4.2 Other Semiconductor Crystal Growth Technologies

4.2.1 Single Crystals Other Than Si

Gallium Arsenide

We know that we need **III-V** semiconductors for <u>optoelectronic products</u>, so we have to make single crystals as input to our optoelectronics factory.

- We may use those single crystal for the product or we may just use them as the substrate for the thin film that will contain the product. Whatever they better be as perfect as possible because any defect present at the surface (e.g. a dislocation ending a the surface) will simply continue in any <u>epitaxial layer</u> we might produce.
- Let's look at GaAs single crystal production as en example for other III-V's like InP or GaP. However, not all III-V's can be grown on a similar way; we will come to that.

The <u>Czochralski method</u> for growing crystals should work for **GaAs**, too. So let's try it and see why it is far more difficult to grow compound single crystals than **Si** (or **Ge**) single crystals

- The melting point of GaAs, however, is T_m(GaAs)=1.238 °C. This is a first indication that things might not be quite as simple as they look. Obviously, we now need a phase diagram of the Ga As system. Here it is: We mix Ga and As in a (molar) ratio of 50:50 and melt it in a crucible. That should be easy, because both elements have low melting points (Ga: 30 °C; As: ? °C it actually doesn't melt but evaporates at a sublimation or boiling point of 615 °C).
- So what happens if our mixture is not 50 : 50 but 49.999 : 50.001? We will have little droplets of liquid in our growing crystal, that solidify later and probably cause defects at a concentration that is large in comparison to what we call "low defect concentration".
- To make things worse, whatever you mix together and melt, at the required 1.238 °C the As in the melt will evaporate off quickly; the concentration thus changes as a function of time.

The catchword now is "liquid encapsulation technique" (*LEC*).

- The melt in the crucible will be enclosed in B₂O₃, a viscous liquid at the growth temperature. It encloses the entire melt, i.e. sits between crucible and melt and on the surface. Obviously, we have <u>another</u> <u>case</u> of getting surface energies right here. The seed crystal is now dipped through that layer into the melt and so on.
- There is no way that the crystal you get will come close in quality to Si crystals. The present state of the art is given in this relatively <u>short</u> <u>article</u> given at that Czochralski-Symposium 2003 in Poland
- You will not understand some of the topics in the article, but you will get a feeling for the complexity involved in growing good GaAs single crystals.

Of course, any other **III-V** compound crystals grown in a similar way runs into all the difficulties hinted at above plus some of its own. This is particularly true for **GaN**, which will be shortly discussed below.

Less of course, there are some other ways of growing GaAs (and other) single crystals but we cam't go for that here.



Gallium Nitride

In the (slightly changed) words of Dr. Yoke Khin **Yap**; Department of Physics, Michigan Technological University:

High-quality GaN single-crystals are still not possible because the melt growth of GaN single-crystal is prohibited due to the extremely high decomposition pressure (~ 45000 atm) involved at the melting point (~2500 °C). At present, GaN single-crystals can be grown by the solution method at (1300 - 1600) °C and N₂ gas pressure of (10000 - 17000) atm. Because of the need of high temperatures and gas pressures, the solution growth method is not suitable for mass-production of GaN single-crystals.

We have been growing high-quality **GaN** single-crystals by a **Na** flux method at **800** °C and **<50 atm**. This technique is promising for mass production of **GaN** single-crystals at significantly low temperatures and pressures. A similar approach is also applicable for growing other nitride crystals like **w-AIN** and **h-BN**.



From Yap Research Lab, Michigan Tech

So how come we have a flourishing GaN industry, making UV - blue LED's and Lasers?

- Because the input to a "commercial" GaN factory consists of substrates with a thin GaN layer on it. The substrate is either Al₂O₂ with a lattice misfit of 13.8% and a thermal expansion coefficient mismatch of 25.5%
- Not good. So you look for a better substrate and find SiC. Not so hot either, look at our "master graph". Any you now need SiC single crystal (see below).

Nevertheless, **GaN** made it into the ranks of serous semiconductors with billion €or so business attached. What we learn form this is that semiconductor technology, like politics, is the art of the possible (*"Politik ist die Kunst des Möglichen*; Otto von **Bismarck**)

Silicon Carbide

- We already know that **SiC** comes in many <u>polytypes</u>. That makes us suspect that growing a single crystal of just one polytype may not be so easy.
 - The fact that its melting point is < 2.500 °C and that SiC is extremely hard does not make the task to produce "perfect" single-crystalline wafers any easier.
 - On the other hand, inside the huge reactors designed for making absolutely imperfect poly-crystalline SiC of any polytype for products like grinding paper, one does find nice single crystals hanging on the wall on occasion, so single crystal growth should be possible.

Here is how it is done.

In most cases, large single crystals are grown from a melt or some solution (e.g. quartz, or sugar if you leave you coffee cup around too long), but this is not a feasible option for **SiC** single crystal growth since **SiC** does not have a liquid phase under normal conditions (i.e. without applying a large pressure). **SiC** is also extremely hard (close to diamond) and therefore has a high melting point (or is it the other way around?).

There is also, in principle, no crucible material that could contain molten SiC at is nominal melting point temperature of < 2.500 °C. Nevertheless, SiC was grown from a melt at 2200 °C and 150 bar in a recent study, but this is probably not a commercially viable process.</p>

We need a basically new method of crystal growth, and the main method used nowadays is **physical vapor transport** (*PVT*) also known as seeded sublimation growth or modified Lely method.

A piece of **SiC** is heated to **(1800-2600)** ^o**C** at low pressure. Due to the high sublimation rate, **SiC** vapor forms and deposits itself on a cooler single-crystalline seed crystal. Straightforward and basically simple, as shown in the schematic picture below.

However, pondering the situation, some questions should come to mind:

- What materials can you use for the crucible and everything else that gets hot? After all, not many materials can cope with temperatures above 2000 °C! Well, you are basically stuck with graphite, and maybe a bit of Ta here and there. That means, of course, that you are forming SiC also on your crucible walls and everywhere else. If it flakes off, you will have a defect problem.
- What kind of growth rate can you get? As you would expect: Not much! Growth rates depend on many parameters, but are in the range of 0.2 2 mm/hr. That's about a factor of 50 slower than the growth rates for Si crystal pulling and that makes SiC crystal growing automatically expensive

What polytype will you get (hoping that it will not be a mixture)? What determines what you get? Can you control it and, if yes, how?

Good questions! First, you might get mixtures as shown in the picture (*courtesy of the Erlangen group*). Otherwise, the following parameters are essential:

- Polytype of the seed crystal (as you might have guessed).
- "Face" of the seed crystal; i.e if the surface is a C- or a Si layer. If you start with a 4H-SiC seed crystal, for example, you tend to get 4H-SiC if you have a C-face, and 6H-SiC if you have a Si face. Why? Nobody really knows.
- Temperature difference and gradient betwen SiC source and seed. Small values tend to favor 4H-SiC, larger values 6H-SiC growth.
- Gas composition. Whatever gas you add will influence the polytype you obtain. **C**-rich gases, for example, promote **4H-SiC** growth
- The pressure, oddly enough, seems not to have a large influence on polytypie.

Note that while the polytype 6H is the easiest to grow, 4H would be favored by the power electronics industry.

Last not least: What kind of crystal quality do you get? What is the dislocation density?

The bad news is: the dislocation density is high. The good news is, you do not worry too much about that - you worry about something weird called "micropipe (and mixtures of polytypes, and all kinds of stacking faults or special boundary faults, and carbon inclusion, or Si inclusion, or big voids, ...).

- What are micropipes? Well, micropipes are hollow channels running through the lattice; the diameter of these pipes is (0.1 5) μm. One might consider them to be screw dislocations with a gigantic Burgers vector and a hollow core.
- Micropipes also will definitely kill any device that contains one of them. They thus must be avoided as much as possible! Micropipes are also somehow connected to the growth mechanism of the crystal. This is neatly illustrated in the picture on the right (taken with a scanning force microscope, courtesy of H. Strunk; Uni Erlangen) where typical growth spirals are visibly centered around a micropipe.



Let's look at the state of the art of what is around. To quote from the product sheet of the major **SiC** supplier Cree, Inc. (located somewhat ironically in Silicon Drive 4600 in Durham, North-Carolina, **USA**):

At present, wafer diameters are **50.8 mm** or **76.2 mm**; doping (usually with **N** for **n**-type and **AI** for **p**-type) at high levels produces resistivities in the **0.0x m**Ω**cm** region. Or there is no doping for semi-insulating stuff. **4H**- and **6H**-**SiC** polytypes are sold

The **2003** state of the art (mostly in the laboratories and not necessarily on the market) is summarized in the following table:

Diameter		100 mm "Four-inch"	For Si , 100 mm was the standard back in the late 70 ties/early 80 ties).
Defects	Micropipes	< 1 cm ⁻² for 3" < 30 cm ⁻² for 100 mm	Increasing wafer size usually dramatically increases micropipe density
	Dislocations	3 · 10 ³ cm ⁻² achieved	Factor 10 reduction

Of course, in the many laboratories (university and industrial) devoted to SiC, some data might be even better.

The Rest

Just kidding. All we can do here is to look at a list of what is left over

🦉 <u>Germanium</u>.

- Obviously, Ge wafer technology can follow the Si lead in principle. It should be even somewhat easier to grow Ge single crystals because the melting point of Ge is lower (938 °C vs. 1410 °C, resp.)
- However, while there is a lot of SiO₂ around to start from, where do you find Ge-minerals? Indeed, the price of germane,GeH₄, needed for starting, is orders of magnitudes higher that that of silane, SiH₄.
- All things considered, you can get pretty good dislocation free **Ge** wafers up to one or two **300 mm** diameters but for a price. The total quality, however, is not as good as the mass-produced **300 mm** Si wafer.

II-VI Semiconductors

- There are many kinds, and you may have to find a specific way of growing single crystals for each kind.
- The state of the art in terms of what you can buy are small and rather imperfect single crystals of CdSe, ZnO and
- Everything else mentioned in <u>chapter 2</u>.
 - Forget it .No single crystals to speak of yet.

Special Wafers

- Some Si microelectronic factories have switched from regular Si wafers as input to "SOI" wafers; "Silicon on Insulator"; e.g. AMD in Dresden
- Read the <u>article</u> in the link if you want to find out what SOI is all about