Structural and morphological analysis of hydrothermally fabricated ternary palladium alloys for use as efficient catalysts in dye sensitized solar cell counter electrodes

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### Outline



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- Aims & Objectives
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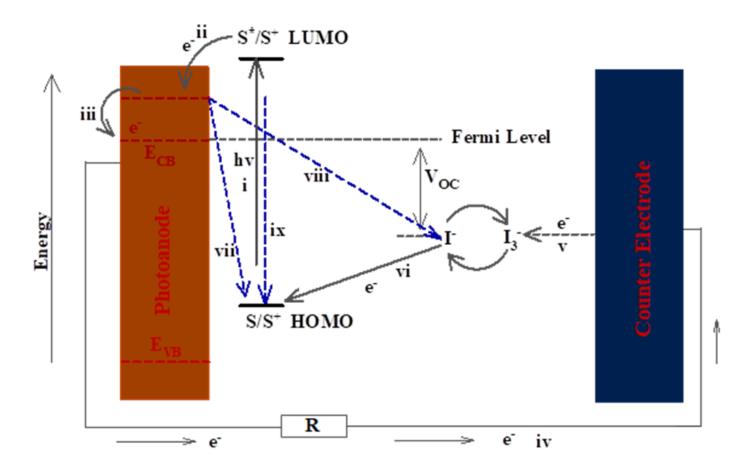




- Greater demand for more energy necessitated by the rising world population and high global economic development[1].
- The need to limit the emission of toxic materials into the atmosphere
- Geopolitical instability causes fluctuation of oil prices
- Turn to renewable energy sources to fill the void created by depletion of fossil fuels as well as to fulfil the stringent environmental regulations enacted by various governments
- Potential alternatives include the DSSC technology which was invented by Gratzel in 1991 mimicking the conversion of sunlight into energy by plants [2].







#### Fig 1: Operational procedure for a DSSC



#### **Advantages** Produce energy from DSSC Disadvantages cleaner sources. **Reduce** emissions High fabrication costs Possible improvements **Reduce deforestation** associated with the Simple operating expensive ruthenium and Elimination of liquid procedure platinum metals. electrolyte. Replace ruthenium dye Low conversion efficiency with other inorganic dyes 13.4% Replace platinum counter electrode with core shell alloys whose efficiency is higher than that of platinum.





• Pure transition metal elements that have been explored up to date have only given low efficiencies leading to development of higher multi-metallic systems

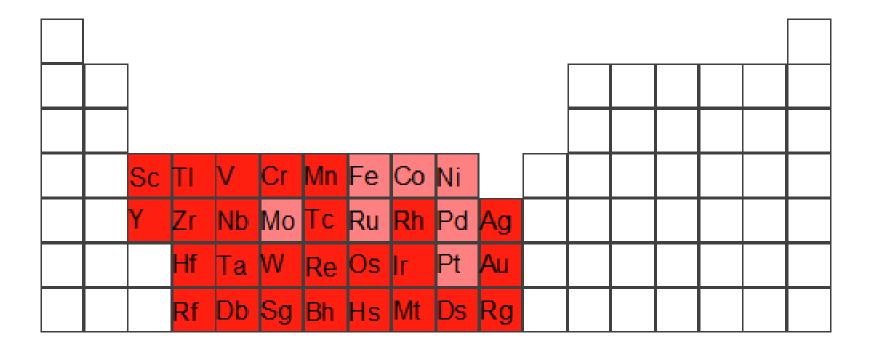


Fig 2:Periodic Table of elements





- Use of ternary transition metal composites or alloys where each component serves a particular function. The more reactive metal would ideally occupy the outer part so as to partake in interfacial reactions.
- Ternary alloys possess excellent catalytic qualities which are due to the synergistic effect produced by the existence of three elements in a crystal structure.
- Synergy created by the interaction of two metals results in electron transfer which increases surface interaction with the electrolyte[3].



# Aims and objectives



#### Aim

- Synthesize ternary palladium alloy CE for DSSC use.
  Objectives
- To synthesise PdNiCo, PdNiZn, ternary alloys
- Determine the structure and morphology of the synthesised counter electrodes using XRD and SEM respectively
- Evaluate the electrochemical properties of the developed catalyst samples using CV EIS and CD
- Determine the effect of varying the composition of the alloy on the performance of the counter electrode.



# **Methodology**



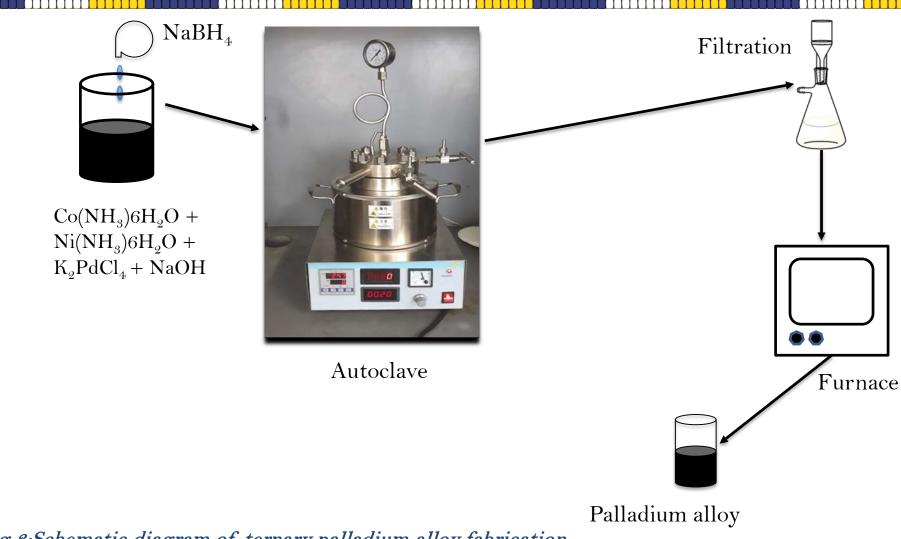


Fig 3:Schematic diagram of ternary palladium alloy fabrication

#### **Results:XRD**



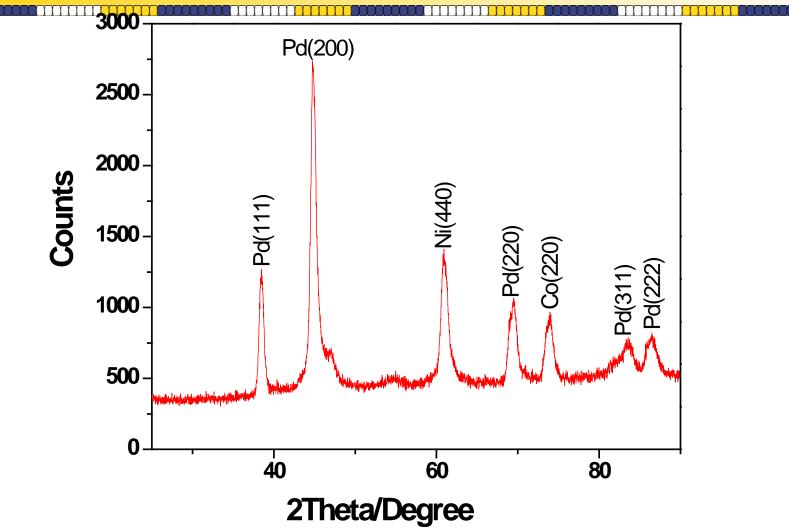


Figure 4: XRD image of PdNiCo

### **Results:XRD**



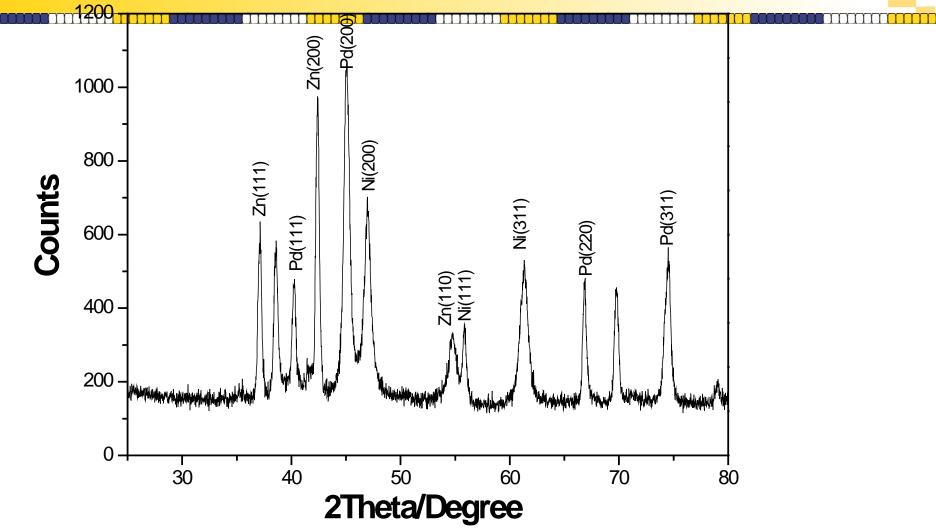


Figure 5: XRD image of PdNiZn

#### **Results: SEM**



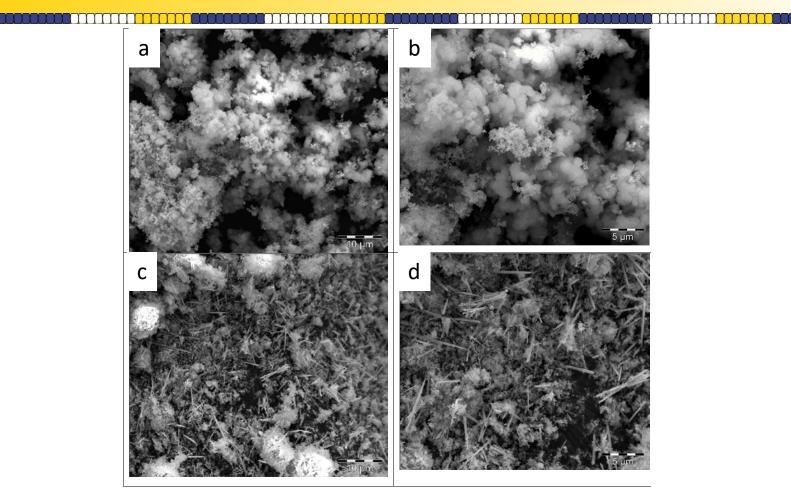
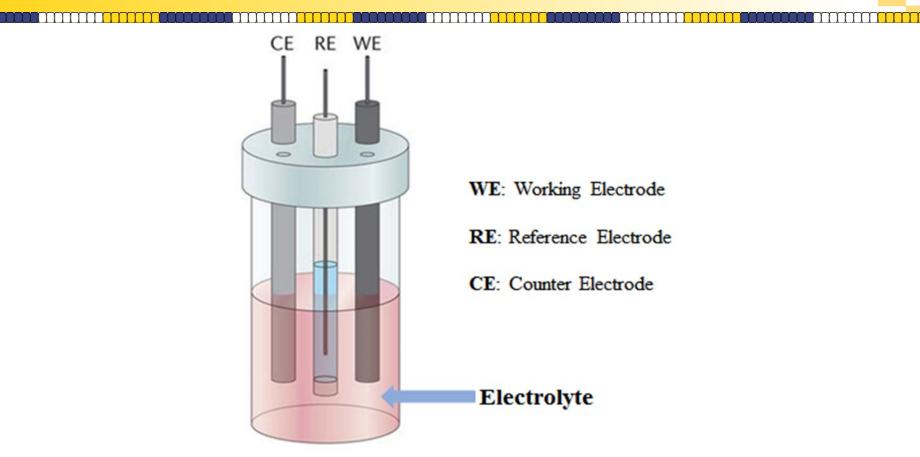


Figure 6: SEM images of (a and b) low and high magnification of PdNiCo respectively (c) and (d)PdNiZn



# Results





#### Fig:7 Three electrode cell configuration



#### **Results:** CV



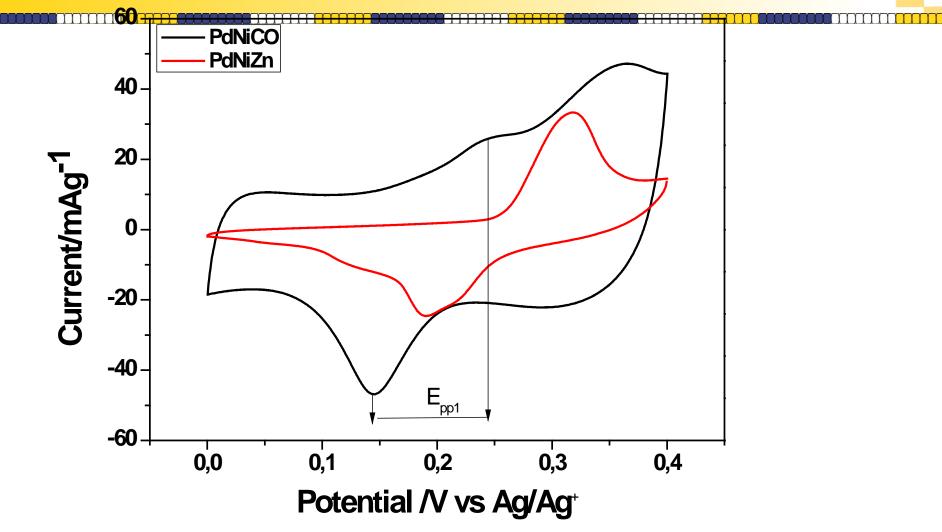


Fig 8: Cyclic Voltammetry graphs for ternary palladium alloys at 5omVs







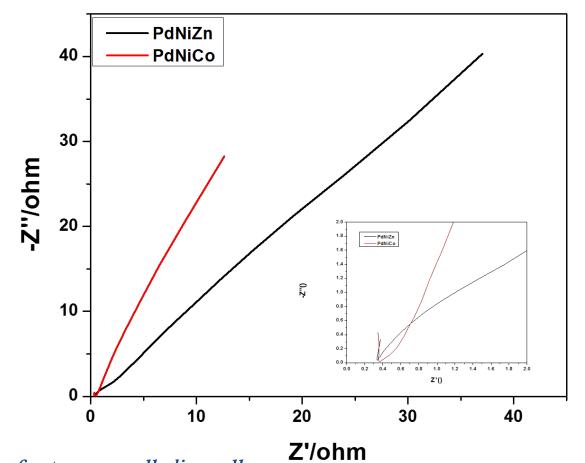


Fig 9: EIS graphs for ternary palladium alloys

#### **Results** :CD



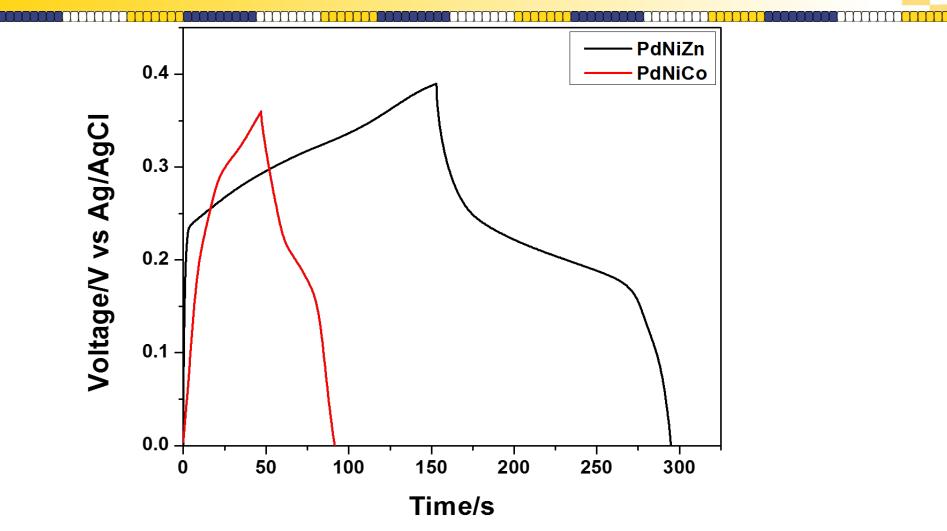


Fig 10: CD curves for PdNiZn and PdNiCo

### **Conclusion**



- XRD has shown 5 dominant peaks at 2 theta values of 40°, 46.2°, 63.2,°
  71.2°,75.1°,83.3° and 86.1° for PdNiCo whereas PdNiZn had peaks at 43.2°
  , 46.4°,45.6°, 62.3°, 67.4° and 75.2°.
- SEM identified spherical densely packed PdNiCo particles whereas PdNiZn was composed of nanoneedles.
- CV results showed that PdNiZn had more reduction current density however it also possessed a higher peak to peak potential difference signifying a lower rate of reduction intensity.
- From EIS analysis PdNiZn possessed the least charge transfer resistance at 0.320hm with PdNiCo not performing any worse at 0.36 ohm.
- Charge discharge graphs show that PdNiZn slowly discharges electrical energy hence it is a potential material for use in capacitors.



### References



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# **Acknowledgment**







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