

Lecture 7 - Epitaxy Techniques and Considerations - Outline □

- **Lattice-matching considerations**

 - Natural lattice-matching**

 - 1. Review of lattice-matched material systems (Lect. 1 discussion)
 - 2. Lattice pulling

 - Forced lattice matching** □

 - 1. Pseudomorphic layers
 - 2. Matthews-Blakeslee limit, other models □
 - 3. Examples - devices using pseudomorphic layers □

 - Mismatched epitaxy** □

 - 1. Step-grading
 - 2. Linear grading
 - 3. Examples - devices using graded heterostructures

- **Epitaxy techniques - overview** (survey of commonly used techniques)

 - Liquid phase epitaxy** □

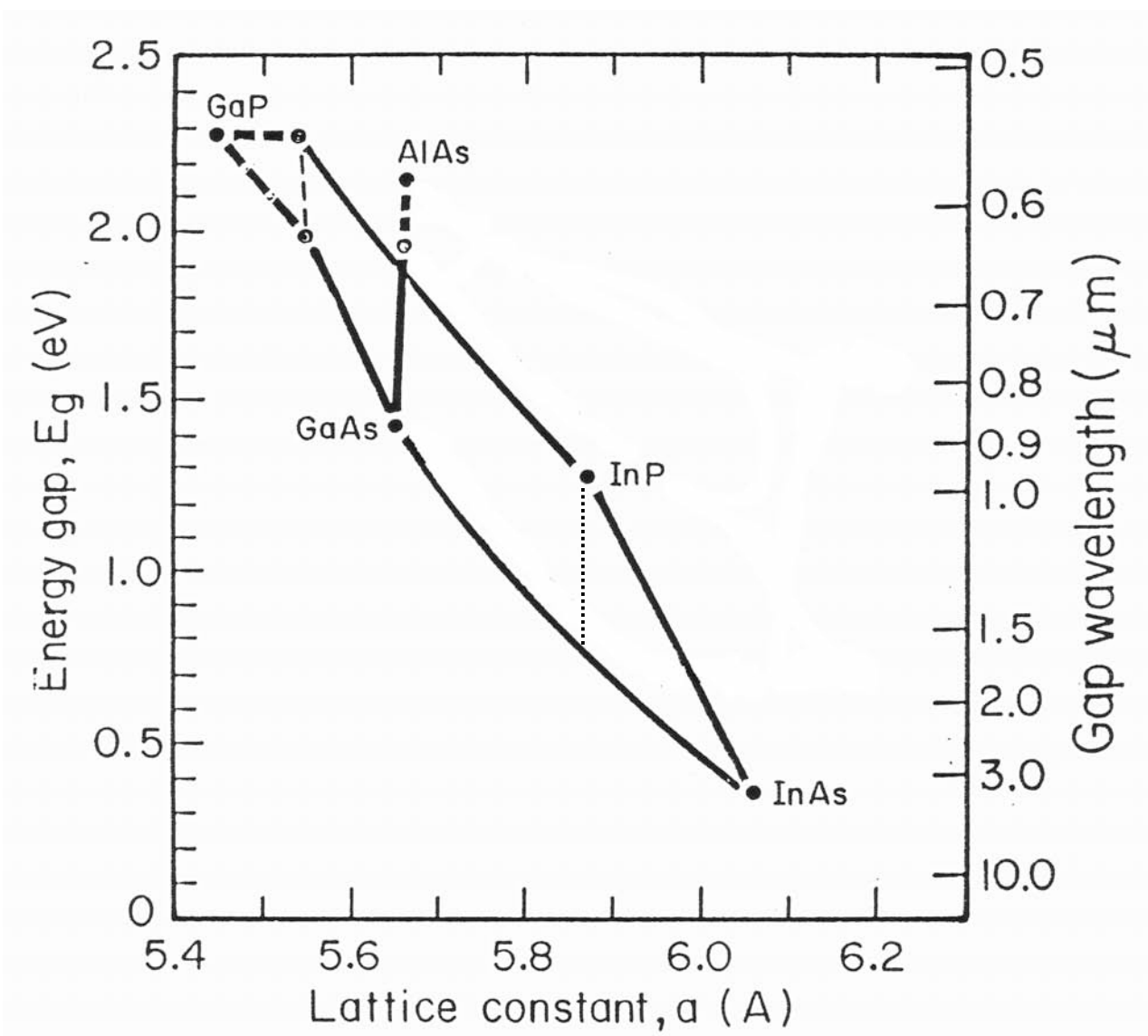
 - Gas phase epitaxy** □

 - 1. Vapor phase epitaxy; 2. Metallorganic chemical vapor deposition

 - Molecular beam epitaxy**

 - 1. Solid source; 2. Gas source; 3. Metalloganic; 4. Chemical beam

III-V systems: InGaAsP and AlGaAs



Insert 1 - Strained layers □

Strained layers

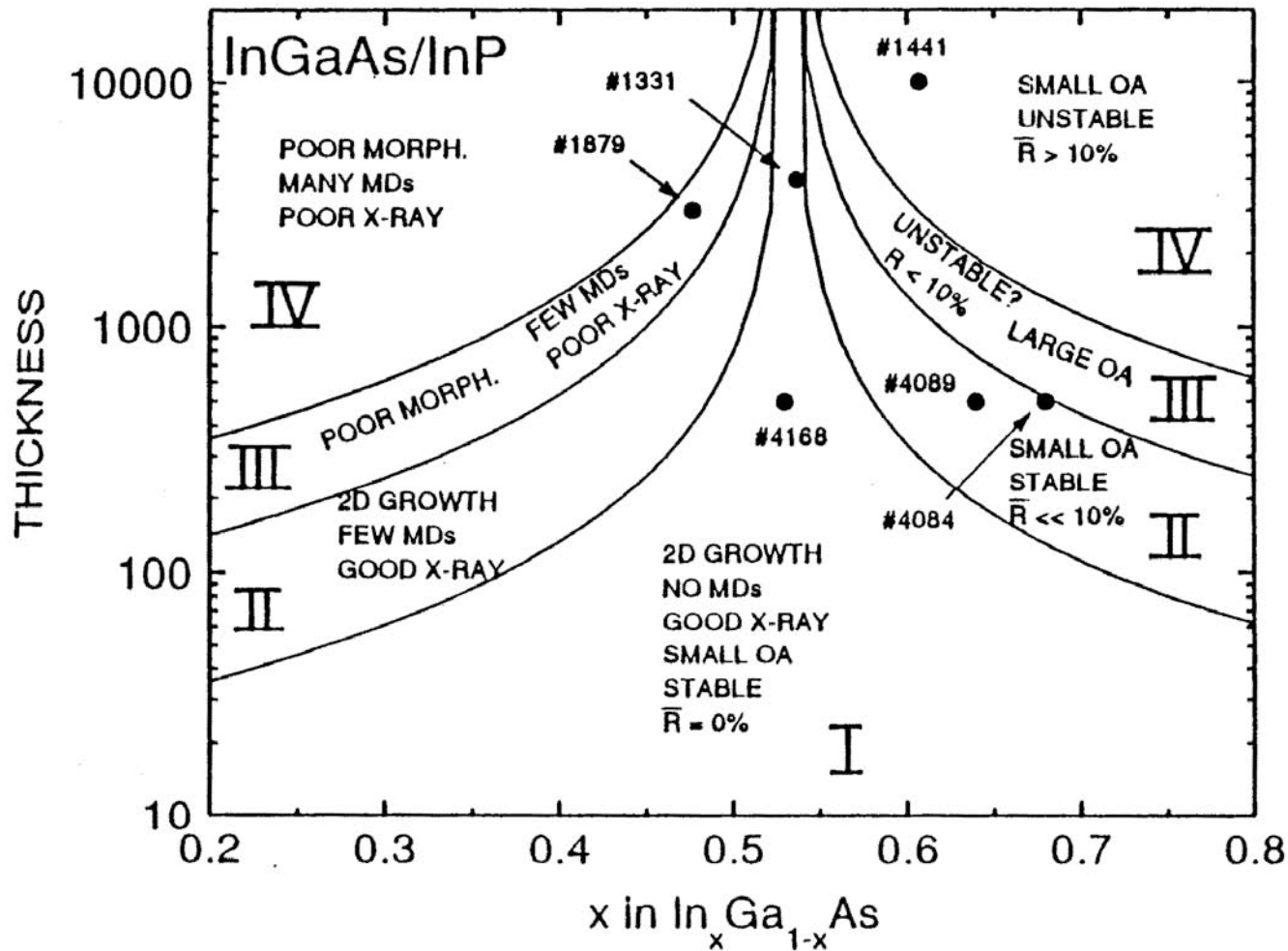
- 1. accommodating mismatch**
- 2. dislocation generation**
- 3. models**
- 4. parameter values**
- 5. effects of strain**

Impact of strain and QW quantization on bands □

15 nm wide wells
GaAs/In_{0.06}Ga_{0.57}Al_{0.37}As

Critical layer thicknesses - a final comment \square

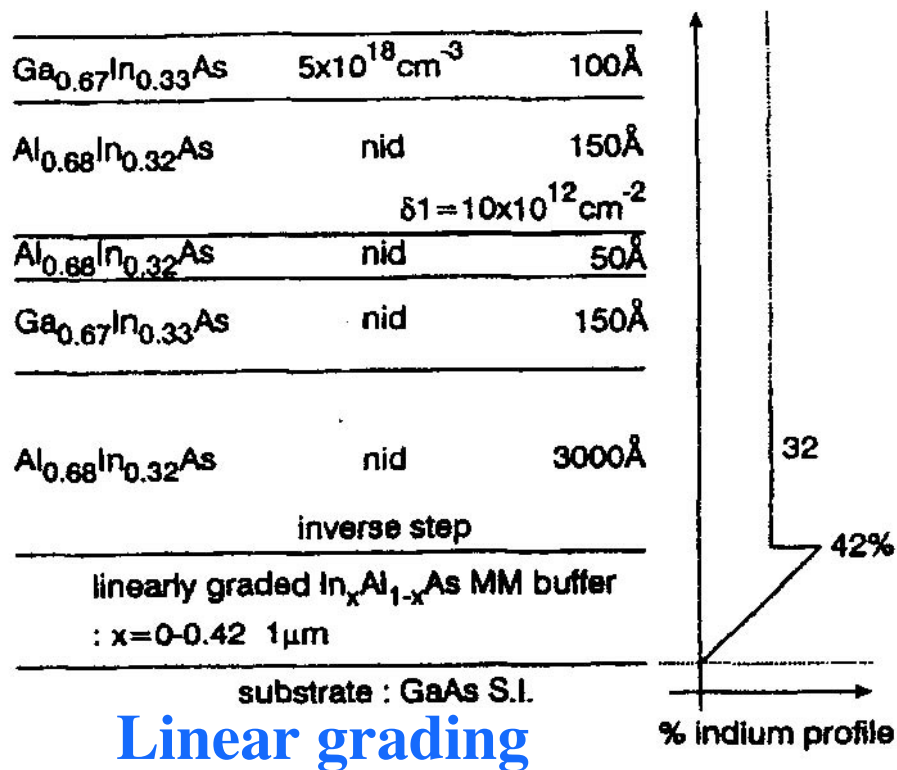
Courtesy of Jesus Del Alamo, Professor in Electrical Engineering and Jagdeep Bahl, PhD candidate; Used with Permission.



The critical thickness depends on the need -
work of Professor Jesus del Alamo and Jagdeep Bahl, MIT

Mismatched epitaxy - letting layers relax □

Graded lattice constant structures



n^+	$\text{In}_{0.30}\text{Ga}_{0.70}\text{As}$	13 nm	5×10^{18}
i	$\text{In}_{0.29}\text{Al}_{0.71}\text{As}$	20 nm	
n^+	$\text{In}_{0.29}\text{Al}_{0.71}\text{As}$	30 nm	2×10^{18}
i	$\text{In}_{0.29}\text{Al}_{0.71}\text{As}$	4 nm	
i	$\text{In}_{0.30}\text{Ga}_{0.70}\text{As}$	40 nm	
i	$\text{In}_{0.29}\text{Al}_{0.71}\text{As}$	0.4 μm	
i	$\text{In}_{0.20}\text{Al}_{0.80}\text{As}$	0.2 μm	
i	$\text{In}_{0.10}\text{Al}_{0.90}\text{As}$	0.2 μm	
i	GaAs		

Step grading

- There is no general agreement on which approach is superior and □ the choice often one of convenience and/or practicality.
- Because the last layer is often not fully relaxed, it is common to grade to a certain level and then step back, as seen in the structure on the left.

Epitaxy techniques - the spectrum of options □

Liquid Phase Epitaxy □

Vapor phase techniques

(hydrodynamic flow)

Hydride transport

Chlorine transport

Metallo-organic CVD

Molecular beam epitaxy

(ballistic flow)

solid source

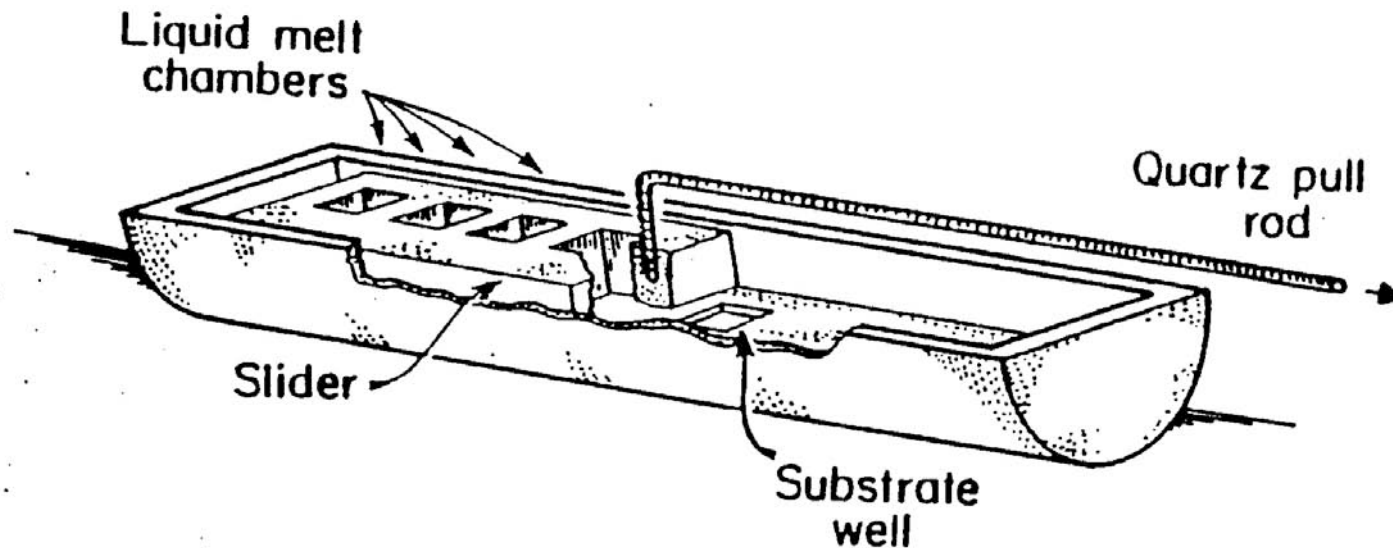
gas source

chemical beam

organo-metallic source

Liquid Phase Epitaxy - LPE

Growth from a Ga or In solution in a hydrogen ambient



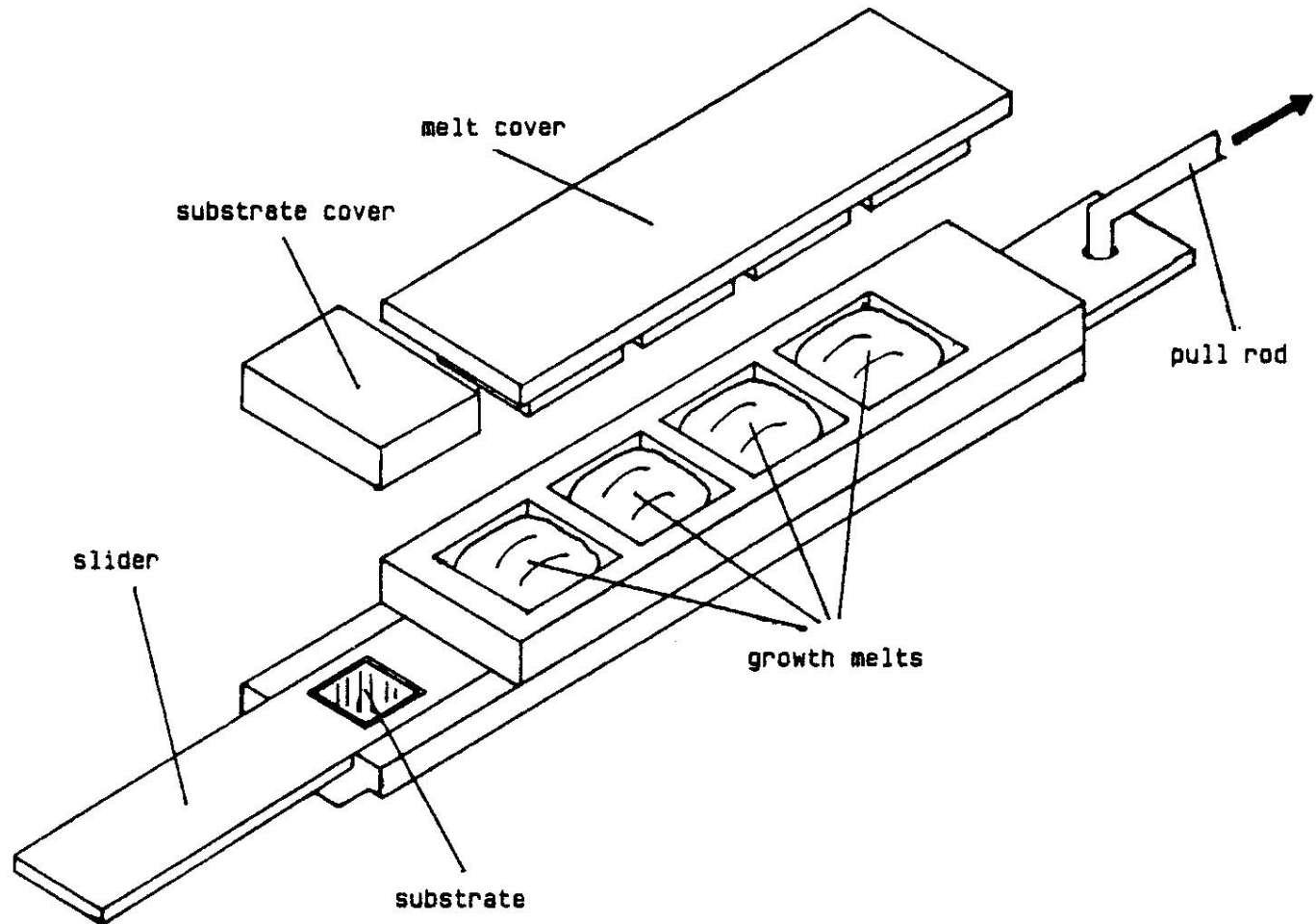
Advantages: Near-equilibrium growth, excellent crystal quality
Inexpensive; Fast

Disadvantages: Difficult to scale up for production
Dimensional control poor
Structure complexity limited

Current status: Widely used for LEDs and laser diodes in well established processes. Rarely used in new installations.

Liquid Phase Epitaxy - □

Explode view



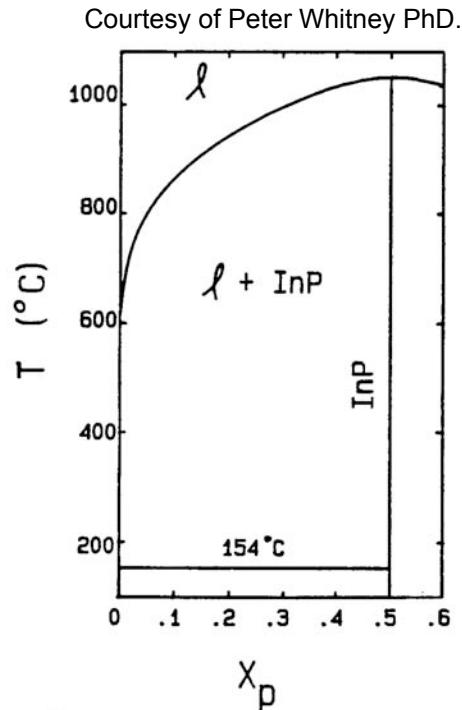
Material: □

Machined pyrolytic graphite

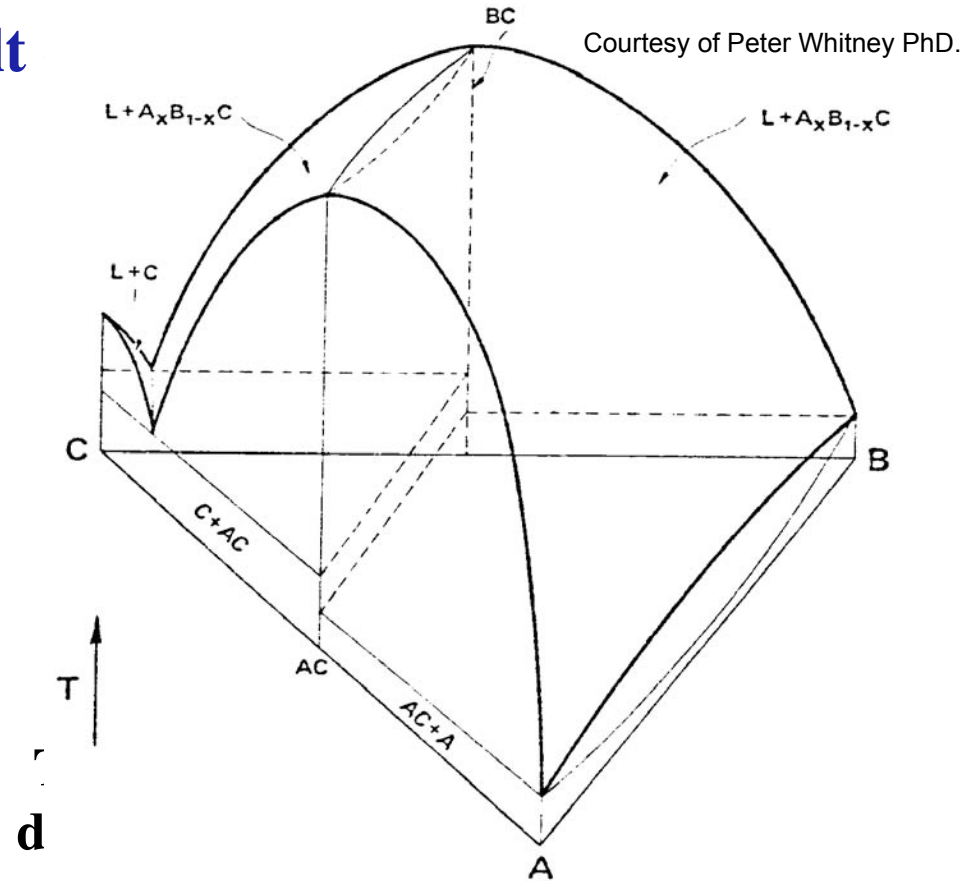
Growth ambient: □ Purified hydrogen at atmospheric pressure

Within a quartz tube in a resistance heated furnace

Liquid Phase Epitaxy - Melt



Binary phase diagram, In-P

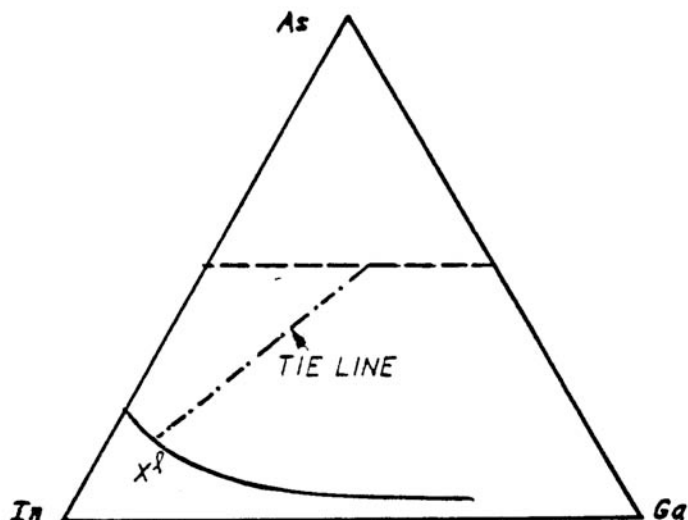


As T is reduced the binary (or ternary) solidifies from the melt:

- In the case of the binary shown on the left the liquid is $\text{In}_{1-x}\text{P}_x$, and the solid is InP.
- In the case of the ternary shown on the right, the situation is more complicated. The liquid is $\text{A}_y\text{B}_z\text{C}_{1-y-z}$ and the solid is $\text{A}_x\text{B}_{1-x}\text{C}$, but the relationship between the liquid composition and x is not shown in the diagram. (cont. next foil)

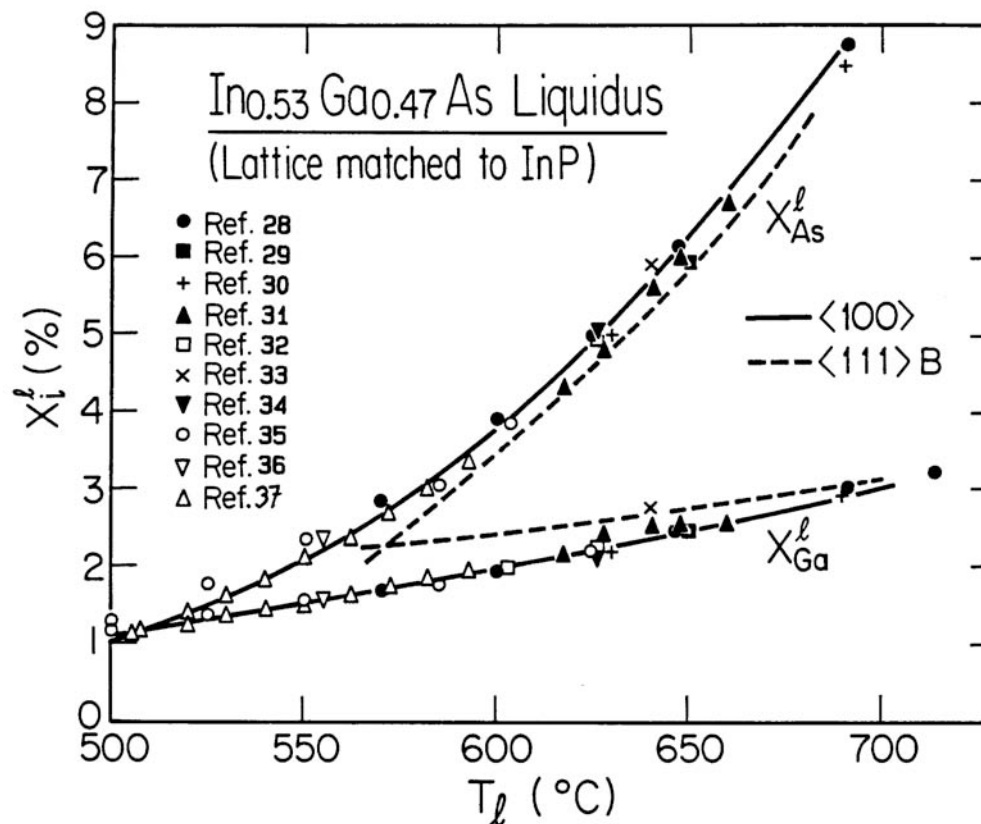
Liquid Phase Epitaxy - Melt calculation, cont.

Courtesy of Peter Whitney PhD.



Projected isotherm and tie line for In-Ga-As system

Courtesy of Peter Whitney PhD.

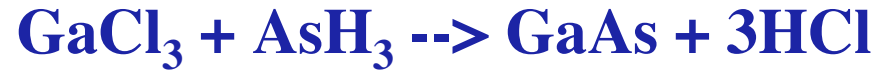


Equilibrium liquidus for InGaAs lattice-matched to InP

The relationship between the ternary liquid composition and the ternary solid composition is displayed by a tie line between a projected isotherm and ternary solid line, as shown on the left,

Or by a plot of liquid composition versus temperature for a specific solid composition of interest, as shown on the right.

Chlorine Transport Vapor Phase Epitaxy - VPE □



(Image deleted)

See Enda, H., Japanese Journal of Applied Physics. 18, 2167, (1979).

Advantages: □ Conceptually simple

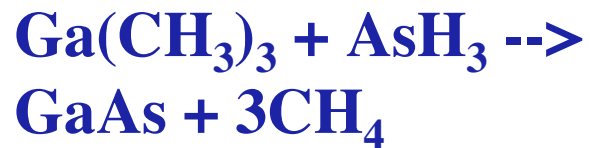
Disadvantages: □ Messy
Difficult to control group III supply
Uses toxic gases (AsH_3 , PH_3)

Current status: Used for some GaAs epitaxy where high purity (low background doping) needed but largely superceded by MOCVD

Metallorganic Chemical Vapor Deposition - MOCVD □

Group III elements transported
as a metallorganic compound

on a carrier gas: i.e.



Advantages: □ All sources gaseous
Precise composition and dimension control

Disadvantages: □ Involves complex chemistry □
Uses toxic gases (AsH_3 , PH_3) □

Current status: Viewed as the standard production process for
many epitaxial heterostructures

Metallorganic Chemical Vapor Deposition - MOCVD □

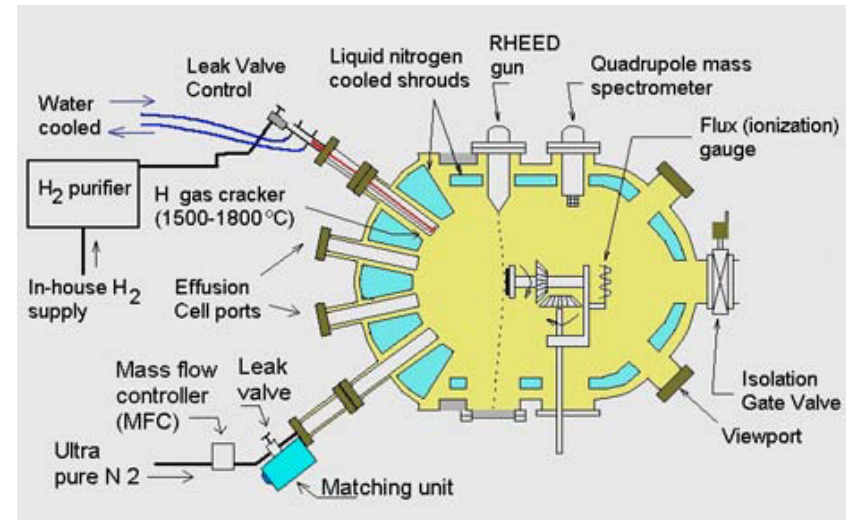
A group III precursor bubbler: □

Operation: Bubbler held in constant-temperature bath
Hydrogen "carrier" saturated with precursor (group
III organo-metallic) at fixed T
Precursor transported through heated lines as a
"gas"
Flow controlled by mass flow meters

Molecular Beam Epitaxy - □

MBE □

Slow vacuum deposition □
on a heated substrate □
under ultra-high vacuum □
conditions; growth a □
layer at a time □



Courtesy of Professor Yoon Soon Fatt, NTU.

Advantages: Extremely flexible, simple chemistry
Insitu monitoring; Atomic layer control
Non-equilibrium technique

Disadvantages: No in situ cleaning or purifying reactions
Expensive (to assemble and operate)
Non-equilibrium technique

Current status: A research workhorse; increasingly used heavily □
in production

Insert 2 - Comparison of techniques □

Comparison of epitaxy techniques

- 1. Table of techniques**
- 2. Timeline of techniques**
- 3. The spectrum from VPE to MBE**
- 4. Deviation for equilibrium**
- 5. Further comparison of VBE and MBE**

Insert 3 - MOCVD

MOCVD

- 1. Precursors**
- 2. MOCVD reactor**
- 3. Flow concerns in MOCVD reactor**
- 4. More flow issues**
- 5. MOCVD surface reactions**
- 6. Optical reflection oscillations used in MOCVD**