

Crystalline Silicon PV Technology

Note:

Take-home Midterm exam starts Feb. 28

Due March 13th @ beginning of class

February 23, 2012

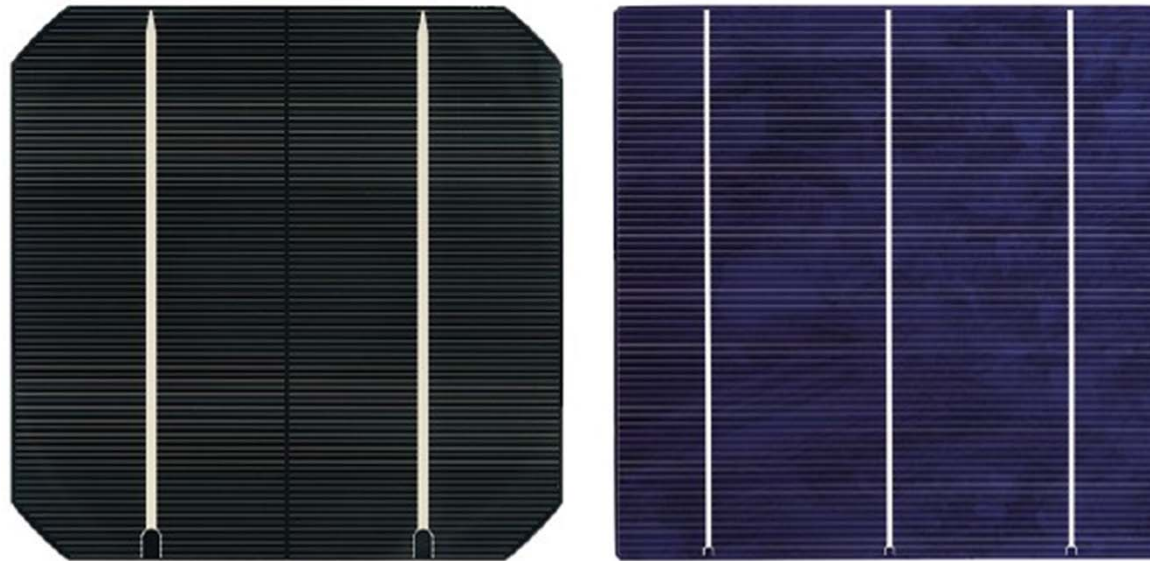
The University of Toledo, Department of Physics and Astronomy
SSARE, PVIC

Principles and Varieties of Solar Energy (PHYS 4400)
and
Fundamentals of Solar Cells (PHYS 6980)

Crystalline/Polycrystalline Silicon PV Technology (some facts)

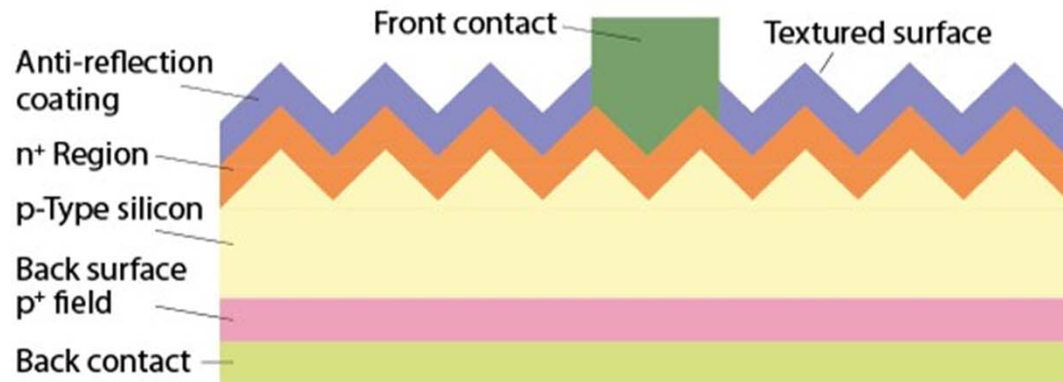
- Crystalline silicon PV cells are used in the largest quantity of all types of panels on the market, representing about 90% of the world total PV cell production in 2008.
- The highest energy conversion efficiency reported so far for research crystalline silicon PV cells is 25%.
- Standard industrial cells are limited to 15–18% with the exception of certain high-efficiency cells capable of efficiencies greater than 20%.
- High-efficiency research PV cells have advantages in performance but are often unsuitable for low-cost production due to their complex structures and the lengthy manufacturing processes required for fabrication.
- World annual production of PV cells reached more than 7.9 GWp in 2008, 10.6 GWp in 2009, 18.6 GWp in 2010, 28 GW in 2011.
- Average annual growth rate in PV cell production/installation over the last decade has been ~40% (CAGR). Cumulative installed capacity is ~ 67 GW.
- Yet the electrical power generated by all PV systems around the world is less than 0.1% of the total world electricity generation

T. Saga, NPG Asia Mater. 2(3) 96–102 (2010)



Monocrystalline silicon solar cell

Polycrystalline silicon solar cell



Typical mono- and polycrystalline silicon solar cells (upper), and simplified cross-section of a commercial monocrystalline silicon solar cell (lower) (© 2010 Sharp).

Production of “Standard” Silicon PV Cells

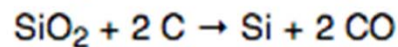
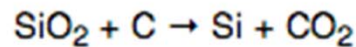
- Standard cells are produced using monocrystalline and polycrystalline boron-doped p-type silicon substrates. Cells are typically 125 mm (5 inches) or 156 mm (6 inches) square, respectively.
- Monocrystalline solar cells are typically produced from pseudo-square silicon wafer substrates cut from column ingots grown by the Czochralski (CZ) process.
- Polycrystalline cells, on the other hand, are made from square silicon substrates cut from polycrystalline ingots grown in quartz crucibles. The front surface of the cell is covered
- with micrometer-sized pyramid structures (textured surface) to reduce reflection loss of incident light.
- An anti-reflection coating (ARC) of silicon nitride (SiN_x) or titanium oxide (TiO_x) is overlaid on the textured silicon surface to further reduce reflection loss.
- A highly phosphorous doped n⁺ region is produced on the front surface of boron-doped p-type substrates to form p–n junctions.

The value chain for crystalline silicon solar cells and modules is longer than that for thin-film solar cells.

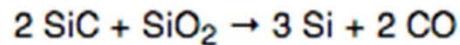
- There are generally three industries related to crystalline silicon solar cell and module production:
 1. metallurgical and chemical plants for raw material silicon production,
 2. monocrystalline and polycrystalline ingot fabrication and wafer fabrication by multi-wire saw, and
 3. solar cell and module production.
- The cost of PV production is roughly divided in half between solar cell module production and balance-of-system fabrication, which includes the inverter, cables and installation.
- The fabrication cost for solar cell modules includes the cost of the silicon substrate (50%), cell processing (20%) and module processing (30%).
- The cost share is therefore strongly affected by the market price for poly-silicon feedstock, and reducing the cost of the silicon substrate remains one of the most important issues in the PV industry.

Production of metallurgical grade Si

Silicon is commercially prepared by the reaction of high-purity [silica](#) with wood, charcoal, and coal, in an [electric arc furnace](#) using carbon [electrodes](#). At temperatures over 1,900 °C (3,450 °F), the carbon reduces the silica to silicon according to the [chemical equations](#):



Liquid silicon collects in the bottom of the furnace, and is then drained and cooled. The silicon produced via this process is called *metallurgical grade silicon* and is at least 98% pure. Using this method, [silicon carbide](#) (SiC) may form. However, provided the concentration of SiO₂ is kept high, the silicon carbide can be eliminated:



In September 2008, [metallurgical](#) grade silicon cost about [USD 1.45](#) per pound (\$3.20/kg),^[9] up from \$0.77 per pound (\$1.70/kg) in 2005.^[10]

Pure silicon (>99.9%) can be extracted directly from solid silica or other silicon compounds by molten salt electrolysis.^{[11][12][13][14]} This method, known from 1854^[15] (see also [FFC Cambridge Process](#)) has the potential to directly produce solar grade silicon without any CO₂ emission and at much lower energy consumption.

Purification by Physical Methods

Early silicon purification techniques were based on the fact that if silicon is melted and re-solidified, the

The earliest method of silicon purification, first described in 1919 and used on a limited basis to make radar components during World War II, involved crushing metallurgical grade silicon and then partially dissolving the silicon powder in an acid. When crushed, the silicon cracked so that the weaker impurity-rich regions were on the outside of the resulting grains of silicon. As a result, the impurity-rich silicon was the first to be dissolved when treated with acid, leaving behind a more pure product.

In zone melting, also called zone refining, the first silicon purification method to be widely used industrially, rods of metallurgical grade silicon are heated to melt at one end. Then, the heater is slowly moved down the length of the rod, keeping a small length of the rod molten as the silicon cools and re-solidifies behind it. Since most

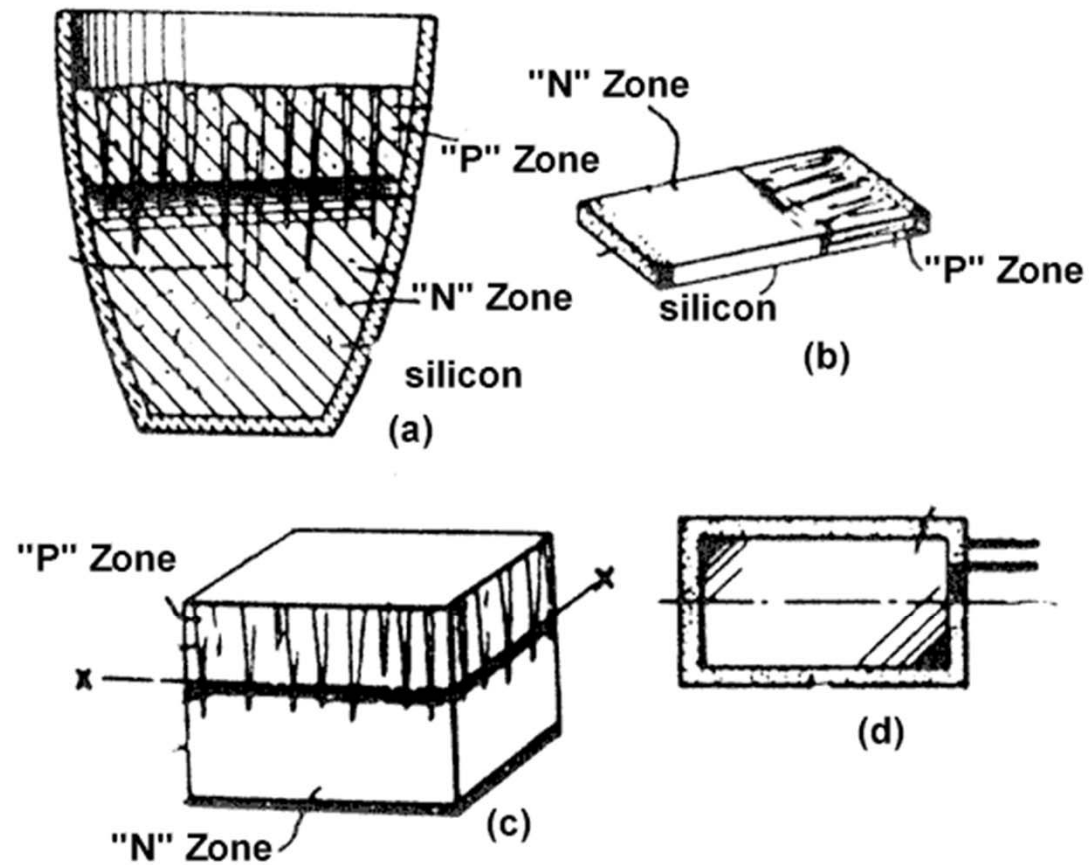
when the process is complete, most of the impurities in the rod will have been moved into the end that was the last to be melted. This end is then cut off and discarded, and the process repeated if a still higher purity is desired.

400 kgs



From Wikipedia

In studies of recrystallized melts of pure silicon prepared for this purpose, Ohl [2] discovered the presence of a well-defined barrier in ingots grown from commercially available, high-purity silicon.

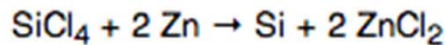


(a) Cast ingot showing natural junction formed by impurity segregation during melting; (b) photovoltaic device cut perpendicular to junction; (c) device cut parallel to junction; (d) top surface of device cut parallel to junction.

Purification by Gas Phase Chemical Methods

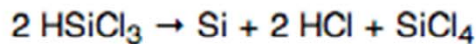
Today, silicon is purified by converting it to a silicon **compound** that can be more easily purified by distillation than in its original state, and then converting that silicon compound back into pure silicon. **Trichlorosilane** is the silicon compound most commonly used as the intermediate, although **silicon tetrachloride** and **silane** are also used. When these gases are blown over silicon at high temperature, they decompose to high-purity silicon.

At one time, **DuPont** produced ultra-pure silicon by reacting silicon tetrachloride with high-purity **zinc** vapors at 950 °C, producing silicon:



However, this technique was plagued with practical problems (such as the **zinc chloride** byproduct solidifying and clogging lines) and was eventually abandoned in favor of the Siemens process.

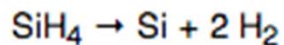
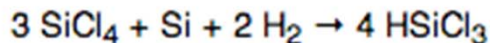
In the **Siemens process**, high-purity silicon rods are exposed to trichlorosilane at 1150 °C. The trichlorosilane gas decomposes and deposits additional silicon onto the rods, enlarging them:



Silicon produced from this and similar processes is called **polycrystalline silicon**.

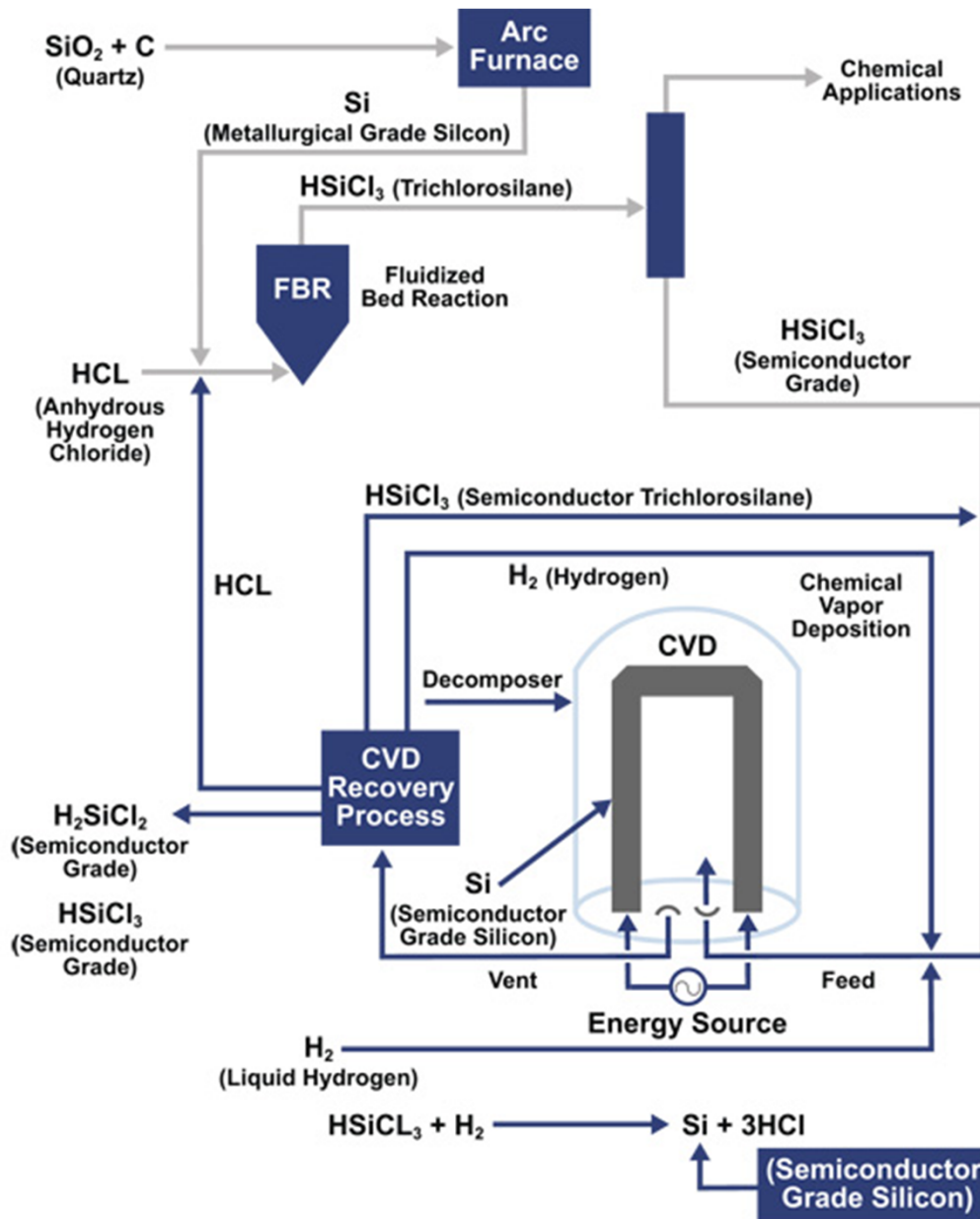
Polycrystalline silicon typically has impurity levels of less than 10^{-9} .

In 2006 **REC** announced construction of a plant based on fluidized bed technology using silane:^[17]



From Wikipedia

From quartz rock to hyper-pure poly Si (99.999999999%)



Hemlock Semiconductor Group (HSC) manufactures high-purity poly Si feedstock for the manufacture of solar and semiconductor products. HSC's manufacturing uses trichlorosilane, which is converted to polysilicon in a chemical vapor deposition (CVD) reactor employing an U-rod. All chemicals are captured and recycled.

Impurities are 1 part in 10^{11} , "intrinsic" (i.e. impurity) dopant concentration of $\sim 10^{12} \text{ cc}^{-1}$, or 1 ppt (i.e. 1 part per trillion)

Hemlock Semiconductor Group



- Owned by Dow Corning Corporation (63.25%), Shin-Etsu Handotai (24.5%), and Mitsubishi Materials Corporation (12.25%)
- Began operation in 1961 in Hemlock, Michigan
- Leading world supplier of high purity polycrystalline silicon to the semiconductor and solar industries
- Currently completing several major expansions in Michigan and Tennessee
- 36,000 metric ton capacity by 2010

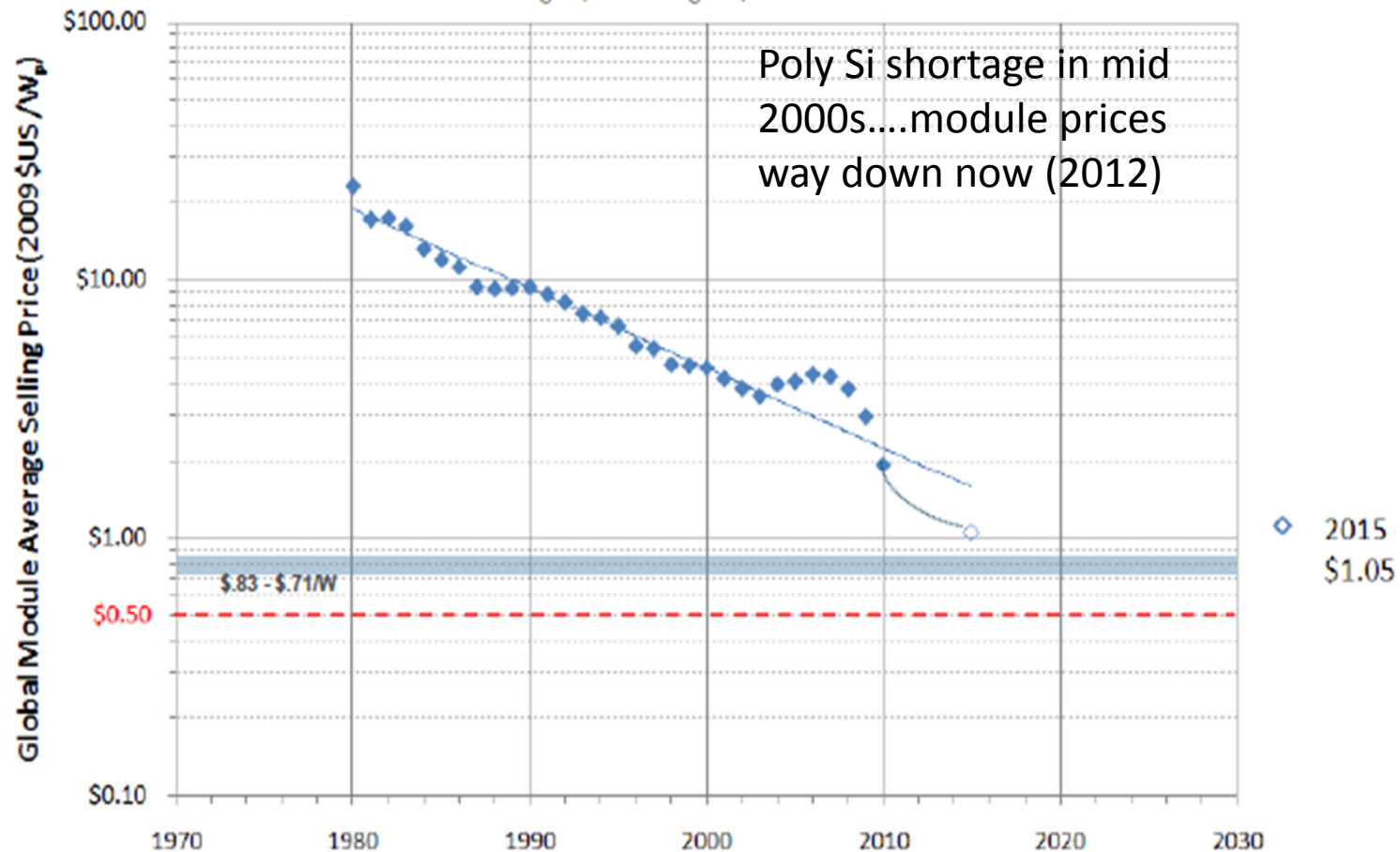


http://www.hscpoly.com/content/hsc_home/default.aspx?bhcp=1

Solar PV Experience Curves:

Crystalline Silicon (c-Si)

Sources: Navigant, Bloomberg NEF, NREL internal cost models



US net exporter in PV in 2010 (+1.9 B\$); \$2.5 billion was polysilicon.

<http://www.greentechmedia.com/articles/read/stat-of-the-week-positive-us-solar-trade-balance/>

The Float-Zone and Czochralski Crystal Growth Processes

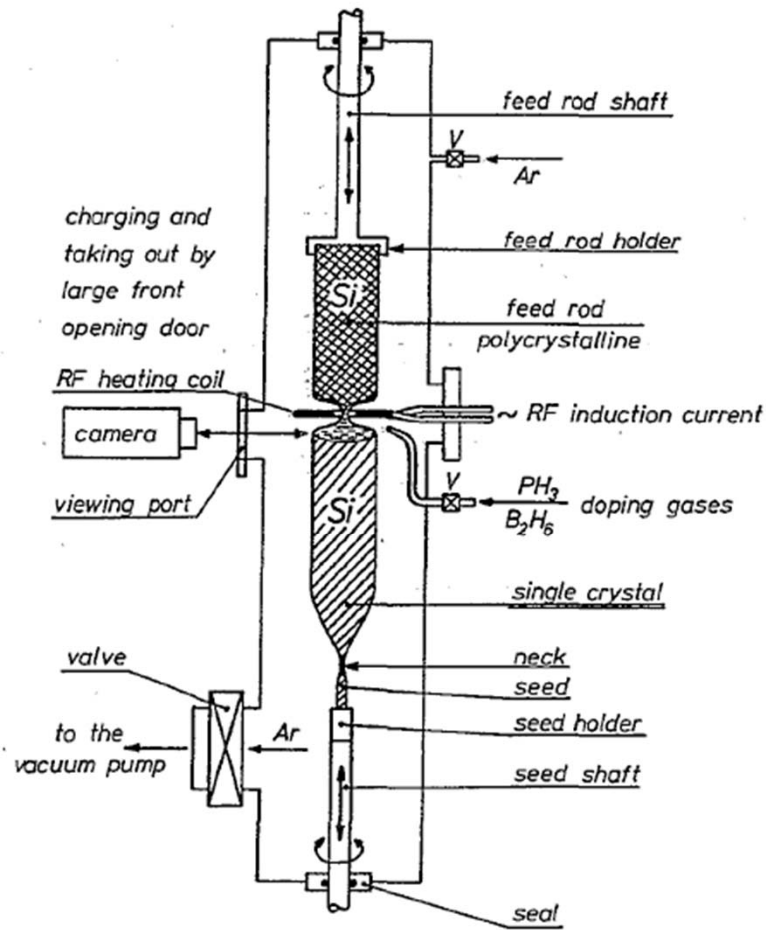


Fig. 9. Schematic of a modern FZ-Si single crystal puller for the growth of large crystals.

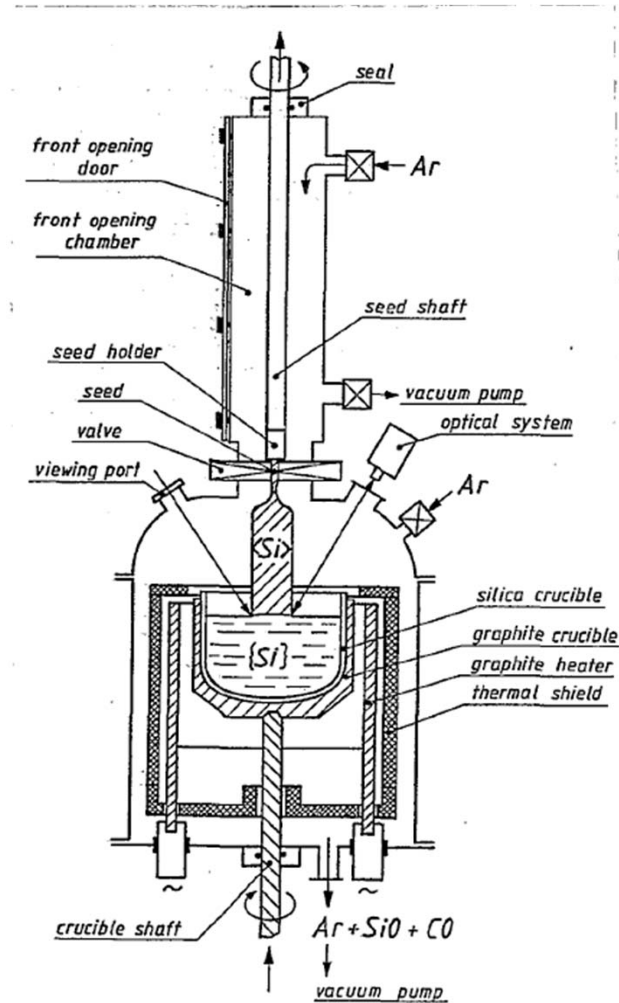
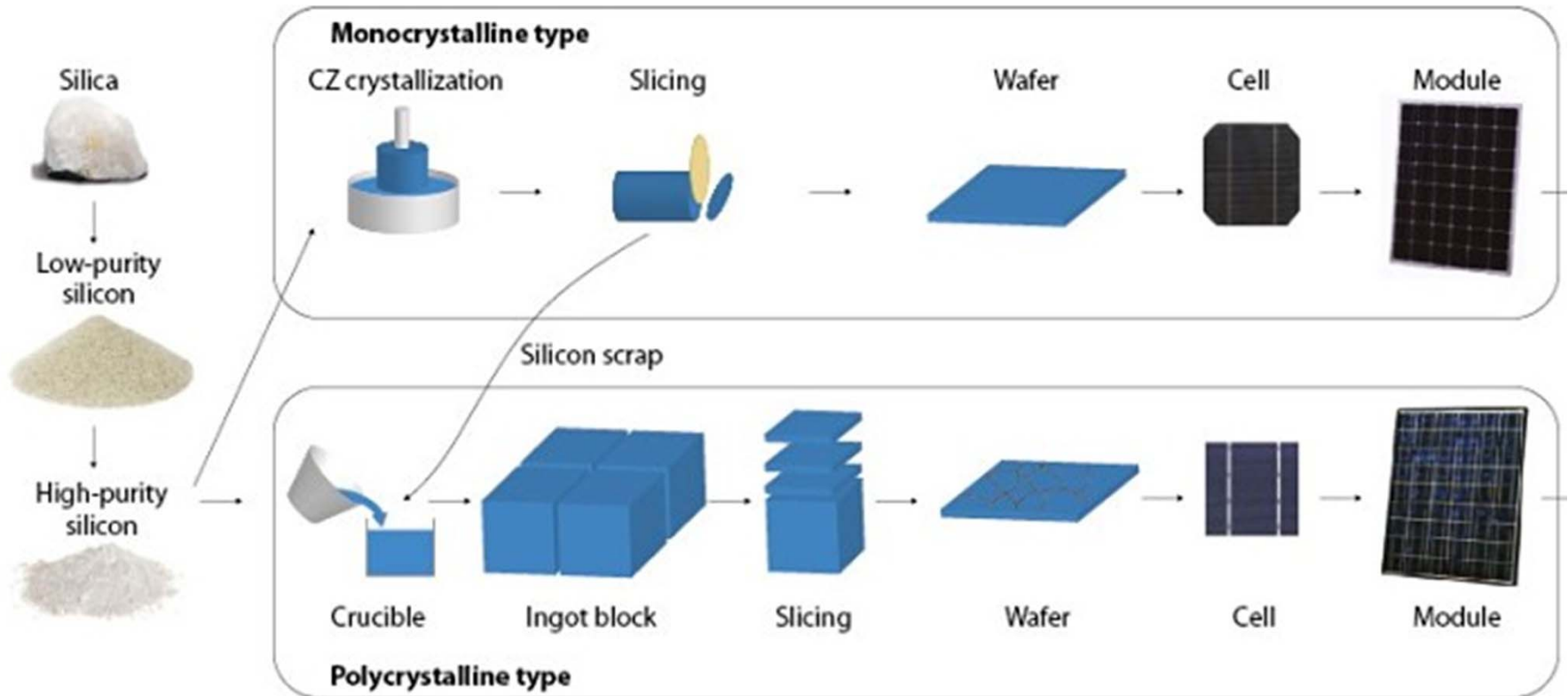


Fig. 6. Schematic of a modern CZ-Si single crystal puller for the growth of large crystals.

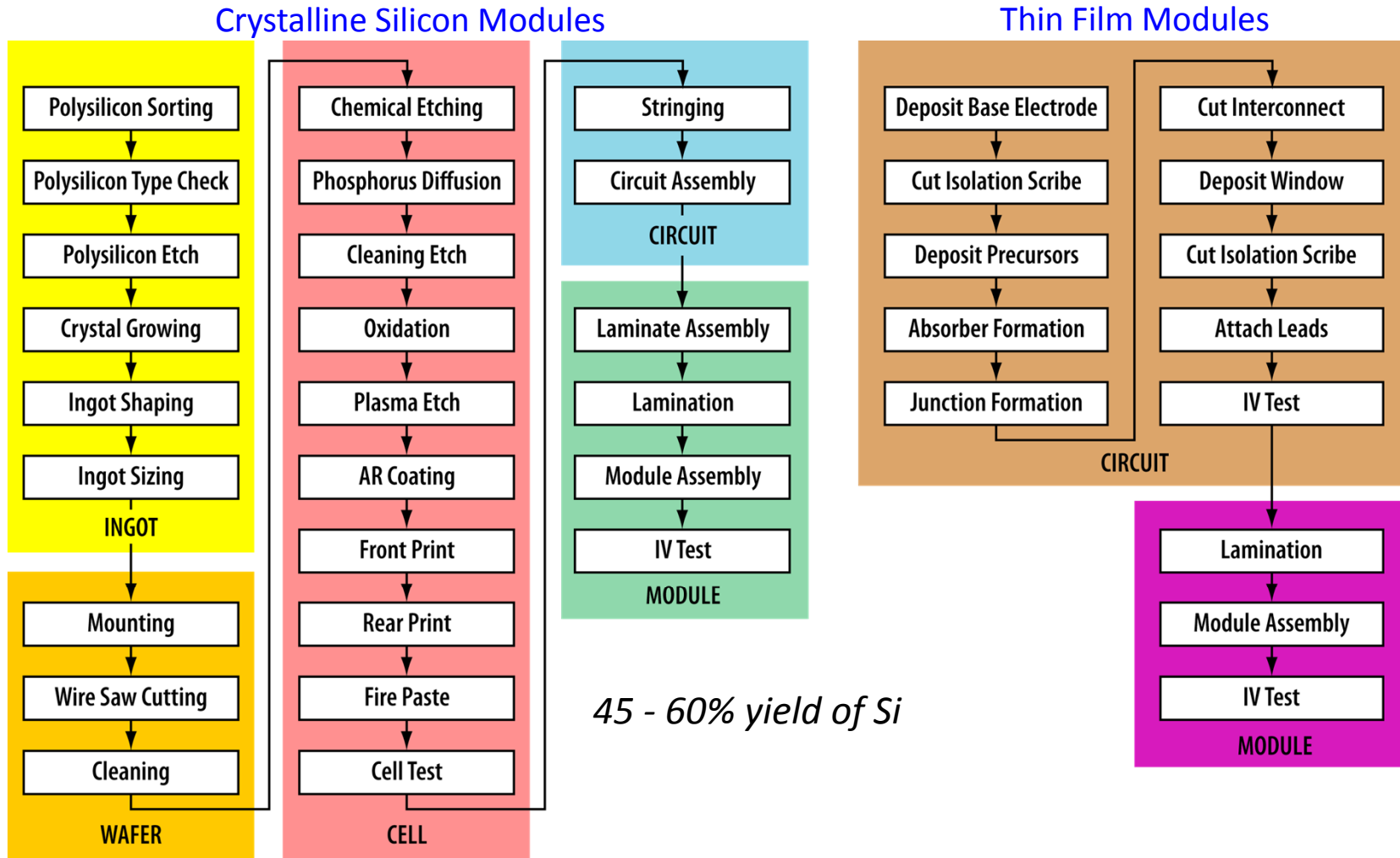
W. Zulehner, Wacker Siltronic AG

From Raw Silica to Final Cell



Comparison of Manufacturing Sequence For X-Si and Thin Film

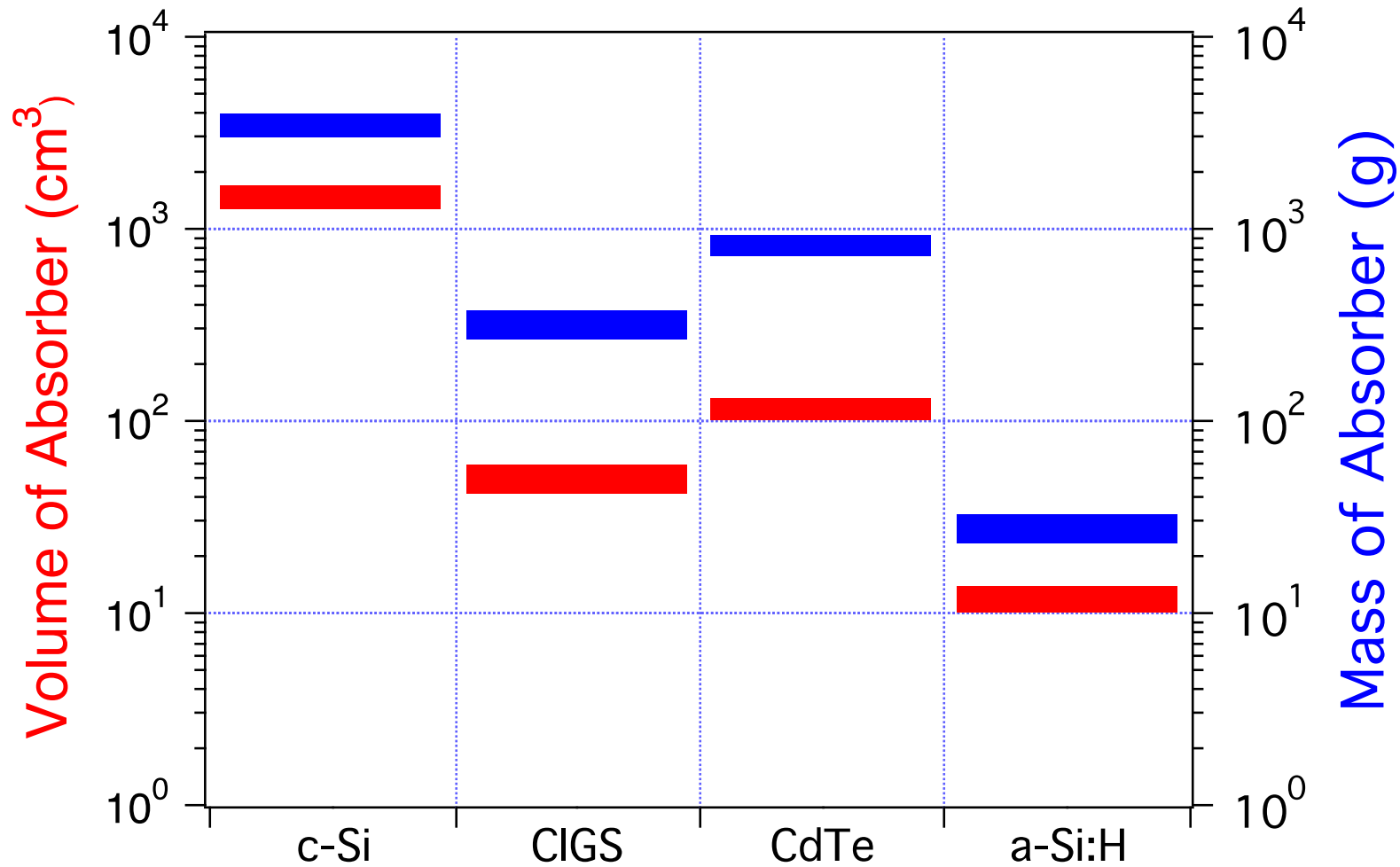
Courtesy of Brian Keyes, NREL



Ref: 'Overview and Challenges of Thin Film Solar Electric Technologies', Harin S. Ullal, Ph.D., NREL 2008

Thin-Films Use Less SC Material

Courtesy of Brian Keyes, NREL

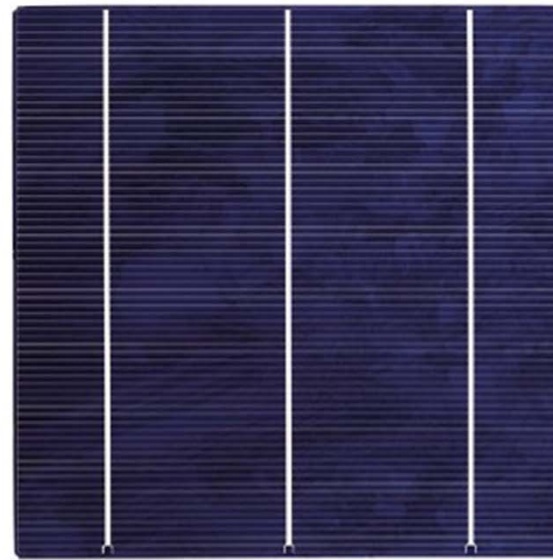


Amount of material needed for 1 kW output (more area for less efficiency).

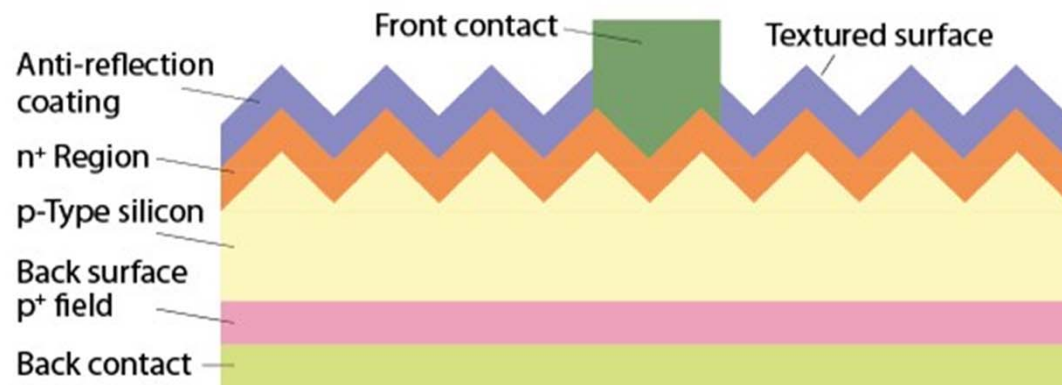
“Standard” Silicon PV Cells



Monocrystalline silicon solar cell



Polycrystalline silicon solar cell



Typical mono- and polycrystalline silicon solar cells (upper), and simplified cross-section of a commercial monocrystalline silicon solar cell (lower) (© 2010 Sharp).

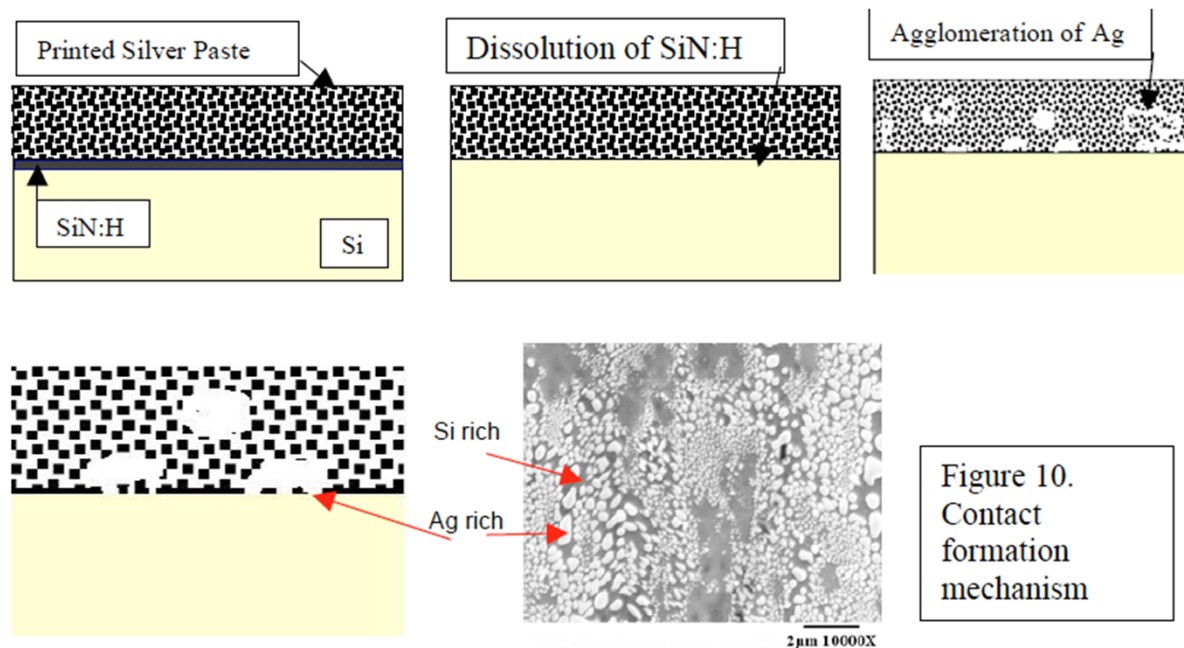
“Standard” Silicon PV Cells (cont.)

- Back-surface p+ field (BSF) regions are formed on the back surface to suppress recombination of minority carriers. The BSF regions are usually formed by firing screen-printed aluminum paste in a belt furnace.
- The minority carriers (electrons) generated in the silicon bulk and diffusion layers are collected by silver contacts (electrodes) formed as gridlines connected by a bus bar to form a comb-shaped structure.
- The back contact is usually an aluminum layer which is connected the adjacent cell via soldered copper or printer silver interconnects.
- The front contact is formed using screen-printed silver paste applied on top of the ARC layer, with “fire through” technology, which also produces the BSF
- The screen-printed front silver contact prepared by firing to penetrate the ARC is one of the most important techniques for large-volume fabrication of modern xstal Si cells.

Metallization-firing of Si solar cells

Simple to perform, but it involves a complex synergism of many mechanisms:

- (1) dissolution of (SiN:H) antireflection coating by the constituents of the Ag ink, followed by Si-Ag interactions at the front contact;
- (2) diffusion of hydrogen from the interface into the bulk of the cell for impurity and defect passivation; and
- (3) alloying of the back Al contact to form a back-surface field (BSF).

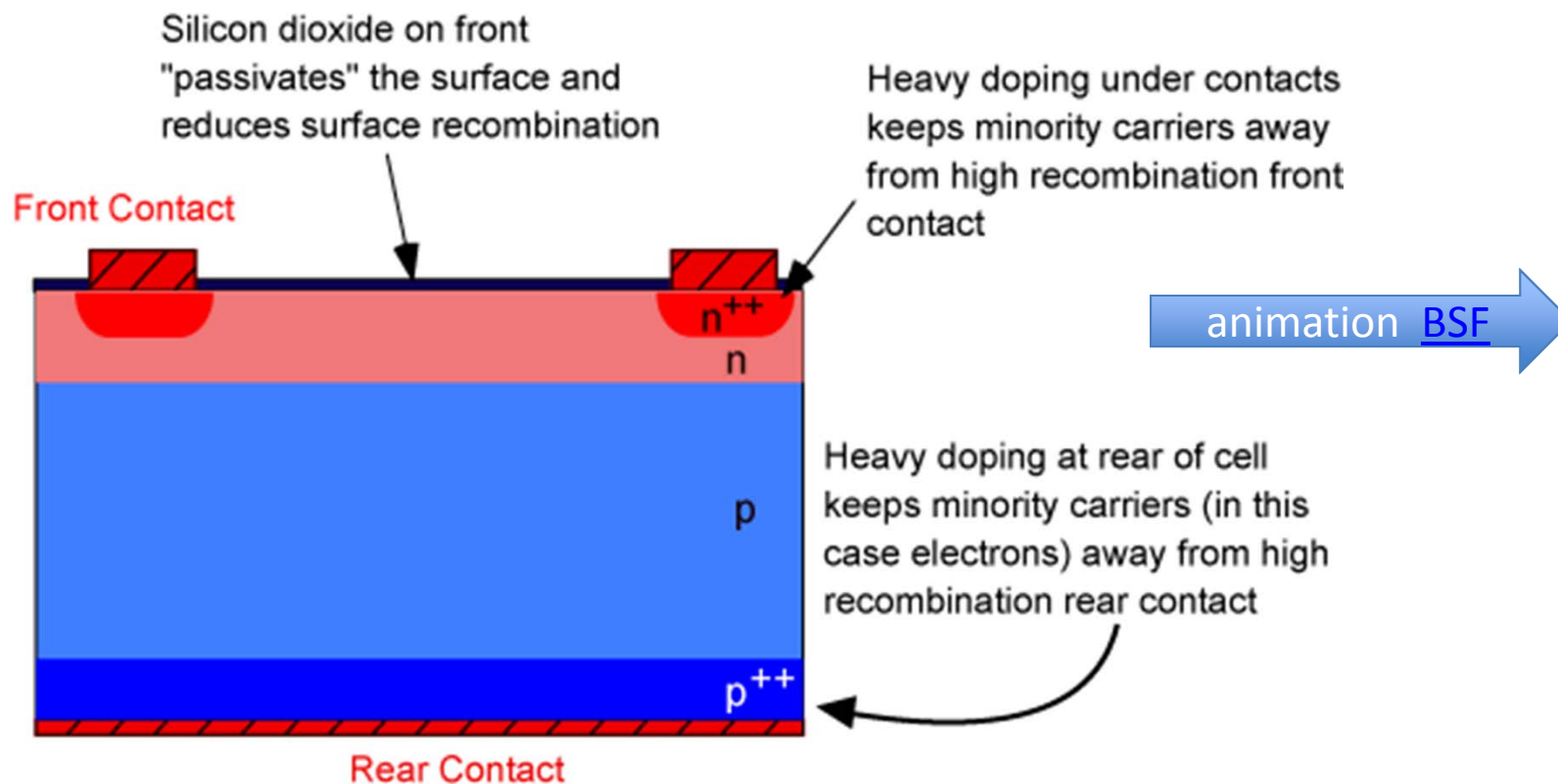


Sopori et al.

Presented at: 17th Workshop on Crystalline Silicon Solar Cells & Modules: Materials and Processes, August 5 - 8, 2007, Vail Cascade Resort, Vail, Colorado USA

BSF in Silicon PV Cells

Surface recombination can have a major impact both on the short-circuit current and on the open-circuit voltage. High recombination rates at the top surface particularly impact J_{sc} since the top surface generates a high concentration of carriers. Lowering the high top surface recombination is typically accomplished by reducing the number of dangling silicon bonds at the top surface by growing a "passivating" layer (usually silicon dioxide) on the top surface. Recombination at the back surface become increasingly important for thin cells with long minority carrier lifetimes.



“Standard” Silicon PV Cells

The industrial goal for PV power is to reduce the electricity generation cost to the equivalent of that for commercial grid electricity. The energy conversion efficiency of solar cells is another important issue because the efficiency influences the entire value-chain cost of the PV system, from material production to system installation.

The solar cell efficiency is limited by the three loss mechanisms:

- photon losses due to surface reflection, silicon bulk transmission and back contact absorption;
- minority carrier (electrons in the p region and holes in the n region) loss due to recombination in the silicon bulk and at the surface; and
- heating joule loss due to series resistance in the gridlines and busbars, at the interface between the contact and silicon, and in the silicon bulk and diffusion region.

In the design of solar cells and processes, these losses are minimized without lowering the productivity of the solar cells.

“Standard” Silicon PV Cells

The basic cell structure used in current industrial crystalline solar cells were developed for space and terrestrial use in the 1970s:

- lightly doped n+ layer (0.2–0.3 μm) for better blue-wavelength response
- a BSF formed by a p/p+ low/high junction on the rear side of the cell
- a random pyramid-structured light-trapping surface
- an ARC optimized with respect to the refractive index of the glue used to adhere to it.
- The efficiency of monocrystalline cells for space use is in the range of 14–16% under ‘1 sun’ AM0 test conditions, equivalent to 15–17% at AM1.5.
- These standard structures for crystalline silicon cells are still used in standard industrial crystalline cells, which offer efficiencies in the range of 14–17%.

The key technologies needed to realize efficiencies of higher than 20% were developed in the 1980s and '90s, and the latest high-efficiency crystalline silicon cells possess most of these features (Table 1).

Table 1. Key technologies for high-efficiency crystalline silicon solar cells

Minimizing carrier loss

Passivation of front electrode (partly in contact with highly doped silicon layer) to reduce carrier recombination under front electrode

Shallow-doped p–n junction with front surface dielectric passivation layer to reduce carrier recombination in the n⁺-doped region and at the surface; heterojunction with thin amorphous layers on a crystalline silicon base in a heterojunction cell; front surface field and surface passivation for back contact in a back-junction cell

Locally p⁺-doped back surface field and point contact structure to reduce carrier recombination in highly doped p⁺ back region

Back surface passivation by a dielectric layer or heterojunction structure to reduce back surface recombination

Minimizing photon loss

Front textured surface of random pyramid or inverted pyramid structures to reduce surface reflection loss

Single- or double-layer ARC to reduce surface reflection loss

Back-contact cell structure to reduce front contact shadow loss

Flat back surface by chemical etching of silicon to improve back reflectivity and reduce photon absorption

Back surface reflector consisted of a dielectric layer and high-reflectivity thin metal layer to reduce photon absorption

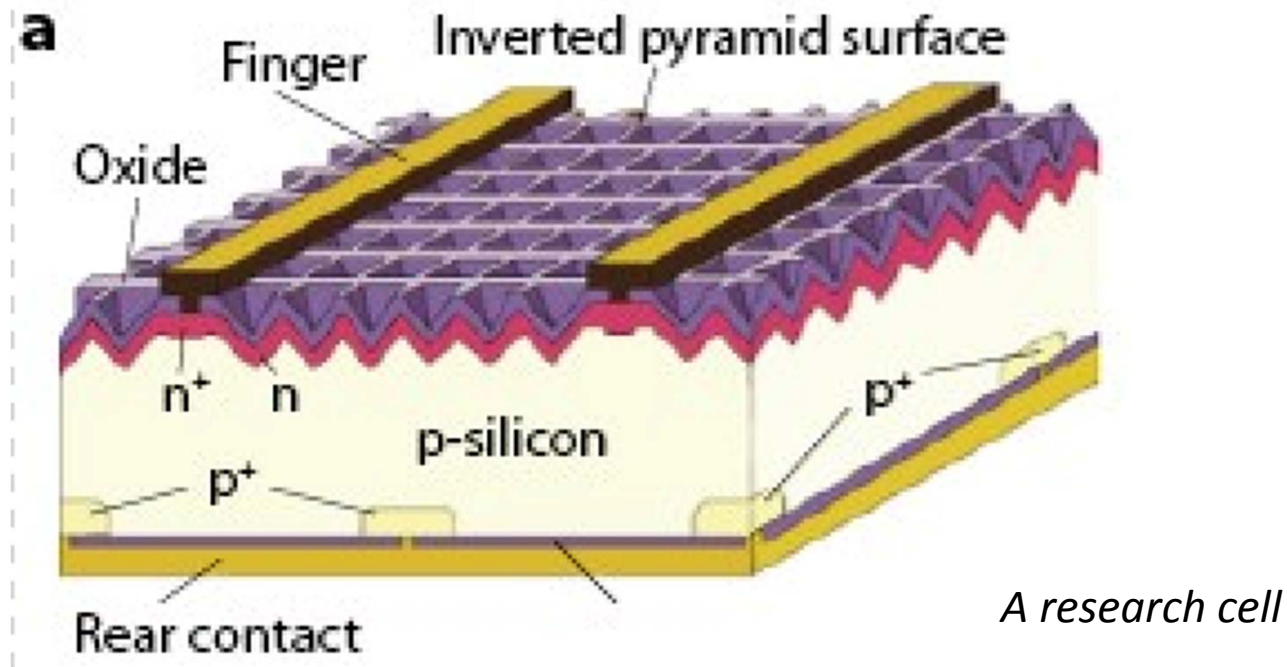
Minimizing electrical loss

Fine gridline front contact to reduce series resistance of n^+ -doped region

Selective emitter (deep and highly doped emitter under the contact) to reduce contact resistance of front contact with silicon surface

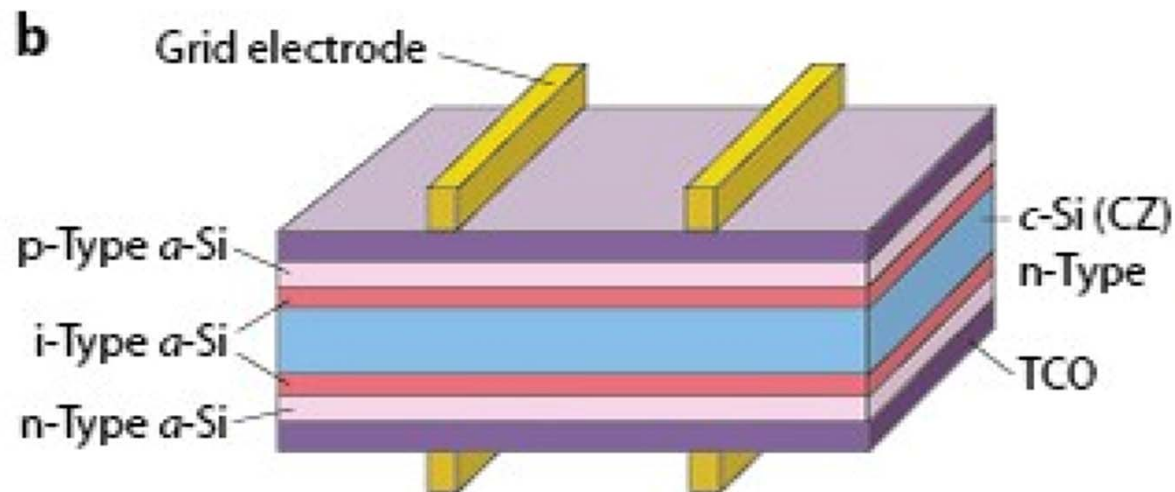
n-Type or p-type silicon substrates with minority carrier diffusion lengths longer than the base thickness

Passivated Emitter Rear Localized (PERL) cell:



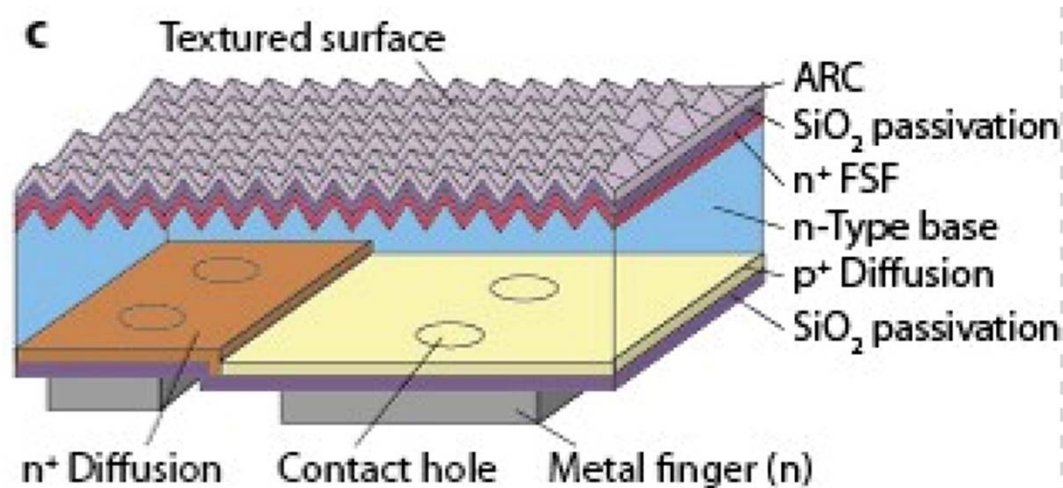
Front and rear surface passivation, inverted-pyramid light-trapping surface, a rear localized p⁺ layer (BSF), a double-layer ARC, p-type float zone monocrystalline substrate. The bulk minority carrier lifetime > 1 ms. $V_{oc} = 706$ mV, $J_{sc} = 42.7$ mAcm⁻², FF = 0.828 and $\eta = 25.0\%$ for a 4 cm² cell [4]. J_{sc} is close to limit for absorption/collection in the emitter and base. $\eta = 24.7\%$ was reported almost 10 years ago, and record of 25.0% (University of New South Wales - UNSW) in 2009 was from remeasurement of the same cell. Full PERL design is not easy to apply to low-cost industrial production because of the necessity for multiple photolithography steps. Expensive silicon PV cells for space applications have a similar structure to the PERL cell.

Heterojunction with intrinsic thin layer (HIT) cell:



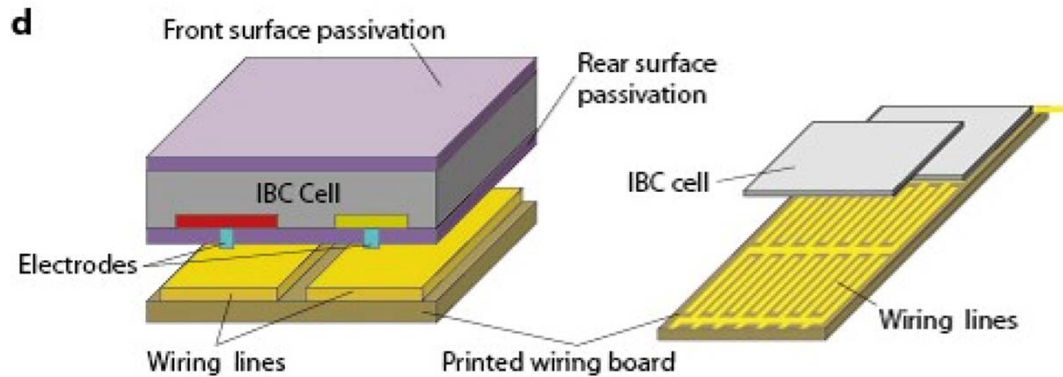
Developed for manufacturing; has thin amorphous p- and n- layers and intrinsic amorphous layers on the front and rear surfaces of a CZ n-type monocrystalline-silicon substrate. Best output parameters are $V_{oc} = 729$ mV, $J_{sc} = 39.5$ mA-cm⁻², FF = 0.800, and $\eta = 23.0\%$ for a large 100.4 cm² cell. V_{oc} is improved by large bandgap of the front amorphous silicon layer and the high interface quality between the a-Si and x-Si substrate. Low temperature coefficient of 0.30% K⁻¹ at P_{max} is advantage ($\sim 0.45\%$ K⁻¹ for other x-Si cells). TCO ARC reduces sheet resistivity of front a-Si layers. Lower J_{sc} in comparison to other high-efficiency Si cells due to weaker blue response. Generation/collection in the front a-Si layers and bulk Si is lower due by the effects of the lower transparency of the TCO layer compared to other ARCs and/or the lower internal quantum efficiency of the amorphous layers.

Back Contact Back Junction (BC-BJ) cell:



Interdigitated n- and p-doped regions and n and p contacts on the back surface. Original BC-BJ cell, called the FSF cell or interdigitated back contact (IBC) cell developed for space PV in late 1970s. The BC-BJ-structured point contact (PC) cell developed by Stanford University in the 1980s had $\eta >$ than 20%. SunPower made BC-BJ cells for unmanned aircraft and solar race cars in the 1990s, followed by large-scale PV plants in 2000s. Best efficiency for a large-area industrial BC-BJ cell is 23.4%. The BC-BJ cell has front and rear surface passivation layers, a random-pyramid light-trapping surface, FSF, interdigitated n- and p-doped regions on the back surface, n and p contact gridlines on n- and p-doped regions, a single-layer ARC and CZ n-type single-crystalline silicon substrate with a minority carrier lifetime > 1 ms. Of all silicon PV modules on market at this time, only those based on BC-BJ cells provide the possibility of module efficiencies exceeding 20%. BC-BJ cells have no gridlines or busbars shading, a front surface with good passivation and improved aesthetics due to the absence of front electrodes, which permits freedom in the design of back contacts.

T. Saga, NPG Asia Mater. 2(3) 96–102 (2010)



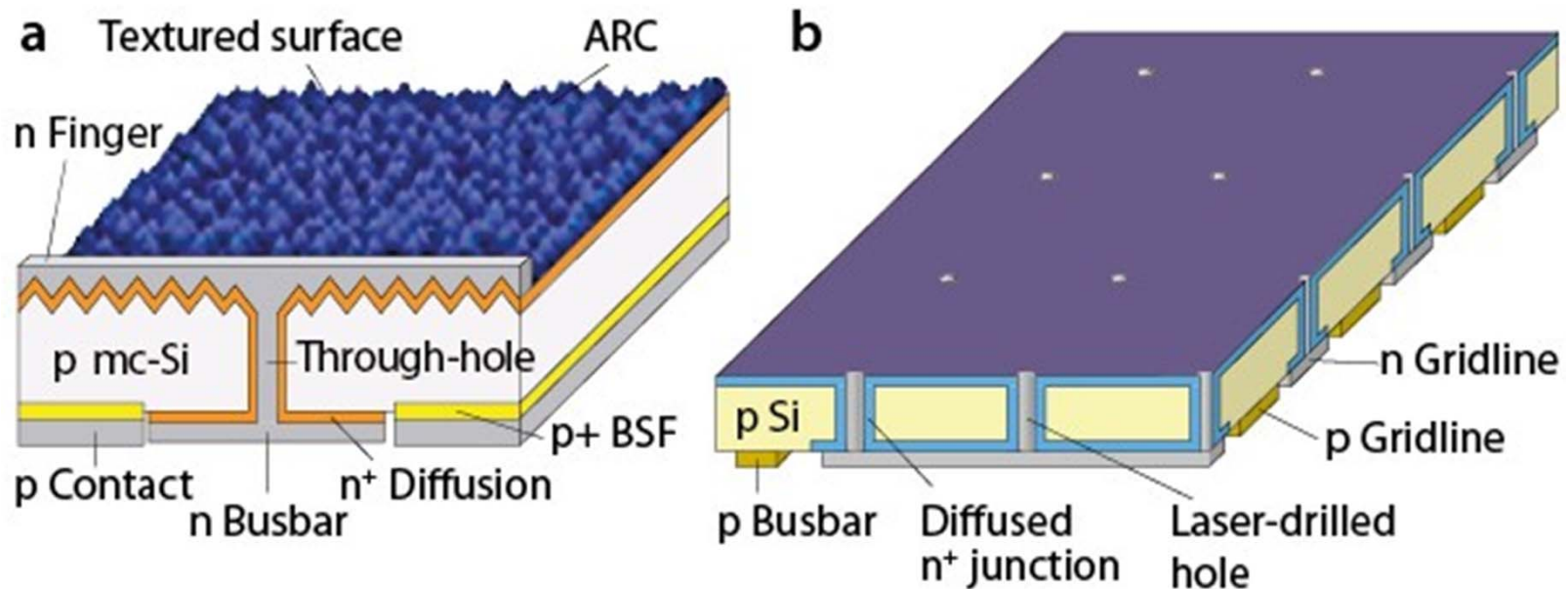
57 K panels, 100 acres, 10 MW, largest in the world when completed in 2004

Freedom in design of back contact also provides advantages in module assembly, allowing the simultaneous interconnection of all cells on a flexible printed circuit. The low series resistance of interconnection formed by this type of surface-mount technology results in a high FF of 0.800 , compared with around 0.75 for standard silicon PV cell modules.

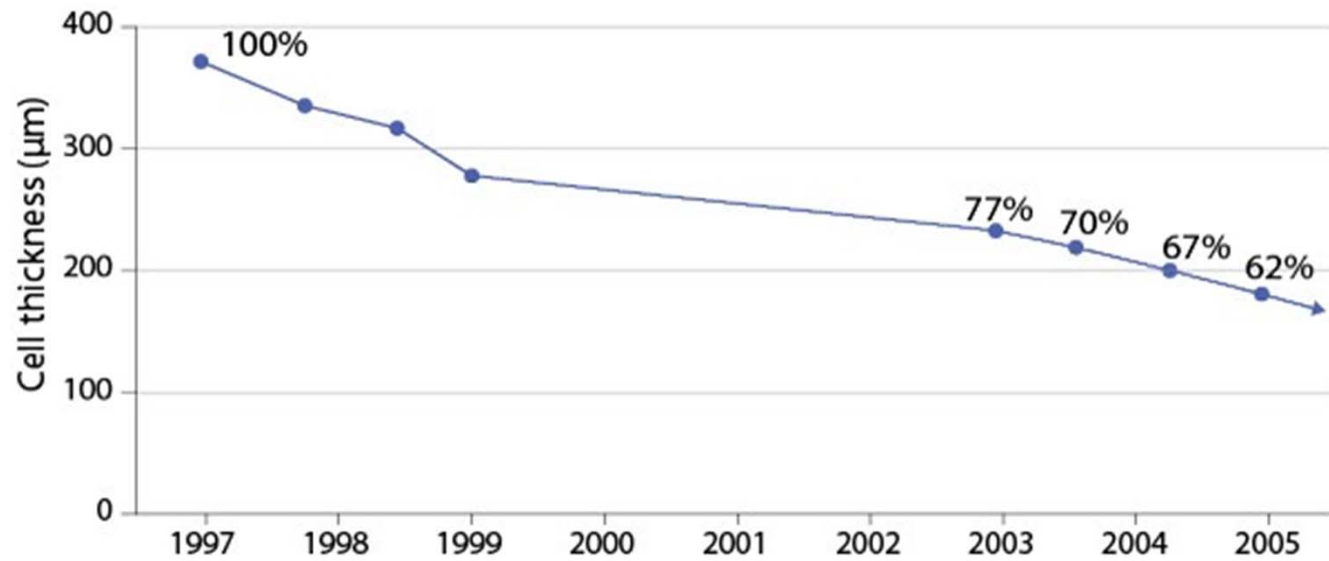


Emitter Wrap Through and Metal Wrap Through Cells

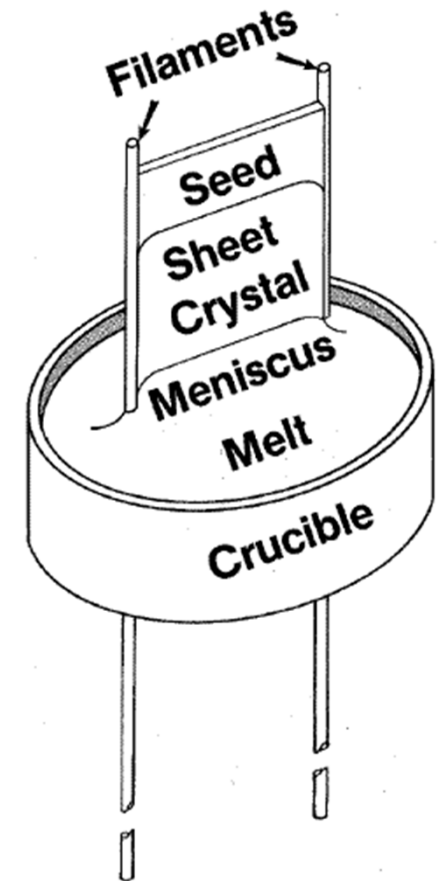
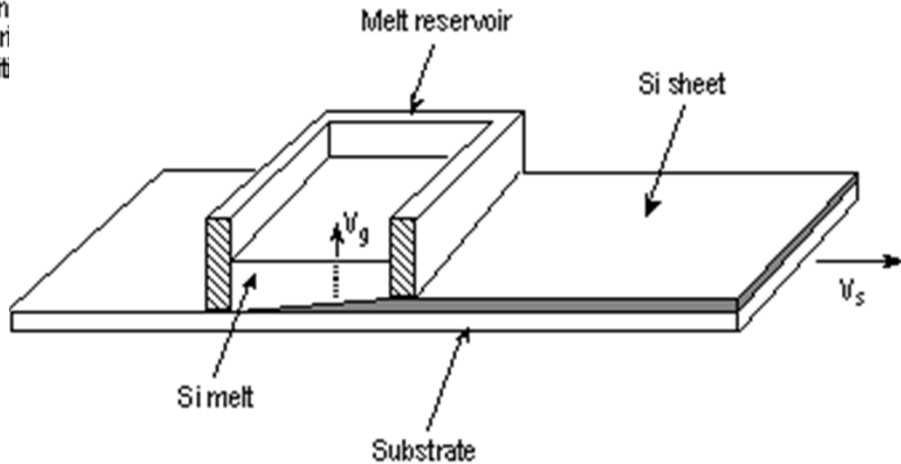
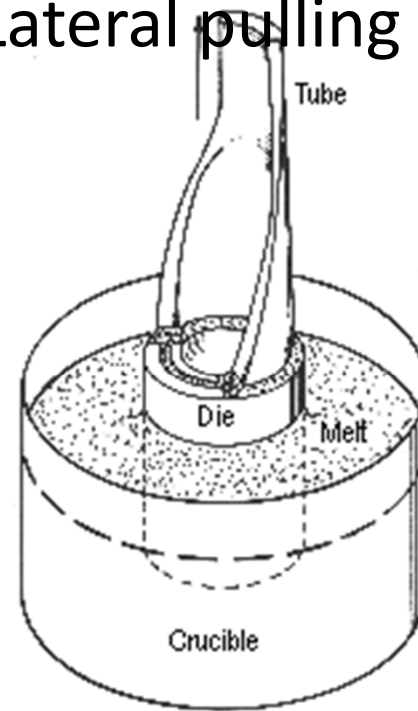
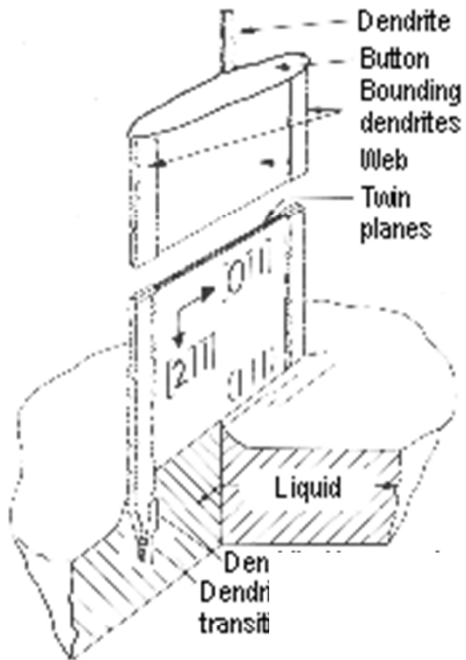
- Schematics of back-contact solar cell structures. (a) MWT. Adapted from Ref. 26 (© 2008 WIP Munich). (b) EWT. Adapted from Ref. 25 (© 2008 IEEE). All figures reproduced with permission.



Reducing the amount of Si in the cell, by reducing the silicon wafer thickness (© 2010 Sharp)



Edge Defined Growth and Lateral pulling

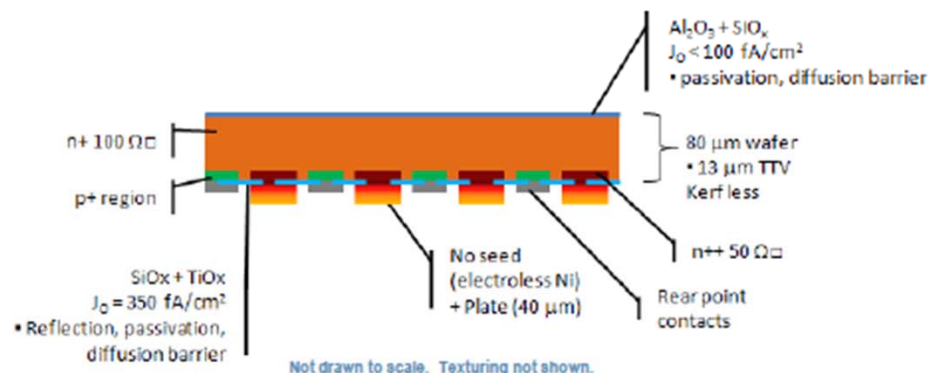


c-Si Cell Description: 2030

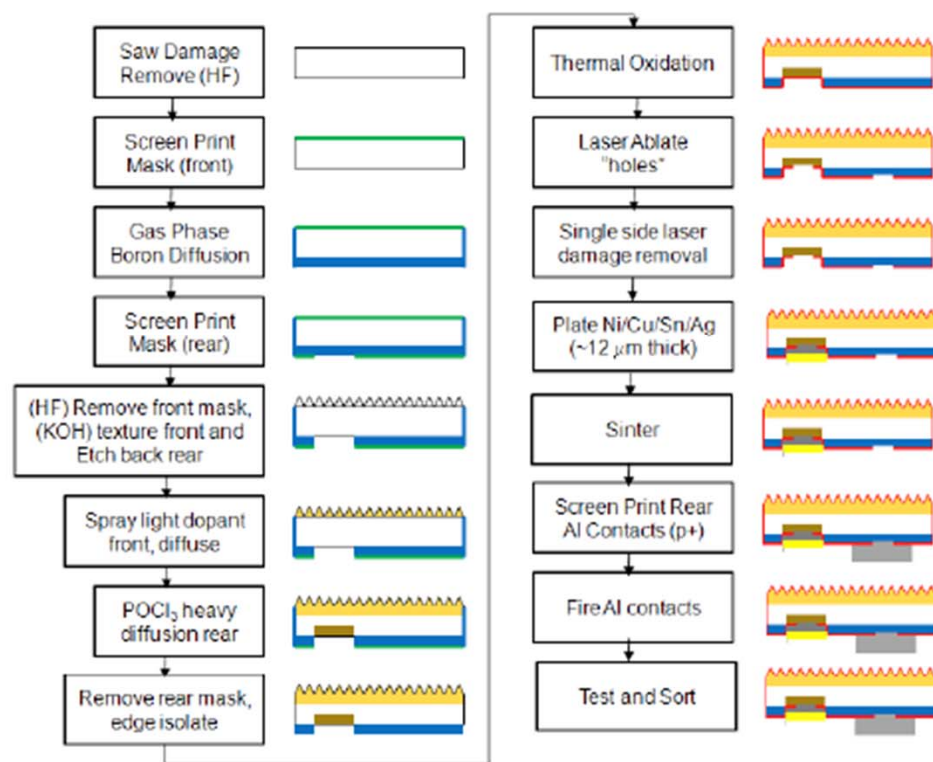
*Based on publicly disclosed (literature) cell designs, not intended to depict proprietary architectures

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ENERGY

Energy Efficiency &
Renewable Energy



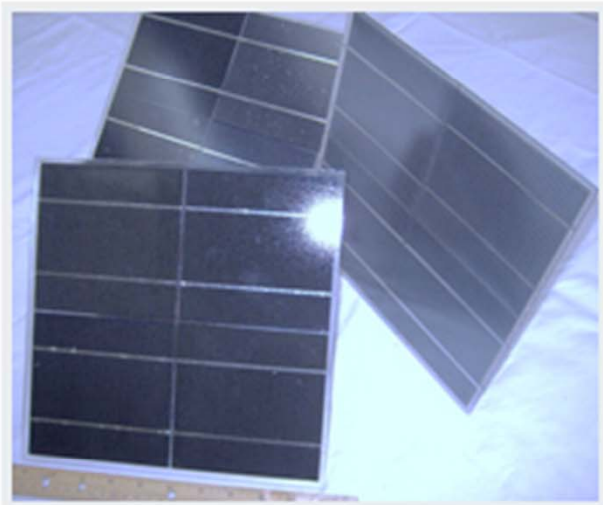
- All Rear (Interdigitated) Contacts
- High lifetime (n- type) wafer
- Ultra thin (80 microns) kerfless wafers
- High quality surface passivation
- Plated emitter contacts
 - Electroless nickel barrier, Cu plating
- Base point contact absorbers
 - Printed Al contacts



ENERGY

Silicon Solar Cells Ditch the Wafers

A startup's silicon solar cells save on costs by using less material.



Skinny silicon: These solar modules, made from silicon films just 40 micrometers thick, have a 15 percent efficiency.

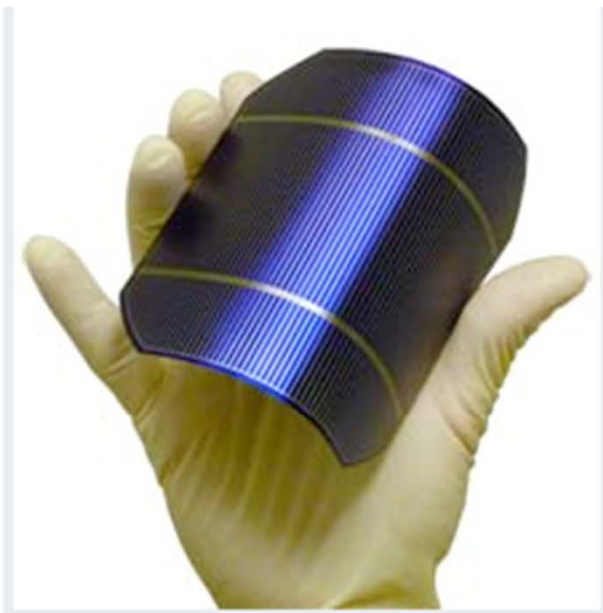
Credit: Crystal Solar

ENERGY

Nano Ink Boost for Silicon Solar

Inkjet-printed silicon increases solar-cell efficiency.

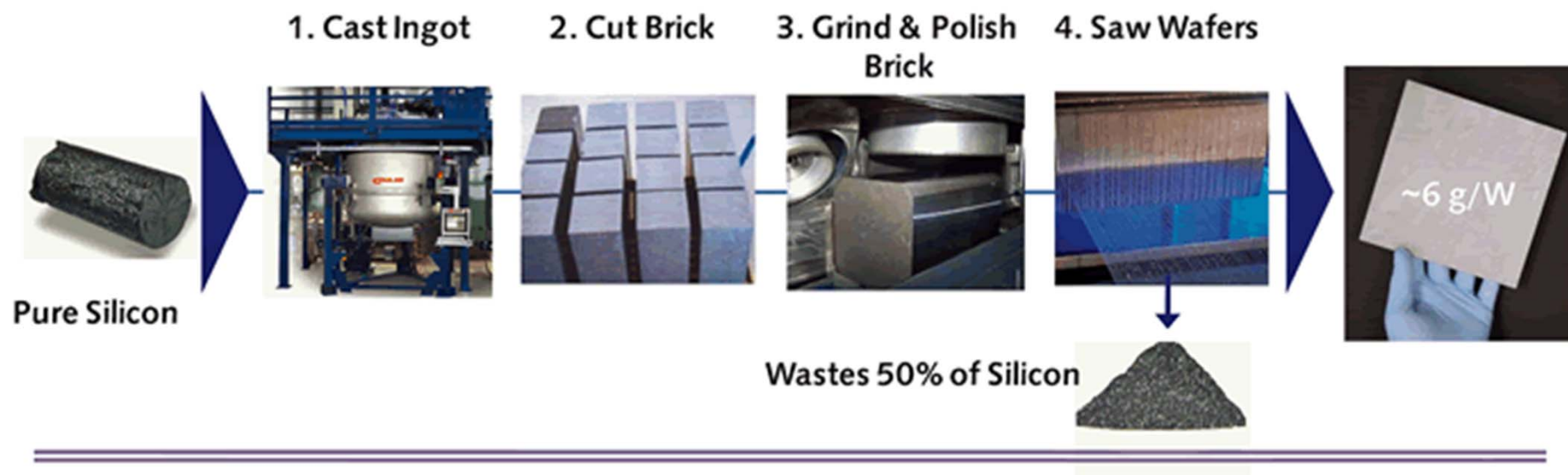
FRIDAY, SEPTEMBER 18, 2009 | BY KATHERINE BOURZAC



Inkjet solar: The inkjet printing process allows Innovalight to make silicon wafers that are thin enough to bend.

Credit: Innovalight

Standard Process



1366 Direct Wafer

