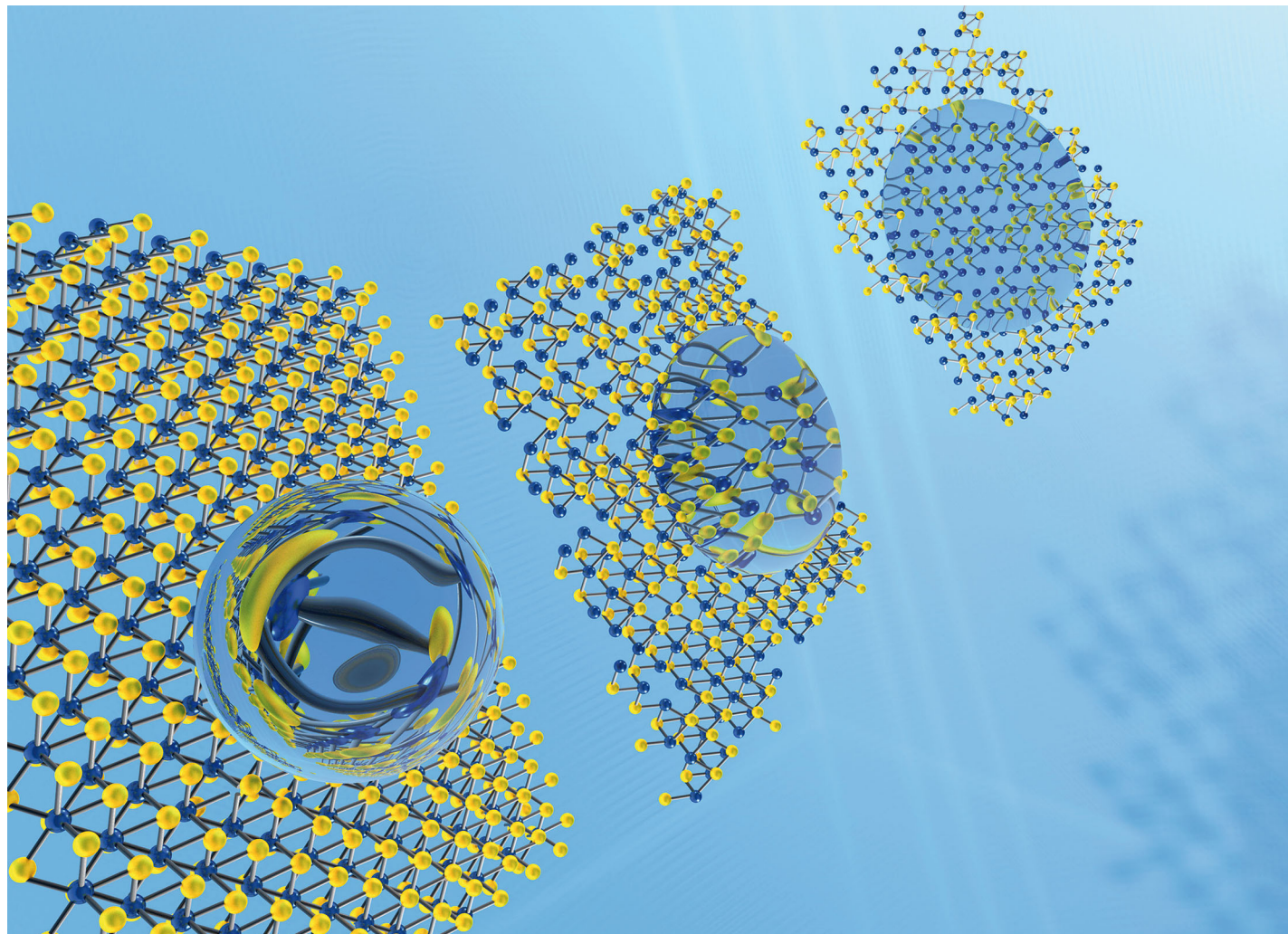




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Department of Biomedical Engineering, Texas A&M University,
United States.

Superhydrophobic states of 2D nanomaterials controlled by
atomic defects can modulate cell adhesion

A new concept to control the wetting characteristics by
modulating atomic defects in 2D-nanomaterials is introduced.
This work shed new light on the role of atomic vacancies
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Superhydrophobic states of 2D nanomaterials controlled by atomic defects can modulate cell adhesion†

Manish K. Jaiswal,^a Kanwar Abhay Singh,^a Giriraj Lokhande^a and Akhilesh K. Gaharwar^{abc}

We introduced a new concept to the control of wetting characteristics by modulating the degree of atomic defects of two-dimensional transition metal dichalcogenide nanoassemblies of molybdenum disulfide. This work shed new light on the role of atomic vacancies on wetting characteristic that can be leveraged to develop a new class of superhydrophobic surfaces for various applications without altering their topography.

Superhydrophobic materials are used extensively in bioelectronics, medical, and self-cleaning devices. These superhydrophobic properties rely on altering topological features such as surface roughness and surface energy. For instance, microstructural features with grooves filled with air-packets govern the wetting behavior of microengineered superhydrophobic surfaces.¹ Such behavior has been explained by Cassie–Baxter wetting theory, where liquid droplets make contact only with tips of the grooves, and intact air-packets support the spherical shape of the droplet to provide superhydrophobic properties.^{2,3} These methods involve a cumbersome fabrication approach and cannot be applied to biosensing, lab-on-a-chip, blood-repellent, anti-fouling, and self-cleaning applications, which demand a non-textured approach to achieve a superhydrophobic state.

Two-dimensional (2D) nanomaterials have received considerable attention due to high degree of structural anisotropy and chemical functionality.^{4–7} For example, 2D transition metal dichalcogenide (TMD) nanosheets have been investigated for nanoelectronics, optical sensors, renewable energy sources, catalysis, and lubrication.^{8–10} In particular, 2D molybdenum disulfide (MoS₂) has been explored in nanoelectronics^{9,11} and as a photothermal agent^{12–14} due to its tunable band gap. The

atomic defects present on the lattice plane of MoS₂ provide an active site for various applications, including catalyst and chemical conjugations.^{10,15} More recently, hierarchical nanoassemblies of defect-rich MoS₂ have been shown to improve the performance of lithium ion storage batteries by 20-fold compared with traditional monolayers of MoS₂,^{16–19} and can be used to form hydrogels using polymeric binders *via* metal-click chemistry.²⁰ MoS₂ is cytocompatible and degrades in an *in vitro* microenvironment and, thus, has been explored for various biomedical applications.²¹

In the past few years, the superhydrophobic characteristics of 2D nanomaterials such as graphene,²² boron nitride,²³ tungsten disulfide (WS₂),²⁴ and MoS₂²⁵ have been reported. The wetting characteristics of these 2D nanomaterials are attributed mostly to the surface roughness or deposition of air-borne hydrocarbons which, ultimately, reduce the surface activity.^{25,26} Atomic defects that can be incorporated into 2D nanomaterials during the synthetic process have not been investigated but which could provide a facile approach to tune the wetting characteristics.

Here, we report the tunable wetting behavior of defect-rich MoS₂ *via* modulating atomic vacancies. Specifically, we demonstrated that MoS₂ with a low degree of atomic defects resulted in superhydrophobic behavior (contact angle > 150°), whereas a high degree of atomic defects resulted in superhydrophilic properties (contact angle ~ 0). The wetting properties controlled by atomic vacancies displayed minimum hysteresis of the contact angle, indicating that the superhydrophobic behavior was independent of the amount of energy dissipated during wetting and dewetting. The superhydrophobic MoS₂ could be coated on various substrates (glass, silica, rubber and paper), thereby indicating high versatility for surface-coating applications.

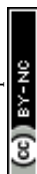
MoS₂ nanoassemblies with predefined atomic defects were synthesized by modulating the ratio of precursors of molybdenum (hexaammonium heptamolybdate):sulfur (thiourea) (1:1, 1:2, 1:4 and 1:6) (see ESI†). The defect-rich MoS₂ lattice had hexagonal crystallographic arrangements of Mo and S atoms (Fig. 1a). Depending on the precursor ratio (Mo:S), the number of atomic

^a Department of Biomedical Engineering, Texas A&M University, College Station, TX-77843, USA. E-mail: gaharwar@tamu.edu, manish@tamu.edu

^b Department of Biomedical Engineering, Texas A&M University, College Station, TX-77843, USA

^c Center for Remote Health Technologies & Systems, Texas A&M University, College Station, TX-77843, USA

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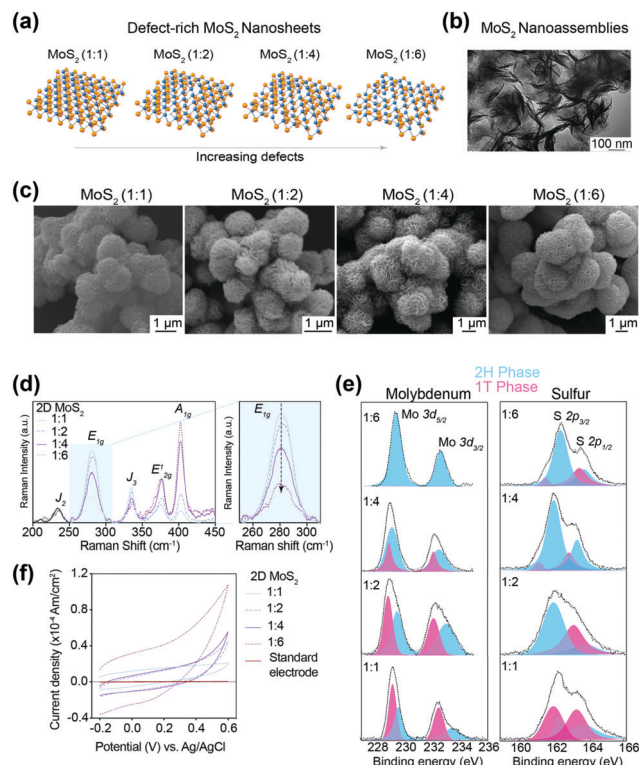


Fig. 1 Defect-rich nanoassembly of 2D MoS₂ sheets. (a) Changing the ratio of molybdenum : sulfur precursors led to the formation of MoS₂ with different degrees of atomic vacancies. (b) TEM image confirmed the order of individual sheets into nanoassemblies. (c) SEM image shows a “flower”-like morphology of nanoassemblies of typical size 1.5–3 μm. No significant change in size or shape was observed due to changes in atomic vacancies for MoS₂ (1 : 1, 1 : 2, 1 : 4 and 1 : 6 samples). (d) Raman spectra confirmed 2H to be the dominant phase with a partial presence of 1T-phase formation. The vibrations of in-plane S atoms confirmed that a highly ordered and pronounced 2H phase was present with increasing amounts of the sulfur precursor. (e) XPS showing the core-level binding energies of Mo and S atoms for MoS₂ samples. (f) Cyclic voltammetry verified the increased defect densities with increasing amounts of the sulfur precursor, thus confirming 1 : 1 with least atomic vacancies and 1 : 6 with maximum atomic vacancies.

defects could be modulated from sample 1:1 to 1:6. The transmission electron microscopy (TEM) image of typical defect-rich MoS₂ shows the nanoassembly of individual 2D sheets (Fig. 1b). The scanning electron microscopy (SEM) images of MoS₂ samples with different ratios of Mo and S demonstrated a well-segregated “flower-like” hierarchical architecture (Fig. 1c). The average size of nanoassemblies was 1.5–3 μm. A significant effect of atomic defects on the size of nanoassemblies was not observed (Fig. S1, ESI†).

In 2D MoS₂, a layer of Mo (oxidative state, 4+) atoms was held between two layers of S (oxidative state, 2–) atoms. Two MoS₂ layers interacted with each other *via* short-range van der Waals interactions. The Mo atoms could have trigonal prismatic co-ordination with S atoms to result in hexagonal symmetry (2H phase) or octahedral co-ordination to result in tetragonal symmetry (1T phase). The Raman spectra showed the formation of a dominant 2H phase with partial presence of the 1T phase (Fig. 1d).

The characteristic E_{2g}¹ (in-plane vibration of two S atoms) and A_{1g} (out-of-plane vibration of S atoms) occurred at ~375 cm^{–1} and ~402 cm^{–1}, respectively. The appearance of the E_{1g} peak at 280 cm^{–1} (which was due to the tetragonal symmetry of the lattice) indicated a metastable 1T phase besides the dominant 2H phase in MoS₂ samples, which was likely due to ammonium cations released during synthesis.²⁷ Furthermore, as expected, a simultaneous attenuation in the intensity of the E_{1g} peak accompanied with gradual amplification in E_{2g}¹ and A_{1g} from sample 1 : 1 to sample 1 : 6 demonstrated the formation of a highly ordered 2H phase with an increase in thiourea concentration. The observation in Raman spectra was corroborated further with X-ray photoelectron spectroscopy (XPS) for the core-level binding energies of Mo and S atoms for MoS₂ samples (1 : 1, 1 : 2, 1 : 4 and 1 : 6) (Fig. 1e). The presence of the 1T phase (tetragonal symmetry) was dominated by the 2H phase (hexagonal symmetry), as represented by blue curves with increments in the sulfur precursor (thiourea content) from 1 : 1 through 1 : 6. These results corroborated the findings of Raman spectroscopy. The specific BE for the 1T-phase Mo was (3d_{5/2}) 229.2 and (3d_{3/2}) 232.4 eV, and for S was (2p_{3/2}) 161.9 and (2p_{1/2}) 163.1 eV. The corresponding BE for the 2H phase was slightly higher for Mo (229.6 and 233.3 eV) and S (162.2 eV and 163.5 eV), respectively.

The number of atomic vacancies for 2D MoS₂ was determined with cyclic voltammetry *via* expanded J/V (current density *vs.* potential) curves (Fig. 1f). An increase in the sulfur precursor (thiourea) resulted in an increase in the area under the J/V curve. The concentration of atomic defects increased with an increase in the Mo:S precursor ratio, 1 : 1 (~0.59 × 10² μM g^{–1}), 1 : 2 (~0.72 × 10² μM g^{–1}), 1 : 4 (~0.80 × 10² μM g^{–1}) and 1 : 6 (~2.1 × 10² μM g^{–1}) (see ESI† for details).

The effect of atomic vacancy on wetting characteristics was evaluated by determining the contact angle of water and polar/non-polar organic solvents on MoS₂-coated surfaces. To determine wetting characteristics, the water contact angle on a MoS₂ (1 : 1, 1 : 2, 1 : 4 and 1 : 6)-coated glass substrate was evaluated (Fig. 2a). The highest contact angle was observed for sample 1 : 1 (157.5 ± 5.2°), which decreased gradually for sample 1 : 2 (137.6 ± 2.0°), sample 1 : 4 (23.7 ± 10.0°) and sample 1 : 6 (3.6 ± 3.0°), thus causing the surface to become superhydrophilic. The associated liquid-droplet images are shown on top of the bar graph. As expected, the droplet shape was spherical for 1 : 1, indicating superhydrophobic characteristics. As the ratio of sulfur precursor increased during MoS₂ synthesis, the water droplet became oblate (1 : 2) and eventually spread (1 : 4) and adsorbed (1 : 6) onto the coated surface, thereby suggesting atomic defect-dependent wetting behavior.

To further verify superhydrophobic characteristics, a rectangular pattern (2 cm × 1 cm) covering the boundary area using MoS₂ (1 : 1) on the glass slide was filled with 1 mL of water (Fig. 2b). The uniform coating of MoS₂ (1 : 1) on the glass substrate was verified using SEM. The rectangular pattern (thickness, 0.5 mm) could hold 10-fold water molecules (height, 5 mm) due to its superhydrophobic characteristics. Even after it was subjected to manual shaking, the movement of water was confined within the rectangle area due to the hydrophobic nature of the MoS₂ (see ESI,† Video V1). In addition, a water droplet was allowed to roll-down



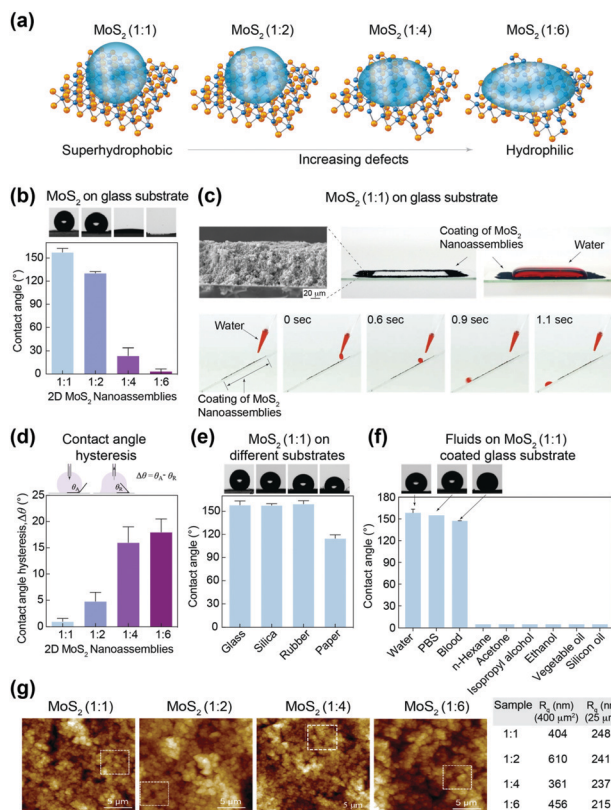


Fig. 2 Wetting characteristics of defect-rich MoS_2 nanoassemblies. (a) Wetting characteristics are controlled by atomic defects. (b) The contact-angle measurements of MoS_2 (1:1, 1:2, 1:4, and 1:6)-coated glass slides exhibited superhydrophobic-to-superhydrophilic transition. (c) A MoS_2 (1:1)-coated rectangular area confined the water droplet without spilling. The water droplets (red) rolled down on the MoS_2 (1:1)-coated slanted surface without being adsorbed. (d) MoS_2 (1:1)- and MoS_2 (1:2)-coated surfaces showed low hysteresis of the contact angle. (e) MoS_2 (1:1) coated on glass, silica, rubber and paper retained its superhydrophobic characteristics. (f) The superhydrophobic behavior of a MoS_2 (1:1)-coated surface was observed for water, PBS and blood, whereas an organic solvent was readily spread on the surface. (g) The roughness (R_q) for all four MoS_2 samples was analyzed by AFM for 400 and 25 μm^2 imaging areas. The values are presented in the table.

a slanted glass slide coated with MoS_2 (1:1) to investigate the superhydrophobic properties in dynamic conditions. The water droplet (red) rolled down without being adsorbed over the surface and remained intact (spherical shape) during rolling. Hence, MoS_2 (1:1) could be used for various coating applications (Fig. 2b and c; see ESI† Video V2).

The wetting behavior of a MoS_2 -coated surface can be understood in terms of the interfacial tension (γ) between the liquid and the coating as given by Young's equation,²⁸ $\cos \theta = (\gamma_{\text{sa}} - \gamma_{\text{sl}})/\gamma_{\text{la}}$, where θ is the contact angle, γ_{sl} is the solid/liquid interfacial free energy, γ_{sa} is the solid surface free energy, and γ_{la} is the liquid surface free energy. As the Mo:S precursor ratio changed from 1:1 to 1:6, the interaction between the MoS_2 -coated surface and water molecules increased due to the presence of defects owing to direct access of the Mo atomic plane, and could potentially form a molybdenum–water complex. This resulted in reduction of the interfacial tension between the liquid and substrate (γ_{sl}) and

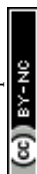
surface tensions (γ_{sa} and γ_{la}) approached each other. This results in θ tended to zero, thus causing the sample to become superhydrophilic.

The contact angle is often a function of whether the liquid is advancing or receding on the substrate.²⁸ To ascertain such possible behavior of the samples, the contact angle hysteresis ($\Delta\theta$) was measured (which is essentially the difference between advancing (θ_{A}) and receding contact angles (θ_{R})). As expected, $\Delta\theta$ was minimal for the MoS_2 samples 1:1 ($1 \pm 0.6^\circ$) and 1:2 ($4.8 \pm 1.7^\circ$), indicating that their superhydrophobic character was independent of roughness or surface contamination (Fig. 2d). Higher $\Delta\theta$ was observed for sample 1:4 ($16 \pm 3^\circ$) and 1:6 ($18 \pm 2.5^\circ$). The low $\Delta\theta$ for MoS_2 (1:1) indicated that the Cassi–Baxter wetting behavior of the liquid caused the droplets to roll-down in a spherical shape over the slightly slanted surface (see ESI†). Such superhydrophobic materials with small $\Delta\theta$ are highly sought after for commercial applications (e.g., designing self-cleaning and spill-resistance coating materials for exterior glass and wind shields).

To further investigate the versatility of MoS_2 (1:1) as a superhydrophobic coating, substrates such as silica, rubber and papers were coated with MoS_2 (1:1) and the water contact angle determined (Fig. 2e). All the surfaces showed superhydrophobic characteristics: silica ($157.8 \pm 5.2^\circ$), rubber ($159.7 \pm 4.0^\circ$), and paper ($115 \pm 4.7^\circ$). In addition, the contact angle for various biological fluids and organic solvents, including phosphate-buffered saline (PBS) (pH 7.4), whole blood, hexane, ethanol, isopropanol, fatty vegetable oil and silicone oil, were investigated (Fig. 2f). The contact angle for PBS ($151.0 \pm 1.07^\circ$) and whole blood ($147.2 \pm 0.41^\circ$) was similar to that of water. For glycerol ($106.0 \pm 28.6^\circ$), silicone oil ($16.4 \pm 2.5^\circ$) and other organic solvents ($\sim 5^\circ$), the contact angle was lower due to enhanced interactions of the sulfur layer (from MoS_2) with hydrocarbons, acetone and mild polar alcohols. Moreover, an aqueous solution of an amphiphile, such as sodium dodecyl sulfate (SDS), spread onto the MoS_2 (1:1 and 1:6)-coated glass substrate.

To investigate the effect of surface roughness on wetting characteristics, topographical analysis was carried out using atomic force microscopy (AFM). The root-mean square value of roughness (R_q) was calculated for an imaging area of 400 and 25 μm^2 for all samples (Fig. 2g). There was no significant difference in R_q (400–600 nm for 400 μm^2 and 220–250 nm for 25 μm^2) for different MoS_2 (1:1 to 1:6) samples. This observation suggested that topographical features did not govern the hydrophobic character of MoS_2 .

We investigated the adhesion of human mesenchymal stem cells (hMSCs) onto superhydrophobic (1:1) and superhydrophilic surfaces (1:6). High adhesion of hMSCs was observed on hydrophilic surfaces. Superhydrophobic surfaces of MoS_2 (1:1) restricted adhesion of hMSCs compared with MoS_2 (1:6) (Fig. 3a). To investigate the role of protein adsorption, we tested adhesion of hMSCs in the presence and absence of serum. We observed significantly higher adhesion of hMSCs in the presence of serum in both MoS_2 surfaces (1:1 and 1:6). Energy-dispersive X-ray spectroscopy (EDS) mapping for C and N elements confirmed the higher adhesion of hMSCs on hydrophilic surfaces of MoS_2 (1:6).



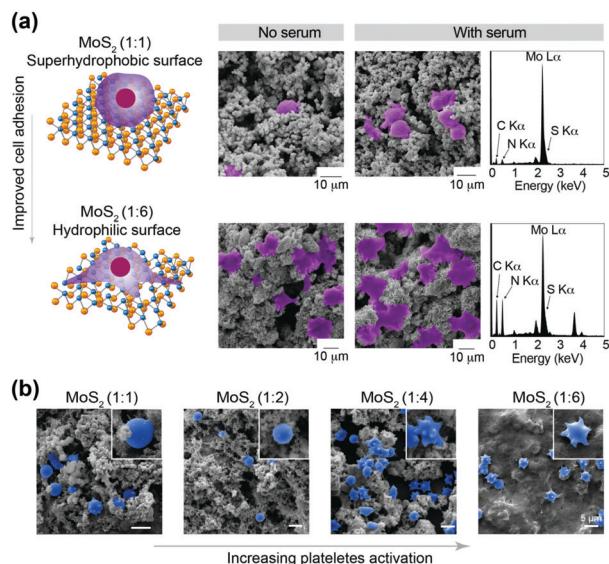


Fig. 3 Cell-adhesion characteristics of defect-rich MoS₂ nanoassemblies. (a) Cellular adhesion on MoS₂-coated substrates (1:1, hydrophobic) and (1:6, hydrophilic) in the presence and absence of serum proteins. The false-color SEM images show adherent cells in purple. Energy-dispersive X-ray spectroscopy (EDS) profile showing stronger peaks of C and N, which further confirmed higher cell adhesion for hydrophilic substrates regardless of serum addition. (b) Adhesion and activation of platelets on MoS₂-coated substrates. The morphological changes of platelets shown using the false blue color in SEM images demonstrated hydrophobicity-controlled activation of platelets.

However, we did not observe a significant change in protein adsorption on MoS₂ surfaces (1:1 and 1:6). We further evaluated the effect of wetting characteristics on platelet activation. The adhesion and activation of platelets was done using bovine blood on MoS₂ surfaces (1:1 through 1:6) (Fig. 3b). The hydrophilic nature of the surface promoted platelet activation.

In summary, we introduced a new concept to control the wetting characteristics of 2D nanomaterials by modulating the degree of atomic defects. Specifically, we synthesized superhydrophobic 2D MoS₂ and demonstrated that atomic defects dictated hydrophilic-to-hydrophobic transition. The highest contact angle was observed for MoS₂ with a Mo:S precursor ratio of 1:1 ($157.5 \pm 5.2^\circ$), which decreased gradually for 1:2 ($137.6 \pm 2.0^\circ$), 1:4 ($23.7 \pm 10.0^\circ$) and 1:6 ($3.6 \pm 3.0^\circ$), thus causing the surface to become superhydrophilic. MoS₂ (1:1) could be coated readily on glass, silica, rubber and paper and retain its superhydrophobic characteristics. Due to low $\Delta\theta$, superhydrophobic MoS₂ (1:1) could be used for biosensing, lab-on-a-chip, blood-repellent, anti-fouling, and self-cleaning applications, which demand a non-textured approach to achieve a superhydrophobic state. The wetting characteristics of the MoS₂ surface directly influence cell-adhesion characteristics that can be leveraged for biomedical applications.

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Conflicts of interest

There are no conflicts to declare.

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