# Improved surface and interface properties of InP

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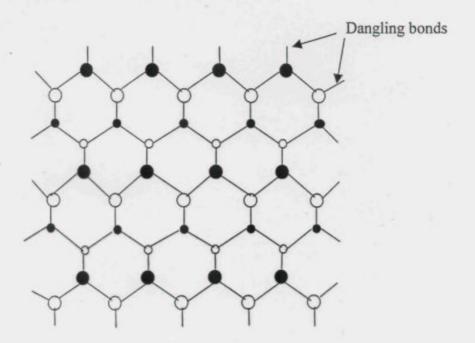


Figure 1.1 Zinc-blende lattice projected on (110) face showing the dangling bonds on the surface

#### Problem addressed:

- Surface Fermi-level pinning is a very serious problem in indium phosphide.
- The Fermi-level pinning limits the rapid development of InP based devices.

# Possible solution for the problem:

To reduce the surface states, thereby unpin the Fermi-level in indium phosphide (InP) by proper surface preparation and effective passivation schemes

## Our approach:

#### Surface polishing:

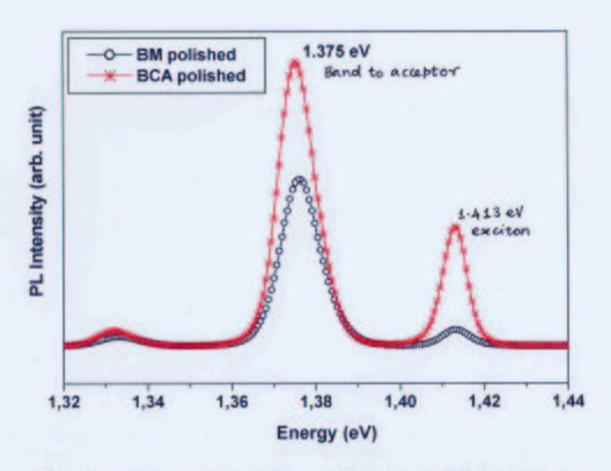
➤ A new polishing solution, HBr:K2Cr2O7:H2O (BCA) has been developed and it has been compared with conventional brominemethanol (BM) solution.

#### Surface passivation:

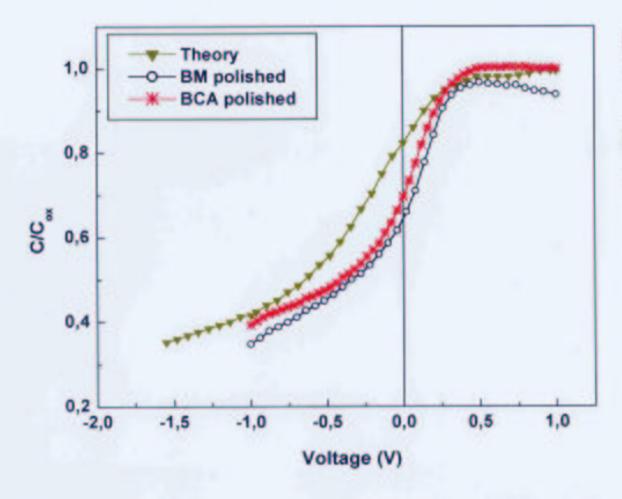
Cadmium sulfide (CdS) has been identified as a better passivating agent for InP and CdS passivation has been carried out using simple chemical bath deposition (CBD) method.

## CdS passivation mechanism:

- As the bonding in InP is mainly covalent, there will be a large perturbation of the valence-state energy at the surface
- But constituents of II-VI compounds have a larger electro negativity difference, indicating that the bonds are ionic in nature
- the electron wave functions of the ions do not overlap sufficiently to induce midgap states
- Thus insulator/II-VI interface should behave more ideally than insulator/III-V counterpart



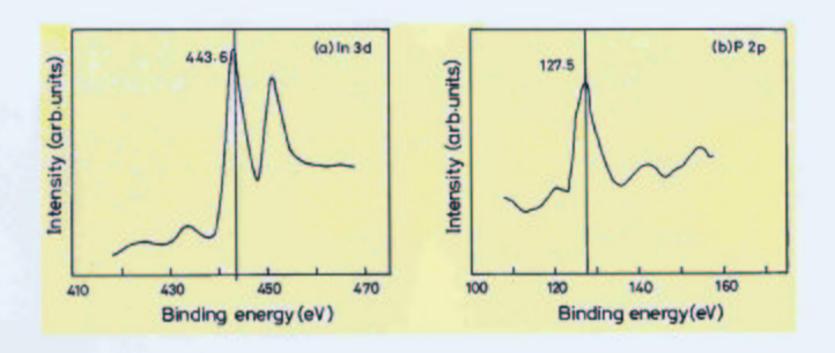
LT PL spectra of (a) BCA and BM polished InP surface



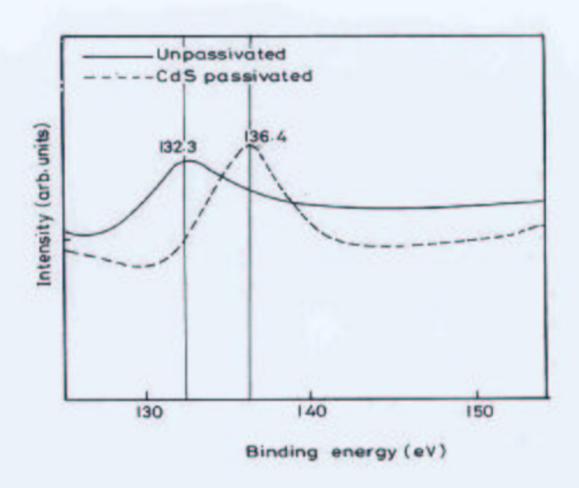
For BM Polished Sample, Nss = 9 x 10 cm 2 ev 1

For BCA Polished Sample,
NSS = 6 × 100 cm² ev²

Theoretical and experimental C-V curves for N<sub>ss</sub> calculation



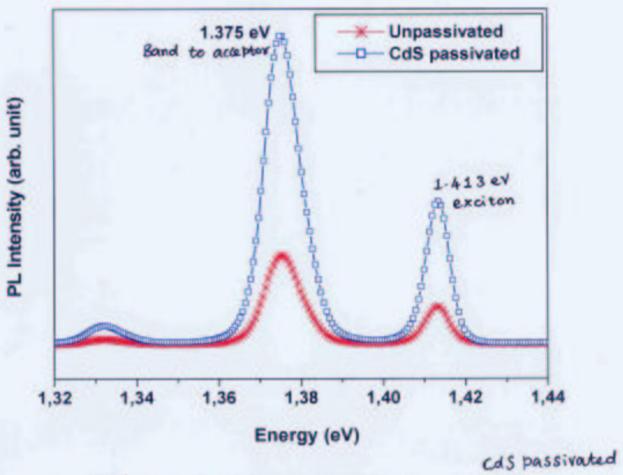
XPS spectra of CdS passivated InP showing the absence of native oxides



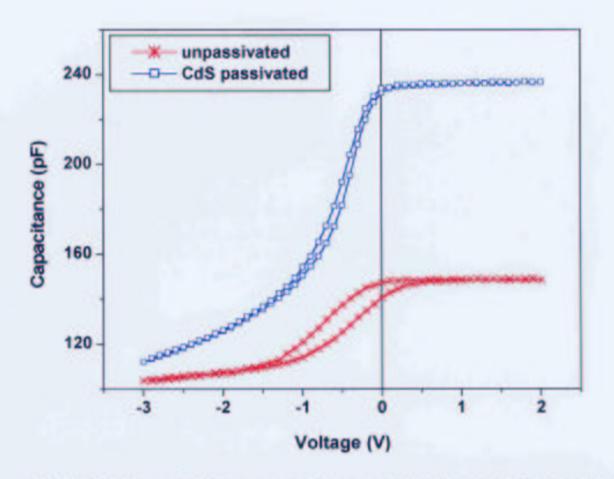
XPS spectra of P 2p peak of unpassivated and CdS passivated InP

# Possible mechanism for formation of $P_2O_5$ :

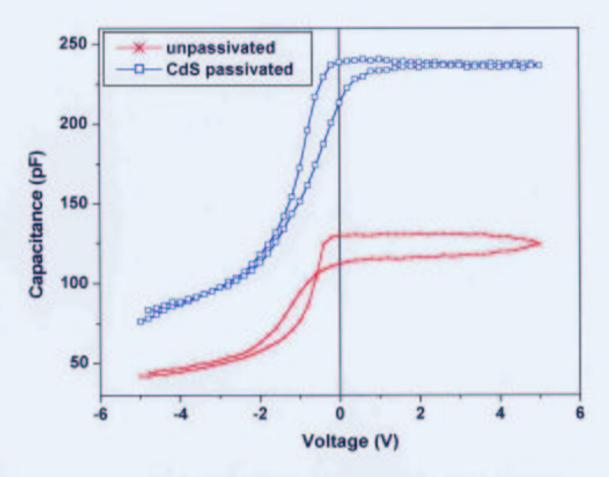
- During anodic oxidation of InP, both indium oxide (In<sub>2</sub>O<sub>3</sub>) and phosphorus oxide (P<sub>2</sub>O<sub>5</sub>) can be formed. In<sub>2</sub>O<sub>3</sub> is conductive oxide, whereas P<sub>2</sub>O<sub>5</sub> is a high resistive oxide.
- After passivation, the sulfur atoms in the CdS substitutes the oxygen atoms present in In<sub>2</sub>O<sub>3</sub> and thereby forming In<sub>2</sub>S<sub>3</sub>.
- Thus the formation of In<sub>2</sub>O<sub>3</sub> is suppressed and further this In<sub>2</sub>S<sub>3</sub> is dissolvable in the electrolyte. Hence P<sub>2</sub>O<sub>5</sub> formation is enhanced and formation of high resistive layer is accomplished



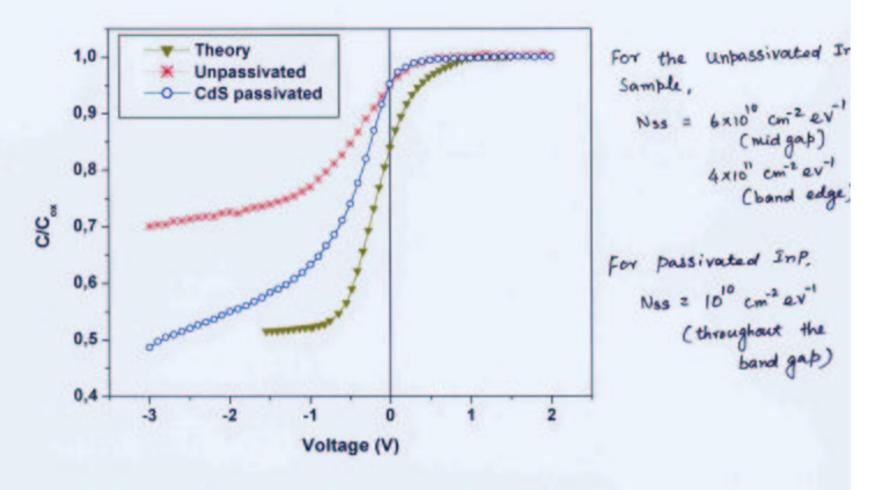
Enhanced PL intensity for BCA polished InP



1 MHz C-V curves for unpassivated and CdS passivated MIS diodes



C-V curve of the passivated sample shows the high stability even in higher bias values



C-V curve of the passivated sample follows very good with the theoretical curve, unlike unpassivated sample

#### Conclusions:

- The surface states have been reduced, thereby reducing the surface Fermi-level pinning in InP by proper surface polishing and effective surface passivation processes
- A new polishing solution (BCA) has been established and it acts as a better self-passivating agent to reduce the surface states (N<sub>SS</sub> = 6 x 10<sup>10</sup> cm<sup>-2</sup> eV<sup>-1</sup>)
- By CdS passivation the surface states have been further reduced (N<sub>SS</sub> = 10<sup>10</sup> cm<sup>-2</sup> eV<sup>-1</sup>), thus reducing the Fermi-level pinning.
- The new polishing and passivation mechanism has been analysed and understood. The methods employed in this work are simple and very cost effective