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Gas-phase deposition of di- and tetra-lithium salts of 2,5-dihydroxyterephthalic acid†

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Thin films of two ambipolar lithium-organic electrode materials, Li₂DHTP and Li₄DHTP, are grown from gaseous precursors, Li(thd) (tetramethyl heptanedione) and DHTP (dihydroxyterephthalic acid). These precursors are pulsed into the reactor in a sequential manner like in atomic/molecular layer deposition, but the reaction product, i.e. the di- or the tetra-lithium salt, is controlled by adjusting the precursor pulse lengths.

Introduction

Metal salts of 2,5-dihydroxyterephthalic acid (H₄-DHTP) comprise an interesting group of coordination polymer (CP) and metal-organic framework (MOF) type metal-organic materials with extraordinarily rich coordination chemistries.^{1–5} The richness arises from the two different reactive groups, carboxylic acid, and phenol, both present as duplicates in the H₄-DHTP molecule. This allows the formation of both di- and tetra-metal salts, depending on the bonding scheme and the degree of deprotonation of H₄-DHTP. A comprehensive account of the coordination modes with different metals is found in a recent paper by Quarez *et al.*⁶ Potential applications proposed for these materials include gas adsorption, detection, and separation,^{2,4,5,7,8} as well as active rechargeable batteries^{3,9,10} and electrochromic devices.¹¹

With lithium, both the H₄-DHTP derivatives, Li₂DHTP and Li₄DHTP (Fig. 1), have been realized in crystalline form through conventional solution-synthesis routes, using respectively, Li₂CO₃ or MeOLi as the lithium source.^{3,9,10} The Li-rich Li₄DHTP is air-sensitive though, yielding Li₂O as a result of auto-oxidation.³ The carboxylate moiety undergoes reversible redox reactions at *ca.* 0.8 V vs. Li⁺/Li and the aryloxide (reacted phenol) at *ca.* 2.6 V vs. Li⁺/Li. This makes Li₄DHTP – similarly

to Na₄DHTP¹² – an interesting ambipolar electrode material for symmetric cells with an average working potential of 1.8 V vs. Li⁺/Li.⁹

Electrochemically active organic materials, in general, are currently gaining significant attention. They may not beat the state-of-the-art inorganic electrode materials in overall battery performance but could provide an attractive alternative when issues related to critical raw materials, energy-efficient synthesis/recycling, or gravimetric and mechanical properties are considered. The major challenges of the organic electrode materials are their intrinsically low electron conductivity and dissolution in liquid electrolytes; these issues can, however, be circumvented by applying the materials in thin-film form in combination with a solid electrolyte.^{13–17} The all-solid-state thin-film configuration would also serve as an optimal model system for investigating the intrinsic properties of organic electrode materials and understanding their solid-state electrochemical behavior and stability.^{15,18–22} Hence, it is important to develop reliable and scalable thin-film fabrication processes for the most interesting organic electrode materials.

In recent years, the combined ALD/MLD technique,^{23–25} derived from the atomic layer deposition (ALD) technology of high-quality inorganic thin films for microelectronics and beyond, has been harnessed for the growth of many intriguing *in situ* crystalline metal-organic materials directly from gaseous precursors.^{18–20,26–32} A prototype ALD process is based on two mutually reactive gaseous (or evaporated with moderate

Fig. 1 Visualization of the chemical compositions and specific bonding features of the targeted materials.

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heating) precursors sequentially pulsed into the reactor chamber to facilitate the atomic layer by atomic layer film growth with excellent control for the film thickness, homogeneity, and conformity. Ideally, the two successive surface reactions should saturate and be self-limiting, meaning that the growth per cycle (GPC) levels off to a constant value when long enough precursor pulses are applied. Molecular layer deposition (MLD) is an extension of ALD, to deposit purely organic thin films from two different organic precursors in a similar fashion; ideally, the process controllability and the film quality should be of the same level as in ALD. Then, in the hybrid ALD/MLD technique a metal-bearing ALD precursor is combined with an organic MLD precursor to deposit metal-organic thin films within the same sequential gas-surface reaction scheme. The precursor pulse times required for perfect saturation are somewhat reactor and reaction condition specific, but usually these pulsing times do not exceed a few seconds for metal precursors and a few tens of seconds for heavy organic precursors (longer pulse lengths needed due to their typically lower vapor pressures).^{24,33–35}

In this communication, we use a similar sequential precursor supply scheme to introduce Li(thd) (thd = 2,2,6,6-tetramethyl-3,5-heptanedione) and H₄-DHTP precursor vapors in an ALD/MLD reactor chamber. Excitingly, the films grow in a highly reproducible manner such that the film thickness is well controlled by the number of ALD/MLD cycles applied (once the other deposition parameters are fixed), but the end product, *i.e.* Li₂DHTP or Li₄DHTP, depends on the lengths of the precursor pulses. The two types of films are readily distinguished from each other with X-ray diffraction, infrared absorption, and cyclic voltammetry data.

Methods

Commercial H₄-DHTP powder (98%, Sigma-Aldrich) and in-house synthesized Li(thd) powder were used as the precursors; the reagents for Li(thd) synthesis were LiOH (98%, Alfa Aesar) and 2,2,6,6-tetramethyl-3,5-heptanedione (Tokyo Chemical Industry Co. Ltd) and the product was purified by sublimation. The films were grown on p-doped silicon test wafers (Okmetic Ltd), and also on stainless steel disks (MTI Corporation) for the electrochemical characterization, using an F-120 flow-type hot-wall ALD reactor (ASM Microchemistry Ltd) functioning at a low pressure of *ca.* 5 mbar and operated under nitrogen (99.999%) atmosphere generated from the air (Parker HPN 5000 N₂ generator). The same N₂ gas was used as a carrier and purge gas. The Li(thd) and H₄-DHTP precursor powders, placed in separate containers within the reactor, were heated for sublimation at 175 and 190 °C, respectively; the latter temperature was chosen based on TG-vacuum analysis.³⁶ Most of the depositions were made at 200 °C, but few depositions of 300 cycles were made at 250 °C, just to confirm that the observations also take place at higher temperatures.

The X-ray diffraction patterns were recorded for the samples in grazing-incidence geometry (GIXRD; X'Pert Pro,

PANalytical; Cu K α ; incidence angle 0.5°). X-ray reflectivity (XRR) patterns were recorded with the same equipment to determine the film thicknesses and densities. Chemical compositions and bonding structures were studied with Fourier transform infrared spectroscopy (FTIR; Bruker Alpha II); the resolution was 4 cm⁻¹ over 32 scans, and the spectrum of a blank reference wafer was subtracted from the measured spectra.

Samples for the electrochemical characterizations were deposited on stainless steel substrates ($\varnothing$ 15.5 mm). These coated steel disks were applied as a working electrode in CR2016 coin cells assembled in a glove box (<1 ppm of O₂ and <0.1 ppm of H₂O). The samples were dried in a vacuum (110 °C) and moved under vacuum into a glovebox (<1 ppm for O₂, <0.1 ppm for H₂O). Lithium foil (Sigma Aldrich, 99.9%) was used as a counter electrode, and 1 M LiPF₆ in 1:1 EC:DMC as electrolyte. The cells were let to stabilize for 24 hours prior to the cyclic voltammetry (CV) measurements (Autolab PGSTA302N potentiostat).

Results

Since H₄-DHTP contains both carboxylic acid and phenol functional groups, we could expect from our previous experiences with terephthalic acid and hydroquinone precursors combined with Li(thd) and LiHMDS (lithium bis(trimethylsilyl)amide), respectively,^{18,19,37} that H₄-DHTP could readily react with these common lithium precursors; for the present study, our choice was Li(thd), but presumably, LiHMDS would work similarly. Based on these previous experiences, we selected the following precursor and purge pulsing sequence for our initial experiments: 4 s Li(thd)/4 s N₂/10 s H₄-DHTP/30 s N₂. As was predicted, this process was found to yield Li-organic thin films in a reproducible manner independent of the pulsing sequence. Further analysis with GIXRD and FTIR (details discussed later) indicated that the films were *in situ* crystalline of the Li₂DHTP phase.

Among the two types of functional groups, the carboxylic acid group is more acidic, and our assumption was that it would react first with Li(thd); indeed, this seems to be the case as the deposition product was found to be Li₂DHTP. Then, the exciting discovery was that apparently the reaction proceeds further to Li₄DHTP if the Li(thd) pulse length is elongated. When the Li(thd) pulse length exceeded 32 s (while keeping the H₄-DHTP pulse length fixed at 10 s, the crystalline films were found to transform into amorphous Li₄DHTP phase. This second reaction step resulting in Li₄DHTP was noticed – besides the GIXRD observation – from the GPC *versus* Li(thd) pulse length plot in Fig. 2, as this plot does not show the ordinary saturation behavior expected for a prototype ALD/MLD process yielding a single product. From Fig. 2, essentially phase-pure Li₂DHTP films could be grown with the rate of *ca.* 4–8 Å per cycle, while the GPC value seen in the case of the essentially phase-pure Li₄DHTP films was *ca.* 13 Å per cycle, *i.e.* roughly twice the value seen for Li₂DHTP. It should

Fig. 2 GPC versus Li(thd) pulse length (H₄-DHTP pulse length fixed to 10 s); the resultant product (Li₂DHTP or Li₄DHTP) is highlighted.

however be noted that compared to the Li₂DHTP films, the Li₄DHTP films were less uniform with some thickness profile along the flow direction, and thus the GPC values were less precise.

The two phases, Li₂DHTP and Li₄DHTP, present in the samples were most easily distinguished by FTIR (Fig. 3a). In both cases, the features seen in the spectra can be interpreted based on the spectra previously reported and interpreted for bulk Li₂DHTP to Li₄DHTP samples (ESI S1–3†).^{3,10} The most important features are the asymmetric (ν_{as}) and symmetric (ν_s) vibrations of the carboxylate group. The separation of these vibrations gives a rough indication of the binding mode between the metal ion and the carboxylate group. The transformation from Li₂DHTP to Li₄DHTP decreases the peak separation from 281 to 193 cm⁻¹ and shifts the position of the ν_{as} peak from 1642 to 1557 cm⁻¹. The straightforward interpretation of these observations would be that the binding mode changes from monodentate for Li₂DHTP to bridging type for Li₄DHTP.³⁸ The coordination chemistry of DHTP is however quite complicated, and verification of this interpretation would require knowledge of the detailed crystal structure which unfortunately is known for neither Li₂DHTP nor Li₄DHTP. Indeed, only the crystal structure of hydrated Li₂DHTP·4H₂O is known,⁶ but this structure is apparently different from those of Li₂DHTP and Li₄DHTP judging from their different XRD patterns. An additional important signature to distinguish Li₄DHTP from Li₂DHTP in FTIR is the wide

absorption feature seen for Li₂DHTP in the 2500–3000 cm⁻¹ area due to intramolecular hydrogen bonding between the phenol and carbonyl groups. These peaks are naturally not seen for Li₄DHTP, as such hydrogen bonds are not possible.

The phase composition of the films was visible also from the XRR and GIXRD data (Fig. 3b and c). The film densities derived from the XRR data were different for the films composed of Li₂DHTP (1.73 g cm⁻³) and Li₄DHTP (1.52 g cm⁻³). From GIXRD, the Li₂DHTP films are well crystalline, with very intense peaks at 27.6° and 28.6°; the intensity ratio and peak positions match the previously reported diffraction pattern for evacuated bulk samples of Li₂DHTP (but not those of Li₂DHTP·4H₂O,⁶ as already mentioned).¹⁰ The as-deposited Li₄DHTP films were amorphous, but were crystallized in ambient conditions over the night, forming – most likely – the quinone variant (no phenol groups) of Li₂DHTP, and eventually – unless protected – Li₂O as reported earlier for bulk samples.³ In contrast to Li₄DHTP, the Li₂DHTP films were stable in ambient conditions, and did not absorb moisture.

We characterized the electrochemical properties of the Li₂DHTP and Li₄DHTP films with cyclic voltammetry measurements (Fig. 4). Firstly, for Li₂DHTP cycled between 1.5–3.5 V,

Fig. 4 Cyclic voltammograms for Li₂DHTP and Li₄DHTP films. The Li₄DHTP voltammogram is offset by 10 µA cm⁻².

Fig. 3 Representative (a) FTIR spectra (most important spectral features highlighted), (b) XRR patterns, and (c) GIXRD patterns to show the differences between the Li₂DHTP and Li₄DHTP films.

no electrochemical activity due to the alkoxide/quinone pair was observed; the irreversible peaks seen are probably due to the electrochemical activity of the phenol ($-\text{OH}$).³⁹ It is known that phenol groups can be lithiated at low potentials coupled with formation of H_2 gas.^{40–42} Here also, the Li_2DHTP film could be electrochemically lithiated into Li_4DHTP by sweeping the potential to 0.1 V and back to OCV. After this lithiation cycle the alkoxide/quinone redox pair appeared at *ca.* 2.6 V vs. Li^+/Li , as expected.^{3,9} The as-grown amorphous Li_4DHTP film behaved in a different fashion; it exhibited only a single redox pair centered around 2.9 V (and unstable behaviour at higher voltages), which does not exactly agree with the expected quinone redox pair value. Whether or not this peak is due to some other redox pair in the molecule remained to be clarified in more detailed studies in the future. However, here we like to refer to a somewhat similar case of ALD V_2O_5 films, grown in both crystalline and amorphous forms depending on the precursors used, and showing different electrochemical characteristics depending on the crystalline/amorphous nature of the films.⁴³ Finally, we should mention that among the two phases, Li_2DHTP and Li_4DHTP , the latter is preferred for the electrochemical applications, as it possesses the required functional groups both for the negative ($-\text{COOLi}$) and the positive ($-\text{OLi}$) electrode redox couples. Moreover, in this compound the phenol groups are already lithiated such that no H_2 is formed during the cell operation.

Discussion

To gain a deeper understanding of the surface chemistry features of our $\text{Li}(\text{thd}) + \text{H}_4\text{-DHTP}$ deposition process, we carried out a set of additional experiments with different combinations of precursor pulse lengths (Table S1†). The clear conclusion was that essentially phase pure (crystalline) Li_2DHTP

films are reproducibly obtained when the $\text{Li}(\text{thd})$ pulse length is short and also clearly shorter than that of $\text{H}_4\text{-DHTP}$. On the other hand, to deposit pure (amorphous) Li_4DHTP films the $\text{Li}(\text{thd})$ pulse needs to be considerably longer than that of $\text{H}_4\text{-DHTP}$. When the $\text{H}_4\text{-DHTP}$ pulse length was of the same magnitude as that of $\text{Li}(\text{thd})$, the GPC value became very high (exceeding even 30 Å per cycle when the pulse length was 32 s for both precursors); it should be noted, however, that the film thickness determination was somewhat ambiguous for the films with the highest GPC values due to their roughness.

Tentatively, we assume that the Li_4DHTP phase forms through a similar lithium-replacing reaction as is known to occur for the transformation of MgF_2 to LiF .⁴⁴ Hence, intuitively one might think that an efficient method to fabricate Li_4DHTP could be to lithiate Li_2DHTP just with a very long $\text{Li}(\text{thd})$ pulse at the end of the film growth process. However, in practice, this may not work as pure Li_2DHTP is difficult to form in the given conditions. This is further explained below.

In Fig. 5 we present a schematic illustration of the reaction process. In **phase 1**, the faster reaction between $\text{Li}(\text{thd})$ and the carboxylic acid group takes place forming initially Li_2DHTP . After this, Li_2DHTP lays on the surface and if the incoming $\text{Li}(\text{thd})$ pulse is long enough, the lithiation of the phenyl moiety also begins, to eventually form Li_4DHTP in **phase 2**. Most likely this second reaction step is less controlled than the first one, being thus the underlying reason why the Li_4DHTP films were less uniform than the Li_2DHTP films. In **phase 3**, the formed Li_4DHTP reacts with the incoming $\text{H}_4\text{-DHTP}$ precursor; here again, the carboxylic acid group reacts more readily than the phenyl group with the $-\text{OLi}$ group. This reaction converts one Li_4DHTP moiety into two Li_2DHTP moieties, which can again react in the “expected” way with the $\text{H}_4\text{-DHTP}$ precursor. The proposed surface reac-

PULSING SCHEMATIC

REACTION SCHEMATIC

Fig. 5 Schematic illustration of the proposed reactions occurring in the $\text{Li}(\text{thd}) + \text{H}_4\text{-DHTP}$ process with different precursor pulse lengths.

tions could be simplified with the following exchange reactions:

Owing to the exchange of $\text{H}_4\text{-DHTP}$ to Li_4DHTP , during one cycle four $\text{H}_4\text{-DHTP}$ moieties may be deposited on the surface. Thus, to make the deposition process for Li_2DHTP films as controllable as possible, the formation of Li_4DHTP during the process should be avoided, as its transformation back to Li_2DHTP leaves extra molecules on the surface.

The Li_4DHTP films, on the other hand, often exhibited gradients over areas where the precursor flux is expected to be the highest; the most straightforwardly expected cause for this is an insufficient precursor pulse time. It seems that Li_4DHTP forms fastest on the substrate areas that saturate first. If the whole film area is not successfully converted into Li_4DHTP , large growth gradients may quickly form due to the extra surface species generated (that would again saturate quicker than the other areas making the problem even worse).

Finally, we like to mention that in principle, it should be possible to perfectly transform the entire film into Li_4DHTP and then back to Li_2DHTP to achieve uniform Li_2DHTP coatings with an extraordinarily high growth rate. However, this was difficult to achieve in practice and the experiments resulted in non-uniform growth along the flow direction. In the future, it might be interesting to play with the reactivity of the lithium precursor. Towards this research direction, we already carried out a few experiments with LiHMDS that is more reactive than $\text{Li}(\text{thd})$, to find out that the growth rate increased but the film uniformity got worsened. The other option, *i.e.* to look for a less reactive lithium precursor, remained to be challenged in future works.

Conclusions

We have presented straightforward and reproducible gas-phase fabrication routes for two intriguing Li-organic materials, Li_2DHTP and Li_4DHTP , for the first time. The processes are based on two precursors only, *i.e.* $\text{Li}(\text{thd})$ and $\text{H}_4\text{-DHTP}$, sequentially pulsed in an ALD/MLD type reactor. The exciting feature – which distinguishes the processes from the ideal ALD/MLD scheme – is that the phase composition of the films is controlled by the precursor pulse lengths, the Li_2DHTP films requiring shorter and the Li_4DHTP films longer $\text{Li}(\text{thd})$ pulses. The $\text{Li}(\text{thd}) + \text{H}_4\text{-DHTP}$ process delivers the Li_2DHTP films in *in situ* crystalline form, while the Li_4DHTP films are initially amorphous but are crystallized during storage in ambient conditions. Preliminary characterization of the films with cyclic voltammetry

experiments demonstrated that both the Li_2DHTP and Li_4DHTP films are electrochemically active, as desired.

We foresee that similar unconventional surface-reaction pathways as found and successfully utilized here for the Li_2DHTP and Li_4DHTP films could be innovatively exploited also with other interesting organic molecules with multiple different reactive groups connected to the same organic moiety.

Author contributions

Juho Heiska: Conceptualization, methodology, investigation, writing – original draft, visualization. Maarit Karppinen: Supervision, funding acquisition, writing – original Draft.

Conflicts of interest

There are no conflicts of interest to declare.

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