

Lecture 6 Junction characterisation

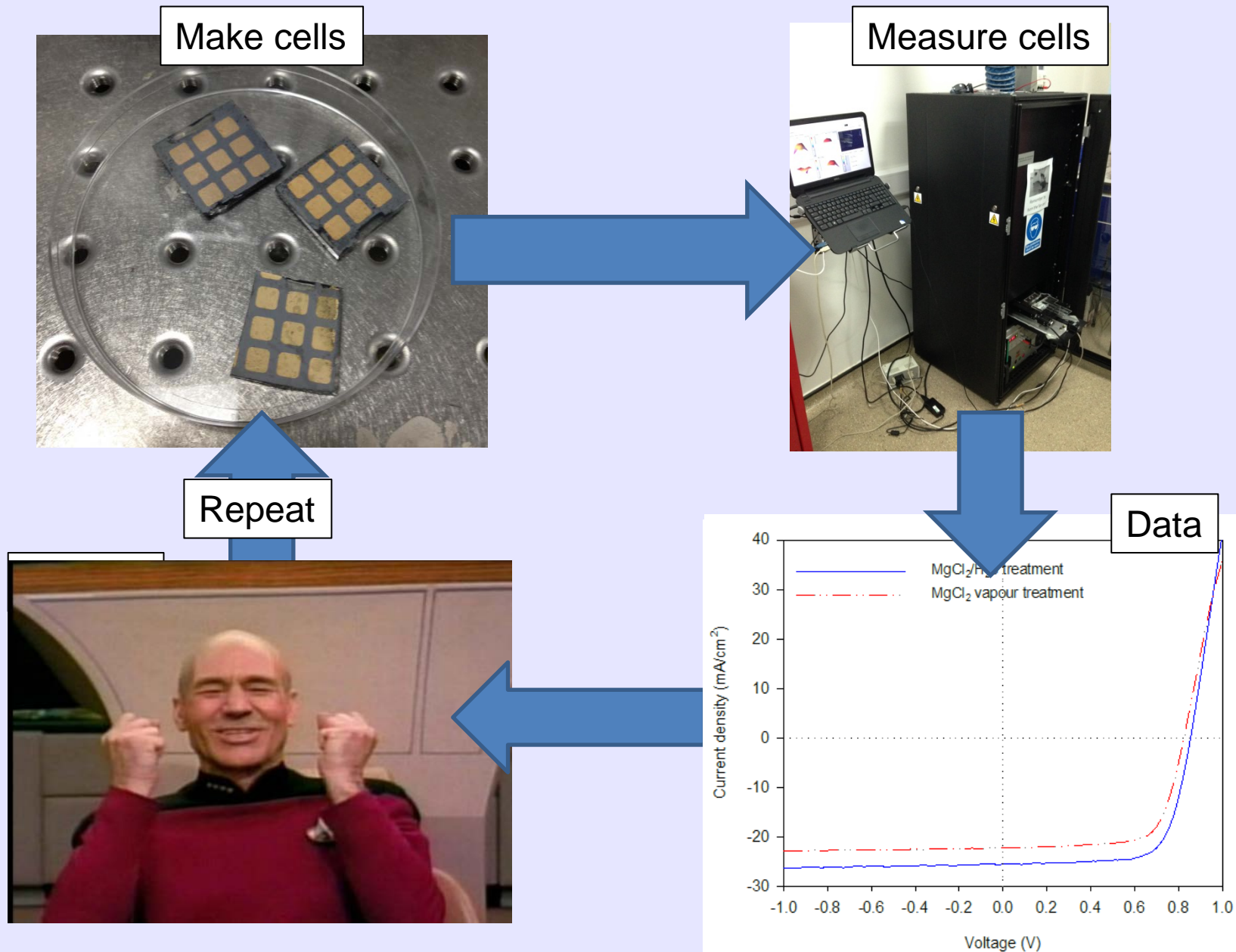
Jon Major

Nov 6th 2014





The PV research cycle





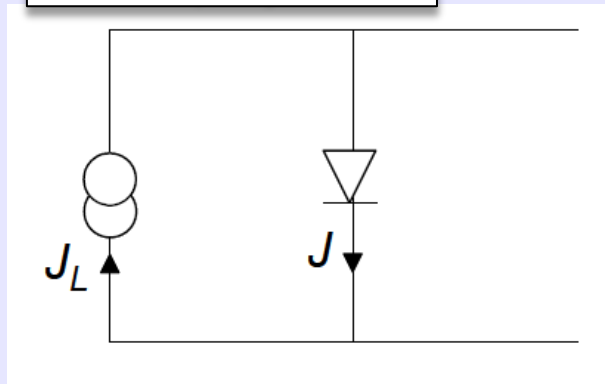
Key junction characterisation techniques

- Analysis of current voltage (J-V) curves
- External quantum efficiency (EQE) measurements
- Capacitance voltage (C-V) measurements

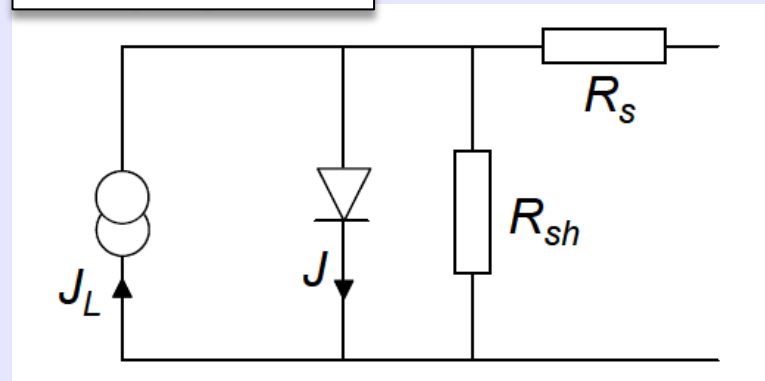


Parasitic resistances

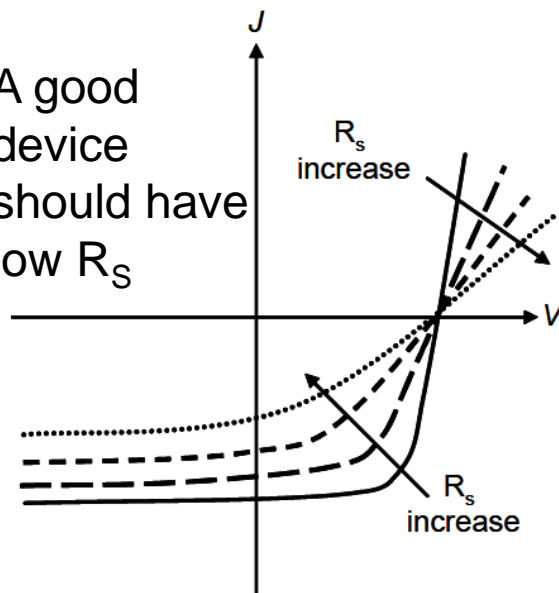
Perfect p-n junction



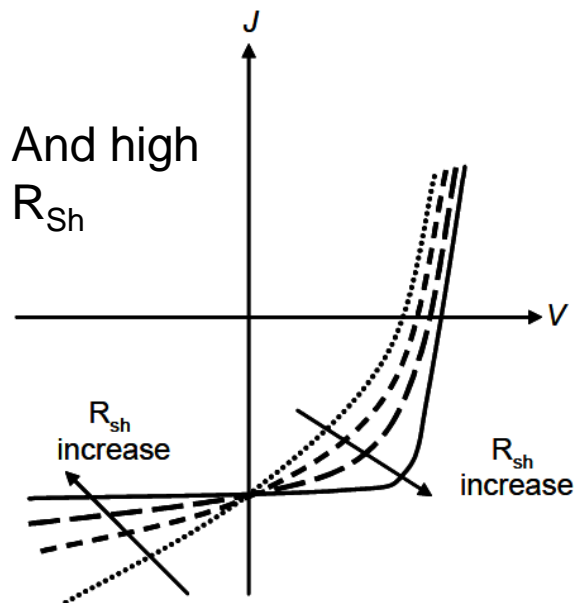
Real p-n junction

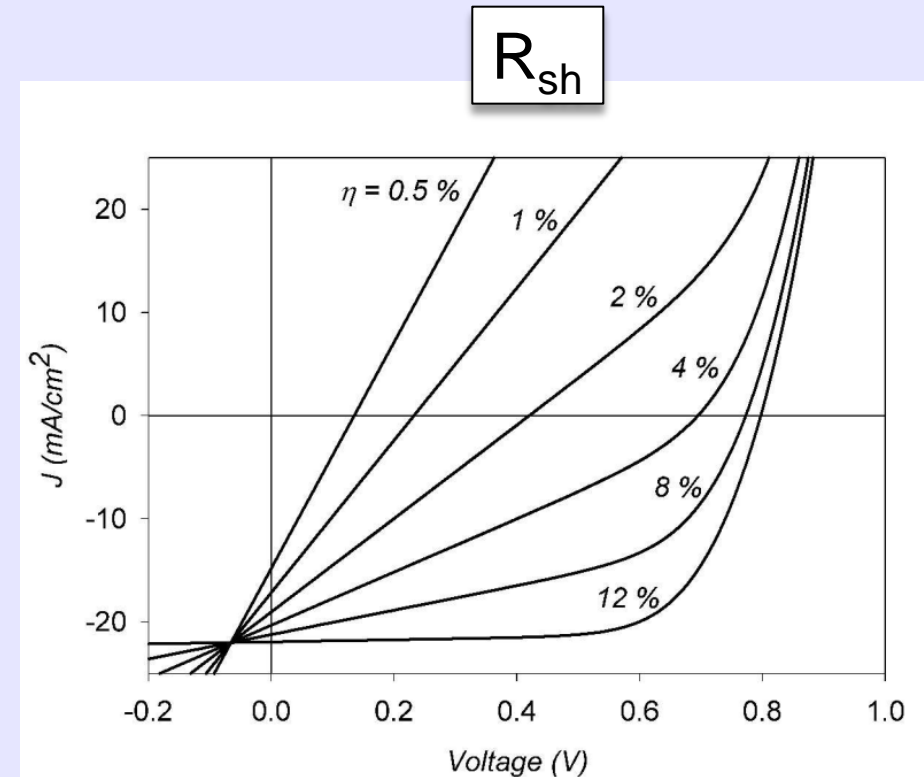
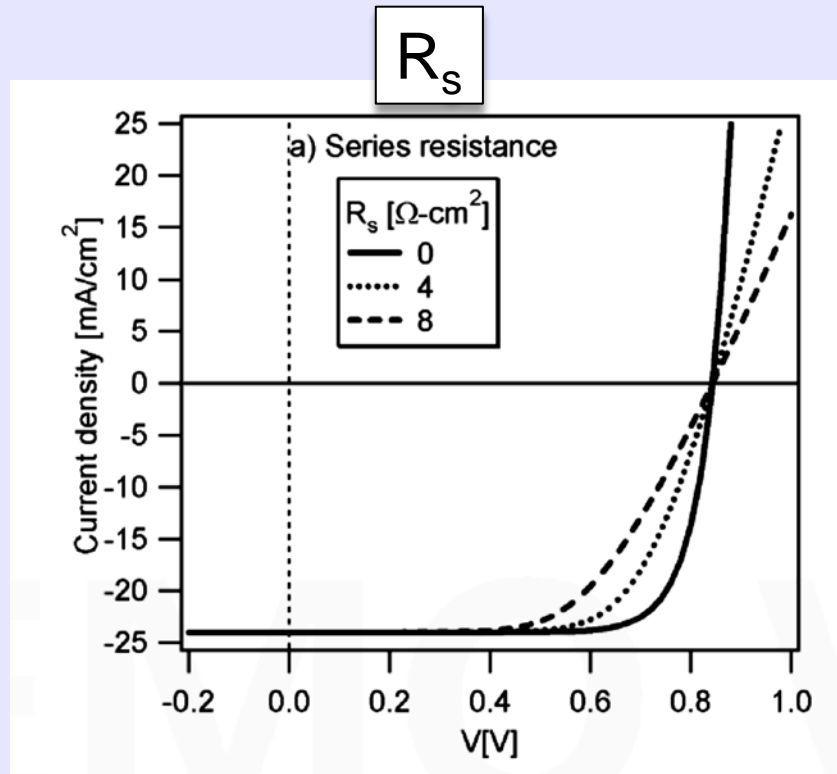


A good device should have low R_s



And high R_{sh}



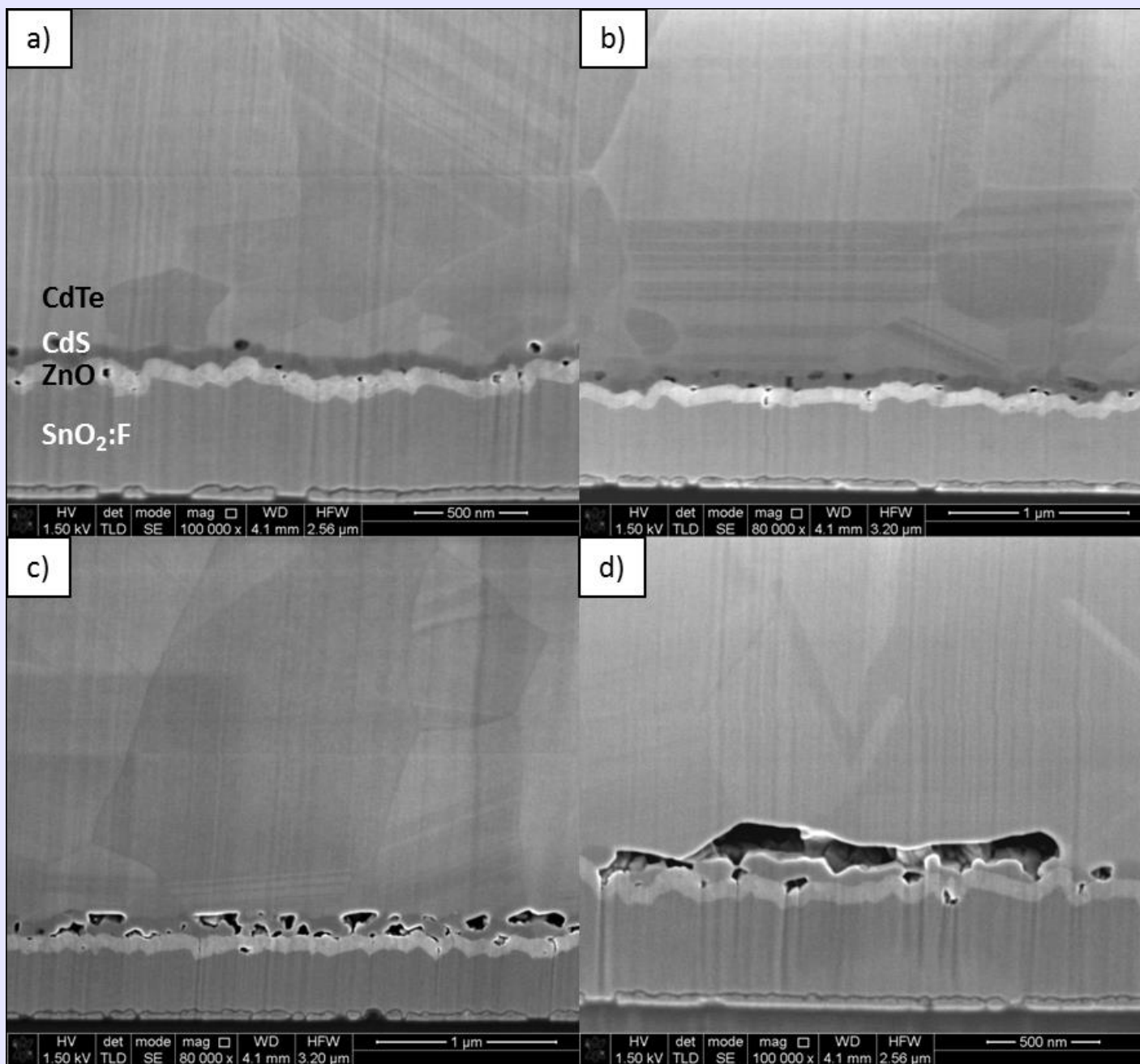


Causes of high R_s values

- Overly thick absorber layer
- Low conductivity TCO
- Low doping levels

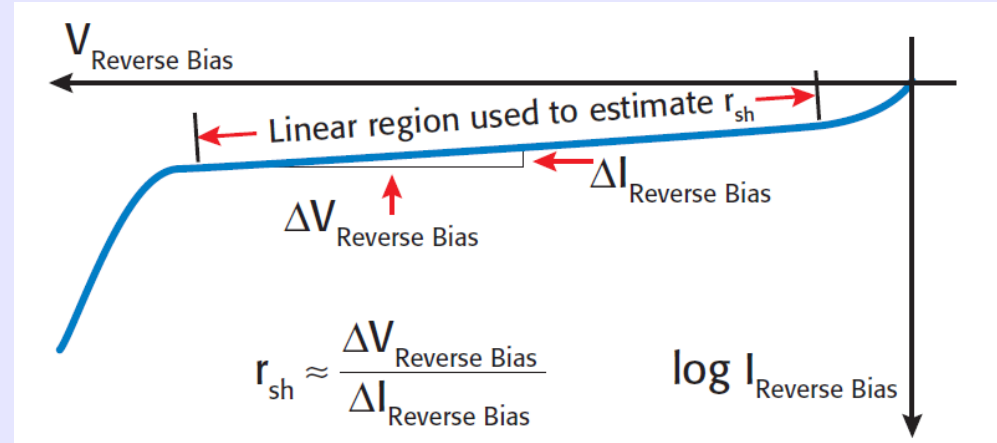
Causes of low R_{sh} values

- Pinholes in layers
- Weak diode regions

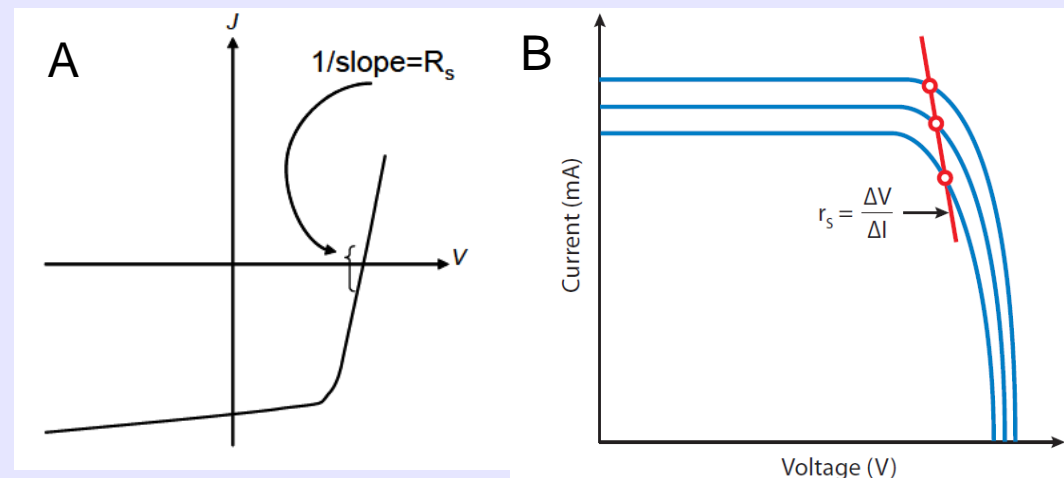




To determine R_{sh} fit to straight line portion of JV curve in reverse bias

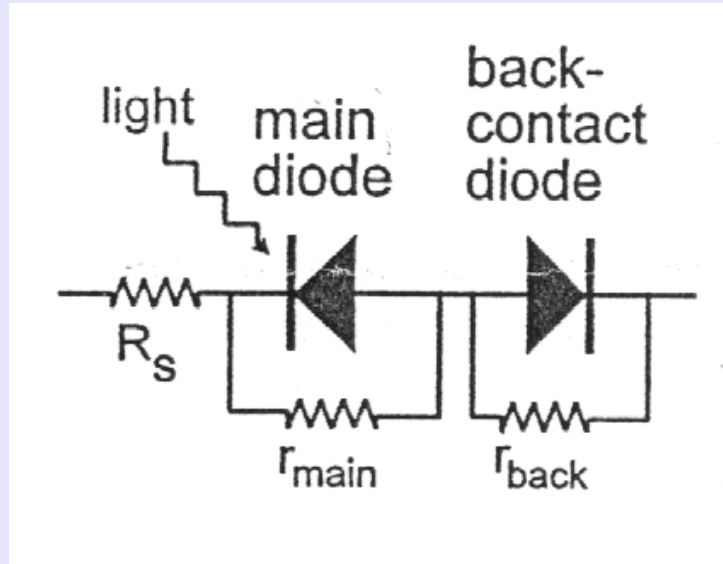


R_s can be more difficult to accurately determine. Two common methods



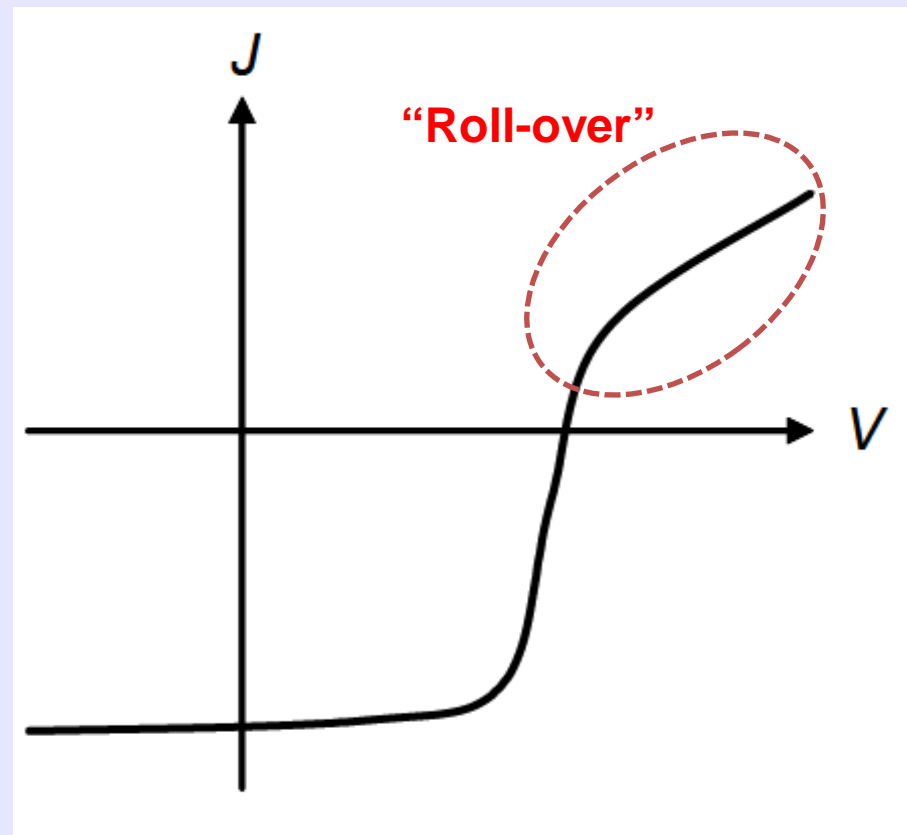
Fit to forward bias region of curve

Measure change as function of light bias



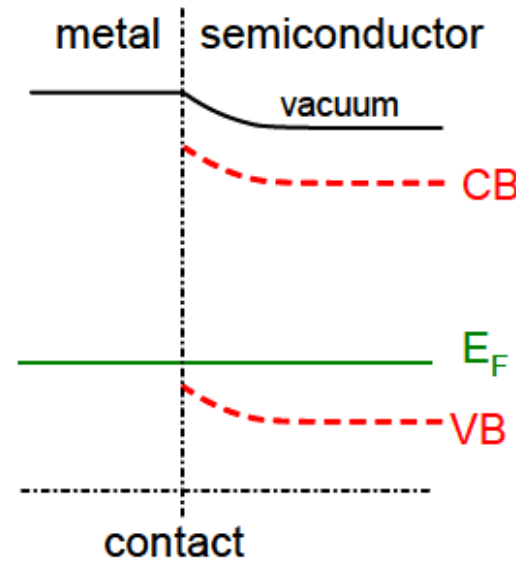
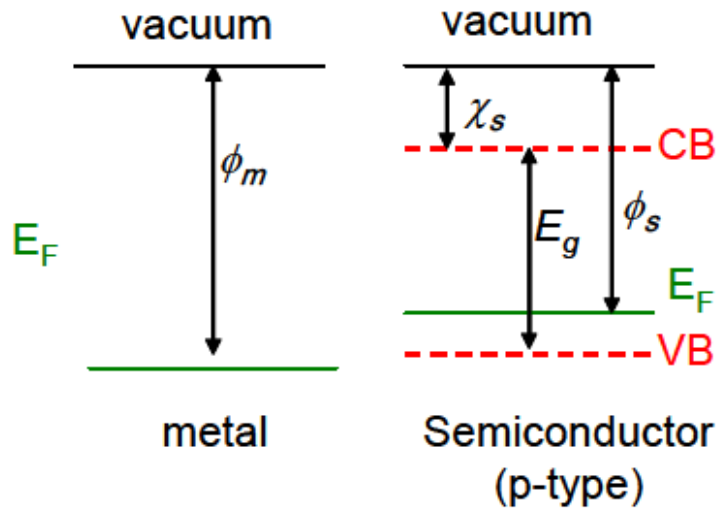
If a non-ohmic contact is formed this results in the creation of a back contact junction diode.

This back-contact diode opposes the main junction diode and leads to the phenomenon of “roll-over” at high forward bias.





Ohmic contact



For a p-type semiconductor can create an ohmic contact if;

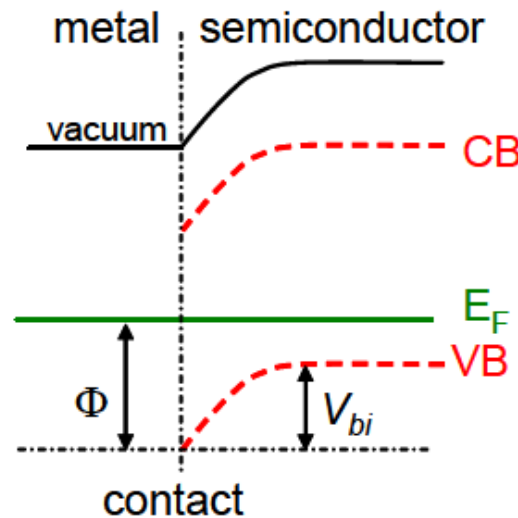
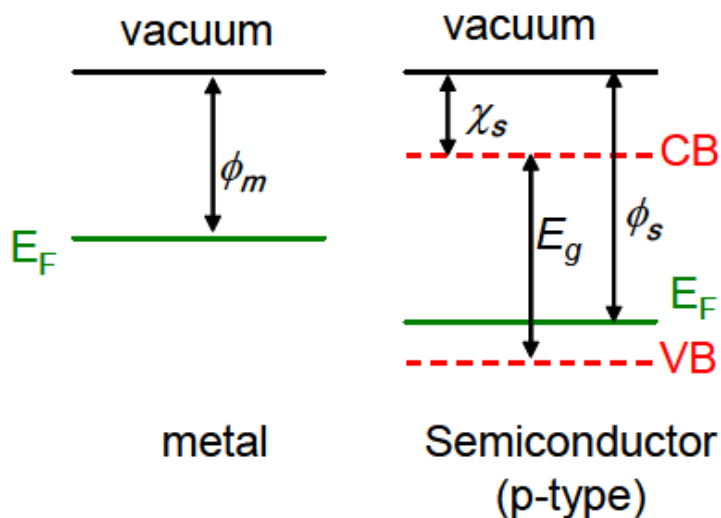
$$\phi_m \geq \chi_s + E_g$$

Metal work function

Electron affinity

Bandgap

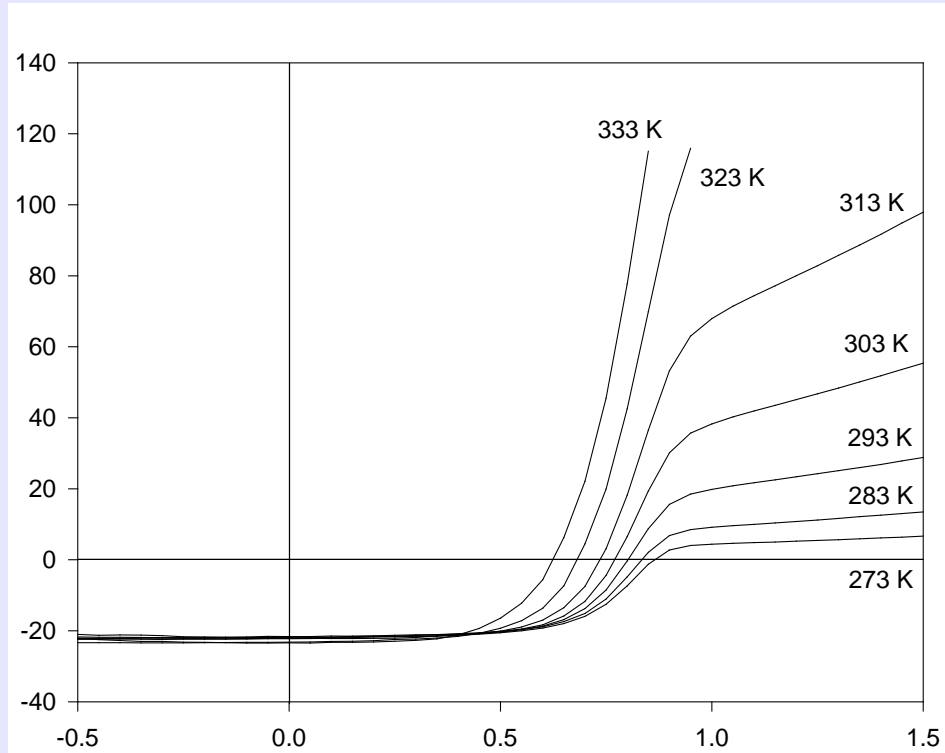
Schottky barrier



If this condition is not met we introduce a back contact Schottky barrier

$$\Phi \approx \phi_m - (\chi_s + E_g)$$

Back contact barrier height



Influence of barrier height changes as a function of temperature

By measuring J-V curves as a function of temperature we can extract the barrier height

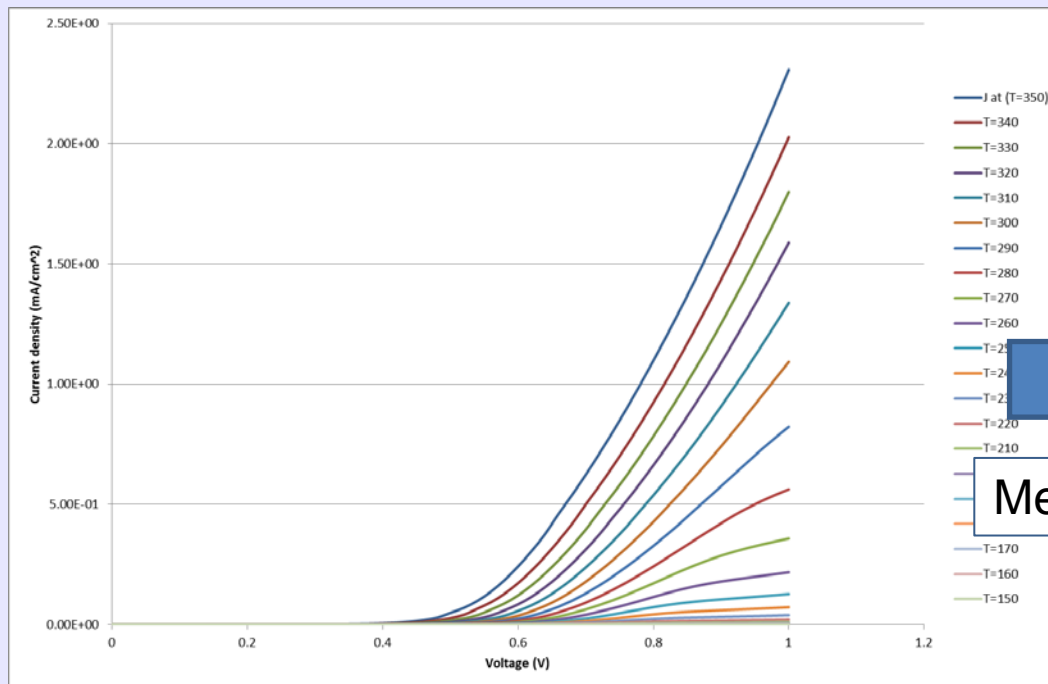


Series resistance varies with temperature via

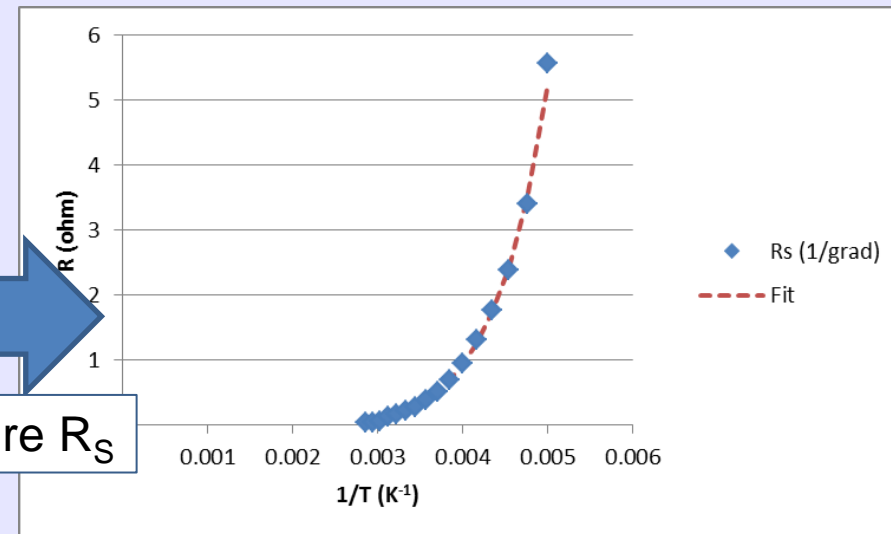
$$R_S = R_{\Omega 0} + \frac{\partial R_{\Omega 0}}{\partial T} T + \frac{C}{T^2} \exp\left(\frac{\Phi_b}{kT}\right)$$

Ohmic resistance

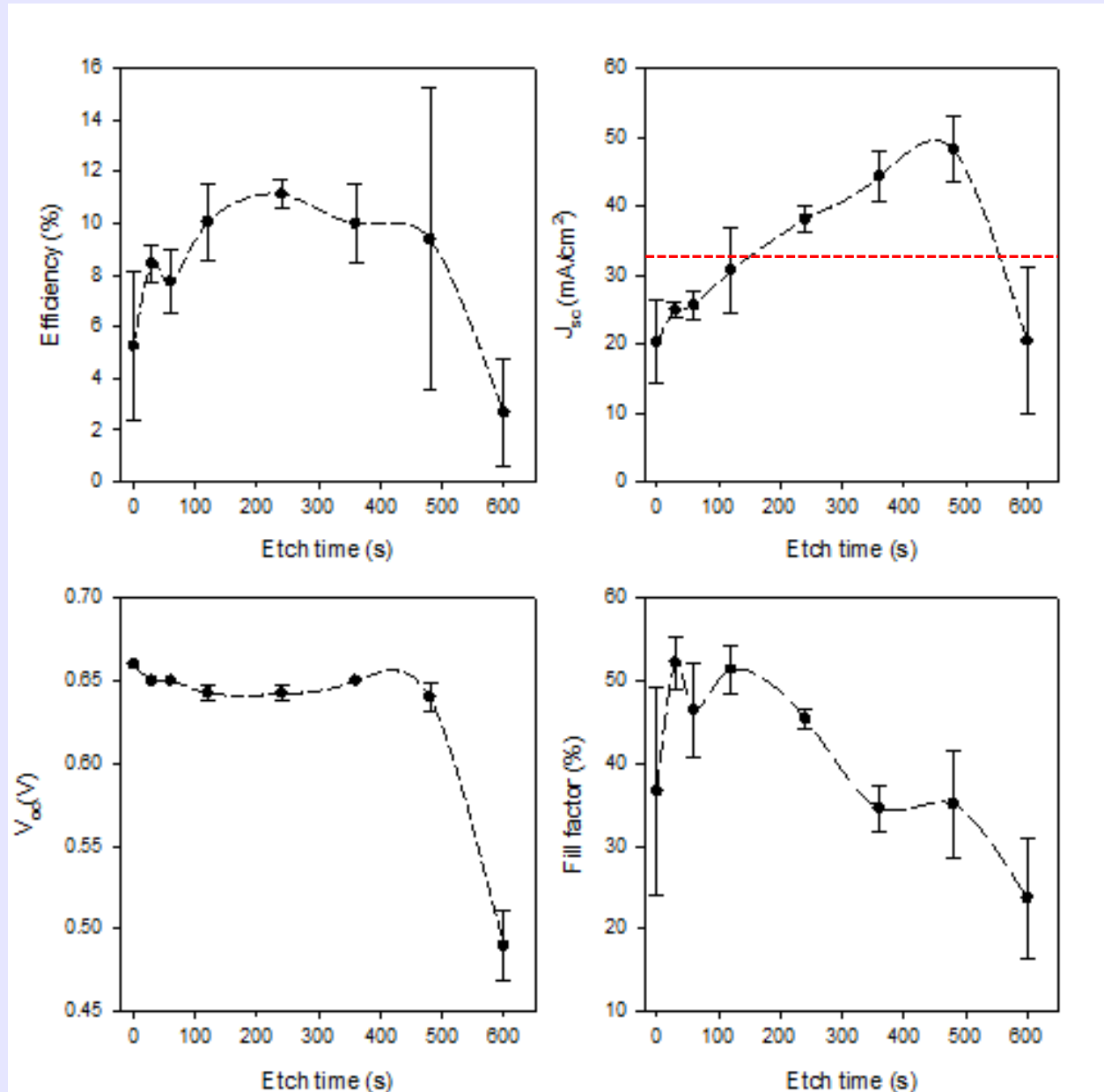
Ohmic temperature dependence



Measure R_S

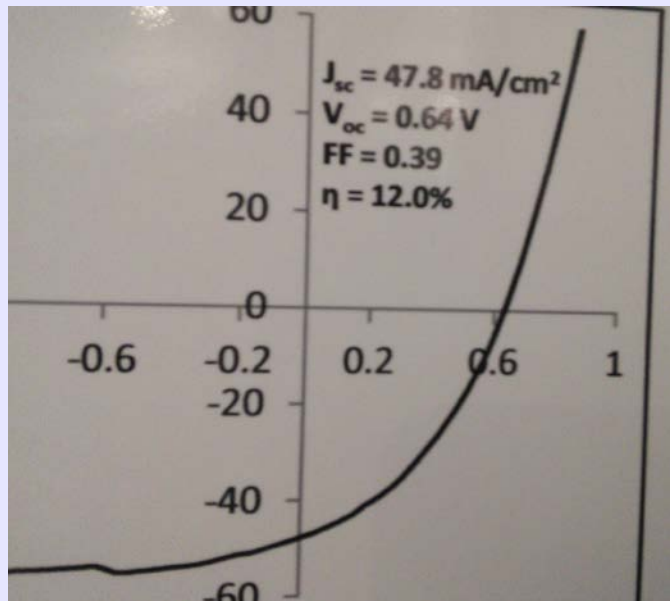
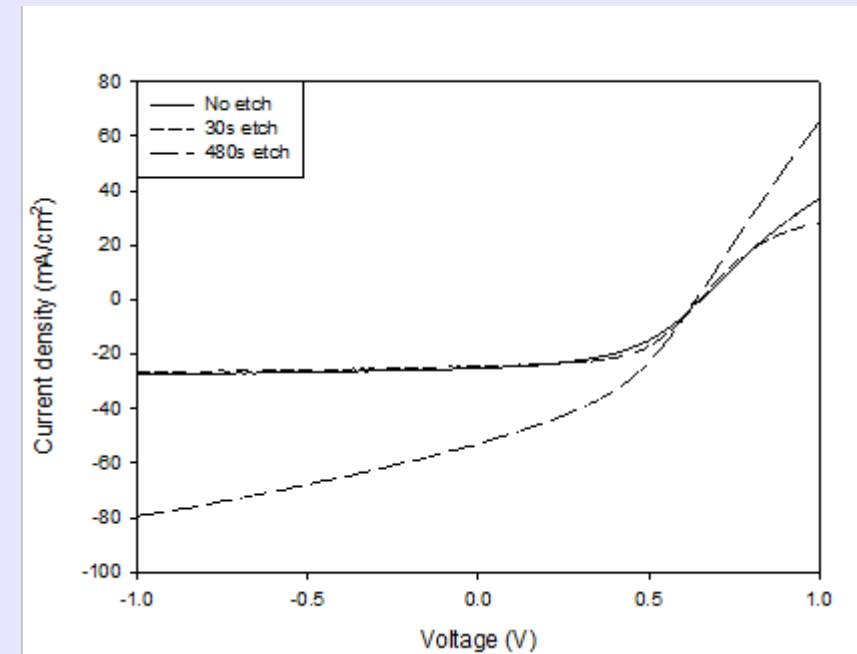
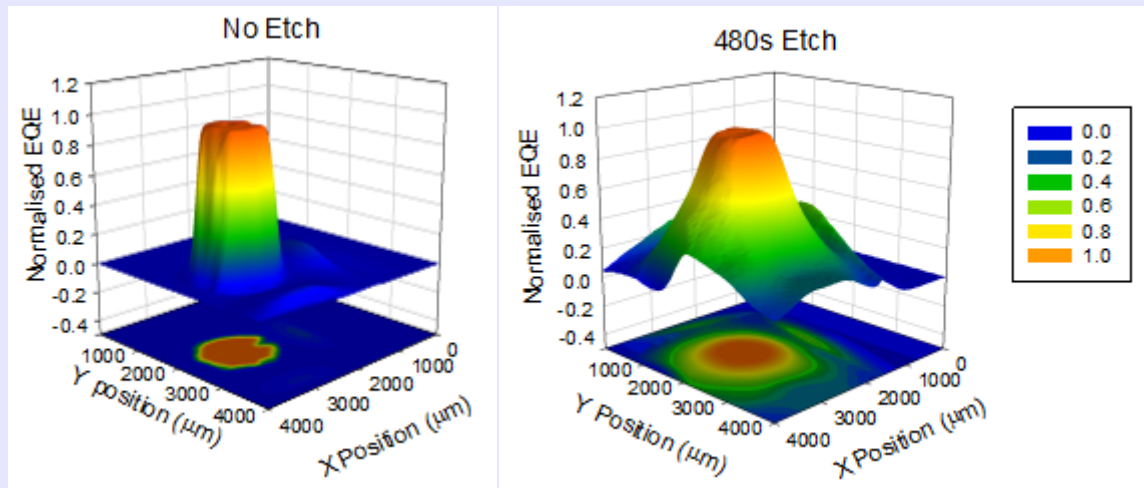


Fitting to exponential region allows us to extract a value in eV for the barrier height. Generally anything < 0.3eV is considered a good contact





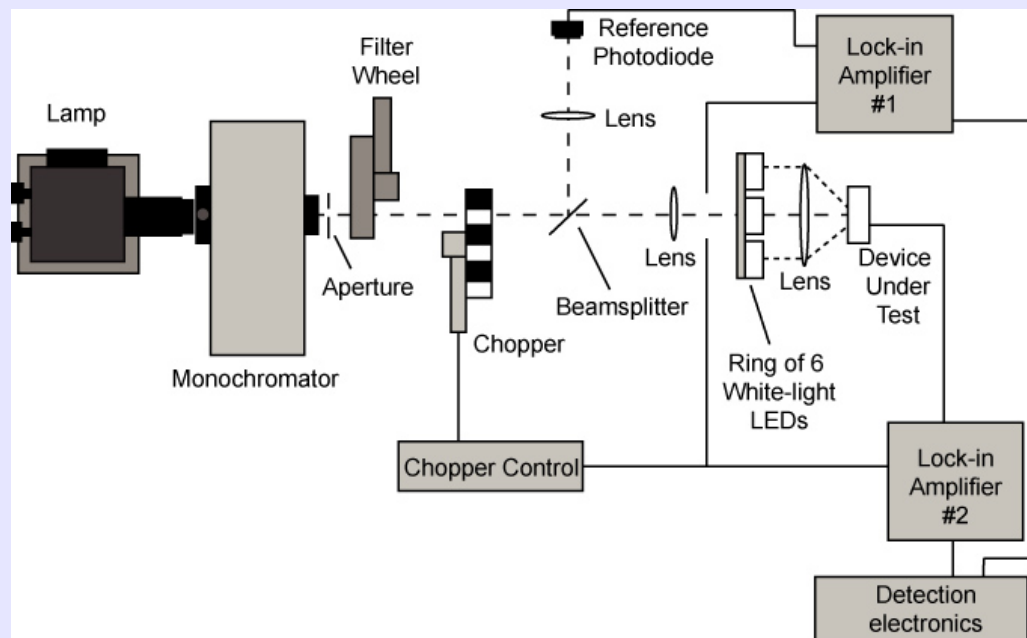
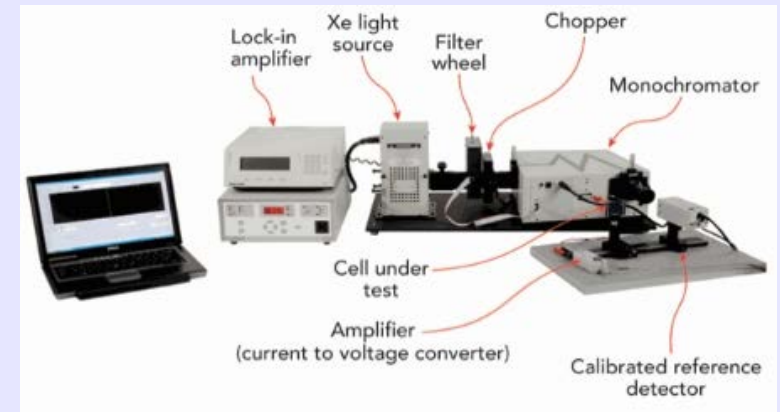
Errors in J-V analysis (J-V)



- High V_{oc} and FF are reliable
- J_{sc} values are very sensitive to calibration or contact size errors
- Contacts should be minimum of 0.25cm^2
- If it looks too good to be true it usually is!

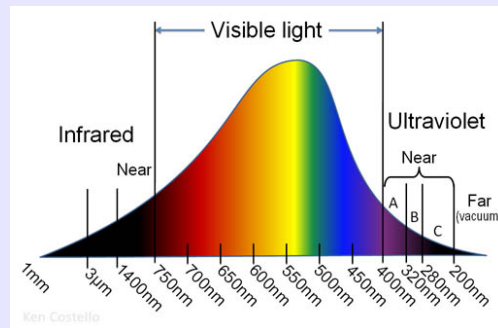
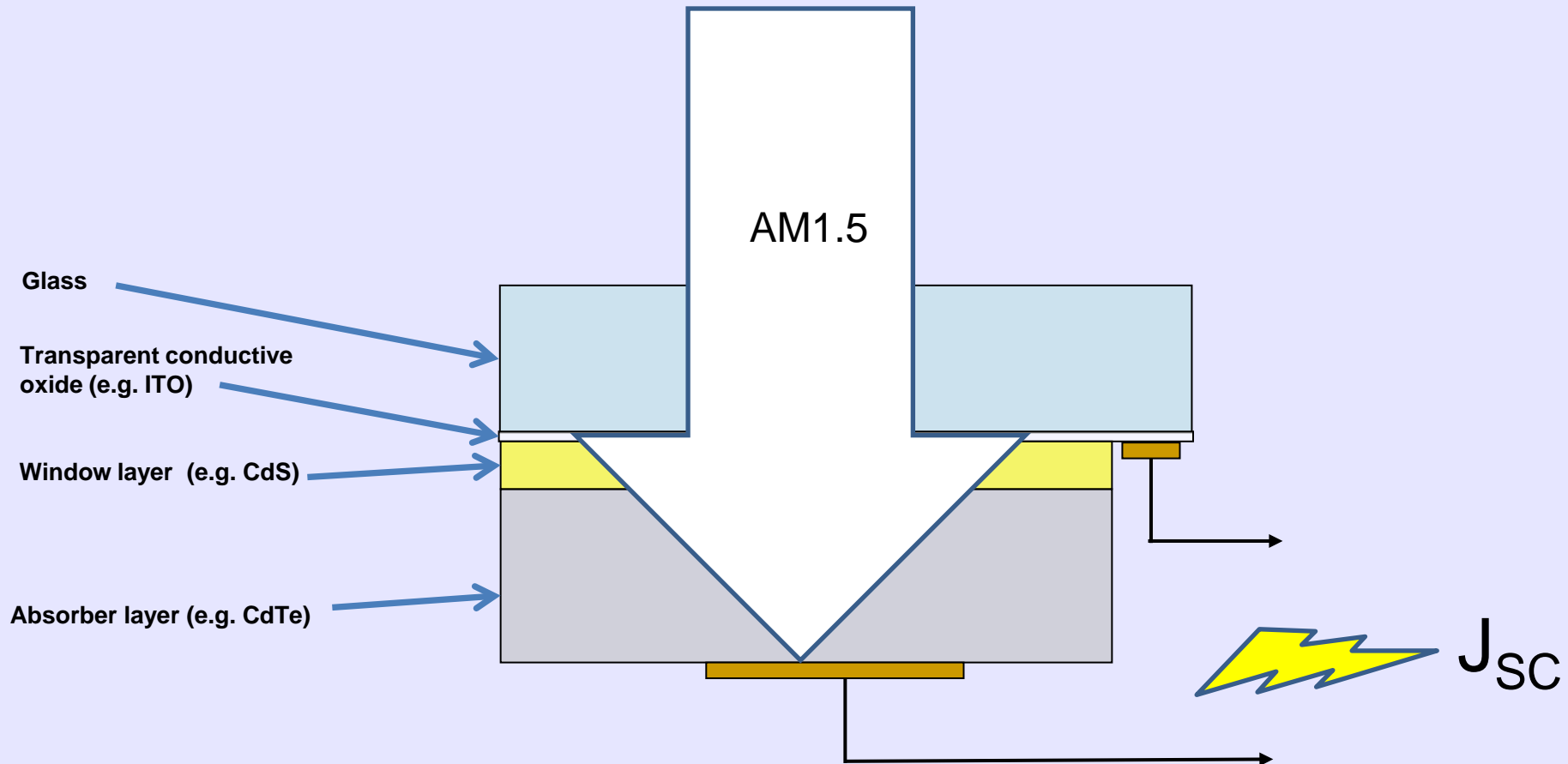


External quantum efficiency (EQE)





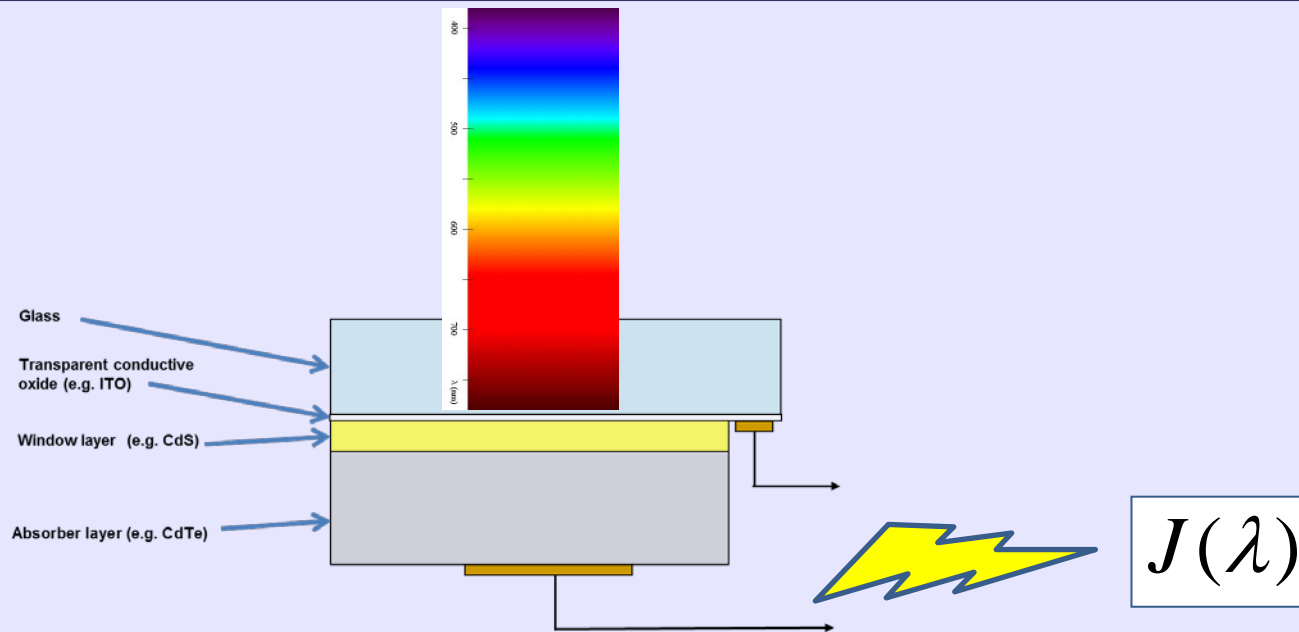
External quantum efficiency (EQE)



$$E(\lambda) = \frac{hc}{\lambda}$$



External quantum efficiency (EQE)



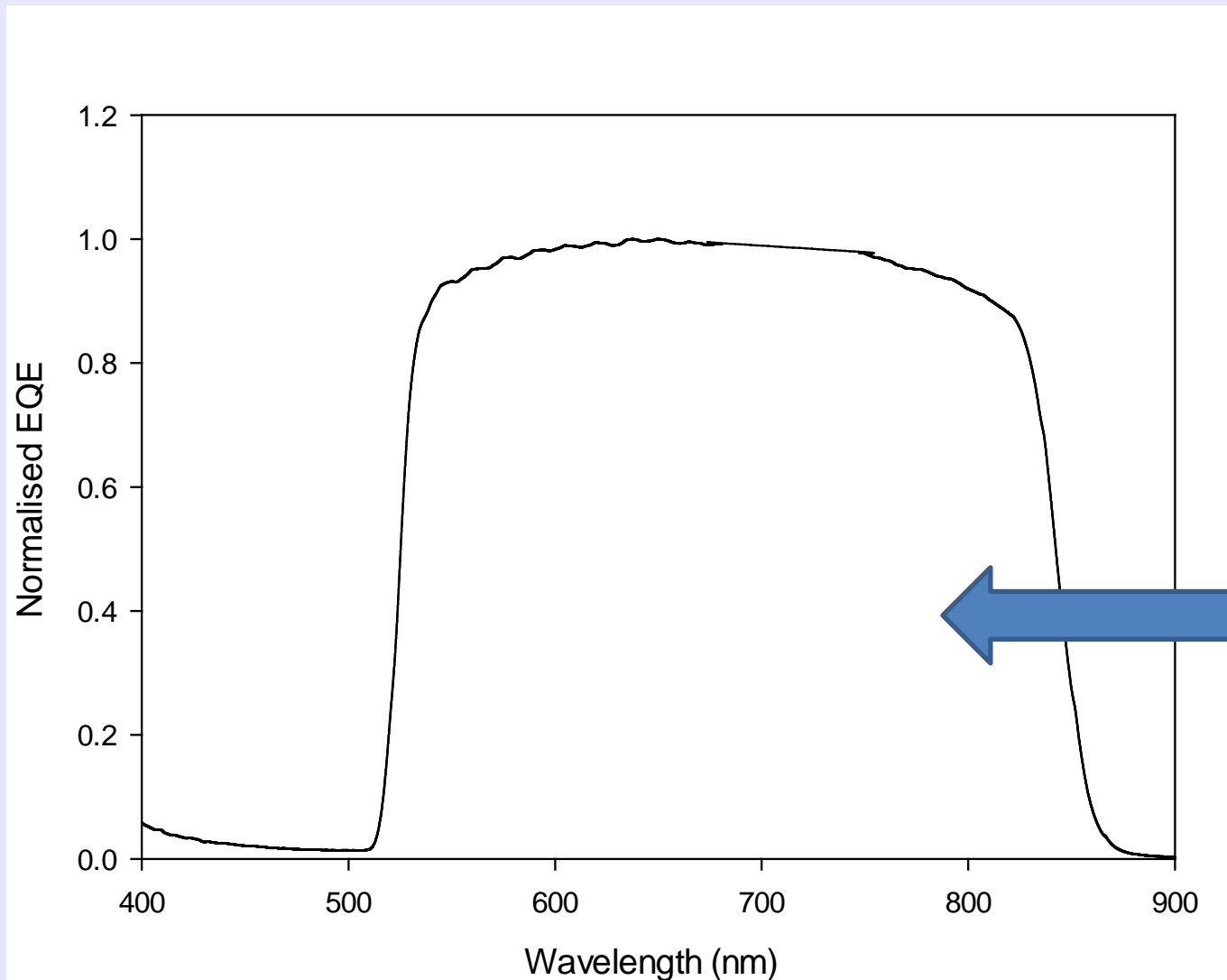
Define the EQE as the ratio of photons in to current generated

$$EQE = \frac{J_{cell}}{J_{photon}}$$

A 100% EQE means every photon that strikes the cell generates an electron hole pair which flows through the external circuit



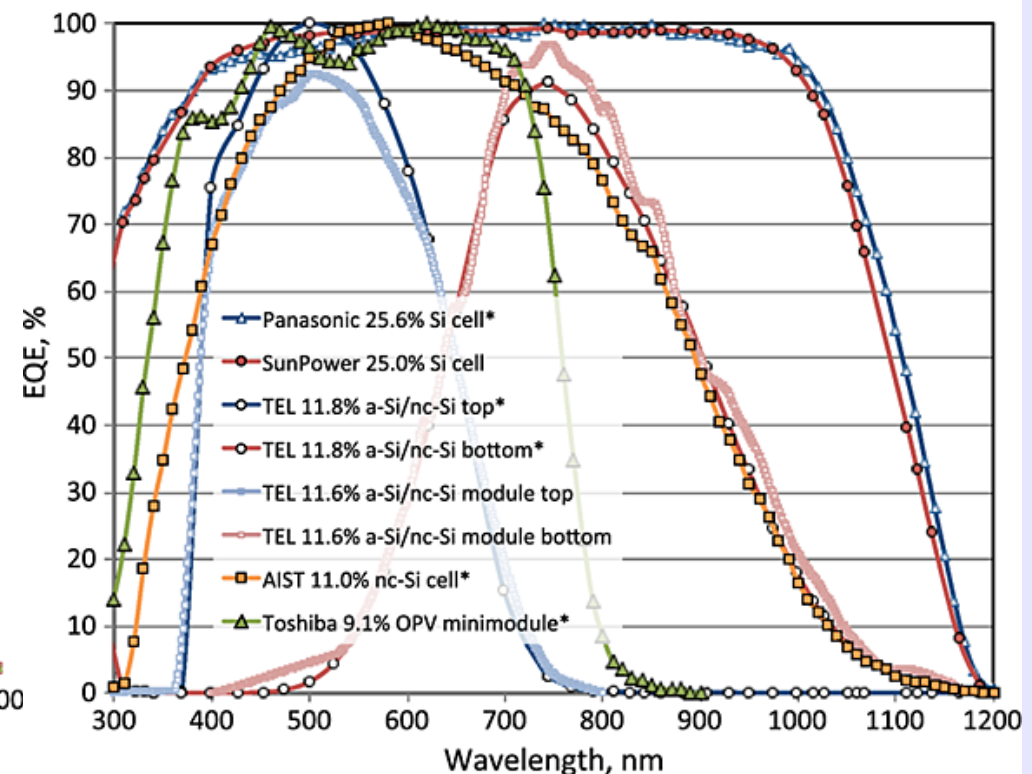
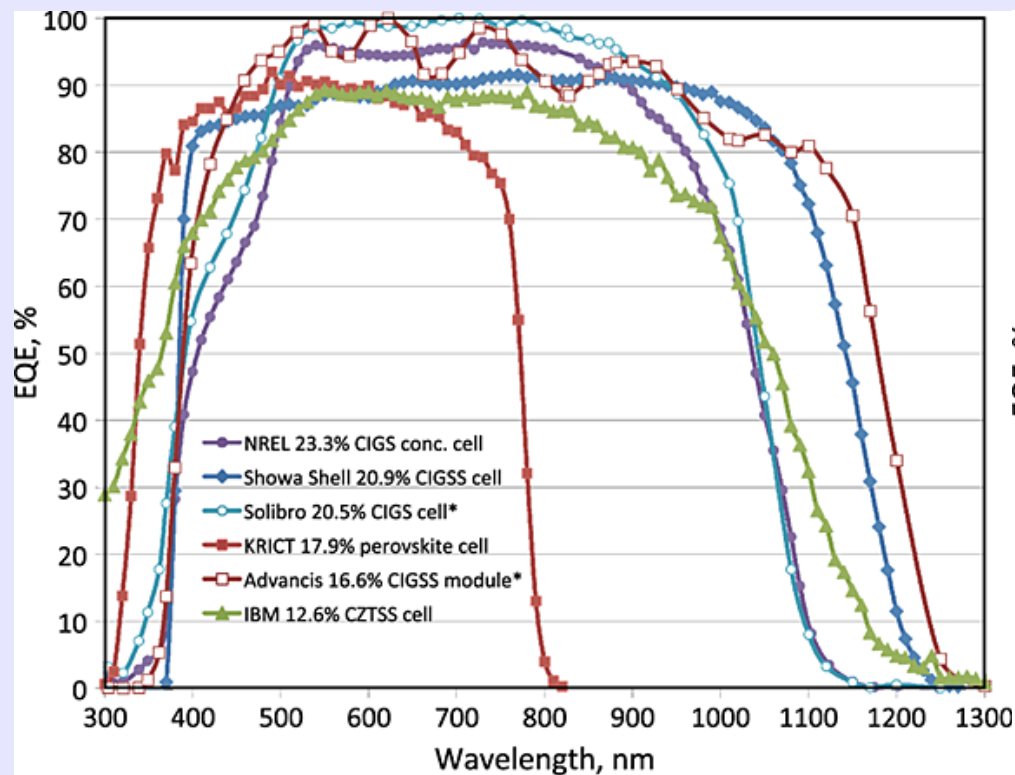
Theoretical case



Area under
curve
proportional to
 J_{sc}



Most solar cell technologies display same typical “top hat” EQE curve shape





Progress in PHOTOVOLTAICS

PROGRESS IN PHOTOVOLTAICS: RESEARCH AND APPLICATIONS

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ACCELERATED PUBLICATION

Solar cell efficiency tables (version 44)

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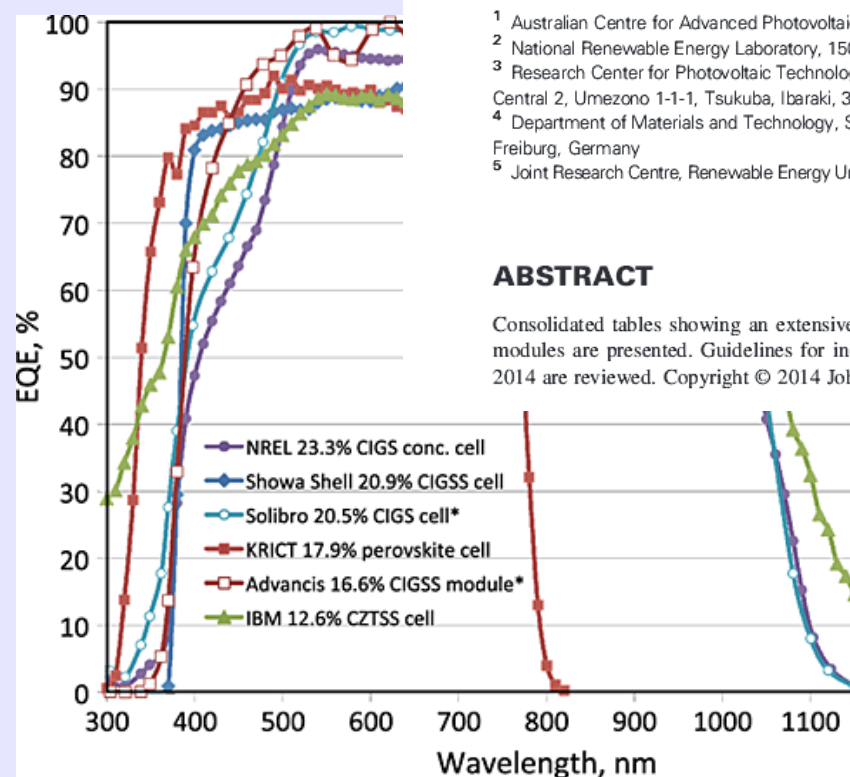
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ABSTRACT

Consolidated tables showing an extensive modules are presented. Guidelines for in 2014 are reviewed. Copyright © 2014 Jo



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Table I. Confirmed terrestrial cell and submodule efficiencies measured under the global AM1.5 spectrum (1000 W/m²) at 25 °C (IEC 60904-3: 2008, ASTM G-173-03 global).

Classification ^a	Efficiency (%)	Area ^b (cm ²)	V _{oc} (V)	J _{sc} (mA/cm ²)	Fill factor (%)	Test centre ^c (date)	Description
Silicon							
Si (crystalline)	25.6 ± 0.5	143.7 (da)	0.740	41.8 ^d	82.7	AIST (2/14)	Panasonic HIT, rear-junction [4]
Si (multicrystalline)	20.4 ± 0.5	1.002 (ap)	0.664	38.0	80.9	NREL (5/04) ^e	PhG-ISE [21]
Si (thin film transfer)	20.1 ± 0.4	242.6 (ap)	0.682	38.14 ^f	77.4	NREL (10/12)	Solixel (43 μm thick) [22]
Si (thin film minimodule)	10.5 ± 0.3	94.0 (ap)	0.492 ^g	29.7 ^g	72.1	PhG-ISE (8/07) ^g	CSG Solar (<2 μm on glass; 20 cells) [23]
III–V Cells							
GaAs (thin film)	28.8 ± 0.9	0.9927 (ap)	1.122	29.68 ^h	86.5	NREL (5/12)	Alta Devices [24]
GaAs (multicrystalline)	18.4 ± 0.5	4.011 (b)	0.994	23.2	79.7	NREL (11/95) ^e	RTI, Ge substrate [25]
InP (crystalline)	22.1 ± 0.7	4.02 (b)	0.878	29.5	85.4	NREL (4/90) ^e	Spire, epitaxial [26]
Thin film chalcogenide							
CIGS (cell)	20.5 ± 0.6	0.9882 (ap)	0.752	35.3 ^d	77.2	NREL (3/14)	Solibro, on glass [5]
CIGS (minimodule)	18.7 ± 0.6	15.892 (da)	0.701 ^g	35.29 ^g	75.6	PhG-ISE (9/13)	Solibro, four serial cells [27]
CdTe (cell)	19.6 ± 0.4	1.0055 (ap)	0.8573	28.59 ^g	80.0	Newport (8/13)	GE Global Research [28]
Amorphous/microcrystalline Si							
Si (amorphous)	10.1 ± 0.3 ^k	1.036 (ap)	0.886	16.75 ^l	67.8	NREL (7/09)	Oerlikon Solar Lab, Neuchâtel [29]
Si (microcrystalline)	11.0 ± 0.3 ^m	1.045 (da)	0.542	27.44 ^d	73.8	AIST (1/14)	AIST [9]
Dye sensitised							
Dye sensitised	11.9 ± 0.4 ⁿ	1.005 (da)	0.744	22.47 ^l	71.2	AIST (8/12)	Sharp [30]
Dye sensitised (minimodule)	29.9 ± 0.4 ⁿ	17.11 (ap)	0.719 ^g	19.4 ^g	71.4	AIST (8/10)	Sony, eight parallel cells [31]
Dye (submodule)	8.8 ± 0.3 ⁿ	398.8 (da)	0.697 ^g	18.42 ^g	68.7	AIST (8/12)	Sharp, 26 serial cells [32]
Organic							
Organic thin film	10.7 ± 0.3 ^o	1.013 (da)	0.872	17.75 ^l	68.9	AIST (10/12)	Mitsubishi Chemical (4.4 × 23.0 mm) [33]
Organic (minimodule)	9.1 ± 0.3 ^o	25.04 (da)	0.794 ^g	17.06 ^g	67.5	AIST (2/14)	Toshiba (four series cells) [10]
Organic (submodule)	6.8 ± 0.2 ^o	395.9 (da)	0.798 ^g	13.50 ^g	62.8	AIST (10/12)	Toshiba (15 series cells) [10]
Multijunction devices							
InGaP/GaAs/InGaAs	37.9 ± 1.2	1.047 (ap)	3.065	14.27 ^l	86.7	AIST (2/13)	Sharp [34]
a-SiHc-SiHc-Si (thin film)	13.4 ± 0.4 ^p	1.006 (ap)	1.963	9.52 ^l	71.9	NREL (7/12)	LG Electronics [35]

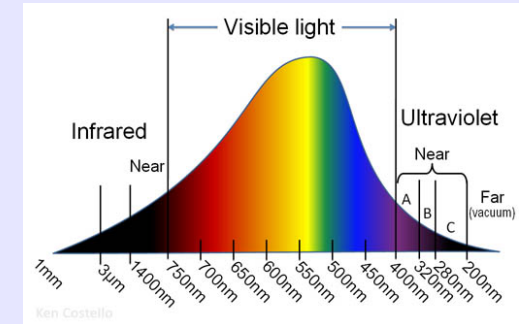
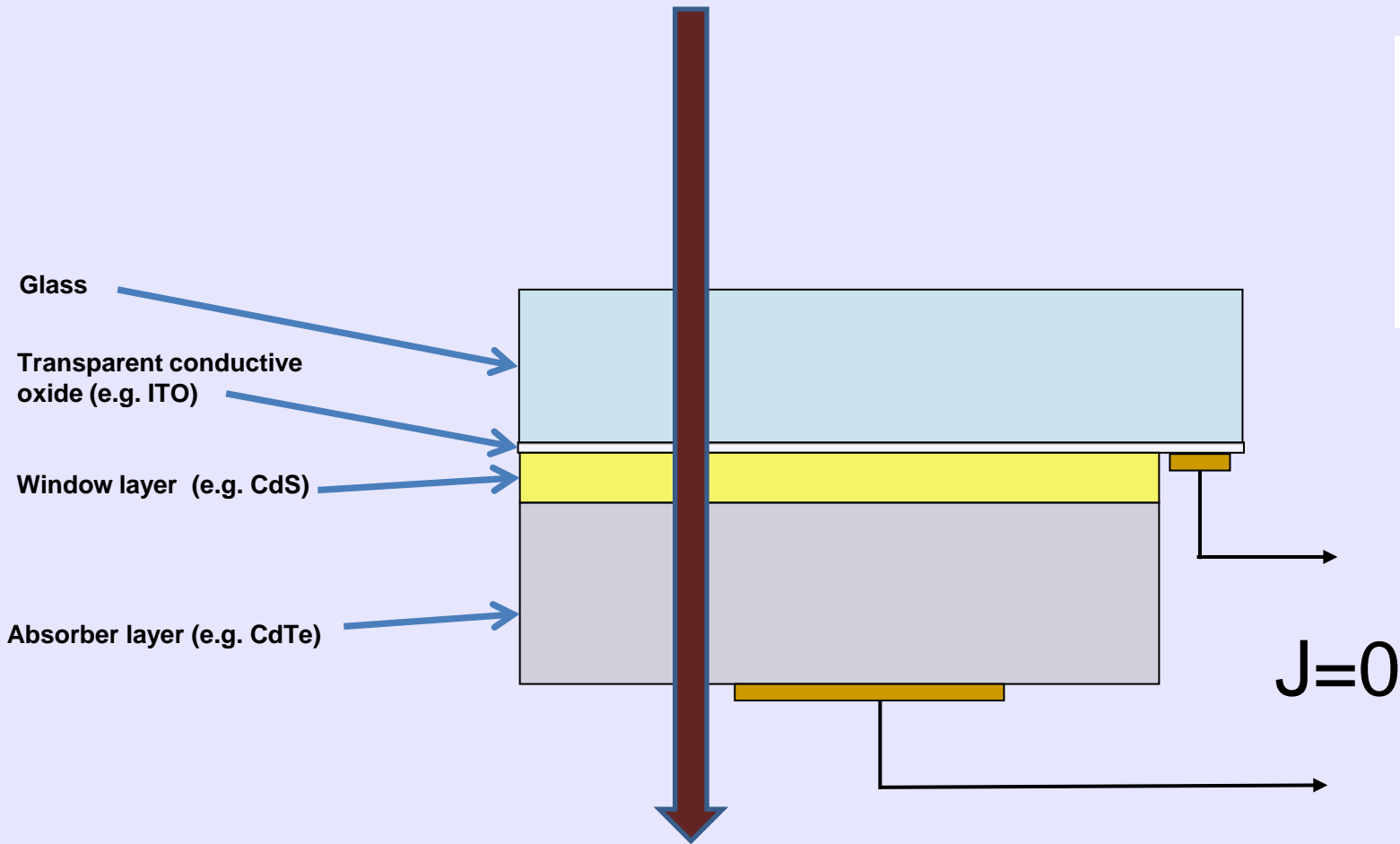
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Solar cell efficiency tables

M. A. Green et al.



External quantum efficiency (EQE)

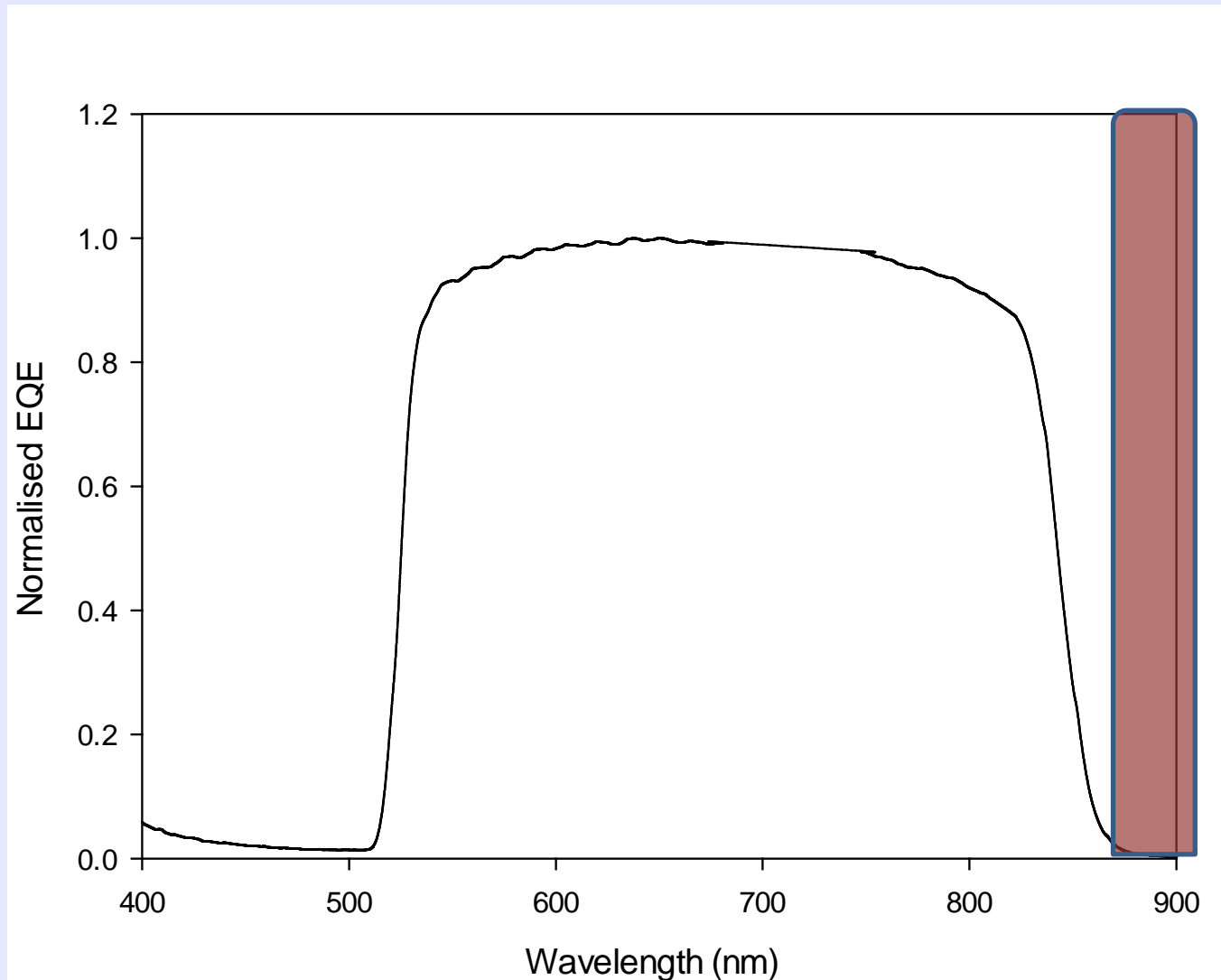


$$E(\lambda) = \frac{hc}{\lambda}$$

Region 1: $E_{\text{photon}} < E_{\text{Asorber}}$

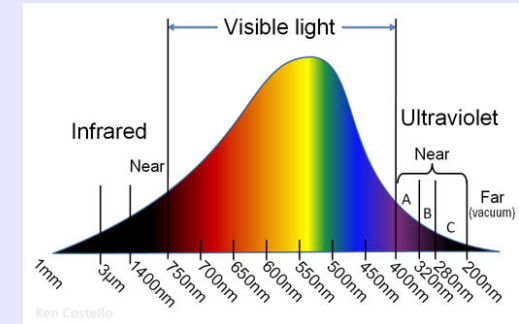
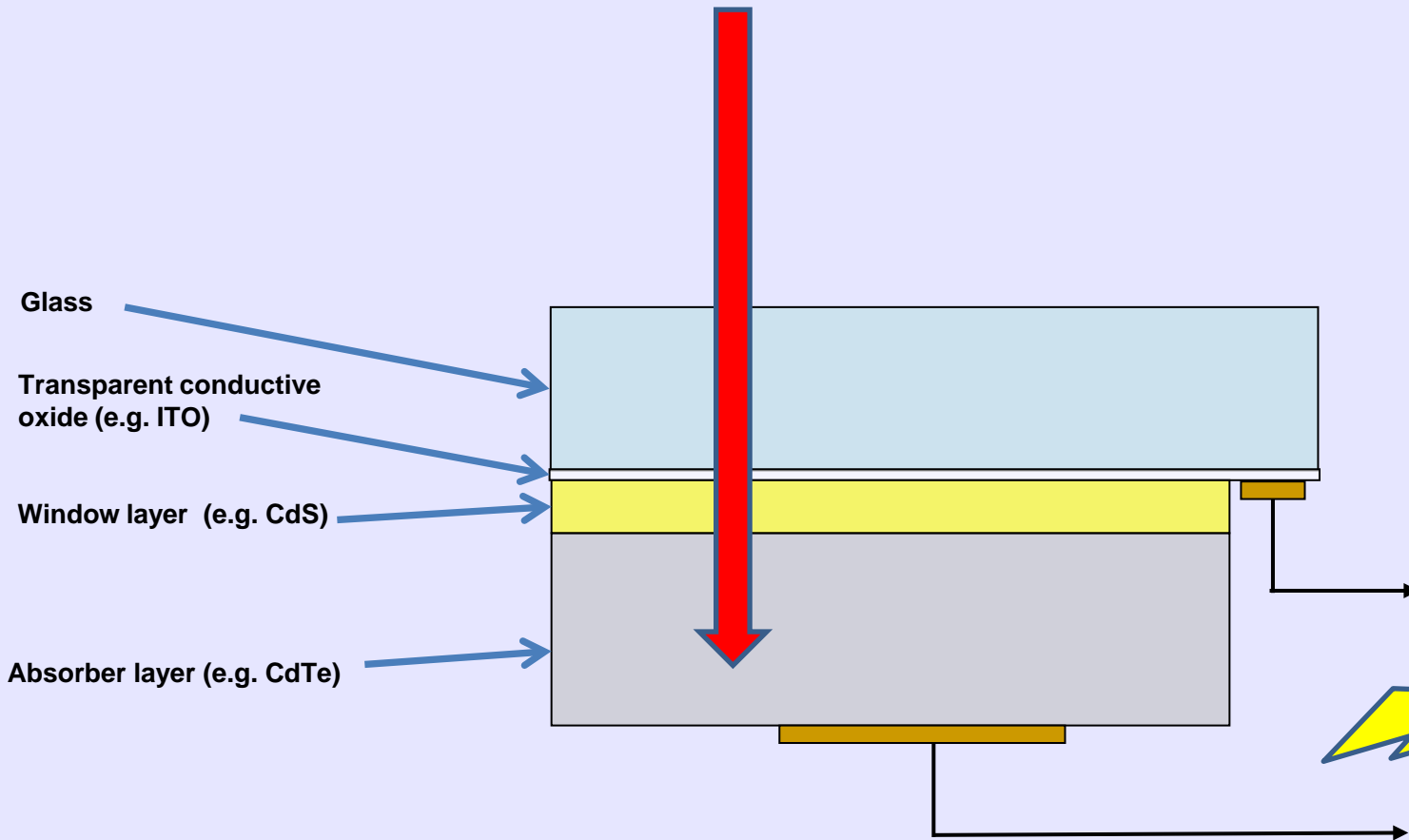


External quantum efficiency (EQE)





External quantum efficiency (EQE)

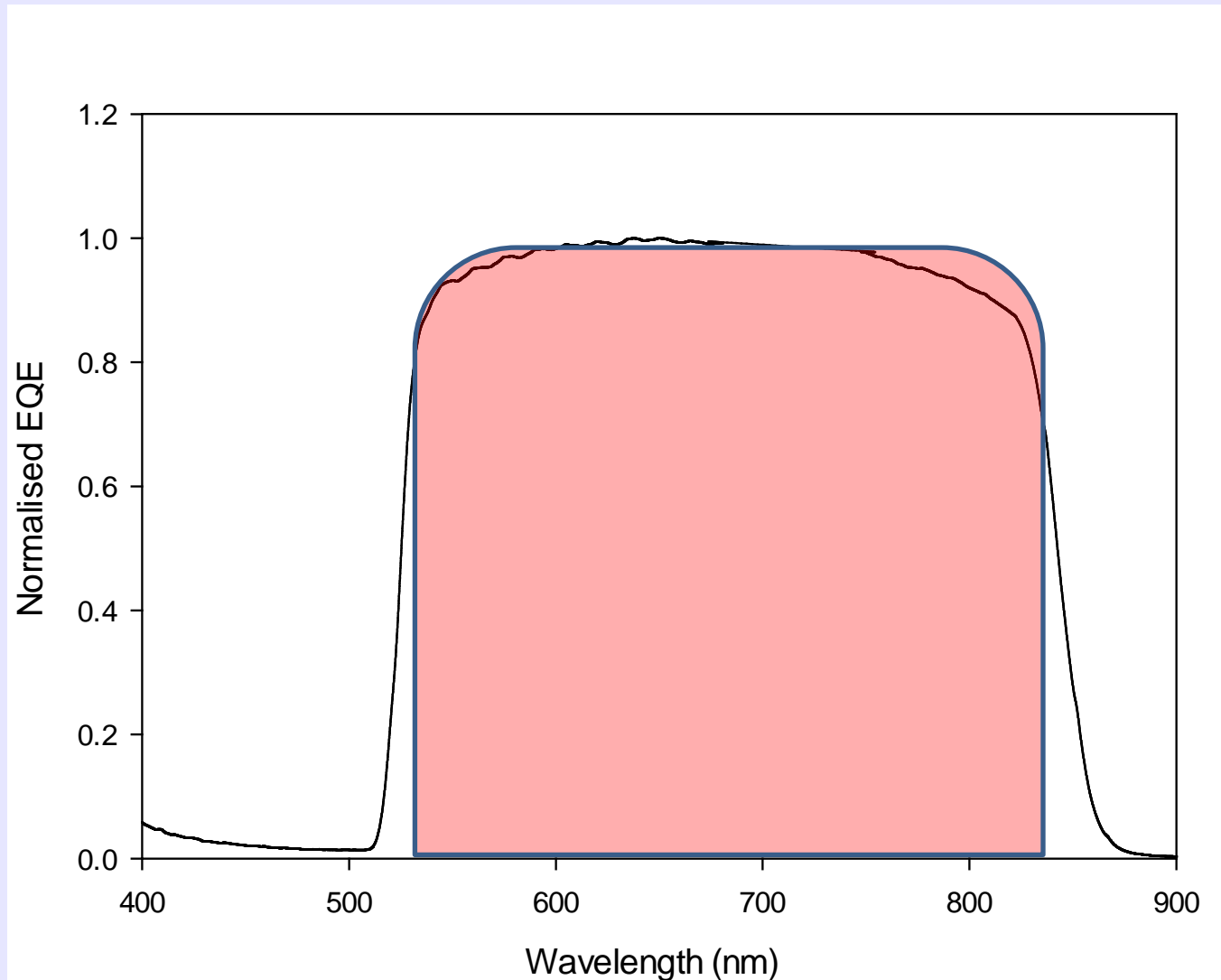


$$E(\lambda) = \frac{hc}{\lambda}$$

Region 2: $E_{\text{photon}} > E_{\text{Asorber}}$

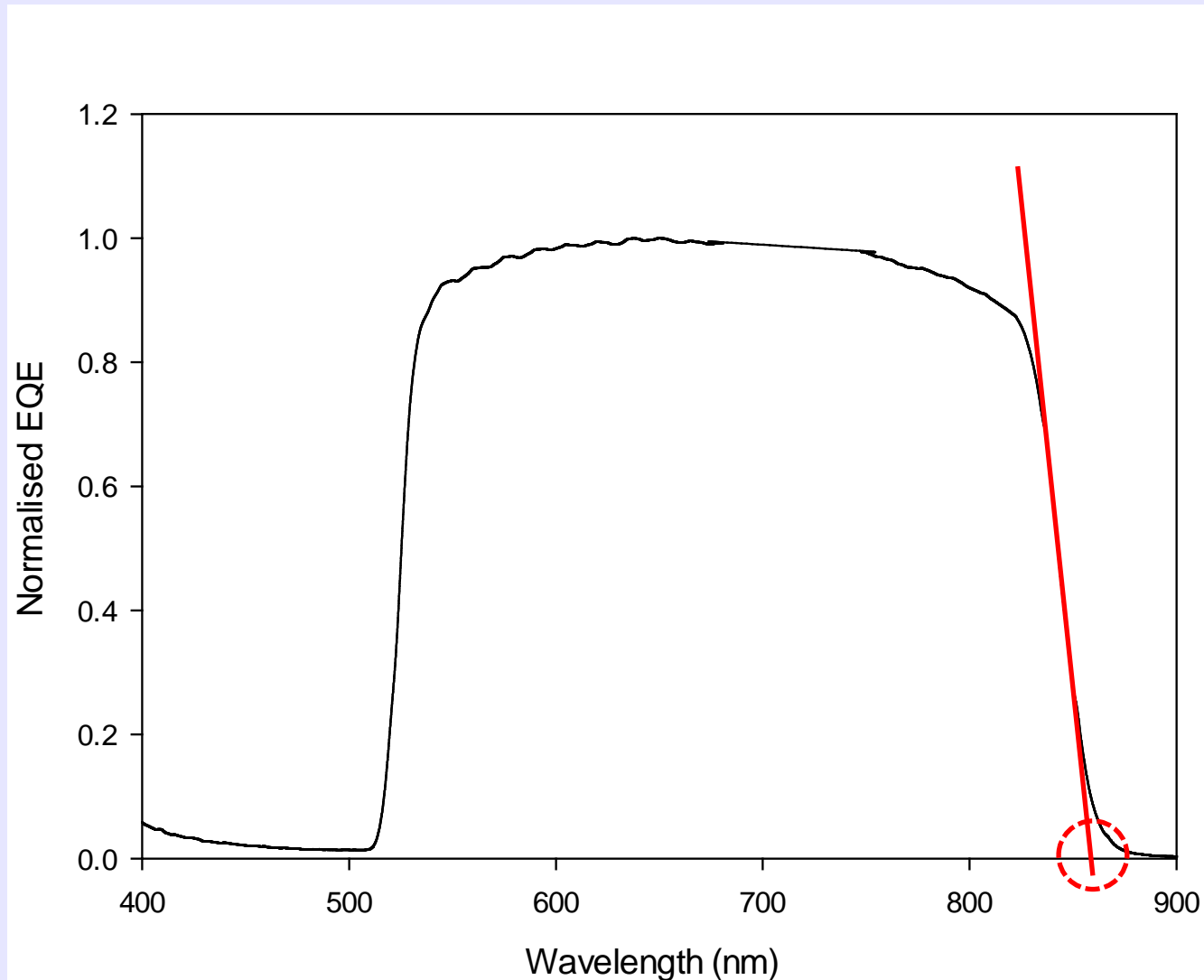


External quantum efficiency (EQE)





External quantum efficiency (EQE)

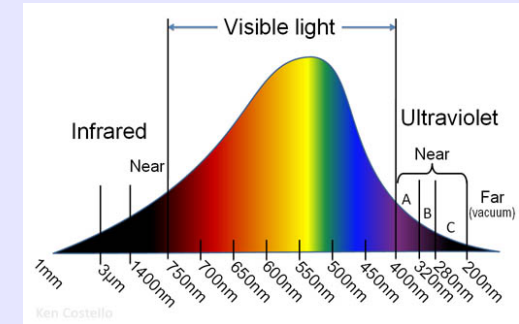
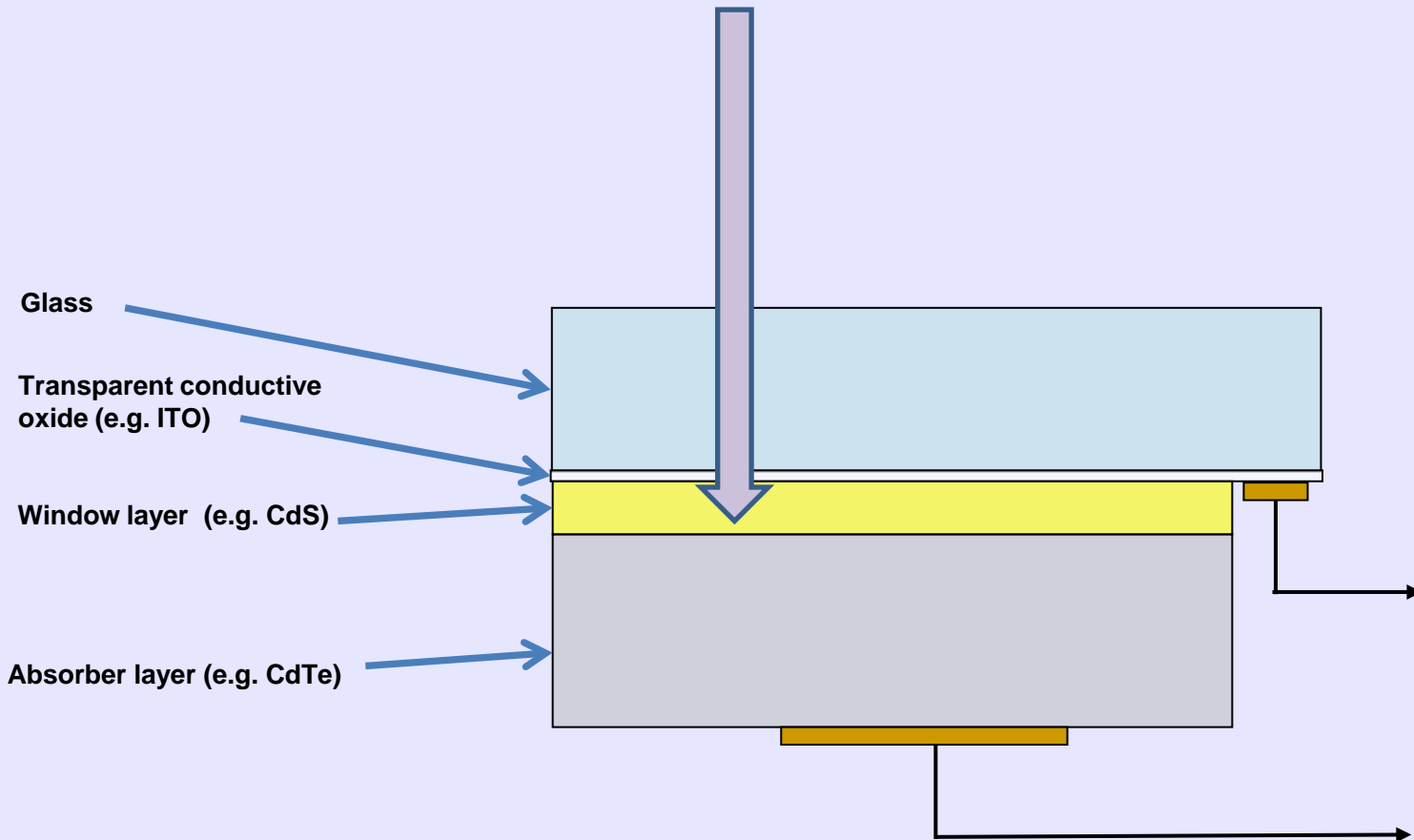


Absorber bandgap

$$E(\lambda) = \frac{hC}{\lambda}$$



External quantum efficiency (EQE)

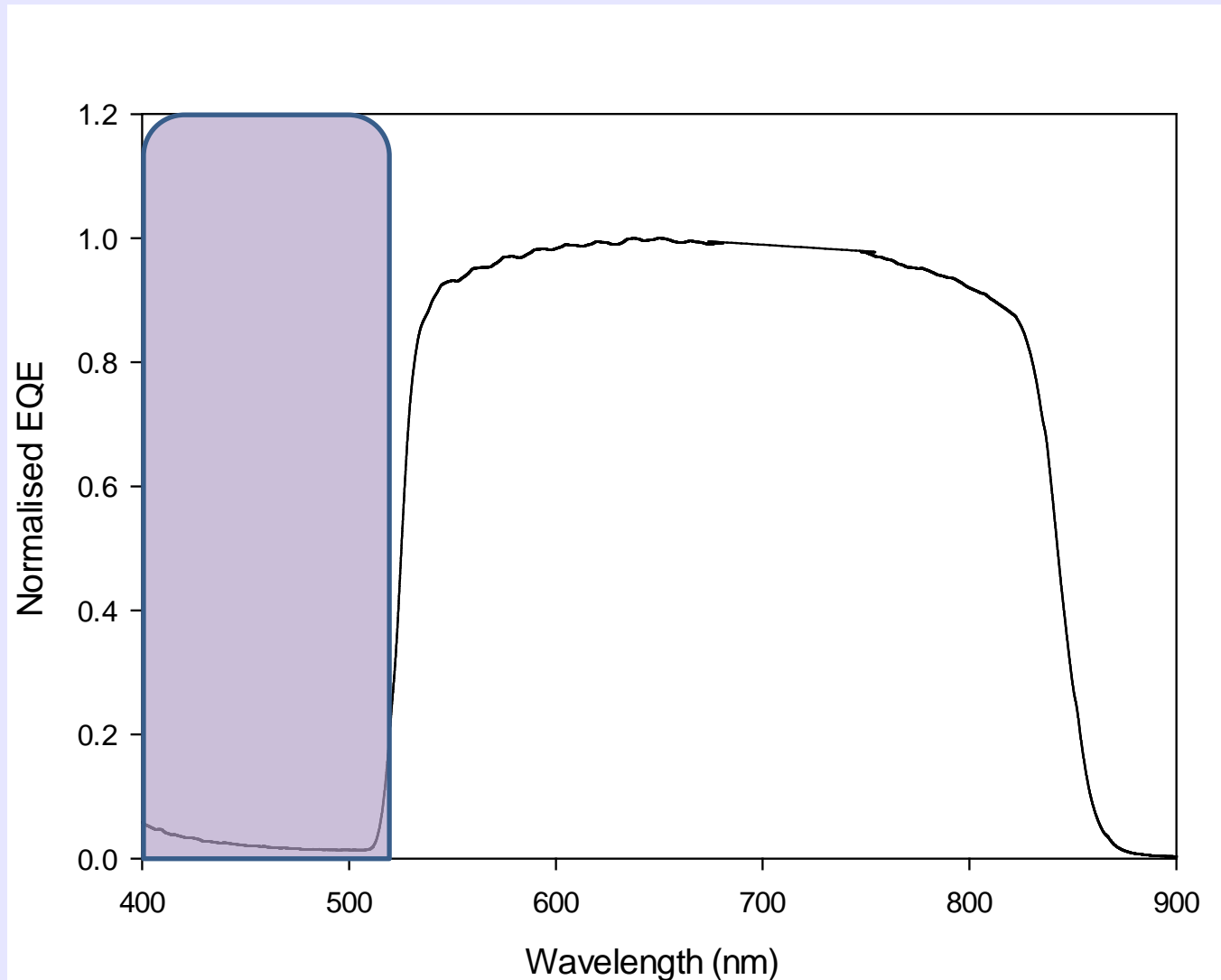


$$E(\lambda) = \frac{hc}{\lambda}$$

Region 3: $E_{\text{photon}} > E_{\text{window}}$

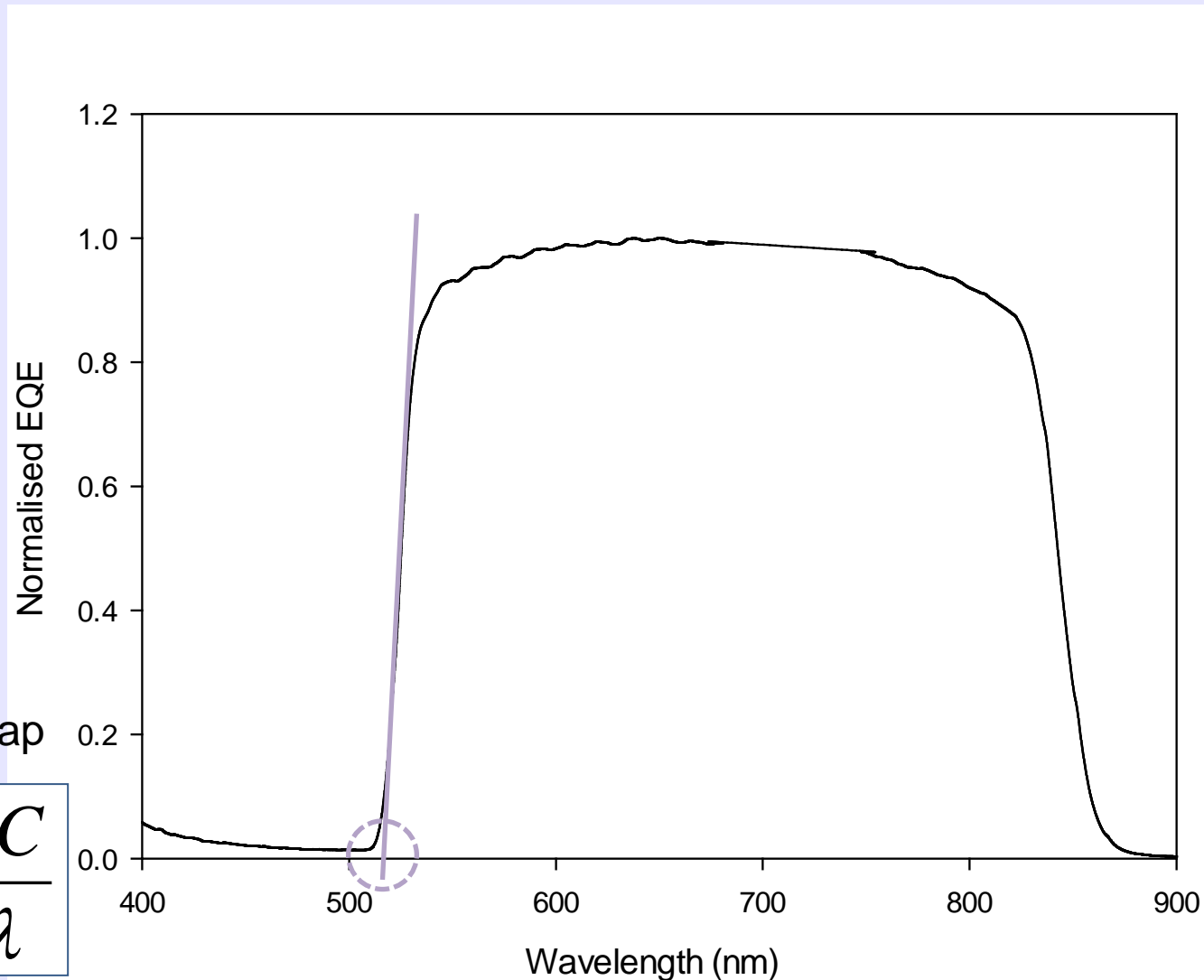


External quantum efficiency (EQE)





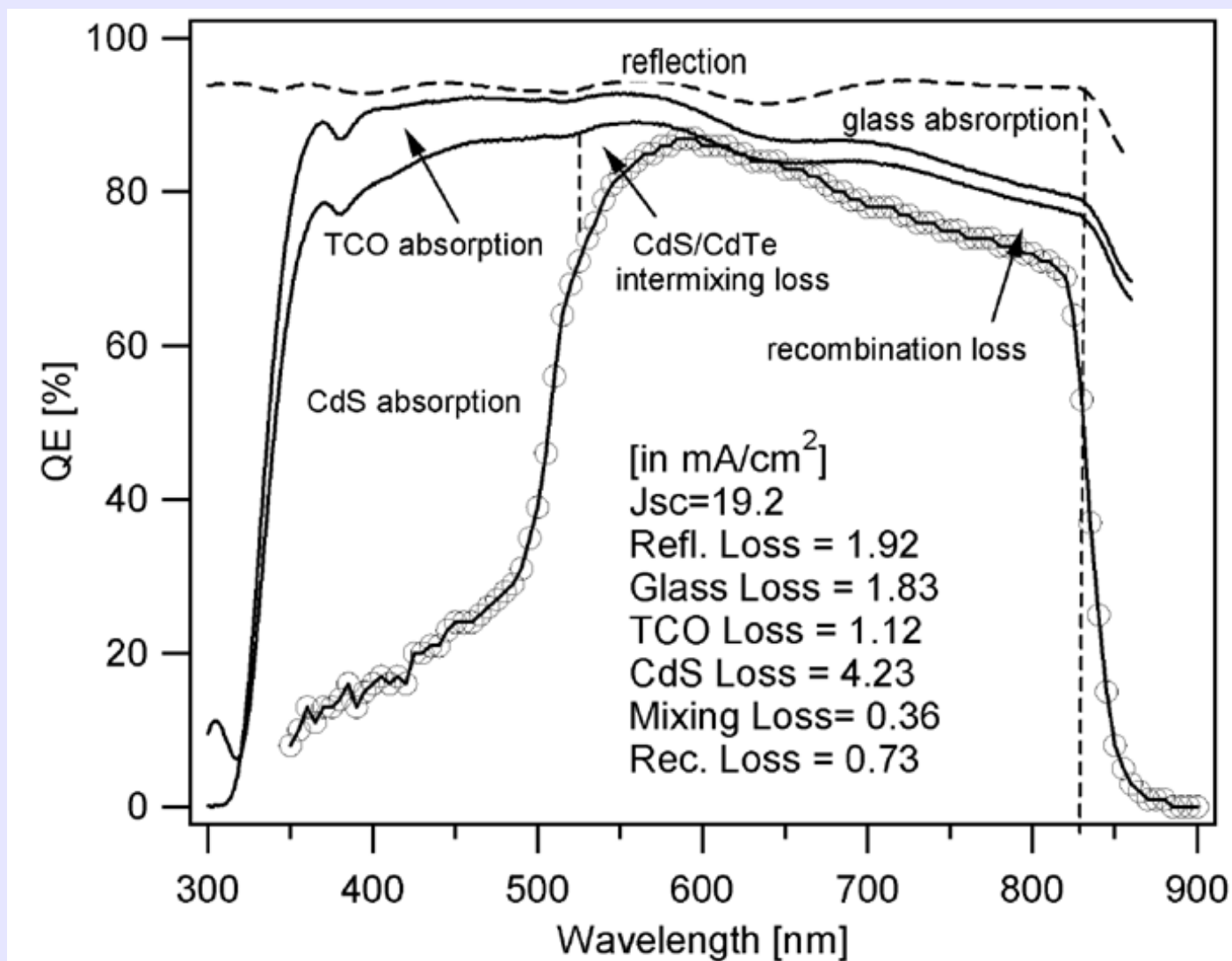
External quantum efficiency (EQE)



$$E(\lambda) = \frac{hc}{\lambda}$$

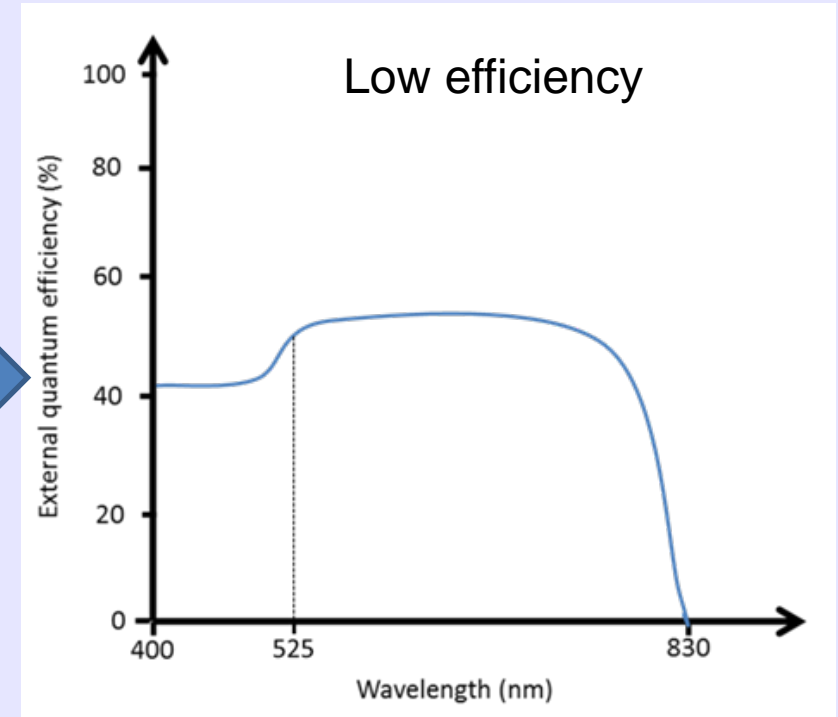
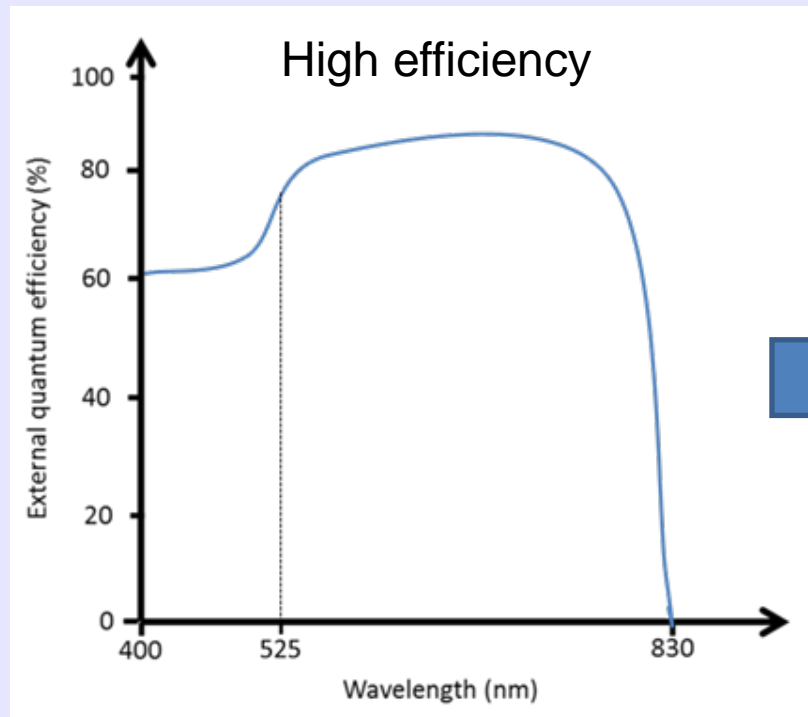


Real cells





Optical losses

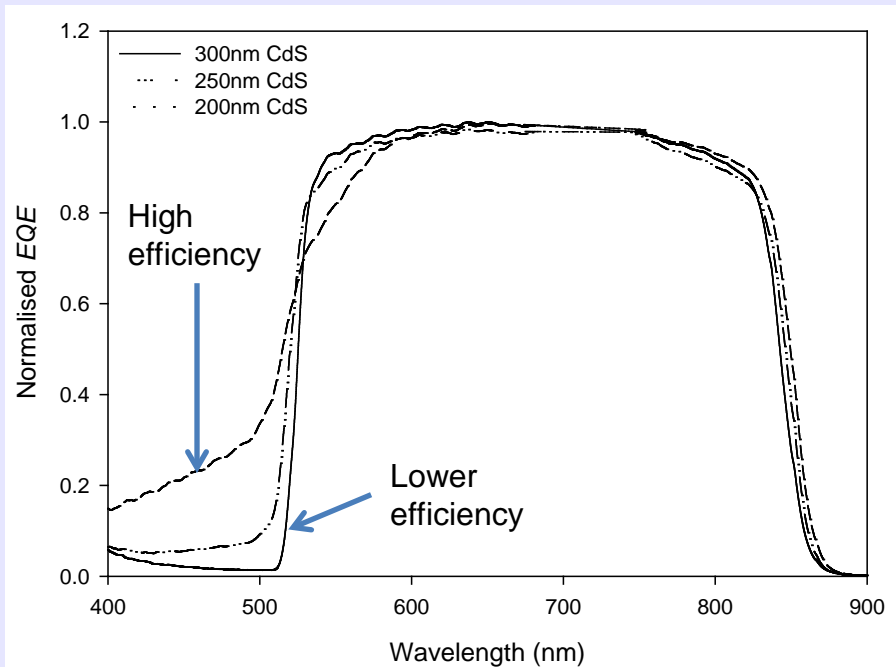


Uniform reduction in EQE signal across wavelength range is an indication of optical losses i.e. reflection loss, optical blockages etc.

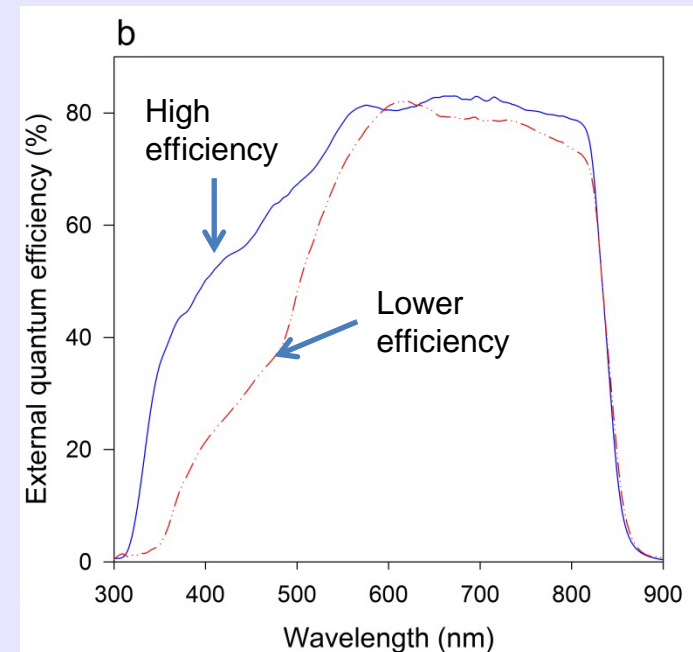
Can also be an indication of poor system calibration.



window/ n-type layer losses



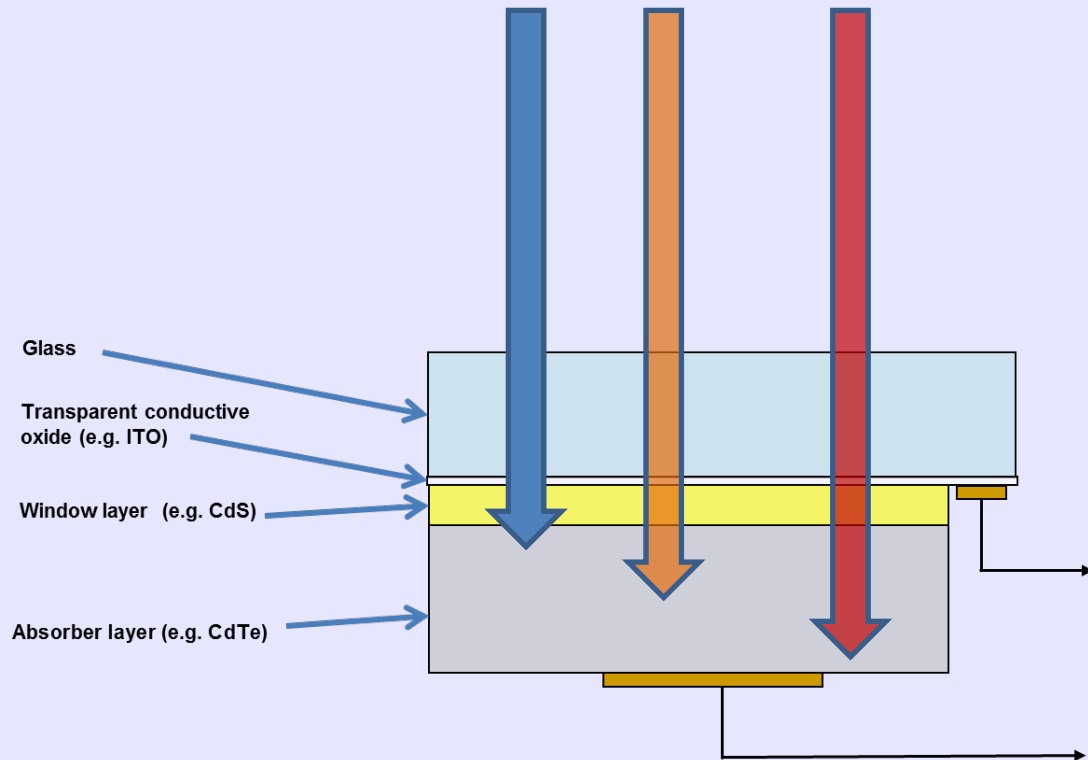
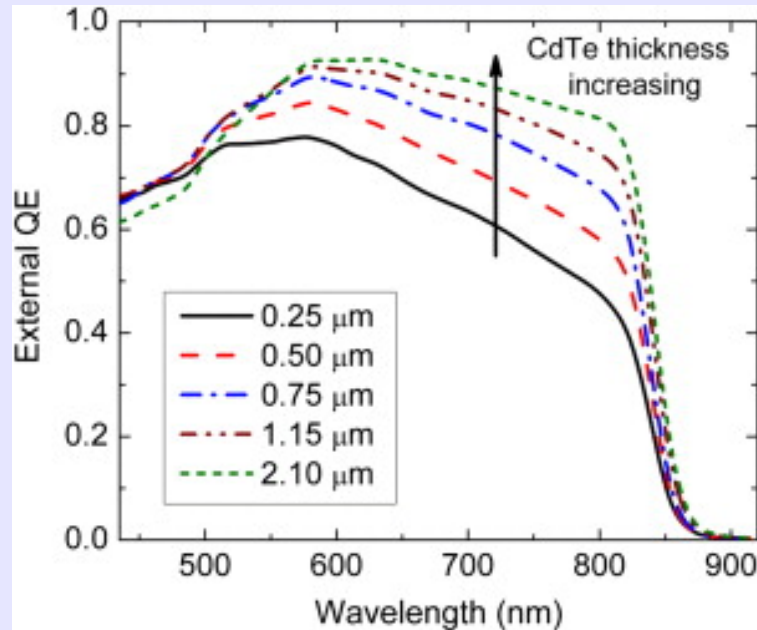
Reduced window layer thickness allows more light transmission to absorber



Higher bandgap n-type layer shifts absorption edge



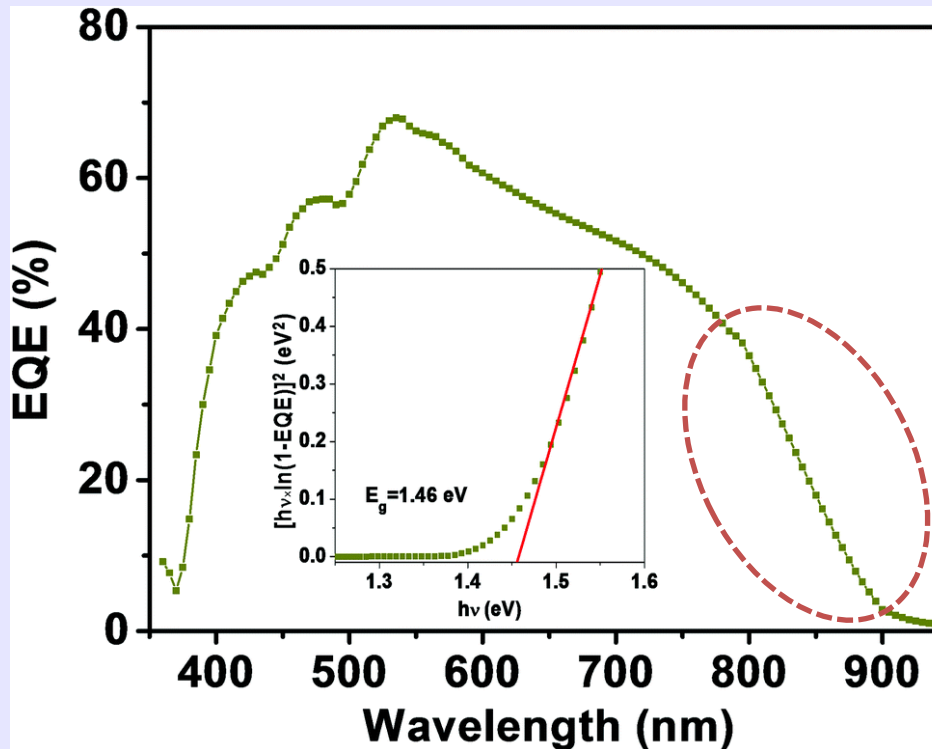
Deep penetration losses



If absorber layer is thin or depletion width is narrow then longer wavelength photons have an increased probability of not contributing to photocurrent. See a decrease in EQE for longer wavelengths.



Graded bandgap/back surface recombination



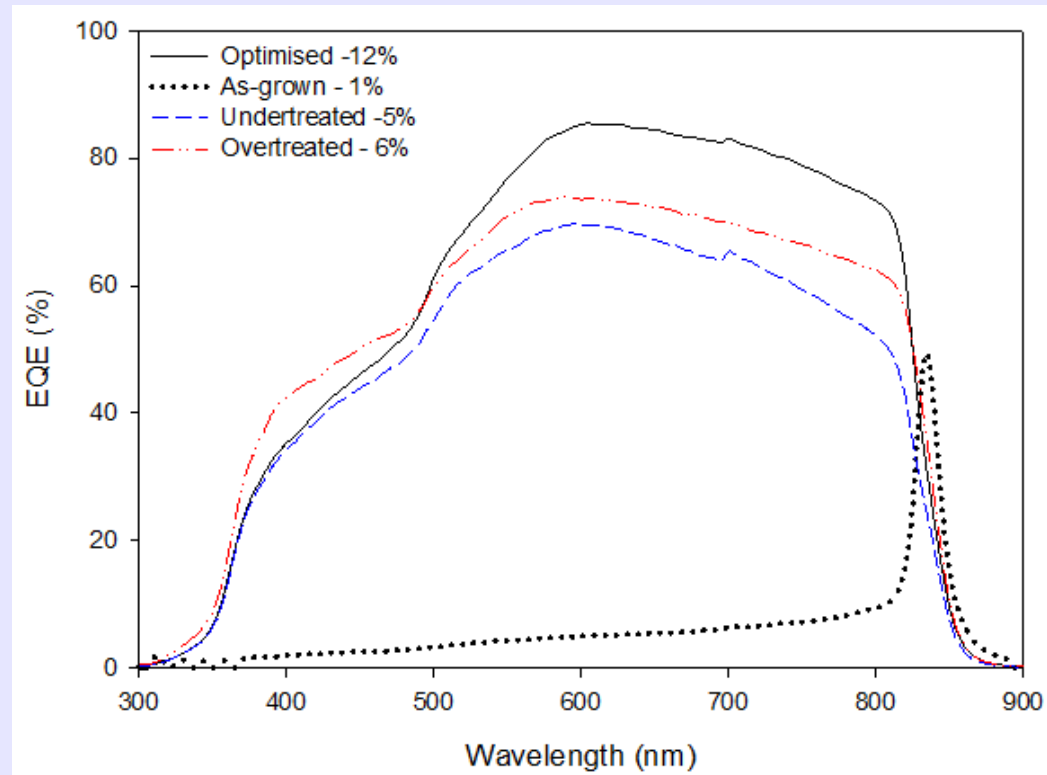
In multi component materials such as CZTS can get a broad cut-off region due to changes in the absorber bandgap.

This often signifies the material isn't single phase and reduces the efficiency.

Can also indicate enhanced recombination close to the back surface.



External quantum efficiency (EQE)

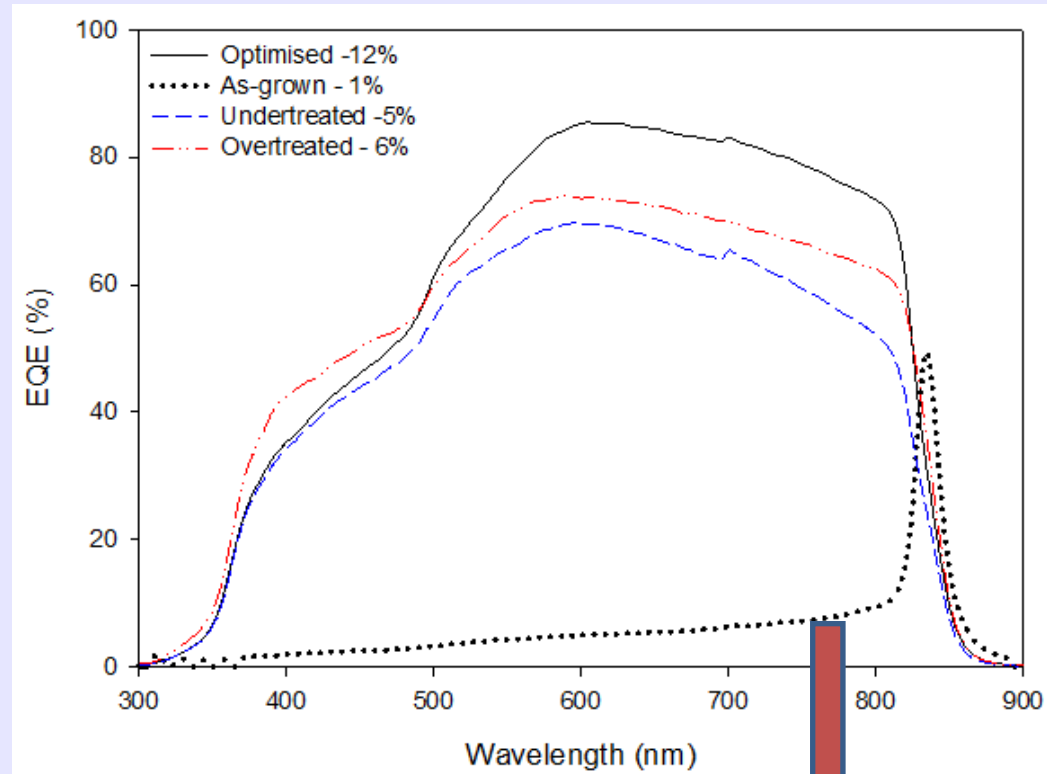


In certain situations see a complete change in EQE that corresponds to very low performance

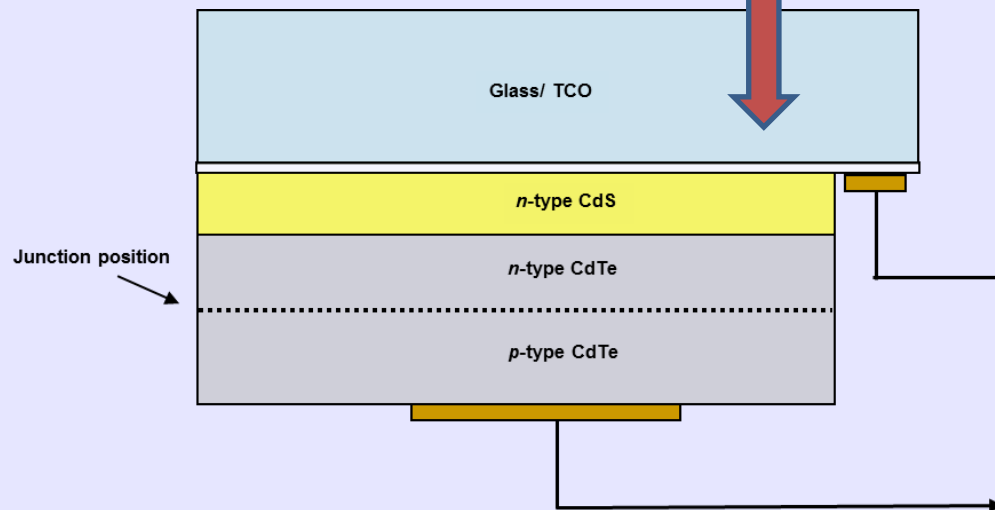
See reasonable EQE response near absorber band-edge but low response at all other wavelengths



External quantum efficiency (EQE)

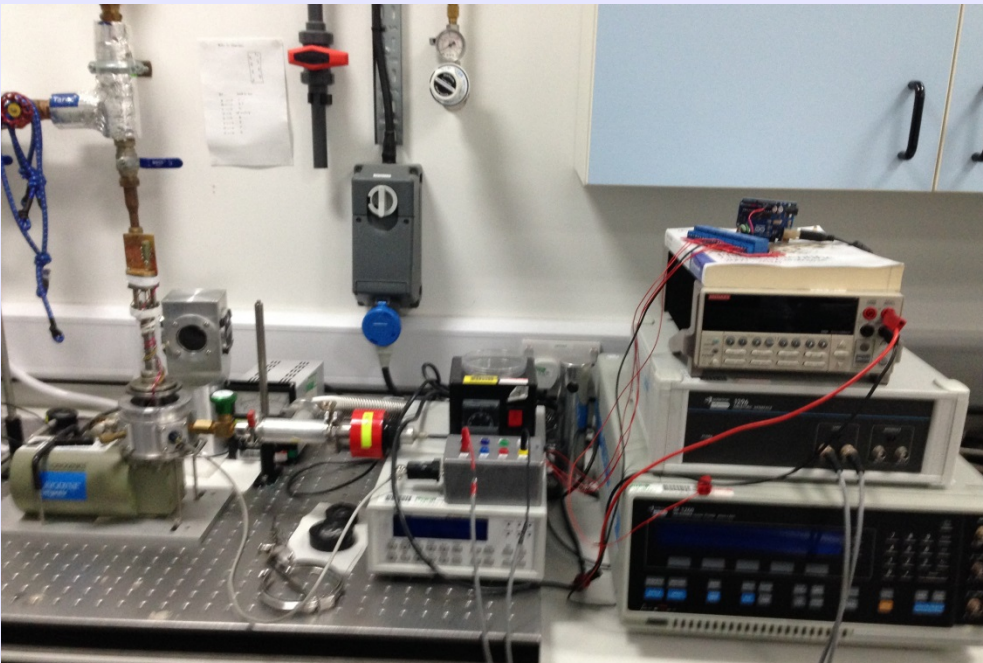


This is a **buried junction response** due to both p and n type regions being present in the absorber layer





Capacitance Voltage analysis (C-V)



Liverpool CV/Admittance spectroscopy system

Capacitance-voltage measurements are useful in deriving particular parameters about PV devices.

Depending on the type of solar cell, capacitance-voltage (C-V) measurements can be used to derive parameters such as the doping concentration and the built-in voltage of the junction.

A capacitance-frequency (C-f) sweep can be used to provide information on the existence of traps in the depletion region.



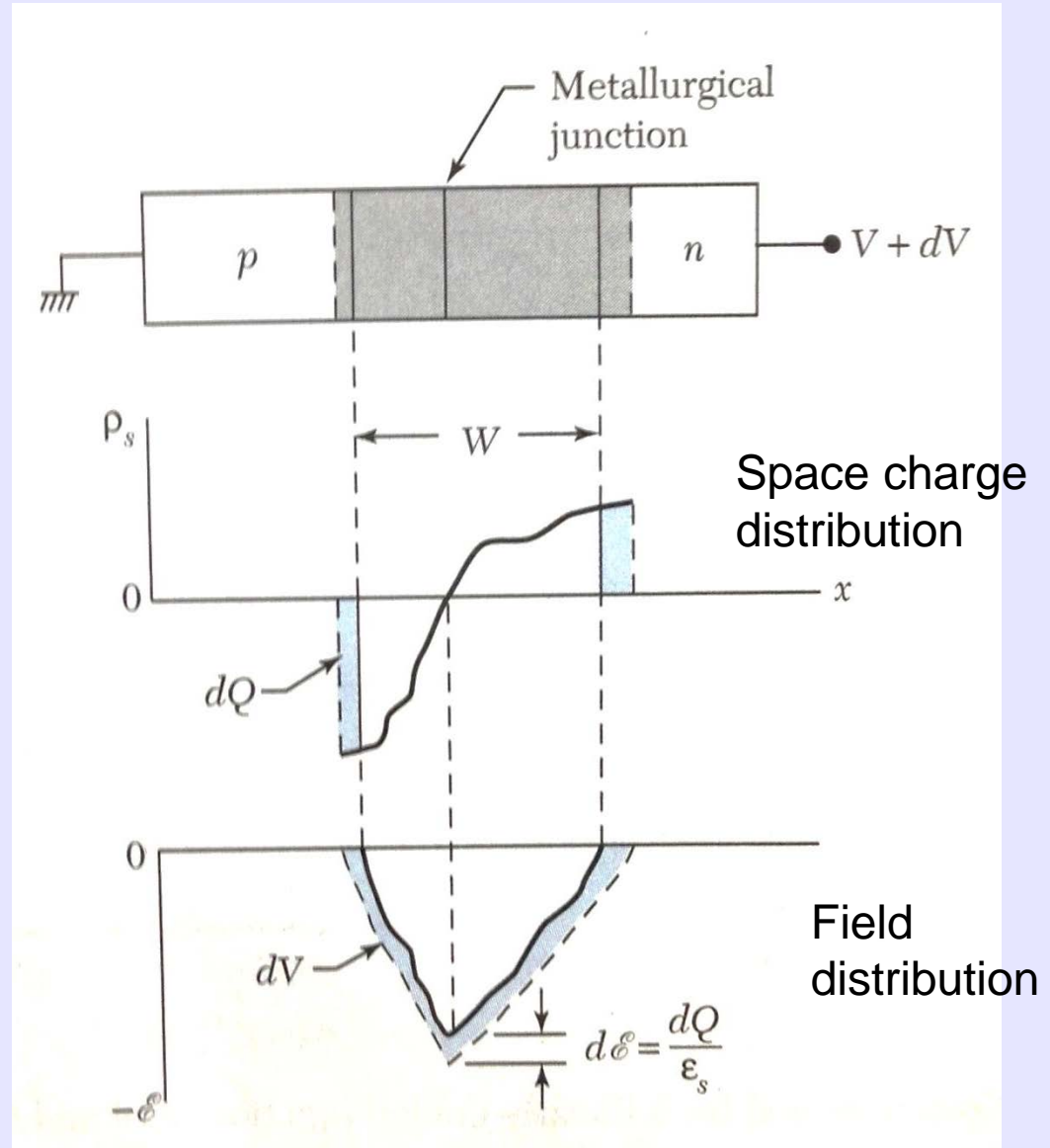
Capacitance Voltage analysis (C-V)

Can treat a p-n junction as a capacitor - depletion region sandwiched between two plates.

Can define the junction capacitance per unit area as

$$C_j \equiv \frac{dQ}{dV} = \frac{dQ}{W \frac{dQ}{d\epsilon_s}} = \frac{\epsilon_s}{W}$$

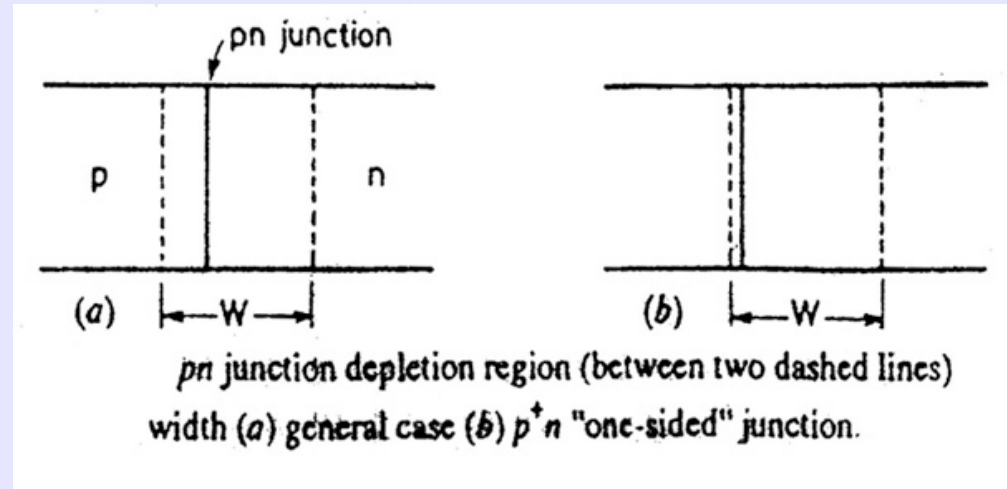
Where W is the width of the depletion region and ϵ_s is the semiconductor permittivity





Capacitance Voltage analysis (C-V)

We generally assume a one sided abrupt junction for calculations. This is due to the higher doping densities in the n-type layer meaning the depletion region lies within the p-type layer



For a one sided junction we can determine the depletion width as

$$W = \sqrt{\frac{2\epsilon_s V}{qN_a}}$$

Electron charge

Acceptor doping density

Using previous equation we get the relation

$$\frac{1}{C^2} = \frac{2V}{q\epsilon_s N_a}$$

Hence we can get N_a from a plot of $1/C^2$ vs V



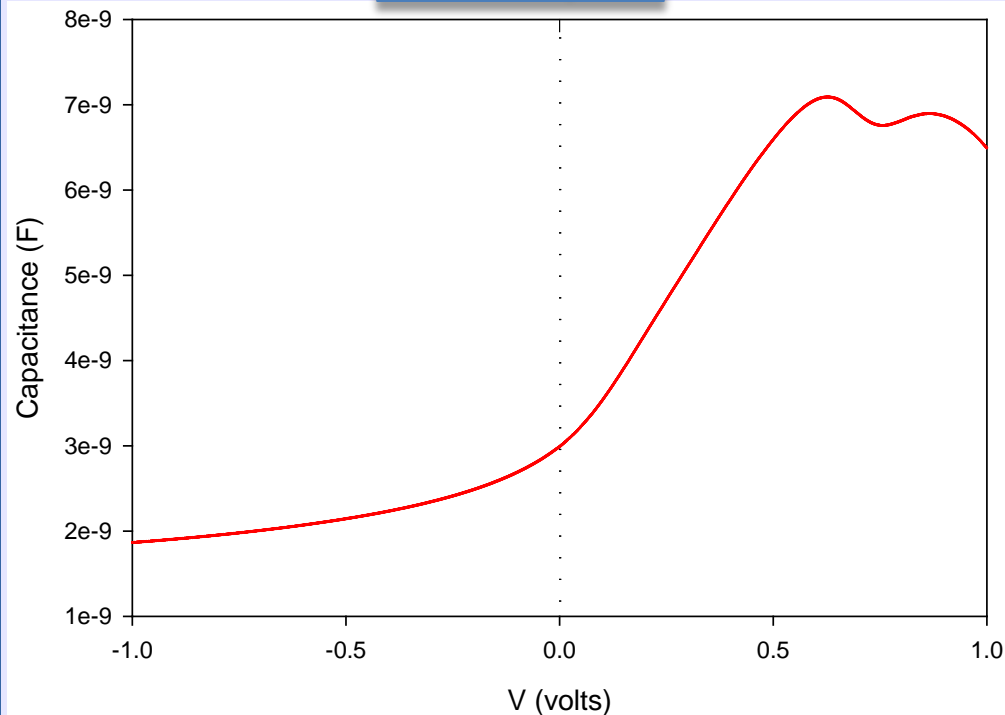
Capacitance Voltage analysis (C-V)

Worked example – CdTe solar cell

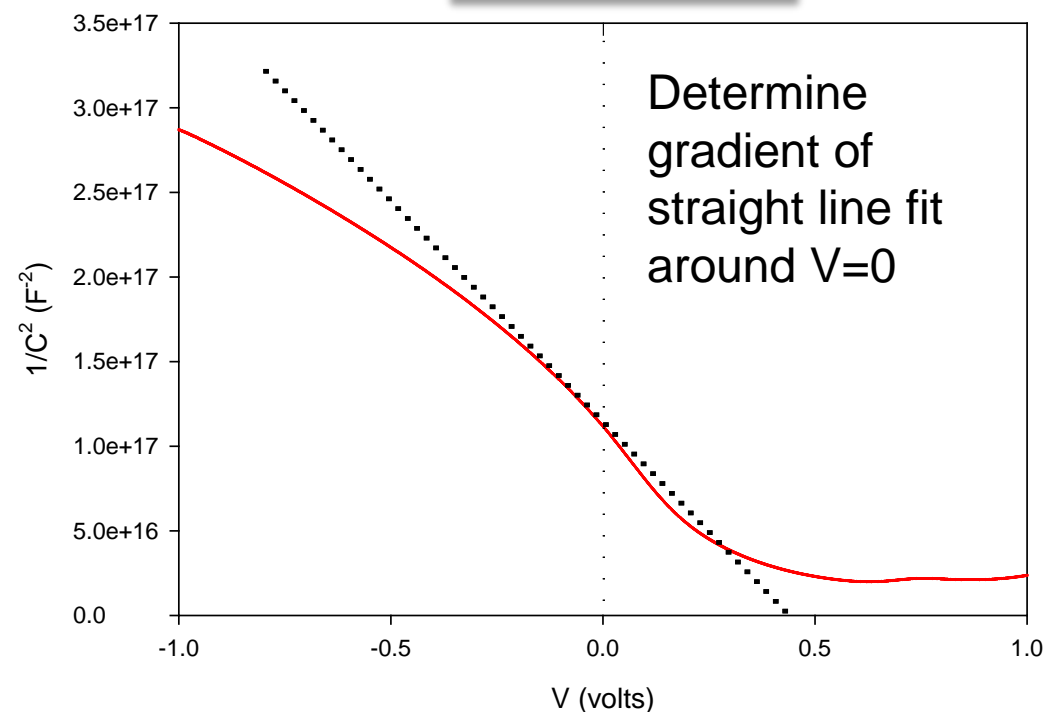
Measure the capacitance response of the cell as a function of applied bias.

C-V measurements can be made either forward-biased or reverse biased. However, when the cell is forward-biased, the applied DC voltage must be limited; otherwise non-ohmic back contacts can alter the signal.

C vs. V plot



1/C² vs. V plot





Capacitance Voltage analysis (C-V)

p-type doping
density is given by

$$N_a = \frac{2}{q\epsilon_s A^2 \frac{d(\frac{1}{C^2})}{dV}}$$

We have included a
contact area term A

So for a CdTe cell with 5mm diameter square contacts we measured a the slope of the $1/C^2$ plot to be 2.7×10^{17} .

$$\frac{d(\frac{1}{C^2})}{dV} = 2.7 \times 10^{17} V^{-1} F^{-2}$$

$$A = 0.005 \times 0.005 = 0.25 \times 10^{-5} m^2$$

$$\epsilon_s = 10.36 \epsilon_0 = 9.2 \times 10^{-11} Fm^{-1}$$

$$q = 1.6 \times 10^{-19} C$$

So CdTe doping
density is

$$N_a = 8.1 \times 10^{20} m^{-3} = 8.1 \times 10^{14} cm^{-3}$$



Key junction characterisation techniques

- JV – R_S and R_{SH} → can infer the issue
- J-V-T – Back contact barrier height measurements
- EQE – Layer behaviour and optical losses
- CV - doping density of p-type layer