

Supporting Information

Controllable p-type doping of 2D MoS₂ via Sodium intercalation for optoelectronics

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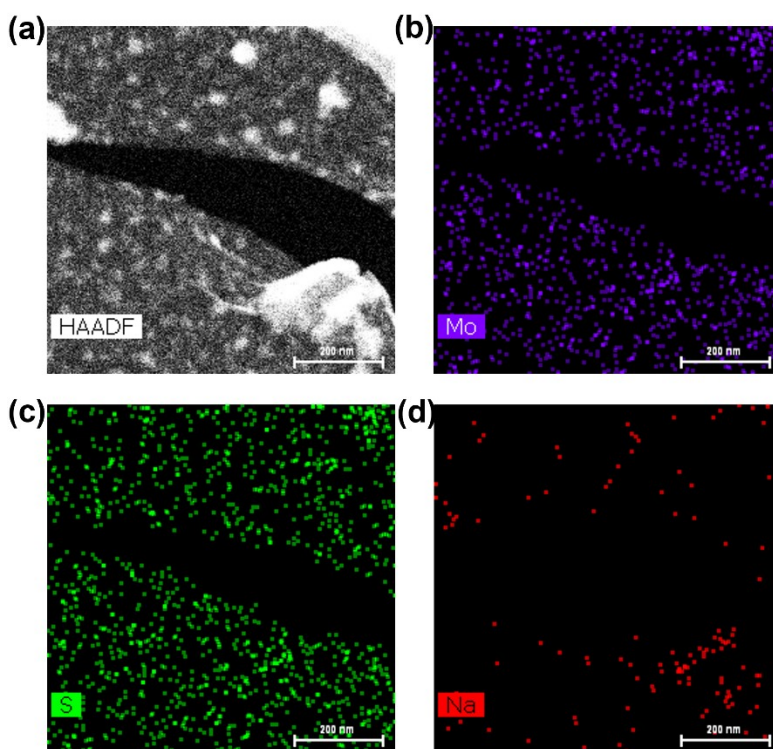


Figure S1. STEM EDS mapping of thin MoS₂ with NaOH grown on SiO₂/Si (a) High-angle-annular-dark-field (HAADF) image of the MoS₂ with NaOH on SiO₂/Si substrate. (b), (c), and are the elemental maps of MoS₂ on SiO₂/Si, showing the location of Mo, S, and Na respectively.

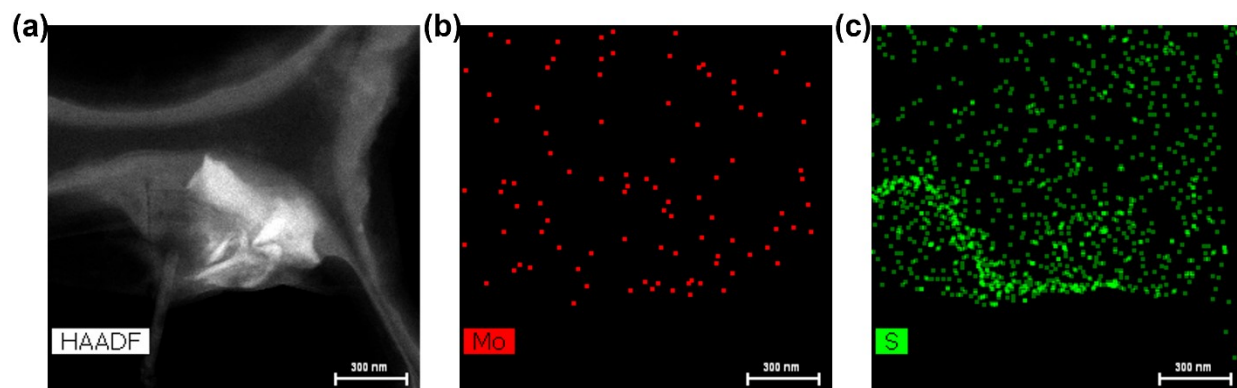


Figure S2. STEM EDS mapping of thin MoS₂ without NaOH grown on SiO₂/Si (a) High-angle-annular-dark-field (HAADF) image of the MoS₂ without NaOH on SiO₂/Si substrate. (b) and (c) are the elemental maps of MoS₂ on SiO₂/Si, showing the location of Mo and S respectively.

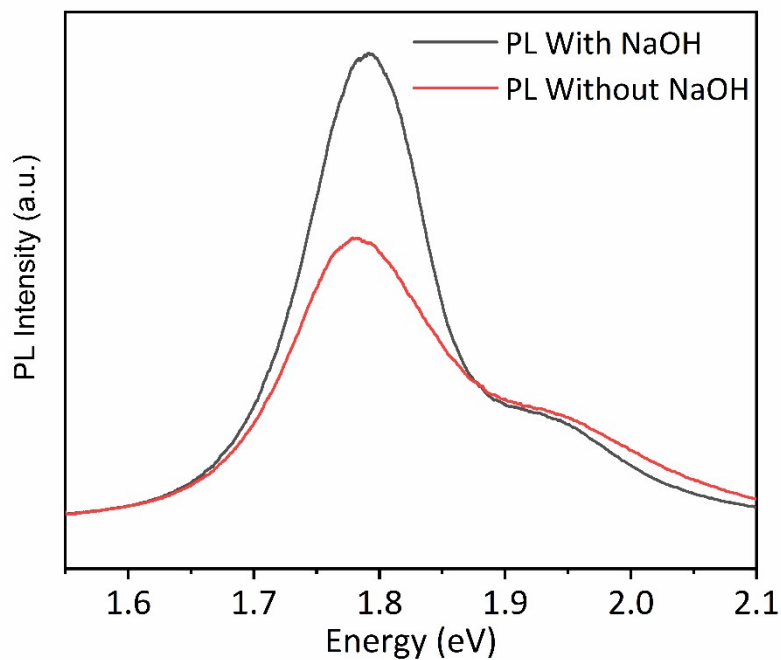


Figure S3. PL spectra of MoS₂ with and without NaOH

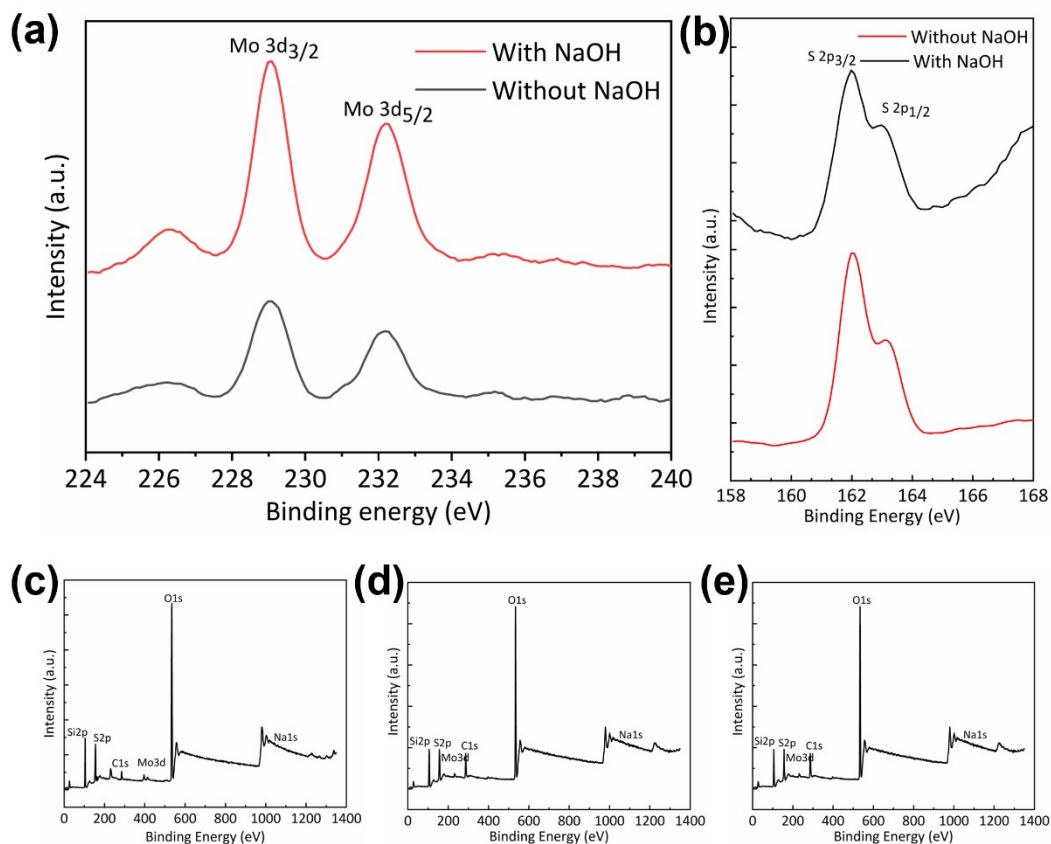


Figure S4. XPS spectra of (a) Mo 3d and (b) S 2p of the sample with and without NaOH. Survey spectrum of Na-doped MoS₂ (c) Low doped concentration (d) medium concentration (e) High concentration

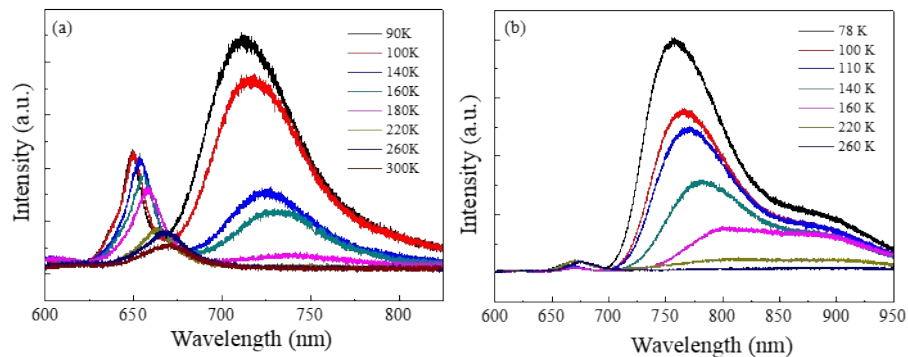


Figure S5. PL spectra of MoS₂ with (a) and without (b) NaOH

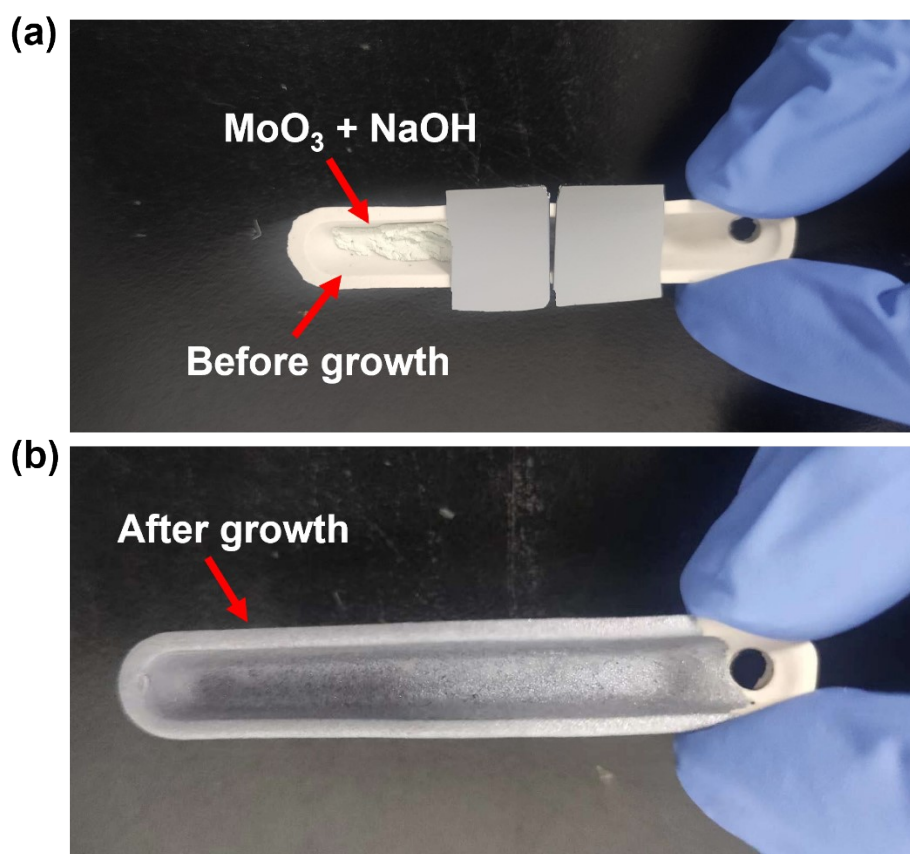


Figure S6. Influences of NaOH precursor during CVD synthesis of MoS₂ flakes, sodium clusters were observed on the alumina boat.

A TEM grid is placed on the targeted MoS₂ flakes grown on the SiO₂/Si substrate is shown in figure S7 (a). As displayed in figure S7 (b) a drop of isopropyl alcohol (IPA) is placed next to the

TEM grid. After the IPA evaporates completely, a drop of potassium hydroxide (KOH) placed next to the TEM grid is shown in figure S7 (c). After etching away, the SiO_2 , the TEM grid with MoS_2 flakes is detached from the Si substrate. The TEM grid is then rinsed in de-ionized (DI) water several times is shown in figure S7 (d).

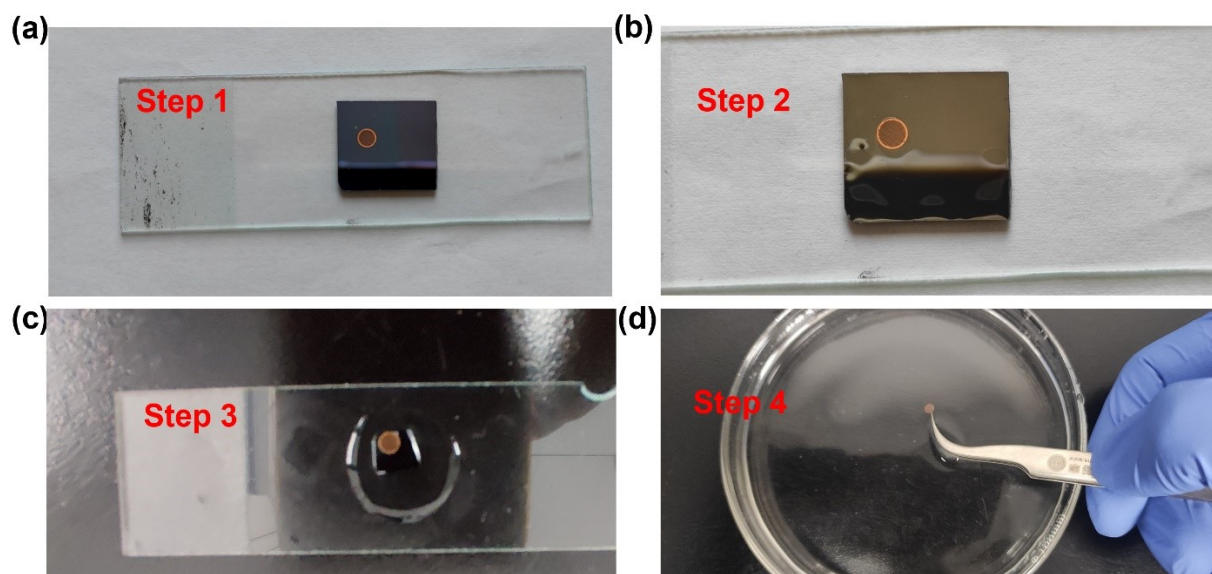


Figure S7. Transfer of CVD synthesis MoS_2 flakes to the TEM grid.

we observed the from the SEM result, when the growth temperature reduce from $800\text{ }^{\circ}\text{C}$ to $700\text{ }^{\circ}\text{C}$ the morphology of MoS_2 flakes changes from equilateral trinagle to three-point star is shown in **Figure S8**. This is possibly due to lowering the temperature the evaporation of MoO_3 reduces, resulting in a Mo:S ratio lower than the higher growth temperature under $800\text{ }^{\circ}\text{C}$, which makes the difference in the growth between Mo-zz terminations and S-zz terminations larger.

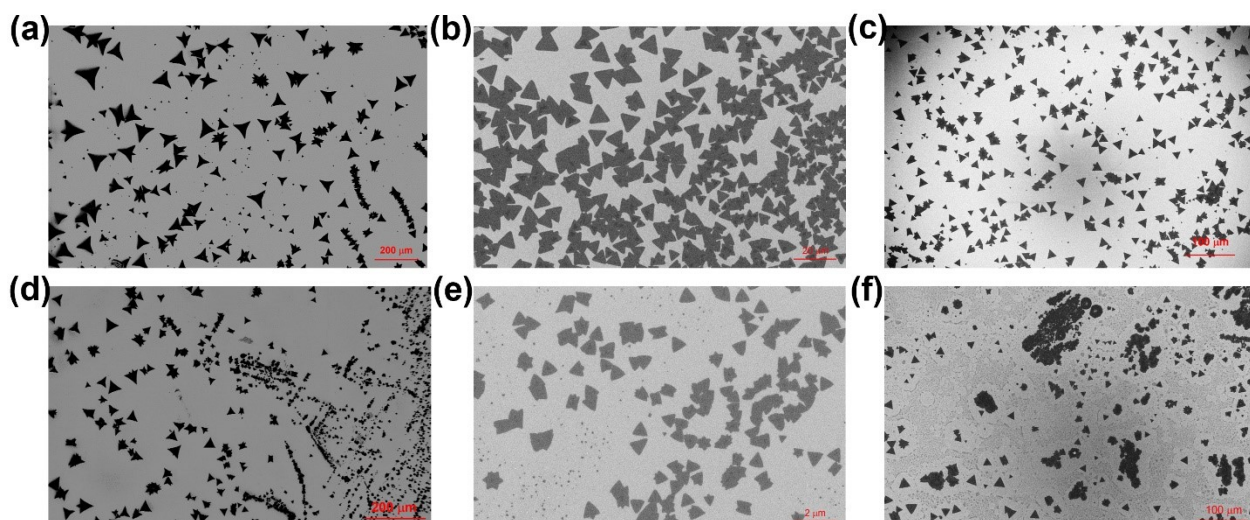


Figure S8. Scanning electron microscopy (SEM) images of MoS₂ flakes: With NaOH Chemical vapor deposition (CVD) of MoS₂ grown on SiO₂/Si at different growth condition is displayed in figure S8. (a-c) i.e., 700 °C, 750 °C and 800 °C respectively. Without NaOH CVD of MoS₂ flakes synthesis on SiO₂/Si as shown in figure S8. (d-f) at different growth temperature i.e., 700 °C, 750 °C and 800 °C respectively.

We used different amount of NaOH as shown in Table S1. The relative composition NaOH i.e., 8 mg, 12mg, 15 mg, was used with a fixed amount of MoO₃ and Sulfur (S). The details parameter is given below.

Growth parameter of MoS ₂	NaOH	MoO ₃	Sulfur	Temperature	Sccm
Low doping	8 mg	50 mg	500 mg	700-800 °C	20
Moderate doping	10 mg	50 mg	500 mg	700-800 °C	20
High doping	15 mg	50 mg	500 mg	700-800 °C	30

Table S1