

#### Ivan Bazarov

Physics Department, Cornell University

Fundamental processes in III-V photocathodes; application for high-brightness photoinjectors

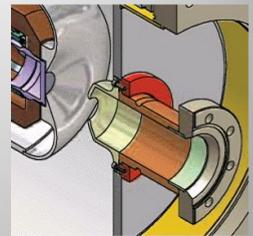
#### Contents

- Motivation
- NEA photoemission
- Some practical aspects
- Study cases: GaAs, GaAsP, GaN
- Summary



## Why are we interested?

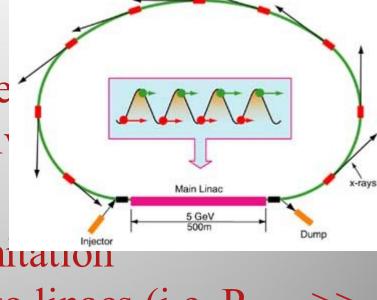
- Photoinjectors: a photocathode in high electric field (>> MV/m), either DC or RF
- Relativistic electrons can be further accelerated in a linac (linear accelerator) without degradation of beam brightness:
  - CW ultra-bright x-ray sources; high power FELs
  - Electron-ion colliders and ion coolers
  - Ultrafast electron diffraction, etc.





# Energy recovery linac

• Energy recovery linac: a new class of accele in activ



• Essentially removes the average current limitation

typical to linacs (i.e. P<sub>beam</sub> >>

• Average currents 10's to 100's of mA can be efficiently accelerated (and de-accelerated)



# Cathode figures of merit

QE and photon excitation wavelength

$$i(\text{mA}) = \frac{\lambda(\text{nm})}{124} \times P(\text{W}) \times \text{QE}(\%)$$

- E.g. 1W of 775 nm (Er-fiber  $\lambda/2$ )  $\Rightarrow$  6.2 mA/% 520 nm (Yb-fiber  $\lambda/2$ )  $\Rightarrow$  4.2 mA/% 266 nm (Nd-glass  $\lambda/4$ )  $\Rightarrow$  2.1 mA/%
- Transversely cold (thermalized) electron distribution
  - Directly sets the solid angle of the emitted electrons; an upper limit on achievable beam brightness



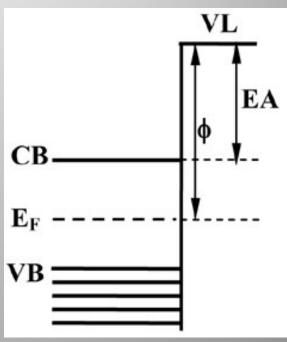
# Figures of merit (contd.)

- Prompt response time
  - A picosecond response is essential to take advantage of the space charge control via laser pulse shaping
- Long lifetime and robustness
  - Extraction of many 100's to 1000's of C between
    activations are necessary to make the accelerator practical



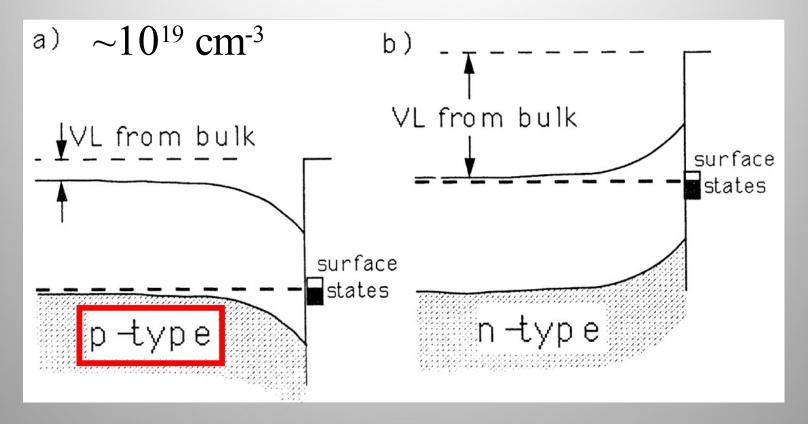
# Negative electron affinity

- Defined as vacuum level  $E_{vac}$  relative to the conduction band minimum
- Negative affinity: the vacuum level lies below the CBM
   ⇒ very high QE possible
- NEA:
  - -1) band bending
  - -2) dipole layer





### Strong p-doping

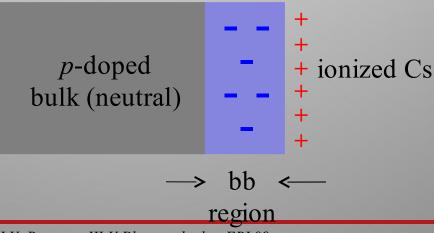


 Alperovich et al., Phys. Rev. B 50 (1994) 5480: clean p-doped GaAs has Fermi level unpinned and shows little band bending



# NEA: Cs ~monolayer

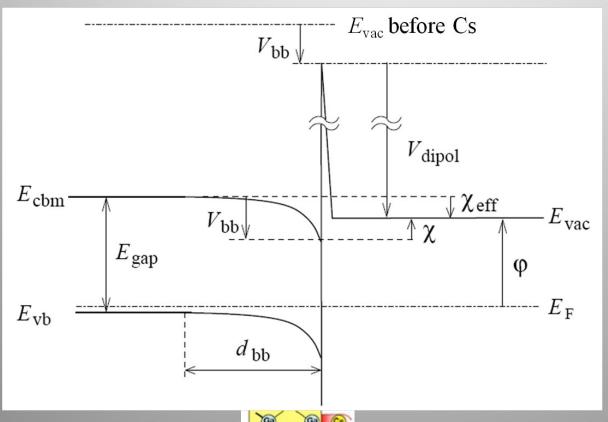
- Cs was found to play a larger role for NEA instead:
  - 1) band bending through donor surface states, and
  - 2) dipole surface layer from polarized Cs adatoms
    - Cs-induced donor-like surface states contribute their electrons to the bulk
  - Hole depleted region (negatively charged acceptors) lead to band bending region





## NEA: ~Cs monolayer (contd.)

 Majority of Cs atoms become only polarized (not ionized), forming a dipole layer (e- Cs+)



GaAs

$$E_{gap} = 1.42 \text{ eV}$$

Before Cs

$$\chi = 4 \text{ eV}$$

After Cs

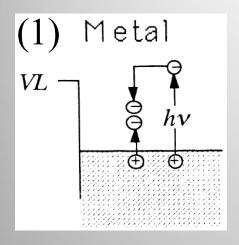
$$\chi_{\rm eff} \sim -0.1~{\rm eV}$$

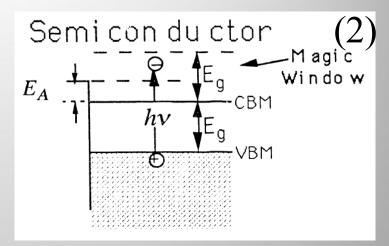
$$V_{bb} \sim 0.4 \; eV$$

$$d_{bb} \sim 10 \text{ nm}$$



# Spicer's magic window

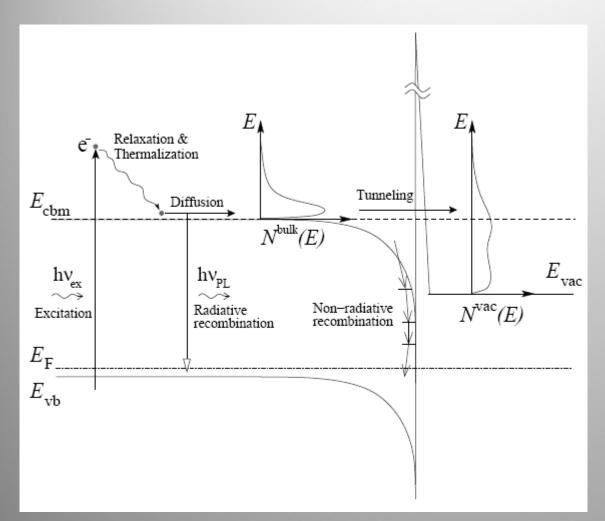




- (1) *electron-electron* scattering: typical of metals, large energy loss per collision
- (2) *electron-phonon* scattering: slowly depletes excessive energy of excited electron (LO phonons in GaAs  $\sim$  35 meV) "*Magic window*": in semiconductors, one needs excess KE > E<sub>gap</sub> for e-/e- scattering. Thus, electrons excited with E<sub>vac</sub> < KE < E<sub>VBM</sub> + 2E<sub>gap</sub> have excellent chances of escape



#### Electron transport processes



CBM thermalization time: 0.1-1ps

Electron-hole recombination: ~ns

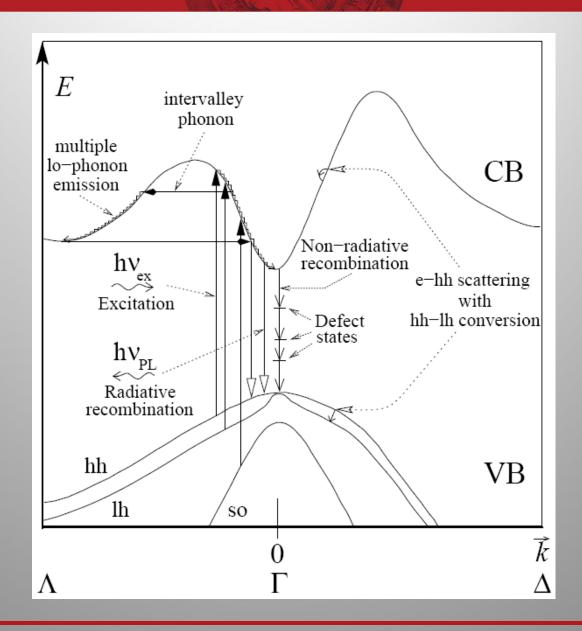
**Emission time:** 

 $\propto 1/(\alpha^2 D)$ 

strong wavelength dependence



## Energy vs. momentum

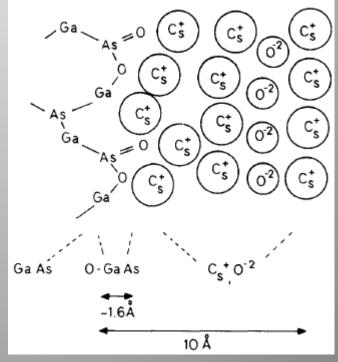




# Role of fluorine/oxygen

- Routine "yo-yo" activation employs O<sub>2</sub> or NF<sub>3</sub>
- Further reduction of affinity
   consistent
   with a double dipole model
- Stabilizes Cs on the surface; no lifetime or otherwise apparent advantage for either gas
- Bonded unstable nitrogen is found on Cs-NF<sub>3</sub> activated surfaces (APL 92, 241107)

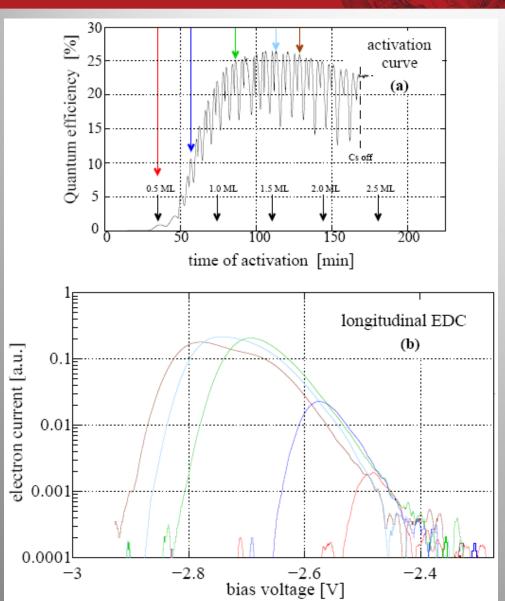




JAP 54 (1983) 1413



### GaAs: Optimal Cs coverage



laser wavelength: 670 nm

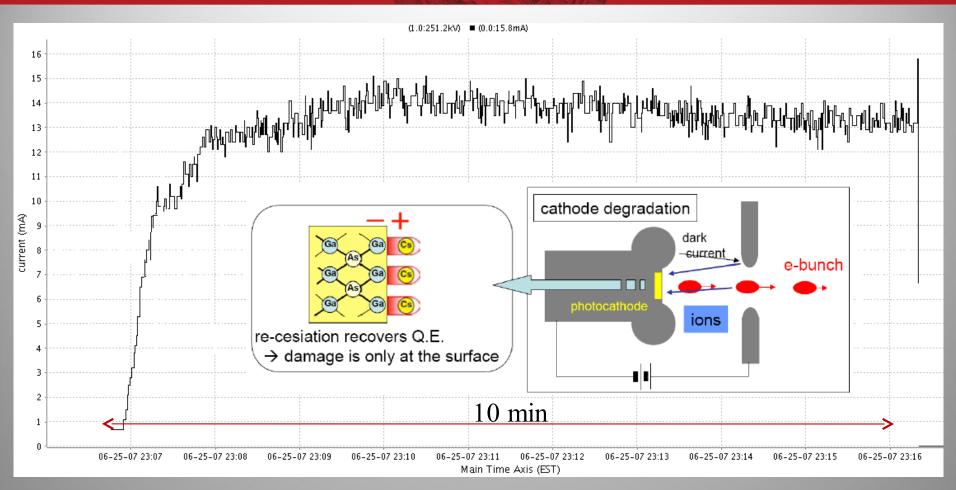
Ugo Weigel, PhD thesis



#### Lifetime matters

- H<sub>2</sub>O, CO<sub>2</sub> and O<sub>2</sub> can lead to chemical poisoning of the activated layer
- Low current ( $\sim 1\mu A$ ) 1/e lifetimes  $\sim 100$  hours typical in our prep chambers
- 3-5 times better in the DC gun (low 10<sup>-12</sup> Torr vacuum)
- High average current (mA's) lifetime limited by ion backbombardment





 ~5 hour lifetime (limited by gas backstreaming from the beam dump), i.e. 20 hours 1/e for 5 mA



# Study cases

- Our group has been evaluating III-V photocathodes
  - Transverse energy of electrons (thermal emittance)
  - Measure the photoemission response time
- Materials studied so far
  - GaAs @ 450-850nm: JAP 103, 054901; PRST-AB 11, 040702
  - GaAsP @ 450-640nm: Ibid
  - GaN @ 260nm: JAP 105, 083715

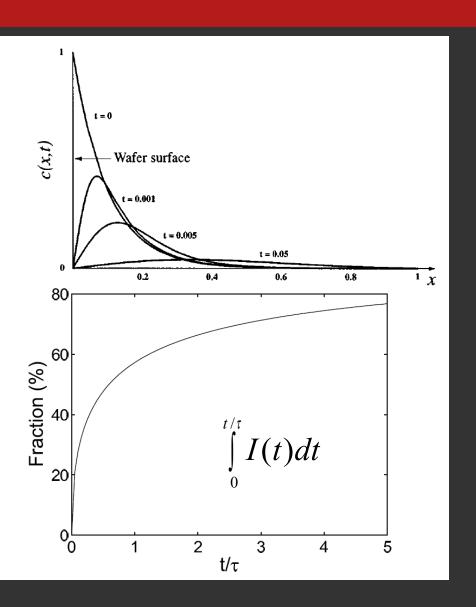
#### Diffusion model

$$\frac{\partial c(h,t)}{\partial t} = D \frac{\partial^2 c(h,t)}{\partial h^2}$$

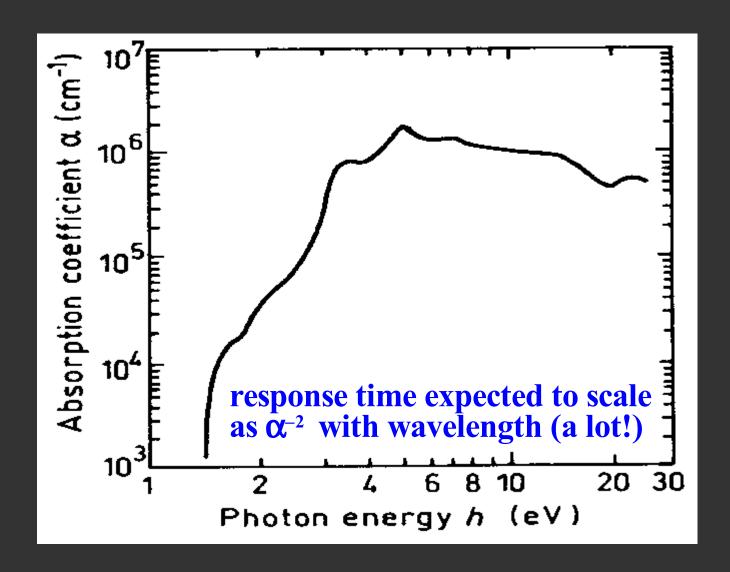
#### subject to:

$$c(h, t = 0) = c_0 e^{-\alpha h}$$
$$c(h = 0, t) = 0$$

$$\begin{split} &I(t) \propto \frac{\partial}{\partial t} \int\limits_0^\infty c(h,t) dh. \\ &I(\kappa) \propto \frac{1}{\sqrt{\pi \kappa}} - \exp(\kappa) \operatorname{erfc}(\sqrt{\kappa}) \\ &\kappa \equiv t/\tau, \text{ where } \tau \equiv \alpha^{-2} D^{-1} \end{split}$$



### **GaAs absorption**

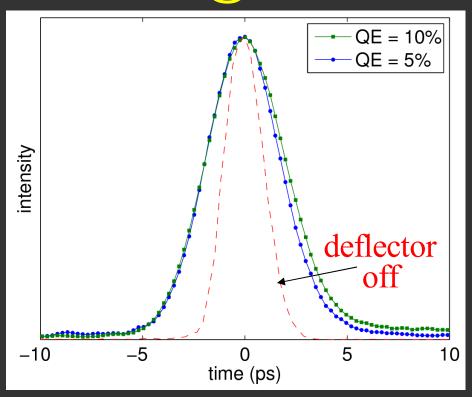


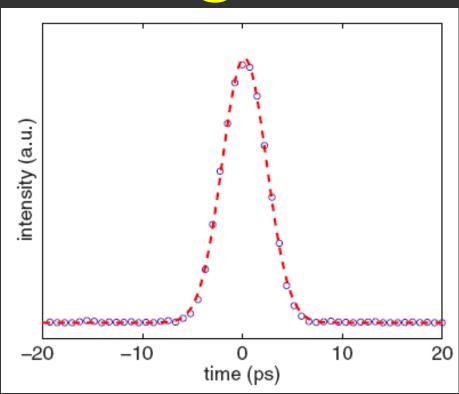


## Prompt emitters

#### GaAs @ 520 nm

## GaN @ 260 nm



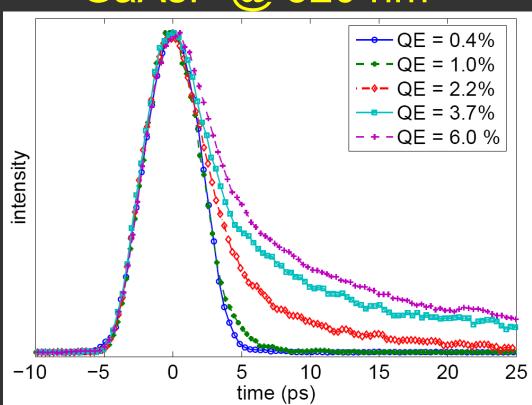


Measurements done by transverse deflecting RF cavity Limited by 1.8 ps rms resolution dominated by laser to RF synchronization



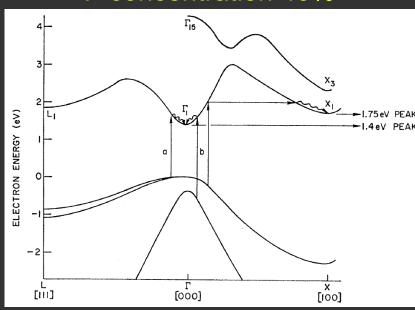
#### Diffusion tail

#### GaAsP @ 520 nm



# Strong QE dependency

#### P concentration 45%

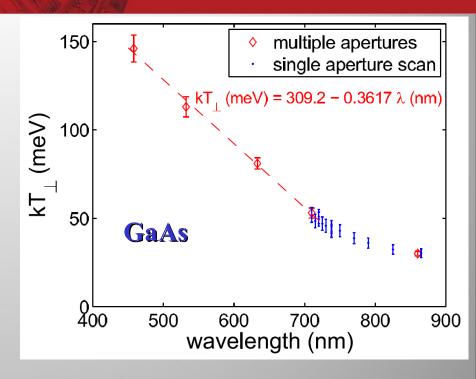


Two valleys:  $\Gamma$  (direct) and X (indirect) involved in the process



# Transverse energy distributions

- No surprises for bulk
  GaAs: cold electrons
  with a near band-gap
  excitation
- Surprisingly large transverse energy spread for GaN and GaAsP:



- GaAsP:  $kT_{\perp}$  = 130-240 meV for photons 0-780 meV photons above the band-gap
- GaN:  $kT_{\perp} = 0.9$  eV for photons with 1.4 eV above the band-gap



# Summary

- Transverse energy of photoelectrons remain poorly understood for III-V semiconductors (other than GaAs)
- More carefully controlled experimental data on transverse energy distributions/time response needed
- Predictive codes and models need to be developed and benchmarked with experiments
- This will allow photocathode engineering with the desired characteristics such as cold electrons with a ps response



# Acknowledgements

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