

Fabrication of a Low-Cost CuInS_2 Solar Cell by Ink Processing

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Outline

- Objective

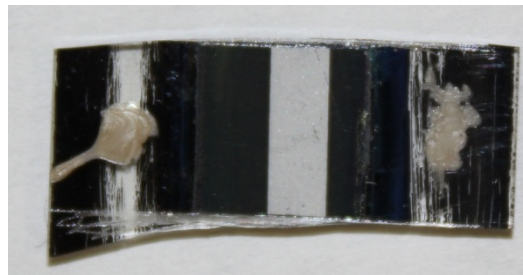
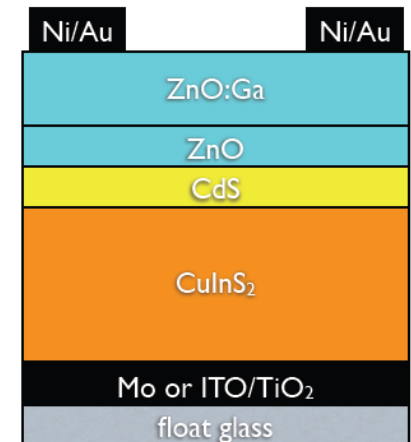
Develop scalable fabrication steps to make CuInS_2 (CIS) photovoltaic cells using solution-processable inks.

- Project Progress

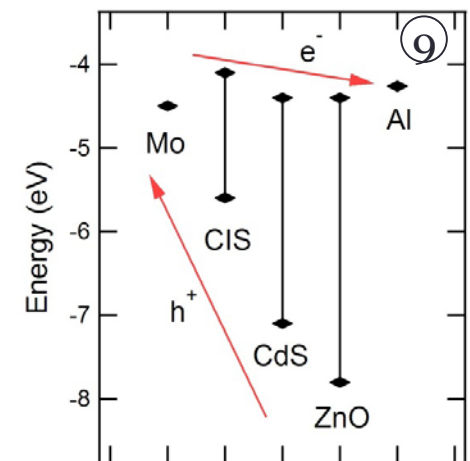
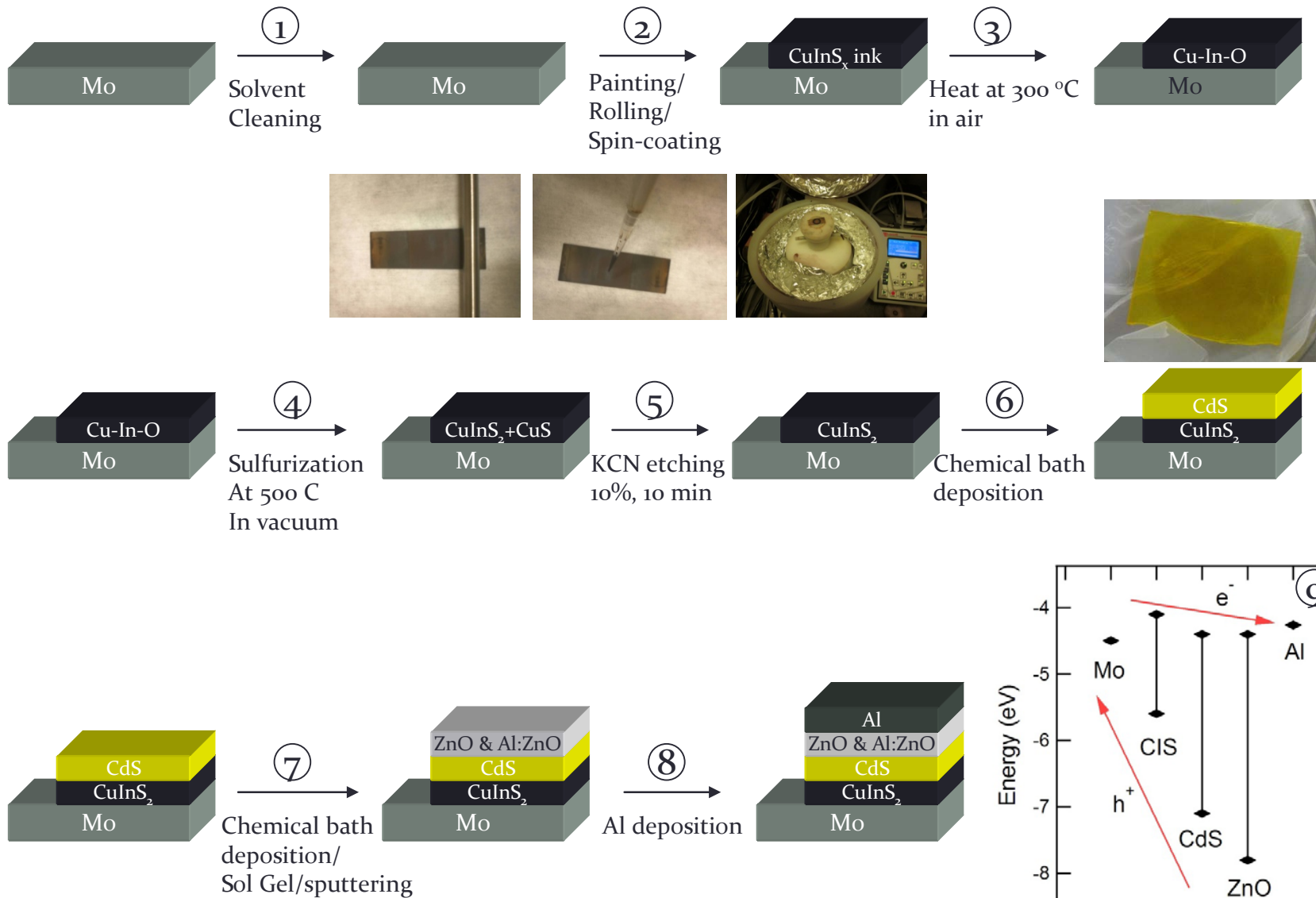
1. Solution-based deposition and characterization of CIS, CdS, ZnO, Ga-doped ZnO
2. Integration of layers into a working cell
3. Control of fabrication steps

- Initial Results

IV under 1.5 air mass yielded $V_{oc} = 0.07 \text{ V}$ and $J_{sc} = 4.3 \text{ mA/cm}^2$, with a fill factor of 0.26 and efficiency of 0.1% (0.15 cm^2 area).



Fabrication Process



Surface Composition of CuInS_2 Layer



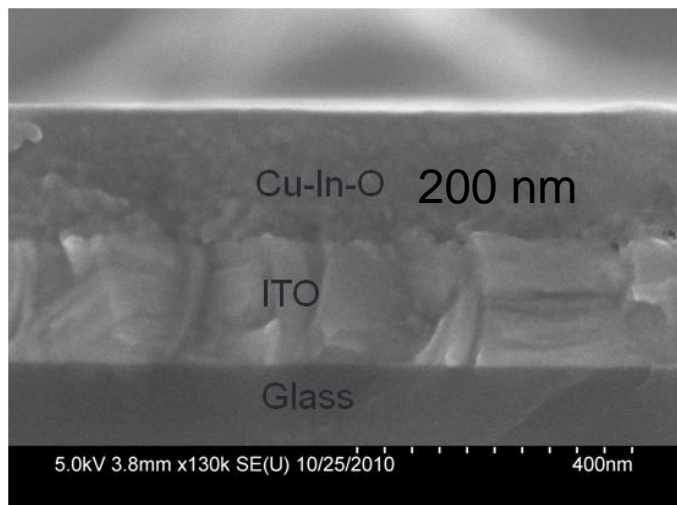
Cu:In ratio (from left to right): 1:2, 1:1.5, 1:1, 1.5:1, 2:1
at 400 °C oxidation temperature



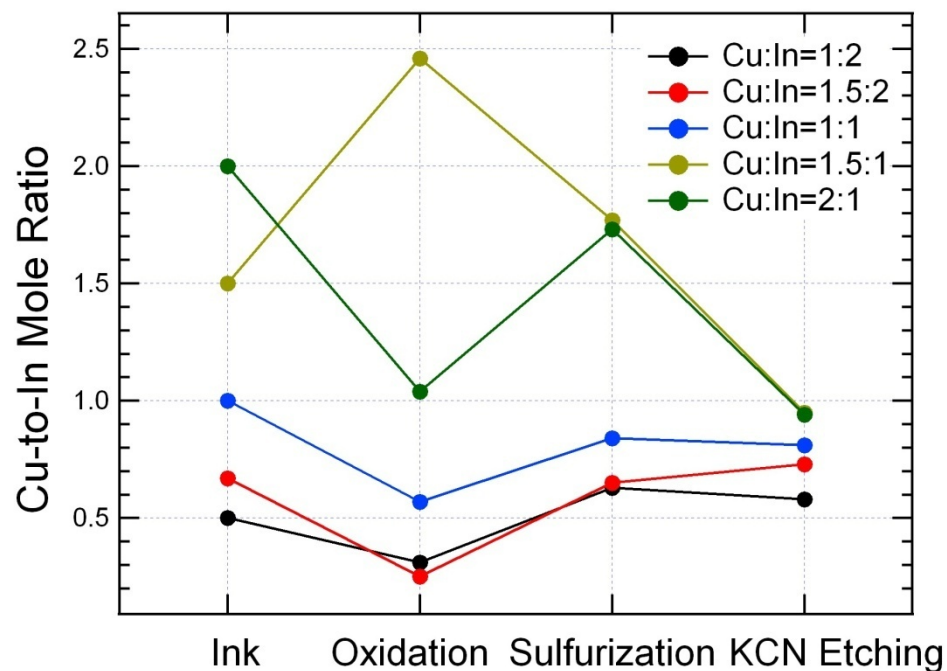
Cu:In ratio (from left to right): 1:2, 1:1.5, 1:1, 1.5:1, 2:1
at 500 °C sulfurization temperature for 1 hour



Cu:In ratio (from left to right): 1:2, 1:1.5, 1:1, 1.5:1, 2:1
10 wt% KCN etching 5 min for each

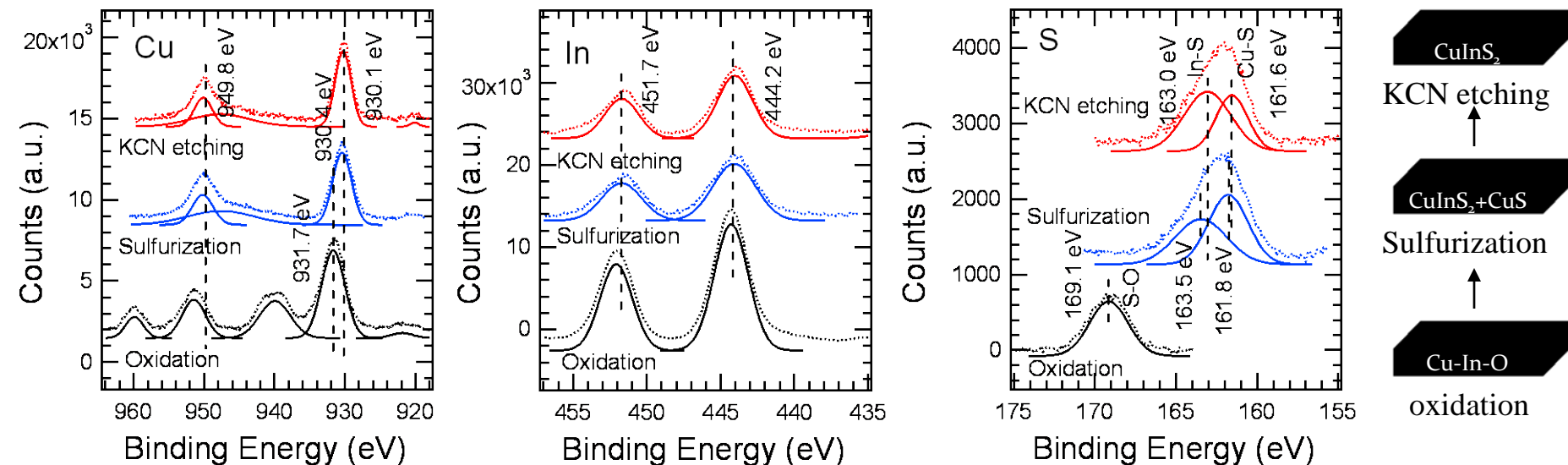


SEM image of Cu-In-O layer deposited by spin coating



Cu/In mole ratio at different steps. Ratio in ink varied from 1:2 to 2:1. After oxidation, Cu/In ratio decreased comparing to the initial ratio in ink for all of the samples, except the 1.5:1 one, suggesting that surface is In rich. After sulfurization, Cu/In ratio increased again, showing that Cu merged to the surface. After KCN etching, excess CuS was removed and final stoichiometry was reached for the samples with initial Cu/In ratio larger than 1.

Surface Composition of CuInS₂ Layer

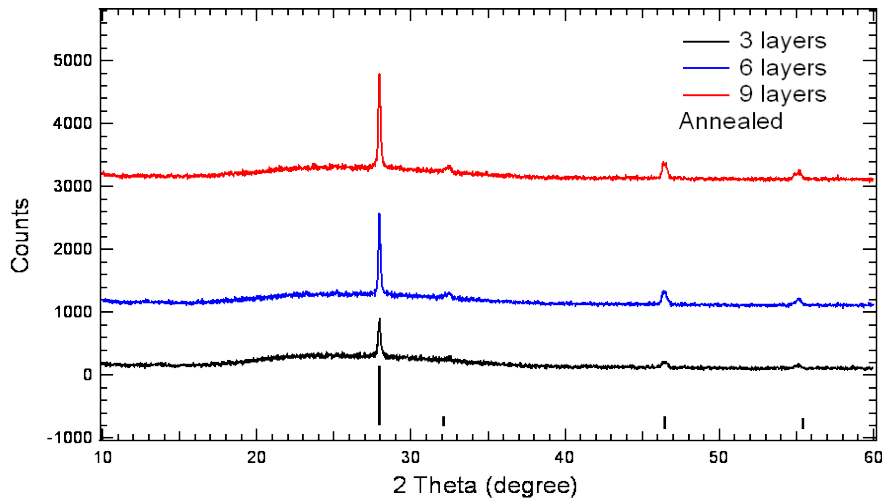


Cu 2p_{3/2} shifted from 931.7 eV after oxidation to 930.4 eV after sulfurization, and further shifted to 930.1 eV, indicating the transition from CuO to CuInS₂ with CuS to CuInS₂.

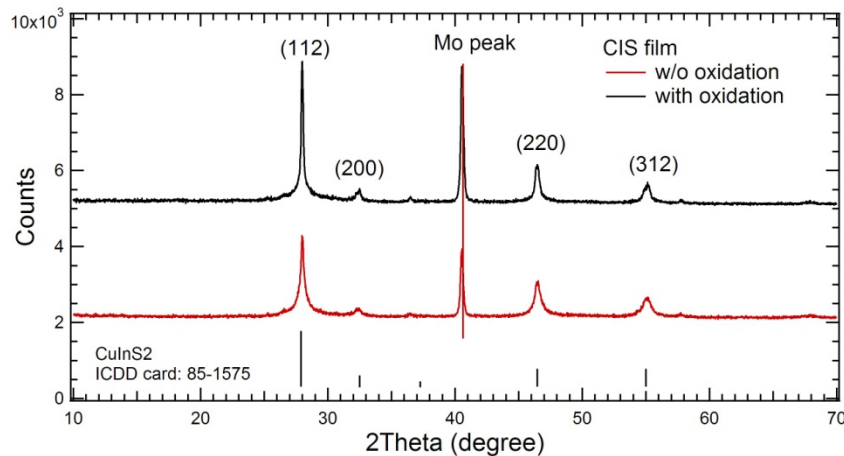
In 2p_{3/2} shifted 0.2 eV to the lower binding energy when changing from oxides to sulfides, then stayed at 444.2 eV, corresponding to In in CuInS₂.

After oxidation, a peak corresponding to sulfate was shown at 169.1 eV. After sulfurization, a broad peak for metal sulfides appeared from 161 to 164 eV. The de-convolution of this peak shows that KCN etching did remove extra CuS phase formed on the surface.

Crystal Structure



CIS layer made by spin-coating
Substrate=glass

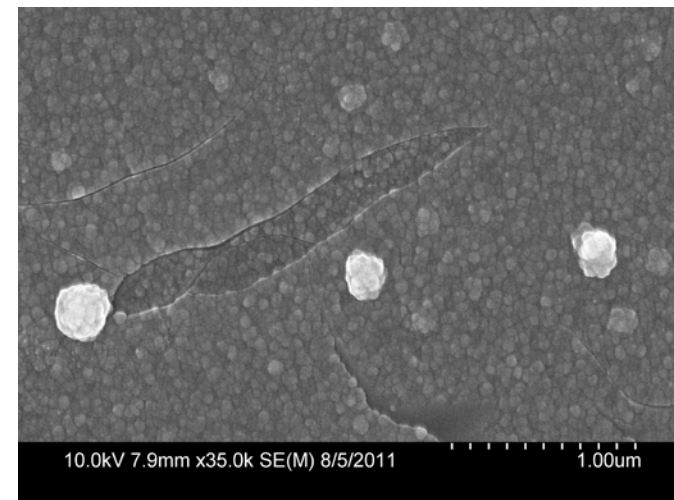
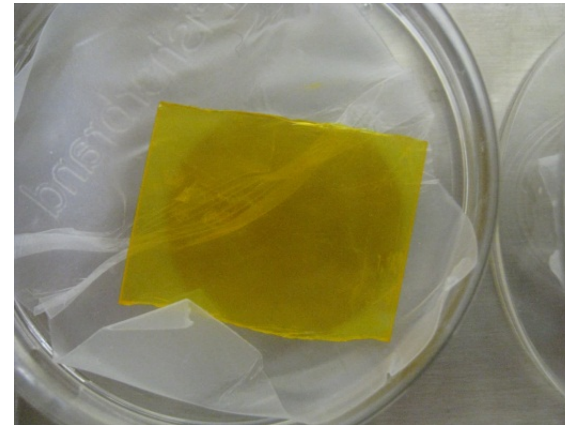


CIS layer made by painting
Substrate=Mo coated glass

CdS Deposition Using CBD

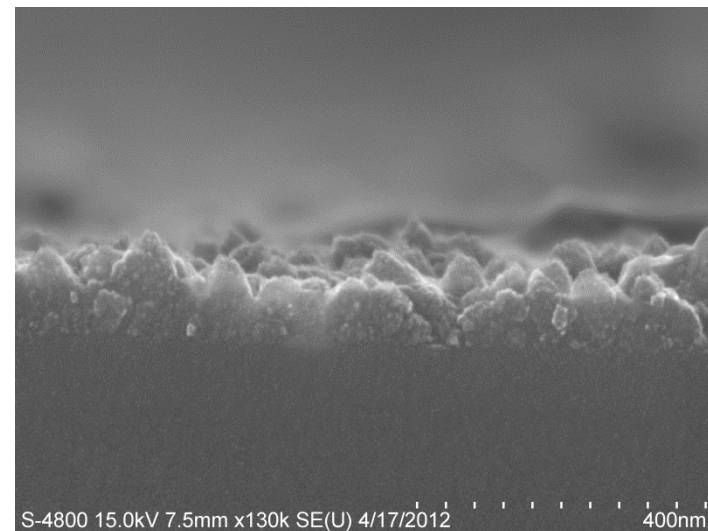
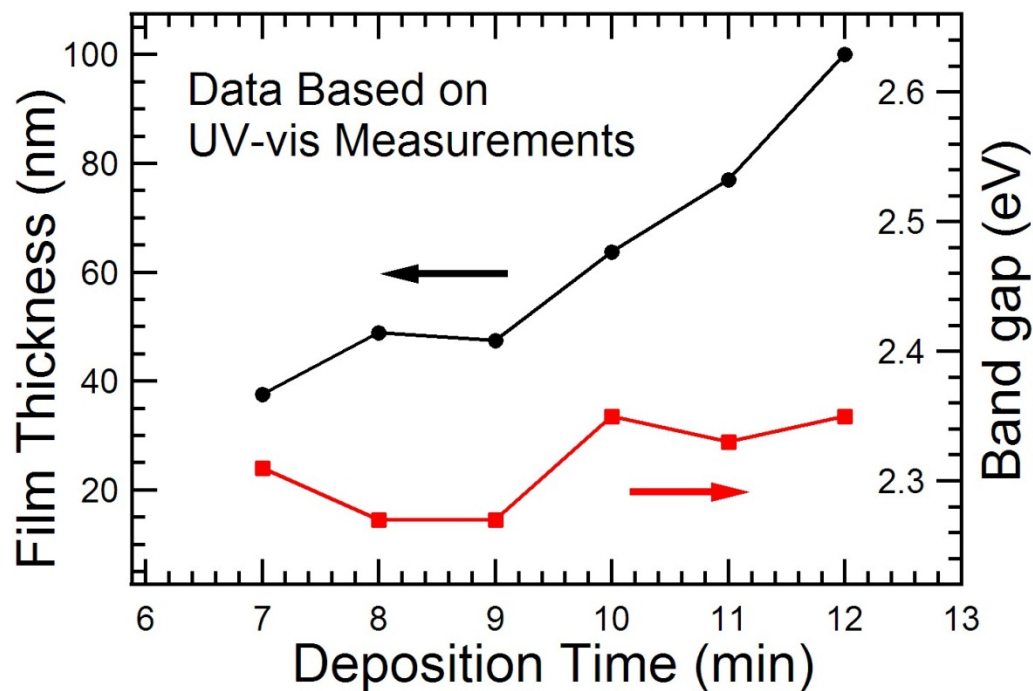


The Chemical Bath Deposition Setup

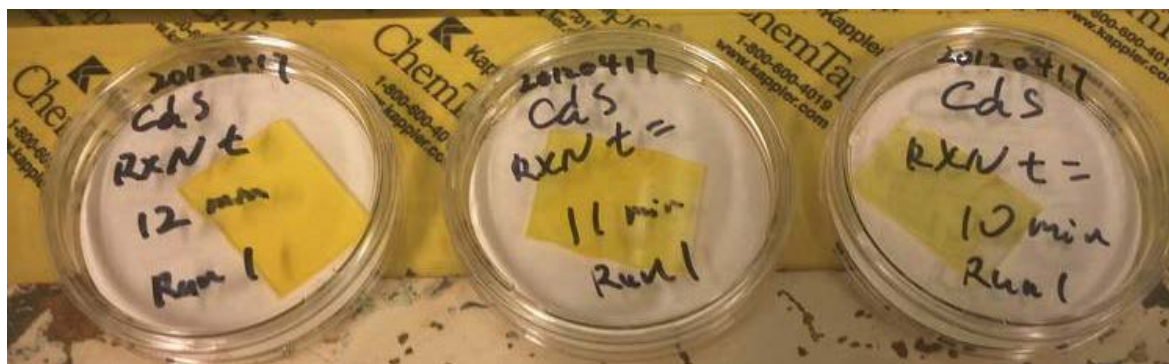


Small grains with an average size of 50 nm were obtained, as shown in the SEM image.

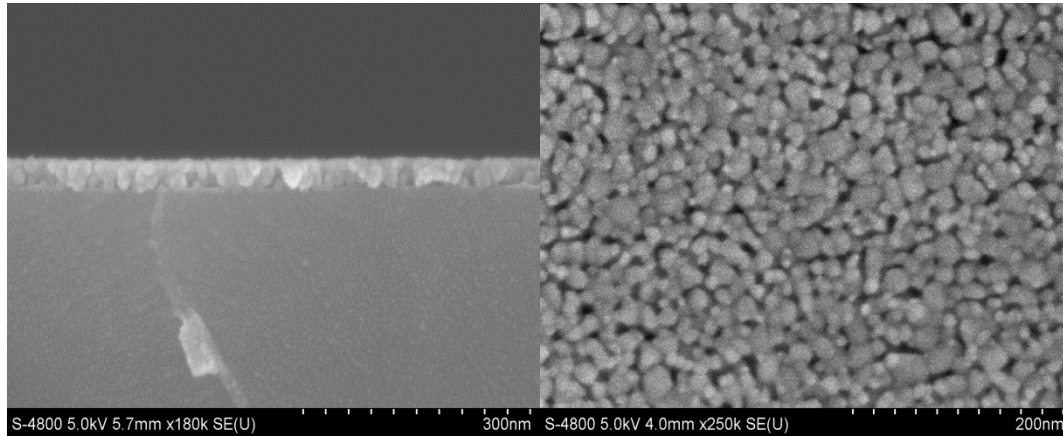
Thickness/Band gap vs. Deposition Time



SEM image of CdS film with deposition time of 11 min. Film thickness is about 80 nm

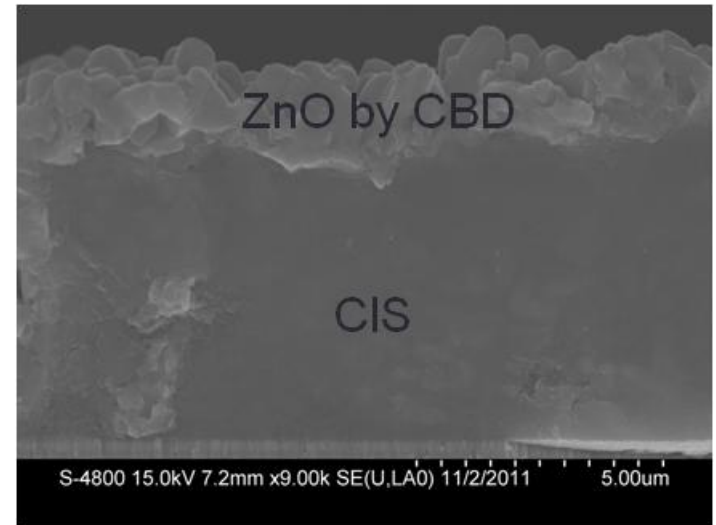
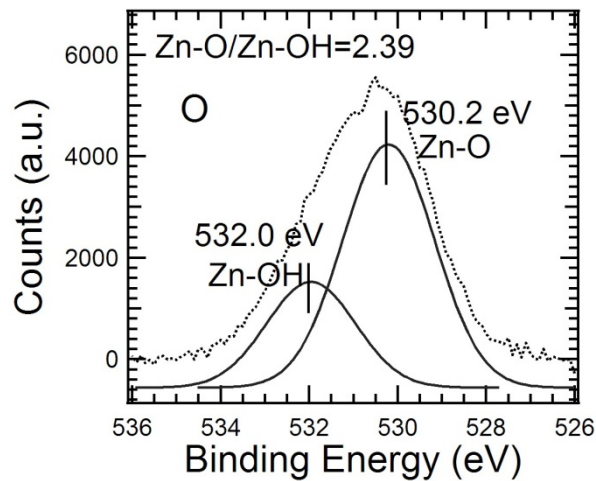


ZnO by Sol-Gel and CBD

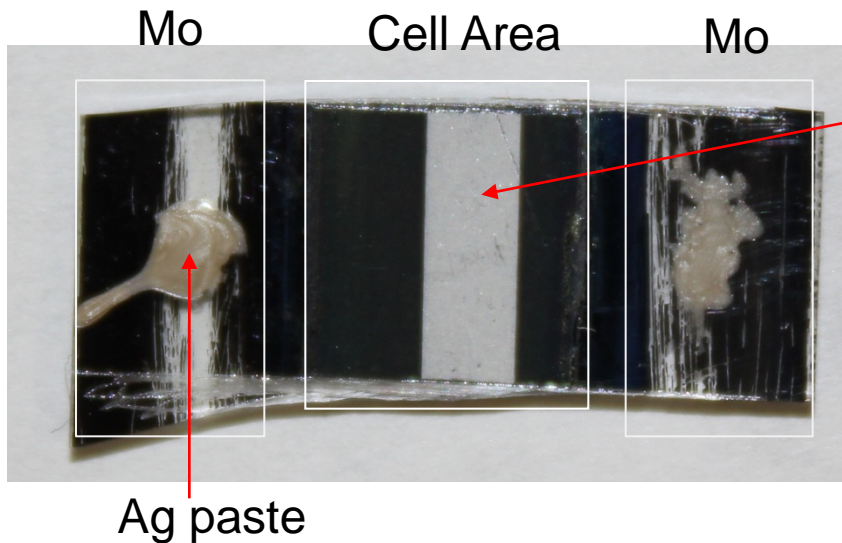


Cross-section

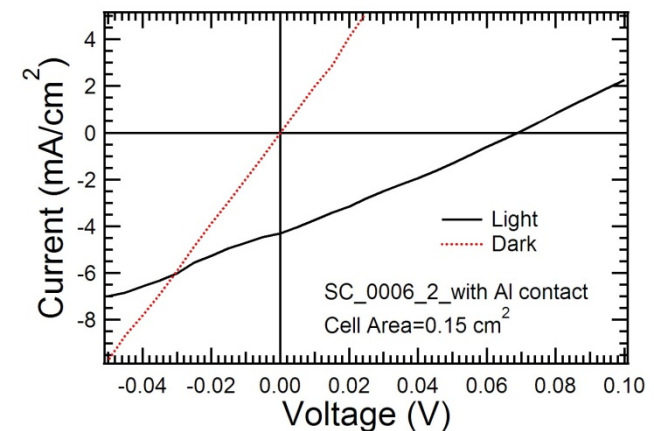
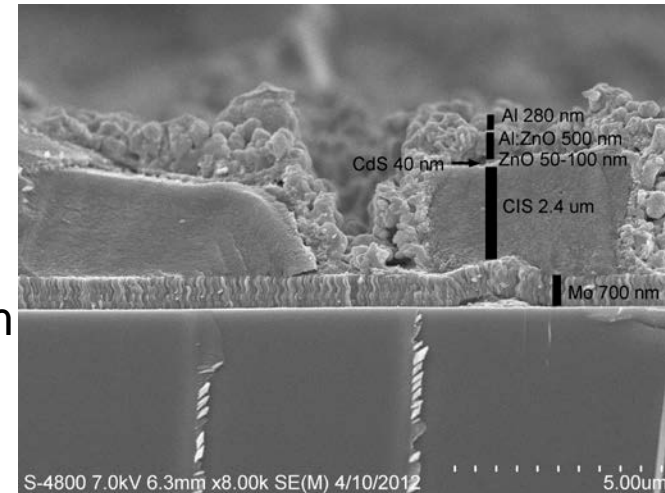
Topography



The First Working Cell



Al: 280 nm
 Al:ZnO: 500 nm
 ZnO: 50 nm
 CdS: 40 nm
 CIS: 2400 nm
 Mo: 700 nm



Sample dimension: 1.5 by 0.5 cm
 Cell dimension: 0.5 by 0.5 cm
 Al contact 0.2 by 0.5 cm
 Cell Area: $0.25 \times 0.10 = 0.15 \text{ cm}^2$

Conclusion

- Demonstrated solution-based deposition of CuInS_2 , CdS , and ZnO
- Process integration highlighted the need for good control of layer thickness
- First working cell with efficiency of 0.1% provides a path to improve the performance of future devices

Acknowledgements

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