

Electronic Supplementary Information

Optical Readout of Charge Carriers Stored in a 2D Memory Cell of Monolayer WSe₂

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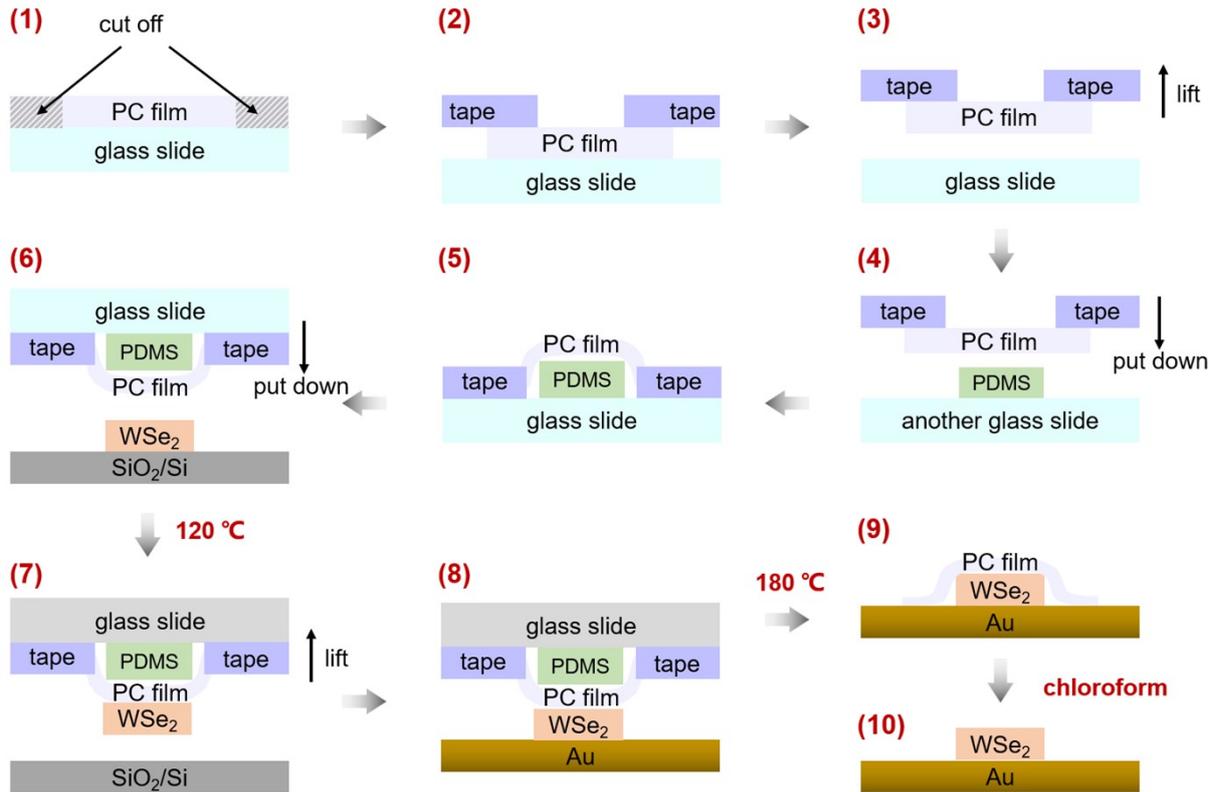


Fig. S1 Schematic diagram for the fabrication process of a 2D WSe₂ memory cell (see detailed descriptions in the sample preparations section of the main text). Of special note is that the PDMS film used here is pre-cleaned by plasmas to increase the surface viscosity, so that it can be tightly attached to the PC film for the eventual obtainment of a wrinkle-free WSe₂ monolayer. In the plasma-cleaning process, the glass substrate with a PDMS film on top was first placed inside a reaction cavity, which was then pumped to the vacuum with a pressure value of 100 pa. Under the continuous O₂ flow at the speed of 50 mL/min, the plasma generator was started to create a high-voltage AC (alternate-current) field between the two electrodes of the reaction cavity. This electric field could accelerate the free electrons to a high energy, which was sufficient to excite the O₂ molecules for the generation of plasmas. During the 3-5 s cleaning time, the high-energy plasmas reacted with the organic contaminants or collided with the micro-sized particles on the PDMS surface to yield various volatile materials. These volatile materials would be pumped out of the cavity together with the O₂ flow, leaving behind a PDMS film with the high surface cleanness and activity.

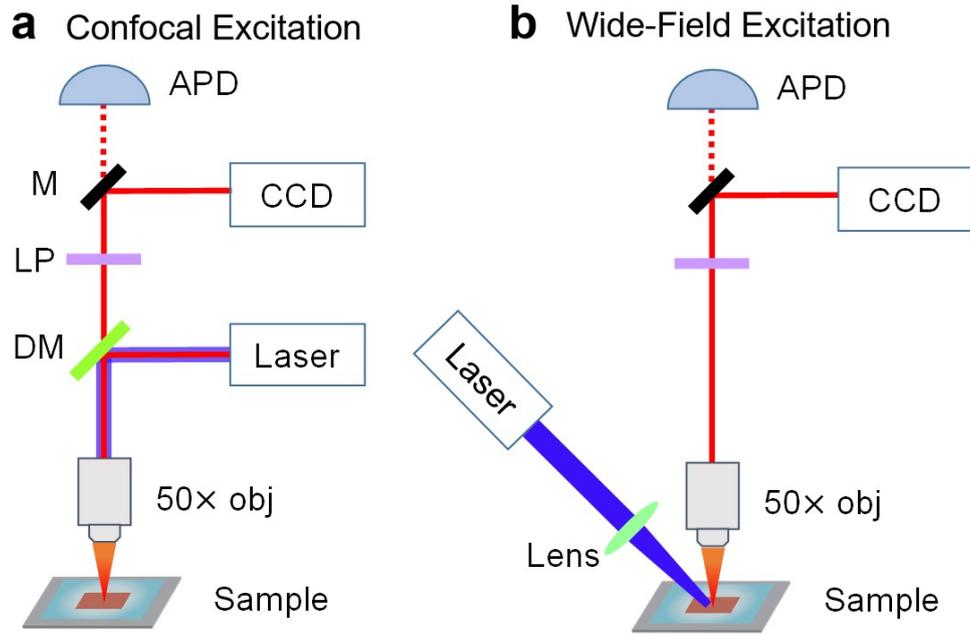


Fig. S2 Schematic diagrams for the optical measurements of 2D WSe₂ memory cells using the (a) confocal and (b) wide-field setups (see detailed descriptions in the optical measurements section of the main text). APD: avalanche photodetector; M: mirror; CCD: charge-coupled device; LP: long-pass filter; DM: dichroic mirror; obj: objective.

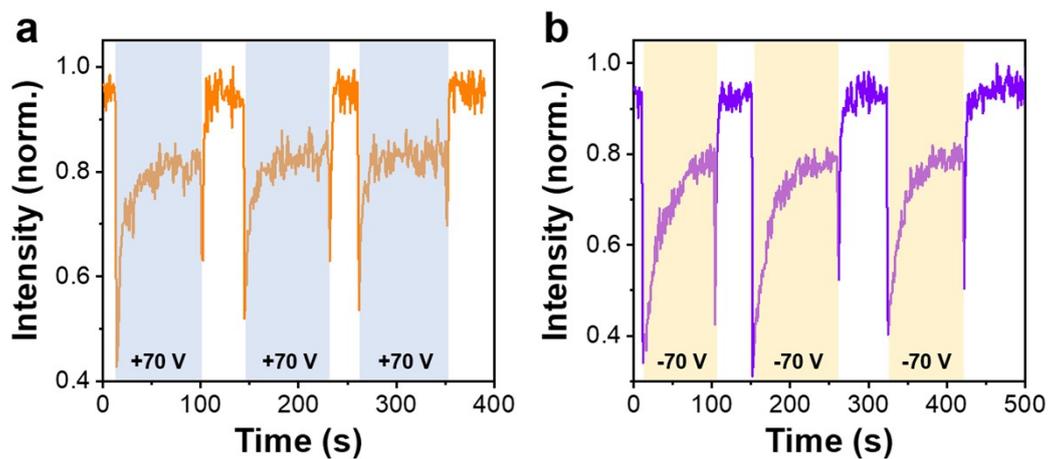


Fig. S3 Time-dependent PL intensities of WSe₂ measured when the external biases of (a) +70 V and (b) -70 V are first applied to and then removed from the electrode for several times. The PL intensity of WSe₂ stays at almost the same level when there is no external bias on the electrode, showing stable operation of the memory cell during the fast charging and discharging processes.

Positions	A_1	$\tau_1(\text{ns})$	A_2	$\tau_2(\text{ns})$
A	1.00	0.25		
B	0.77	0.43	0.23	1.45
C	0.85	0.16	0.15	0.92
D	0.80	0.40	0.20	1.43

Table S1 Fitting parameters for the PL decay curves of monolayer WSe₂. For position A, the PL decay curve is fitted by a single-exponential function of $A_1 e^{-t/\tau_1}$. For positions B-D, the PL decay curves are each fitted by a bi-exponential function of $A_1 e^{-t/\tau_1} + A_2 e^{-t/\tau_2}$.

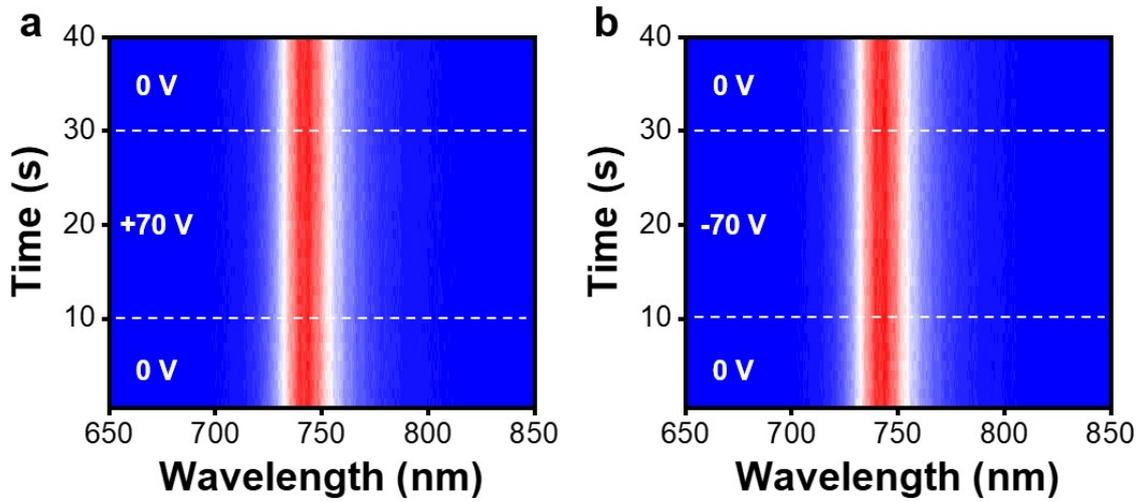


Fig. S4 Time-dependent PL spectral image measured for monolayer WSe₂ placed between two non-contacting electrodes. (a) A +70 V bias is applied to and then removed from the two electrodes at 10 and 30 s, respectively. (b) A -70 V bias is applied to and then removed from the two electrodes at 10 and 30 s, respectively.

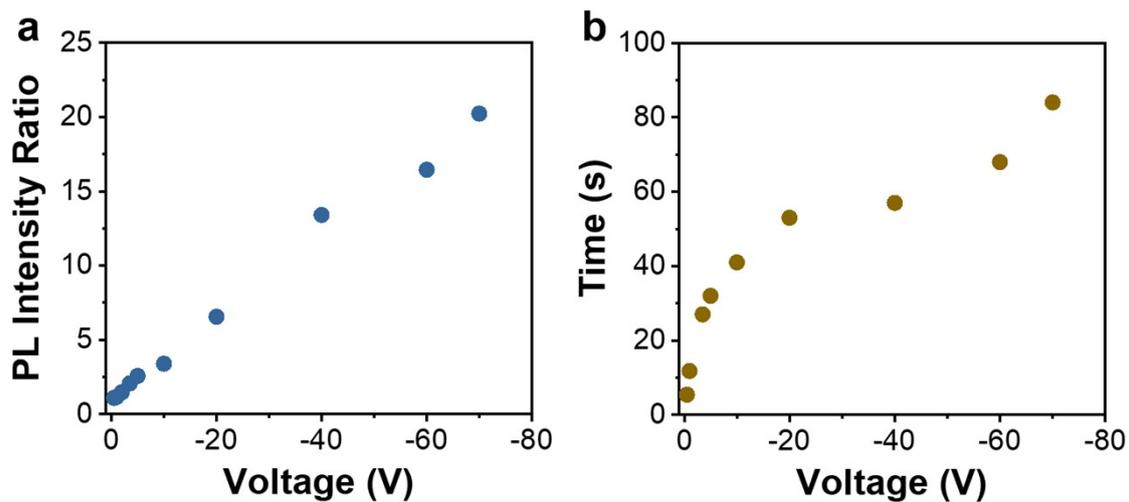


Fig. S5 (a) The ratio of WSe₂ PL intensities measured before and right after an external bias from -0.5 to -70 V has been applied to the electrode. A larger PL intensity ratio corresponds a more negatively-charged WSe₂ monolayer. (b) Electron retention time measured after an external bias from -0.5 to -70 V has been applied to the electrode.

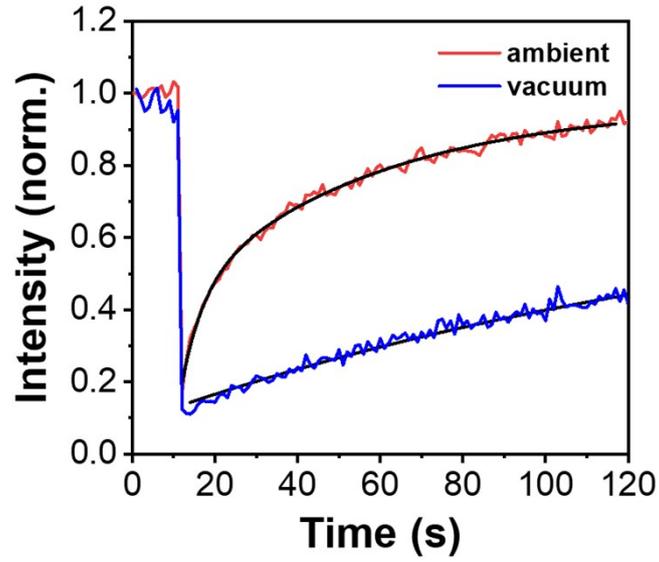


Fig. S6 Time-dependent PL intensities measured under the ambient and vacuum conditions for the same monolayer WSe₂ before and after an external bias of -10 V is applied to the electrode at 10 s, where the exponential fittings yield the electron retention times of ~27 and ~163 s, respectively. The above measurements are performed at room temperature, with the monolayer WSe₂ being excited at ~30 W/cm² by a pulsed 405 nm laser.

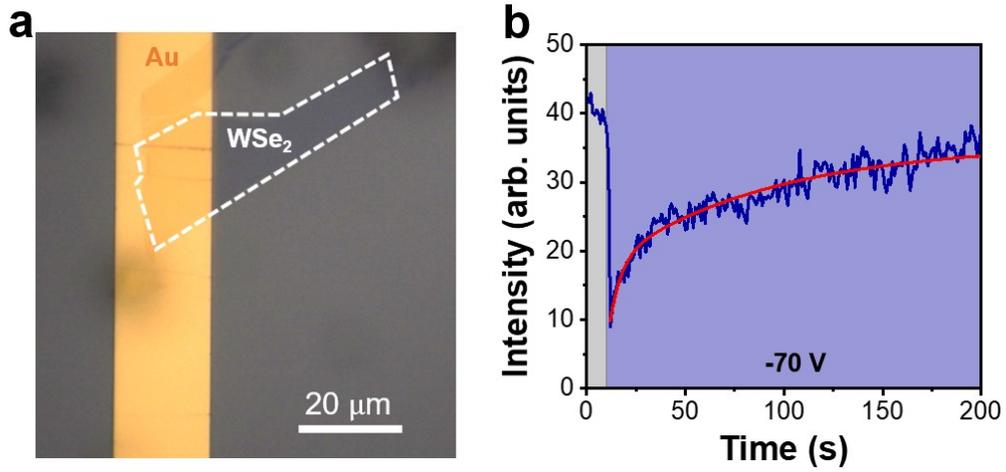


Fig. S7 (a) Optical image of a memory cell with only one electrode, above which a WSe₂ monolayer is partially deposited to serve as the channel material. (b) Time-dependent PL intensity measured at room temperature for monolayer WSe₂ before and after an external bias of -70 V is applied to the electrode at 10 s. The PL intensity rise curve measured during 10-200 s is exponentially fitted to yield an electron retention time of ~81 s.

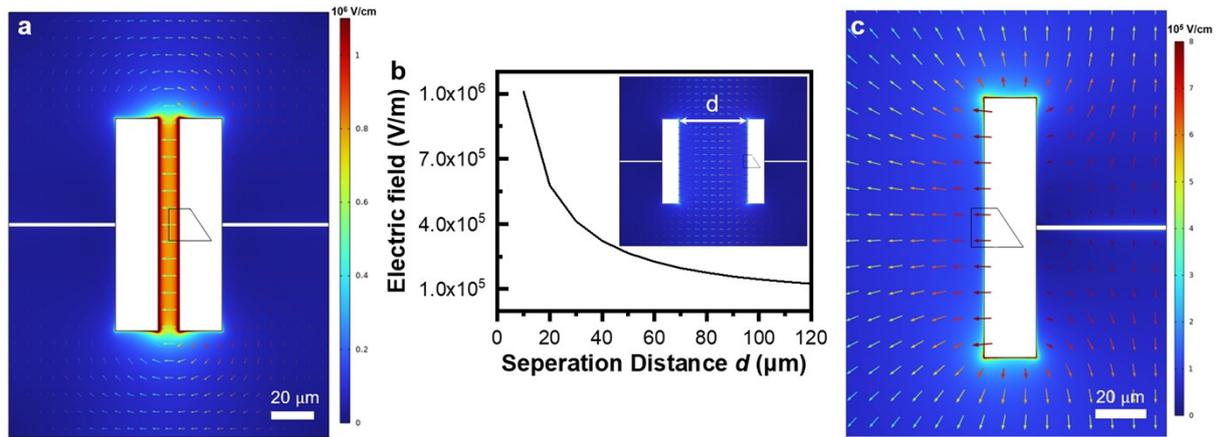


Fig. S8 (a) Simulated electric-field distribution around the two electrodes (white regions) being grounded (left) and biased at +10 V (right), respectively. The two electrodes are separated by 10 μm , and each of them has the width, length and thickness of 20 μm , 100 μm and 130 nm, respectively. The monolayer WSe_2 (black right-angled trapezoid) is deposited partially on top of the right electrode, while the average electric field estimated within its non-contacting area is $\sim 1.00 \times 10^6$ V/m. (b) Average electric field simulated within the non-contacting area of monolayer WSe_2 as a function of the separation distance d between the two electrodes being grounded and biased at +10 V, respectively. Inset: Representative electric-field distribution simulated at $d = 80$ μm . According to these simulated results, the average electric field starts from $\sim 10^6$ V/m at $d = 10$ μm and decreases exponentially with the further increase of d . However, once d is larger than ~ 100 μm , this electric field would stay at almost a constant value of $\sim 10^5$ V/m. (c) Simulated electric-field distribution around the one electrode being biased at +10 V, with the grounded position being set at the source meter used in our experiment. The average electric field estimated within the non-contacting area of monolayer WSe_2 is $\sim 1.46 \times 10^5$ V/m, which is confirmed by us to have the same charge injection efficiency as that of the $\sim 10^6$ V/m case (see Figure S7). The above simulations are performed by a finite-element method using the Comsol Multiphysics software.

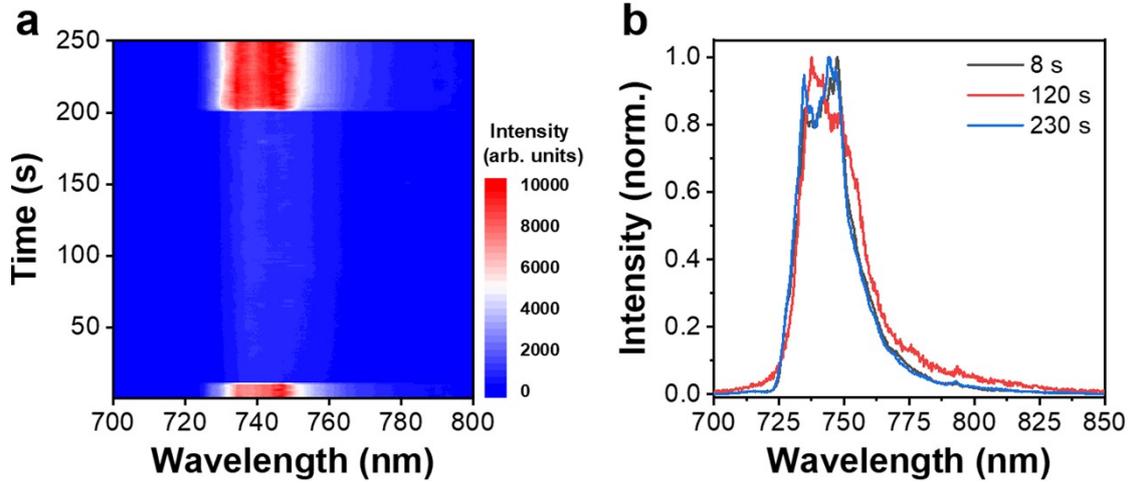


Fig. S9 (a) Time-dependent PL spectral image measured for monolayer WSe₂ at 4 K, wherein a -70 V bias is first applied to and then removed from the electrode at 10 and 200 s, respectively. (b) PL spectrum extracted from (a) at 120 s (solid red line), showing the emergence of a red-shifted component due to the more negatively-charged condition of monolayer WSe₂. For comparison, the PL spectra extracted from (a) at 8 (solid black line) and 230 s (solid blue line) are also provided, corresponding to the originally n-doped condition when no external bias is applied to the electrode.

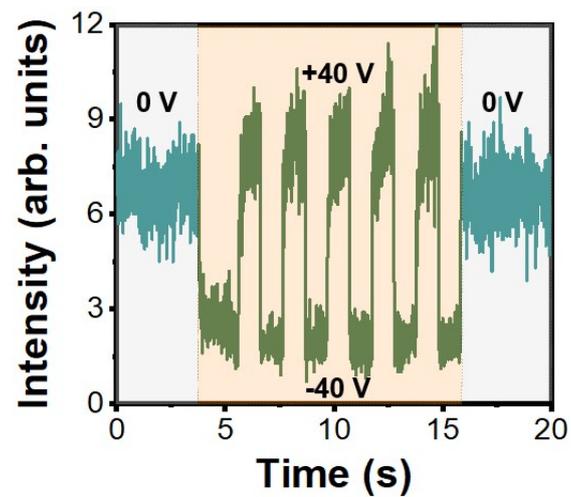


Fig. S10 Time-dependent PL intensity measured for monolayer WSe₂ under alternating applications of the ± 40 V biases to the electrode.