




Cite this: *RSC Adv.*, 2018, 8, 2837

Polymer/oxide bilayer dielectric for hysteresis-minimized 1 V operating 2D TMD transistors†

Minho Yoon,  ‡ Kyeong Rok Ko, ‡ Sung-Wook Min and Seongil Im*

Despite their huge impact on future electronics, two-dimensional (2D) dichalcogenide semiconductor (TMD) based transistors suffer from the hysteretic characteristics induced by the defect traps located at the dielectric/TMD channel interface. Here, we introduce a hydroxyl-group free organic dielectric divinyl-tetramethyldisiloxane-bis (benzocyclobutene) (BCB) between the channel and conventional SiO₂ dielectric, to practically resolve such issues. Our results demonstrate that the electrical hysteresis in the n-channel MoS₂ and p-channel MoTe₂ transistors were significantly reduced to less than ~20% of initial value after being treated with hydrophobic BCB dielectric while their mobilities increased by factor of two. Such improvements are certainly attributed to the use of the hydroxyl-group free organic dielectric, since high density interface traps are related to hydroxyl-groups located on hydrophilic SiO₂. This concept of interface trap reduction is extended to stable low voltage operation in 2D MoTe₂ FET with 30 nm BCB/10 nm Al₂O₃ bilayer dielectric, which operates well at 1 V. We conclude that the interface engineering employing the BCB dielectric offers practical benefits for the high performance and stable operation of TMD-based transistors brightening the future of 2D TMD electronics.

Received 21st November 2017
Accepted 8th January 2018

DOI: 10.1039/c7ra12641g

rsc.li/rsc-advances

Introduction

Two-dimensional (2D) dichalcogenide semiconductors (TMD) have been extensively studied during recent years due to their massive potential for next-generation electronics.^{1,2} Compared to the gapless semiconductor graphene,³ TMD such as molybdenum disulfide (MoS₂),⁴ molybdenum ditelluride (MoTe₂),⁵ and tungsten diselenide (WSe₂)⁶ provide the tunable electronic bandgap which is dependent on the thickness of the layer itself. Due to their discrete bandgap, the TMD-based field effect transistors (FETs) often exhibit clear switching operation with its relatively high values of mobility and ON/OFF current ratio.^{7–10} However, several reports reveal that the electrical performances of the devices are significantly degraded by the interface and surface states of the devices.^{11,12} For a good candidate to minimize the interfacial defect states in FET, the hydrophobic hexagonal boron nitride (h-BN) as dielectric and passivation layer has been proposed.^{13,14} Although this approach has been encouraging, h-BN is basically expensive due to its special growth processes for good crystalline quality control,^{15,16} and furthermore, additional elaborate processes are still requested for h-BN flake to be incorporated in device fabrications. Hence, alternatives to replace h-BN have been suggested: self-assembled monolayers,¹⁷ organic

insulating materials,¹⁸ high-k oxide dielectrics,¹⁹ and *etc.* Among these candidates, an organic insulating polymer, divinyl-tetramethyldisiloxane-bis(benzocyclobutene) (BCB) could be an appropriate option for TMD-channel FETs. BCB is a well-known non-polar organic polymer with hydroxyl-group free chemical structure and it has been widely used as a dielectric for stable high-performance organic transistors.^{20–22} To the best of our limited knowledge, BCB has never been attempted for the TMD-based field effect transistors while it is easily formed by spin-casting.

In the present study, we report n-channel MoS₂ and p-channel MoTe₂ transistors with the BCB gate dielectric on oxide gate dielectric. According to our results, initial hysteresis in the MoS₂ and MoTe₂ transistors on 285 nm-thick SiO₂/p⁺-Si back gate without BCB were ~15 V in their transfer curve characteristics, but it reduced to ~4 V in the devices with BCB. Based on these hysteresis minimization effects by BCB, we successfully extended our results to a more practical device application, 1 V operation of MoTe₂ FET with 30 nm-thin BCB on 10 nm-thin atomic layer deposited (ALD) Al₂O₃. It is thus regarded that BCB dielectric layer offers benefits for the high performance and stable operation of TMD-based transistors. The main advantage of the BCB would be its non-polar hydrophobicity and process conveniences.

Experimental detail

For basic investigations on 2D TMD device stability, 300 nm-thick organic insulating material, divinyl-tetramethyldisiloxane-bis

Institute of Physics and Applied Physics, Yonsei University, 50 Yonsei-ro, Seodaemun-gu, Seoul 120-749, Korea. E-mail: semicon@yonsei.ac.kr; Fax: +82-2-392-1592; Tel: +82-2-2123-2842

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra12641g

‡ M. Y. and K. K. contributed equally to this work.



(benzocyclobutene) (BCB, CYCLOTENE, Dow Chemical) was formed on the 285 nm-thick $\text{SiO}_2/\text{p}^+\text{-Si}$ substrate by spin-casting and subsequent thermal annealing at 300 °C for 10 minutes in nitrogen ambient. (But for extended practical device fabrication, 30 nm-thin BCB on 10 nm-thin atomic layer deposited Al_2O_3 was formed on patterned Au gate). Then, MoS_2 and MoTe_2 flakes for n and p-type transistors were micromechanically exfoliated with polydimethylsiloxane (PDMS) stamps, respectively. Those TMD flakes are subsequently dry-transferred onto the two types of dielectric substrates: BCB/ SiO_2 and SiO_2 . Then, 50 nm-thick Au for MoS_2 and 100 nm-thick Pt for MoTe_2 were sputter-deposited on the substrates for source/drain electrodes and patterned by conventional lift-off processes. The thickness of the flakes were confirmed with the atomic force microscopy (AFM) and also identified with Raman spectroscopy.

The current–voltage (I – V) measurements of the transistors were performed with a semiconductor parameter analyser (Model HP4155C, Agilent Technologies) and the capacitance–voltage (C – V) measurements of the devices were conducted with a LCR meter (Model HP4284A, Agilent Technologies). Electrical characterisations of the devices were carried out entirely in air ambient (relative humidity RH \sim 45%) at room temperature, but for one MoTe_2 device sample was also measured in vacuum (less than 1 Torr) at 300 K, to investigate the hysteresis-induction effects of air molecules adsorbed on TMD channel surface.

Results and discussion

Fig. 1a and d show bottom-gate top-contact MoS_2 and MoTe_2 transistors on 285 nm thick SiO_2 dielectric, respectively. Their thickness appears to be 4 nm for MoS_2 and 3 nm for MoTe_2 as measured by atomic force microscopy (AFM) scan in Fig. 1b and e. Identity of the two flakes was again confirmed by Raman spectroscopy whose results are seen in ESI section (Fig. S1†). Fig. 1c and f show the transfer characteristics (I_{DS} vs. V_{GS}) of our bottom-gate top-contact MoS_2 and MoTe_2 transistors, respectively. Their insets are schematic cross sections of those devices. The MoS_2 transistor with Au source/drain electrodes displays n-type conduction behaviour with the electron mobility of $11.5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and high on/off current ratio ($>10^5$). The MoTe_2 transistor with Pt source/drain electrodes presents p-type conduction with the hole mobility of $9.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and high on/off current ratio ($>10^5$) as well. Although the electrical performances of the devices are promising as reported previously by others,^{23,24} it is very apparent that the devices much suffer from the gate bias-induced hysteresis ($\Delta V \approx \sim 10.2 \text{ V}$ for n- MoS_2 and $\sim 14.3 \text{ V}$ for p- MoTe_2 FET) in the present transfer characteristics, which should be minimized for practical device applications in electronics.

Numerous reasons count for the hysteresis in TMD-based 2D transistors as discussed enormously in the organic and inorganic transistors for decades,^{25–27} and the reasons were classified into four main categories: the adsorption of water and oxygen molecules on semiconductor channel surface, defects in the semiconductor material, mobile trap charges in the dielectric, and the interface trap charges between the

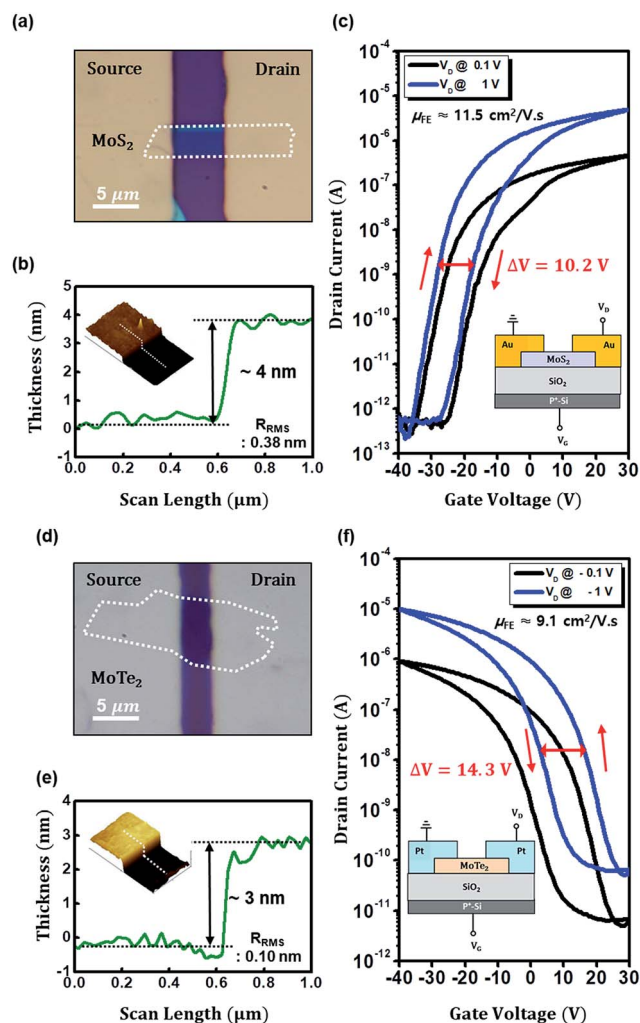


Fig. 1 (a–c) Optical microscopy (OM) image, AFM image along with channel thickness profiles ($\sim 4 \text{ nm}$), and transfer characteristics (I_{DS} vs. V_{GS}) of n-type MoS_2 FET on 285 nm-thick SiO_2 dielectric with cross section scheme (inset). The channel width and length of the MoS_2 FET are 4.1 and 5.4 μm , respectively, and the device shows the electron mobility of $11.5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. The value of the clockwise hysteresis is around 10.2 V. (d–f) OM image, AFM image along with thickness profiles ($\sim 3 \text{ nm}$), and the transfer characteristics of p-type MoTe_2 FET on 285 nm-thick SiO_2 dielectric with cross section scheme (inset). The channel width and length of the MoTe_2 FET are 8.1 and 3.4 μm , respectively, and the device shows the hole mobility of $9.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. The value of the anticlockwise hysteresis is around 14.3 V.

semiconductor and the contacting dielectric. Here, we mainly focused on the interface effects because the single-crystalline-like 2D semiconducting channel would meet hydrophilic oxide dielectric in general and their interface becomes to contain high density charge traps which originate from the surface hydroxyl-group of the oxides dielectric.^{12,28}

To this end, a hydrophobic organic insulator, BCB was conceived as an excellent option for the dielectric for the transistors because the BCB film has a hydroxyl-group free chemical structure and can be deposited by simple spin-casting.

Prior to exploiting the BCB as a dielectric layer for MoS_2 and MoTe_2 transistors, the dielectric properties and hydrophobicity

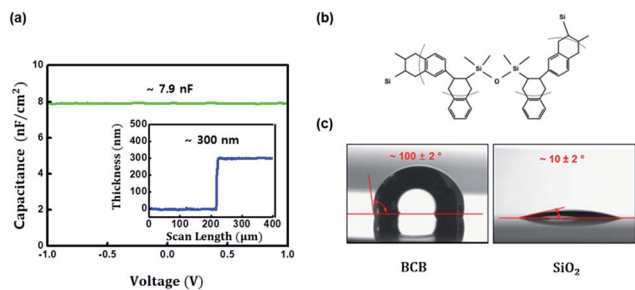


Fig. 2 (a) The geometrical capacitance of the BCB dielectric in Au/BCB/Au structure with the thickness of BCB dielectric (inset). (b) Chemical structure of the BCB. (c) The contact angle of D.I. water on the BCB and SiO₂, respectively. The contact angle of D.I. water on the hydrophobic BCB dielectric was $\sim 100^\circ$ while that on a cleaned SiO₂ hydrophilic substrate was $\sim 10^\circ$.

of the BCB layer were initially investigated. The capacitance of our BCB dielectric was measured to be $\sim 7.9 \text{ nF cm}^{-2}$ at 1 kHz with top and bottom Au electrodes (Au/BCB/Au) in the Fig. 2a, where the thickness of the spin-casted BCB layer was $\sim 300 \text{ nm}$ (inset of the Fig. 2a) as scanned with a surface profiler. Deduced dielectric constant of the BCB dielectric at the thickness was ~ 2.67 , which is in a good agreement with the previously reported value.²⁹ Furthermore, as shown in the Fig. 2c, the contact angle of D.I. water on the BCB dielectric was $\sim 100^\circ$ while that on a cleaned SiO₂ substrate was as low as $\sim 10^\circ$; such contact angle measurements clearly indicate the hydrophobicity of the BCB layer surface due to its chemical structure (Fig. 2b).³⁰

Fig. 3a and d show bottom-gate top contact MoS₂ and MoTe₂ transistors with BCB (300 nm)/SiO₂ dielectric, respectively. Their thickness appears to be 6 nm for MoS₂ and 8 nm for MoTe₂ as measured by atomic force microscopy (AFM) scan in Fig. 3b and e. Fig. 3c and f show the transfer characteristics ($I_{\text{DS}}-V_{\text{GS}}$) of the bottom-gate top contact MoS₂ and MoTe₂ transistors on the BCB/SiO₂ dielectric, respectively. Their insets are schematic cross sections of those devices. With BCB dielectric, the values of the hysteresis in the MoS₂ and MoTe₂ transistors were dramatically reduced to $\sim 4.5 \text{ V}$ and $\sim 2.3 \text{ V}$. (Similar hysteresis reduction is also shown in the output characteristics of Fig. S2†). Furthermore, the field-effect mobility of the devices was significantly increased by the factor of two in both cases (from 11.5 to 15.8 $\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ in MoS₂ transistors and from 9.1 to 18.2 $\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ in MoTe₂ transistors). We thus regard that the BCB dielectric effectively minimizes the trap sites or passivate the electro-active hydroxyl groups at the dielectric/TMD channel interface improving the electrical performance and the stability of the devices. On the one hand, the linear mobility of our FETs was estimated with the following equation.³¹

$$\mu_{\text{LIN}} = \frac{1}{C_i V_{\text{DS}}} \frac{L}{W} \frac{\partial I_{\text{D}}}{\partial V_{\text{GS}}} \quad (1)$$

where C_i is the geometric dielectric capacitance, L is channel length and W is its width.

Based on aforementioned transfer characteristics from TMD-devices with and without BCB dielectric, we considered

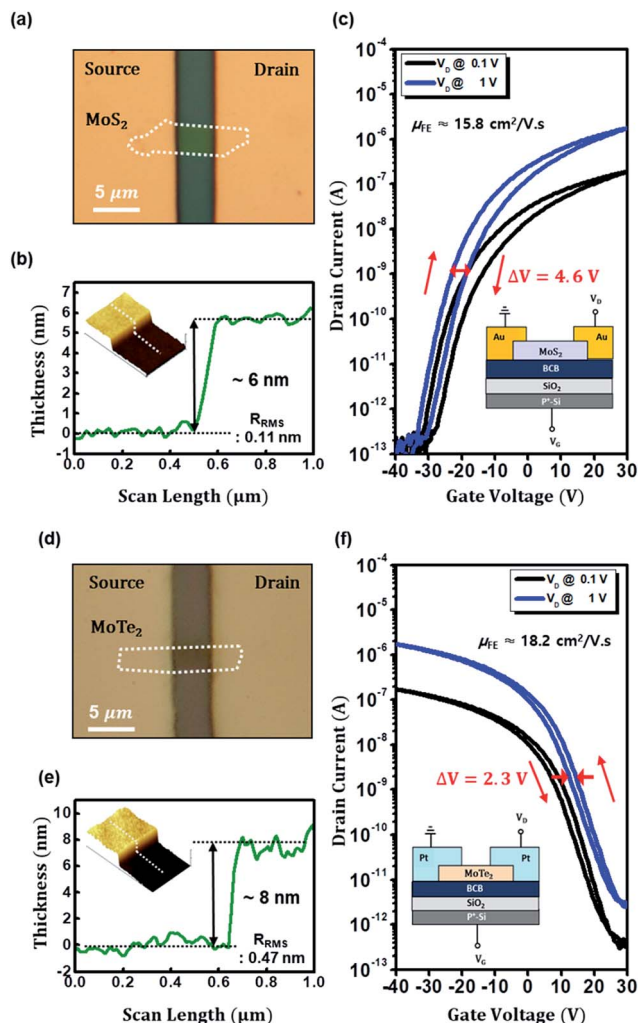


Fig. 3 (a–c) Optical microscopy image, AFM image along with channel thickness profile ($\sim 6 \text{ nm}$), and the transfer characteristics (I_{DS} vs. V_{GS}) of n-MoS₂ FET with 300 nm-thick BCB/SiO₂ dielectric with cross section scheme (inset). The channel width and length of the MoS₂ FET are 2.5 and 2.9 μm , respectively, and the device shows the electron mobility of 15.8 $\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. The value of the clockwise hysteresis is around 4.6 V. (d–f) OM image, AFM image along with channel thickness profile ($\sim 8 \text{ nm}$), and the transfer characteristics of p-MoTe₂ FET with BCB/SiO₂ dielectric with cross section scheme (inset). The channel width and length of the MoTe₂ FET are 2.8 and 3.9 μm , respectively, and the device shows the hole mobility of $\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. The value of the anticlockwise hysteresis is around 2.3 V.

quantifying the effective interface trap density, D_{it} by deducing the values from the subthreshold swing (SS) of the transistors, which is expressed in the following equation.³¹

$$\text{SS} = \left(\frac{d(\log_{10} I_{\text{DS}})}{dV_{\text{GS}}} \right)^{-1} \approx \ln 10 \frac{kT}{q} \left(1 + \frac{q^2 D_{\text{it}}}{C_i} \right) \quad (2)$$

where k is the Boltzmann constant, T is the temperature in Kelvin, q is the electronic charge, and C_i is the geometric capacitance of the dielectric.

We thought this type of approach is at least worthy even though this SS equation ignores any effects caused by contact resistance which is sometimes not ignorable at all.³² The

subthreshold swing (SS) values of the MoS₂ transistors with BCB/SiO₂ and SiO₂ dielectrics seem similar each other as about 2.2 V dec⁻¹ and 2.0 V dec⁻¹, respectively, but their geometric capacitances are quite different, to be 4.8 nF cm⁻² and 12.1 nF cm⁻². Thus, the estimated values of D_{it} become 1.1×10^{12} cm⁻² eV⁻¹ and 2.4×10^{12} cm⁻² eV⁻¹ for the n-MoS₂ transistors with and without BCB. Similarly, the SS values of p-MoTe₂ transistors with and without BCB dielectric were 3.6 V dec⁻¹ and 3.4 V dec⁻¹, resulting in D_{it} values of 1.9×10^{12} cm⁻² eV⁻¹ and 4.4×10^{12} cm⁻² eV⁻¹, respectively. Likewise, devices with BCB dielectric appear to contain 2–3 times lower number density of traps at the interface in SS-based estimation. However, such SS-based approach and D_{it} results must be still unclear because SS behaviour cannot ignore contact resistance effects from TMD/source-drain electrode contact.³³ In addition, since the gate hysteresis of $I_{DS}-V_{GS}$ transfer characteristics ($I-V$ measurement system) unavoidably contains source/drain contact resistance effects in general.³⁴ Hence, it might be necessary to find other measurement scheme which is immune from contact effects.

We thus conceived to perform capacitance–voltage ($C-V$) measurements for evaluating the trapped interface charges, because $C-V$ measurements are oriented to focus on the interface by gate bias without any interference from the contact resistance effects.³⁵ For the $C-V$ measurements on FET structure at high and low frequencies, DC and small signal AC voltages are applied to the gate electrode while source/drain (S/D) electrodes are grounded in general.³⁶ However, general $C-V$ method

on FET structure would not be effective if the channel area is too small compared to the S/D-to-gate overlap area; real capacitance signals from channel should be overridden by parasitic capacitance. Our device with small 2D channel would definitely meet such parasitic capacitance issue, so we modified the general $C-V$ measurements by grounding the source electrode only, considering that DC voltage sweep would eventually induce the capacitance of channel and drain electrode area by channel accumulation or channel conducting.¹⁵ Since the trapped charges cannot respond at high frequency, the interface trap density can be extracted at a low frequency of 100 Hz, using the sample transistor architecture as shown in Fig. 4a. According to the schematic circuits of Fig. 4a, we meet with two cases during DC sweep: channel depletion and accumulation which are dependent on the DC bias. In the case of channel depletion, a capacitance (C_S) is measured from the dielectric area under only one electrode (source), however such capacitance should be doubled-up by channel accumulation, which would connect the source and drain (the other electrode; now total capacitance becomes $C_S + C_D = 2C_S$). According to the $C-V$ plots of Fig. 4b–e, the capacitance value is indeed doubled-up from 55 to 112 pF (for FETs without BCB but SiO₂) or from 22 to 45 pF (for FETs with BCB). In our device (photo images of Fig. 1a and 4a), the channel area for channel capacitance (C_{ch}) is too small (incomparable to electrode area) to be visibly counted for total capacitance. In spite of that, the TMD channels successfully performed as connecting/disconnecting (on/off) switches.

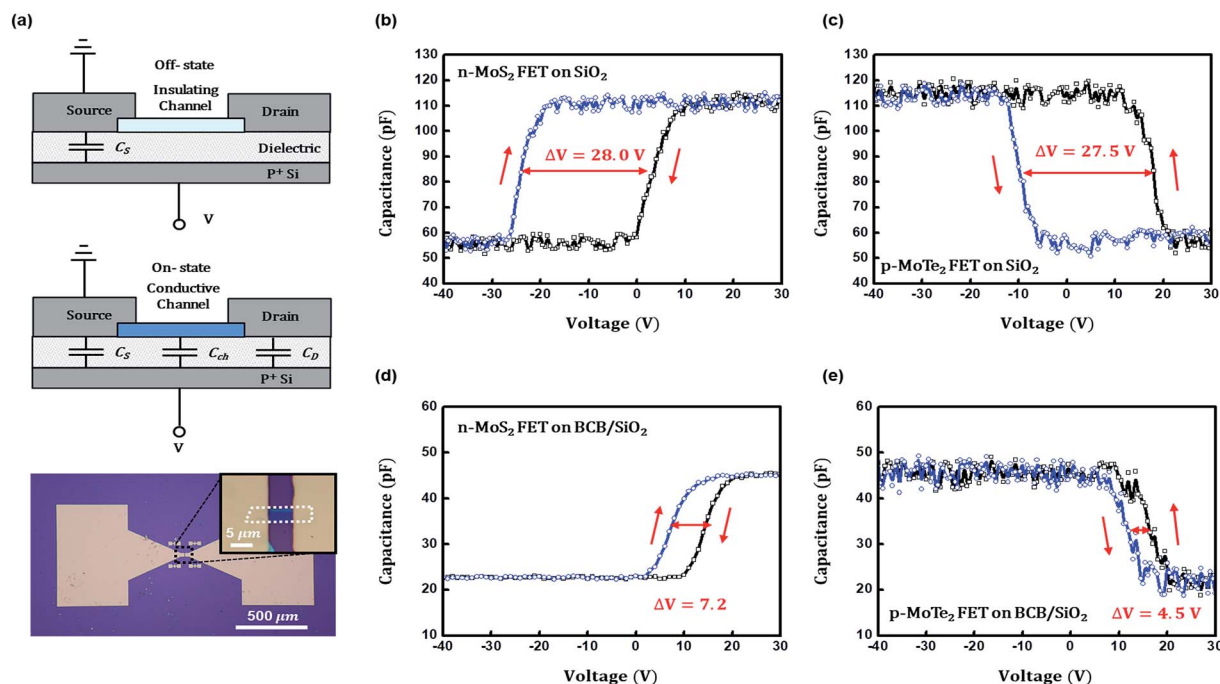


Fig. 4 (a) Schematic circuit and illustration of the modified $C-V$ measurements on our TMD FETs. In channel depletion, a capacitance (C_S) is measured from the dielectric area under only one electrode (source), however such capacitance is doubled-up by channel accumulation (note the plots in (b–e)), which would connect the source and drain [the other electrode for $C_D (=C_S)$]. The channel area (\sim less than $25 \mu\text{m}^2$ as seen in the inset OM) for C_{ch} is incomparably smaller than that of one electrode which is around $4.55 \times 10^5 \mu\text{m}^2$. According to the $C-V$ characteristics obtained from (b) MoS₂ on SiO₂, (c) MoTe₂ on SiO₂, (d) MoS₂ on BCB/SiO₂, and (e) MoTe₂ on BCB/SiO₂, the initial voltage hysteresis is dramatically reduced from ~ 28 V to 4.5–7.2 V due to the BCB effects.

Table 1 Summary of the mobility, subthreshold swing, hysteresis, and estimated trap densities of the devices

Semicon.	Dielec.	Capacit. (nF cm ⁻²)	Mobility (cm ² V ⁻¹ s ⁻¹)	S.S (V dec ⁻¹)	Hysteresis (ΔV) from		Trap density ($\times 10^{12}$ cm ⁻²) from	
					Transfer curves	Modified <i>C-V</i>	S.S	Modified <i>C-V</i>
MoS ₂	SiO ₂	12.1	11.5	2.0 ± 0.1	10.2 ± 0.1	28.0 ± 0.1	2.9 ± 0.1	2.12 ± 0.01
MoS ₂	BCB/SiO ₂	4.8	15.8	2.2 ± 0.1	4.6 ± 0.1	7.2 ± 0.1	1.30 ± 0.03	0.22
MoTe ₂	SiO ₂	12.1	9.1	3.4 ± 0.1	14.3 ± 0.1	27.5 ± 0.1	4.4 ± 0.1	2.08 ± 0.01
MoTe ₂	BCB/SiO ₂	4.8	18.2	3.6 ± 0.1	2.3 ± 0.1	4.5 ± 0.1	1.86 ± 0.03	0.14

Noticeable in Fig. 4b and c is that those TMD channels clearly show hysteresis from forward-to-backward sweep. Because trapping and de-trapping of charges at the interface are certainly related to the forward and backward sweep, respectively, it is anticipated that the interfacial trap density would be quite precisely estimated using the hysteresis voltage. For both n-MoS₂ and p-MoTe₂ FETs with only SiO₂ dielectrics, almost the same amount of large voltage hysteresis was obtained to be 27.5–28 V in Fig. 4b and c, respectively. The hysteresis was significantly reduced to be 7.2 and 4.5 V of for n- and p-FETs as seen in Fig. 4d and e. This result is readily expected from the related transfer curves in Fig. 3c and f. Small amount of hysteresis still remains as 4.5–7.2 V due to the small trap density at the BCB dielectric/TMD channel interface. But such number is only about 15–25% of previous trap density on SiO₂. Based on our *C-V* results, it is very likely that the hysteresis observed from *I*_{DS}-*V*_{GS} transfer characteristics is mainly from gate dielectric/channel interface, not from the TMD surface effects involved with air molecules. For further confirmation on the dielectric/channel interface-induced hysteresis, we subsidiary performed *I*_{DS}-*V*_{GS} transfer characteristics of another MoTe₂ device with

BCB dielectric in vacuum ambient at 300 K. As shown in Fig. S3 of ESI,[†] already-reduced hysteresis by BCB application seems not decreased any further even in vacuum, which supports our assumption that such hysteresis is mainly related to the amount of interfacial traps.

With the results from *C-V* characteristics in Fig. 4a–e, we could easily estimate the interfacial trap densities of MoS₂ and MoTe₂-based FETs with and without BCB, using the following simple equation, $Q_{it} = C_i \Delta V / q$, where ΔV is the voltage hysteresis and C_i is the geometric capacitance (Farad cm⁻²) of the dielectric. C_i value can be obtained from the *C-V* curves in Fig. 4b and d because we already know the electrode area. From the equation, the estimated values of the interface trap charge density were 2.08×10^{12} cm⁻² eV⁻¹ and 2.11×10^{12} cm⁻² eV⁻¹ in the MoS₂ and MoTe₂ transistors on SiO₂ dielectric. But those values were an order of magnitude reduced to the values of 2.2×10^{11} cm⁻² eV⁻¹ and 1.4×10^{11} cm⁻² eV⁻¹ when BCB was inserted as a dielectric layer. Table 1 summarizes all the values on interface trap densities and device performances. Here, we assumed that the trap charges only stem from the dielectric/channel interface, and the trap density values by SS and D_{it}

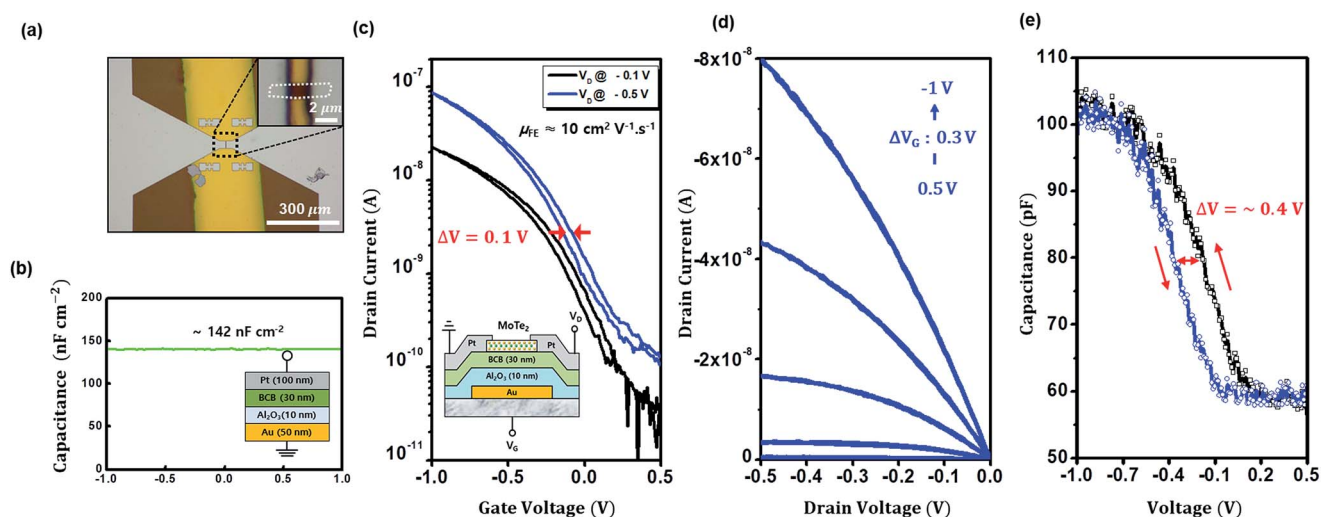


Fig. 5 (a) Optical image of our MoTe₂ FET with BCB/Al₂O₃ dielectric and Au bottom gate, along with inset magnified image of 5 nm-thin MoTe₂ channel (*W/L* ratio $\sim 1 \mu\text{m}/2 \mu\text{m} = 0.5$). (b) The geometrical capacitance of the thin BCB/Al₂O₃ dielectric. (Inset: schematic illustration of device cross section) (c) transfer characteristics of MoTe₂ FET. ON/OFF *I*_D current ratio appears to be ~ 1000 . (Inset: schematic illustration of device cross section. Au gate was patterned on thick SiO₂/p⁺-Si wafer substrate). (d) Output characteristics show a very low operational voltage of 0.5–1 V. (e) *C-V* curve hysteresis voltage was measured, to be ~ 0.4 V which leads to an estimated interface trap density of $\sim 3.5 \times 10^{11}$ cm⁻².

measurement were worked out as overall average trap density ($D_{it} \times \text{energy gap}$).²⁴ According to Table 1, SS-driven method always results in higher values of trap density than the values by C - V method whether the device has BCB dielectric or not, because any I - V method implicitly reflects the contact resistance effects.³⁴ The difference between SS-driven and C - V methods is very clear in the FETs with BCB dielectric, but such difference becomes relatively quite small in the other devices with high density traps (with only SiO_2 dielectric).

Although we have mainly focused on minimizing the interfacial trap density in the present study, we also extended our results to a more practical device application as our final effort: low voltage operational MoTe_2 FET with 30 nm-thin BCB on 10 nm-thin atomic layer deposited (ALD) Al_2O_3 , Fig. 5a and its inset show optical images of our MoTe_2 FET with BCB/ Al_2O_3 dielectric and Au bottom gate, where width-to-length ratio of the device was $1 \mu\text{m}/2 \mu\text{m}$. Capacitance of BCB/ Al_2O_3 was $\sim 142 \text{ nF cm}^{-2}$ as obtained from metal-insulator-metal (MIM) C - V measurement (Fig. 5b). We also confirmed the BCB thickness of 30 nm with a mechanical profiler in Fig. S4a† and the MoTe_2 thickness of 5 nm by AFM scan (Fig. S4b†). From the transfer characteristics, device mobility (linear regime) turned out to be $\sim 10 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, and ON/OFF I_D current ratio appears to be ~ 1000 (Fig. 5c). Transfer and output characteristics (Fig. 5c and d) show a very low operational voltage of 0.5–1 V. The C - V curve hysteresis was measured in Fig. 5e, to be $\sim 0.4 \text{ V}$ which leads to an estimated interface trap density of $\sim 3.5 \times 10^{11} \text{ cm}^{-2}$. Since the trap density is comparable to those from thick BCB/ SiO_2 in Table 1, it is well regarded that 30 nm-thin BCB polymer on 10 nm Al_2O_3 keeps the function of trap minimization ensuring the practical low voltage operation as well. Such hysteresis-reduced low voltage operation in 2D TMD FET has rarely been demonstrated.^{17,37}

Conclusions

We have fabricated a few layer n- MoS_2 and p- MoTe_2 channel FETs with and without BCB dielectric on $\text{SiO}_2/\text{p}^+\text{-Si}$. Our results from C - V measurements and I - V transfer characteristics display that the hysteresis in the MoS_2 and MoTe_2 transistors were significantly reduced to less than $\sim 20\%$ of initial value after treated with hydrophobic BCB dielectric, while the linear mobilities of both p- and n-FETs increased by factor of two. Such improvements are certainly attributed to the hydroxyl-group free organic dielectric, BCB on SiO_2 , since high density interface traps are related to hydroxyl-groups located on SiO_2 . In particular, our modified C - V measurements turned out to be a more useful tool than I - V characteristics for the quantification of interface trap density. Our concept of interface trap reduction was successfully applied to stable low voltage operation in 2D MoTe_2 FET with 30 nm BCB/10 nm Al_2O_3 bilayer dielectric. We thus conclude that the interface engineering employing the BCB dielectric offers practical benefits for the high performance and stable operation of TMD-based transistors and brightens the future of 2D TMD electronics.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors acknowledge the financial support from NRF (NRL program: Grant No. 2017R1A2A1A05001278, SRC program: Grant No. 2017R1A5A1014862, vdWMRC center), Creative Materials Discovery Program through NRF funded by the Ministry of Science, ICT and Future Planning (Grant No. 2015M3D1A1068061). This research was supported by Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education (NRF-2017R1A6A3A11034195).

Notes and references

- 1 D. Jariwala, V. K. Sangwan, L. J. Lauhon, T. J. Marks and M. C. Hersam, *ACS Nano*, 2014, **8**, 1102–1120.
- 2 Q. H. Wang, K. Kalantar-zadeh, A. Kis, J. N. Coleman and M. S. Strano, *Nat. Nanotechnol.*, 2012, **7**, 699–712.
- 3 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, M. I. Katsnelson, I. V. Grigorieva, S. V. Dubonos and A. A. Firsov, *Nature*, 2005, **438**, 197–200.
- 4 K. F. Mak, C. Lee, J. Hone, J. Shan and T. F. Heinz, *Phys. Rev. Lett.*, 2010, **105**, 2–5.
- 5 I. G. Lezama, A. Arora, A. Ubaldini, C. Barreateau, E. Giannini, M. Potemski and A. F. Morpurgo, *Nano Lett.*, 2015, **15**, 2336–2342.
- 6 S. B. Desai, G. Seol, J. S. Kang, H. Fang, C. Battaglia, R. Kapadia, J. W. Ager, J. Guo and A. Javey, *Nano Lett.*, 2014, **14**, 4592–4597.
- 7 H.-J. Chuang, X. Tan, N. J. Ghimire, M. M. Perera, B. Chamlagain, M. M.-C. Cheng, J. Yan, D. Mandrus, D. Tománek and Z. Zhou, *Nano Lett.*, 2014, **14**, 3594–3601.
- 8 K. Kang, S. Xie, L. Huang, Y. Han, P. Y. Huang, K. F. Mak, C.-J. Kim, D. Muller and J. Park, *Nature*, 2015, **520**, 656–660.
- 9 S. H. H. Shokouh, P. J. Jeon, A. Pezeshki, K. Choi, H. S. Lee, J. S. Kim, E. Y. Park and S. Im, *Adv. Funct. Mater.*, 2015, **25**, 7208–7214.
- 10 S. Kim, A. Konar, W.-S. Hwang, J. H. Lee, J. Lee, J. Yang, C. Jung, H. Kim, J.-B. Yoo, J.-Y. Choi, Y. W. Jin, S. Y. Lee, D. Jena, W. Choi and K. Kim, *Nat. Commun.*, 2012, **3**, 1011.
- 11 J. P. Shu, G. T. Wu, Y. Guo, B. Liu, X. L. Wei and Q. Chen, *Nanoscale*, 2016, **8**, 3049–3056.
- 12 M. Y. Chan, K. Komatsu, S.-L. Li, Y. Xu, P. Darmawan, H. Kuramochi, S. Nakaharai, A. Aparecido-Ferreira, K. Watanabe, T. Taniguchi and K. Tsukagoshi, *Nanoscale*, 2013, **5**, 9572–9576.
- 13 G.-H. Lee, X. Cui, Y. D. Kim, G. Arefe, X. Zhang, C. Lee, F. Ye, K. Watanabe, T. Taniguchi, P. Kim and J. Hone, *ACS Nano*, 2015, **9**, 7019–7026.
- 14 X. Cui, G.-H. Lee, Y. D. Kim, G. Arefe, P. Y. Huang, C.-H. Lee, D. A. Chenet, X. Zhang, L. Wang, F. Ye, F. Pizzocchero, B. S. Jessen, K. Watanabe, T. Taniguchi, D. A. Muller,

- T. Low, P. Kim and J. Hone, *Nat. Nanotechnol.*, 2015, **10**, 534–540.
- 15 S. M. Kim, A. Hsu, M. H. Park, S. H. Chae, S. J. Yun, J. S. Lee, D.-H. Cho, W. Fang, C. Lee, T. Palacios, M. Dresselhaus, K. K. Kim, Y. H. Lee and J. Kong, *Nat. Commun.*, 2015, **6**, 8662.
- 16 K. H. Lee, H. Shin, J. Lee, I. Lee, G. Kim, J. Choi and S. Kim, *Nano Lett.*, 2012, **12**, 714–718.
- 17 T. Kawanago and S. Oda, *Appl. Phys. Lett.*, 2016, **041605**, 108.
- 18 W. Bao, X. Cai, D. Kim, K. Sridhara and M. S. Fuhrer, *Appl. Phys. Lett.*, 2013, **042104**, 102.
- 19 H. Y. Chang, S. Yang, J. Lee, L. Tao, W. S. Hwang, D. Jena, N. Lu and D. Akinwande, *ACS Nano*, 2013, **7**, 5446–5452.
- 20 L. Chua, J. Zaumseil, J. Chang, E. C.-W. Ou, P. K.-H. Ho, H. Sirringhaus and R. H. Friend, *Nature*, 2005, **434**, 194–199.
- 21 T. D. Anthopoulos, B. Singh, N. Marjanovic, N. S. Sariciftci, A. Montaigne Ramil, H. Sitter, M. Cölle and D. M. De Leeuw, *Appl. Phys. Lett.*, 2006, **89**, 119–121.
- 22 S. R. Saudari, P. R. Frail and C. R. Kagan, *Appl. Phys. Lett.*, 2009, **023301**, 95.
- 23 H. S. Lee, S.-W. Min, Y.-G. Chang, M. K. Park, T. Nam, H. Kim, J. H. Kim, S. Ryu and S. Im, *Nano Lett.*, 2012, **12**, 3695–3700.
- 24 K. Choi, Y. T. Lee, J. S. Kim, S. W. Min, Y. Cho, A. Pezeshki, D. K. Hwang and S. Im, *Adv. Funct. Mater.*, 2016, **26**, 3146–3153.
- 25 J. Veres, S. Ogier and G. Lloyd, *Chem. Mater.*, 2004, **16**, 4543–4555.
- 26 H. Sirringhaus, *Adv. Mater.*, 2009, **21**, 3859–3873.
- 27 J. K. Jeong, *Semicond. Sci. Technol.*, 2011, **26**, 34008.
- 28 Y. Guo, X. Wei, J. Shu, B. Liu, J. Yin, C. Guan, Y. Han, S. Gao and Q. Chen, *Appl. Phys. Lett.*, 2015, **103109**, 106.
- 29 L. L. Chua, P. K. H. Ho, H. Sirringhaus and R. H. Friend, *Appl. Phys. Lett.*, 2004, **84**, 3400–3402.
- 30 L. Kong, Y. Cheng, Y. Jin, Z. Ren, Y. Li and F. Xiao, *J. Mater. Chem. C*, 2015, **3**, 3364–3370.
- 31 H. Klauk, *Chem. Soc. Rev.*, 2010, **39**, 2643.
- 32 J. S. Lee, S. Chang, H. Bouzid, S. M. Koo and S. Y. Lee, *Phys. Status Solidi A*, 2010, **207**, 1694–1697.
- 33 K. Choi, S. R. A. Raza, H. S. Lee, P. J. Jeon, A. Pezeshki, S. Min, J. S. Kim, W. Yoon, S. Ju, K. Lee and S. Im, *Nanoscale*, 2015, **7**, 5617–5623.
- 34 E. G. Bittle, J. I. Basham, T. N. Jackson, O. D. Jurchescu and D. J. Gundlach, *Nat. Commun.*, 2016, **7**, 10908.
- 35 M. Kimura, *Solid-State Electron.*, 2011, **63**, 94–99.
- 36 N. L. Cohen, R. E. Paulsen and M. H. White, *IEEE Trans. Electron Devices*, 1995, **42**, 2004–2009.
- 37 H. S. Lee, S. S. Baik, K. Lee, S. Min, P. J. Jeon, J. S. Kim, K. Choi, H. J. Choi, J. H. Kim and S. Im, *ACS Nano*, 2015, **9**, 8312–8320.