 0.58 nm [73] than DQ²⁺ with kinetic diameter of about 0.7 nm [74]) but this is expected to be quite limited if considering that the bulkier cation (Ru(bpy)₃²⁺, characterized by a size of 1.2-1.3 nm [75]) led to enhancement values close to that of DQ²⁺ (compare green and orange curves in Fig. 1).

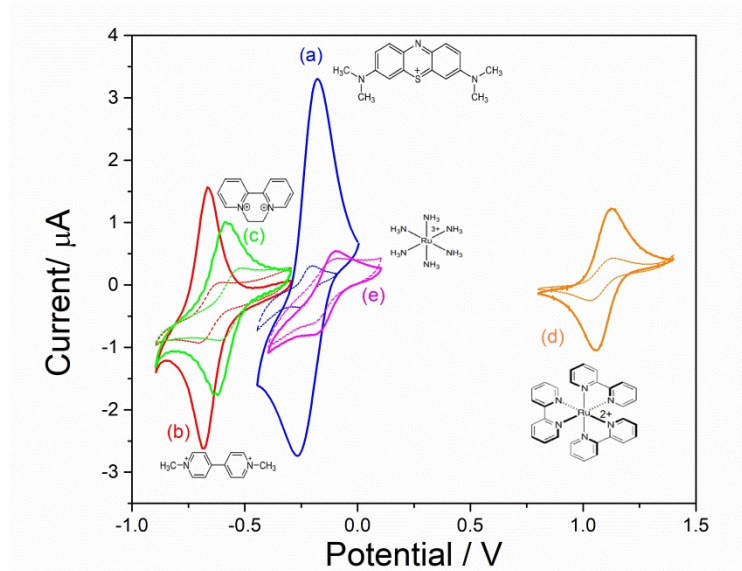


Figure 1: Cyclic voltammograms of various redox probes recorded at bare FTO (dashed line) and at mesoporous silica film modified FTO (straight line) electrodes: (a) MB⁺; (b) PQ²⁺; (c) DQ²⁺; (d) Ru(bpy)₃²⁺; (e) Ru(NH₃)₆³⁺; the curves correspond to the 50th cycle obtained after multiple potential scanning at 20 mV s⁻¹; The concentration for all probes was 20 µM, in a supporting electrolyte solution made of 70 mM NaNO₃.

Figure 2 showed the evolution of the absolute peak current value of the forward scan as a function of the number of cycles for both the silica film electrode (part A) and the bare FTO electrode (part B). Peak currents at the bare electrode were constant (independent on the number of cycles) for the five redox probes, and much lower than those recorded for the film electrode, confirming that the accumulation behavior is due to the presence of mesoporous silica thin films. For MB^+ , PQ^{2+} , DQ^{2+} and $\text{Ru}(\text{bpy})_3^{2+}$, the forward peak current rapidly increased in the first few cycles (10-20 cycles), before reaching a plateau. Even, the first cycle for each probe led to current values much larger than at bare FTO, with magnitudes of more than 50% of the maximal steady-state values, indicating quite fast accumulation processes (thanks to the vertical channels). The increase in current in the initial stage is due to transport-limited by diffusion accompanied by rapid adsorption onto the silica substrate [76–78]. Once the full coverage of the silica surface is attained, redox probes will no longer adsorb as shown by the plateau in the curve. For $\text{Ru}(\text{NH}_3)_6^{3+}$, the peak current remained more or less constant throughout the series of cycles, suggesting that its adsorption is limited, despite a charge of +3. Actually, a main difference between this probe and the four above ones is the absence of aromatic rings; accordingly, it seems that the presence of aromatic moieties in the cationic probes contributed to improve their binding to the silica surface, which might be due to non-covalent forces between electron deficient aromatic systems of the redox cations and silanolate anions (i.e., π -anion interactions [79]), in addition to the more classical electrostatic interactions.

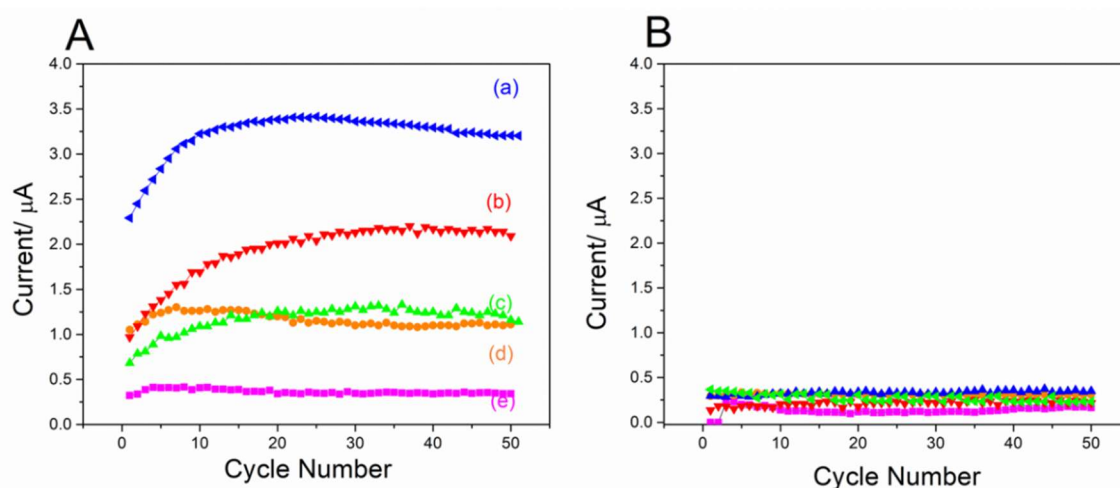


Figure 2: Forward peak current as a function of the number of cycles for the various cationic redox probes at (A) a mesoporous silica film and (B) a bare FTO electrode for MB^+ (a, blue); PQ^{2+} (b, red); DQ^{2+} (c, green); $\text{Ru}(\text{bpy})_3^{2+}$ (d, orange); and $\text{Ru}(\text{NH}_3)_6^{3+}$ (e, magenta). All the experimental conditions were the same as Figure S2.

3.2. Thin-layer versus semi-infinite diffusion

In order to investigate further the behaviour of various redox probes in mesoporous silica thin films, cyclic voltammetry studies were conducted at two different concentrations (20 and 1000 μM) after accumulation at open-circuit potential for 30 min. Control experiments showed that the peak current increased with accumulation time, before reaching a plateau (Figure S3). Figure 3 shows the cyclic voltammograms recorded for DQ^{2+} at the two concentrations for scan rates varying from 10 to 100 mV s^{-1} . At the lowest concentration (Fig. 3A), the peak-to-peak separation was approximately 0.03 V, indicating an electrochemical response mainly controlled by the redox molecules adsorbed on the silica and corresponding to thin layer diffusion. When the concentration raised to 1000 μM (Fig. 3B), the peak-to-peak separation increased to 0.09 V (at 10 mV s^{-1}) suggesting a semi-infinite diffusion-controlled reaction. These observations were further supported by the scan rate studies. When the redox probe concentration was 20 μM , the peak current varied linearly with the scan rate (Fig. 3A, inset), confirming that the electrochemical response is dominated by adsorbed species. On the other hand, diffusion of redox probes is dominating the electrochemical response at the highest concentration as established by the linear variation of the peak current with the square root of the scan rate (Fig. 3B, inset). Similar behaviour was observed for the other redox probes (Figure S4).

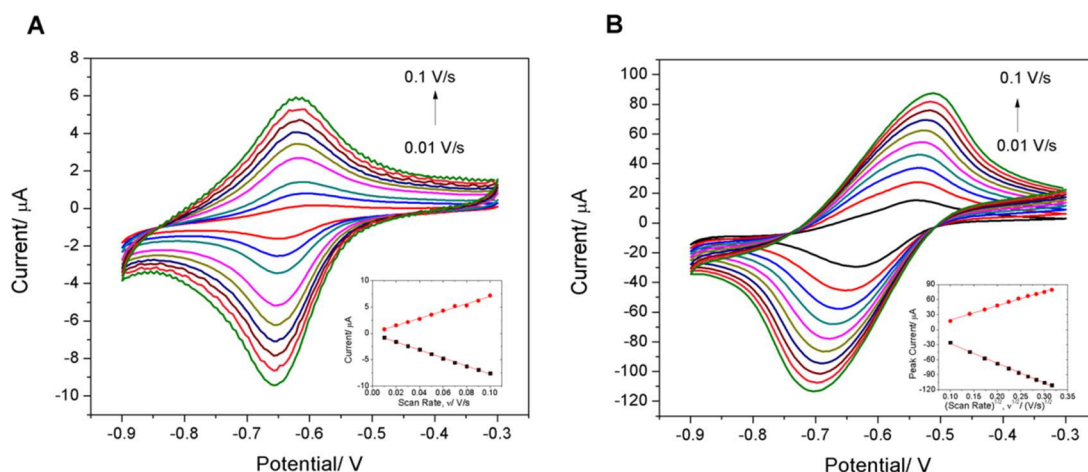


Figure 3: Cyclic voltammograms of (A) 20 μM and (B) 1000 μM DQ^{2+} on mesoporous silica film with scan rates ranging from 10 – 100 mV s^{-1} . Insets show the peak current as a function of (A) scan rate, v , and (B) square root of scan rate, $v^{1/2}$.

Logarithmic plots of current vs scan rates are shown in Figure 4 for the different redox probes. A slope of 1 indicates that the electrochemical response is confined to a thin-layer behaviour whereas a slope of 0.5 is due to a semi-infinite diffusion process according to Randles-Sevcik equation [80]. At the lower concentration of DQ^{2+} and PQ^{2+} , adsorbed species are dominating the electrochemical response as indicated by the slope of nearly 1 recorded. However, for the same concentration of $\text{Ru}(\text{NH}_3)_6^{3+}$ and $\text{Ru}(\text{bpy})_3^{2+}$, a mixed behaviour, with both diffusion and thin-layer processes, was observed (slopes were measured at 0.85 and 0.80 for $\text{Ru}(\text{NH}_3)_6^{3+}$ and $\text{Ru}(\text{bpy})_3^{2+}$, respectively). These slope values lower than unity (Figure 4A), supported that geometry of the cationic probes may play a role in the electrochemical behaviour observed in such confined channels. In the case of $\text{Ru}(\text{bpy})_3^{2+}$, this is further demonstrated by the deviation from the linear behaviour observed when the scan rate increased. This differences of behaviour could be attributed to either an effect of the pseudo-diffusion of electrons or to the limitation of counter-ions diffusion in the mesochannels. When probe concentrations were increased, the slope values approached closer to 0.5, which indicated the transition from thin layer behaviour to semi-infinite diffusion. At these higher concentrations, there was no significant behaviour between the different redox probes (Figure 4B).

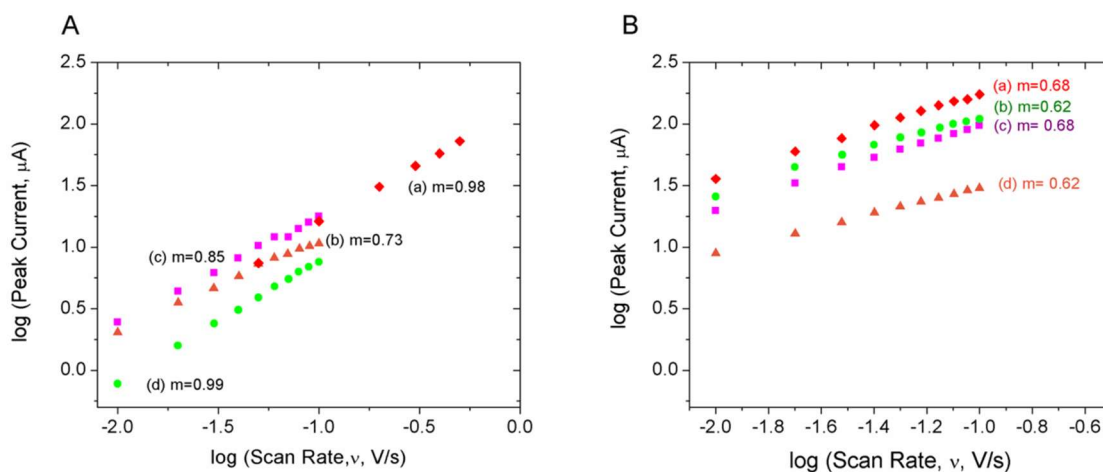


Figure 4: Logarithmic plots of peak current vs scan rate at low concentration for PQ^{2+} (a), DQ^{2+} (b), $\text{Ru}(\text{NH}_3)_6^{3+}$ (c), and $\text{Ru}(\text{bpy})_3^{2+}$ (d) at (A) low ($20 \mu\text{M}$) and (B) high concentration ($1000 \mu\text{M}$ for PQ^{2+} , DQ^{2+} and $\text{Ru}(\text{NH}_3)_6^{3+}$, and $500 \mu\text{M}$ for $\text{Ru}(\text{bpy})_3^{2+}$). All parameters are the same as for Figure 3.

In the case of mixed behaviour, both diffusion-controlled and thin layer-confined reactions contributed to the recorded current, which was assessed using the following equation [81]:

$$i = k_1\nu + k_2\nu^{1/2} \quad (1).$$

In equation (1), i is the current measured at a given potential, ν was the scan rate and k_1 and k_2 were empirical parameters representing the thin layer and the diffusion contributions, respectively. Normalising equation (1) led to equation 2:

$$\frac{i}{\nu^{1/2}} = k_1\nu^{1/2} + k_2 \quad (2).$$

The values for the parameters k_1 and k_2 were extracted from the plots of normalized current values vs the square root of potential scan rate. The thin-layer confined and semi-infinite diffusion contributions to the current were plotted for different scan rates and concentrations (Figure 5). At the low concentration values, the electrochemical response is dominated by the reaction confined in the thin layer and its contribution increased slightly with the scan rate (Figure 5A). When the redox probe concentration is higher, the contribution of semi-infinite diffusion became dominant (Figure 5B). However, a complete comparison on the effect of the thin-layer diffusion the different redox probes is difficult, because of pseudo-diffusion of electrons [72] and of the counter ion diffusion through the film [47,82]. Nevertheless, the dominating thin-layer behaviour for the lower concentration (i.e. 20 μM), was observed for the redox probes investigated (as shown in Figure S5).

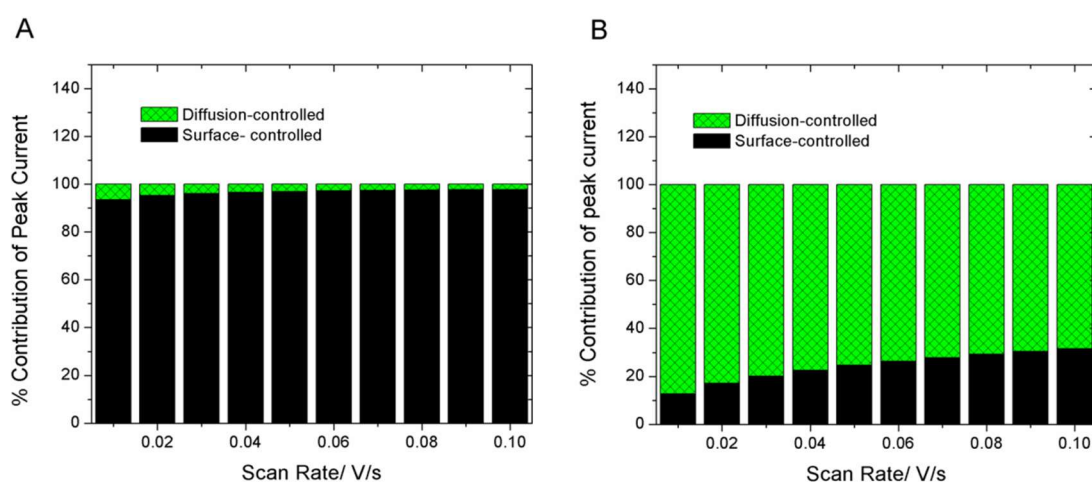


Figure 5: Contribution of thin-layer and semi-infinite diffusions to the current at mesoporous silica thin film electrodes for DQ^{2+} at (A) 20 and (B) 1000 μM . Scan rates were ranging from 0.01 to 0.1 V s^{-1} . Supporting electrolyte was 70 mM NaNO_3 . Similar graphs for the other redox probes are shown in supplementary information.

3.3. Effect of redox probe concentration on its accumulation in the mesochannels

The concentration of redox probes was varied between 10 μM and 8 mM and the forward peak current of cyclic voltammograms obtained at both bare and mesoporous silica film electrodes was plotted as a function of concentration (Figure 6). A linear response was observed for low concentrations of PQ^{2+} (10 - 200 μM , $R^2 = 0.999$), but exhibited two other different linear ranges at higher concentrations: 300 - 1000 μM ($R^2 = 0.985$) and 2000 - 8000 μM ($R^2 = 0.996$). As the concentration increased, the sensitivity of the modified electrode towards the redox probe decreased as shown by the results in Table 1; unlike the sensitivity at the bare electrode, which remained relatively constant throughout the 10 - 8000 μM concentration range. At the lower concentration range (10 – 200 μM), the sensitivity for PQ^{2+} at the mesoporous silica film electrode was three times higher than at the bare electrode, while the ratio of sensitivity of modified / bare electrode dropped to 1.8-fold for the 300 -1000 μM . This decrease of sensitivity with the concentration observed at the mesoporous silica electrode is consistent with the fall of the contribution of thin-layer diffusion and the increasing contribution of semi-infinite diffusion. At the highest concentration (8000 μM), the signal recorded at the modified electrode was even lower than at bare FTO, which is ascribed to restricted mass transport through mesochannels containing such large amounts of PQ^{2+} . For the lowest concentrations, improved sensitivities for the mesoporous silica film electrodes were also observed for $\text{Ru}(\text{NH}_3)_6^{3+}$ (1.5-fold improvement) and for $\text{Ru}(\text{bpy})_3^{2+}$ (3.2-fold improvement).

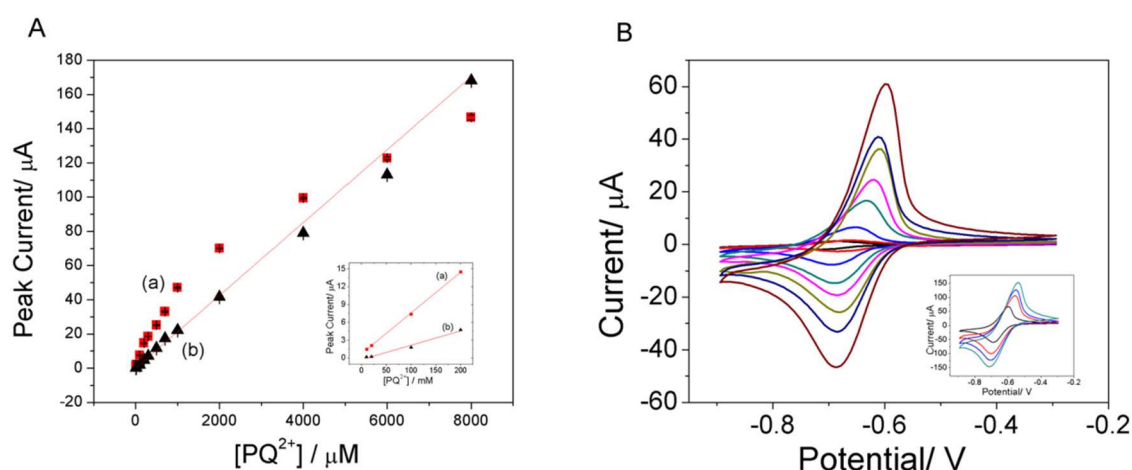


Figure 6: (A) Cyclic voltammograms for 10 – 1000 μM PQ^{2+} at a mesoporous silica film electrode (50th cycles); supporting electrolyte was 70 mM NaNO_3 ; $\nu = 20 \text{ mV s}^{-1}$. Inset shows cyclic voltammograms in the 2000 - 8000 μM concentration range. (B) Calibration curves of

PQ²⁺ at a mesoporous silica film electrode (a, red squares) and at bare FTO electrode (b, black triangles).

Table 1: Sensitivities for PQ²⁺, Ru(bpy)₃²⁺, and Ru(NH₃)₆³⁺ at mesoporous silica films and bare FTO electrodes

Electrode	Probe	Dynamic range / μM	Sensitivity / $\text{A M}^{-1} \text{cm}^{-2}$	R^2
Mesoporous silica film	PQ ²⁺	10 – 200	0.274	0.999
		300 – 1000	0.162	0.986
		2000 – 8000	0.052	0.996
	Ru(bpy) ₃ ²⁺	2 – 20	0.192	0.991
	Ru(NH ₃) ₆ ³⁺	10 – 100	0.095	0.999
Bare FTO electrode	PQ ²⁺	10 – 200	0.089	0.998
		300 – 1000	0.092	0.997
		2000 – 8000	0.080	0.998
	Ru(bpy) ₃ ²⁺	2 – 20	0.065	0.977
	Ru(NH ₃) ₆ ³⁺	10 – 100	0.060	0.998

4. Conclusions

The aim of this study was to characterize the accumulation of cationic redox probes at electrodes modified with mesoporous silica films for the improvement of electrochemical sensors. The electrochemical signal of the cationic redox probes was enhanced in the following order: MB⁺ > PQ²⁺ > DQ²⁺, Ru(bpy)₃²⁺ > Ru(NH₃)₆³⁺. Monocationic species are accumulated more efficiently, probably due to the necessity of more counter ions to diffuse with multi-charged species to maintain the charge balance. Scan rate studies showed that thin-layer diffusion dominated at the lower concentration for all redox probes, even for Ru(NH₃)₆³⁺, which accumulation was rather limited compared to the other probes. As the redox probe concentration increased, semi-infinite diffusion contribution increased, cancelling out the benefit of accumulation. Mesoporous silica film electrodes exhibited higher sensitivities than bare electrodes at lower concentrations, which can give benefits for sensor applications at trace levels.

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