

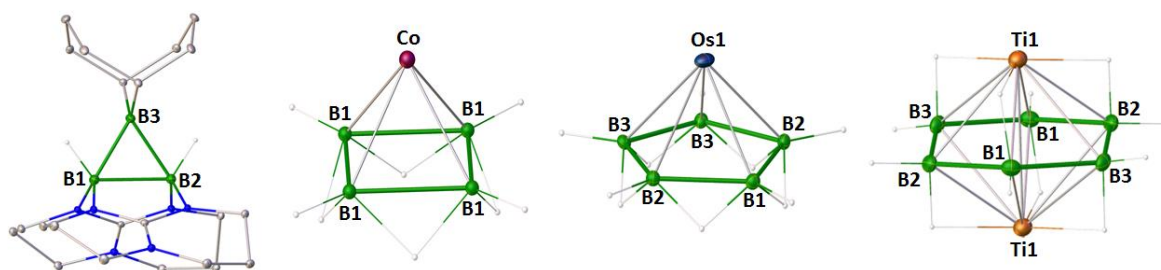
## Seminar

### Flat Boron Rings Stabilised by Transition Metal Templates

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Unlike carbon, boron does not usually form ring compounds due to its electron deficiency-driven affinity towards polyhedral geometries. Therefore, the planar boron rings were ignored for a long time. However, recent developments by us and others demand further attention in stabilising planar boron rings. For example, Himmel synthesised a derivative of a doubly base-stabilised  $[B_3H_6]^+$ , the structural equivalent of  $[B_3H_8]^-$  and  $[B_3H_9]$ . Although the smaller  $B_3$  and  $B_4$ -rings are mostly isolated as individual species, larger  $B_5$  and  $B_6$ -rings are stabilised by TMs, depending on the ring size and electronic requirements of the rings. In this regard, we have recently synthesised and structurally characterised tetraborane ring  $[B_4H_8]^{2-}$  and pentaborane ring  $[B_5H_{10}]^-$  stabilised by Os-template in  $\eta^4$  or  $\eta^5$ -fashion which mimics  $[C_4H_4]^{2-}$  and  $[C_5H_5]^-$  rings, respectively. We have also isolated the first examples of a nearly planar  $[B_6H_6]$  unit, stabilised as a part of  $[(Cp^*Ti)_2(\mu-\eta^6:\eta^6-B_6H_6)(\mu-H)_6]$ , where significant electron delocalisation was observed from the Ti-Ti bonding orbital to the  $\mu$ -H atoms and  $B_6$  skeleton. The selection of transition metals mostly depends on their diffuse orbitals, valence electrons, and ring sizes, while the bridging hydrogens balance the electron deficiency of the borane rings.



**Friday, Sep 20<sup>th</sup> 2024**

**11:30 Hrs (Tea / Coffee 11:15 Hrs)**

**Auditorium, TIFR-H**