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HfS₂ thin films deposited at room temperature by an emerging technique, solution atomic layer deposition†

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As a member of the two-dimensional metal dichalcogenide family, HfS₂ has emerged as a promising material for various optoelectronic applications. Atomic layer deposition is widely used in microelectronics manufacturing with unique properties in terms of accurate thickness control and high conformality. In this work, a simple and versatile method based on the atomic layer deposition principles is presented to generate hafnium disulfide from the solution phase ('solution ALD' or sALD). For ease of comparison with the traditional gaseous atomic layer deposition (gALD) method, the same precursors are used, namely tetrakis-(dimethylamido) hafnium(IV) and H₂S. The deposit is characterized on several different oxide substrates by spectroscopic ellipsometry, scanning electron microscopy, and X-ray photoelectron spectroscopy. In the saturated regime, the growth rate depends on the substrate nature and is between 0.4 and 0.6 Å per sALD cycle. This growth rate determined at room temperature is lower than with the gALD process reported at 100 °C recently. At those low deposition temperatures, the films remain in an amorphous state. This success in sALD expands the range of material classes available by the new method, adding transition metal dichalcogenides to the list containing oxides, cubic sulfides, hydrides, and organics so far. It promises to overcome the precursor constraints associated with the traditional gALD method, in particular the volatility requirement.

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Introduction

HfS₂, as a member of the transition metal dichalcogenides, has recently emerged as a promising material for electronics and energy conversion applications in the semiconductor community due to its sizeable bandgap and other favorable physical properties.^{1–9} In comparison to its bulk counterpart, thin film HfS₂ has shown further intriguing properties.^{1,10} The methods reported so far for the synthesis of thin film HfS₂ have been mechanical exfoliation and chemical vapor deposition.^{11–14} Mechanical exfoliation is not applicable to the systematic variation of film thickness and study of the physical properties as they depend on it. Furthermore, its use in practical applications is debatable. Chemical vapor deposition yields continuous films over sizeable areas, which can be of high quality after treatment at elevated temperature. The film thick-

ness homogeneity, however, becomes limited for large substrates. Accordingly, significant effort has been dedicated to developing atomic layer deposition methods for the generation of HfS₂ thin films over large areas.^{15–17} Atomic layer deposition (ALD) relies on self-limiting surface chemical reactions of two distinct precursors introduced in the vapor phase to react with the solid surface in a sequential manner, avoiding direct contact between both precursors. Repeating the alternating pulses of the two precursors allows one to deposit films with high conformality even in deep pores and with accurate thickness control.¹⁸ In this traditional, gas-based ALD variant (gALD), the precursors need to fulfill several characteristics simultaneously: volatility, reactivity, and thermal stability to avoid thermal decomposition.^{19,20} These requirements constrain the list of ALD-grown materials. Moreover, vacuum conditions and high processing temperatures limit the range of thin film materials and substrates for which it is adequate.

Recently, the availability of deposition methods relying on self-limiting surface chemical principles^{21–27} inspired the development of solution atomic layer deposition (sALD) as a general equivalent of gALD in solution processing.^{28–31} The simplicity of its experimental realization represents one additional advantage of the novel method. Here, the surface chemical reactions of the precursors need to provide sufficient

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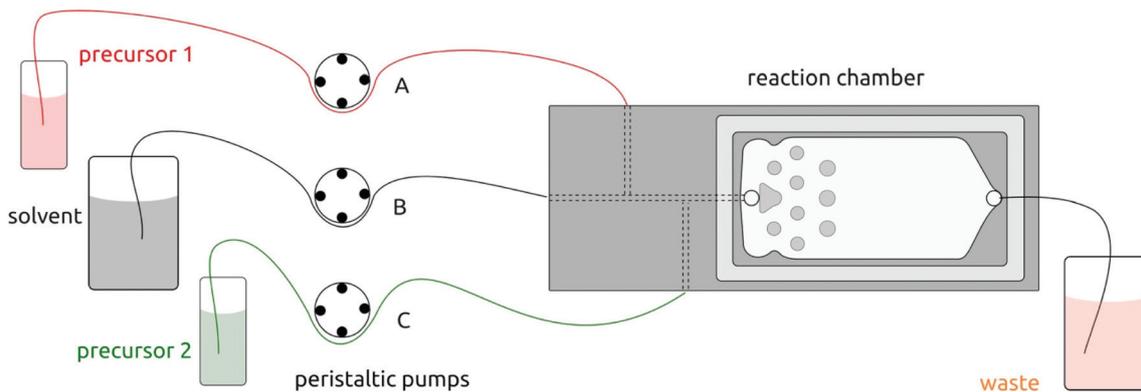


Fig. 1 Schematic illustration of the solution atomic layer deposition (sALD) setup. Three peristaltic pump channels drive the precursors and solvent to the reaction chamber in alternating fashion.

sufficient precursor delivery to the surface, avoid direct precursor contact, and minimize solvent consumption, so that an efficient trade-off must be found between those specific constraints.

As substrates for this study, we will consider various oxides in order to address potential nucleation difficulties: Si wafers with 200 nm thermal SiO₂ and optionally coated with either SnO₂, Al₂O₃, or ZnO (by ALD). Anhydrous *n*-hexane is used as the purging solvent. In a first control experiment, no solid is deposited when a hexane solution of Hf(NMe₂)₄ flows over the substrate in the absence of complementary reagent for the equivalent of 30 sALD cycles. Fig. S3 in the ESI† exhibits the formation of a layer on the order of 0.1 nm thickness, corresponding to the one adsorbed monolayer of Hf complex that must be expected. This demonstrates that Hf(NMe₂)₄ does not decompose thermally or in the presence of adventitious water.

For deposition tests, 1 mM Hf(NMe₂)₄ and 2 mM H₂S *n*-hexane solutions prepared under inert atmosphere are injected into the chamber in alternating manner, after a preliminary flush with pure solvent. One standard sALD cycle is defined as follows: (i) Hf(NMe₂)₄ solution is pumped into the chamber for 10 s. (ii) The chamber is purged with *n*-hexane for 60 s. (iii) The H₂S solution is pumped for 10 s. (iv) The chamber is purged again for 60 s. After the desired number of cycles, *n*-hexane is used for a final purge for 3 minutes. After a 100-cycle test, the thickness of the layer determined by spectroscopic ellipsometry is 5.8 nm (Fig. 2). Samples look homogeneous even after 410 cycles (Fig. S4a†).

The demonstration of self-limiting surface chemistry which defines ALD growth is provided by a series of tests in which the precursor pulse duration is varied (while the purge duration is maintained, Fig. 3). The saturating behavior is found for pulses of 10 to 20 s, proving ALD growth. For extreme pulse durations, the purge no longer suffices to prevent uncontrolled deposition (of the chemical vapor deposition or chemical bath deposition type, last datapoint of Fig. 3 and Fig. S4b†). The linear growth behavior is demonstrated on the four types of substrates tested in Fig. 4. In this experiment, care must be taken to reproduce the same placement of substrates in each

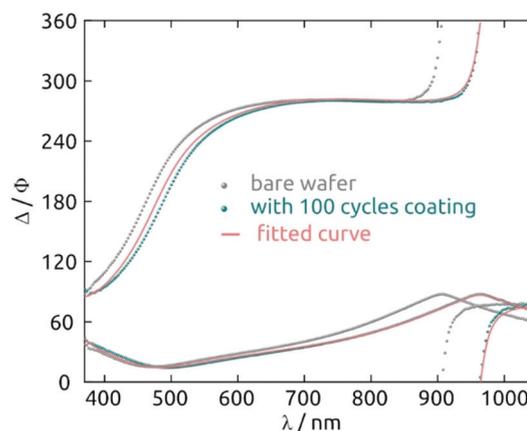


Fig. 2 Hafnium sulfide film thickness characterization: spectroscopic ellipsometry data obtained from a bare silicon wafer (in gray), 100 cycles HfS₂ coated wafer (in dark cyan), and curves fitted with an air/HfS₂/SiO₂/Si stacks model (in pink).

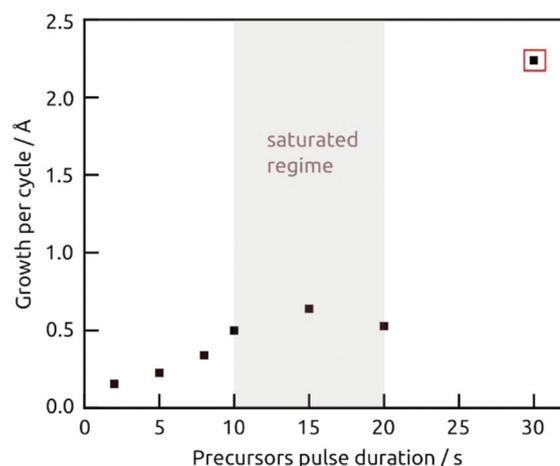


Fig. 3 Growth rate as a function of the precursors pulse duration, the concentrations for Hf(NMe₂)₄ and H₂S are 1 mM and 2 mM, respectively, and the solvent purging is 60 s for each half cycle. Note that the thickness for the marked data point was measured after removing the dusty precipitate with sonication in hexane.



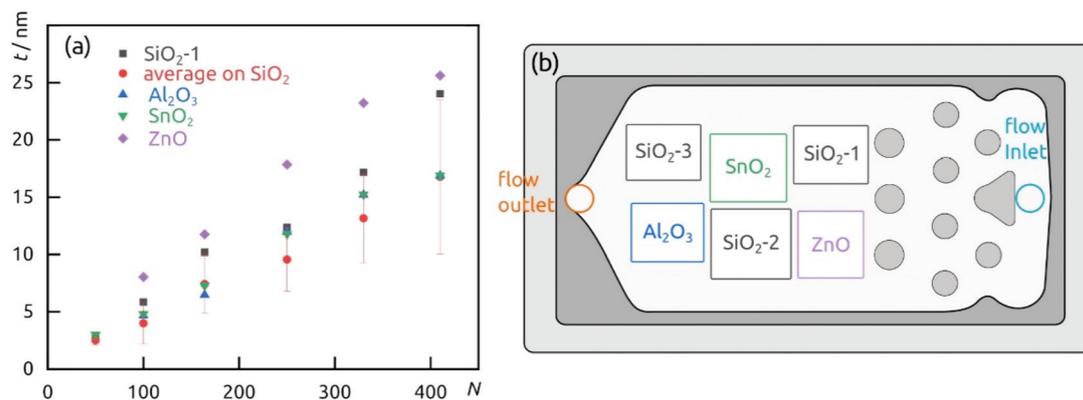


Fig. 4 (a) Film thickness dependence on the number of cycles performed on different substrates. (b) Wafers distribution in the reaction chamber.

individual run (Fig. 4b) in order to account for deviations in growth rate due to placement and substrate chemistry. Although all substrates exhibit linear growth, the growth rate varies significantly, with samples situated close to the inlet experiencing more deposition. This observation indicates that the flow dynamics of the microfluidic chamber design can be improved further. In addition to this, ZnO seems to give rise to a slightly faster growth, perhaps associated with its higher surface roughness.

Characterization of the deposit

The morphology of a 24 nm HfS₂ film deposited on a bare wafer is continuous with some roughness observable in scanning electron micrographs, Fig. 5. X-ray diffraction measurements performed in grazing incidence do not yield any hint of a crystalline structure (Fig. S6†). We conclude that the deposit is amorphous, as expected given the very low processing temperature.

A strong indication for the identity of the deposit as HfS₂ is provided by the spectroscopic ellipsometry data. Fig. 6 exhibits the $n(\lambda)$ and $k(\lambda)$ spectra obtained from the fit to the experimental data. As a comparison, we also provide the database spectra of HfO₂. The material deposited is clearly not HfO₂. Not only do the refractive indices diverge, especially in the UV, but the solid deposited with our method absorbs strongly over

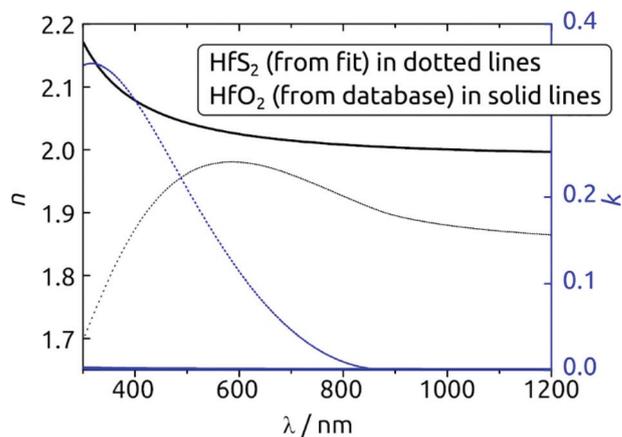


Fig. 6 Refractive index and extinction coefficient spectra obtained for the HfS₂ deposition (dotted lines), as compared with the database reference values for HfO₂ (solid lines). The clearly distinct curves furnish a strong argument for the chemical identity of the deposit.

a large fraction of the visible range, whereas HfO₂ is of course perfectly transparent. The bandgap obtained from the Tauc-Lorentz optical model is 1.4 eV, a value smaller than that reported for crystalline HfS₂ (1.8 eV)³⁶ but possible given the amorphous nature of our material and the range of values computed for different configurations of HfS₂.³⁷

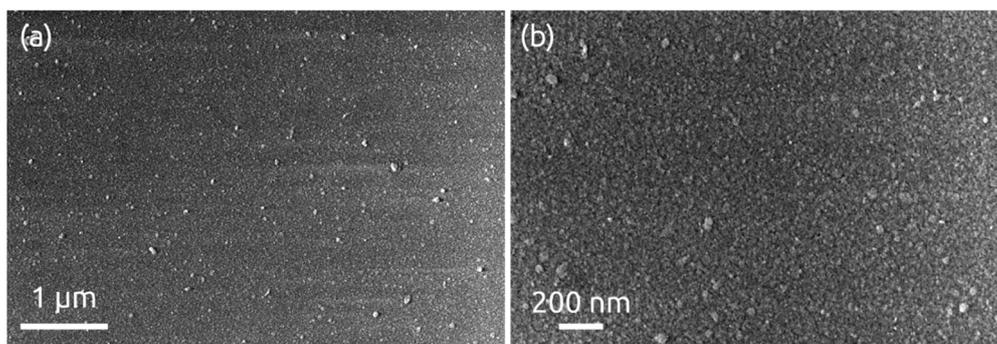


Fig. 5 SEM images of 24 nm of HfS₂ deposition grown on a bare Si/SiO₂ wafer at (a) low and (b) high magnification levels.



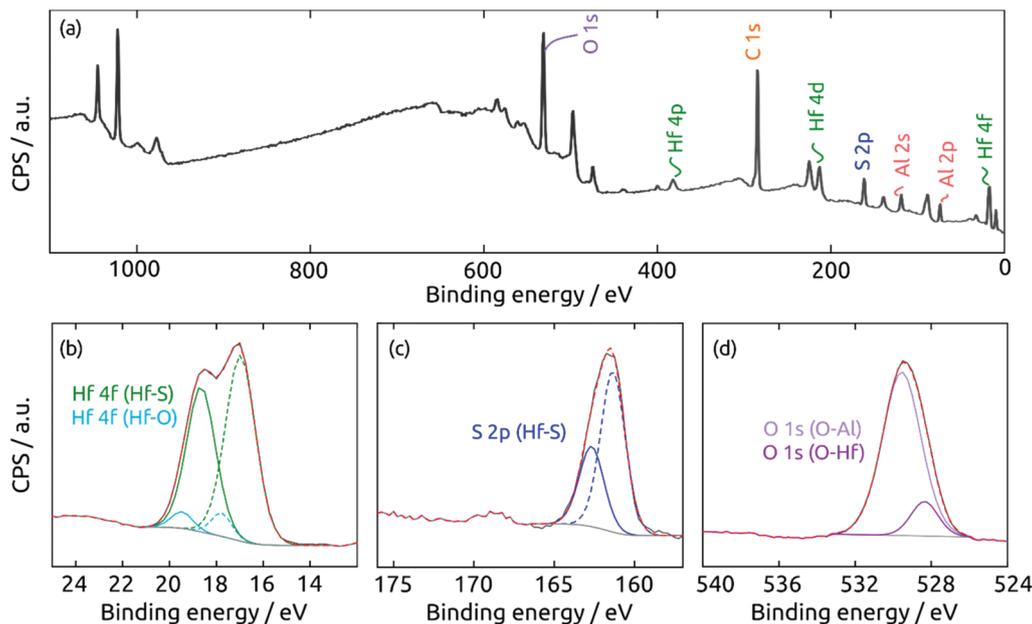


Fig. 7 XPS spectra of a HfS_2 film on a Al_2O_3 substrate. (a) Survey spectrum, (b) Hf 4f, (c) S 2p, and (d) O 1s regions.

The surface composition of the coating exposed to air is determined by X-ray photoelectron spectroscopy (XPS). The survey spectrum (Fig. 7a) of a $\text{HfS}_2/\text{Al}_2\text{O}_3/\text{SiO}_2/\text{Si}$ sample exhibits the desired elements Hf and S, together with Al and O from the substrate. Not surprisingly, the surface is partly oxidized in air. The high-resolution Hf 4f region in Fig. 7b can be deconvoluted into two doublets (Hf 4f_{7/2}/4f_{5/2}) centered at 17.0 and 18.6 eV and at 17.8 and 19.3 eV, which can be ascribed to Hf bonded to sulfide and to oxide, respectively. The S 2p region in Fig. 7c can be fitted to two components at 161.3 and 162.7 eV, which are associated with the 2p_{3/2}/2p_{1/2} doublets of sulfide in HfS_2 . Finally, the O 1s region in Fig. 7d confirms the presence of HfO_2 (O 1s at 528.4 eV) and Al_2O_3 (529.6 eV). To sum up, the XPS measurements prove the success of HfS_2 growth by sALD, and remind us of the highly oxyphilic nature of the element Hf.^{16,38–40}

Discussion

A comparison of the HfS_2 sALD results with our gALD process published recently provides interesting insight.

The growth rate in this sALD is 0.4 Å–0.6 Å per cycle at room temperature. Compared to the gALD process (1.2 Å per cycle at 100 °C in gALD),¹⁶ this value is significantly smaller. This result points to an influence of the solvent on the surface reactions. A molecular-level explanation of this effect is suggested by the somewhat unusual shape of the growth rate dependence on temperature in gALD (shown in Fig. 3 of that paper).¹⁶ Indeed, a constant growth rate of 1.2 Å per cycle from 65 °C to 100 °C is followed by another plateau from 130 °C to 180 °C at the lower value 0.6 Å per cycle in gALD. This behavior contrasts with the most common curve, which typically com-

bins a plateau up to the decomposition temperature of a precursor, after which the growth rate increases. The facts that in gALD (a) two seemingly self-saturating regimes exist, and (b) the high-temperature regime exhibits half of the growth rate obtained at lower temperature may be related to the formation of a $\text{Hf}(\text{NMe}_2)_4$ dimer, the stability of which is observed in DFT computations (ESI of the publication).¹⁶ This dimer could form on the surface during gALD growth at low temperature and be responsible for the elevated growth rate. It would, however, be disrupted by temperatures in excess of 100 °C. Interestingly, the formation of this dimer seems to also be prevented at low temperature by the presence of solvent.

On a more general level, our results show that obtaining crystalline TMDCs at low temperature remains an elusive target. Furthermore, the extreme air sensitivity of HfS_2 may prevent it from being the best suited model system in which to study the details of nucleation and growth. However, the fact that one ALD reaction can be performed both from the gas phase and the liquid phase demonstrates the continuity that exists between sALD and gALD, and it opens new avenues of research towards low-temperature deposition of crystalline 2D materials. Indeed, the fine-tuning of interfacial energies is crucial for the control of the film morphology, and it can be adjusted best from the liquid phase. To this goal, the choice of solvents, the variation of ligands in the metal-organic precursors, and the use of additives such as surfactants represent tools that are available to the experimentalist from the gas phase only.

Conclusions

In this study, we developed an sALD method for the generation of HfS_2 thin films from $\text{Hf}(\text{NMe}_2)_4$ and H_2S as precursors. The



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