

## Effect of long selenization time on Co films under a low temperature of 300 °C to synthesize a nanostructure CoSe<sub>2</sub> and optical properties

Po-Feng Wu<sup>\*</sup>, Jen-Bin Shi

Department of Electronic Engineering, Feng Chia University, No.100, Wenhwa Rd., Seatwen, Taichung 40724, Taiwan

Accepted for publication on 25<sup>th</sup> March 2015

This work investigates a simple method to transform pre-deposited amorphous Co film into  $CoSe_2$  films at a fixed, low temperature of 300 °C. Single  $CoSe_2$ -phase films having good crystallinity were obtained at a selenisation time  $\geq 24$  hours. A nanostructure  $CoSe_2$  having two different nano-morphological layers was observed. The  $CoSe_2$  films (72 hours) observed a large absorption and a direct band gap.

Transition metal chalcogenides/dichalcogenides  $MX_2$  (M = Mn, Fe, , Ni, Co; X = S, Se, Te) has been investigated on their optical, magnetic, and electrical properties. Many theoretical and experimental reports have been done on the pyrites  $MX_2$ . Due to the application in, rechargeable battery electrodes and their potential applications in such areas as spintronic devices pyrites  $MX_2$ . CoSe<sub>2</sub> and CoSe have received increasing appeal as potential cathodic catalysts for fuel cells. Fuel cells can directly reduce our dependence on oil, energy use, and harmful emissions, which are used for stationary, portable power, and transportation. In the other hand, the oxygen reduction reaction (ORR), used in an acid medium Pt-based material, is the best and most used catalyst. However, Pt materials are too expensive, while Fe and Co metals are much cheaper and serve equally as potential catalytic centers towards ORR in the creation of solar cell material. The cathodic oxygen reduction in chalcogenides of various transition metals has been further reported.

Key words: Nanostructure, Selenisation, Transition metal dichalcogenides