Large-Area Epitaxial Monolayer MoS₂

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ABSTRACT

Two-dimensional semiconductors such as MoS₂ are an emerging material family with wide-ranging potential applications in electronics, optoelectronics, and energy harvesting. Large-area growth methods are needed to open the way to applications. Control over lattice orientation during growth remains a challenge. This is needed to minimize or even avoid the formation of grain boundaries, detrimental to electrical, optical, and mechanical properties of MoS₂ and other 2D semiconductors. Here, we report on the growth of high-quality monolayer MoS₂ with control over lattice orientation. We show that the monolayer film is composed of coalescing single islands with limited numbers of lattice orientation due to an epitaxial growth mechanism. Optical absorbance spectra acquired over large areas show significant absorbance in the high-energy part of the spectrum, indicating that MoS₂ could also be interesting for harvesting this region of the solar spectrum and fabrication of UV-sensitive photodetectors. Even though the interaction between the growth substrate and MoS₂ is strong enough to induce lattice alignment via van der Waals interaction, we can easily transfer the grown material and fabricate devices. Local potential mapping along channels in field-effect transistors shows that the single-crystal MoS₂ grains in our film are well connected, with interfaces that do not degrade the electrical conductivity. This is also confirmed by the relatively large and length-independent mobility in devices with a channel length reaching 80 μm.

KEYWORDS: two-dimensional materials, MoS₂, epitaxial growth, electronic transport, grain boundaries, Kelvin probe force microscopy

The most investigated member of the transition metal dichalcogenide (TMD) family, molybdenum disulfide (MoS₂), has attracted widespread attention for a variety of next-generation electrical and optoelectronic device applications because of its unique properties. In the bulk, this material has a crystalline structure consisting of covalently bonded layers weakly coupled to each other by weak van der Waals (vdW) forces. Owing to the weak coupling, two-dimensional (2D) monolayer MoS₂ can be easily obtained by exfoliation using Scotch tape or liquid-phase exfoliation.2,3 Whereas bulk MoS₂ is a semiconductor with an indirect band gap of 1.2 eV,4 monolayer MoS₂ is a direct gap semiconductor with a band gap of at least 1.8 eV5–8 due to the 2D confinement. It also has Raman-active modes that are very sensitive to the thickness, which provides a convenient method for determining the number of layers with a reliable precision in addition to atomic force microscopy and optical detection techniques.9 The dramatic difference in the electronic structure of monolayer MoS₂ in comparison with its bulk counterpart offers many opportunities.

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for diverse applications. Using 2D crystals of MoS₂ mechanically exfoliated from bulk geological samples, versatile devices including field-effect transistors with high on/off current ratio, memory cells, ultrason sensitive photodetectors, and nanopores have been reported. Because MoS₂ has a high mechanical flexibility and breaking strength, all these devices can be implemented on flexible substrates. However, exfoliation of geological material and samples grown using chemical vapor transport lacks the systematic control of the thickness, size, and uniformity of the 2D film and is not scalable for large-scale device fabrication. Because of this, several methods, such as decomposition of thiophosphates and sulfuration of Mo metal or molybdenum oxides, have been exploited to synthesize MoS₂ on diverse substrates. Among them, chemical vapor deposition (CVD) is the most promising method to synthesize monolayer MoS₂ triangular islands tens of micrometers in size. In most of these reports, SiO₂ was used as the growth substrate, resulting in the random orientation of MoS₂ domains because of the amorphous nature of the substrate and its relatively high surface roughness. This inevitably results in a large concentration of grain boundaries that can be detrimental to the electrical and mechanical properties of the grown films over length scales exceeding several micrometers. In order to avoid this, it is necessary to control the crystallographic orientation of MoS₂ islands during growth so that they can coalesce into a uniform layer with a reduced density of grain boundaries.

Such control could be achieved using a suitable atomically smooth crystalline substrate. In the case of classical three-dimensional materials this is normally achieved using epitaxial growth. Recently, van der Waals epitaxy has shown its potential for circumventing the need for a lattice-matched substrate and still keeping an ordered relation with the substrate. Van der Waals interaction between the substrate surface and the grown material governs the formation of the initial nuclei (islands), which constitute the precursors of the layers. Only energetically stable nuclei are able to grow by lateral spreading through the lateral facets. This leads to an overall preferential orientation of the grown layers. The relatively low strength of the van der Waals interaction and the relaxed requirement for lattice matching for substrates and overlayers without dangling bonds make it difficult in practice to control the lattice orientation of the deposited films. This was the case in previously reported deposition of centimeter-scale monolayer MoS₂ films that have been deposited on other layered materials such as mica or graphene, where MoS₂ grains showed a wide distribution of orientations with respect to each other and the substrate lattice.

RESULTS AND DISCUSSION

Here, we use highly polished, EPI-ready grade sapphire substrates to achieve control over lattice orientation during CVD growth of monolayer MoS₂. Such crystals are commonly used as growth substrates for GaN growth in LED manufacturing and for the growth of III–V high electron mobility transistors. Even though MoS₂ and sapphire interact only via the relatively weak van der Waals interaction, commensurability of the sapphire lattice with MoS₂ allows the van der Waals interaction to control the lattice orientation of MoS₂ in a similar way to the lattice matching requirement in covalent semiconducting materials. While growth of MoS₂ and TMDs on sapphire substrates has been reported before, this is the first time control over lattice orientation with centimeter-scale uniformity has been achieved during the growth of a monolayer 2D semiconductor, and it opens the way to large-area growth of high-quality MoS₂. Key to achieving such lattice orientation is in the preparation of atomically smooth sapphire terraces on the surface of sapphire by annealing it in air for 1 h at a temperature of 1000 °C just prior to the growth process (Supporting Figure S1). An AFM-based analysis shows that the sapphire surface is characterized by atomically flat terraces ~50–70 nm wide with a step height of 0.22 nm due to a small miscut angle estimated to be ~0.2°. The terrace edges follow the (11̅20) direction on average. Annealed samples are transferred to the CVD system, and growth occurs. The growth process is based on the gas-phase reaction between MoO₃ and sulfur evaporated from solid sources using ultra-high-purity argon as the carrier gas.

The growth procedure results in characteristic single-crystal domains in the shape of well-defined equilateral triangles that merge into a continuous monolayer film covering a typical area of 6 mm × 1 cm in the middle portion of the growth substrate, Figure 1a. Figure 1b presents optical images of regions showing partial (top image) and almost full (bottom) coverage. A careful examination of a region with incomplete coverage, Figure 1c, reveals that most of the single-crystal domain edges are oriented along dominant directions. A reflection high-energy electron diffraction (RHEED) pattern is shown in the inset of Figure 1c. The appearance of streaks typical of 2D materials and the MoS₂ (1×1) pattern indicate the growth of a homogeneous and well-structured film over a large area. The majority (91.5% of islands shown in Figure 1c) of single-crystal domains are well aligned with the relative orientation of edges that can be expressed as multiples of 60°. This is confirmed by the orientation histogram presented in Figure 1f for the same area as in Figure 1c, showing that the dominant edge orientations are 0° and ±60°. A small fraction of domains (6% of islands in Figure 1c) show edges with a
relative angle of \( \pm 30^\circ \), while the rest (2.5%) show other orientations.

Figure 1d shows an AFM image of a single crystal of monolayer MoS\(_2\). According to previous STM studies, these MoS\(_2\) edges are expected to be (1010) Mo zigzag edges likely terminated with sulfur.\(^{36}\) The sapphire terrace step height of 0.22 nm is sufficiently low to ensure the growth of continuous MoS\(_2\) single crystals, as confirmed by TEM imaging and electrical transport measurements. Figure 1e shows a high-resolution TEM image of a suspended MoS\(_2\) film showing the crystallinity of the sample. View of the structural model is overlaid. Scale bar is 0.5 nm long. (f) Orientation histogram based on the area shown in part c confirms that the majority of MoS\(_2\) grain edges are oriented along \( 0^\circ \) and \( \pm 60^\circ \) angles.

The observed frequency difference confirms that the deposited material is monolayer MoS\(_2\). The ratio between the \( A_1 \) and \( E' \) mode intensities can be used as an indicator of doping levels,\(^{38}\) and it indicates that the CVD material is less doped than the exfoliated counterpart. Another indication of a smaller structural disorder\(^{39}\) in the CVD material is the smaller width of the \( A_1 \) mode.

Figure 2b shows a photoluminescence spectrum acquired at room temperature on CVD-grown and adhesive tape exfoliated monolayer MoS\(_2\). We can clearly resolve the intense A excitonic peak at 659 nm (1.88 eV), while the B exciton is not expected to be visible at low excitation intensities due to state-filling effects.\(^{40}\) Typical peak widths are \(~26\) nm (~72 meV) and are smaller than in exfoliated MoS\(_2\) samples (~40 nm or ~111 eV), indicating that our CVD MoS\(_2\) has superior optical qualities to the exfoliated material. Relative photoluminescence (PL) intensities also indicate a reduced doping level in the CVD material. Detailed photoluminescence mapping of single domains did not resolve any internal structure, indicating the absence of internal grain boundaries.

Figure 1. Monolayer MoS\(_2\) growth with controlled lattice orientation. (a) Photograph of centimeter-scale monolayer MoS\(_2\) grown on sapphire. (b) Optical microscopy images from different regions of the sample showing incomplete coverage close to the edges (upper image, position marked by the circle in a) and full coverage close to the center of the growth substrate (lower image, position marked by the rectangle in a). Scale bar length is 20 \( \mu \)m on the top image and 10 \( \mu \)m for the bottom image. Original optical images were converted to grayscale, and the contrast was enhanced. (c) Optical microscopy image of monolayer MoS\(_2\) grains grown on atomically smooth sapphire. Scale bar length is 50 \( \mu \)m. Inset: RHEED pattern acquired on the CVD-grown sample showing a film with long-range structural order. (d) Atomic force microscope image of a monolayer MoS\(_2\) grain. Scale bar is 2 \( \mu \)m long. Inset: Line scan showing the thickness profile along the blue line in the AFM image. (e) High-resolution TEM image of a suspended MoS\(_2\) film showing the crystallinity of the sample. View of the structural model is overlaid. Scale bar is 0.5 nm long. (f) Orientation histogram based on the area shown in part c confirms that the majority of MoS\(_2\) grain edges are oriented along \( 0^\circ \) and \( \pm 60^\circ \) angles.
Thanks to the optical transparency of the sapphire substrate and the large area of the sample covered by monolayer MoS$_2$, we can perform UV–visible absorption characterization using a simple benchtop spectrophotometer. The resulting spectrum shown in Figure 2c represents broadband absorbance measurements on MoS$_2$. The spectrum shows the well-known A and B excitonic absorption bands at 695 nm (1.78 eV) and 611 nm (2.02 eV). This demonstrates the high optical quality and uniformity of our monolayer MoS$_2$ over a large area. In addition, we can also clearly observe the recently reported C peak at 430 nm (2.88 eV) and the D peak at 303 nm (4.08 eV). Whereas the A and B peaks are associated with optical absorption by band-edge excitons, peaks C and D are associated with van Hove singularities of MoS$_2$.

Figure 3c schematically illustrates the most likely relative orientations for monolayer MoS$_2$ growing epitaxially on the atomically smooth surface of sapphire. These calculations confirm that the interactions between MoS$_2$ and the sapphire surface are of vdW nature and magnitude (see Supporting Section 4). DFT calculations need to be performed in the coincident-site lattice of common supercells in which the MoS$_2$ monolayers are in registry with the substrate. Since these cells exhibit a limited size, some amount of artificial strain needs to be introduced in order to limit the size of the supercells used. In reality, the MoS$_2$ overlayers are not expected to show much strain because of their weak vdW interactions with the substrate. The strain energy turns out to be of similar magnitude to the vdW interaction energy with the substrate. As a consequence, it becomes difficult to ascertain the stability of one orientation with respect to another from DFT calculations alone. On the other hand, these results also clearly point to a vdW nature for the interactions. This prompted us to construct a simple model of adsorption accounting only for pairwise atom–atom interactions of a $-C_6/r_6$ nature. To our surprise, this model shows an extremely rich adsorption phenomenology that is in agreement with the experimental findings, notably, pointing out that

![Figure 2. Optical properties of large-area monolayer MoS$_2$. (a) Raman spectra of as-grown monolayer MoS$_2$ on sapphire and monolayer MoS$_2$ exfoliated from bulk crystals and transferred onto sapphire. (b) Photoluminescence spectra of as-grown monolayer MoS$_2$ on sapphire and monolayer MoS$_2$ exfoliated from bulk crystals onto sapphire. Black lines in parts a and b correspond to fits. (c) UV–vis optical absorbance spectra acquired from large-area monolayer MoS$_2$ showing the A and B absorption peaks due to band-edge excitons as well as C and D peaks associated with van Hove singularities of MoS$_2$.](image)
only MoS$_2$ configurations at either 0° or 60° equivalent orientations (see Figure 3c) provide a structured potential energy surface that is sensitive to arbitrary in-plane translations of the MoS$_2$ plane, and with well-defined maxima and minima in the adsorption energies (Figure 4). All other orientations yield instead a flat energy surface where the adsorption energy is completely independent of translations and rotations of MoS$_2$ with respect to the surface. From this, we can conclude that the MoS$_2$ layer would be free to rotate and slide along the surface of sapphire until it would find the orientation and positions of strongest adsorption. The oriented growth of MoS$_2$ would proceed as follows: using low supersaturation conditions very few MoS$_2$ nuclei are formed and grow laterally on the substrate. Weak (vdW) interactions with the substrate guarantee a certain degree of rotation/translation with respect to the surface in the initial phase. When the nucleus reaches a certain size, not only is it stable against decomposition but it gets pinned down in the most stable orientation with the substrate. Growth proceeds then forming a continuous and oriented layer. We cannot help but see the analogy of this pinning process with the process of attaching a Lego brick onto another larger Lego brick surface, in an analogy introduced by A. Geim in ref 43. Last, the presence of surface defects could be at the origin of less frequently observed lattice orientations.

We characterize the electrical quality of constituent grains by performing electrical transport measurements on single grains (Supporting Section 5). Results show that the mobility reaches a value of 43 cm$^2$/Vs, comparable to results from previous two-terminal measurements on CVD MoS$_2$ (ref 44) and slightly higher than in devices based on exfoliated MoS$_2$ prepared in the same way. Good stitching of such high-quality single crystals into continuous films is necessary in order to realize large-area MoS$_2$ with the same quality as that of the single crystal. We first study the influence of grain boundaries on the electrical
the local variation of the local potential of the surface, relative to the metallic probe. Grain boundaries that contribute additional electrical resistance and degrade device mobility readily show up on such scans (Supporting Section 6). Figure 5 shows local potential maps (upper panels) and potential line scans (lower panels) across the three types of conjoined single-crystal MoS₂ grains that account for pairs formed by orientations shown in Figure 3c. All other possible combinations involving pairs composed of any of the MoS₂ lattice and triangle orientations shown in Figure 3c can be reduced to these three misorientation angles. Figure 5a shows measurements acquired across two grains sharing the same crystalline orientation. In this case, the two grains can join without the appearance of a grain boundary, while in the case of merged single crystals shown in Figure 5b and c with relative lattice orientations characterized by the misorientation angles \( \theta_{\text{rel}} = 60^\circ \) and \( \theta_{\text{rel}} = 30^\circ \), a grain boundary is expected to be present at their interface. Potential line scans, acquired along the red lines overlaid on the potential maps for all three configurations, show a clear monotonic drop of the voltage between the contacts, revealing the absence of an additional voltage drop (resistance) due to the presence of a grain boundary. This shows that for misorientation angles \( \theta_{\text{rel}} = 60^\circ \) and \( \theta_{\text{rel}} = 30^\circ \) occurring in our samples, the single crystals are joined by twin grain boundaries, previously shown not to degrade the electrical properties of polycrystalline MoS₂. This proves that the epitaxial growth of MoS₂ on sapphire, providing these predominant domain

Figure 4. Binding energies for MoS₂ and sapphire. Data are shown as a 3D surface and contour plot of relative binding energies for MoS₂ on sapphire, for different relative lattice orientations and as a function of relative shift in x- and y-directions, corresponding to the lateral motion of the MoS₂ monolayer on the surface of atomically smooth sapphire. Only the 0° and 60° orientations result in significant corrugation and substantial stable minima. The distance between MoS₂ and sapphire is 3.24 Å, as obtained from DFT calculations.
increasing length from 4 μm.

Figure 6. We observe no degradation in mobility with variations in various lead configurations, is shown in Figure 6. We observe no degradation in mobility with increasing length from 4 μm to ~80 μm, within the measurement error. We compare our findings with recent results on polycrystalline continuous CVD MoS2 films, where the material was grown on SiO2 with random relative orientation of grains as well as bilayer formation on grain boundaries influencing electrical transport properties. In our case these sources of scattering are eliminated and mobility stays constant in the devices built on multiple merged grains. We estimate that there are ~16 junctions between single crystals along the length of the 82 μm channel (Supporting Section 7). This shows that the superior electrical properties of the single crystals are preserved over length scales larger than those corresponding to single grains.

CONCLUSIONS

In conclusion, we have achieved epitaxial chemical vapor deposition growth of monolayer MoS2 with a high degree of control over lattice orientation by using atomically smooth surfaces of sapphire as the growth substrate. The large-area MoS2 film is formed from merging single-crystalline domains, with the majority of the domains having lattice orientations being restricted to high-symmetry cases. This shows that even the relatively weak van der Waals interaction can control the crystalline orientation of the overlayer in a similar way to chemical bonds in covalently bonded semiconductors such as GaAs or Si. Here, the preferred MoS2 orientation corresponds to the one with the highest commensuration. While DFT calculations had difficulty explaining the existence of a preferred orientation due to complications related to applying periodic boundary conditions to an incommensurate structure, a remarkably simple model based on pairwise interactions results in predictions in good agreement with the experiment. We believe that this result is general for all vdW heterostructures, and hence we suggest this method as an excellent tool for prediction of such structures.

Local electrical measurements across grain boundaries reveal good stitching between the single crystals in our film with the absence of electrical resistance at most common types of grain boundaries in our sample. The use of sapphire substrates can therefore suppress the formation of tilted grain boundaries, which are detrimental for the electrical properties of the polycrystalline material, and that epitaxial growth of MoS2 is crucial for the formation of nonresistive grain boundaries in a polycrystalline film. We now turn to polycrystalline films and examine their electrical properties by performing measurements on a section of a continuous film with a length L = 81.2 μm and width W = 36.6 μm, contacted by multiple electrodes (Supporting Section 7).

The effective field-effect mobility as a function of channel length, extracted from two contact measurements in various lead configurations, is shown in Figure 6. We observe no degradation in mobility with increasing length from 4 μm to ~80 μm, within the measurement error. We compare our findings with recent results on polycrystalline continuous CVD MoS2 films, where the material was grown on SiO2 with random relative orientation of grains as well as bilayer formation on grain boundaries influencing electrical transport properties. In our case these sources of scattering are eliminated and mobility stays constant in the devices built on multiple merged grains. We estimate that there are ~16 junctions between single crystals along the length of the 82 μm channel (Supporting Section 7). This shows that the superior electrical properties of the single crystals are preserved over length scales larger than those corresponding to single grains.

METHODS

Growth Procedure. Monolayer MoS2 has been grown by chemical vapor deposition on c-plane sapphire. After consecutive cleaning by acetone/2-propanol/DI-water and piranha solution, the substrates were annealed for 1 h at 1000 °C in air. After that, they were placed face-down above a crucible containing ~5 mg of MoO3 (≥99.998% Alfa Aesar) and loaded into a split-tube three-zone CVD furnace with a 32 mm outer diameter quartz tube. CVD growth was performed at atmospheric pressure...
using ultra-high-purity argon as the carrier gas. A second crucible containing 350 mg of sulfur (≥99.99% purity, Sigma-Aldrich) was located upstream from the growth substrates. The growth recipe is as follows: set the temperature of 300 °C with 200 sccm of 10 min, ramp to 700 °C at a rate of 50 °C/min and 10 sccm of carrier gas flow, set the temperature to 700 °C for 10 min, cool to 570 °C with 10 sccm gas flow, increase the gas flow to 200 sccm, and open the furnace for rapid cooling. For more details, please refer to Supporting Sections 1 and 2.

**TEM and AFM Imaging.** CVD MoS₂ was transferred from sapphire using the simultaneous transfer method. TEM grids were used, while for high-resolution TEM (HR-TEM) we used PELCO holey silicon nitride support film with 2.5 um holes in a 200 nm thick Si₃N₄ support. Transmission electron microscopy was performed using a JEOL 2200 FS operated in the 120–200 keV energy range. HR-TEM images were recorded at a magnification of ×200,000. A series of 10–30 images was recorded and stacked with drift correction and averaging using the Stackreg plugin in Fiji (ImageJ). Island orientations in optical images were analyzed using the direlctionality plugin in Fiji (ImageJ). Samples were also imaged using an atomic force microscope (Asylum Research Cypher) operating in ac mode.

**Density Functional Theory Modeling.** All ab initio simulations have been performed within density functional theory, using Quantum-ESPRESSO as implemented in PWscf code of the Quantum ESPRESSO distribution⁵⁹ and VASP.⁶⁰ For calculations using Quantum-ESPRESSO, in order to account for dispersion interactions, we used a revised version¹¹ of the nonlocal density functional by Vydrov and Van Vooths⁶² (rVV10). Ion–electron interactions were taken into account by means of ultrasoft pseudopotentials,⁵³,⁵⁵ with energy cutoffs of 60 Ry for wave functions and 400 Ry for density. Pseudopotentials have been generated using parameters taken from the PSLibrary repository (v.0.3.0) of the QEforge distribution and setting the rVV10 parametrization.⁵² Surface calculations were performed by considering Al₂O₃ slabs with six oxygen O₃ layers and 10 (12) aluminum layers in the case of OH (Al) termination. In order to minimize the interaction between periodic replicas, supercells were built including 20 Å of vacuum. A series of 10–30 images was recorded and stacked with drift correction and averaging using the Stackreg plugin in Fiji (ImageJ). Island orientations in optical images were analyzed using the direlctionality plugin in Fiji (ImageJ). Samples were also imaged using an atomic force microscope (Asylum Research Cypher) operating in ac mode.

**Conflict of Interest:** The authors declare no competing financial interest.

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**Supporting Information Available:** Substrate preparation, growth setup, SAED analysis, sapphire-MoS₂ van der Waals interaction, electrical transport measurements on a single crystal, scanning Kelvin probe microscopy, and electrical transport measurements on large-area films. This material is available free of charge via the Internet at http://pubs.acs.org.

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