Supporting Information

Title Metal-Agglomeration-Suppressed Growth of MoS₂ and MoSe₂ Films with Small Sulfur and Selenium Molecules for High Mobility Field Effect Transistor Applications

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Figure S1. Change in chamber pressure with respect to time at $T_{C-zone} = 700$ and 950 °C.



Figure S2. Mo 3d XPS spectrum of Mo film that was deposited to be 2 nm-thick using ebeam evaporation and exposed to air. The component peaks indicate that elemental Mo, oxidized Mo (Mo^{5+} and Mo^{6+}) bonding states coexist in the film. The ratio of different oxidation states of Mo was metallic Mo : Mo^{5+} : $Mo^{6+} = 35 : 52 : 13$, and the partially oxidized Mo-precursor film was approximately 2.5 nm-thick.



Figure S3. AFM images (2 μ m x 2 μ m) and rms roughness (σ_{rms}) values of MoS₂ films (T_G=500 °C) of (a) procedure A and (b) procedure B. The σ_{rms} value of a region with no protrusions (procedure A) was also compared with that of procedure B in a selected small area

(500 nm x 500 nm).



Figure S4. (a) Photograph of a high quality MoS_2 film on a 6 inch SiO_2/Si wafer. Height profiles of the MoS_2 film at points (b) B, (c) C, (d) D, and (e) E.



Figure S5. XPS peaks of (a) Mo and (b) S in a 6 nm-thick MoS_2 film fabricated at $T_{R-zone} = 120$ °C.



Figure S6. (a) A cross-sectional TEM image of a 3.1 nm-thick MoS₂ film. EDS maps of (b) Mo and (c) S in a 3.1 nm-thick MoS₂ film.



Figure S7. Histogram of field-effect mobility of MoS₂-FETs for 20 devices fabricated on a substrate.