Ultrahigh resistance of hexagonal boron nitride to mineral scale formation

Interfacial interactions play a fundamental role in many aqueous processes including adsorption, catalytic reaction, corrosion, filtration, and scale formation. In particular, scale formation, i.e., the development of mineral deposits on a material surface due to precipitation from the bulk solution and/or crystal formation initiated by surface nucleation, has great impacts on interfacial transfer of mass, heat, electrons, and light. It causes profound performance decline in numerous industrial processes, such as impaired heat transfer in heat exchangers and boilers, increased pressure drop in pipes, flow blockage in filtration membranes, corrosion damage of steam turbines, decreased conductivity and activity of electrodes, premature failure of heating and electrochemical components etc.\(^1\)\(^-\)\(^4\), all leading to higher operation cost and safety risk. It is reported that economical loss due to mineral scaling in boilers, turbines, and heat exchangers accounts for 0.17–0.25% of the gross domestic product (GDP) in industrialized nations\(^5\). Understanding mineral scaling behavior is important for the development of next generation materials and technologies that address these critical challenges.

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Formation of mineral scale on a material surface has profound impact on a wide range of natural processes as well as industrial applications. However, how specific material surface characteristics affect the mineral-surface interactions and subsequent mineral scale formation is not well understood. Here we report the superior resistance of hexagonal boron nitride (hBN) to mineral scale formation compared to not only common metal and polymer surfaces but also the highly scaling-resistant graphene, making hBN possibly the most scaling resistant material reported to date. Experimental and simulation results reveal that this ultrahigh scaling-resistance is attributed to the combination of hBN’s atomically-smooth surface, in-plane atomic energy corrugation due to the polar boron-nitrogen bond, and the close match between its interatomic spacing and the size of water molecules. The latter two properties lead to strong polar interactions with water and hence the formation of a dense hydration layer, which strongly hinders the approach of mineral ions and crystals, decreasing both surface heterogeneous nucleation and crystal attachment.
Scale formation can occur through the deposition of mineral crystals formed in the bulk solution, as well as through surface-induced heterogeneous nucleation with crystals growing from nucleation sites on a surface. Both processes are strongly influenced by material surface properties. Similar to deposition of other particles, material properties affect the attachment of mineral crystals via hydrophobic and electrostatic interactions. Surface-induced heterogeneous nucleation is a more thermodynamically favorable process, but it is poorly understood because it happens on very small time and length scales. Few previous studies have investigated the different surface properties that influence surface-induced heterogeneous nucleation: roughness, charge, and hydrophobicity. Surface roughness is directly related to the number of nucleation sites; it is generally recognized that mineral crystallization increases with surface roughness. However, findings on the impact of charge and surface hydrophobicity have been inconsistent. For example, some studies found that surface charge influenced the heterogeneous nucleation via electrostatic interactions or complexation reactions with the mineral ions, while others reported similar nucleation rates on surfaces of different charges. Contradictory results have also been reported on the role of surface hydrophobicity in mineral scaling. Hydrophilic coatings such as graphene oxide (GO), graft polymers, and polyethylene glycol, were shown to delay the onset of CaCO₃ scaling in some studies, while other studies showed that hydrophilic surfaces promoted CaCO₃ nucleation, and GO had no anti-scaling effect. One possible reason for such apparent contradiction is that modification of a surface property (e.g., hydrophobicity or charge) often leads to inevitable changes in other surface properties, which makes it difficult to discern the role of an individual surface property for development of anti-scaling materials.

Two dimensional (2D) materials possess atomically smooth surfaces, and have attracted tremendous interest for their potential applications in processes where interfacial interactions play a critical role. For example, graphene was shown to effectively inhibit nucleation of metals and metal oxides in nonaqueous phases. Theoretical and experimental research on water-surface interactions and related phenomena, such as ultrafast water transport, de-icing, and anti-fouling, also suggest that the atomically smooth morphology plays an important role in preventing adhesion. However, no studies have investigated the scaling behavior, especially heterogeneous nucleation, on the surface of 2D nanomaterials such as graphene and hexagonal boron nitride (hBN) in aqueous solutions. Furthermore, 2D materials vary widely in surface chemistry despite their common feature of atomic-level smoothness: Graphene consists of a single layer of carbon atoms arranged in a honeycomb lattice nanostructure, featuring a small lattice constant, low in-plane polarity, and high hydrophobicity; hBN, another important 2D material with its lattice structure and lattice constant similar to those of graphene, has high in-plane polarity due to the boron-nitrogen bond and higher hydrophilicity than graphene. It is unknown how such differences in surface chemistry influence mineral scale formation on these atomically smooth surfaces.

Here we investigate mineral scale formation on graphene and hBN surface, and compare it to that on metal (i.e., titanium (Ti)) and polymer (i.e., polyvinylidene fluoride (PVDF)) surfaces commonly used for high scaling potential applications (Supplementary Note 1, Supplementary Figs. 1 and 2). Surface induced heterogeneous nucleation on these surfaces are investigated using both real time, in situ measurements as well as ex situ characterization methods. We also directly quantify the binding force of mineral crystals grown from surface-induced heterogeneous nucleation. Experimental measurements combined with density functional theory (DFT) illustrate the effect of surface chemistry on the formation of the hydration layer and its key role in mineral ion-surface interactions. Very importantly, we discover hBNs outstanding anti-scaling properties and demonstrate its potential application as an anti-scaling coating in practical engineering systems.

**Results**

**Anti-scaling behavior of hBN**

To evaluate the scaling behaviors of graphene and hBN, graphene and hBN nanocoatings grown on flat Cu substrate were dipped in supersaturated CaSO₄ solution (50 mmol L⁻¹, saturation index (SI) of 3.28) and characterized using a video camera for 220 min (Fig. 1A). As the induction time of homogeneous nucleation for CaSO₄ at this concentration (<40 min) is much shorter than the operation time, both bulk precipitation due to homogeneous nucleation and surface induced heterogeneous nucleation contribute to scale formation. As
shown in Fig. 1B and Supplementary Movie 1, CaSO4 crystals were found on the pristine Cu surface after 20 min, and severe scale formation (0.123 ± 0.014 mg mm⁻²) was observed after 220 min of testing. On the graphene surface, CaSO4 crystals occurred after 40 min, and increased to 0.015 ± 0.001 mg mm⁻² at the end of the experiment (Fig. 1C). However, the hBN-coated Cu surface exhibited almost no scale formation (0.001 ± 0.001 mg mm⁻²) during the 220 min experiment, except at its edges where the hBN nanocoating was damaged due to sample cutting (Fig. 1B).

After illustrating the macroscopic anti-scaling behavior of graphene and hBN using video camera, we investigated the scale formation caused by surface-induced heterogeneous nucleation on graphene, hBN, Ti, and PVDF using microscopic ex situ and in situ measurements with supersaturated CaCO3. CaCO3 and CaSO4 are both common scalants in water, but CaCO3 has a solubility (0.12 mmol L⁻¹) two orders of magnitude lower than CaSO4 (19.10 mmol L⁻¹). It allows using much lower concentration to prepare supersaturated CaCO3 solution with saturation index similar to supersaturated CaSO4, which would generate smaller amount of scalants on the surfaces and is conducive to performing microscopic ex situ or in situ characterizations. As shown in Fig. 2A, supersaturated CaCO3 (SI of 1.18) was introduced to the test surfaces immediately after preparation before homogeneous nucleation occurred (Supplementary Note 2, Supplementary Fig. 3). Vaterite was the main CaCO3 crystal formed before homogeneous nucleation occurred (Supplementary Note 2, Fig. 2F). In over 15 h, almost no vaterite was observed on Ti (Fig. 2B), followed by PVDF (Fig. 2C), graphene (Fig. 2D), and hBN (Fig. 2E). The same trend was observed at different reaction time (Fig. 2F). In over 15 h, almost no vaterite was observed on hBN (0.4 ± 0.2 mm²), while 55.6 ± 3.3, 13.2 ± 4.7, and 5.6 ± 0.8 vaterite crystals mm⁻² were found on Ti, PVDF, and graphene surfaces, respectively (Fig. 2G). The very few vaterite crystals formed on hBN were much smaller in size than those formed on the other surfaces. These results suggest prolonged vaterite heterogeneous nucleation induction time and slower vaterite formation kinetics on the hBN surface.

Real time, in situ measurement of CaCO3 nucleation and growth rates using the quartz crystal microbalance with dissipation (QCMD) technique (Fig. 2A) further confirms that hBN greatly hinders surface-induced heterogeneous nucleation of CaCO3 (Fig. 2H, I). Upon introduction of an influent solution supersaturated with CaCO3, adsorption of ions (e.g., Ca²⁺, CO₃²⁻) led to an immediate albeit small change in frequency and dissipation, which reached equilibrium in ~1 h. The larger frequency and dissipation changes observed for the Ti and PVDF surfaces suggested more adsorption of Ca²⁺ and CO₃²⁻ ions than that on the graphene and hBN surfaces. The similar frequency and dissipation changes on graphene and hBN surface suggested that they had similar ion adsorption, a result of their similar surface charge. After the initial ion adsorption, greater frequency and dissipation changes occurred, and exhibited two distinct stages. The first stage (stage i) featured relatively slow changes in frequency (dF/dt) and dissipation (dD/dt), signaling the formation of vaterite nuclei, i.e., the induction process. This was followed by a much faster frequency shift (stage ii), which signaled crystal growth from the nuclei previously formed on the surface (Fig. 2I). CaCO3 mass accumulation rate calculated using the Sauerbrey equation (Supplementary Fig. 5) showed a very large increase in stage ii compared to that in stage i on Ti, PVDF
and graphene surfaces. As shown in Fig. 2I, J, the Ti surface showed the shortest induction time and the fastest crystal growth kinetics, followed by PVDF. The heterogeneous nucleation induction time on graphene was significantly longer compared to Ti and PVDF (Fig. 2I), and a much slower crystal growth kinetics was observed (Fig. 2J and Supplementary Fig. 5). On the hBN surface, the frequency and dissipation shifts throughout the experiment were very small. It did not exhibit a distinct stage II with fast frequency and dissipation changes. This suggests that the induction time on the hBN surface is longer than the approximately 8h contact time with the test solution in the experiment; no detectable crystal growth occurred during this period (Fig. 2I, J and Supplementary Fig. 5). This may be attributed to the low ion adsorption on hBN (Fig. 2I), which prolongs the induction of heterogeneous nucleation and hinders vaterite crystal growth. After 10 h of operation, the system was flushed with ultrapure water. Frequency and dissipation signals approached the original DI water baseline at different rate for all samples (Fig. 2I), confirming that the frequency and dissipation changes observed earlier resulted from the nucleation and growth of vaterite, which was removed or dissolved when flushed with ultrapure water.

**Atomic smoothness decreases nucleation and binding**

Although the four surfaces are all nominally flat and smooth at the macroscopic length scale, their nanoscale morphologies differ significantly (Fig. 3A). The root-mean-square roughness values of Ti (2.0 ± 0.2 nm) and PVDF (1.2 ± 0.2 nm) are almost an order of magnitude higher than those of graphene (0.3 ± 0.1 nm) and hBN (0.2 ± 0.1 nm), which are atomically smooth with no observable sags, crests, or other surface irregularities. During the nucleation and crystal growth stages, the nanoscale rough features on Ti and PVDF serve as heterogeneous nucleation sites, provide cratered surface for nucleus attachment, and increases friction that may hinder local flow velocity, all conducive to vaterite nucleation and crystal growth, resulting in a shorter induction stage and faster vaterite growth (Fig. 2I, J).

During the crystal growth stage, the higher roughness of Ti and PVDF surfaces increases the contact area with the vaterite crystals, resulting in stronger binding forces. We quantify the lateral force required to detach vaterite crystals from the surface using a nanoindenter inside a SEM (Supplementary Note 3, Supplementary Fig. 6). The applied load increases linearly with the lateral travelling distance of the nanoindenter tip relative to the vaterite crystal, until the vaterite crystal is pushed off the surface (Fig. 3E–H, and Supplementary Movies 2 and 3), i.e., the “detaching point”. The applied load at the detaching point is defined as the detaching force, which reflects the binding force between the vaterite crystal and the sample surface. The contact area (Fig. 3G) normalized detaching force followed the order of Ti (44.9 ± 12.7 μN μm⁻²) > PVDF (27.6 ± 3.4 μN μm⁻²) > hBN (12.1 ± 4.3 μN μm⁻²) > graphene (11.1 ± 6.1 μN μm⁻²), correlating well with the surface roughness (Figs. 2J and 3D). The atomically smooth graphene and hBN exhibit notably lower binding forces. Note that these forces are much higher than typical colloidal adhesion forces, a notable distinction between scaling due to crystal formation from surface-induced heterogeneous nucleation and that due to deposition of mineral crystals formed in the bulk solution.

The lower binding force on the atomically-smooth graphene and hBN surfaces measured in our study is consistent with findings from previous studies that show atomically smooth carbon and boron-nitride materials exhibit super-lubricity at solid/solid rigid junctions and peculiar interactions with water in 1D nanotubes or 2D structures at their solid/liquid interfaces. The reduced binding force allows easier detachment of the mineral crystals and decreases the formation of scales under hydraulic shear.

**Dense hydration layer mitigates nucleation**

The similarity in their atomically smooth surface morphology is apparently responsible for the similarly in low vaterite detachment forces on graphene and hBN. The lower nucleation rate on hBN compared to graphene, however, suggests the difference in surface chemistry may play an important role. This is supported by DFT calculations. As a result of the delocalized π system in graphene, the charge distribution on graphene surface is much more uniform than on hBN, with a small dipole moment (0.036 D) and moderate carbon atom electronegativity (2.55) (Fig. 4A, B, Supplementary Fig. 7, Supplementary Note 4). In hBN, N atoms share their lone pair electrons with B, and the electrons of the π system are more localized on the N atoms. The electronegativity of N (3.04) is therefore notably higher than B (2.04), leading to corrugated charge distribution on hBN surface with a dipole moment of 0.047 D (Fig. 4A). The highest occupied molecular orbital (HOMO) is more localized on N atoms, and the

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**Fig. 3 | Atomically-smooth morphology yields low scalant binding force.** AFM image of A Ti, B PVDF, C graphene, D hBN surfaces; E–H Measurement of vaterite detaching force from various coatings. E Sectional view schematic showing the measurement method by adding a lateral force on a vaterite crystal grown on a sample surface. Top view SEM image of a vaterite crystal (F) before and G after being pushed off from the sample surface by the nanoindenter tip. The scale bars in F and G represent 10 μm. The dashed circle in G shows the contact area between the vaterite crystal and the underlying sample surface after detachment; H Applied force load increases linearly with the moving depth of the nanotip upon contact to the vaterite crystal; I The average vaterite detaching force and surface roughness of the four nanocoatings. Error bars represent the standard deviation of corresponding results.
lowest unoccupied molecular orbital (LUMO) is more concentrated on B atoms. The non-uniform charge distribution of the hBN surface has an important influence on its interaction with polar water molecules. The alternating negative and positive charge regions on hBN attract H and O atoms in water, respectively, leading to stronger interaction with water molecules than on graphene surface, where the charge is more uniform. As a result, water molecules are more closely packed and located closer to the hBN surface: the average distance between water molecules is 2.45 Å, and the closest distance between water molecule and the hBN surface is around 2.92 Å (Fig. 4C–E and Supplementary Fig. 8), consistent with previously reported values from quantum Monte Carlo simulations. In comparison, on graphene surface, the closest water molecules are located 2.95 Å away from the surface, and the distance between water molecules measures 2.55 Å on average. Similar results have been reported in previous studies. Utilizing ab initio molecular dynamics simulation, it was reported that the friction coefficient on hBN was about 3 times larger than that on graphene, and the adsorbed water molecule had faster slippage on graphene than on hBN.

The water molecule packing on the surface has direct impact on ion-surface interactions. Simulation of interactions between Ca²⁺, CO₃²⁻, or CaCO₃ with hBN or graphene in the presence and absence of water (Supplementary Figs. 9–11) show that hBN binds more strongly with Ca²⁺, CO₃²⁻, and CaCO₃ than graphene in the absence of water molecules (Supplementary Fig. 9), due to the locally charged B and N having higher attraction to the charged Ca²⁺, CO₃²⁻, and polar CaCO₃ species. However, when water molecules are introduced, the distance between the surface and Ca²⁺ or CaCO₃ is larger on hBN than on graphene (Fig. 4E and Supplementary Fig. 11). This is attributed to the denser hydration layer on hBN, which hinders the approach of Ca²⁺ and CaCO₃ to the surface. More energy (1.0 eV) is required for a Ca²⁺ ion to penetrate the hydration layer on hBN and graphene surface calculated by DFT. Detailed calculation and simulation processes and results can be seen in Supplementary Note 4. The formation of hydration layer on hBN and graphene surface characterized by QCMD. G QCMD experimental setup. H Frequency and dissipation change on hBN and graphene surfaces with changes in gas flow humidity; I The effect of gas flow humidity on average frequency decrease and the absolute ratio of dissipation change over frequency decrease (ΔD/ΔF).

Fig. 4 | Effect of surface chemistry on nucleation behavior of CaCO₃ on hBN and graphene surface. a) DFT simulated charge distribution on hBN and graphene surfaces. Blue and red spheres in a) represent boron and nitrogen atoms, respectively. Silver, light blue, and light red spheres in b) represent carbon atoms (Supplementary Fig. 7). The edge of both hBN and graphene flakes are passivated with H atoms; DFT simulated interaction of H₂O, Ca²⁺, CO₃²⁻, and CaCO₃ with c) hBN and d) graphene surface (Supplementary Fig. 11); e) DFT calculated distance between water and surface, between water molecules, and between solutes (Ca²⁺, CO₃²⁻, or CaCO₃) and hBN or graphene surfaces. Gra: graphene; F Minimum energy needed for a Ca²⁺ ion to penetrate the hydration layer on hBN and graphene surface calculated by DFT. Detailed calculation and simulation processes and results can be seen in Supplementary Note 4.
Experimental measurements of water vapor adsorption confirm the simulation results. Adsorption of water vapor on graphene and hBN surfaces was measured using the QCMD technique at varying relative humidity of the influent N₂/H₂O mixed gas (Fig. 4G). Both graphene and hBN coated quartz sensors exhibit a decrease in the resonate frequency with the increase in the relative humidity of the influent gas (Fig. 4H), indicating increasing adsorption of water molecules on graphene and hBN surfaces. The hBN coated sensor experiences larger frequency decreases than the graphene coated sensor at all humidity levels (Fig. 4H, I), suggesting more adsorption of water on hBN than on graphene. Furthermore, the ratio of dissipation increase over frequency change (ΔD/Δf) is smaller on hBN than on graphene (Fig. 4I). As water slippage on surfaces has little impact on ΔD and ΔhBN, the measured ΔD/Δf directly reflects the viscoelastic properties of the adsorbed water layer. The smaller ΔD/Δf on hBN indicates that the water layer adsorbed on the polar hBN surface is more compact and viscous compared to that on the uniformly charged graphene, consistent with the simulation results that a denser and more compact water layer forms on hBN. Although several studies have hinted the higher affinity of hBN for water54,55, high dielectric constant56, high thermal conductivity57,61, make hBN possibly the most anti-scaling material known, with notable advantages over graphene. The atomic strength54,55, high dielectric constant56, high thermal conductivity57,61, make hBN possibly the most anti-scaling material known, with notable advantages over graphene. The atomic

The close match between the distribution of hBN surface energy corrugation (charge nonuniformity) and the dipole of water molecules leads to strong polar interactions between the hBN surface and polar water molecules, and hence compact packing of water molecules on hBN surface. Combining an atomically smooth surface, large in-plane polarity, and a lattice constant matching best with the size of water molecules (Supplementary Table 1), hBN is possibly one of the most anti-scaling materials. The understanding of how hBN interacts with water can also provide insights to other processes that occur in aqueous solutions, such as absorption, lubrication, catalysis, and corrosion etc.

**Anti-scaling applications**

Besides the flat Cu substrate, we grew hBN coating on both inside and outside surfaces of stainless steel (SS) pipes using the chemical vapor deposition (CVD) method (Fig. 5A), and assessed scale formation behaviors of the pristine and hBN-coated SS pipes by pumping real produced water collected from an oil & gas production site in Texas through the pipes (Fig. 5B, Supplementary Note 5). These experiments represent performance of the hBN coating on more realistic substrates in complex solutions. As shown in Fig. 5C, significant scale formation occurred inside the pristine SS tube after 7 days of operation, and its effective inner diameter decreased by ~17.2% after 14 days of operation, resulting in a 31.3% reduction of cross-sectional area for water passage. In contrast, the hBN-coated SS tube only experienced slight fouling by organic matter in the produced water at the outlet (Fig. 5D), with no measurable scale formation throughout the tube. These results demonstrate the excellent anti-scaling potential of the hBN nanocoating for real water and wastewater applications.

Overall, the unprecedented scaling resistance of hBN combined with its other unique properties, including superior mechanical strength45–47, high dielectric constant48, high thermal conductivity47, and chemical and thermal stability48–51, make hBN an excellent candidate for multi-functional coatings in various industrial processes (e.g. fluid transmission, heat exchange, membrane separation). These coatings can protect substrate materials from scaling or corrosion52, as well as ensure energetically efficient fluid flow, high heat transfer rate, and long material lifetime.

In summary, we discover that the unique combination of atomically smooth surface, high in-plane polarity, and proper interatomic spacing (lattice constant) makes hBN possibly the most anti-scaling material known, with notable advantages over graphene. The atomic
smoothness of both hBN and graphene surfaces reduces the number of surf ace heterogeneous nucleation sites as well as the binding force between the mineral crystal and the surface, resulting in much greater resistance to mineral scaling compared to conventional scale-resistant materials. Interestingly, hBN exhibits even greater resistance to surface-induced heterogeneous nucleation than graphene, which stems from the in-plane dipole of the boron-nitrogen bond at a scale closely matching that of water molecules, leading to in-plane atomic energy corrugation that favors interaction with water. Direct experimental measurements and DFT calculations show that such chemical structure results in the formation of a dense hydration layer on the hBN surface, which strongly hinders mineral ions from approaching the surface and hence suppresses the heterogeneous nucleation process. The hBN nanocoating grown on a stainless-steel tube exhibits outstanding anti-scaling properties in real oil & gas produced water, demonstrating its potential application in practical engineering systems. The results of the study provide important insights for future development of novel functional materials by manipulating their interactions with surrounding media. On the other hand, the scalability, durability, and long-term scaling resistance of the hBN coating as well as the specific roles of substrate materials and defects in large scale coatings need to be evaluated before practical applications can be possible.

Methods

Synthesis and characterization of the coatings

Ti, PVDF, graphene and hBN coatings were formed on Cu foil (25 μm in thickness, McMaster-CARR, USA). Before coating, the Cu substrate was electrochemically polished. Ti films of 40 nm in thickness were formed on the Cu substrate using a sputter coater (Denton Desk V, Denton Vacuum, USA) at a current of 30 mA. PVDF was coated on the Cu substrate via spin coating. A 1 wt% PVDF in DMF solution was applied at a rotation speed of 2000 rpm for 1 min, followed by overnight drying at 60 °C before use.

Graphene and hBN films were formed on Cu substrate by chemical vapor deposition (CVD). During growth, the electrochemically polished Cu foil was first loaded into a tube furnace and heated to 1000 °C, followed by annealing at 1 Torr 15% H2/Ar for 20 min. The graphene growth was then carried out by feeding 15% H2/Ar (flow rate of 100 sccm) and methane (flow rate of 10 sccm) to the furnace for 20 min. For hBN growth, ammonia borane was used as the precursor. After Cu foil annealing, ammonia borane was heated to ~85 °C for evaporation upstream of the tube furnace, and was carried by H2 gas to the Cu substrate to grow hBN at 1000 °C for 30 min. After growth with graphene or hBN, the Cu foil was rapidly cooled down to room temperature for further use.

After fabrication, the morphology of the four coatings were characterized by scanning electron microscopy (SEM, Quanta FEG 250, Thermo Fisher Scientific, USA) and atomic force microscopy (AFM) (N200, Park, Suwon, Korea). The chemical property of the graphene and hBN were characterized by Raman spectroscopy (Renishaw inVia, UK) and X-ray photoelectron spectroscopy (XPS) (PH Quantera II, Physical Electronics, USA) (Supplementary Note 1, Supplementary Figs. 1 and 2).

Evaluation of scale formation on graphene and hBN using video camera

The graphene and hBN nanocoatings were formed on flat Cu substrates, and their scaling resistance was evaluated in a solution supersaturated with CaSO4, a common scalar in ground water that was very difficult to remove. The supersaturated CaSO4 solution (30 mmol L−1) was prepared by mixing 100 mmol L−1 CaCl2 with 100 mmol L−1 Na2SO4 at a 1:1 volume ratio. The prepared solution had a pH of 7.0 and a saturation index (SI) of 3.28. SI is defined as the ratio between the chemical activity product of the mineral ions and their solubility product63. During the experiments, the pristine Cu foil, hBN- and graphene-grown Cu foils were immersed in the CaSO4 solution. Scale formation was monitored using a video camera for 220 min (Supplementary Movie 1). The mass of mineral precipitation was determined by measuring the sample mass before and after the experiment.

Ex situ measurement of scaling caused by heterogeneous nucleation on various coatings

Coated samples were cut into round wafers and installed in a flow cell. Supersaturated CaCO3 solution was prepared by mixing CaCl2 and Na2CO3 solutions, which were prepared using ultrapure water and aerated with air for 24 h to reach equilibrium before use. As shown in Fig. 2A, the equilibrated CaCl2 and Na2CO3 solutions were continuously fed into an inline mixer at 0.1 mL min−1 and then the flow cell at a combined flow rate of 0.2 mL min−1. The chemistry of the mixed feed solution is simulated using Visual MINTEQ (version 3.1, KTH): Ca2+ and CO32− activity are 1.23 × 10−8 and 1.24 × 10−4 mol L−1, respectively; pH is 8.54, ionic strength is 5.6 × 10−3, and calcite, aragonite, and vaterite saturation indexes ([Ca2+]2[CO3]2/Ksp) are 4.45, 3.18, and 1.18 respectively. The induction time for homogeneous nucleation was determined by dynamic light scattering (DLS) measurements using a NanoBrook Omni (Brookhaven Instrument, Holtsville, NY, USA) (Supplementary Note 2, Supplementary Fig. 3). Since the hydraulic retention time from the mixer to the samples is much shorter than the induction time for CaCO3 homogeneous nucleation, formation of CaCO3 crystals on the sample surface is attributed to surface-induced heterogeneous nucleation. After the experiments, samples were retrieved and characterized using scanning electron microscopy (SEM, Quanta FEG 250, Thermo Fisher Scientific, USA) (Supplementary Note 2, Fig. 2B–E and Supplementary Fig. 4). The number and size of the crystals formed were analyzed using ImageJ software. More than 10 SEM images were analyzed for each surface to collect the crystal number and size data. The particle size distribution data were fitted with a Gaussian distribution function to obtain the average and standard deviation of particle size.

Real-time, in situ characterization of scale formation caused by heterogeneous nucleation using the QCMD technique

The heterogeneous nucleation and crystal formation of CaCO3 was also characterized in situ and in real time using the quartz crystal microbalance with dissipation (QCMD) technique (Qsense E4 analyzer, Bolin Scientific, Sweden), which has a mass sensitivity and a dissipation sensitivity of 1.8 ng cm−2 and 0.1 × 10−4 in liquid, respectively64. To perform the QCMD experiments, the four test materials were first coated on QCMD sensors. Ti and PVDF were coated on QCMD sensors using the same methods as those used for coating the Cu substrate. Graphene and hBN were first grown on Cu substrate, and then transferred onto QCMD sensors using a PMMA-assisted method. In this method, a PMMA film was spin-coated on the graphene- or hBN-grown on the Cu foil. The sample was then immersed in a FeCl3 solution to dissolve the Cu substrate. The graphene or hBN layer immobilized on the PMMA film was then transferred to DI water, and subsequently collected onto the QCMD sensor surface with the graphene or hBN side attaching to the sensor surface. Finally, acetone and IPA were used to dissolve the PMMA layer, exposing the graphene or hBN surface. The coated sensors were then mounted in 4 parallel QCMD cells, and characterized for CaCO3 nucleation and growth at a temperature of 22 °C using the same supersaturated solution and flow rates as the offline scaling experiments described above. The frequency and dissipation data were continuously monitored for each coated sensor.

The increase of adsorbed or precipitated mass (Δm, ng cm−2) on the sensors was calculated from the frequency change using the
Sauerbrey equation:

\[ \Delta m = - C \frac{\Delta f}{n} \]  

(1)

Here \( C \) is the mass sensitivity constant, which is related to the properties of the quartz sensors and equals 17.7 ng cm\(^{-2}\) Hz\(^{-1}\) in this study. \( \Delta f \) is the frequency change (Hz). \( n \) is the harmonic number.

**Binding force measurement**

To measure the binding force between the vaterite crystals and the various surfaces that they grow from, samples were prepared by flowing the mixed CaCl\(_2\) and Na\(_2\)CO\(_3\) solution over the four surfaces for 15 h to allow vaterite growth. After crystal growth, the samples are transferred into a SEM assembled with nanoindenter equipment. To perform the measurement, the tip of the nanoindenter approaches and pushes a vaterite crystal grown on sample surface at a constant speed of 0.03 \( \mu \)m s\(^{-1}\), until the vaterite crystal is detached from the underlying surface (Supplementary Note 3, Supplementary Fig. 6, Supplementary Movies 2 and 3). During this process, the applied load is recorded, and the applied load at the detaching point for the vaterite crystal is referred to as the detaching force. After normalizing the detaching force by contact area between the vaterite crystal and the surface, the obtained force reflects the binding force between the vaterite crystal and the underlying coating.

As shown in Supplementary Fig. 6 and Supplementary Movies 2 and 3, the nanotip can push the vaterite crystal at a direction perpendicular (Supplementary Fig. 6a, Supplementary Movie 2) or parallel (Supplementary Fig. 6b, Supplementary Movie 3) to the plane of the semi-hexagonal vaterite crystal. The detaching force differs greatly depending on the direction in which the load is applied (Supplementary Fig. 6a4, b4). This is due to the anisotropy structure of the vaterite crystal. As shown in Supplementary Fig. 6a, when pushing the crystal at the perpendicular direction, the detaching force is also greatly affected by the location of the contact point (e.g., distance from the coating surface). Therefore, data reported in the manuscript are all obtained by pushing the vaterite crystal in the direction parallel to the vaterite plane as shown in Supplementary Fig. 6b.

**Simulation and calculation**

In this study, we used density function theory (DFT) to evaluate (i) the charge distribution on graphene and hBN, (ii) the interaction between water molecules and hBN or graphene surface, (iii) the interaction of Ca\(^{2+}\), CO\(_3\(^{2-}\)\), and CaCO\(_3\) with bare hBN and graphene, and (iv) the interaction between Ca\(^{2+}\), CO\(_3\(^{2-}\)\), and CaCO\(_3\) and hydrated hBN or graphene. Detailed calculation and simulation methods can be found in the Supplementary Note 4 and Supplementary Figs. 7–11.

In addition to calculation, we also used the QCMD technique to quantitively verify the simulation results by measuring the adsorption of water vapor on graphene and hBN surfaces utilizing N\(_2\)/H\(_2\)O mixed gas at different humidity levels. In these experiments, a customized humidity control device was installed between a N\(_2\) gas tank and the QCMD analyzer (Fig. 4G). N\(_2\) gas with different humidity was continuously flown into the QCMD analyzer at a flow rate of 7.5 mL min\(^{-1}\). Corresponding frequency and dissipation change were recorded at 22 \( ^\circ\)C.

**Evaluating scale formation potential using real produced water**

We also grew a hBN nanocoating inside a stainless-steel tube, and compared its scale forming behavior with an uncoated stainless-steel tube when exposed to a real oil & gas produced water taken from Texas. The produced water has a total organic carbon (TOC) concentration of 108 mg L\(^{-1}\) and conductivity of 148 mS cm\(^{-1}\). Detailed water quality data are shown in Supplementary Table 2 (Supplementary Note 5). As minerals have precipitated out from the solution during transportation of the produced water sample, the received produced water is no longer supersaturated (Supplementary Table 2) and cannot be directly utilized for scaling experiment. Instead, the oil & gas produced water is spiked with 50 mM CaCl\(_2\) or Na\(_2\)SO\(_4\) to prepare the respective Ca\(^{2+}\)- or SO\(_4\(^{2-}\)\)-rich solutions. During the experiment, the two solutions were continuously fed into an inline mixer at 0.1 mL min\(^{-1}\) and then flowed through the pristine and hBN coated SS tubes at a combined flow rate of 0.2 mL min\(^{-1}\). Scale formation was monitored using a camera during 14 days of operation.

**Data availability**

All data generated or analyzed during this study are reported in this published article and its supplementary information files, and are available from the author upon request.

**Code availability**

All codes supporting the findings of this study are available from the corresponding author upon request.

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Author contributions
All authors contributed intellectually to this paper. K.Z., X.H. designed and performed the experiments, analyzed data, and drafted the manuscript. Q.L., J.L., P.M.A., W.W., M.E. and P.J.J.A. conceived the idea, designed the experiments, revised the manuscript, and led the project. X.Z., T.Z., H.G. assisted sample growth and characterization. E.F.O. performed the simulation.

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Competing interests
The authors declare no competing interests.

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