

## Seminar

#### Exploring Atomic Layers based Hybrid 2D Materials for Photocatalysis and Photodetection

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Two dimensional (2D) layered semiconductor atomic layers, such as transition metal dichalcogenides (MX<sub>2</sub>, where M=Mo, W etc, and X=S, Se, Te etc.) and their van der Waals (vdW) heterostructures have attracted considerable interest. Heterostructures with atomically sharp interfaces and alloyed versions of these exhibit widely tunable band gaps as a function of composition and stacking sequence. In this talk, I will demonstrate aberration corrected high angle annular dark field (HAADF) - scanning transmission electron microscopy (STEM) of MoS<sub>2</sub>, WS<sub>2</sub> and their alloys with selenium 'Se' such as  $MoS_{2(1-x)}Se_{2x}$  etc and their enhanced optical properties for visible light photoelectrocatalysis (PEC). Further, the recent achievements in fabrication and characterization of atom-by-atom 'Se' doping of MoS<sub>2</sub> and WS<sub>2</sub> atomic layers and seeding of platinum 147 (Pt<sub>147</sub>) size selected nanoclusters on WS<sub>2</sub> atomic layers using cluster beam deposition. Pt<sub>147</sub> atomic clusters were deposited at 0.1 eV per atom kinetic energy with and without Ar ion defects on WS<sub>2</sub> atomic layers for the first time. The statistical analysis and morphological features of size distribution and defect mediated seeding of Pt147 clusters are quantified using HAADF-STEM and complimented with Raman/photoluminescence (PL). Layer dependency and defect mediated interaction of Pt<sub>147</sub> clusters on WS<sub>2</sub> is quantitatively estimated from HAADF–STEM intensity profiles. Furthermore, electrochemical Hydrogen Evolution Reaction Activity (HER) of WS<sub>2</sub> and WS<sub>2</sub>-Pt<sub>147</sub> clusters before and after electrochemical activity will be discussed. These kind of 2D and 0D nanocluster hybrids are very good candidates for the catalysis and energy harvesting applications.

# Friday, Oct 25<sup>th</sup> 2019 11:30 AM (Tea/Coffee at 11:00 AM) Seminar Hall, TIFR-H