



Boosting the electronic and catalytic properties of 2D semiconductors with supramolecular 2D hydrogen-bonded superlattices

Can Wang, Rafael Furlan de Oliveira, Kaiyue Jiang, Yuda Zhao, Nicholas Turetta, Chun Ma, Bin Han, Haiming Zhang, Diana Tranca, Xiaodong Zhuang, et al.

► To cite this version:

Can Wang, Rafael Furlan de Oliveira, Kaiyue Jiang, Yuda Zhao, Nicholas Turetta, et al.. Boosting the electronic and catalytic properties of 2D semiconductors with supramolecular 2D hydrogen-bonded superlattices. *Nature Communications*, 2022, 13 (1), pp.510. 10.1038/s41467-022-28116-y . hal-03633307

HAL Id: hal-03633307

<https://hal.science/hal-03633307>

Submitted on 6 Apr 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

ARTICLE


<https://doi.org/10.1038/s41467-022-28116-y>

OPEN

Boosting the electronic and catalytic properties of 2D semiconductors with supramolecular 2D hydrogen-bonded superlattices

Can Wang¹, Rafael Furlan de Oliveira¹, Kaiyue Jiang², Yuda Zhao¹, Nicholas Turetta ¹, Chun Ma¹, Bin Han¹, Haiming Zhang ³, Diana Tranca², Xiaodong Zhuang ², Lifeng Chi ³, Artur Ciesielski ¹✉ & Paolo Samorì ¹✉

The electronic properties of two-dimensional semiconductors can be strongly modulated by interfacing them with atomically precise self-assembled molecular lattices, yielding hybrid van der Waals heterostructures (vdWHs). While proof-of-concepts exploited molecular assemblies held together by lateral unspecific van der Waals interactions, the use of 2D supramolecular networks relying on specific non-covalent forces is still unexplored. Herein, prototypical hydrogen-bonded 2D networks of cyanuric acid (CA) and melamine (M) are self-assembled onto MoS₂ and WSe₂ forming hybrid organic/inorganic vdWHs. The charge carrier density of monolayer MoS₂ exhibits an exponential increase with the decreasing area occupied by the CA·M unit cell, in a cooperatively amplified process, reaching $2.7 \times 10^{13} \text{ cm}^{-2}$ and thereby demonstrating strong n-doping. When the 2D CA·M network is used as buffer layer, a stark enhancement in the catalytic activity of monolayer MoS₂ for hydrogen evolution reactions is observed, outperforming the platinum (Pt) catalyst via gate modulation.

¹Université de Strasbourg, CNRS, ISIS, 8 allée Gaspard Monge, 67000 Strasbourg, France. ²The Meso-Entropy Matter Lab, The State Key Laboratory of Metal Matrix Composites, Shanghai Key Laboratory of Electrical Insulation and Thermal Ageing, School of Chemistry and Chemical Engineering, Frontiers Science Center for Transformative Molecules, Shanghai Jiao Tong University, 200240 Shanghai, P. R. China. ³Institute of Functional Nano & Soft Materials (FUNSOM), Jiangsu Key Laboratory for Carbon Based Functional Materials & Devices, Soochow University, 215123 Suzhou, P. R. China. ✉email: ciesielski@unistra.fr; samori@unistra.fr

Since the first isolation of graphene¹, the family of two-dimensional materials (2DMs) has expanded rapidly providing access to materials holding different properties. Their combination, by arbitrary stacking dangling-bond-free 2D nanosheets to form van der Waals heterostructures (vdWHs)², enabled the further diversification of the properties^{3,4} as a result of the charge redistribution at the interfaces of different 2DMs' crystals⁵.

Artificial vdWHs can be produced by using diverse strategies, such as the pick-and-lift technique and wet-transfer from chemical vapor deposition (CVD) samples. However, all these methods suffer from the lack of control over the lattice alignment at the interface between two 2D nanosheets. Moreover, the transferred layers can exhibit microcracks, wrinkles, and contaminations by residues from the often-employed sacrificial layers. Epitaxial growth techniques are frequently exploited for the fabrication of 2D heterostructures via lateral overgrowth⁶. Additionally, lattice and thermal expansion coefficient mismatches limit the possibility for growth and integration of high-efficiency electronic and photonic devices based on dissimilar 2DMs. Most importantly, neither of these conventional methods can meet the requirements for programming highly ordered electrostatic superlattices in vdWHs, to ultimately efficiently modulate the electronic properties of 2DMs. In this context, the use of the supramolecular networks represents a simple, yet remarkably effective solution to overcome the aforementioned problems. 2DMs provide atomically flat surfaces for molecular assembly governed by non-covalent interactions^{7,8}. The physisorption of suitably designed molecules allows the formation of self-assembled monolayers (SAMs) under thermodynamic control which represents true superlattices on top of the crystalline 2DMs. The presence of electron-donating/withdrawing groups in the molecule employed to form the SAMs can lead to doping of 2DMs through the controlled local modifications of their surface potential^{8–10}. Compared to molecules randomly adsorbed on the 2DMs, the use of highly ordered superlattices may maximize the electronics variation via collective or even cooperative effects. Pioneering works demonstrating the feasibility of 2DMs' surface decoration with molecular monolayers include organic/inorganic vdWHs of graphene, *h*-BN, and MoS₂⁷. While the band structures of the pristine layered materials are modified by the additional periodic potential introduced by SAM lattices, hitherto the examples are limited to monolayers formed via unspecific weak intermolecular interactions between alkyl chains that result in small domains (<10⁵ nm²) featuring random orientations^{11–13}. Additionally, SAMs formed via lateral vdW interactions may undergo dynamic rearrangements upon humidity^{14,15} or high temperature exposure¹⁶ further leading to instabilities in 2DM-based (opto)electronic devices under operation. The use of 2D supramolecular structures held together by directional and specific non-covalent interactions appears as an ideal solution to attain higher stability and structural control over the micrometer scale.

Intermolecular hydrogen bonds exhibit higher energy (25–40 kJ/mol) compared to intermolecular vdW forces (5 kJ/mol), and they have been widely exploited to guide the self-assembly of suitably designed molecular modules into a variety of supramolecular architectures including 1D, 2D, and 3D arrangements¹⁷. This type of interaction is unique as it offers a high level of control over the molecular assembly process, along with high specificity, directionality, and reversibility. H-bonding interactions can benefit from a wide range of interaction energies, depending on the number and position of consecutive strong H-bonds. In this framework, melamine (M) represents an archetypal system for the formation of stable non-covalent architectures, in the form of bi-component networks with

cyanuric acid (CA)¹⁸, perylene-3,4,9,10-tetracarboxylic diimide¹⁹, and other di-imide molecules^{20,21}. Micrometer-size extended CA·M networks self-assembled on 2D substrates, including HOPG, Au(111), and black phosphorous^{22–25}, have been shown to exhibit a low amount of structural defects and elevated thermal stability²³.

Here, we study the effect of the physisorbed CA·M bi-component self-assembled network on the electronic properties of the underlying TMDCs. We found the occurrence of a strong charge-transfer process taking place on TMDCs as characterized by a positive cooperative effect emerging in densely packed CA·M bi-component tri-hapto hydrogen-bonded networks. Hybrid van der Waals heterostructures based on molybdenum disulfide (MoS₂) functionalized with CA·M supramolecular lattices exhibit an increase in charge carrier density, exceeding 10¹³ cm⁻². Moreover, the same superlattice grown on tungsten diselenide (WSe₂) also displays similar significant improvement of the electronic transport through the 2D material. Further insights into the electronic effect of this functionalization approach are gained with density functional theory (DFT) calculations and Kelvin Probe Force Microscopy (KPFM). The efficient charge transfer from the physisorbed CA·M superlattice to the 2DMs determines an upward shift of the Fermi level (*E_F*) of the 2DM. The studies on MoS₂ as an earth-abundant and inexpensive material are essential for the implementation of clean energy technologies using hydrogen. The formation of MoS₂ based 2D heterostructures, e.g., graphene/MoS₂²⁶ and WS₂/MoS₂²⁷, induces a built-in electric field formed among the dissimilar layers, enhancing the MoS₂ catalytic performance. The highly crystalline nature of the 2D CA·M network and its significant *n*-doping effect on the 2DMs induces a strong enhancement of the hydrogen evolution reaction (HER) activity of MoS₂, surpassing the performance of the standard polycrystalline platinum catalyst. The effect the CA·M superlattice shifts the HER overpotential of MoS₂ over 100 mV, yielding a small Tafel slope of 40 mV/dec.

Results

Positive cooperative effect on charge transfer. Cooperativity is a hallmark of molecular self-assembly, with the ensemble behaving differently as a whole from the sum of the isolated individual molecules. DFT calculations revealed that the CA and M co-assembly on MoS₂ surface exhibits a positive cooperativity on the charge transfer process.

Tightly packed 2D assemblies of M, CA, and CA·M onto the surface of monolayer MoS₂ have been modeled (Supplementary Fig. 1). The different charge density images based on such superlattices (Fig. 1a–c) indicate the charge transfer process from the organic layer to the MoS₂, and the calculated magnitude of charge transfer is summarized in Fig. 1d. M molecules are found to adopt a bent conformation in tightly packed monolayers on MoS₂, which leads to a lower magnitude of charge transfer (2.47 × 10¹³ cm⁻²) compared to nearly perfectly flat CA assemblies (2.70 × 10¹³ cm⁻²). The bi-component CA·M assembly onto MoS₂ reveals a charge transfer magnitude as high as 3.62 × 10¹³ cm⁻², hence outperforming the mono-component assemblies. To gain deeper insight into the role of the admolecules on the local electronics of the MoS₂, the magnitude of the charge transfer is computed for mono-component M and CA as well as for the bi-component CA·M superlattice as a function of the tightness of the packing at the supramolecular level (Fig. 1e). We find a linear growth of the charge transfer with the decreasing area of the unit cell of the superlattice for mono-component M and CA assemblies-based hybrids, highlighting a collective nature of the effect²⁸. In stark contrast, the bi-component CA·M superlattice exhibits a positive cooperativity

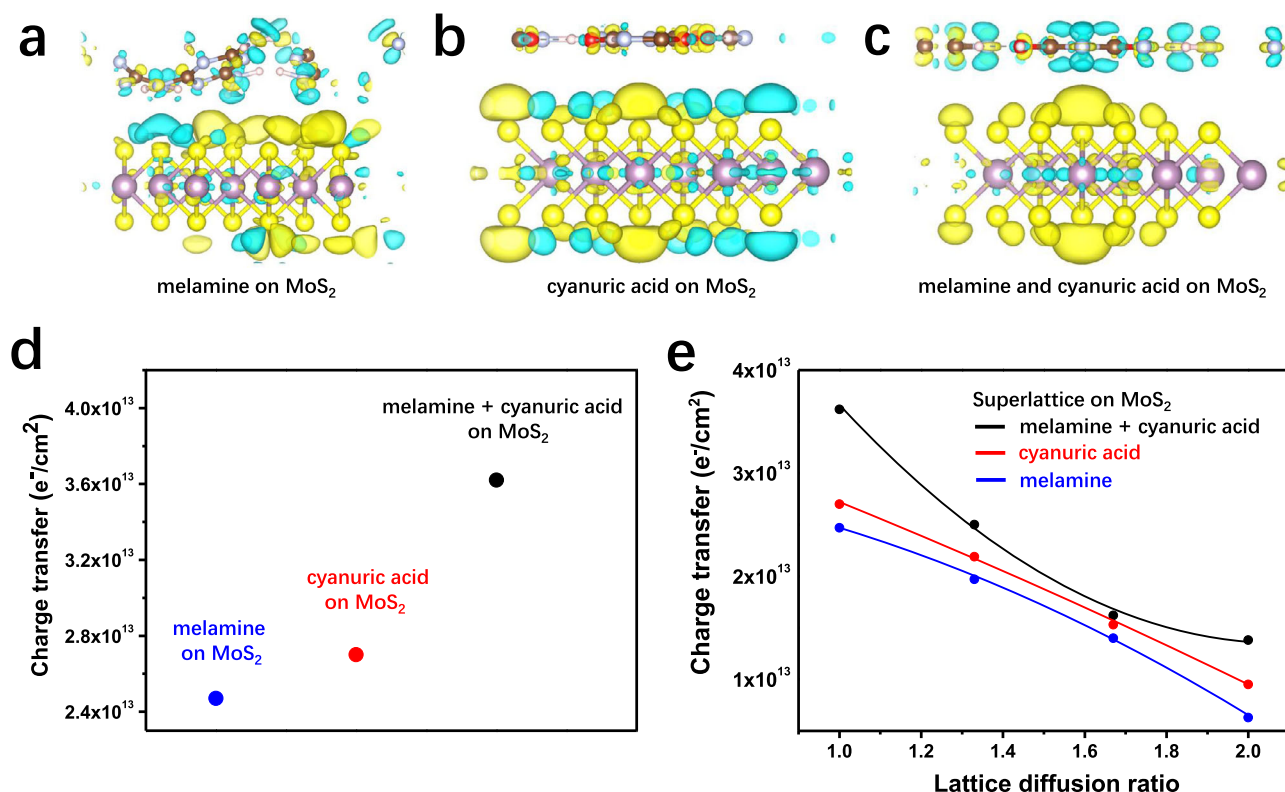


Fig. 1 Simulated charge transfer from melamine (M), cyanuric acid (CA), and co-assembled structure to MoS₂ monolayer. **a–c** The side-view differential charge density images of **a** melamine (M), **b** cyanuric acid (CA), and **c** co-assembled network (CA-M) on monolayer MoS₂. The yellow and light blue regions in panels indicate the charge accumulation and depletion, respectively. **d** The magnitude of the charge transfer in tightly packed molecule adlayers. **e** Charge transfer as a function of unit cell size for melamine, cyanuric acid, and co-assembled networks on monolayer MoS₂.

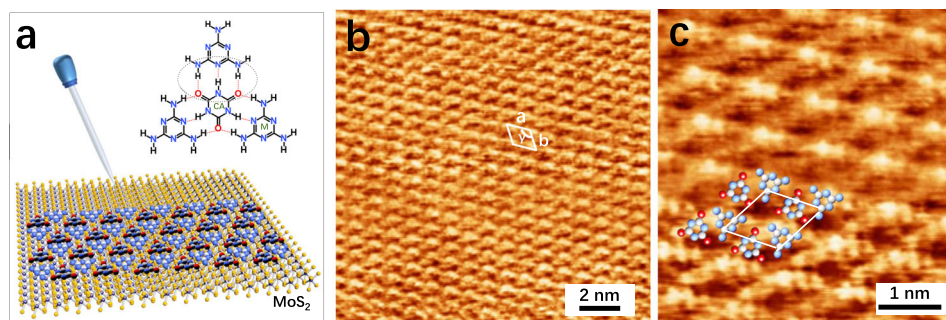


Fig. 2 Supramolecular network of co-assembled CA-M monolayer onto monolayer MoS₂. **a** Schematic representation of the network formation, and the molecular structure of CA-M formed through hydrogen bonding between on monolayer MoS₂. **b, c** Scanning tunneling microscope height image of co-assembled network adsorbed onto MoS₂ surface, and its unit cell is denoted by a white rhombus. Unit cell parameters, i.e., vectors $a = b = 0.96 \pm 0.02$ nm, and the angle between them $\gamma = 120 \pm 2^\circ$ (**b**) survey (**c**) zoom-in with proposed molecular packing motif of CA-M. Tunneling parameters: **b** tip bias voltage (V_t) = -930 mV, average tunneling current (I_t) = 60 pA; **c** V_t = -950 mV, I_t = 60 pA.

with an exponential growth of the magnitude of the charge transfer upon reduction of the area of the unit cell. At all values of lattice areas, the magnitudes of the charge transfer for the bi-component CA-M are greater than those determined for the mono-components M and CA superlattices on MoS₂.

Hybrid structures formation and characterization. The formation of CA-M network on the TMDCs surface is schematically illustrated in Fig. 2a. A 1:1 mixture of CA and M molecules is prepared with Milli-Q water at a concentration of 1×10^{-7} M. A 20 μ l drop of the warm solution (ca. 350 K) is deposited onto a freshly cleaved MoS₂ crystal surface which is subsequently transferred into a desiccator to promote the solvent evaporation.

The modified surface is then rinsed abundantly with water to remove the excess of physisorbed molecules not involved in the formation of the 2D CA-M network, followed by thermal annealing at 430 K in a vacuum for 12 h. Atomic force microscopy (AFM) imaging is employed to monitor the evolution of the surface morphology. It reveals that annealing under vacuum promotes molecular rearrangement on the MoS₂ surface yielding a drastic increase in the size of the CA-M domains up to tens of μ m² forming a continuous crystalline film (Supplementary Fig. 3b). Such 2D CA-M network exhibits an average thickness of 0.397 ± 0.103 nm.

The structure of the dry self-assembled 2D CA-M network is further investigated with a sub-nm resolution by scanning

tunneling microscopy (STM) imaging under ambient conditions. In general, the on-surface molecular assembly is driven by the interplay between molecule-substrate and molecule-molecule interactions. In the present case, the 2D CA-M network formation is mainly governed by intermolecular interaction through the generation of three tri-hapto hydrogen bonds between one CA and three M molecules (Fig. 2a). Such self-assembly pattern further expands across the surface to form a large-area 2D supramolecular structure, where the CA-M network physisorbs onto the underlying MoS₂ support as stabilized by vdW interactions. Figure 2b reveals that the CA-M network displays a hexagonal symmetry similar to that of the underlying MoS₂ lattice. The unit cell parameters are measured as $a = b = 0.96 \pm 0.02$ nm, and $\gamma = 120 \pm 2^\circ$. As the CA-M structure observed on the MoS₂ surface corresponds well to the periodic distance observed in bulk CA-M crystals (measured as 0.964 nm)²⁹, we consider that the packing structure of the formed 2D CA-M monolayer is completely dominated by the intermolecular interactions. The detailed structural information gathered from high-resolution STM imaging (Fig. 2c) reveals the presence of hexagonal cavities in the CA-M network. DFT simulations of the optimized CA-M structure confirms that the CA-M network adsorbs flat on MoS₂ surface, with the CA and M molecules interacting through N-H...O and N-H...N hydrogen bonds (Supplementary Fig. 1c). The simulated SAM structure perfectly matches the STM results.

The same preparation procedure of this hybrid supramolecular adlayer was also applied by using the WSe₂ crystal as support, and a similar surface morphology and structure have been monitored by AFM and STM, respectively (Supplementary Fig. 4). Due to the strong multiple hydrogen bonds holding adjacent molecules together, the nature of the underlying surface has little effect on this bi-component assembly. The large-scale highly ordered CA-M monolayer can be easily obtained on the atomically flat surfaces, demonstrating that this should be a universal approach to prepare hybrid TMDC/CA-M vdWHs.

Electrical and spectroscopic characteristics. The influence of the 2D CA-M network on the electronic characteristics of TMDCs has been investigated by means of electrical and spectroscopic characterizations. Monolayer MoS₂ back-gated FETs (Fig. 3a) are fabricated on n⁺⁺-Si/SiO₂ (270 nm oxide thickness) with patterned top Au source and drain electrodes (channel length (L)/width (W) amounting to 1.77 μ m/1.79 μ m, respectively). Transfer curves are recorded at source-drain bias $V_{DS} = 100$ mV before and after the CA-M network deposition to assess its effect on the device electrical characteristics (Fig. 3b). Upon coating with CA-M monolayer, a significant shift of the MoS₂ FET threshold voltage (ΔV_{th}) to the negative V_{GS} is observed (Fig. 3b and Supplementary Fig. 5). The off-state current increases from 0 to 1.1 μ A at $V_{GS} = -80$ V. This can be attributed to a strong electrostatic *n*-doping effect³⁰: the functionalized monolayer MoS₂ is degenerately doped showing reduced V_{GS} dependence in the transfer curves. The *n*-doping effect can be quantified from the shift of the ΔV_{th} , which linearly correlates to the charge carrier density change (Δn). Here, ΔV_{th} is estimated as 125 V (Supplementary Fig. 5a) after CA-M formation, leading to a Δn of 1.1×10^{13} cm⁻² for the transfer curve shown in Fig. 2b. The measurements are reproducible on five different devices, reaching a maximum Δn of 2.7×10^{13} cm⁻² and average Δn of $(1.9 \pm 0.8) \times 10^{13}$ cm⁻². The calculated magnitude of the charge transfer matches well with the simulation result, 3.6×10^{13} cm⁻² in Fig. 1a. To our knowledge, previously reported changes in electron density caused by organic molecules physisorbed onto monolayer TMDCs falls usually in the range of 10^{10} – 10^{12} cm⁻²,

even when thick molecular films are used^{31–34}. Significantly, our results indicate that when the CA-M molecules are arranged into crystalline H-bonded self-assembled 2D supramolecular structures the doping effect is enhanced. Meanwhile, the field-effect electron mobility (μ) of the CA-M functionalized MoS₂ monolayer results ca. 38 cm² V⁻¹ s⁻¹, being notably larger than the 14.3 cm² V⁻¹ s⁻¹ measured in pristine device. After CA-M coating, the increased electron density induces a decreased contact resistance and screening of Coulomb scatterings, yielding a μ increase. Further analysis of the device transfer curves indicate that hysteresis decreases from 27.5 V to 8.1 V upon CA-M coating (Supplementary Fig. 5b, c), which is translated into a significant reduction of scattering from the interfacial traps (from 2.2×10^{12} cm⁻² eV⁻¹ to 6.4×10^{11} cm⁻² eV⁻¹)³⁵. This provides the evidence for the screening of coulomb scattering from interfacial traps. Finally, the reversibility of our supramolecular functionalization strategy is confirmed by desorbing the molecules from the MoS₂ surface with hot water (>350 K), as evidenced by the FET transfer curves returning to the original state.

Raman and PL spectroscopies are powerful techniques to assess and quantify the electronic properties of 2DMs and monitor fundamental processes like doping³⁶. Here, we observed that the Raman signal from out-of-plane vibrations (A_{1g}) of monolayer MoS₂ is sensitive to the presence of the CA-M superlattices. The $E_{2g}^{1/2}$ mode at 386 cm⁻¹ and A_{1g} mode at 405 cm⁻¹ display a <20 cm⁻¹ difference from the bare MoS₂ flake, confirming its monolayer nature³⁷. CA-M coating results in a 1.3 ± 0.1 cm⁻¹ redshift of the A_{1g} peak and nearly no change in the $E_{2g}^{1/2}$ peak position (Fig. 3c). Doping in monolayer MoS₂ is reported to affect its A_{1g} vibration mode more significantly than $E_{2g}^{1/2}$, where A_{1g} redshift is a clear indication of the increase in the surface electron concentration of MoS₂³⁸. The *n*-doping of MoS₂ upon CA-M coating is also confirmed by PL spectroscopy. For monolayer MoS₂, the direct bandgap at K point enables strong PL emission^{37,39}, as displayed as a black curve in Fig. 3d for the non-coated flake. Two major peaks originate from the spin-orbit splitting of transition metal Mo, viz. the A exciton peak at lower energies (≈ 1.84 eV) and the B exciton peak at higher energies (≈ 2.05 eV). Upon CA-M deposition, the MoS₂ A exciton peak redshifts of ca. 25 ± 2 meV and undergoes significantly quenching, while the B exciton peak remains unchanged. Our observations agree with the literature for chemical doping of monolayer MoS₂ by charge-transfer^{37,38}. Similar results from the electrical and spectroscopical characterization of WSe₂ flakes confirm that the CA-M network acts as an *n*-type dopant on monolayer TMDCs (Supplementary Figs. 6 and 7). The reported assembly of H-bonded supramolecular networks as dopant of TMDC monolayers represents a simple, efficient, and reversible strategy to markedly improve device performance for various applications, such as in FETs and HER, as discussed hereafter.

Doping and surface morphology relationships. Frequently, the doping of 2DMs is achieved upon physisorption of thick molecular layers^{40,41}. Only a few previous works report the use of physisorbed SAMs to adjust the electrical properties of TMDCs, with the best Δn values amounting ca. 10^{12} cm⁻²^{12,13,32,42}. Here, the use of H-bonded 2D CA-M supramolecular networks yields an improvement of the electron density in monolayer MoS₂, with carrier density being at least one order of magnitude superior. To shed light on the role played by the CA-M monolayer on such doping, we studied the morphology of pristine and functionalized device's channel by AFM (Supplementary Fig. 8). The intermolecular multiple H-bonds determine a high stability of the CA-M 2D structure, which grows forming highly ordered molecular film with micrometer-scale crystalline domains, fully

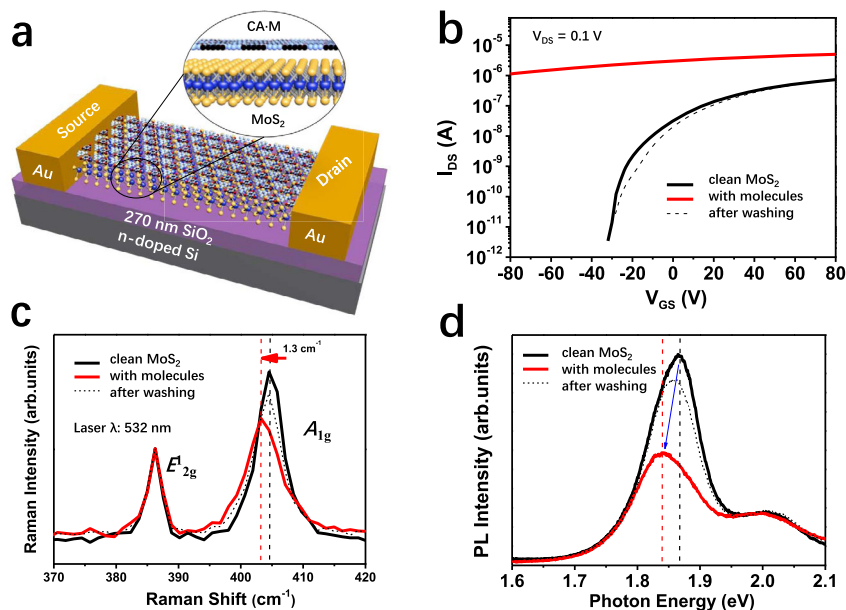


Fig. 3 Electrical and spectroscopic characteristics of monolayer MoS₂ FET with and without the functionalization of melamine and cyanuric acid co-assembled supramolecular network (CA-M). **a** Schematics of the MoS₂/CA-M hybrid device. **b** Transfer characteristics of MoS₂ FET. I_{DS} is the drain current and V_{GS} is the gate potential. **c** Raman and **d** Photoluminescence (PL) spectra of the device before (black curve) and after (red curve) CA-M functionalization, acquired under ambient conditions. As prepared MoS₂ samples (black solid), MoS₂/CA-M (red solid), device after wiping the CA-M network (black dotted). The peak position is labeled with vertical dashed lines, MoS₂ in black and MoS₂/CA-M in red. The arrows give the direction of the spectra shift: Raman A_{1g} peak shift in red and PL A exciton peak shift in blue.

covering the TMDC flake's surface. The surface roughness displays a very minor increase upon the physisorption of the CA-M film on both TMDCs, with values of 0.81 nm onto MoS₂ monolayer and 0.68 nm onto WSe₂ bilayer as quantified on a 1 μm^2 area. The defect-free, long-range crystalline structure of CA-M turns out to be crucial to promote such elevated doping. CA-M films produced at lower annealing temperatures (373 K instead of 430 K) onto WSe₂ FETs assemble into several smaller crystalline domains (of ca. 0.25 μm^2), resulting in a weaker doping effect (Supplementary Fig. 9).

Additionally, we analyze the contributions of the individual components of the supramolecular network, i.e., M and CA molecules on the morphological and electrical characteristics of TMDCs (Supplementary Figs. 10–13). Individual M molecules are observed to self-assemble into multiple small domains on MoS₂ (with an average size of $2.5 \times 10^5 \text{ nm}^2$), which are easily disassembled during thermal annealing (Supplementary Fig. 11). On the other hand, individual CA molecules form aggregates on the MoS₂ surface at room temperature (Supplementary Fig. 12). From AFM phase images, while the molecular film of individual M covers almost half of the device channel's surface, CA molecules display a greater propensity to undergo aggregation, leaving most of the flake surface uncoated. Such substantial dissimilarity in morphology between mono-component and bi-component films determines a different influence on the electrical characteristics of MoS₂ based FETs. For M-only coated devices the negative V_{th} shift is more pronounced than using CA molecules, viz. 28.1 V and 3.4 V, respectively (Supplementary Fig. 13). This leads to Δn of $2.2 \times 10^{12} \text{ cm}^{-2}$ for M-only and $3.7 \times 10^{11} \text{ cm}^{-2}$ for CA-only coatings on the MoS₂ FETs. For measurements taken on five similar devices, the average Δn amounts to $(1.9 \pm 0.3) \times 10^{12} \text{ cm}^{-2}$ and $(4.7 \pm 1.5) \times 10^{11} \text{ cm}^{-2}$ for M-only and CA-only, respectively, which are both significantly smaller than the values measured on H-bonded CA-M networks with a $\Delta n = (1.9 \pm 0.8) \times 10^{13} \text{ cm}^{-2}$. The tight packing of CA-M monolayer on TMDC flakes may also

act as an encapsulation layer to protect the device from the environment and subsequent degradation of performance.

Density of states and Work function modulation. Further insights into the electronic properties of the MoS₂/CA-M hybrid system can be obtained by elucidating the DOS and WF. The doping of a semiconductor shifts its E_F position, ultimately influencing the material's WF^{43,44}. Alongside, E_F of a 2D semiconductor is particularly sensitive to modification with organic molecules^{45,46}. E_F is known to shift towards the conduction band minimum (CBM) in the case of *n*-type doping, leading to a lower WF, while *p*-type doping causes the opposite effect by shifting E_F towards the valence band maximum (VBM).

The evolution of the WF of a monolayer MoS₂ surface is measured by Kelvin probe force microscope (KPFM), which allows to record a contact potential map. As shown in Fig. 4a, d, freshly evaporated Au electrodes are used to ground the single MoS₂ flakes deposited on a 270 nm SiO₂ substrate. Darker regions in the KPFM images (Fig. 4b, e) correspond to larger WF values, indicating that the WF of both pristine and CA-M-doped MoS₂ surfaces is smaller compared to the Au electrodes, as shown by the brighter contrast of the flakes. The contact potential difference related to Au increases from 50 mV to 170 mV with CA-M coating. The presence of the CA-M network decreases the WF by 120 meV compared to the pristine MoS₂ surface, evidencing a *n*-doping effect, as determined by averaging the SP values on five monolayer MoS₂ flakes and six CA-M coated monolayer MoS₂ samples (Supplementary Table 1). Note here that SP differences of nano-objects determined via amplitude modulation KPFM are usually underestimated due to the finite size of the probe, which limits the spatial and potential resolution of the measurement⁴⁷. The calculated WF values (Supplementary Fig. 14a) revealed a negative shift in WF by 220 meV, in good agreement with the KPFM results. These results indicate that due

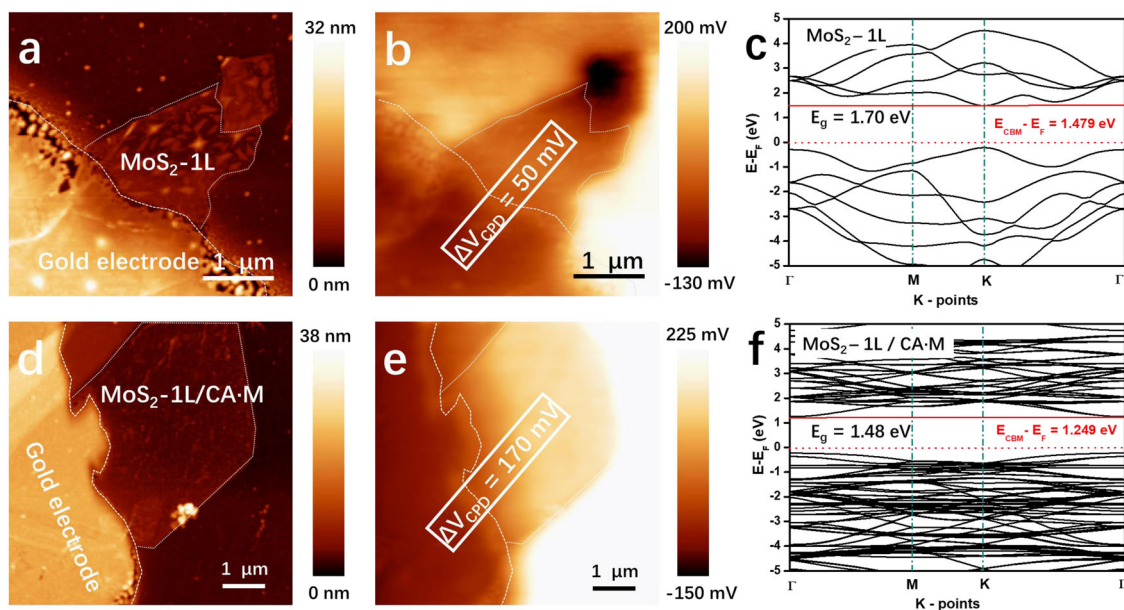


Fig. 4 Morphology and energetics of coated and uncoated MoS₂ monolayers. The area of monolayer MoS₂ flake (1L-MoS₂) is circled by white dotted lines. **a** Atomic force microscope (AFM) height image and **b** Kelvin probe force microscope (KPFM) image of a monolayer MoS₂ flake. **c** Calculated electronic band structure of monolayer MoS₂. **d** AFM image and **e** KPFM image of a CA-M doped MoS₂ monolayer flake. **f** The band structure of monolayer MoS₂/CA-M heterostructure. Contact potential difference (ΔV_{CPD}) is the contact potential difference between the E_F of the flake and that of the Au electrode ($E_{F,MoS_2} - E_{F,Au}$)/e. The energy level of conduction band minimum (E_{CBM}) is marked with red solid lines, and the Fermi level (E_F) is marked with the red dashed lines. E_g is the energy gap between conduction band minimum and valence band maximum.

to the interaction between the CA-M and MoS₂ surface, the removal of an electron from the whole hybrid system is much easier than from the CA-M SAM or MoS₂ surface separately.

The DOS simulations are also in good agreement with the literature⁴⁴ and corroborate the experimental results, proving that the CA-M network determines n-type doping on monolayer MoS₂ surface. The band structure maps are shown in Fig. 4c, f: the energy gap from E_F to CBM decrease from 1.479 to 1.249 eV after introducing the CA-M SAM. The upward shift of E_F leads to an increased probability of exciting electrons to CBM. The n-type doping on MoS₂ mostly originates from the charge transfer effect of CA-M molecular network, which can be read from the partial density of states (PDOS) plots-derived energy level map (Supplementary Fig. 15).

The MoS₂ monolayer flakes coated with the individual molecular components (either M or CA) were also measured via KPFM, as depicted in Supplementary Fig. 17. Coating with M causes a reduction of the WF of pristine MoS₂ of $\Delta WF = -80$ meV, thus being of smaller magnitude compared to the CA-M system. The WF does not change significantly in CA-coated flakes because of the propensity of the molecules to undergo strong aggregation. When coated with a CA-M superlattice the bilayer WSe₂ displays a negative WF shift of 95 meV (Supplementary Fig. 18), which is in good agreement with the simulated monolayer WSe₂, exhibiting a $\Delta WF = -170$ meV (Supplementary Fig. 14b). The experimental WF results from KPFM are summarized in Supplementary Table 1.

High-efficiency HER catalytic performance. Electrochemical water splitting is considered one of the most promising approaches towards the production of hydrogen fuel. Platinum (Pt) is so far the best performing and most employed electrocatalysts for the hydrogen evolution reaction (HER)⁴⁸, yet its high cost and scarcity greatly limit large-scale applications. Among various possible alternatives to Pt, MoS₂ – material abundant in nature –

exhibits Gibbs free energy adequate for atomic hydrogen adsorption⁴⁹. Although the MoS₂ catalytic activity has been significantly improved over the last decade^{50,51}, it is still far from the HER performance of Pt. The catalytic activity of MoS₂ is known to be affected by its interaction with supporting substrates via charge transfer^{52,53}. CA-M co-crystal nanosheets have been reported as supports for CoP nanoparticles catalyst, giving embedded composite with high hydrogen evolution activity and stability⁵⁴. Since the CA-M network has been proved as a strong n-dopant to MoS₂, it is also expected to act as a suitable buffer layer to boost the catalytic performance of MoS₂. In HER, the adsorbed protons sequester electrons from the MoS₂ surface thereby forming hydrogen molecules through a process of desorption from the surface. Therefore, the electron density on the MoS₂ surface can indirectly affect the rate of hydrogen evolution to a certain extent. The electron transfer in the CA-M/MoS₂ heterostructure occurs from CA-M to MoS₂, hence increasing the charge density on MoS₂ surface and further enhancing the HER activity. Multilayer CA-M film is prepared by drop-casting 200 μ L of CA and M mixture solution at 1×10^{-6} M concentration on a 1 cm² 270-nm-thick SiO₂ wafer, and then annealed overnight under vacuum at 450 K. The obtained CA-M multilayer film exhibits roughness of 0.29 nm, allowing the easy deposition of mechanically exfoliated monolayer MoS₂ flake on its surface (Supplementary Fig. 19). Raman and PL characterizations confirm that upon inverting the position of the supramolecular network with respect to MoS₂ (i.e., MoS₂ on CA-M) a strong n-doping effect is still induced on the upper MoS₂ monolayer.

The typical three-electrode electrochemical system is employed to evaluate the HER performance of CA-M/MoS₂. A micron-sized window is opened on a photoresist passivation layer on the device surface to expose only the MoS₂ flake to the electrolyte (0.5 M H₂SO₄). Here, monolayer MoS₂ flake is used as working electrode (WE), while Ag/AgCl and pencil graphite are used as reference (RE) and counter (CE) electrodes, respectively (see Fig. 5a and Supplementary Fig. 20 for details). The catalytic performance of

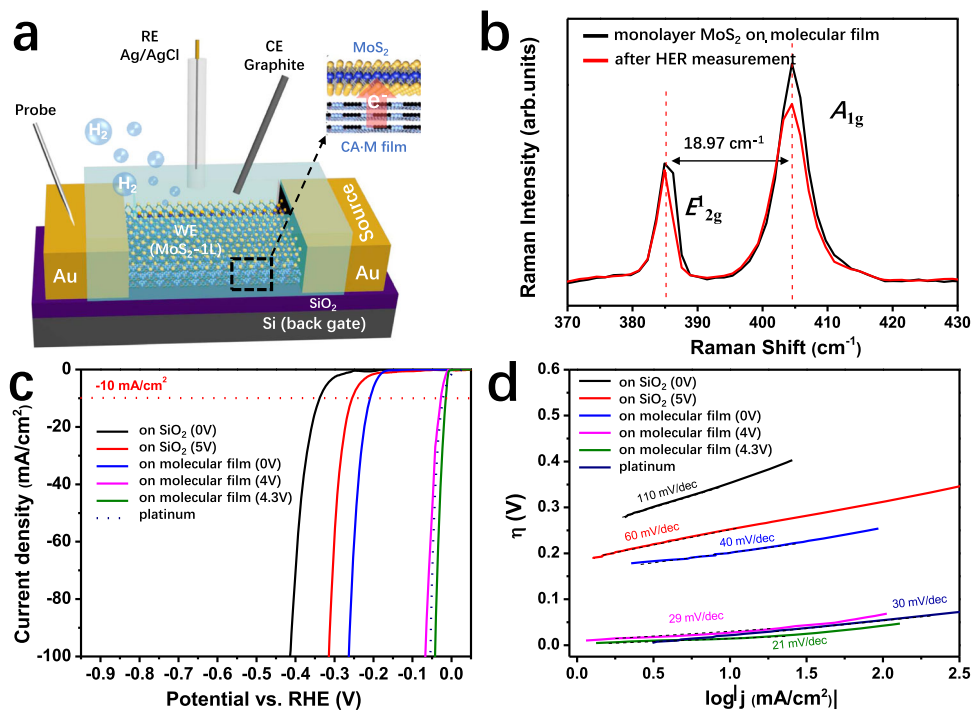


Fig. 5 Hydrogen evolution reaction (HER) in the co-assembled supermolecule network (CA-M)/MoS₂ heterostructure. **a** Monolayer MoS₂ on CA-M molecular film device for HER measurements. The red arrow indicates the charge transfer direction in the heterostructure. **b** Raman spectra of the monolayer MoS₂ flake on CA-M before (black) and after (red) HER measurements, and Raman peak positions are marked by the dashed lines. **c** The polarization curves for HER on Platinum (dashed blue line) and monolayer MoS₂ flakes under different conditions (SiO₂ substrate with/without 2D CA-M buffer layer, and the respective voltages applied gate). The -10 mA cm^{-2} current density position is marked by the dashed red line. **d** Corresponding Tafel curves.

neat monolayer MoS₂ flake shows a clear improvement upon addition of the underlying CA-M thin film, as exhibited in Fig. 5c. The HER overpotential (η) is defined as the potential at the standard current density (j) -10 mA cm^{-2} . Here, the CA-M/MoS₂ hybrid device reaches $\eta = 208 \text{ mV}$, $\sim 130 \text{ mV}$ smaller than that for the pristine MoS₂ surface ($\eta = 340 \text{ mV}$), indicating a great enhancement of energy conversion efficiency. The participation of CA-M film improves the electrical coupling between the substrate and the MoS₂ catalyst, therefore the injection of electrons from the electrode and their transport to the catalyst active site is facilitated. Moreover, the CA-M strong n -doping effect induces an excessive negative charge density on the MoS₂ surface that boosts the HER performance. Figure 5d depicts the Tafel slope, an important parameter that quantifies the amount of additional applied potential is required to observe a logarithmic increase in current density during HER. The CA-M/MoS₂ hybrid device shows a Tafel slope of 40 mV/dec , which is significantly superior to the values reported for most 2H MoS₂ related compounds (typically $41\text{--}120 \text{ mV/dec}$)⁵⁵, and even comparable with that for metallic 1T MoS₂ (43 mV/dec)⁵⁶. The integrity of the MoS₂ flakes is confirmed by Raman spectroscopy carried out after the HER measurements, which endorses the material as a very stable high-performing catalyst.

The conductivity of the MoS₂ channel undergoes a sharp increase when a positive V_{GS} is applied to the FETs, which is considered being a powerful strategy for enhancing the HER catalytic performance^{57,58}. The FET characteristics of CA-M/MoS₂ validates that the bottom multilayer CA-M film can induce significant electron density increases on MoS₂ upon back-gate V_{GS} modulation (Supplementary Fig. 21). Moreover, when MoS₂ surface is immersed in a sulfuric acid electrolyte, the applied positive gate voltage has also been confirmed to induce the excess electron density on the Mo atoms near the S vacancy sites, leading

to enhanced Mo-H bond strengths (stabilizing H proton adsorption)⁵⁹. The HER catalytic property of our field-tuned CA-M/MoS₂ devices is largely boosted with V_{GS} increasing (Supplementary Fig. 23c). In particular, the CA-M/MoS₂ channel achieves a better performance than standard polycrystalline Pt catalyst when the CA-M/MoS₂ vdWHs is gated at $V_{\text{GS}} = +4.3 \text{ V}$, exhibiting a $\eta = 14 \text{ mV}$ (20 mV for Pt) and a small Tafel slope of 21 mV/dec (30 mV/dec for Pt). Interestingly, the field effect on the bare monolayer MoS₂ surface is much weaker than on the CA-M/MoS₂ hybrid surface. When applying V_{GS} from 0 V to $+4 \text{ V}$, the over potential decreases by 62 mV for monolayer MoS₂ and by impressive 180 mV for the CA-M/MoS₂ hybrid structure (Fig. 5c). The strong n -doping effect from the underlying CA-M film enables directional electron flow to MoS₂ active sites, which determines the excellent catalytic performance of this hybrid structure to produce molecular hydrogen. Such a strategy of utilizing the electric field to improve the catalytic activity of HER could also be used in the other electrochemical processes. However, the use of more complicated four electrodes system in cyclic voltammetry experiments may hinder to some extent this gate-enhanced HER strategy for large-scale technological applications.

Discussion

vdWHs based on 2DMs is attracting ever-growing attention due to their unique electronic features that can be engineered on demand. In this context, the use of the small organic building blocks capable of forming supramolecular lattices through non-covalent interactions represents a simple, yet effective solution towards the modulation of 2DM electronic properties. Here, we show that the bi-component, hydrogen-bonded CA-M 2D network can be used for boosting the electron density of TMDCs through a positive cooperative effect in the charge transfer

process as a function of the packing density of the molecular adlayer. We have found that the physisorption of a tightly packed 2D supramolecular network of CA-M onto the monolayer MoS₂ channel determines an electron transfer up to $2.72 \times 10^{13} \text{ cm}^{-2}$, thereby reaching state-of-the-art performance compared to the reported molecule-TMDCs hybrid devices. Optical and electrical analyses corroborated with DFT calculations reveal that the CA-M 2D network induces a charge transfer effect to MoS₂, regardless if placed on top or underneath the 2DMs. Owing to the excellent charge transfer effect, CA-M 2D network is used as a substrate to enhance the HER catalyst performance of monolayer MoS₂. Remarkably, the field-effect enhanced HER performance of the CA-M/MoS₂ hybrid device outperforms the catalytic efficiency of Pt. 2DMs decorated with supramolecular 2D networks may expand the library of heterostructures with on-demand electronic characteristics. Moreover, such an approach can be further employed in multilayer vdWHs displaying electrical, optical, and magnetic functionalities. Compared with classical vdW 2DMs heterostructures, the structural programmability of 2D supramolecular networks enables optimal matching of the crystal structures to form ad hoc superlattices. Importantly, supramolecular chemistry can offer a choice of 2D networks and patterns, with tunable electronic properties, periodic potentials, and atom packing density. The atomic precision achieved through controlled molecular self-assembly can therefore be instrumental for the optimization of charge carriers in 2DMs, and be used to impart novel properties, enabling the realization of multifunctional, high-performance optoelectronic devices.

Methods

STM and AFM. STM measurements were carried out by using a Veeco scanning tunneling microscope (multimode Nanoscope III, Veeco). Bulk MoS₂ and WSe₂ crystals (HQ graphene) were used as substrates before and after CA-M coating. The employed STM tips were mechanically cut from a Pt/Ir wire (80/10, diameter 0.25 mm). The images were obtained at room temperature in air, and the drift of the piezo was corrected using the underlying chalcogen atomic lattice as a reference. AFM imaging was performed using a Bruker Dimension Icon set-up operating in air, in tapping mode, by using tip model TESPA-V2, tip stiffness K is 42 N m⁻¹.

Raman and photoluminescence spectra. Raman spectra were carried out in air by Renishaw inVia spectrometer equipped with 532 nm laser, and the wavenumber resolution was 1 meV. The Si Raman peak at 520.5 cm⁻¹ was used for normalization. The PL spectra were acquired with a Renishaw inVia spectrometer equipped with a 532 nm laser. TMDC flakes were mechanically exfoliated from crystals onto SiO₂/Si substrate. The excitation power was kept below 1 mW to avoid the local heating and damaging effect.

FET fabrication and characterization. MoS₂ and WSe₂ flakes were mechanically exfoliated down to monolayer by the Scotch-tape-based method and further transferred to CA-M/SiO₂ and bare SiO₂ substrates (270 nm SiO₂ thickness onto *n*-doped Si from Fraunhofer IPMS, Germany). Top-gated FET devices were fabricated by maskless optical lithography, with Au source and drain electrodes thermally evaporated onto the patterned substrate. Warm acetone was used for the lift-off process, and the as-fabricated devices were annealed at 430 K in vacuum to remove atmospheric adsorbates. All the FET devices were measured in an N₂-filled glovebox with a probe station connected to a Keithley 2636 A source-meter unit.

The carrier mobility μ was determined from the following Eq. (1):

$$\mu = \frac{dI_{\text{DS}}}{dV_{\text{DS}}} \times \frac{L}{WC_i V_{\text{DS}}} \quad (1)$$

L and W are the channel length and width, and C_i is the insulator capacitance per unit of area.

The dopant-induced charge carrier density increase (Δn) was calculated from Eq. (2):

$$\Delta n = C_i \frac{\Delta V_{\text{th}}}{e} \quad (2)$$

Where ΔV_{th} is the difference in threshold voltage (V_{th}) before and after doping effect, and e is the elementary charge ($1.6 \times 10^{-19} \text{ C}$).

KPFM. Topography and surface potential images were simultaneously collected with Pt/Ir coated silicon probes (Bruker SCM-PIT-V2, resonant frequency $\approx 75 \text{ kHz}$, $k \approx 3 \text{ N m}^{-1}$) at ambient conditions using a Bruker Icon AFM employed in amplitude modulation KPFM (AM-KPFM) mode. For work function (WF) referencing and electrical grounding purposes, a polycrystalline Au electrode was thermally evaporated with a shadow mask on the flakes, before and after the molecule adsorption. The reported surface potential values are referred to the average SP of the non-treated Au electrode ($0.16 \pm 0.10 \text{ eV}$) obtained from >30 samples. The calibration of the Au WF ($4.85 \pm 0.09 \text{ eV}$) was determined on the polycrystalline Au electrodes via macroscopic Kelvin Probe (KP) at ambient conditions (Ambient Kelvin Probe Package from KP Technology Ltd, 2-mm-diameter gold tip amplifier)⁶⁰. The calibration of the KP probe was performed against a freshly cleaved HOPG surface (4.475 eV). The SPM image processing has been done on WSxM software.

Device fabrication for HER measurements. Devices for HER measurements were produced by spin coating a layer of photoresist (AZ1505) onto monolayer MoS₂ transferred to n^{++} -Si/SiO₂ substrate. A window was opened on the photoresist by mask-less optical lithography to expose the desired flake area to the solution during measurements. The HER catalytic performance of monolayer MoS₂ was measured in a three-electrode configuration in 0.5 M H₂SO₄ electrolyte (purged with Ar gas), with an Ag/AgCl reference electrode and pencil graphite counter electrode (diameter 3 mm). The MoS₂ flake acts as the working electrode during linear sweep voltammetry (LSV) from 0.1 to -0.6 V , with a scan rate of 3 mV s^{-1} . All LSV curves were collected after 5 times scan in the same potential range. HER measurements were also performed while biasing the n^{++} -Si gate electrode, as depicted in Fig. 4a. The electrochemical current density is calculated by normalizing the current to the area of the exposed window on the MoS₂. The data has been referenced to $E_{\text{RHE}} = E_{\text{Ag/AgCl}} + 0.210 \text{ V}$.

DFT calculations. First-principles calculations within the density functional theory (DFT) framework were performed to investigate different types of self-assembled monolayers (CA and M) adsorbed on the MoS₂ surface structures. By means of DFT calculations the adsorption of CA-M SAM on the different surface structures was investigated. Here an important role was played by the small distance ($\sim 3.3 \text{ \AA}$) between the SAM and the surface, which can influence the electronic properties of the whole system. The WFs were simulated by DFT. In order to identify the most stable configurations, we performed first-principles total-energy calculations using the gradient corrected (PBE) density functional theory as implemented in the Vienna ab initio simulation package^{61,62}. The geometry of the MoS₂/CA-M hybrid structure was optimized by relaxing the atomic positions of all the atoms. The equilibrium geometry was assumed to be reached when the forces on the relaxed atoms of the system were less than $0.025 \text{ eV/\AA}$. A $(2 \times 2 \times 1)$ Monkhorst-Pack k -point mesh has been used to span the surface Brillouin zone. The system was modeled by using a periodic supercell of $9.81 \times 9.81 \times 18 \text{ \AA}^3$. The bond length parameters as well as the unit cell of the structure were chosen to fit the experimental observations. The electronic wave functions were expanded into plane waves up to an energy cutoff of 400 eV, and a projected-augmented-wave scheme⁶³ was used in order to describe the interactions between the valence electrons and the nuclei(ions).

Data availability

The data that support the findings of this study are available from the corresponding authors on request.

Received: 19 June 2021; Accepted: 6 December 2021;

Published online: 26 January 2022

References

- Novoselov, K. S. et al. Electric field effect in atomically thin carbon films. *Science* **306**, 666–669 (2004).
- Frisenda, R. et al. Recent progress in the assembly of nanodevices and van der Waals heterostructures by deterministic placement of 2D materials. *Chem. Soc. Rev.* **47**, 53–68 (2018).
- Novoselov, K. S., Mishchenko, A., Carvalho, A. & Castro Neto, A. H. 2D materials and van der Waals heterostructures. *Science* **353**, aac9439 (2016).
- Liu, X. & Hersam, M. C. Interface characterization and control of 2D materials and heterostructures. *Adv. Mater.* **30**, 1801586 (2018).
- Frisenda, R., Molina-Mendoza, A. J., Mueller, T., Castellanos-Gomez, A. & van der Zant, H. S. J. Atomically thin p–n junctions based on two-dimensional materials. *Chem. Soc. Rev.* **47**, 3339–3358 (2018).
- Kum, H. et al. Epitaxial growth and layer-transfer techniques for heterogeneous integration of materials for electronic and photonic devices. *Nat. Electron.* **2**, 439–450 (2019).

7. Daukiya, L., Seibel, J. & De Feyter, S. Chemical modification of 2D materials using molecules and assemblies of molecules. *Adv. Phys.* **4**, 1625723 (2019).
8. Wang, Q. H. & Hersam, M. C. Room-temperature molecular-resolution characterization of self-assembled organic monolayers on epitaxial graphene. *Nat. Chem.* **1**, 206–211 (2009).
9. Cho, K., Pak, J., Chung, S. & Lee, T. Recent advances in interface engineering of transition-metal dichalcogenides with organic molecules and polymers. *ACS Nano* **13**, 9713–9734 (2019).
10. Huang, Y. L. et al. The organic–2D transition metal dichalcogenide heterointerface. *Chem. Soc. Rev.* **47**, 3241–3264 (2018).
11. Phillipson, R. et al. Tunable doping of graphene by using physisorbed self-assembled networks. *Nanoscale* **8**, 20017–20026 (2016).
12. Gobbi, M. et al. Periodic potentials in hybrid van der Waals heterostructures formed by supramolecular lattices on graphene. *Nat. Commun.* **8**, 14767 (2017).
13. Gobbi, M. et al. Collective molecular switching in hybrid superlattices for light-modulated two-dimensional electronics. *Nat. Commun.* **9**, 3689 (2018).
14. Cai, Z.-F., Yan, H.-J., Wang, D. & Wan, L.-J. Potential- and concentration-dependent self-assembly structures at solid/liquid interfaces. *Nanoscale* **10**, 3438–3443 (2018).
15. Kudernac, T., Lei, S., Elemans, J. A. A. W. & De Feyter, S. Two-dimensional supramolecular self-assembly: nanoporous networks on surfaces. *Chem. Soc. Rev.* **38**, 402–421 (2009).
16. Blunt, M. O. et al. Temperature-induced structural phase transitions in a two-dimensional self-assembled network. *J. Am. Chem. Soc.* **135**, 12068–12075 (2013).
17. Yan, X. et al. Supramolecular polymers with tunable topologies via hierarchical coordination-driven self-assembly and hydrogen bonding interfaces. *Proc. Natl. Acad. Sci. USA* **110**, 15585 (2013).
18. Seto, C. T. & Whitesides, G. M. Self-assembly based on the cyanuric acid-melamine lattice. *J. Am. Chem. Soc.* **112**, 6409–6411 (1990).
19. Theobald, J. A., Oxtoby, N. S., Phillips, M. A., Champness, N. R. & Beton, P. H. Controlling molecular deposition and layer structure with supramolecular surface assemblies. *Nature* **424**, 1029–1031 (2003).
20. Piot, L. et al. Selective formation of Bi-component arrays through H-bonding of multivalent molecular modules. *Adv. Funct. Mater.* **19**, 1207–1214 (2009).
21. Llanes-Pallas, A. et al. Engineering of supramolecular H-bonded nanopolygons via self-assembly of programmed molecular modules. *J. Am. Chem. Soc.* **131**, 509–520 (2009).
22. Zhang, H.-M. et al. One-Step preparation of large-scale self-assembled monolayers of cyanuric acid and melamine supramolecular species on Au(111) surfaces. *J. Phys. Chem. C* **112**, 4209–4218 (2008).
23. Korolkov, V. V. et al. Supramolecular networks stabilise and functionalise black phosphorus. *Nat. Commun.* **8**, 1385 (2017).
24. Zhang, H.-M. et al. Preparing self-assembled monolayers of cyanuric acid and melamine complex on HOPG surfaces. *J. Phys. Chem. C* **113**, 13940–13946 (2009).
25. Liu, Z. et al. Electronic decoupling of organic layers by a self-assembled supramolecular network on Au(111). *J. Phys. Chem. Lett.* **10**, 4297–4302 (2019).
26. Biroju, R. K. et al. Hydrogen evolution reaction activity of graphene–MoS₂ van der Waals heterostructures. *ACS Energy Lett.* **2**, 1355–1361 (2017).
27. Shi, J. et al. Temperature-mediated selective growth of MoS₂/WS₂ and WS₂/MoS₂ vertical stacks on Au foils for direct photocatalytic applications. *Adv. Mater.* **28**, 10664–10672 (2016).
28. Hunter, C. A. & Anderson, H. L. What is cooperativity? *Angew. Chem. Int. Ed.* **48**, 7488–7499 (2009).
29. Ranganathan, A., Pedireddi, V. R. & Rao, C. N. R. Hydrothermal synthesis of organic channel structures: 1:1 hydrogen-bonded adducts of melamine with cyanuric and trithiocyanuric acids. *J. Am. Chem. Soc.* **121**, 1752–1753 (1999).
30. Bertolazzi, S., Gobbi, M., Zhao, Y., Backes, C. & Samori, P. Molecular chemistry approaches for tuning the properties of two-dimensional transition metal dichalcogenides. *Chem. Soc. Rev.* **47**, 6845–6888 (2018).
31. Liu, B. et al. High-performance chemical sensing using schottky-contacted chemical vapor deposition grown monolayer MoS₂ transistors. *ACS Nano* **8**, 5304–5314 (2014).
32. Wang, Y., Gali, S. M., Slassi, A., Beljonne, D. & Samori, P. Collective dipole-dominated doping of monolayer MoS₂: orientation and magnitude control via the supramolecular approach. *Adv. Funct. Mater.* **30**, 2002846 (2020).
33. Ji, H. G. et al. Chemically tuned p- and n-type WSe₂ monolayers with high carrier mobility for advanced electronics. *Adv. Mater.* **31**, 1970301 (2019).
34. Chen, K. et al. Air stable n-doping of WSe₂ by silicon nitride thin films with tunable fixed charge density. *APL Mater.* **2**, 092504 (2014).
35. Park, R. S. et al. Hysteresis in carbon nanotube transistors: measurement and analysis of trap density, energy level, and spatial distribution. *ACS Nano* **10**, 4599–4608 (2016).
36. Luo, P. et al. Doping engineering and functionalization of two-dimensional metal chalcogenides. *Nanoscale Horiz.* **4**, 26–51 (2019).
37. Li, H. et al. From bulk to monolayer MoS₂: evolution of Raman scattering. *Adv. Funct. Mater.* **22**, 1385–1390 (2012).
38. Dhakal, K. P. et al. Confocal absorption spectral imaging of MoS₂: optical transitions depending on the atomic thickness of intrinsic and chemically doped MoS₂. *Nanoscale* **6**, 13028–13035 (2014).
39. Splendiani, A. et al. Emerging photoluminescence in monolayer MoS₂. *Nano Lett.* **10**, 1271–1275 (2010).
40. Han, C. et al. Surface functionalization of black phosphorus via potassium toward high-performance complementary devices. *Nano Lett.* **17**, 4122–4129 (2017).
41. Cheng, L. et al. Sub-10 nm tunable hybrid dielectric engineering on MoS₂ for two-dimensional material-based devices. *ACS Nano* **11**, 10243–10252 (2017).
42. Wang, Y. et al. Doping of monolayer transition-metal dichalcogenides via physisorption of aromatic solvent molecules. *J. Phys. Chem. Lett.* **10**, 540–547 (2019).
43. Park, S. et al. Demonstration of the key substrate-dependent charge transfer mechanisms between monolayer MoS₂ and molecular dopants. *Commun. Phys.* **2**, 109 (2019).
44. Sun, J. et al. Lateral 2D WSe₂ p–n homojunction formed by efficient charge-carrier-type modulation for high-performance optoelectronics. *Adv. Mater.* **32**, 1906499 (2020).
45. Almadori, Y., Bendiab, N. & Grévin, B. Multimodal kelvin probe force microscopy investigations of a photovoltaic WSe₂/MoS₂ type-II interface. *ACS Appl. Mater. Interfaces* **10**, 1363–1373 (2018).
46. Tosun, M. et al. MoS₂ heterojunctions by thickness modulation. *Sci. Rep.* **5**, 10990 (2015).
47. Liscio, A., Palermo, V. & Samori, P. Nanoscale quantitative measurement of the potential of charged nanostructures by electrostatic and kelvin probe force microscopy: unraveling electronic processes in complex materials. *Acc. Chem. Res.* **43**, 541–550 (2010).
48. McCrory, C. C. L. et al. Benchmarking hydrogen evolving reaction and oxygen evolving reaction electrocatalysts for solar water splitting devices. *J. Am. Chem. Soc.* **137**, 4347–4357 (2015).
49. Jaramillo, T. F. et al. Identification of active edge sites for electrochemical H₂ evolution from MoS₂ nanocatalysts. *Science* **317**, 100 (2007).
50. Li, H., Jia, X., Zhang, Q. & Wang, X. Metallic transition-metal dichalcogenide nanocatalysts for energy conversion. *Chem* **4**, 1510–1537 (2018).
51. Yang, L., Liu, P., Li, J. & Xiang, B. Two-dimensional material molybdenum disulfides as electrocatalysts for hydrogen evolution. *Catalysts* **7**, 285 (2017).
52. Li, G. et al. Engineering substrate interaction to improve hydrogen evolution catalysis of monolayer MoS₂ films beyond Pt. *ACS Nano* **14**, 1707–1714 (2020).
53. Hong, M., Shi, J., Huan, Y., Xie, Q. & Zhang, Y. Microscopic insights into the catalytic mechanisms of monolayer MoS₂ and its heterostructures in hydrogen evolution reaction. *Nano Res.* **12**, 2140–2149 (2019).
54. Wu, J. et al. Graphene-like hydrogen-bonded melamine–cyanuric acid supramolecular nanosheets as pseudo-porous catalyst support. *Adv. Mater.* **33**, 2007368 (2021).
55. Ding, Q., Song, B., Xu, P. & Jin, S. Efficient electrocatalytic and photoelectrochemical hydrogen generation using MoS₂ and related compounds. *Chem* **1**, 699–726 (2016).
56. Lukowski, M. A. et al. Enhanced hydrogen evolution catalysis from chemically exfoliated metallic MoS₂ nanosheets. *J. Am. Chem. Soc.* **135**, 10274–10277 (2013).
57. Wang, J. et al. Field effect enhanced hydrogen evolution reaction of MoS₂ nanosheets. *Adv. Mater.* **29**, 1604464 (2017).
58. Liu, X. et al. The critical role of electrolyte gating on the hydrogen evolution performance of monolayer MoS₂. *Nano Lett.* **19**, 8118–8124 (2019).
59. Wang, Y. et al. Field effect modulation of electrocatalytic hydrogen evolution at back-gated two-dimensional MoS₂ electrodes. *Nano Lett.* **19**, 6118–6123 (2019).
60. Kahn, A. Fermi level, work function and vacuum level. *Mater. Horiz.* **3**, 7–10 (2016).
61. Kresse, G. & Hafner, J. Ab initio molecular dynamics for liquid metals. *Phys. Rev. B* **47**, 558–561 (1993).
62. Kresse, G. & Joubert, D. From ultrasoft pseudopotentials to the projector augmented-wave method. *Phys. Rev. B* **59**, 1758–1775 (1999).
63. Blöchl, P. E. Projector augmented-wave method. *Phys. Rev. B* **50**, 17953–17979 (1994).

Acknowledgements

We acknowledge funding from European Commission through the ERC project SUPRA2DMAT (GA-833707), the Graphene Flagship Core 3 project (GA- 881603), and the Marie Curie ITN project UHMob (GA-811284), the Agence Nationale de la Recherche through the Labex projects CSC (ANR-10-LABX-0026 CSC) and NIE (ANR-11-LABX-0058 NIE) within the Investissement d’Avenir program (ANR-10-120 IDEX-0002-02), the International Center for Frontier Research in Chemistry (icFRC) and the Institut Universitaire de France (IUF).

Author contributions

The experiment was conceived by C.W., L.C., A.C. and P.S.; C.W. performed the AFM, STM characterization; C.W. and Y.Z. carried out FET device study and Raman characterization; K.J., D.T. and X.Z. performed the DFT calculations; C.W., R.F.O. and H.Z. performed the HER measurements; N.T. worked on KPFM; C.M. worked on the capacitance measurement; B.H. performed the Au electrode transfer; P.S. and A.C. supervised the entire study. C.W. wrote the manuscript with comments and suggestions from all co-authors.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41467-022-28116-y>.

Correspondence and requests for materials should be addressed to Artur Ciesielski or Paolo Samori.

Peer review information *Nature Communications* thanks the anonymous reviewers for their contribution to the peer review of this work. Peer reviewer reports are available.

Reprints and permission information is available at <http://www.nature.com/reprints>

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2022