

## DIELECTRICS

# High energy density in artificial heterostructures through relaxation time modulation

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Electrostatic capacitors are foundational components of advanced electronics and high-power electrical systems owing to their ultrafast charging-discharging capability. Ferroelectric materials offer high maximum polarization, but high remnant polarization has hindered their effective deployment in energy storage applications. Previous methodologies have encountered problems because of the deteriorated crystallinity of the ferroelectric materials. We introduce an approach to control the relaxation time using two-dimensional (2D) materials while minimizing energy loss by using 2D/3D/2D heterostructures and preserving the crystallinity of ferroelectric 3D materials. Using this approach, we were able to achieve an energy density of 191.7 joules per cubic centimeter with an efficiency greater than 90%. This precise control over relaxation time holds promise for a wide array of applications and has the potential to accelerate the development of highly efficient energy storage systems.

Managing high energy density has become increasingly important in applications ranging from electric power systems to portable electronic devices (1–3). Electrostatic capacitors have been widely used for high energy storage and release owing to their ultrafast charge and discharge rate, but their performance is limited by the low maximum polarization ( $P_m$ ) of conventional dielectric materials (4, 5). By contrast, ferroelectric materials such as  $\text{HfO}_2$ ,  $\text{ZrO}_2$ , and  $\text{BaTiO}_3$  (BTO) can achieve higher maximum polarization because of their higher electric susceptibilities, which are related to dielectric constants (6, 7). However, their high remnant polarization ( $P_r$ ) limits the effectiveness of energy storage and release during the discharging process (8). To overcome this limitation, relaxor-ferroelectric materials have been studied for their ability to achieve high energy densities with low remnant polarization. Through compositional and defect design, nanodomains have been introduced into ferroelectric materials to realize relaxor ferroelectricity, offering a potential avenue for developing high-performance electrostatic capacitors (9, 10). These methodologies lead to a

redistribution of domain walls that can serve as effective relaxor-like defects to suppress the formation of large polar domains and reduce the remnant polarization. Although recent advances have shown great promise in realizing relaxor ferroelectricity by inducing nanodomains in ferroelectric materials through composition and defect engineering, these approaches result in the loss of crystallinity, which leads to reduced permittivity and sacrifices the maximum polarization. Moreover, some approaches are restricted from achieving a high polarization saturation limit, which results in polarization saturation at a low electric field and leads to low energy density ( $U_e$ ), despite substantial breakthroughs otherwise (11). These limitations highlight the need for a different approach that can supplement previous methods and lead to the development of electrostatic capacitors with extremely high energy density.

We introduce a strategy for precise control of the relaxation time of polarization that maintains minimal energy loss by using monolayer two-dimensional (2D) materials produced by the layer-splitting technique (12). We achieved this by using artificially designed 2D/single-

crystalline 3D/2D (2D/C-3D/2D) heterostructures. We used a layer-transfer technique to produce freestanding single-crystalline  $\text{BaTiO}_3$  (C-BTO) in which both interfaces can be manipulated, and we formed 2D/C-3D/2D heterostructures by the addition of various 2D materials. Unlike previous approaches that deteriorate ferroelectric materials by involving structural changes, our approach preserves the single-crystal nature of the BTO. Instead, we sandwiched a C-BTO layer with 2D materials in the form of a freestanding membrane, such that the Maxwell-Wagner (MW) effect, a relaxation by charge accumulation at heterogeneous interfaces (13, 14), takes place at the interfaces to change the relaxation time. Atomically precise control of the thickness of 2D materials by layer-resolved splitting (15, 16) enables minimal energy loss and tangent delta ( $\tan \delta$ ), a dielectric loss due to electrical phase difference (12, 17), while controlling relaxation time. Using this strategy, we effectively suppressed the remnant polarization of ferroelectric materials while maintaining maximum polarization. We show that this allows for an energy density of 191.7 J/cm<sup>3</sup> with an efficiency greater than 90%. We believe our approach has the potential to enhance the performance of dielectric materials and be of use in other related applications that require high-energy storage systems.

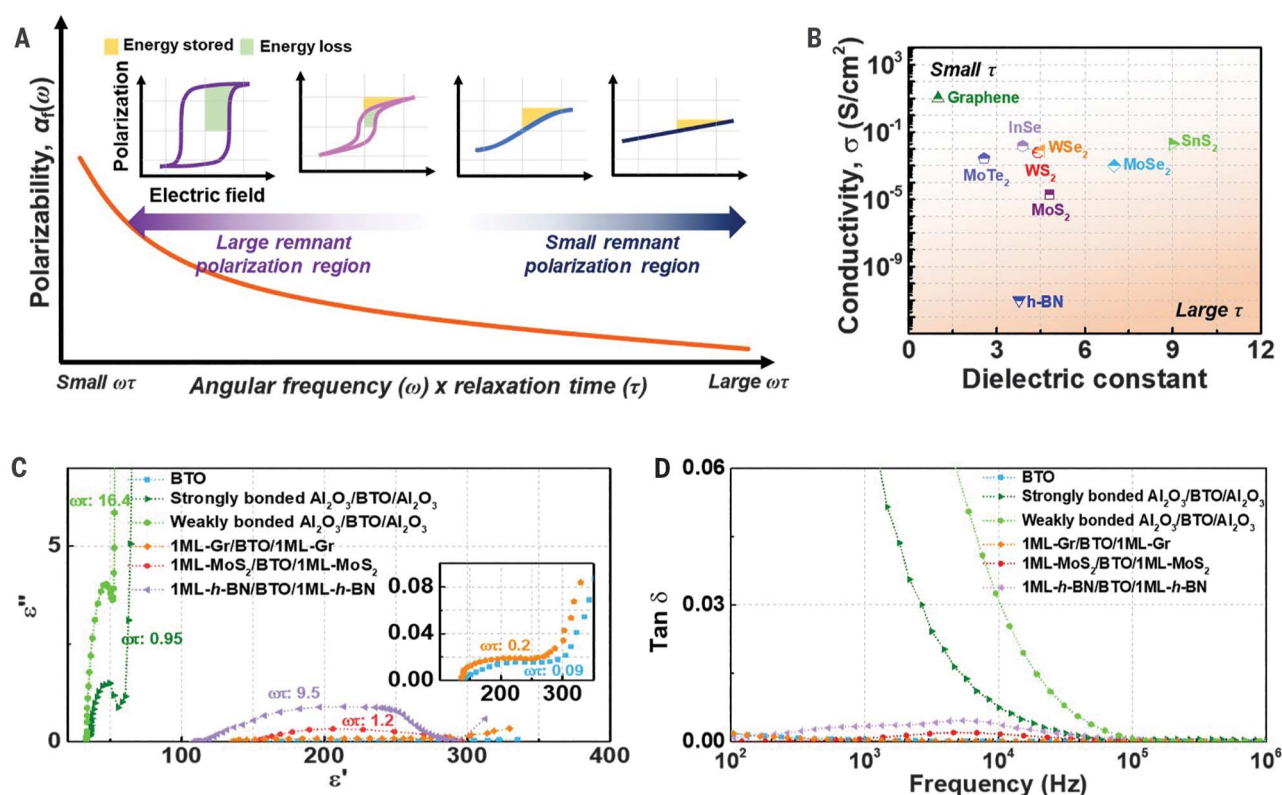
## MW relaxation at 2D/3D interfaces

The Miller model inspired by classic Debye relaxation provides a theoretical framework for controlling spontaneous polarization by manipulating the relaxation time (Fig. 1A). One approach for regulating the relaxation time involves the creation of heterostructures comprising two distinct materials, typically a ferroelectric material and a dielectric material, owing to their disparities in electrical conductivity and permittivity (18). These heterostructures enable the accumulation of charge at the interfaces between the phases when subjected to an alternating electric field, a phenomenon known as the MW relaxation effect. This effect offers a means to influence and modulate the relaxation time within the heterostructures. However, effective modulation of the relaxation time has been difficult to achieve while conserving maximum polarization in conventional heterostructures through the MW effect

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**Fig. 1. Management of spontaneous polarization using the Miller model inspired from classic Debye relaxation.** (A) Polarizability versus  $\omega\tau$  curve of the Miller model. (B) Summary of the conductivity versus dielectric constant curve of several 2D materials. (C and D) Dielectric Cole-Cole plots (C) and  $\tan \delta$  versus frequency curves (D) of the C-BTO, 2D/C-3D/2D, and 3D/C-3D/3D structures.  $\epsilon'$ , real part of the dielectric constant;  $\epsilon''$ , imaginary part of the dielectric constant.

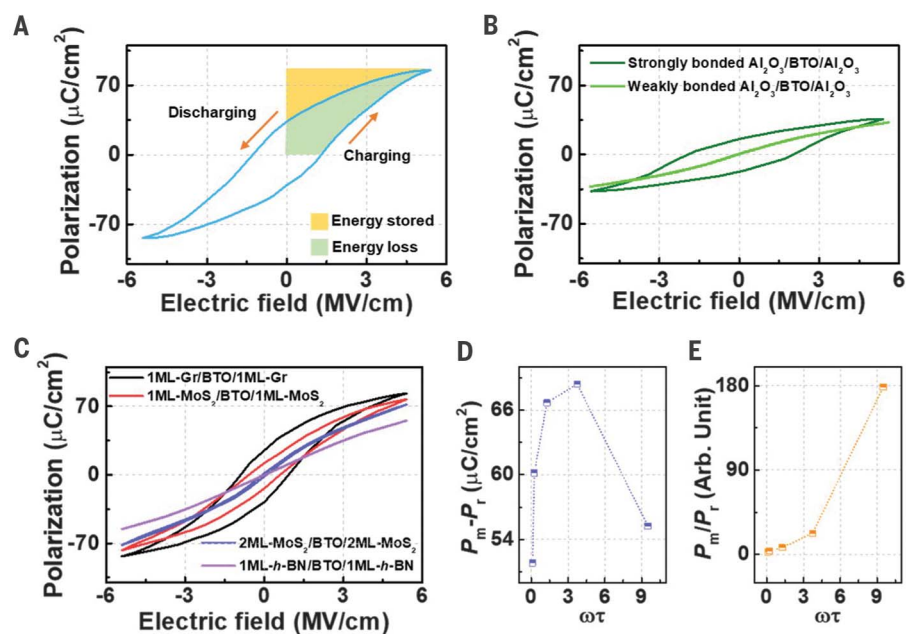
for two reasons: (i) Previous attempts mainly led to the deterioration of ferroelectricity, generating strain or creating multidomain structures when the additional layers form chemical bonds with the ferroelectric material (19, 20). (ii) As the thickness increases because of the additional layers that are needed for the MW effect, this inevitably leads to additional energy loss and a higher  $\tan \delta$ , which is closely associated with dielectric loss (21–23). We circumvented these problems by using 2D/C-3D/2D heterostructures formed using a layer-transfer technique (24–26). We first produced freestanding C-BTO with a thickness of 30 nm. The film is sufficiently high quality that we observed only the (001) orientation in electron back-scattering diffraction mapping. We used a variety of characterization techniques to show the C-BTO quality (figs. S1 to S4). A schematic representation of the 2D/C-BTO/2D fabrication process is shown in fig. S5. The freestanding nature of these C-BTO nanomembranes then allows us to coat both sides with other materials to produce artificial heterostructures.

We anticipated that choosing a 2D material for the coating layer would induce an efficient MW effect while avoiding substantial energy loss because of both the atomically thin nature of the 2D layers and the lack of strong

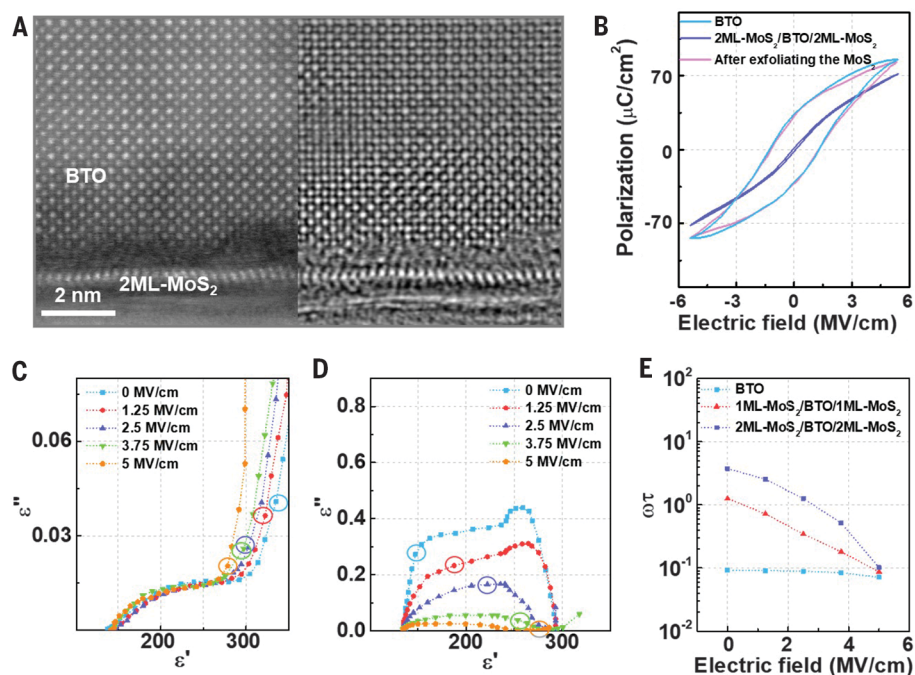
chemical bonding at the heterointerfaces. To aid in 2D material choice, we summarize the conductivity versus dielectric constant of representative 2D materials in Fig. 2B (27–33). The higher the dielectric constant and the lower the conductivity, the stronger the MW relaxation induced at the interface. From the 2D candidates, we chose graphene (a 2D semimetal), MoS<sub>2</sub> (a 2D semiconductor), and hexagonal boron nitride (h-BN) (a 2D insulator). Although some studies of MoS<sub>2</sub> have shown ferroelectricity due to structural deformation, our previous measurements (12) of devices that included similar MoS<sub>2</sub> did not show such behavior, so we do not expect or include any ferroelectric behavior of the MoS<sub>2</sub> layer in our analysis. We selected these materials because we anticipated the possibility of observing diverse variations in relaxation time and  $\tan \delta$  within different artificial heterostructures. For comparison purposes, we also fabricated two different heterostructures of 3-nm-thick Al<sub>2</sub>O<sub>3</sub>/C-BTO/Al<sub>2</sub>O<sub>3</sub>. One heterostructure was formed by atomic layer deposition, for which we anticipated strong chemically bonded BTO/Al<sub>2</sub>O<sub>3</sub> interfaces (fig. S6). We formed the other heterostructure by layer transfer without an annealing process and expected that it would lack strong chemically bonded interfaces (fig. S7).

After fabrication of this set of heterostructures, we measured the dielectric Cole-Cole plot at a frequency range of  $10^2$  to  $10^6$  Hz (Fig. 1C) and calculated the relaxation times at 10 kHz (fig. S8) (34, 35). In the bare freestanding C-BTO nanomembranes, we obtained a value of  $\sim 0.09$  for  $\omega\tau$ , which is defined as the product of angular frequency ( $\omega$ ) of 10 kHz and relaxation time ( $\tau$ ). We calculated the  $\omega\tau$  values of the Al<sub>2</sub>O<sub>3</sub>/C-BTO/Al<sub>2</sub>O<sub>3</sub> samples to be 0.95 and 16.4 for the strongly and weakly bonded samples, respectively. We expect that some internal strain in the Al<sub>2</sub>O<sub>3</sub> and C-BTO may be present, owing to the difference in thermal expansion coefficients, and any such effects are included in the relaxation times. This result indicates that the weakly bonded and discontinuous interface provides a higher relaxation time than that of a strongly chemically bonded interface (36, 37). However, we measured the  $\tan \delta$  values of the strongly and weakly bonded Al<sub>2</sub>O<sub>3</sub>/C-BTO/Al<sub>2</sub>O<sub>3</sub> to be  $7.4 \times 10^{-3}$  and  $31.0 \times 10^{-3}$  at 10 kHz, respectively, which are much higher than those of bare C-BTO (Fig. 1D). We expected substantial energy loss and a decrease in the maximum polarization density based on these values (21, 38, 39). By contrast, the heterostructures we fabricated using layer-resolved splitting of





**Fig. 2. Polarization of the C-BTO, 3D/3D/3D, and 2D/3D/2D heterostructures.** (A)  $P$ - $E$  loop of C-BTO. (B) Comparison of  $P$ - $E$  loops of  $\text{Al}_2\text{O}_3/\text{C-BTO}/\text{Al}_2\text{O}_3$  with strongly and weakly bonded interfaces. (C)  $P$ - $E$  loop of the 2D/3D/2D heterostructure. The  $P$ - $E$  loops were measured at 10 kHz. (D and E)  $P_m - P_r$  (D) and  $P_m/P_r$  (E) calculated from the  $P$ - $E$  loops.



**Fig. 3. Atomic-scale polarization distribution and additional electrical performance under bias.** (A) STEM-HAADF (left) and iDPC (right) images of  $2\text{ML-MoS}_2/\text{C-BTO}/2\text{ML-MoS}_2$ . Separation between the  $\text{MoS}_2$  and BTO and interface roughness of the BTO are visible. (B)  $P$ - $E$  loop of C-BTO,  $2\text{ML-MoS}_2/\text{C-BTO}/2\text{ML-MoS}_2$ , and  $2\text{ML-MoS}_2/\text{C-BTO}/2\text{ML-MoS}_2$  after mechanically exfoliating the  $\text{MoS}_2$  layers. (C and D) Dielectric Cole-Cole plots of C-BTO (C) and  $2\text{ML-MoS}_2/\text{C-BTO}/2\text{ML-MoS}_2$  (D) under dc bias electric field from 0 to 5 MV/cm. The circled values mean the  $\epsilon'$  and  $\epsilon''$  measured at 10 kHz. (E) The  $\omega\tau$  versus electric field of C-BTO,  $1\text{ML-MoS}_2/\text{C-BTO}/1\text{ML-MoS}_2$ , and  $2\text{ML-MoS}_2/\text{C-BTO}/2\text{ML-MoS}_2$ .

2D materials, which enabled thickness control of 2D materials at atomic precision (15, 26), yielded lower  $\tan \delta$  and  $\omega\tau$  values. One-monolayer (ML)  $h$ -BN ( $1\text{ML-}h\text{-BN}$ )/C-BTO/ $1\text{ML-}h\text{-BN}$  had a  $\omega\tau$  of 9.5 and a  $\tan \delta$  of  $3.7 \times 10^{-3}$  at 10 kHz. More effective control of relaxation time is clearly achievable using  $h$ -BN compared with conventional dielectric materials such as  $\text{Al}_2\text{O}_3$ . Additionally, the lower  $\tan \delta$  with  $h$ -BN suggests the potential for relatively smaller energy loss using 2D materials. We attribute the benefits of the 2D material to its atomic thickness, compared with the relatively large  $\text{Al}_2\text{O}_3$  layer thickness. To further explore our hypothesis, we repeated the experiment by creating an artificial heterostructure using  $\text{MoS}_2$ , which is known for its lower carrier density and dielectric constant, as well as graphene, which exhibits much lower values. The one-monolayer  $\text{MoS}_2$  ( $1\text{ML-MoS}_2$ )/C-BTO/ $1\text{ML-MoS}_2$  heterostructure had a  $\omega\tau$  value of 1.2 and a  $\tan \delta$  of  $1.61 \times 10^{-3}$ , whereas the one-monolayer graphene ( $1\text{ML-Gr}$ )/C-BTO/ $1\text{ML-Gr}$  heterostructure had a  $\omega\tau$  value of 0.2 and an even lower  $\tan \delta$  of  $7.6 \times 10^{-5}$ . Unlike the  $1\text{ML-}h\text{-BN}/\text{C-BTO}/1\text{ML-}h\text{-BN}$  heterostructure, which displayed two distinct peaks, indicating the presence of MW relaxation (40, 41), the  $1\text{ML-MoS}_2/\text{C-BTO}/1\text{ML-MoS}_2$  heterostructure did not exhibit clear peak separation, instead displaying an asymmetric Gaussian distribution. This still proves the presence of MW relaxation. However, the  $1\text{ML-Gr}/\text{C-BTO}/1\text{ML-Gr}$  heterostructure did not show similar behavior in the  $\tan \delta$  measurement despite its extremely low  $\tan \delta$ . These findings highlight the role of the 2D materials in influencing and modulating relaxation time and  $\tan \delta$  within the heterostructures.

### Polarization of artificial heterostructures

We expected that the large  $\omega\tau$  values that are caused by the large relaxation time would induce a strong reduction in the remnant polarization. We measured the polarization-electric field ( $P$ - $E$ ) loop of the unmodified C-BTO and the strongly and weakly chemically bonded  $\text{Al}_2\text{O}_3/\text{C-BTO}/\text{Al}_2\text{O}_3$  heterostructures, all at 10 kHz (Fig. 2, A and B). Thirty-nanometer-thick BTO was chosen because it has a higher  $P_m$ , a lower  $P_r$ , and lower electrical leakage than other thicknesses that we tested (fig. S9). The maximum polarizations of both  $\text{Al}_2\text{O}_3/\text{C-BTO}/\text{Al}_2\text{O}_3$  samples were substantially decreased because of the  $\tan \delta$ . However, the remnant polarization of the strongly bonded sample is much higher than that of the weakly bonded sample. These results agree with our expectation, driven by the  $\tan \delta$  and relaxation time values. Nevertheless, the  $\text{Al}_2\text{O}_3/\text{C-BTO}/\text{Al}_2\text{O}_3$  with weakly bonded interfaces still provides low maximum polarization and thus poor energy storage performance because of the high  $\tan \delta$  attributed to the thick nature of  $\text{Al}_2\text{O}_3$ .

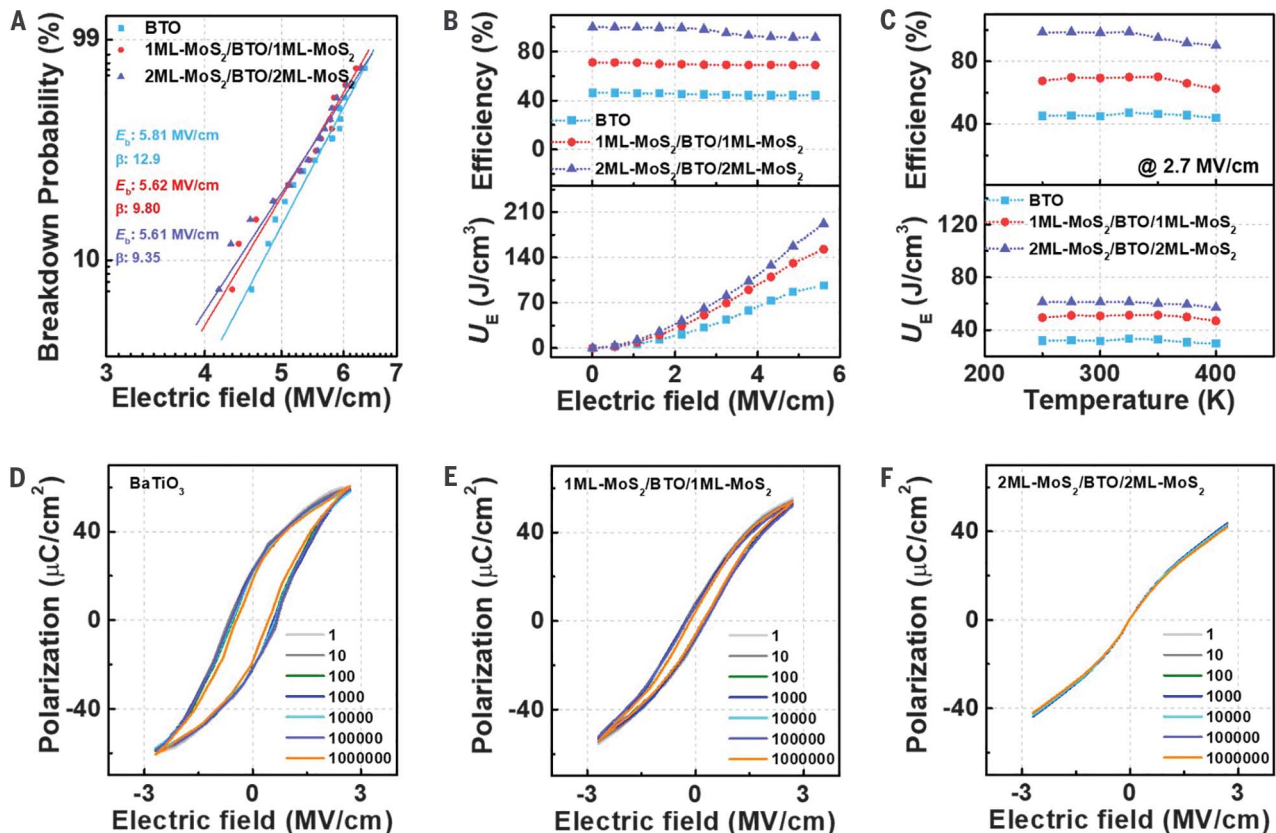
Based on our understanding that relaxation time can be controlled effectively while minimizing  $\tan \delta$ , we conducted  $P$ - $E$  loop measurements on the various 2D/C-BTO/2D heterostructures (Fig. 2C). First, we examined  $h$ -BN/C-BTO/ $h$ -BN structures and observed a reduction in remnant polarization. However, this reduction also led to a decrease in maximum polarization, resulting in a notable decline in energy storage density and efficiency. To mitigate the decrease in maximum polarization, we used  $\text{MoS}_2$ , which possesses a higher conductivity than  $h$ -BN and is thus expected to minimize the MW relaxation. Indeed, when C-BTO was coated with 1ML- $\text{MoS}_2$ , we observed a smaller decrease in maximum polarization. However, we also noted an insufficient decrease in remnant polarization, leading to energy loss. This insufficient decrease is attributed to the increase in conductivity of 1ML- $\text{MoS}_2$ , for which we confirmed, through density functional theory calculations, the Fermi level shifts due to the metal contacts (fig. S10). Meanwhile, the successful suppression of the remnant polarization was observed in heterostructures using bilayer (2ML)- $\text{MoS}_2$ , formed by sequentially transferring 1ML- $\text{MoS}_2$  twice,

and C-BTO. We propose that the free electrons of the electrodes do not affect the  $\text{MoS}_2$  layer adjacent to the C-BTO, which provides a sufficiently low conductivity to allow for screening of the dielectric polarization (fig. S11). We further investigated heterostructures containing graphene, which has a smaller dielectric constant. However, because of its high charge density, we were unable to induce an effective relaxation time delay. Instead, the  $P$ - $E$  loop slightly decreased because of the small increase in the dielectric loss at the non-chemically bonded interface of the graphene/C-BTO (#2). We attribute such performance, with small remnant polarization and high maximum polarization, to several factors: (i) The artificial heterostructures do not sacrifice the crystallinity of single-crystalline ferroelectric materials, unlike conventional heterostructures that experience lattice and thermal mismatch issues. (ii) The atomically thin 2D layers provide extremely low  $\tan \delta$ , preventing a high dielectric loss and decrease in the dielectric constant. (iii) The weakly bonded interface facilitates a substantial increase in dielectric relaxation even when the layer is atomically thin, effectively decreasing the remnant polarization.

Together, we expect the small remnant polarization and large maximum polarization to lead to high energy density and high efficiency. To analyze which of the artificial heterostructures are most suitable for high energy density (Fig. 2, D and E), we subtracted the maximum polarization from the remnant polarization ( $P_m - P_r$ ), which is related to the energy density, and divided the maximum polarization by the remnant polarization ( $P_m/P_r$ ), which is related to the efficiency, according to the  $\omega\tau$ . Up to a  $\omega\tau$  value of 3.4, the  $P_m - P_r$  and  $P_m/P_r$  values increased, indicating improved energy density and efficiency. For  $\omega\tau > 3.4$ , even though  $P_m/P_r$  increased,  $P_m - P_r$  rapidly decreased. Considering these results, we therefore anticipated that the 2D/C-BTO/2D heterostructures with  $\omega\tau$  around 3.4 (1ML- $\text{MoS}_2$ /C-BTO/1ML- $\text{MoS}_2$  and 2ML- $\text{MoS}_2$ /C-BTO/2ML- $\text{MoS}_2$ ) provide the largest energy storage system in our samples (Cole-Cole plot of 2ML- $\text{MoS}_2$ /C-BTO/2ML- $\text{MoS}_2$ ; fig. S12)

#### Atomic-scale polarization distribution

To gain a more comprehensive understanding of polarization behavior in the artificially designed 2D/C-3D/2D structures, we conducted



**Fig. 4. Energy storage performance of C-BTO and  $\text{MoS}_2$ /C-BTO/ $\text{MoS}_2$  with respect to the number of  $\text{MoS}_2$  layers. (A and B)** Two-parameter Weibull distribution analysis of the characteristic breakdown fields (A) and voltage-dependent energy density and efficiency of the C-BTO and  $\text{MoS}_2$ /C-BTO/ $\text{MoS}_2$  heterostructures (B). **(C)** Temperature dependence of energy densities and efficiencies at temperatures ranging from 250 to 400 K. **(D to F)**  $P$ - $E$  loops of C-BTO (D), 1ML- $\text{MoS}_2$ /C-BTO/1ML- $\text{MoS}_2$  (E), and 2ML- $\text{MoS}_2$ /C-BTO/2ML- $\text{MoS}_2$  (F) at temperatures ranging from 250 to 400 K.



additional electrical measurements and obtained atomic-scale structural information for the most effective structure, MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub>. Figure 3A shows high-angle annular dark field (HAADF) and integrated differential phase contrast (iDPC) images of this sample measured by scanning transmission electron microscopy (STEM). We characterized the sample after hysteresis tests for 10 cycles under an electric field of  $-5$  to  $5$  MV/cm at  $10$  kHz. We measured the lattice spacings of the C-BTO in all samples to be about  $0.409$  nm, corresponding to the (001) lattice plane of the BTO perovskite structure and indicating preferred growth along the [001] direction. Our HAADF and iDPC images clearly show the gap between the lattice structures of the 2ML-MoS<sub>2</sub> and C-BTO as well as some interfacial roughness (further example images are shown in fig. S13). Thus, structural discontinuity at the top and bottom surfaces of the C-BTO can work as a screener for dielectric polarization in the C-BTO crystal (43, 44), leading to a relatively small remnant polarization. To further confirm the role of the 2D layer, we measured the  $P$ - $E$  curve after mechanically exfoliating both the top and bottom MoS<sub>2</sub> layers from the 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> heterostructure. The hysteresis after removing the MoS<sub>2</sub> layers is similar to that of bare C-BTO (see the comparison in Fig. 3B), indicating that the pseudo-relaxor ferroelectric behavior exhibited by the 2D/3D/2D heterostructure is driven by charge compensation rather than relaxor ferroelectricity and that the 2D layers did not directly affect the dipoles inside the C-BTO crystal, preserving the high polarization.

Another intriguing aspect of our artificial 2D/3D/2D heterostructure is the observation that as the dc bias electric field increases, the polarization correspondingly increases, with the maximum polarization approaching that of C-BTO. This phenomenon indicates a strong correlation between the applied electric field and the relaxation time within the artificial heterostructure. We show the dielectric Cole-Cole plots of the C-BTO and 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> samples at a dc bias electric field of  $0$  to  $5$  MV/cm (Fig. 3, C and D), where the frequency-dependent dielectric constants are described in fig. S14. In the C-BTO sample, the  $\omega\tau$  measured at  $10$  kHz is observed at the right side of the center of the semicircle regardless of the electric field. By contrast, the  $\omega\tau$  of 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> measured at  $10$  kHz was shifted to the right with increased field, indicating an increase in the relaxation time. We summarize the  $\omega\tau$  calculated from the dielectric Cole-Cole plot as a function of electric field for the C-BTO, 1ML-MoS<sub>2</sub>/C-BTO/1ML-MoS<sub>2</sub>, and 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> samples in Fig. 3E. The dc bias electric field-dependent Cole-Cole plot of MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub> is described in fig. S15. With increasing field, the relaxa-

tion times of the 2D/C-3D/2D become similar to those of the bare C-BTO. We attribute this result to the decrease in the accumulated charge by MW relaxation due to the electric field. This interplay between the electric field and relaxation time provides useful insights into the ferroelectric properties of the artificial 2D/3D/2D heterostructures.

### Performance of MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub>

To investigate the complete energy storage performance of the C-BTO and MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub> heterostructures, we measured their statistical breakdown strengths ( $E_b$ ) by Weibull distribution fitting (Fig. 4A and fig. S16). We calculated the  $E_b$  values of the MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub> structures to be  $5.62$  and  $5.61$  MV/cm, which are comparable with that of C-BTO ( $5.81$  MV/cm) owing to the low thickness of the C-BTO layer. Furthermore, our result is consistent with the negligible effective permittivity change arising from the atomically thin 2D layers ( $I$ ). We observed nonzero leakage currents in the C-BTO and MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub> heterostructures (fig. S17), which are attributed to the narrow bandgap nature of BTO and the probability of oxygen vacancies (45, 46). Nevertheless, the leakage current values are comparable to those of previously reported capacitors with high energy capability (3, 10, 11), indicating high reliability and  $E_b$  under high electric field conditions. From the  $P$ - $E$  loops at  $10$  kHz, we further calculated the  $U_e$  and efficiencies of the samples at an electric field of  $\sim 5.6$  MV/cm (close to their  $E_b$ ) (Fig. 4B). As expected, we obtained a higher energy density from the 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> samples ( $191.7$  J/cm<sup>3</sup>) than from the C-BTO ( $96.9$  J/cm<sup>3</sup>) and 1ML-MoS<sub>2</sub>/C-BTO/1ML-MoS<sub>2</sub> ( $152.4$  J/cm<sup>3</sup>) samples. The energy density we observed in the 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> heterostructure exhibits a high value (fig. S18). These ultrahigh  $U_e$  values are the combined results of high polarization density, low hysteresis, and high efficiency (greater than 90%). The high efficiency is critical to address the energy dissipation of dielectrics for high-power applications, facilitating reliable operation. At higher frequency, smaller  $P_m$  and  $P_r$  were observed because there is insufficient time to align the dipoles at the higher frequency (fig. S19). Nevertheless, high energy storage performance is still achieved. We also conducted stability and reliability tests, which are crucial for electrostatic energy storage. For a thermal stability test, we measured the  $P$ - $E$  loops of the MoS<sub>2</sub>/C-BTO/MoS<sub>2</sub> structures, which showed good stability of the polarization (fig. S20) and energy storage performance (Fig. 4C) under a wide temperature range ( $250$  to  $400$  K) with a small degradation in  $U_e$  ( $<6\%$ ) and efficiency ( $<6\%$ ). We attribute this  $U_e$  and efficiency reduction to the complex thermal scattering of charges accumulated at the non-chemically bonded

interfaces (47). Nevertheless, the degradation rates are similar to previously reported results, indicating good thermal stability (9, 48, 49). They survived over  $10^8$  cycles during an accelerated charge-and-discharge test, as shown in Fig. 4, D to F. Moreover, the degradation rate of 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> is  $<5\%$  after  $10^8$  cycles, indicating that our samples are stable during the cycling test. To assess the practical feasibility of our approach, we fabricated a 2ML-MoS<sub>2</sub>/C-BTO/2ML-MoS<sub>2</sub> array (fig. S21). The average  $P_m$  is  $36.3 \pm 9.3$   $\mu\text{C}/\text{cm}^2$  at  $2.7$  MV/cm with suppressed  $P_r$ , indicating the potential for practical application.

### Conclusions

Our findings highlight the complex interplay between different materials in artificially designed 2D/C-3D/2D heterostructures and their impact on remnant polarization, maximum polarization, energy loss, and efficiency. These insights contribute to the understanding of how to optimize the design and performance of high-energy electrostatic capacitors using ferroelectric materials. By using this approach, we anticipate that future advances can be achieved in the development of more efficient and giant energy storage systems.

### REFERENCES AND NOTES

1. J.-K. Huang et al., *Nature* **605**, 262–267 (2022).
2. A. J. Yang et al., *Nat. Electron.* **5**, 233–240 (2022).
3. H. Pan et al., *Science* **374**, 100–104 (2021).
4. Q. Li et al., *Nature* **523**, 576–579 (2015).
5. H. Luo et al., *Adv. Sci.* **9**, e2202438 (2022).
6. X.-K. Wei et al., *Adv. Mater.* **32**, e2003479 (2020).
7. J. P. B. Silva, K. C. Sekhar, H. Pan, J. L. MacManus-Driscoll, M. Pereira, *ACS Energy Lett.* **6**, 2208–2217 (2021).
8. L. Yang et al., *Prog. Mater. Sci.* **102**, 72–108 (2019).
9. T. Li et al., *Adv. Funct. Mater.* **32**, 2202307 (2022).
10. B. Yang et al., *Nat. Mater.* **21**, 1074–1080 (2022).
11. J. Kim et al., *Science* **369**, 81–84 (2020).
12. J. Shim et al., *Science* **362**, 665–670 (2018).
13. Z. Wu et al., *Adv. Mater.* **34**, e2107538 (2022).
14. M. Qin, L. Zhang, H. Wu, *Adv. Sci.* **9**, e2105553 (2022).
15. S.-H. Bae et al., *Proc. Natl. Acad. Sci. U.S.A.* **114**, 4082–4086 (2017).
16. M. T. Sebastian, R. Ueb, H. Jantunen, *Int. Mater. Rev.* **60**, 392–412 (2015).
17. T. Rahman, M. Vargas, R. Chintalapalle, *J. Alloys Compd.* **617**, 547–562 (2014).
18. S. S. Cheema et al., *Nature* **604**, 65–71 (2022).
19. H. W. Park et al., *Adv. Funct. Mater.* **33**, 2206637 (2023).
20. Y. J. Kim et al., *Sci. Rep.* **6**, 19039 (2016).
21. A. K. Saha, M. Si, P. D. Ye, S. K. Gupta, *Appl. Phys. Lett.* **119**, 122903 (2021).
22. J. P. B. Silva et al., *J. Mater. Chem. A Mater. Energy Sustain.* **8**, 14171–14177 (2020).
23. Z. Wu et al., *ACS Appl. Mater. Interfaces* **10**, 11108–11115 (2018).
24. J. Shin et al., *Nature* **614**, 81–87 (2023).
25. H. Kim et al., *Nat. Nanotechnol.* **18**, 464–470 (2023).
26. S.-H. Bae et al., *Nat. Mater.* **18**, 550–560 (2019).
27. A. Laturia, W. G. Vandenberghe, in *2017 International Conference on Simulation of Semiconductor Processes and Devices (SI-SPAD)* (IEEE, 2017), pp. 337–340.
28. H. Zhou, Y. Cai, G. Zhang, Y.-W. Zhang, *Nanoscale* **10**, 480–487 (2017).
29. J.-H. Ahn et al., *Nano Lett.* **15**, 3703–3708 (2015).
30. S.-I. Kim et al., *Synth. Met.* **299**, 117464 (2023).
31. A. Perepeliciu et al., *Appl. Phys. Lett.* **122**, 263503 (2023).
32. S. Hwangbo, L. Hu, A. T. Hoang, J. Y. Choi, J.-H. Ahn, *Nat. Nanotechnol.* **17**, 500–506 (2022).
33. Y. Ahn, M. Shin, *IEEE Trans. Electron Dev.* **64**, 2129–2134 (2017).

34. T. A. Hanafy, K. Elbanna, S. El-Sayed, A. Hassen, *J. Appl. Polym. Sci.* **121**, 3306–3313 (2011).
35. H. Wang, L. Yang, X. Zhang, M. H. Ang Jr., *Results Phys.* **29**, 104781 (2021).
36. B.-H. Fan *et al.*, *Compos. Sci. Technol.* **80**, 66–72 (2013).
37. K. Venkateshan, G. P. Johari, *J. Chem. Phys.* **125**, 014907 (2006).
38. M. H. Park, H. J. Kim, Y. J. Kim, T. Moon, C. S. Hwang, *Appl. Phys. Lett.* **104**, 072901 (2014).
39. M. Liu *et al.*, *ACS Appl. Mater. Interfaces* **4**, 5761–5765 (2012).
40. M. Samet *et al.*, *J. Chem. Phys.* **142**, 194703 (2015).
41. H. Hammami, M. Arous, M. Lagache, A. Kallel, *J. Alloys Compd.* **430**, 1–8 (2007).
42. Y. G. Lee *et al.*, *Appl. Phys. Lett.* **98**, 183508 (2011).
43. T. Akamatsu *et al.*, *Science* **372**, 68–72 (2021).
44. Y. Liu, Y. Huang, X. Duan, *Nature* **567**, 323–333 (2019).
45. S. H. Kim *et al.*, *Phys. Status Solidi Rapid Res. Lett.* **13**, 1900373 (2019).
46. M. Pawlak *et al.*, *Appl. Phys. Lett.* **101**, 042901 (2012).
47. J. Yang *et al.*, *Nat. Nanotechnol.* **7**, 91–95 (2012).
48. H. Qi *et al.*, *Adv. Funct. Mater.* **29**, 1903877 (2019).
49. D. Li *et al.*, *Small* **19**, e2206958 (2023).

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characterization under the supervision of F.M.R. S.Se. and J.-H.P. analyzed the 2D/3D and 3D/3D interfaces. S.B., C.K., J.Z., C.W., J.Kw., and D.-H.K. performed the electrical measurement. J.H., H.C., and H.-S.K. conducted the breakdown performance. S.-H.B. and S.H. wrote the first draft of the manuscript. All authors discussed the results and revised the manuscript. **Competing interests:** S.H. and S.-H.B. are inventors on patent application no. 63/617,314 assigned to Washington University that covers heterostructures that have a van der Waals interface for high-energy density capacitors. **Data and materials availability:** All data are available in the main text or the supplementary materials. **License information:** Copyright © 2024 the authors, some rights reserved; exclusive licensee American Association for the Advancement of Science. No claim to original US government works. <https://www.science.org/about/science-licenses-journal-article-reuse>

## SUPPLEMENTARY MATERIALS

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Materials and Methods

Supplementary Text

Figs. S1 to S21

References (50–76)

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