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Unusual oxidation behavior of light metal hydride by tetrahydrofuran solvent molecules confined in ordered mesoporous carbon

Markus Klose^{a)}

IFW Dresden, Institute for Complex Materials, D-01171 Dresden, Germany; and Institut für Werkstoffwissenschaft, Technische Universität Dresden, D-01069 Dresden, Germany

Inge Lindemann and Christian Bonatto Minella

IFW Dresden, Institute for Metallic Materials, D-01171 Dresden, Germany; and Institut für Werkstoffwissenschaft, Technische Universität Dresden, D-01069 Dresden, Germany

Katja Pinkert, Martin Zier, and Lars Giebeler

IFW Dresden, Institute for Complex Materials, D-01171 Dresden, Germany; and Institut für Werkstoffwissenschaft, Technische Universität Dresden, D-01069 Dresden, Germany

Pau Nolis

Servei de Resonància Magnètica Nuclear (SeRMN), Universitat Autònoma de Barcelona, 08193 Bellaterra, Spain

Maria Dolors Baró

Departament de Física, Universitat Autònoma de Barcelona, E-08193 Bellaterra, Spain

Steffen Oswald

IFW Dresden, Institute for Complex Materials, D-01171 Dresden, Germany

Oliver Gutfleisch

Materials Science, Technische Universität Darmstadt, 64287 Darmstadt, Germany

Helmut Ehrenberg

Karlsruhe Institute of Technology (KIT), Institute for Applied Materials (IAM), D-76344 Eggenstein-Leopoldshafen, Germany

Jürgen Eckert

IFW Dresden, Institute for Complex Materials, D-01171 Dresden, Germany; and Institut für Werkstoffwissenschaft, Technische Universität Dresden, D-01069 Dresden, Germany

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Confining light metal hydrides in micro- or mesoporous scaffolds is considered to be a promising way to overcome the existing challenges for these materials, e.g. their application in hydrogen storage. Different techniques exist which allow us to homogeneously fill pores of a host matrix with the respective hydride, thus yielding well defined composite materials. For this report, the ordered mesoporous carbon CMK-3 was taken as a support for LiAlH_4 realized by a solution impregnation method to improve the hydrogen desorption behavior of LiAlH_4 by nanoconfinement effects. It is shown that upon heating, LiAlH_4 is unusually oxidized by coordinated tetrahydrofuran solvent molecules. The important result of the herein described work is the finding of a final composite containing nanoscale aluminum oxide inside the pores of the CMK-3 carbon host instead of a metal or alloy. This newly observed unusual oxidation behavior has major implications when applying these compounds for the targeted synthesis of homogeneous metal–carbon composite materials.

I. INTRODUCTION

Light metal hydrides such as LiAlH_4 or NaAlH_4 are currently extensively investigated for their possible application in hydrogen storage systems due to their high hydrogen content of 10.6 and 7.4 wt%, respectively. Among the challenges that still need to be overcome to

introduce these materials into commercially available systems, hampered hydrogen desorption and reabsorption kinetics are probably the most pressing issue.^{1,2} It has been proposed that confining light metal hydrides into micro- and mesoporous scaffolds can help to solve some of the current limitations. For that purpose, different lightweight carbon-based materials such as carbon nanotubes^{3,4} or high surface area⁵ graphite were used as well as microporous metal–organic frameworks.^{6,7} Reducing the size of the active material down to several nanometers can dramatically alter its stability leading to vastly reduced

^{a)} Address all correspondence to this author.

e-mail: m.klose@ifw-dresden.de

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activation energies for hydrogen desorption as well as enhanced desorption kinetics. Moreover, due to reduced mobility of the resulting decomposition products, phase separation and sintering occur less pronounced.^{8–13} The result of these efforts is a notably improved homogeneity of the obtained composite materials.

Two general pathways exist for infiltrating light metal hydrides into micro- or mesoporous matrices. When using the melt infiltration technique the hydride is molten into the porous host after being heated under elevated hydrogen pressure. The latter is required to shift the chemical equilibrium toward the hydride and to prevent preliminary decomposition.¹⁴ While this procedure is somewhat feasible for sodium aluminum hydride, lithium aluminum hydride is metastable at room temperature and would require an equilibrium pressure that is beyond the technological means available.¹⁵ An alternative can be found in the solution impregnation procedure. Here the hydride is dissolved in a suitable solvent and then added to, e.g., the carbon compound. Followed by a drying step to evaporate the solvent, the impregnation step can be repeated several times to increase the loading. Lithium aluminum hydride decomposes in three steps according to the following equations:

Thus, by introducing lithium aluminum hydride into porous carbon such as CMK-3 and then desorbing the hydrogen entirely should allow for the formation of highly homogeneous aluminum metal–carbon composites. These composites are currently of great interest, e.g., for applications in lithium ion batteries.^{16–18} However, as already reported in earlier studies on the direct synthesis of complex metal hydrides by Ashby et al.,^{19,20} it is rather difficult if not entirely impossible to remove the last traces of tetrahydrofuran (THF) once in contact with LiAlH_4 . Nevertheless, more recent investigations showed the importance of THF as an adduct to LiAlH_4 , which dramatically improves the thermodynamics of both hydrogen desorption and reabsorption. Wang et al.²¹ pointed out that the regeneration of LiAlH_4 in the presence of THF as an adduct to the hydride was about seven times more effective compared with experiments without THF. In contrast, there exist no reports on the fabrication of oxidic compounds by using light metal hydrides in THF.

Therefore, this study focuses on tetrahydrofuran as it is one of the most commonly used solvents for light metal hydrides and its effect on the decomposition of LiAlH_4 when confined in mesoporous carbon CMK-3. A compre-

hensive investigation of both the desorption process as well as the resulting final compounds using solid-state magic angle spinning nuclear magnetic resonance (MAS-NMR), depth profile auger electron spectroscopy, and x-ray photoelectron spectroscopy is presented.

II. METHODS

A. Preparation procedure

All materials were used and stored under argon inert gas atmosphere in a glove box with water and oxygen levels below 3 ppm at all times. The borosilicate glassware used for handling the hydrides was dried under vacuum at 373 K for 24 h to remove traces of water adsorbed to the surface. Quartz glassware was dried at 1073 K. Mesoporous carbon CMK-3 was prepared according to a procedure described elsewhere.²² Prior to the impregnation process, it was heated to 873 K under 0.9 bar hydrogen over pressure to reduce any oxidizing surface functionalities, which might react with the hydride. A two molar solution of LiAlH_4 (LAH) in THF was obtained from Sigma-Aldrich and used without further purification. For a typical impregnation, 100 mg CMK-3 was used together with 80 μL LiAlH_4 solution. The evaporation of THF was performed at 373 K under vacuum for at least 3 h. This impregnation procedure was repeated several times with stepwise reduced amounts of solution until the carbon material was loaded with ~ 17 wt% LiAlH_4 , hereby denoted as “LAH@CMK-3.” In accordance with the results obtained by nitrogen physisorption experiments, the amount of solution added never exceeded the available inner pore volume of CMK-3, thus ensuring that no hydride recrystallized outside of the carbon pore system. LAH@CMK-3 was then heated under dynamic vacuum with a heating rate of 1 K/min up to 673 K and subsequent holding at this temperature for 23 h. Afterward, the material was cooled naturally. The resulting compound is referred to as “LA@CMK-3.”

For x-ray powder diffraction (XRD) experiments, powder samples were fixed as a thin layer on an acetate film. Data were collected in flat sample transmission mode on a STOE STADI P powder diffractometer equipped with a 6° -position sensitive detector (PSD) and a Ge(111)-monochromator. The samples were measured with $\text{Cu K}\alpha_1$ -radiation with a step size of $\Delta 2\theta = 0.01^\circ$.

B. Nitrogen physisorption experiments

The nitrogen sorption experiments were carried out using a Quantachrome Quadrasorb SI apparatus. Prior to the measurement, the samples were degassed under dynamic vacuum at 393 K for 24 h. Specific surface areas were calculated at a relative pressure $p/p_0 = 0.3$ using the single-point Brunauer–Emmett–Teller (BET) method according to the BET theory. The total pore volume was

determined at $p/p_0 = 0.9$. The pore size distributions were obtained using the Quenched Solid Density Functional Theory (QSDFT) equilibrium model implemented in the Quadrawin 5.05 software by Quantachrome assuming a cylindrical pore geometry which in agreement with the actual pore geometry of the CMK-3.

C. MAS-NMR experiments

Solid-state MAS-NMR spectra were obtained using a Bruker Avance 400 MHz spectrometer with a wide bore 9.4 T magnet and a Bruker 4 mm CP-MAS probe. The spectral frequency was 104.27 MHz for the $^{27}\text{Al}\{^1\text{H}\}$ nucleus. The NMR chemical shifts are reported in parts per million (ppm) externally referenced to 1.0 M $\text{Al}(\text{NO}_3)_3$ aqueous solution. The material was packed into a 4 mm ZrO_2 rotor in an argon-filled glovebox and was sealed with a tight fitting Kel-F cap. Sample spinning of 10 kHz was chosen and it was performed using dry nitrogen gas. The one-dimensional $^{27}\text{Al}\{^1\text{H}\}$ MAS-NMR spectrum was acquired after a 4 μs single $\pi/2$ pulse (corresponding to a radiofield strength of 62.5 kHz) with application of a strong ^1H signal decoupling by using the two-pulse phase modulation (TPPM) scheme. Spectra were acquired at 20 °C controlled by a BRUKER BCU unit.

D. Thermoanalysis coupled with mass spectrometry

Simultaneous TG and mass spectrometry analysis (DTA-MS) was carried out in a Sensys Evo (Setaram) placed inside an argon-filled glovebox. The measurements were carried out using a 1 bar argon flow with a heating rate of 1 K/min. The TG-DSC system was coupled to a mass spectrometer from Pfeiffer Vacuum (Omnistar GSD 320 O1) located outside the glovebox via a heated stainless steel capillary (1 m, 423 K). The mass spectrometer uses an yttriated iridium filament and a QMG 220 quadrupole. The collected masses were monitored with the integrated C-SEM detector.

E. Depth profiling auger electron spectroscopy (DP-AES)

A JEOL JAMP-9500 F field emission auger microprobe with primary electrons of 10 keV at an electron current of 10 nA was used to conduct AES measurements and to obtain scanning electron micrographs (SEM). For depth profiling, the sample surfaces were sputtered by a scanned beam of 1.5 keV Ar^+ -ions at an incident angle of about 30°. Spectra were recorded after intervals of 0.3 min. The corresponding sputter rate calibrated to silicon dioxide was determined to be 1.4 nm for each cycle. The quantification of element concentration was performed by using the differentiated signals and the standard single element sensitivity factors.

F. X-ray photoelectron spectroscopy

The XPS measurements were carried out on a PHI 5600 CI (Physical Electronics) spectrometer equipped with a hemispherical analyzer operated at a typical pass energy of 29 eV and with an analysis area of 800 μm in diameter. Mg K_α excitation (350 W) was used without applying an additional energy electron charge neutralizer. To avoid any contact of the samples with air and moisture, a transfer chamber (Physical Electronics) was used for the sample transport from the argon-filled glove box to the XP spectrometer. All spectra were calibrated with respect to sp^2 -hybridized carbon at 285 eV. More details on the experimental procedures can be found in previous works.^{23–25}

III. RESULTS AND DISCUSSION

Figure 1 shows the x-ray patterns of both pure mesoporous carbon CMK-3 as well as LA@CMK-3 after the infiltration–desorption process described above. The pattern of LA@CMK-3 exhibits two reflections which are ascribed to metallic aluminum. Considering the narrow pore size of the carbon host (radii around 20 Å, see Fig. 2) as well as the sharp nature of those reflections, it is reasonable to assume that the aluminum phase is located on the outer surface of the carbon particles rather than inside the pore system. In the latter case, the aluminum would be expected to be nanocrystalline or even amorphous yielding much more broadened reflections in XRD. Such formation of a separate phase of pristine aluminum upon desorption of light metal hydrides confined in micro- or mesoporous carbon scaffolds is a commonly observed effect being characteristic for this particular combination of materials.^{26,27} Within the detection limit of XRD, no long-range ordered hydride is observed.

FIG. 1. X-ray diffraction patterns of pure CMK-3 (black) and the final composite material LA@CMK-3 (red). Reflections from Al metal are marked by the corresponding hkl indices.

Furthermore, nitrogen physisorption experiments were carried out to supplementarily demonstrate a successful infiltration process. Figure 2 shows the adsorption isotherms of pure CMK-3 and LA@CMK-3 and also the respective pore size distributions obtained by QSDFT calculations assuming cylindrical pore geometry. The initial pristine CMK-3 infiltrated sample exhibits a much reduced capacity for nitrogen uptake. Also, while the pore size of the carbon matrix itself is virtually unchanged, the quantity of accessible pores is notably reduced. Additionally, the total surface area and total pore volume decrease by 40 and 25%, respectively, as summarized in Table I, which indicates a partial filling of the pore system.

To further confirm the homogeneous filling of the porous matrix for LA@CMK-3, DP-AES was applied. By definite sputtering through the nanocomposite particle, the inner chemical composition is accessible for AES analysis. Figure 3(a) shows scanning electron micrographs of the as-synthesized nanocomposite. Figure 3(b) shows the SEM image of the nanocomposite after 75 sputter cycles to verify material stripping. The calculated depth profiles in relation to the number of sputtered cycles are depicted in Figs. 3(c) and 3(d). The two marked areas in Figs. 3(a) and 3(b) were analyzed by AES before the measurement [cycle no. 0, Figs. 3(c) and 3(d), bottom] and after each sputter cycle [cycle number 1-75, Figs. 3(c) and 3(d), bottom] to obtain information on the homogeneous distribution of LA@CMK-3. All three depth profiles show

FIG. 2. Pore size distributions of CMK-3 (black) and LA@CMK-3 (red) obtained by QSDFT-Method. The inset shows the respective N_2 physisorption isotherms.

TABLE I. BET surface areas and pore volumes of pure CMK-3 and LA@CMK-3.

	Surface area [m^2/g]	Pore volume [cc/g]
Pure CMK-3	1202	0.91
LA@CMK-3	715	0.68

very similar characteristics revealing the very homogeneous structure of the final composite, thereby proving that the chosen impregnation route is suitable and yields the desired homogenous material. Carbon is present as the main element, followed by aluminum and interestingly, also nonnegligible amounts of oxygen, which seems to trail the aluminum concentration profile. The detection of constantly present and homogeneously distributed oxygen throughout the final composite material is surprising since the decomposition of the LiAlH_4 precursor according to Eqs. (1)–(3) is not expected to yield any form of an oxygen-containing compound.

An indication of an unusual decomposition of the THF solvent molecules attached to the lithium aluminum hydride is provided by the curves obtained from the thermoanalysis (DTA-MS) shown in Fig. 4. Here, the mass and the respective amounts of fragments evolving from the sample during heating are given. This technique allows for in situ monitoring of both the desorption process and also the respective evolving gaseous species. Table II lists the measured mass numbers and the corresponding fragments.²⁸ The heat treatment of the sample was conducted under argon atmosphere with a heating rate of 1 K/min. Two major stages of decomposition are visible in the graph of Fig. 4. The first step reaches peak intensities of ion current for the mass numbers 27, 39, 42, 71, and 72 at around 400 K and 500 K for m/z 2. A second step is observed with m/z of 27, 39, 42, and 2, but this time without 71, 72 but instead 55 and 56 at around 570 K. This difference in the relative amounts of fragments that are detected at the respective temperatures clearly indicates that the products evolving from each stage are resulting from different types of decomposition reactions. Even more notably, these results differ significantly from the fragmentation processes of pure THF.²⁸ As described in the experimental section, each impregnation step of CMK-3 was followed by a drying step at 373 K under vacuum to remove excess solvent. Hence, taking into account the small size of the resulting LiAlH_4 , which leads to the overall downward shift of the respective desorption temperatures for each step, it can be assumed that the reaction described in Eq. (1) already took place during the preparation of LAH@CMK-3. This explains the observation of only two maxima during the desorption process. However, it is interesting to note that the mass numbers 71 and 72 corresponding to the “intact” THF molecule and a THF molecule without one α -H atoms do not appear in the second stage of fragments.^{28,29} Those fragments in the first stage result from solvent molecules which were not coordinated to the hydride. It can be concluded that after surpassing the threshold of approximately 523 K, those remaining coordinated THF molecules are not able to evaporate without substantial bond cleavage and consequently leave oxygen atoms attached to aluminum. It is also a matter of fact that metallic aluminum is highly attractive for absorption and bonding of oxygen. At this

FIG. 3. SEM images before (a) and after (b) auger electron spectroscopy experiments of LA@CMK-3. Depth profiles (c and d) were obtained at the positions 1 and 2, respectively.

point, it is therefore crucial to determine whether or not the detected oxygen is in fact bound to the aluminum.

For that reason, solid state MAS-NMR spectroscopy was performed to gain additional information on the composition of the newly formed compounds located inside the pores of the host matrix. Figure 5 represents two regions of ^{27}Al NMR spectra of the as-infiltrated sample LAH@CMK-3 (black line) as well as the final composite material LA@CMK-3 (red line). In Fig. 5(a), the signals centered at 1640 ppm belong to pure aluminum metal. This further supports the findings from XRD experiments, which also evidenced a fraction of pure aluminum being located outside of the CMK-3 carbon matrix. Lower chemical shifts of the ^{27}Al -NMR-spectra of the same samples are shown in Fig. 5(b). LAH@CMK-3 exhibits a broadened signal (#) between -25 and -50 ppm that corresponds to the hexahydride species Li_3AlH_6 , which is the result of the first decomposition stage of LiAlH_4 according to Eq. 1.³⁰ Hence, the detection of this hexahydride phase additionally confirms the results of the DTA-MS experiments discussed above, in which only two maxima of hydrogen concentration were observed, indicating that the first hydrogen desorption step from LiAlH_4 already took place during the infiltration step. For LA@CMK-3, two signals are located between 40–80 ppm and 0–10 ppm corresponding to aluminum–oxygen bonds with the distinction of both tetrahedral and also octahedral aluminum oxide sites being present.^{30,31} This finding is concurrent with earlier reports that studied the oxidation products of NaAlH_4 . First, a

FIG. 4. DTA-MS curves including the mass numbers of the respective fragments evolving during the heat treatment of LAH@CMK-3. See also Table II for more details.

highly disordered aluminum oxide/hydroxide is formed in which the aluminum is coordinated tetrahedrally by the oxygen atoms. This compound can then later be transformed into a more ordered aluminum oxide where oxygen coordinates aluminum octahedrally.³¹ However in our case, we clearly observe the NMR signals for both of these compounds. The fact that the line shape of the MAS NMR signals at low chemical shifts in Fig. 5(b) of the ^{27}Al -nuclei is broadened to this extent clearly suggests that the respective aluminum-containing compounds exhibit a high degree

TABLE II. Mass numbers and the respective fragments corresponding to Fig. 4.

Mass number	Fragment	Relative abundance according to Ref. 28
2	H ₂	-
27	C ₂ H ₃ ⁺	17%
39	C ₃ H ₃ ⁺	10%
42	C ₃ H ₆ ⁺	100%
55	C ₄ H ₇ ⁺	-
56	C ₄ H ₈ ⁺	-
71	C ₄ H ₇ O ⁺	30%
72	C ₄ H ₈ O ⁺	32%

of structural disorder, thus being either nanocrystalline or even amorphous. This is expected since those compounds are prevented from establishing a long-range order due to the spatial confinement at the nanometer scale imposed by the CMK-3 carbon matrix.^{32,33}

The assessment that indeed oxidic compounds result from the reaction of LiAlH₄ with the THF molecules is additionally supported by the results of the XPS investigations that are discussed as follows. Figure 6 shows the XP spectra of the infiltrated material as well as the final composite LA@CMK-3. For comparative reasons, also the C1s spectrum [Fig. 6(c)] of the pure untreated CMK-3 was added. It is clearly evident that the overall tendency is a strong shift toward higher binding energies of aluminum, oxygen, and a little less pronounced also for lithium when comparing the infiltrated sample with the desorbed one. The Al 2p peak [Fig. 6(a)] of LAH@CMK-3 is centered at 74.7 eV and the Li 1s signal is located mainly around 56.2 eV, which is consistent with values reported for pristine lithium alanate.^{34,35} Moreover, the O 1s peak [Fig. 6(d)] measured at 531.7 eV is at the expected position for molecules containing ether groups.³⁶ Please note that the signals in the C 1s spectrum at 285 eV, which are attributed to the sp²-hybridized carbon of the graphitic parts of the CMK-3, only marginally increase in width throughout the entire process, which likely happens due to local charging effects during the XPS measurement. Nevertheless, the consistency of that signal also points out that the structure of the CMK-3 host remains intact during the entire process of infiltration and desorption. After the heat treatment of LAH@CMK-3, both the Al 2p peak as well as the O 1s signal are shifted toward higher binding energies of 75.7 and 532.8 eV, respectively. This finding provides clear evidence that in fact aluminum oxides are among the final reaction products that evolve from the decomposition of the LiAlH₄ – THF species. In comparison to that, the Li 1s signal [Fig. 6(b)] only shifts to about 56.8 eV suggesting that the change in binding state for lithium overall is less severe than for the other elements. More notably, the C 1s XP spectrum of the final composite material features a second peak arising at 291.7 eV, which most likely accompanies the formation of a carbonate species or a similar organic residue, as re-

FIG. 5. MAS-NMR spectra of the ²⁷Al{¹H} experiments of LAH@CMK-3 (black) and LA@CMK-3 (red) revealing the presence of different aluminum species. (a) Signals of bulk aluminum centered at ~1640 ppm including sidebands. (b) Evidence for Li₃AlH₆ phase (#) in LAH@CMK-3 and aluminum oxides (*). Positions between ~40–80 ppm correspond to tetrahedral Al oxide sites, whereas those at ~0–10 ppm belong to octahedral Al oxide sites³¹. All signals are significantly broadened indicating the nanocrystalline or amorphous nature of the respective material.

ported by Andersson et al.^{34,37,38} A binding energy that is high, basically rules out the presence of any other carbon compound than might emerge considering the elements present in the sample.

Concluding from the afore discussed data, it is clearly evident that THF solvent molecules attached to LiAlH₄ can indeed irreversibly oxidize the hydride at elevated temperatures mainly forming nanosized aluminum oxides and probably also carbonates inside the carbon host. Revealing the detailed mechanism of this unexpected behavior is beyond the scope of this report. However, it shall be noted that the results presented here are in good agreement with previous findings. The destabilization of

FIG. 6. XPS spectra of pure CMK-3 (blue), LAH@CMK-3 (black), and LA@CMK-3 (red) showing the overall shift of binding energies toward more oxidic states.

etheral solvents by lithium compounds is widely known as well as the possibility of THF attaching to aluminum surfaces via its oxygen atom after undergoing a ring-opening reaction.^{39,40} A detailed mechanism for the decomposition of THF is proposed in the supporting information (Scheme S1, ESI). Lacina et al.⁴¹ pointed out that it is energetically most favorable for LiAlH_4 in solution to coordinate three THF molecules forming contact ion pairs at room temperature. It is assumed that the ability of different solvents to form adducts with the alanate is crucial for its successful regeneration.^{21,27,42} To our understanding of the case presented here, the last hydrogen desorption step of LiAlH_4 [Eq. (3)] is crucial for the formation of nanosized aluminum oxides that are confined in the pores of the CMK-3 carbon host.

IV. CONCLUSIONS

It was shown that it is possible to homogeneously infiltrate the lightweight hydride LiAlH_4 into the pore system of ordered mesoporous carbon CMK-3 via a multistep solution impregnation procedure. Analyzing the decomposition behavior of the infiltrated hydride as well as the resulting nanocomposite structure, we revealed an unexpected oxidation behavior of the THF solvent on the confined LiAlH_4 . Nanoscale aluminum oxides were formed

possibly along with carbonate species as a byproduct at temperatures up to 673 K. A reduction of the onset temperature at which hydrogen desorbs from the hydride is achieved by introducing it into the carbon scaffold, consistent with previous studies on complex light metal hydrides confined in mesoporous scaffolds. The presented findings offer an interesting new way toward the targeted formation of carbon-supported nanosized aluminum oxides, which might be promising, e.g., for applications in catalysis.^{43–45} In addition, we demonstrated the benefits of the combined use of multiple state-of-the-art techniques for analyzing the composite materials to gain a comprehensive picture of the structural features of these materials.

NOTES

The authors declare no competing financial interest.

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Supplementary Material

Supplementary materials can be viewed in this issue of the *Journal of Materials Research* by visiting <http://journals.cambridge.org/jmr>.