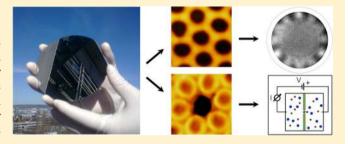
Centimeter-Sized Single-Orientation Monolayer Hexagonal Boron Nitride With or Without Nanovoids

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Supporting Information

ABSTRACT: Large-area hexagonal boron nitride (h-BN) promises many new applications of two-dimensional materials, such as the protective packing of reactive surfaces or as membranes in liquids. However, scalable production beyond exfoliation from bulk single crystals remained a major challenge. Single-orientation monolayer h-BN nanomesh is grown on 4 in. wafer single crystalline rhodium films and transferred on arbitrary substrates such as SiO2, germanium, or transmission electron microscopy grids. The transfer process involves application of tetraoctylammonium bromide before



electrochemical hydrogen delamination. The material performance is demonstrated with two applications. First, protective sealing of h-BN is shown by preserving germanium from oxidation in air at high temperatures. Second, the membrane functionality of the single h-BN layer is demonstrated in aqueous solutions. Here, we employ a growth substrate intrinsic preparation scheme to create regular 2 nm holes that serve as ion channels in liquids.

KEYWORDS: Hexagonal boron nitride, 2D materials, chemical vapor deposition, nanoporous membranes, electrochemical delamination

wo-dimensional (2D) materials are expected to significantly widen the prospects for condensed matter systems¹ and to unlock new functionalities in optoelectronics² or ionic transport.3-5 However, if the materials shall become useful, they have to be produced in a scalable way. This means that the exfoliation procedure of 2D flakes from 3D single crystals has to be replaced by processes that yield large-area material. This also applies to hexagonal boron nitride (h-BN) that emerges to be ideal for the encapsulation of graphene-based electronic devices⁶ or that is a promising membrane material in liquids.^{3,4} Previous reports on transfer of chemical vapor deposition (CVD) h-BN by the H₂ "bubbling" delamination method⁷⁻⁹ show that the transfer size was limited to micrometer flakes.⁷ Recent work reports the transfer of large-area multilayer h-BN via the etching of the sapphire substrates.8 Up to now, the transfer of large-area h-BN from metal growth substrates remained an unsolved problem. Here, we report on the transfer process of large-area single layer h-BN with single orientation that is grown on 4 in. wafer-scale single-crystalline Rh(111) films. The transfer of continuous layers is evidenced for the bare eye with oxidation protection experiments of germanium. The two-step transfer protocol involves electrochemical

tetraoctylammonium ion (TOA+) treatment prior to a hydrogen "bubbling" at the interface of h-BN and metal substrates. It is reliable (high transfer rates above 90%) and gentle because the transferred h-BN layers not only maintain single crystal orientation but also preserve surface modifications such as 2 nm voids that rely on the growth substrates.

As a growth substrate, we employ twin-free single-crystalline Rh(111) thin films on 4 in. Si(111) wafers. A corrugated h-BN layer is grown in ultrahigh vacuum (UHV) by CVD. 10,11 The corrugation appears as a "mesh" superstructure with "pores" of about 2 nm diameter, in which the BN is closely attached to the substrate, and a surrounding "wire" region with a weaker bonding and a larger layer-substrate distance (Figure 1a). 12-14 This nanomesh surface is stable in air and in liquids, ^{13,15} and its corrugation can be switched on and off via hydrogen intercalation, either in vacuum¹⁶ or in liquids.⁴ With additional preparation steps, single 2 nm voids are fabricated with the

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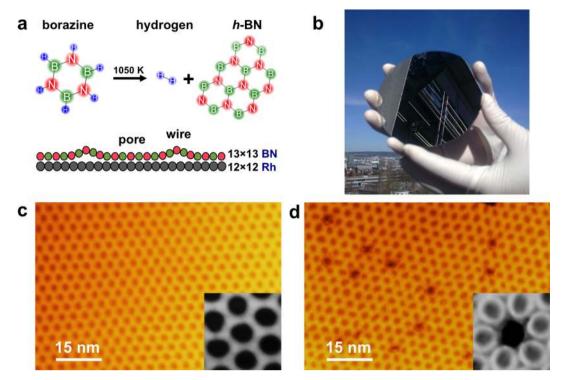


Figure 1. h-BN nanomesh on Rh(111) with and without 2 nm voids. (a) Chemical structure of the borazine (HBNH) $_3$ precursor and its decomposition to form a h-BN monolayer and H $_2$ gas on a hot Rh(111) surface. The h-BN corrugation is caused by the lattice mismatch that accommodates 13 × 13 BN on 12 × 12 Rh units. (b) Photograph of a 4 in. wafer single-crystalline Rh(111) thin film substrate in Zürich air, on which the nanomesh is grown. (c) Room-temperature STM images of pristine h-BN nanomesh on Rh(111), $U_t = -1.20$ V, and $U_t = 0.50$ nA. (d) Multiple 2 nm voids at pore sites generated by the "can-opener" effect after annealing of u-BN/Ar/Rh(111) to 900 K, u-1.20 V, u-1.20

"can-opener" effect to form a surface with uniformly distributed voids in the h-BN layer, 17,18 termed voidal boron nitride (ν -BN). The functionality of the freestanding ν -BN material is demonstrated by immersion of such nanoporous membranes in aqueous solutions and the measurement of ion conductivity.

Results and Discussion. Pristine h-BN monolayers are prepared by the high-temperature CVD method with borazine (HBNH)₃ as precursor on single-crystalline Rh(111)/yttriastabilized zirconia(YSZ)/Si(111) substrates at 1050 K. ^{11,12} The sample fabrication is scaled up to 4 in. wafers (Figure 1b), and the quality of the h-BN is comparable to the one fabricated on Rh(111) single crystals (Figure 1c). ¹¹ The ν -BN is prepared by exposing h-BN/Rh(111) to low-energy Ar ions followed by a subsequent annealing procedure. ^{17,19} The ion bombardment generates vacancy defects in the boron nitride layer, which, upon annealing to 900 K, leads to the "can-opener" effect, i.e., the formation of 2 nm voids in the h-BN layer, ¹⁸ as shown in the scanning tunneling microscopy (STM) image in Figure 1d.

Figure 2 is a schematic illustration of the transfer procedure for single layer *h*-BN. After UHV-CVD growth (Figure 2-1), the *h*-BN samples are introduced into a three-electrode setup in a tetraoctylammonium bromide (TOABr)/acetonitrile solution (Figure 2-2).²⁰ The reference potential is kept at -1.9 V for a thorough diffusion of molecules and ions in the solution and for enabling chemical reactions on the electrodes. This TOA⁺-treatment step ensures transfer rates above 90%, while its omission typically yields transfer rates below 10%, as shown in the Figure S1. Subsequently, the TOA-treated sample (TOA/*h*-BN/Rh) is spin-coated with a polymethylmetacrylate (PMMA) film (Figure 2-3), followed by electrochemical hydrogen intercalation⁴ and subsequent "bubbling" delamination²¹,²²

(Figure 2-4). Finally, the delaminated PMMA/TOA/h-BN film in Figure 2-5 may be transferred to an arbitrary substrate.

After transfer, the remaining Rh substrates, as indicated in Figure 2-6, can be recycled to regrow h-BN, in which the quality of the regrown h-BN monolayer stays high (Figure S2). In the following, we report transfer results on transmission electron microscopy (TEM) grids (Figure 2-7), on SiO₂ and on Ge(111) wafers (Figure 2-8).

To characterize the transferred PMMA/TOA/h-BN layer, they are placed onto SiO₂/Si substrates and annealed in vacuum to 650 K. The thickness of the 80 nm SiO₂ is chosen to maximize the optical contrast of h-BN²³ and to still allow good electron spectroscopy results by avoiding strong sample charging. Figure 3a,b shows the X-ray photoelectron spectroscopy (XPS) spectra of B1s and N1s before (black) and after (red) transfer (the survey spectra are shown in Figure S3). We define the transfer rate as the ratio of the XPS intensities of transferred BN and pristine BN (B1s and N1s) (see section 4 in the Supporting Information). The XPS results indicate that 95% of the h-BN monolayer is transferred, which fits well with the optical microscopy observation of intact layers (Figure S5). The angle scanned X-ray photoelectron diffraction (XPD) of B1s and N1s in Figure 3c,e indicate a flat single-orientation h-BN monolayer transferred onto amorphous SiO₂. The 3-fold symmetry with the 3 grazing angle scattering intensity maxima that are rotated by 60 degree between nitrogen and boron is known from h-BN/Ni(111)²⁴ and h-BN/Rh(111).²⁵ Comparison of the XPD patterns of B1s and N1s with the atomic structure of h-BN in Figure 3d thus proves that the transferred layer consists of large-area single-orientation h-BN. The monolayer h-BN is consistent with Raman measurements, i.e.,

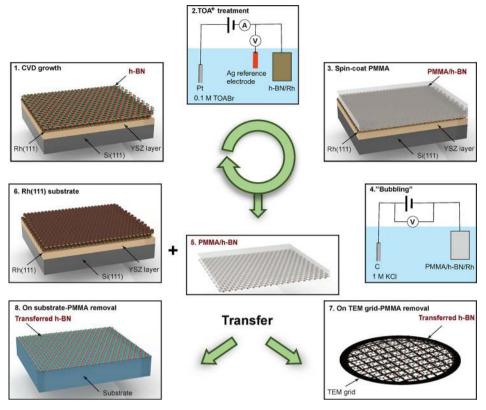


Figure 2. Two-step TOA⁺-assisted electrochemical transfer process of *h*-BN. (1) Structure of *h*-BN/Rh(111) film sample. On a Si(111) wafer, an yttria-stabilized zirconia layer (YSZ) (80 nm), Rh(111) (150 nm), and a *h*-BN monolayer (with or without nanovoids) are grown. (2) First step of the transfer procedure: TOA⁺-treatment in a three-electrode setup, consisting of a *h*-BN/Rh sample as working electrode, a Pt wire as counter electrode, and an Ag wire as reference electrode in 0.1 M TOABr/acetonitrile solution. (3) The TOA/*h*-BN/Rh sample is spin-coated with PMMA. (4) The second step includes a two-electrode setup with the PMMA/TOA/*h*-BN/Rh sample as working electrode and a graphite rod as counter electrode in a 1.0 M KCl solution. H₂ bubble evolution between *h*-BN and Rh substrate lifts the PMMA/TOA/*h*-BN monolayer (shown in panel 5). (6) Rh(111) substrate after *h*-BN transfer can be used for another cycle of *h*-BN growth (panel 1). The PMMA/*h*-BN films are transferred on TEM grids (panel 7) and on SiO₂/Si or Ge substrates (panel 8), followed by PMMA removal.

the characteristic Raman peak at 1368 cm⁻¹ with a full width at half-maximum (fwhm) of 20 cm⁻¹ (Figure S6).²³

To demonstrate the protection functionality of the transferred h-BN, we developed a simple "litmus" test that uses the oxidation of germanium in air as an indicator. Germanium is known to oxidize at temperatures above 370 °C in air, ²⁶ where it changes color from gray to light blue. In Figure 4, the color-change of a piece of Ge(111) wafer is shown during annealing from room temperature up to 610 °C. We track 3 different 2 × 2 mm² regions, which are (i) pristine Ge, (ii) covered with PMMA, and (iii) with PMMA/TOA/h-BN exfoliate that is folded to a double layer. The single fold minimizes the coincidence of defects in the single layer and thus enhance the quality of protection.

After sublimation of the PMMA at about 350 °C, the color is the same for all 3 spots. Upon further temperature increases, the optically visible oxidation process begins. However, the rates of color change are significantly different for the 3 regions, where it becomes clear that the h-BN covered spot resists oxidation up to 610 °C. It is also observed that the PMMA-covered Ge area keeps the memory of having been covered, even though PMMA is no longer optically visible between 350 and 500 °C. This test reflects the superior sealing behavior of atomically thin h-BN in extreme environments. Such membranes, for example, allow the design of packing electronic components with tightest space constraints and best thermal energy dissipation properties. Furthermore, the Ge protection

test signifies a viable low-cost tool for the characterization of the quality of macroscopic BN exfoliates. In particular, we point out that the test reveals sample inhomogeneities, therefore minimizing the need for analysis with expensive surface science methods. As such, it appears to be a useful benchmark method for optimization and comparison of material production and transfer processes.

Finally, we demonstrate the application of freestanding h-BN and v-BN exfoliates in membrane applications. State of the art TEM and scanning TEM (STEM) are known for their capability to measure h-BN and graphene with atomic resolution.²⁷ The h-BN layers are suspended over TEM grids (Cu quantifoil and SiN_x membranes) with diameters of the holes from 1.0 µm down to self-made 50 nm. Figure 5 shows spherical aberration-corrected TEM and STEM images, acquired at room temperature and with electron energies of 80 keV, which is between the boron and the nitrogen knock-on threshold in monolayer h-BN.^{28,29} Figure 5a depicts a TEM grid with 1 μ m holes covered by a single layer h-BN (dark-field TEM). The further zoom-in TEM image in Figure 5b shows freestanding monolayer h-BN with remaining PMMA and TOA traces. Atomically clean single layer h-BN patches can easily be found (Figure 5c). Figure 5d-f demonstrates the application of single layer v-BN with 2 nm voids as permeable membranes in a potassium chloride (KCl) solution. Such membranes are potential candidates for blue energy harvesting like osmotic

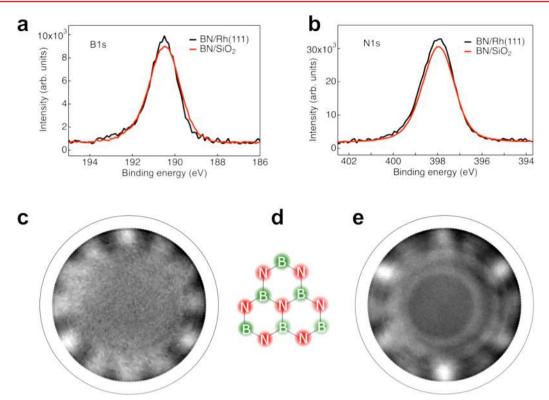


Figure 3. Mg Kα XPS and XPD ($\hbar\omega$ = 1253.6 eV) of h-BN before and after TOA⁺-assisted transfer. (a, b) XPS of B1s and N1s core levels on Rh before (black) and after transfer on 80 nm SiO₂/Si (red). The transfer rate is above 95%. (c, e) X-ray photoelectron diffraction patterns of (c) B 1s and (e) N 1s of transferred h-BN layer on amorphous SiO₂. The 3-fold symmetry indicates a single orientation of the h-BN lattice as depicted in panel d.

power generation as recently demonstrated for MoS₂ membranes⁵ or may be used for ion separation.³⁰

In our ion conductivity setup,³¹ a circular 50 nm hole in 20 nm thick SiN membrane is covered with v-BN membrane (Figure 5d). In Figure 5e, a 5 nm × 5 nm STEM image with one void is shown. Measurements of current versus voltage (I-V) characteristics demonstrate that ion conductivity across such voidal membranes can be measured. In Figure 5f, conductivities of 0.01 M KCl, pH = 8, at room temperature are displayed for four different v-BN membranes. Below cell voltages of ± 200 mV, conductivities of $(-1 \pm 3) \times 10^{-3}$, 0.48 \pm 0.07, 0.36 \pm 0.18, and 4.24 \pm 0.07 nS are obtained (for details, see part 7 in the Supporting Information). 32-34 From the conductance of the KCl solution and the void dimensions, we estimate the number of voids or ion channels to be 0, 2, 1, and 16 for the 4-membrane I-V curves shown in Figure 5f. The hole quantities are in the same order of magnitude as expected from the nominal ν -BN void density before transfer of 2.3 \times 10⁻³ nm⁻² or 3 to 6 voids on a membrane with diameter of 50 nm. Given the uncertainties in relating the measured conductance to the void density we consider the conductivities of membrane samples (#1 and #2 in Figure 5f) to be close to the expectation. At variance to solid-state electrolyte boron nitride membrane applications, 35 we also find ion tight membranes (#0). #3 with 4 nS conductivity likely corresponds to a broken membrane.

Conclusions. A method for transfer of centimeter-sized single-orientation h-BN with or without surface modifications (nanovoids) from single-crystalline Rh(111) thin films is presented. The process involves two steps, a TOA⁺-treatment of the h-BN surface, followed by hydrogen driven delamination.

Advantages of this nondestructive isolation of CVD-grown h-BN include: (1) scalability; (2) substrate recycling; and (3) the fact that functionalizations that rely on the metal substrate like the "can-opener" effect are sustained. To confirm the application potential of these 2D layers, we demonstrate their thermal oxidation protection capability in air and the use as monatomic membranes with and without ion channels in aqueous solutions.

Methods. Instrumentation for Sample Fabrication. The CVD growth of h-BN was performed in an UHV system with a base pressure of 1×10^{-10} mbar. For the high-temperature CVD fabrication of the h-BN nanomesh, we employed a standard procedure on 150 nm single-crystalline Rh(111)/YSZ/Si films at 4 in. wafer scale. For the 2 nm void generation, we used a Specs IQP 10/35 Penning-type ion source run at lowest acceleration potential. The integrated sputter charge density was 250 nC/cm².

Two-Step TOA⁺-Assisted Electrochemical Process. The first step involves a three-electrode electrochemical setup. It consists of a *h*-BN/Rh sample as working electrode, a Pt wire as counter electrode, and an Ag-wire as reference electrode in Ar degassed 0.1 M TOABr/acetonitrile. The reference potential versus Agwire was kept at −1.9 V for 10 min, followed by a potential of −0.2 V versus Ag-wire for 20 s. Subsequently, the sample was rinsed with acetonitrile and spin-coated with 4 wt % PMMA (495 K). The second step involves the PMMA/TOA/*h*-BN/Rh sample as working electrode and a graphite rod as counter electrode in 1.0 M KCl solution. A negative voltage between −3 and −6 V was applied to the sample while it was gradually immersed into the solution. H₂ bubbles form at the *h*-BN/Rh interface and delaminate the *h*-BN/TOA/PMMA film from the

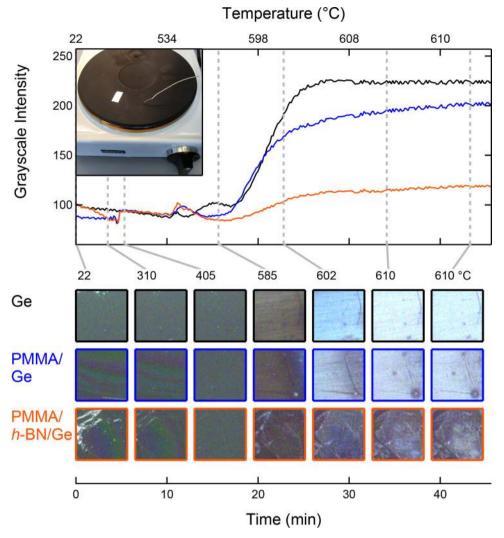


Figure 4. Large-scale high-temperature protection of Ge against oxidation by transferred h-BN. The inset shows the heating plate with a piece of Ge wafer on top. PMMA and transferred PMMA/h-BN are placed on a Ge(111) surface. Top panel: grayscale intensities of the camera images of Ge (black), PMMA/Ge (blue), and PMMA/h-BN/Ge (red) as a function of sample temperature and annealing time. The process involves three stages: (1) PMMA desorption at 300–350 °C. (2) At ~405 °C, the three samples show the same color. (3) At 585 °C, bare Ge and PMMA/Ge (after PMMA removal) start oxidizing, and the colors of the surfaces turn bright blue (Ge bulk oxide formation), while PMMA/h-BN/Ge stays dark even after 10 min at 610 °C. Bottom frame panel: Surface color changes of the three regions (7 frames each) vs time and for temperatures from room temperature up to 610 °C. All squares represent 2 mm × 2 mm. The color codes of the frame-boarders are the same as the traces in the top panel.

substrate. The delaminated h-BN/TOA/PMMA film was then rinsed in ultrapure water (Milli-Q Advantage A10) and transferred on a clean 80 nm SiO₂/Si substrate. In the next step, the PMMA was removed via a sequence of acetone/ ethanol baths at 370 K and gradual anneal in air up to 600 K for 3 h. Two types of TEM grids were used: (1) Cu quantifoil with an empty hole size of 1000 nm and interval distance of holes of 800 nm (Figure 5a) and (2) SiN_x with a thickness of 20 nm and a single hole with 50 nm in diameter in the middle (Figure 5d). The TEM grids were coated with 25 nm Pt prior to the transferred-layer deposition. The Pt acts as a catalyst that reduces the PMMA disintegration temperature and improves the PMMA removal efficiency.³⁶ After transfer, the PMMA/ TOA/h-BN/grids were slowly heated in air to 600 K for 5 h to remove PMMA/TOA. The Rh(111) film substrate was then put back to the UHV system and heated up to 800 K before XPS and low-energy electron diffraction (LEED) measurements.

Sample Characterizations. All sample characterizations were carried out at room temperature. The scanning tunneling microscopy (STM) measurements were accomplished with an Omicron variable temperature STM (VT-STM) with electrochemically etched tungsten tips. All STM images were taken in constant-current mode.

Optical images were acquired using a Leica DMV2500 stereomicroscope with a maximum magnification of 2500×. Sequential frame acquisition was performed using procedures written with the Leica Self-Development Kit package, and the frames were subsequently stitched using the Fiji image-processing package.

XPS, X-ray photoelectron diffraction (XPD), and LEED measurements were carried out on a user-modified Vacuum Generators ESCALAB 220³⁷ with a Mg K α lab source at an energy of $\hbar\omega=1253.6$ eV. X-ray photoemission electron microscopy (X-PEEM) images were recorded at the boron K-edge at the Surface/Interface Microscopy (SIM) beamline of the Swiss Light Source (SLS).

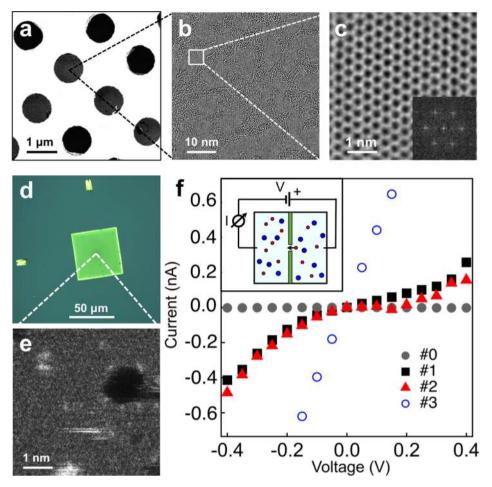


Figure 5. Freestanding h-BN membranes. (a) Dark-field TEM image of a large-area h-BN membrane suspended over a Cu quantifoil TEM grid. (b) Zoomed-in TEM image displays freestanding single-layer h-BN, with some transfer-related residues. (c) Further zoomed-in high-resolution (HR)-TEM image: Single-crystalline BN honeycomb lattice with atomic resolution. Inset: Fast Fourier transform (FFT) of panel c, confirming the crystalline structure with hexagonal symmetry. (d) Optical microscopy image of a h-BN membrane with 2 nm voids on a 20 nm SiN $_x$ membrane with 50 nm hole in the middle. (e) HR-STEM image of a h-BN membrane with a 2 nm void. (f) Current vs voltage characteristic of 4 different v-BN membrane samples in 10 mM KCl solutions at pH 8. The inset shows the sketch of the setup. All TEM and STEM images were taken at 80 keV electron energy and with the sample at room temperature.

Raman measurements were performed with a Thermo Scientific DXRTM 2 Raman Microscope equipped with CCR1000 Linkam at 455 nm excitation wavelength. Scanning electron microscopy in Figure S2 were measured using a Zeiss Supra 50 VP field-emission SEM equipped with secondary, backscattered electron and EDX detectors at energies between 2–2.5 keV.

TEM and STEM images were measured on an aberration-corrected (with double Cs corrector) FEI Titan Themis TEM 60-300 operated at an $80~\rm kV$ acceleration voltage. For HRTEM imaging, a spherical aberration (Cs) of about $-20~\mu m$ was applied, combined with a monochromatic incident illumination to reduce the effects of chromatic aberration, for an optical setup capable of $1~\rm Å$ resolution or better. Images are shown with focus adjusted to give a white atom contrast, 38 which, when imaging a clean uncontaminated monolayer, allows the phase-contrast HRTEM image to be directly interpreted as atomic positions. (S)TEM image processing: Raw TEM images were filtered by using a bandpass filter between 1 and 100 pixels. These filtered images were then blurred out using a Gaussian blur function with a diameter of 1.2 pixels to enhance visual inspections.

Nanofluidic transport experiments were performed using the setup described previously. After mounting in the chamber, chips were wet with $\rm H_2O/ethanol~(1:1)$ solution for 30 min. Nanofluidic measurements were performed by taking the current—voltage response ($\it I-V$ characteristics, respectively) of the nanovoid in salt solutions of potassium chloride (KCl, Sigma-Aldrich) using an Axopatch 200B patch-clamp amplifier (Molecular Devices, Inc. Sunnyvale, CA) with voltage range from -1.0 to 1.0 V.

ASSOCIATED CONTENT

S Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.nanolett.7b04752.

Transfer-methods comparison, reusability of substrates, characterization details, ion conductivity model, and supporting data. (PDF)

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Notes

The authors declare no competing financial interest.

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