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Understanding metal organic chemical vapour deposition of monolayer WS₂: the enhancing role of Au substrate for simple organosulfur precursors†

 Ye Fan, ^{‡a} Kenichi Nakanishi, ^{‡a} Vlad P. Veigang-Radulescu,^a Ryo Mizuta,^a J. Callum Stewart,^a Jack E. N. Swallow,^b Alice E. Dearle, ^a Oliver J. Burton,^a Jack A. Alexander-Webber, ^a Pilar Ferrer,^c Georg Held,^c Barry Brennan,^d Andrew J. Pollard,^d Robert S. Weatherup ^b and Stephan Hofmann ^{*a}

We find that the use of Au substrate allows fast, self-limited WS₂ monolayer growth using a simple sequential exposure pattern of low cost, low toxicity precursors, namely tungsten hexacarbonyl and dimethylsulfide (DMS). We use this model reaction system to fingerprint the technologically important metal organic chemical vapour deposition process by *operando* X-ray photoelectron spectroscopy (XPS) to address the current lack of understanding of the underlying fundamental growth mechanisms for WS₂ and related transition metal dichalcogenides. Au effectively promotes the sulfidation of W with simple organosulfides, enabling WS₂ growth with low DMS pressure (<1 mbar) and a suppression of carbon contamination of as-grown WS₂, which to date has been a major challenge with this precursor chemistry. Full WS₂ coverage can be achieved by one exposure cycle of 10 minutes at 700 °C. We discuss our findings in the wider context of previous literature on heterogeneous catalysis, 2D crystal growth, and overlapping process technologies such as atomic layer deposition (ALD) and direct metal conversion, linking to future integrated manufacturing processes for transition metal dichalcogenide layers.

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Introduction

Tungsten disulfide (WS₂) is a prominent material of the transition metal dichalcogenide (TMDC) family, with a characteristic layered structure of hexagonally packed metal atoms sandwiched between two layers of chalcogen atoms. Monolayer WS₂ exhibits a wide direct bandgap ($E_g \approx 2$ eV), strong spin-orbit coupling and bright room-temperature photoluminescence (PL).^{1–4} As a prototypical 2D semiconductor, WS₂ has raised promise in applications ranging from (opto-) electronics to spintronics and sensors.^{1–4} These emergent applications drive the need for scalable growth of atomically thin, highly crystalline “electronic-grade” 2D layers/films,⁵ which is distinct from

the historic focus on TMDC bulk crystal growth,⁶ and from classic TMDC application areas such as lubricants or catalysis.⁷ A large body of recent literature highlights the progress in such “electronic-grade” TMDC layer growth.⁵ However, the vast parameter space and increased complexity in structure and possible mechanisms, when compared to graphene for instance, leaves the understanding of even the basic processes of TMDC layer growth still severely limited.

The currently most prevalent TMDC growth method employs powder vaporisation, typically based on the sublimation of solid precursors in a hot wall reactor, and has demonstrated large monolayer TMDC crystal domains (>100 μm) on inert substrates like SiO₂/Si,⁸ and sapphire,⁹ as well as on metals such Ni–Ga alloys¹⁰ and Au.^{11,12} For the latter, growth has been referred to as catalytic,¹¹ analogous to graphene and nanowire growth.^{13,14} However, the growth atmosphere is difficult to accurately control and modulate, which restricts uniformity and scalability as well as makes such powder based techniques too complex, as model systems, to advance the required fundamental understanding of crystal growth. Driven by the need for low-cost, high through-put and industrial compatibility, metal–organic chemical vapour deposition (MOCVD) and related atomic layer deposition (ALD) type tech-

^aElectrical Engineering Division, Department of Engineering, University of Cambridge, UK. E-mail: sh315@cam.ac.uk

^bDepartment of Materials, University of Oxford, Parks Road, Oxford OX1 3PH, UK

^cDiamond Light Source Ltd, Harwell Science and Innovation Campus, Didcot OX11 0DE, UK

^dNational Physical Laboratory, Hampton Rd, Teddington, Middlesex, TW11 0LW, UK

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‡Contributed equally.



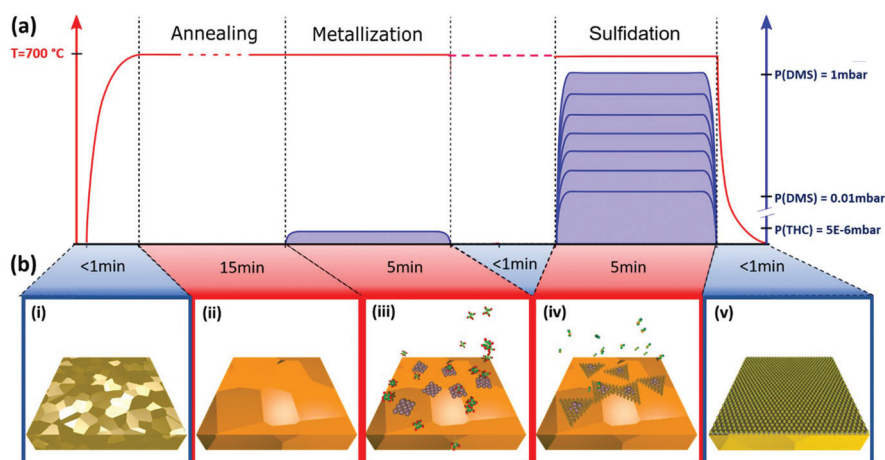


Fig. 1 (a) Schematic of 'deconstructed' MOCVD procedure: Au substrate is heated up to growth temperature in vacuum, followed by exposure to THC, referred to as metallization, and separate exposure to DMS, referred to as sulfidation. (b) Schematic of basic processes: Au foil before loading to reaction chamber (i), grain growth of Au (ii), W deposition (iii), WS_2 nucleation (iv) and domain growth into monolayer WS_2 film (v).



Fig. 2 Characterizations of MOCVD WS_2 film. (a) Raman spectroscopy of WS_2 film transferred onto SiO_2/Si . (i) Typical Raman spectrum normalized by the Raman peak of Si. Measurement data (black dots) overlaps well with the fitted curve (blue). Some major modes (2LA, E_{2g} and A_{1g}) among all the Raman peaks (red) are labelled. (ii) Raman mapping of A_{1g} peak height. (b) PL characterization of WS_2 film transferred onto SiO_2/Si substrate. (i) Typical PL spectrum normalized by the Raman peak of Si. Measurement data (black dots) overlap well with the fitted curve (blue). The exciton (X) and trion (X^-) peaks are labelled. (ii) Integrated PL intensity mapping normalized by the integrated Raman intensity of Si. (c) HRTEM of the WS_2 film. Lattice constant is measured as ~ 3.2 Å. (d) SAED of WS_2 film. The sharp 6-fold diffraction pattern indicates good crystallinity. The inset shows bright field TEM image of the SA. (e) TEM diffraction mapping of WS_2 film. The average grain size of WS_2 of the mapped area is around 10 μm , with a bimodal crystal orientation with 30° rotation (FWHM $< 1^\circ$).

Fig. 2 shows the characterization of a WS_2 film grown by our deconstructed MOCVD process with 0.1 mbar DMS during the sulfidation step. The WS_2 layer is transferred onto a SiO_2/Si wafer support for Raman and photoluminescence (PL) measurements, and Quantifoil support for TEM. Fig. 2(a, i) shows a representative Raman spectrum of the WS_2 with E_{2g} and A_{1g} vibration modes measured at 356.9 cm^{-1} and 417.6 cm^{-1} , respectively, and with a full-width-half-maximum (FWHM) of 4.3 cm^{-1} and 4.2 cm^{-1} , respectively. This is in line with previously reported Raman measurements using 532 nm

laser excitation for WS_2 transferred onto SiO_2 .^{44–46} Fig. 2(a, ii) shows a map of A_{1g} peak height over 108 $\mu m \times 154$ μm area, which confirms the continuity of the as-transferred WS_2 layer. The monolayer nature of as-grown WS_2 is further confirmed by the strong PL peak with an exciton energy of ~ 2.02 eV (Fig. 2 (b, i)). Fig. 2(b, ii) shows a map of normalized integrated PL intensity over the same area as the Raman mapping in Fig. 2 (a, ii). Despite some fluctuations, the PL intensity of WS_2 film is always orders of magnitudes stronger than the Si Raman signal (also stronger than the WS_2 Raman signal), consistent



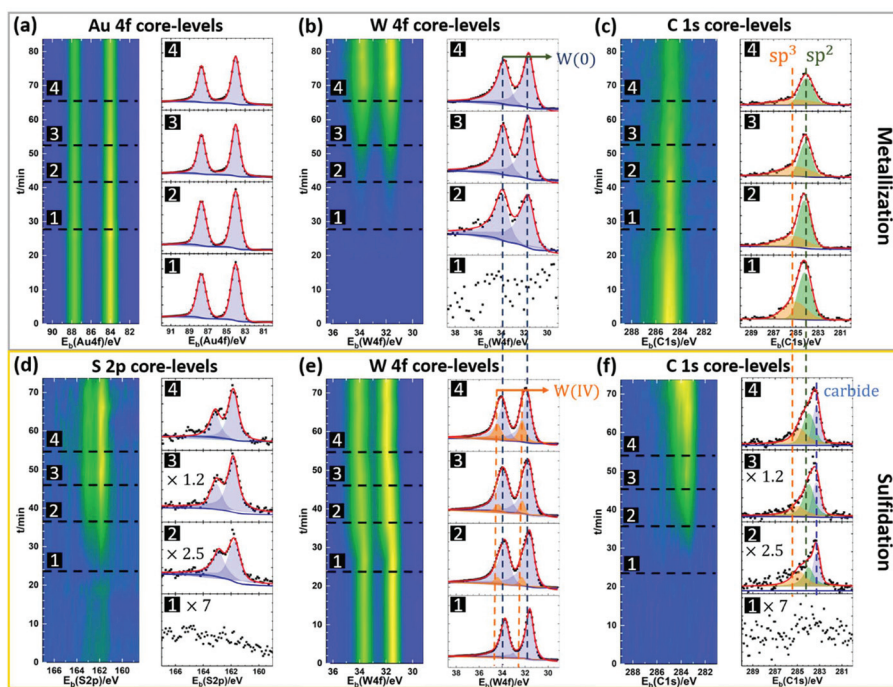


Fig. 3 Operando XPS of the MOCVD process. (a)–(c) Time-resolved Au 4f, W 4f and C 1s core-level spectra, respectively, during metallization step. (d)–(f) Time-resolved S 2p, W 4f and C 1s core-level spectra, respectively, during the sulfidation step. From (a) to (f), the XPS intensity map over reaction time is presented on the left, while the selected spectra of each element showing different reaction stages are presented on the right.

with the monolayer character of the WS_2 and the absence of WS_2 multi-layer regions.³ Similar to previous reports on CVD/MOCVD WS_2 , a significant trion peak around 1.98 eV is detectable.⁴⁷

Fig. 2(c)–(e) show high resolution transmission electron microscope (HRTEM) images and diffraction mapping results for a transferred WS_2 film. The representative HRTEM image in Fig. 2(c) highlights the crystallinity of the WS_2 , confirming a hexagonal lattice structure. The lattice constant is approximately 3.18 Å, which is consistent with both theory,^{48,49} and previously reported measurement results.^{50,51} The selected-area electron diffraction (SAED) analysis in Fig. 2(d) shows a matching clear hexagonal diffraction pattern with sharp spots. In order to determine the crystal orientation over a reasonably large area, over 200 diffraction patterns across a $40 \mu\text{m} \times 40 \mu\text{m}$ WS_2 film area were taken to construct the map shown in Fig. 2(e). We find two dominating crystal orientations rotated by $30^\circ \pm 1^\circ$ as highlighted by the bimodal distribution (Fig. 2(e)). Although the influence of crystal orientation of the Au to the crystal orientation of WS_2 is not our focus in this report, we are aware of the possible heterogeneity in WS_2 arising from the poly-crystallinity of the Au substrate. The data indicates, however, an average WS_2 grain size of the order of 10 μm , which compares favourably to previous literature on MOCVD TMDC materials on non-catalytic substrates with typical domain sizes on the order of a few micrometres.²⁰ The representative transfer curve of a two-terminal WS_2 field effect transistor (FET) device structure is shown in Fig. S3 in ESI† with a positive light response. The MOCVD WS_2 shows n-type

behaviour with a field-effect mobility of approximately $0.5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, which is comparable to previously reported values.^{11,52}

In this paper, we focus on providing insight into the reactions during the ‘deconstructed’ MOCVD process and identifying the key parameters for WS_2 crystal growth control. Fig. 3 shows time-resolved *operando* XPS of the metallization and sulfidation steps. Fig. 3(a)–(c) show the Au 4f, W 4f and C 1s core level data during the metallization step. In Fig. 3(a), the Au 4f_{7/2} core level energy is fixed to 84.0 eV allowing calibration of the other spectra. The peak position of the Au 4f_{5/2} was 87.7 eV, consistent with a spin-orbit splitting of 3.7 eV.⁵³ During the metallization process, the intensities of the Au 4f peaks decrease over time, while the intensities of W 4f peaks increase, (Fig. 3(a) and (b)) indicating the deposition of W onto the Au surface. The binding energies (BEs) of the W 4f doublet are measured at ~ 31.5 eV and ~ 33.7 eV for the W 4f_{7/2} and W 4f_{5/2}, respectively. The long-tail character of the W doublet peaks are fitted well by Doniach–Šunjić (DS) line-shapes, which confirms that the deposited tungsten is metallic.⁵⁴ We note that a W 5p_{3/2} component is expected around 5.5 eV to 6 eV above the W 4f_{7/2}, but is of insufficient intensity to be resolved above the tail. To rule out the formation of tungsten carbide, the C 1s core-level is measured during the metallization process. The broad peaks at 284.2 eV (assigned to sp² carbon) and at 285.3 eV (assigned to sp³ carbon) can be detected prior to exposure as shown in Fig. 3(c), consistent with amorphous carbon (a-C) species.⁵⁵ The C 1s peak associated with tungsten carbide is expected to have a distinctly lower BE (~ 283.3 eV) than what is measured during the metal-



lization process, indicating an absence of tungsten carbide at this stage.⁵⁶ Additionally, during THC exposure the intensity of the C 1s peak decreases without significant change in shape or position of the peak, indicating that the carbon comes from initial Au surface contamination, rather than being introduced by THC dissociation.

Fig. 3(d)–(f) shows the S 2p, W 4f and C 1s core level spectra during the sulfidation step. The formation of WS₂ is marked by an increase in S 2p doublet peak intensity with the BEs of 161.8 eV (S 2p_{3/2}) and 163.0 eV (S 2p_{1/2}), respectively. These BEs agree with previously reported values for WS₂.⁵⁷ The conversion of W to WS₂ is also marked by the shift and broadening of the W 4f lines during sulfidation, as shown in Fig. 3(e) and Fig. S5 in ESI.† W core level signatures from both metallic tungsten and WS₂ are observable during sulfidation. The W 4f peaks for WS₂ are measured at 32.2 eV for W 4f_{7/2} and 34.4 eV for W 4f_{5/2}, with their intensities increasing during DMS exposure, as shown in Fig. 3(e). It is interesting to note that only a fraction of W transforms into WS₂ even for extended DMS exposure (Fig. 3(e)). This indicates the self-limiting nature of the tungsten sulfidation reaction on the surface of Au. Since the W 4f peaks for metallic tungsten overlap strongly with those for carbide,⁵³ we do not separate them when fitting the W 4f signals. The metal carbide and other carbon contamination in as-grown WS₂ is monitored by the C 1s signal (Fig. 3(f)), which clearly increases during sulfidation. We notice a significant C 1s core level signal around 283.3 eV, in addition to the two peaks attributed to a-C in Fig. 3(c). We assign this to tungsten carbide with a distinctively lower BE. This indicates the formation of metal carbide as a side reaction and confirms DMS as the source of carbon contamination for the WS₂ MOCVD process studied here. Although the sulfidation of W is self-limiting, the deposition of a-C contamination proceeds with ongoing DMS exposure (ESI, Fig. S4†).

To further study the role of the Au surface in the MOCVD process, we compare the growth result of WS₂ on bare Au to that on monolayer hexagonal boron nitride (hBN) covered Au

(Fig. 4). The motivation thereby is for the hBN to suppress direct gas precursor contact with the Au surface. hBN has been used as growth substrate in previous reports and is known remain intact after the MOCVD reaction.⁵⁸ We also expect hBN to affect adatom mobilities. Monolayer hBN was grown by CVD on Pt and transferred onto the Au using PMMA.⁵⁹ Post-growth ToF-SIMS imaging is used to map the growth results, as shown in Fig. 4(b). Fig. 4(b, i) shows a 500 μm × 500 μm map of the detected B⁺ ion species, that act as a fingerprint for the presence of the hBN film. The B⁺ map indicates continuous hBN coverage on Au after WS₂ growth and confirms that the hBN film remains intact throughout the MOCVD process. Fig. 4(b, ii) shows a map of the detected WS₃⁻ ion species, used to fingerprint the presence of WS₂. We find that a WS₂ film only forms on the Au surface without hBN coverage. The unreacted W will be oxidized in air at the post-growth stage, so tracking the location of the WO₃⁻ ion species enables us to separate the signals of unreacted W and WS₂, as shown in Fig. 4(b, iii). W is found on both the hBN covered and uncovered Au surface. This is consistent with the high chemical reactivity of the THC precursor and its ready thermal decomposition at the metallization temperature.⁴³ The more sporadic W coverage on the hBN can be due to increased desorption and/or increased W adatom mobility. To rule out the influence of polymer residues on the hBN to the decomposition of DMS, Fig. 2(b, iv) presents a map of the detected C₂⁻ ion species on the surface as a fingerprint for such carbon contamination. No difference in the average carbon level between the hBN covered and uncovered Au surface is observed. We thus rule out polymer residuals as a cause of the different WS₂ growth behaviour on Au and hBN covered Au. Moreover, this result also highlights that although the sulfidation rate is promoted by Au, side reactions such as the formation of carbides (see Fig. 3f) are similar on the different substrates.

Fig. 5 shows Raman spectroscopy and ToF-SIMS data, highlighting the influence of DMS partial pressures during sulfidation on the carbon contamination of as-grown WS₂ on Au.

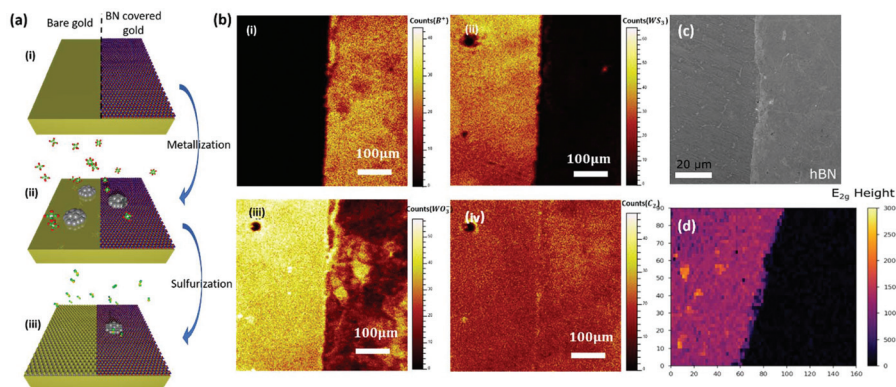


Fig. 4 WS₂ growth on Au partially covered by monolayer hBN. (a) Schematic of sample (i) and MOCVD procedure with metallization (ii) and sulfidation (iii). (b) Select ToF-SIMS surface images of the sample after complete MOCVD process: (i) B⁺ image highlighting hBN coverage; (ii) WS₃⁻ image highlighting WS₂ coverage; (iii) WO₃⁻ image to characterize W coverage; (iv) C₂⁻ image to characterize residual carbon. (c) SEM image of sample. No WS₂ is found to grow on hBN covered Au. (d) Raman intensity map of sample. E_{2g} height is mapped near the edge of hBN after MOCVD of WS₂.



dissolution effects are not significant to such WS_2 growth. Extended THC exposure, *i.e.* more W deposition, does not lead to multi-layer WS_2 growth. Rather, we observe an increasing number of particles (see ESI, Fig. S6†) in addition to the WS_2 monolayer. Previous STM work on epitaxial MoS_2 nucleation on Au(111) highlights the presence of amorphous Mo clusters as a result of insufficient sulfidation.³² In analogy we propose excess W as the source of the observed particles here. This is consistent with the SIMS mapping in Fig. 4 showing W clusters on hBN covered Au. Since W can be detected on both hBN covered Au and bare Au, we determine that the THC decomposition is not surface selective in our reaction conditions. The W clustering indicates a reasonably high W surface mobility, which is required to achieve large WS_2 crystal domain sizes. We observe no W sulfidation on top of the hBN (Fig. 4). Further, we note that as the tungsten thickness is increased the sulfidation to WS_2 at the same conditions becomes less complete, and for W foil ($\sim 25 \mu\text{m}$), we can detect no 2D WS_2 by Raman after sulfidation. This clearly highlights the role of the exposed Au surface in promoting sulfidation of W by DMS to form WS_2 . In contrast to H_2S , which has been shown to promote metal sulfidation on top of an existing TMDC layer, thereby leading to multi-layer TMDC growth,³² DMS presents a lower effective sulfur chemical potential, consistent with the lower TMDC growth rates that have been observed compared to when H_2S is used.³⁸ For our sulfidation process Au allows a significant lowering of the partial pressure of the DMS while still enabling high WS_2 growth rates so that full coverage can be achieved with <10 min sulfidation cycles. Importantly, WS_2 growth in our process is “self-limited” to monolayer thickness. This can be rationalised with the known high affinity of Au for organosulfide adsorption leading to selectively enhanced concentrations on the Au surface.⁴² In turn, less efficient chemisorption and high precursor desorption can be expected for inert surfaces like hBN. Thus, with the lower effective sulfur chemical potential of DMS, the supersaturation required for WS_2 nucleation is only satisfied on the Au surface in our reaction conditions. Once a layer of WS_2 grows and covers the Au, it prevents the DMS molecules from reaching the Au, similar to the hBN layer in Fig. 4. This leads to the observed self-limiting WS_2 growth behaviour.

Thermal decomposition of DMS is reported to be slow and incomplete at our reaction temperature of $700 \text{ }^\circ\text{C}$.^{60,61} The relatively fast WS_2 growth time that we observe is indicative of a clear enhancing effect of the metallic Au surface for the sulfidation with DMS. This is consistent with the known strong chemisorption of organosulfides on Au and their favourable dissociation on Au reported in heterogeneous catalysis literature.^{40–42} There are a range of possible competing reactions including W carbide formation.⁶² Our *operando* XPS data highlights that although the sulfidation of W is self-limiting, the deposition of a-C contamination and formation of tungsten carbide proceeds with ongoing DMS exposure (Fig. 3). This is consistent with excess W being present on top of the formed WS_2 layer, similar to excess Mo clusters reported in previous STM work on top of formed MoS_2 layers on Au.³² It

appears that without direct contact to Au, the DMS exposure of excess W results in a different reaction pathway dominated by carbide/surface carbon formation. The result of DMS exposure of W clusters will further depend on their support, as indicated by the different behaviour on hBN (Fig. 4).

Conclusions

We use a deconstructed MOCVD model process to understand the detailed chemical reaction during MOCVD process and discover the significant enhancing role of Au on the sulfidation step, which results WS_2 monolayer growth with a simple organosulfide like DMS. This opens the possibility of using less-toxic precursors for MOCVD and ALD-type reactions, while achieving low carbon contamination and fast growth times. The much lower chalcogen precursor pressure not only yields self-limiting monolayer WS_2 growth but also makes the overall process much more efficient and gives wider compatibility with standard growth reactors. The reaction scheme also opens future *operando* process characterisation opportunities, to further progress the understanding of the underlying mechanisms. *Operando* characterisation has been crucial to advancing the understanding and thus control of graphene and hBN film growth,¹³ and we expect our results and the growth model presented here to form the basis for more detailed understanding of TMDC MOCVD. In analogy to catalytic graphene growth on Pt, this will include metal surface orientation dependent growth kinetics, and links to the challenge of cost-efficient and scalable metal substrate preparation.^{63,66} While we here focus on Au, which requires subsequent WS_2 transfer for typical opto-electronic device applications, our findings also motivate further studies on catalytic enhancement of dielectric support films, which would allow direct, transfer-free device integration. In contrast, for electro-chemical applications TMDC layers grown directly on Au can serve as powerful model system for new approaches to catalyst design.

Methods

‘Deconstructed’ MOCVD growth of WS_2 on Au

$25 \mu\text{m}$ Au foils with 99.985% purity from Alfa Aesar were used as the growth substrates. All as-received Au substrates were annealed for 6 hours at $1025 \text{ }^\circ\text{C}$ under total pressure of 800 mbar with $H_2 : Ar = 1 : 9$. Right before the growth, samples were treated by oxygen RIE (150 mTorr, 50 W, 5 min) to remove surface carbon contamination. All *ex situ* MOCVD growth was carried out in a cold-wall low pressure MOCVD furnace. An 808 nm continuous wave (CW) IR laser was used for sample heating. Sample temperature during growth was monitored by $1.6 \mu\text{m}$ IR pyrometer assuming transmission 0.9 and emissivity 0.2. The temperature measurement error was estimated as $\sim 50 \text{ }^\circ\text{C}$. In all experiments, the samples were heated to $700 \text{ }^\circ\text{C}$ under base pressure better than 3×10^{-6} mbar for 15 min to



anneal and stabilize the system. After the annealing, $W(CO)_6$ (99.9%+ purity from Strem Chemicals) was sublimed at 120 °C and fed into the system as the W precursor during metallization. The partial pressure of $W(CO)_6$ was controlled by leak-valve, which allow a pressure control precision better than 5×10^{-7} mbar. During the metallization, 5×10^{-6} mbar $W(CO)_6$ was introduced to the chamber and exposed to Au foil at 700 °C for 5 min. After this, 5.5×10^{-3} mbar Ar is introduced to stabilize the base pressure when $S(CH_3)_2$ (99%+ purity from Sigma Aldrich) was fed into the system as S precursor during sulfidation. The partial pressure of $S(CH_3)_2$ was also controlled by a leak-valve. $S(CH_3)_2$ with controlled partial pressure ranging from 0.01 mbar to 1 mbar was introduced for the sulfidation of different samples to compare its influence on WS_2 produced. After sulfidation, the sample is cooled down rapidly for further characterization.

CVD/transfer of hBN

hBN was grown on 25 μ m thick platinum (Pt) foils (99.99%, Alfa Aesar) on a 0.5 mm tantalum (Ta) foil support as previously reported.⁵⁹ Prior to loading, the Pt foils were cleaned by sonication in acetone followed by iso-propanol and dried with a nitrogen gun. The growth recipe consisted of heating the sample in base pressure $\sim 2 \times 10^{-6}$ mbar to 1000 °C then annealing at temperature for 5 minutes. Subsequently, 1×10^{-5} mbar partial pressure of borazine (>97%, Fluorochem) was dosed into the chamber *via* a manually operated leak valve to initiate isothermal growth for 5 minutes. A continuous monolayer BN is produced on Pt by this method. The sample was rapidly cooled under borazine atmosphere.

Electrical delamination, or ‘bubbling’, transfer of hBN is used to peel BN off from Pt.⁶⁴ PMMA (A4, 950k) was spun on the as grown hBN on Pt at 3000 rpm for 45 s as the scaffold layer. PMMA/BN is then peeled off from the Pt by hydrogen bubbling generated by the electrolysis of water in 0.2 M Na (OH) (aq.). The film is then rinsed and fished up by a piece of Au foil to cover half of the surface. After drying overnight, the Au piece with semi-coverage of PMMA/BN is heated to 180 °C for 30 min to soften the PMMA then soaked in acetone to remove the PMMA scaffold.

Transfer of WS_2 on Au by wet etching

PMMA (A4, 950k) was spun on the as grown WS_2 on gold at 3000 rpm for 45 s. The sample is then baked at 120 °C for 3 min to evaporate the solvent of polymer. WS_2 covering the backside of gold is removed by 50 W, 20 s reactive ion etching (RIE) in 150 mTorr CF_4 . After the RIE, gold is etched by KI/I_2 based gold etchant (Sigma-Aldrich) within 4 hours. The sample is then scooped out and rinsed in DI water for several times before being fished out by the target substrate and dried.

Raman/PL

Both Raman and PL spectra of samples are taken by a Renishaw inVia Raman system. The excitation laser is 20 mW, 532 nm. Single spot Raman is measured with 1% laser power 2 s exposure and accumulated 6 times. For Raman mapping,

10% laser power, 0.1 s exposure is used for each spot with a step size of 2 μ m in both x and y direction. Single spot PL is measured by 0.1% laser power 1 s exposure and accumulated 2 times. For PL mapping, 1% laser power, 0.25 s exposure is used for each spot with a step size of 2 μ m in both x and y direction. The sample is kept in the same orientation during all spectroscopy measurements to make sure that the Raman intensity of the silicon reference peak is unchanged.

For the Raman data taken from WS_2 on Au, an Au baseline is taken separately and subtracted from all measured spectra. The integrated intensity for WS_2 is calculated from the peak intensity from 250 cm^{-1} to 500 cm^{-1} . The integrated intensity for a-C is calculated from the peak intensity from 1100 cm^{-1} to 1900 cm^{-1} .

TEM

Quanti-foil TEM grid is used as the support for WS_2 in the associated TEM studies. After transfer of the PMMA/ WS_2 onto the TEM grid, the sample is dried overnight then baked at 150 °C for 30 min to soften the polymer and increase the adherence of the WS_2 to the grid. The PMMA scaffold layer is then removed by dissolving in acetone. Tecnai Osiris TEM is used for imaging and diffraction. All measurement is taken under 80 keV acceleration voltage. For diffraction mapping, selected-area electron diffraction (SAED) is taken over a grid of spots across the large area being mapped. Crystal orientation of the sample is then constructed from relative rotation between the SAEDs.

ToF-SIMS

Ex situ ToF-SIMS measurement were performed using a ToF-SIMS IV instrument (ION-TOF GmbH, Germany) at a base pressure better than 5×10^{-9} mbar. 25 keV Bi_3^+ ions from the liquid metal ion gun (LMIG) are used for imaging. The LMIG spot size is less than 5 μ m in spectroscopy mode and is operated at a current of 0.1 pA. The surface maps (*i.e.* Fig. 4(b)) were acquired directly by randomly rastering the Bi_3^+ ion beam over the surface of the sample in an area of 500 μ m \times 500 μ m in both positive and negative ion mode. For larger area maps, successive 500 μ m \times 500 μ m images are stitched together to create the larger image. 2D depth profiles and 3D depth profile images are acquired by cyclically analysing a 150 μ m \times 150 μ m area (with 256 \times 256 pixel density and a mass spectra taken at each pixel) from the centre of a 400 μ m \times 400 μ m sputtered region during the course of depth profiling to mitigate crater edge effects on the generated spectra. These cyclically analysed images are then combined to create a 3D image and the 2D profile. For sputtering cycles, either 10 keV Cs^+ ions (100 μ s cycle time) with an ion current of 30 nA for bulk characterization, or 10 keV Ar_{2700}^+ (200 μ s cycle time) with an ion current of 0.36 nA from a GCIB for near surface region characterisation, were used. All ion beams are orientated at 45° to the sample surface. No charge compensation was used during these measurements. Data processing was carried out using Surface Lab (v.6.8.99996) software, by selecting relevant



peaks in the ToF-SIMS spectra and monitoring their change in intensity over the course of the sputter profiling.

XPS

Operando XPS measurements were performed at beamline B07-C of the Diamond Light Source (DLS), United Kingdom.⁶⁵ The environmental reaction chamber used in this study, or the 'T-cup', has a small (volume ~ 0.7 L) that allows sample temperature control between room temperature and ~ 1000 °C (*via* a button heater; HeatWave Labs) and local pressures between 10^{-5} mbar and 30 mbar. The sample cell is attached to a differentially-pumped hemispherical electron analyser (SPECS Phoibos 150 NAP) and a differentially pumped beamline interface which enables window-less exposure to synchrotron radiation in the photon energy range 170–2800 eV. Here, a photon energy of 700 eV was used for all measurements. The beamline exit slits were opened to 800 μm in the non-dispersive direction, and 100 μm in the dispersive direction for XPS, which results in a spot size around 75 μm \times 200 μm . The energy resolution of the setup is ~ 0.5 eV for the energies used herein, as determined from the width of the Gaussian component of Voigt functions fitted to Au 4f spectra of gold films. The base pressure is $\sim 8 \times 10^{-7}$ mbar before heating and $\sim 4 \times 10^{-6}$ mbar during heating. All core level spectra were acquired with a pass energy of 20 eV and an energy step size of 0.1 eV.

The BE scale for each spectrum was referenced to simultaneously measured Au 4f_{7/2} core levels, setting the peak position to 84.0 eV. All fittings were performed by using a Shirley background to model inelastically scattered electrons and applying either a Doniach–Šunjić (DS) line-shapes for metallic species, or a Gaussian–Lorentzian (GL) line-shapes for non-metallic species. We do not set artificial constraints on the FWHM of peaks during fitting. During the sulfidation process, we notice that the signal intensity is unstable due to the introduction of DMS. To compare the data recorded at different times, all peaks recorded during the sulfidation process were normalized by the W 4f peak intensity.

Conflicts of interest

There are no conflicts of interest to declare.

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