# TRANSVERSE MOMENTUM MEASURING SYSTEMS: THE CRYOGENIC MOMENTATRON AND TESS

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### Abstract

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Photocathode performance optimization is crucial to the operation of X-ray Free Electron Laser (FEL) facilities such as LCLS-II and the future UK XFEL as the initial photoelectron phase-space distribution directly impacts x-ray generation efficiency in the undulator. Key photocathode parameters such as transverse emittance and quantum efficiency are routinely measured to monitor and improve performance, but other metrics are influenced by the photocathode material such as operating temperature, vacuum and RF environmental conditions and the characteristics of laser illumination (e.g., wavelength, and pulse energy, duration, and repetition rate). To better understand these factors, SLAC has developed a laboratory-based Cryogenic Momentatron system to measure the emission properties of photocathodes at temperatures between 30 K and 300 K using 200 – 800 nm wavelength-tunable light. The system's design and initial commissioning performance have been validated through simulations and error analysis, confirming its reliability for photocathode characterization. We present measurements taken at SLAC, along with complementary data for Cs<sub>2</sub>Te obtained using the TESS system at Daresbury Laboratory a similar photocathode characterization tool. These results provide valuable insights into photoemission physics that may be leveraged to optimize photocathode performance and thus that of X-ray FELs.

### **ENERGY/MOMENTUM MEASUREMENT**

Laser-driven photocathode electron sources are essential components of modern advanced x-ray generation systems, including free-electron lasers (FELs), instruments for ultrafast electron diffraction (UED) and dynamic transmission electron microscopy (DTEM). In all these applications, the ultimate performance is determined and limited by the transverse emittance of the electron beam at the photocathode. The emittance is typically defined as a quantity that remains invariant under acceleration to relativistic energies, referred to as the normalized emittance:

So, 
$$\varepsilon_{n,rms} = \frac{1}{mc} \sqrt{\langle x^2 \rangle \langle p_x^2 \rangle} = \frac{\sigma_x \sqrt{\langle p_x^2 \rangle}}{mc}$$
, (1)

where  $\sigma_x$  is the rms transverse size of the beam,  $p_x$  is the transverse momentum, m is the electron mass and c the speed of light. Eq. (1) suggests that emittance can, in principle,

be reduced indefinitely by decreasing the beam size. However, in practice, this reduction is fundamentally limited by space-charge effects. As a result, for practical applications, lowering the transverse momentum is the only effective route to minimizing emittance. For many light source applications, such as FELs, achieving a normalized emittance below  $0.5 \, \mu \text{m/mm}$  is a critical performance target. Using the three-step framework, Dowell and Schmerge [1] expressed the normalized emittance in terms of the mean transverse energy (MTE) and quantum efficiency (QE) as:

$$MTE(\lambda) = \frac{E_{excess}}{3},$$
 (2)  $QE(\lambda) \propto E_{excess}^2,$ 

where  $E_{excess} = \hbar\omega - \theta$  with  $\hbar\omega$  being the photocathode illumination photon energy and  $\phi$  the photocathode work function. This model was later extended by Vecchione [2] to account for the effects of photocathode temperature and further refined by Saha [3] to include the influence of surface disorder (roughness). The temperature dependence of the rms transverse momentum is illustrated in Fig. 1, reproduced from [2]. The figure shows that reducing the photocathode temperature leads to a lower rms transverse momentum. Exploiting near-threshold photoemission at cryogenic temperatures has enabled record-low energy spreads, such as the 5 meV value reported for a copper photocathode [4].

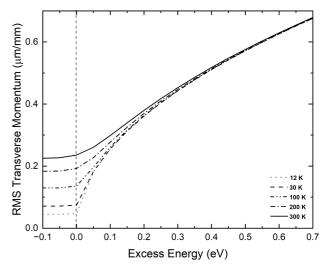


Figure 1: The effect of excess energy and temperature on transverse momentum during photoemission.

The primary motivation for developing these instruments is to identify photocathodes exhibiting low rms transverse

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# LCLS-II PHOTOCATHODE **DEPOSITION CHAMBER**

The LCLS-II photocathode deposition system shown in Fig. 2 includes a particle-free load-lock, air-lock, and a photocathode transport suitcase connected to the air-lock, all maintained under extreme high vacuum (XHV) conditions. The growth chamber is equipped with four physical vapor deposition (PVD) sources that support a variety of chemical species and allow for both sequential and co-deposition processes. Photocathodes are deposited onto molybdenum INFN-type plugs, with a 3 mm diameter active area defined by an aperture positioned in front of the cathode. During Cs<sub>2</sub>Te growth, the photocathode is illuminated at 258 nm to monitor QE, while quartz crystal microbalances (QCMs) are used to estimate the deposited material thickness. The resulting Cs<sub>2</sub>Te photocathodes typically exhibit QE values exceeding 10%, and remain stable for several months in the transport suitcase under XHV conditions. Notably, the most recent Cs<sub>2</sub>Te photocathode, BF01, operated for approximately 1,054 hours, during which its QE gradually degraded from around 8 % to 5 %, but remained sufficient to support beam delivery throughout this period.

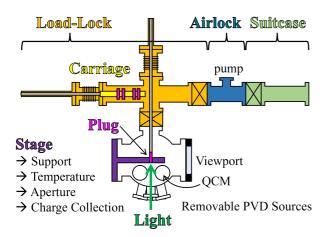


Figure 2: Schematic of the LCLS-II Cs<sub>2</sub>Te photocathode deposition system at SLAC.

## THE CRYOGENIC MOMENTATRON

The Cryogenic Momentatron at SLAC is specifically designed to characterize photocathode performance for the LCLS-II prior to operational deployment. In addition to qualification measurements, the system provides critical data for developing advanced photoemission models, including the influence of phonons and carriers on the emission process, as well as the effects of residual gas molecule physisorption

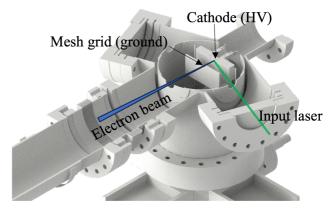


Figure 3: Schematic of the Cryogenic Momentatron.

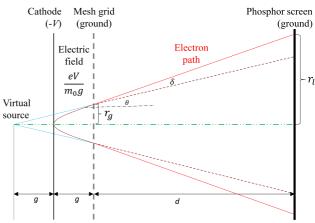


Figure 4: Schematic of the system used to measure transverse momentum. V = applied voltage, 20 kV maximum; g =cathode-anode (grid) gap,  $0.020 \,\mathrm{m}$ ;  $r_{\varrho}$  = radial coordinate on the grid w.r.t. nearest grid hole center;  $\delta$  = angular kick from the mesh-lensing effect;  $d = \text{drift distance}, 0.980 \,\text{m}; r_1$ = photoelectron radial coordinate on the detector, < 5 mm.

on quantum efficiency (QE) and mean transverse energy (MTE) which is directly related to transverse momentum.

The physical layout of the system is shown in Fig. 3. The Momentatron's load-lock chamber allows connection to a photocathode transport suitcase and uses magnetic transfer arms to move cathodes from the suitcase to the plug holder. The measurement head comprises a stretched electroformed mesh grid mounted in a ring, a sapphire electrical isolator, temperature sensors, and a grounded metal cylindrical body that supports the cathode plug receiver. The photocathode holder is connected to a high-voltage feedthrough, with the mesh grid and cylindrical body held at ground potential.

On the mesh side, electrons accelerated through the grid drift to a microchannel plate-phosphor image intensifier, and the resulting light is recorded by a cooled Andor Xion 888 EMCCD camera ( $1024 \times 1024$  pixels,  $13 \mu m$  pixel size). A dipole magnet near the chamber deflects the drifting electron beam to place it within the camera's field of view. Photocathodes are cooled using an Advanced Research Systems DE-215 cryocooler which provides 30 W of cooling at 45 K and 1.5 W at 4.2 K.

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The concept of the measurement system is illustrated in Fig. 4. Electrons photoemitted from the cathode are accelerated by a cathode–anode voltage V applied across a small gap g toward a mesh grid anode. The electrons pass through the grid and then drift over several hundred millimeters to a detector. In this setup, the photocathode is typically held at a negative potential, while the grid and detector are grounded.

Within the cathode–anode gap, electrons follow parabolic trajectories and reach the grid with a lateral position  $r_g$  and an angle  $\theta$ . Upon traversing the grid, the electrons experience an additional angular deviation  $\delta$  due to the mesh grid lensing effect, analogous to the Davisson and Calbick lens phenomenon, as described by Yu  $et\ al.$  [8]. This lensing effect can be approximated by

$$\delta = \frac{r_g}{8\varrho},$$

where  $r_g$  is the radial coordinate relative to the nearest grid hole center, and g is the cathode–anode gap.

After passing through the grid, the electron beam drifts toward the detector. The radial coordinate of electrons on the detector,  $r_l$ , is related to their transverse momentum  $p_x$ , the cathode–grid gap g, the drift distance d, and the applied voltage V by the relation Eq. (3)

$$p_x = \frac{r_l}{2g+d} \sqrt{2m_0 eV}.$$
 (3)

For the chosen parameters, the intrinsic emittance can be estimated from the root-mean-square radial coordinate of electrons on the detector  $(r_{\rm rms})$  as

$$\varepsilon_{\rm int} \ [\mu {\rm m/mm}] \equiv \frac{\varepsilon_{n,{\rm rms}}}{\sqrt{\langle x^2 \rangle}} \approx 0.061 \sqrt{V \ [{\rm kV}]} \ r_{\rm rms} \ [{\rm mm}].$$

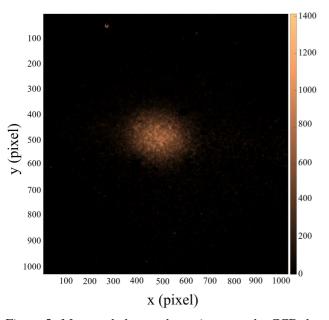


Figure 5: Measured electron beam image at the CCD detector for a copper cathode with a 7 kV accelerating voltage and 4.542 eV photon energy ( $\lambda=273\,\mathrm{nm}$ ),  $g=0.020\,\mathrm{m}$  and  $d=0.980\,\mathrm{m}$ . The Intrinsic emittance is  $0.32\,\mu\mathrm{m/mm}$ .

### CRYOGENIC MOMENTATRON RESULTS

To test the system, a copper cathode was inserted into the chamber. Illumination was provided by light from a high-power xenon arc lamp coupled to a monochromator, which was fiber-coupled to deliver photons onto the cathode, inducing photoemission.

Figure 5 shows a typical measured round electron beam image on the CCD detector at an accelerating voltage of  $7 \, \text{kV}$  and a photon energy of  $4.542 \, \text{eV}$  ( $\lambda = 273 \, \text{nm}$ ). In other measurements, the beam diameter was seen to decrease as the photon energy approached the cathode work function. Additionally, the beam width at the detector can be controlled by varying the acceleration voltage, which is typically adjustable between 1 and  $20 \, \text{kV}$ .

Preliminary data obtained using the Cryogenic Momentatron system are presented in Fig. 6. The experimental results clearly indicate that reducing the temperature results in lower intrinsic emittance, particularly as the excess energy approaches zero in the region close to the photoemission threshold. A substantial amount of additional data has been taken using the Cryogenic Momentatron to study a number of photocathode/photoemissive systems, but this has not yet been fully analyzed. Publication of this work is expected in the coming months.

More comprehensive data will be collected in the future using an NKT Supercontinuum laser which provides illumination wavelengths from 265 nm to over 2000 nm, with pulse lengths under 2 ns and a repetition rate ranging from 1 Hz to  $20\,\mathrm{kHz}$ .

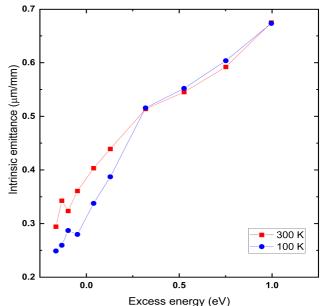


Figure 6: Variation in the intrinsic emittance as a function of excess energy for a copper cathode at temperatures of 100 K and 300 K, with a 7 kV accelerating voltage and a 4.542 eV photon energy ( $\lambda = 273$  nm), g = 0.020 m and d = 0.980 m.

### THE TESS SYSTEM

The TESS system [6, 7] shown conceptually in Fig. 7 has been developed separately to, but bears many similarities to the Momentatron systems [5]. Both systems exploit the transverse displacement from an axial reference path of multiple non-relativistic electrons to generate a measureable photoemission footprint from which the transverse energy (momentum) component is extracted. Notable differences are the typical cathode–anode operating potential differences which are 50-100 V in TESS compared to 1-20 kV for the Momentatron, and the distances through which the electrons propagate. While the Momentatron uses either a xenon arc lamp or a pulsed laser system for cathode illumination, the TESS uses a broadband laser-driven plasma lightsource<sup>1</sup> and monochromator<sup>2</sup> combination to provide a tunable CW light source for photocathode illumination. Like the Cryogenic Momentatron, the TESS system incorporates a liquid nitrogen cooling loop which can reduce the photocathode temperature to around 190 K, so reducing the minimum energy spread (or thermal floor) fixed by  $k_BT$  from around 25 meV at room temperature to around 18 meV. The TESS also features a piezo-electric fine leak valve which has facilitated detailed studies of photocathode response to controlled degradation to poisoning gas species which modify the surface electronic structure, and hence the photoemission properties. TESS has been used to study photocathode physics for semiconductors, both GaAs [9–11] and Cs<sub>2</sub>Te [12, 13], and also for metals [14–17]. The data provided by TESS has improved our understanding of photocathode and photoemission physics, particularly the semiconductor class.

Figure 8 shows the impact on photoemission characteristics for a Cs<sub>2</sub>Te photocathode under illumination at a constant wavelength of  $\lambda = 266 \,\mathrm{nm}$  while subjected to progressive degradation firstly by nitrogen (the control) and then by oxygen (the poison) [13]. The plot demonstrates that the photocathode characteristics do not change under prolonged exposure to nitrogen as this gas species is inert, but that exposure to oxygen (a reactive species) drives an immediate change in both MTE and QE. The data show a progressive evolution in photoemission properties during the integrated exposure to more than 70 L of oxygen, with the MTE increasing while the QE falls. The fundamental emission properties of any Cs<sub>2</sub>Te photocathode must therefore change during prolonged operation in a photoinjector electron source. This is highly undesirable, but exposure to residual oxygen is unavoidable in any practical vacuum system.

Figure 9 illustrates the impact of this degradation on the 'clean' and 'degraded' performance of the  $Cs_2$ Te photocathode [13]. The data show how the MTE and QE vary as a function of illumination wavelength,  $\lambda$ , and includes Dowell-Schmerge fits [1]. These imply a work function of slightly less than 4.0 eV for the 'clean' photocathode which is higher than the accepted value of 3.5 eV for  $Cs_2$ Te, based on a 3.3 eV

bandgap with a 0.2 eV positive electron affinity [18], falling to something close to 3.5 eV once degraded by exposure to oxygen. The plot also shows that photoemission is detected under illumination wavelengths far beyond the expected photo emission threshold of 3.5 eV ( $\lambda = 350 \,\mathrm{nm}$ ), albeit with extremely low and eventually vanishing levels of QE. The data demonstrates two challenges encountered when depositing Cs<sub>2</sub>Te photocathodes. Firstly, the difficulty in creating just the Cs<sub>2</sub>Te compound as Cs and Te are known to form a family of compounds Cs<sub>x</sub>Te<sub>y</sub> which have different work functions [19–21]. Secondly, the consequence of incorrect stoichiometric ratios during deposition resulting in an excess of Cs in the photoemissive layer which duly reacts with residual oxygen to yield CsO. Both Cs and CsO have very low workfunctions and so generate measurable emission in the TESS at 532 nm. This problem can be mitigated through the use of a multi-wavelength QE monitoring system during photocathode deposition [22]. Such a system provides QE at a number of different sequential wavelengths such as 405 nm and 532 nm, with any measurable photocurrent at these longer wavelengths highlighting deviations from correct stoichiometry arising from an excess of caesium.

# **FUTURE WORK AND CONCLUSIONS**

Future upgrades to the TESS would include improved photocathode cooling to access lower temperatures, with increased light flux at shorter illumination wavelengths, specifically in the range 230 to 260 nm. Accurate measurement of the optical power reflected by the photocathode would add the ability to determine photocathode reflectivity and so assess the impact of surface modifications to increase photon absorption, and thereby QE. The ability to directly measure absolute QE in-situ is also a future aspiration.

The results presented in this paper demonstrate the importance of understanding the subtle behaviors of photocathodes of all classes, and recognizing that their performance is not a constant during accelerator operation. We hope that the importance of the continued development of systems such as the Cryogenic Momentatron and TESS is self—evident to support progress towards the generation of electron beams with the highest possible levels of brightness.

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<sup>&</sup>lt;sup>1</sup> Energetiq EQ-99X Laser-Driven Light Source

<sup>&</sup>lt;sup>2</sup> Bentham TMc300 Single Monochromator

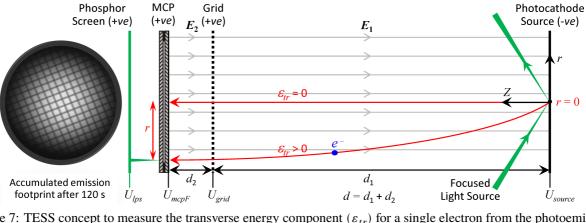


Figure 7: TESS concept to measure the transverse energy component  $(\varepsilon_{tr})$  for a single electron from the photoemission footprint generated by  $\approx 10^7$  electrons accumulated over a period of 120 s with a typical footprint shown to the left [7].

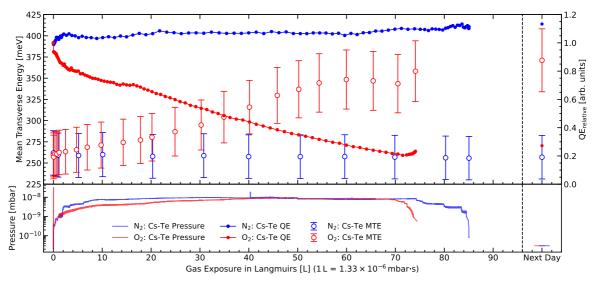


Figure 8: MTE ( $\circ \& \circ$ , left axis) and relative QE ( $\bullet \& \bullet$ , right axis) for a Cs<sub>2</sub>Te photocathode illuminated at  $\lambda = 266$  nm and subjected to progressive degradation by more than 70 L of oxygen (red data) and nitrogen (blue data) [13].

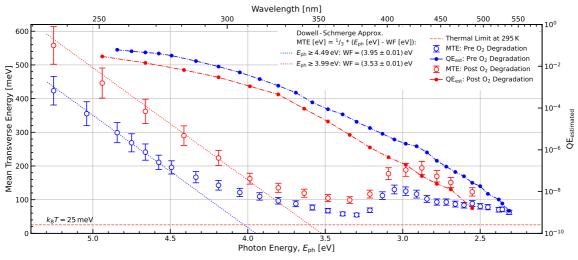


Figure 9: MTE (o & o, left axis) and estimated QE (o & o, right axis) for a Cs<sub>2</sub>Te photocathode under illumination with  $\lambda = 236$  to 536 nm before degradation (blue data) and after degradation (red data) by more than 70 L of oxygen [13].

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