

*Photocathode Physics for Photoinjectors (P3) 2018:*

# **Synthesis and x-ray characterization of Cesium Telluride photocathodes**

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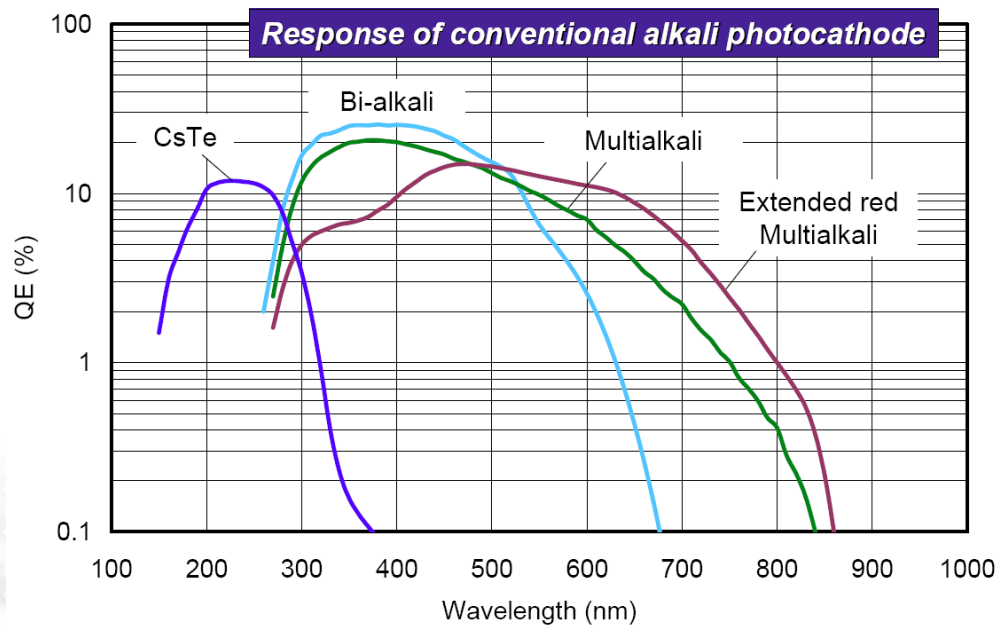
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**BROOKHAVEN**  
NATIONAL LABORATORY



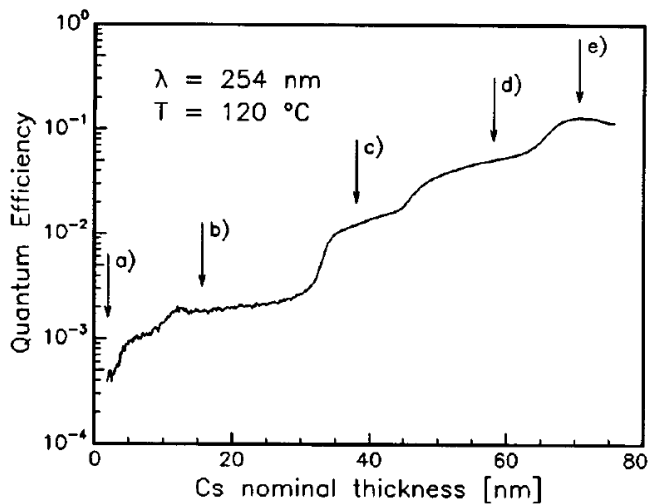
# Introduction: CsTe photocathode as electron source

- Cesium telluride (CsTe) photocathodes has been the first hand choice for electron sources by worldwide accelerators, such as LCLSII, AWA, various FELs, etc...
- Perfect balance between lifetime and quantum efficiency
- Less requirement of vacuum level than GaAs and multialkali photocathodes, robust in high gradient environment



# Introduction: Efforts in characterization of CsTe photocathode

- Growth recipe: mostly sequential. 10 ~ 20 nm of Te + 60 ~ 80 nm of Cs @ 120 °C → QE : 15 % ~ 18% @ 250 nm



A. di Bona *et al*, Auger and x-ray photoemission spectroscopy study on Cs<sub>2</sub>Te photocathodes, *Journal of Applied Physics* 80 3024 (1996)

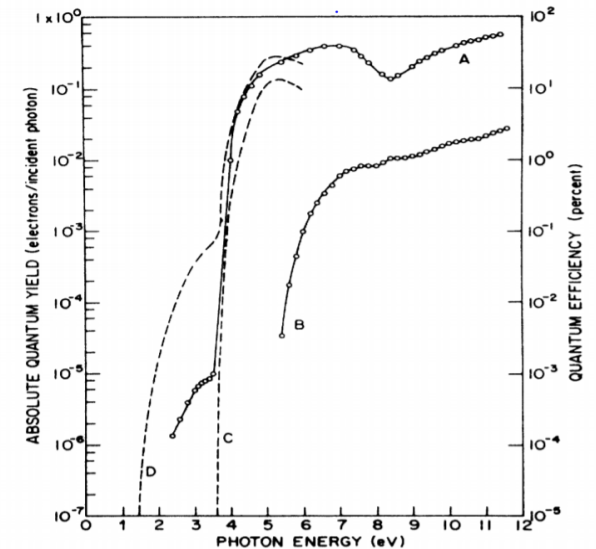
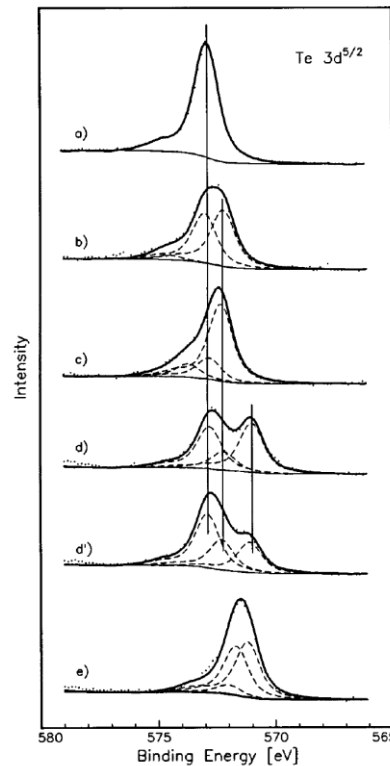


FIG. 1. Spectral distribution of the photoelectric yield from (A) Cs<sub>2</sub>Te; (B) evaporated Te before reaction with Cs. Also shown are the data of Taft and Apker (Ref. 1). In their work, curve C was reported to be from a surface containing excess Cs while curve D was exposed to less Cs.

# Question remained

- Many stable compounds exist besides CsTe and Cs<sub>2</sub>Te: Cs<sub>3</sub>Te<sub>2</sub>, Cs<sub>2</sub>Te<sub>3</sub>, Cs<sub>2</sub>Te<sub>5</sub>, CsTe<sub>4</sub>, Cs<sub>5</sub>Te<sub>4</sub>.....
- The form of each stoichiometric compound specifically depends on the growth conditions:
  - substrate temperature,
  - growth rate of each material
  - Te thickness,
  - etc...

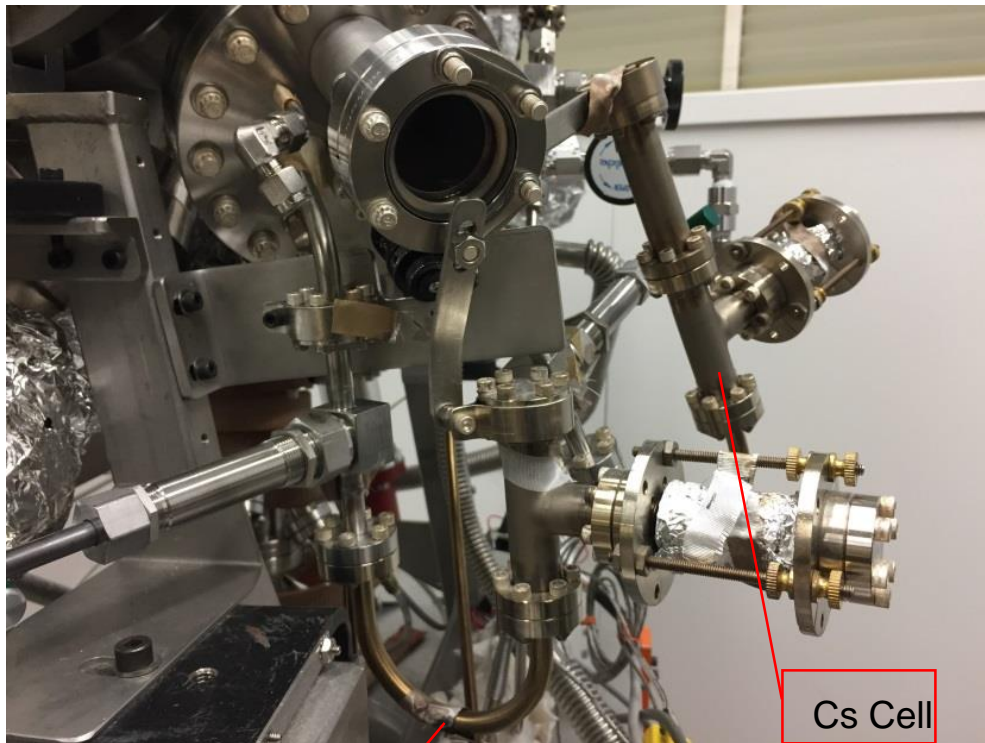
Materials Id	Formula	Spacegroup
mp-8361	CsTe	Pbam
mp-505464	Cs <sub>2</sub> Te <sub>13</sub>	Pbcm
mp-573763	Cs <sub>2</sub> Te	Pnma
mp-505634	Cs <sub>2</sub> Te <sub>3</sub>	Cmc2 <sub>1</sub>
mp-620471	Cs <sub>3</sub> Te <sub>22</sub>	P1

<https://materialsproject.org/>



# Cs-Te cathode growth with effusion cell

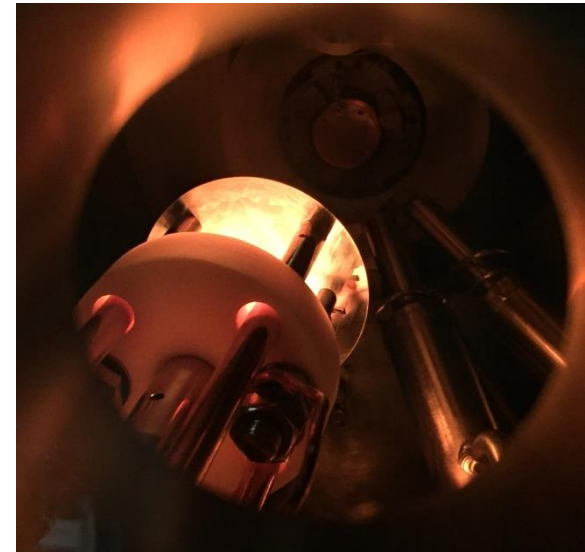
*Cs effusion cell*



Cs Cell

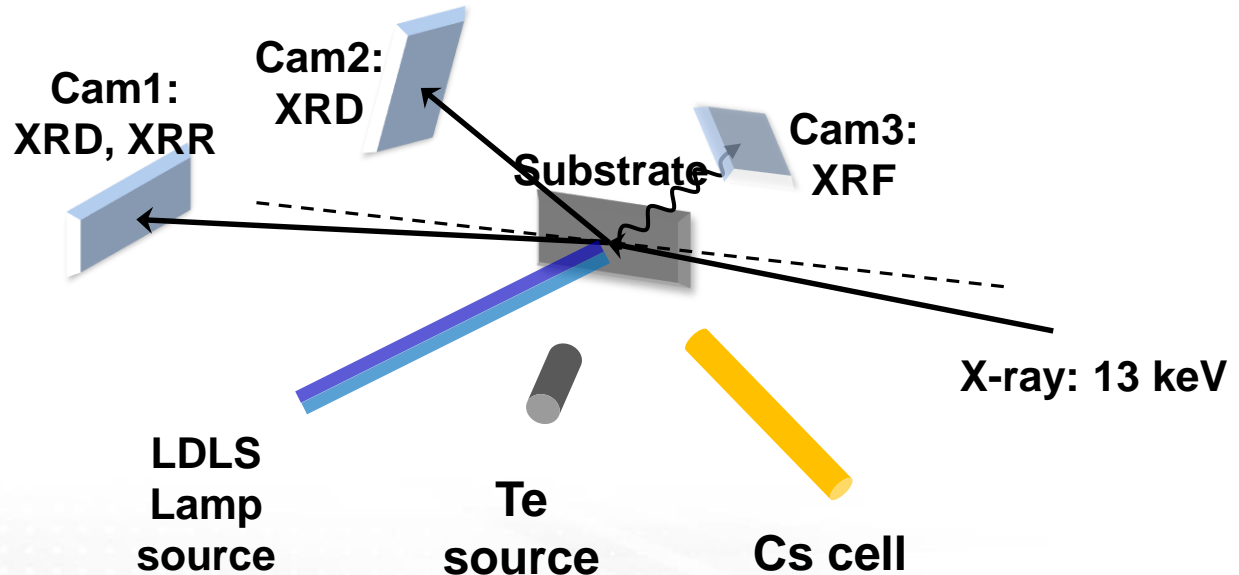
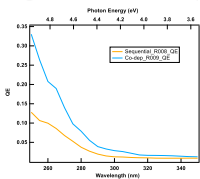
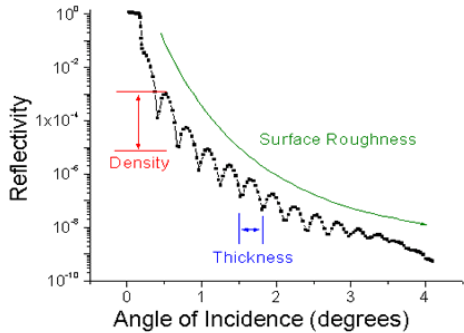
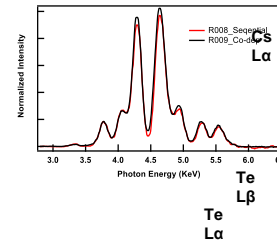
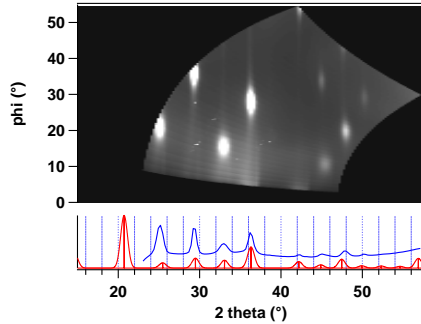
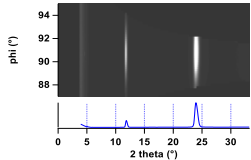
J tube

*Te evaporator*

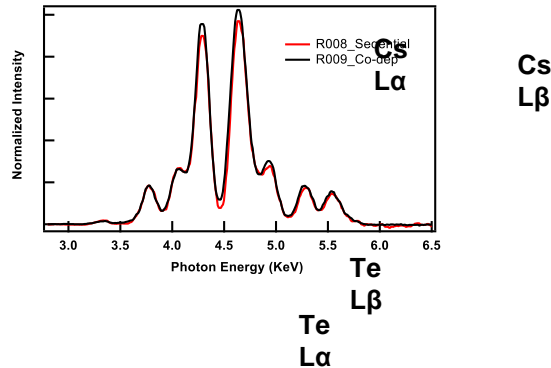
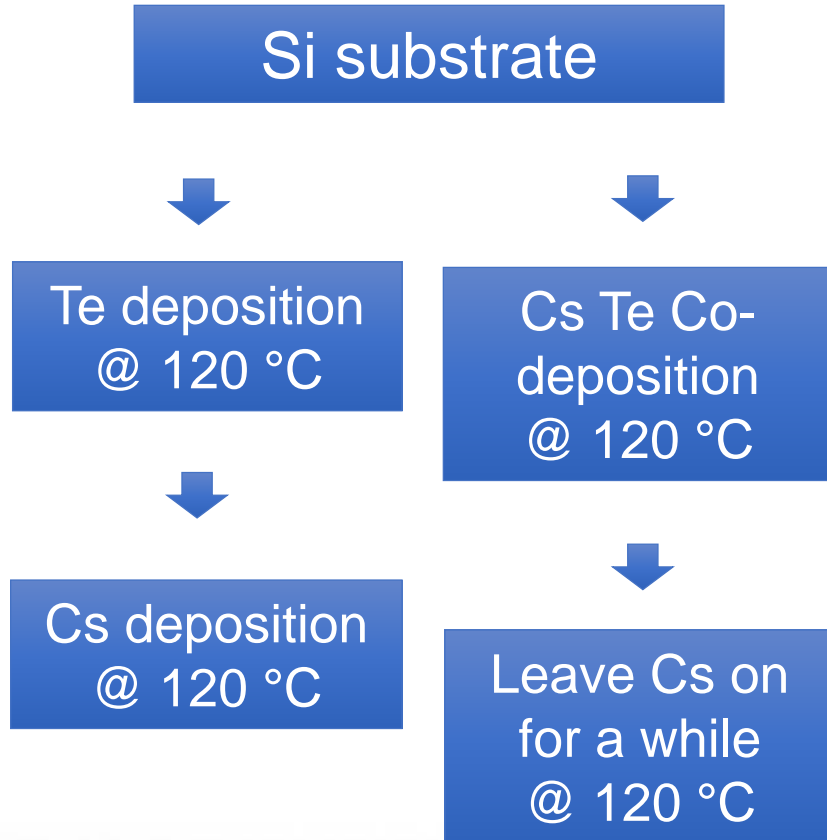


- Growth rate are controlled by J tube temperature, valve and shutter

# Experimental set-up: Cs-Te cathode growth



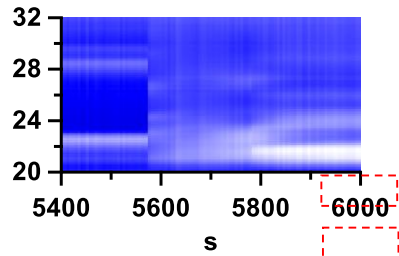
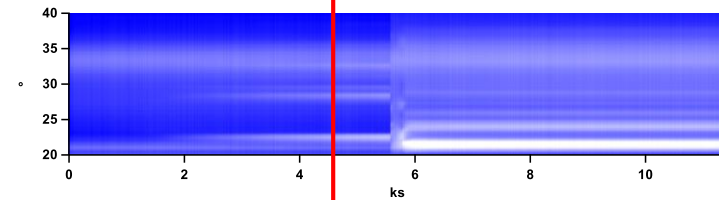
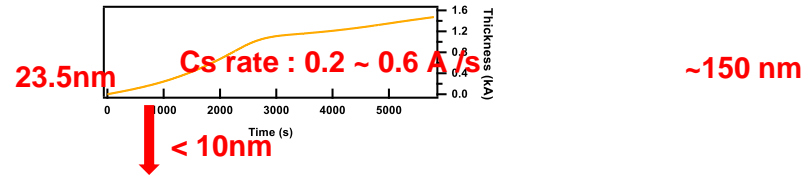
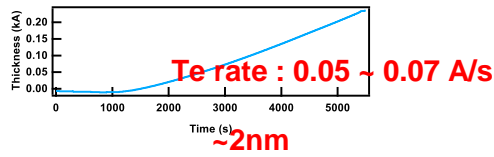
# Sequential vs Co-dep CsTe -- XRF



Sample	Te	Cs (+/-0.05)
R008 Sequential	1.00	1.15
R009 Co-dep	1.00	1.3

- At same substrate temperature; co-dep method incorporates more Cs.

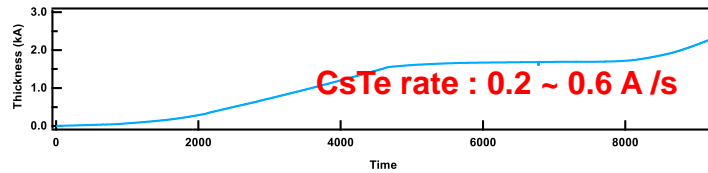
# Realtime analysis : Sequential growth



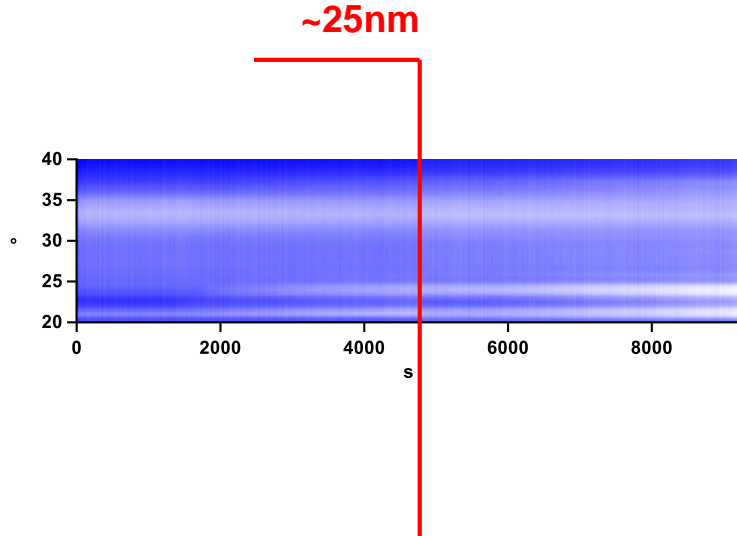
In less than 10 nm of Cs deposition, at least 2 intermediate phases appeared and disappeared, then a stable product of CsTe forms.



# Realtime analysis : Co-deposition



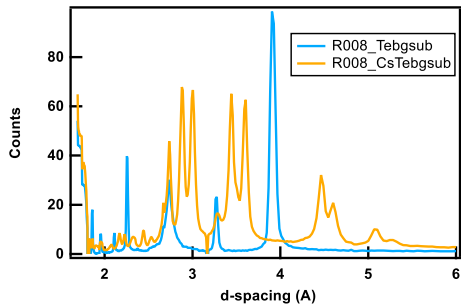
~230nm



- Starts to crystallize ~ 25 nm of total thickness.
- Same phase throughout the growth.

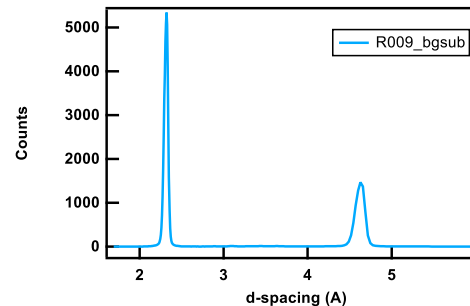
# CsTe cathode structure: XRD analysis

## Sequential growth



- Nearly all of the crystalized Te are dissolved.
- Low counts in diffraction peaks
- Multiple phase of Cs-Te compound co-exist

## Co- deposition



- Well crystalized.
- Single phase

## Sequential growth

Te: 23.5 nm

HKL	Theory_d spacing (Trigonal)	Exp_d spacing
2-1-10	2.256	2.259
10-11	3.268	3.27
10-10	3.908	3.91

Cs : > 150 nm

HKL	Theory_d spacing	Exp_d spacing
132 ( $\text{Cs}_2\text{Te}_3$ )	2.85	2.87
203 ( $\text{Cs}_2\text{Te}$ )	3.005	3.005
131 ( $\text{Cs}_2\text{Te}_3$ )	3.45	3.44
103 ( $\text{Cs}_2\text{Te}$ )	3.59	3.60
201 ( $\text{Cs}_2\text{Te}$ )	4.41	4.46
111 ( $\text{Cs}_2\text{Te}$ )	4.61	4.60
021 ( $\text{Cs}_2\text{Te}_3$ )	5.08	5.08
011 ( $\text{Cs}_2\text{Te}$ )	5.27	5.28

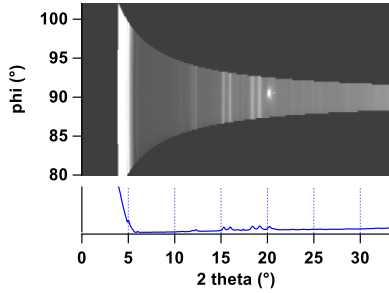
## Co- deposition

~ 100 nm

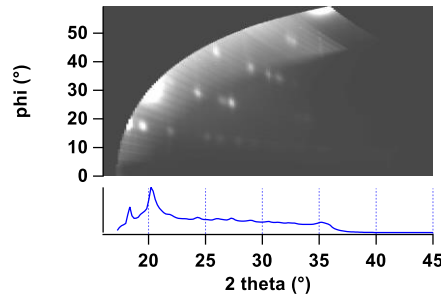
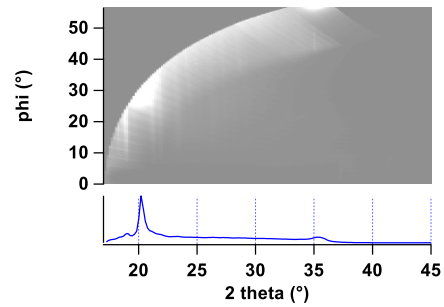
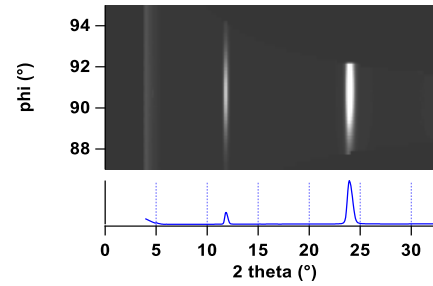
HKL	Theory_d spacing	Exp_d spacing
222 $\text{Cs}_2\text{Te}$	2.307	2.315
111 $\text{Cs}_2\text{Te}$	4.613	4.615

- Sequential cathode is consist of possibly  $\text{Cs}_2\text{Te}_3$  and  $\text{Cs}_2\text{Te}$
- The crystalized phase in the co-dep cathode is  $\text{Cs}_2\text{Te}$

# Sequential growth

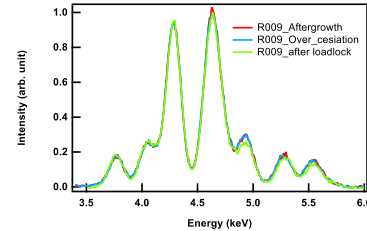
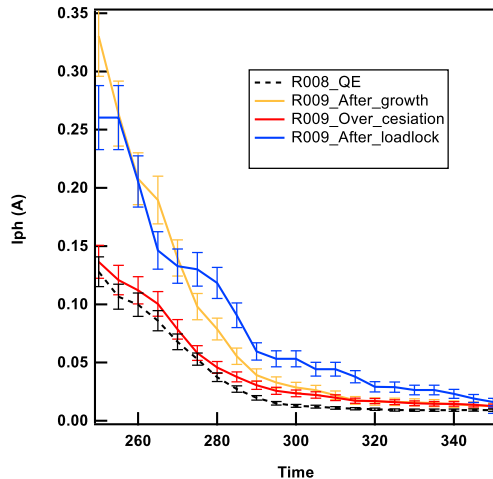


# Co-deposition



**Highly textured CsTe phase!**

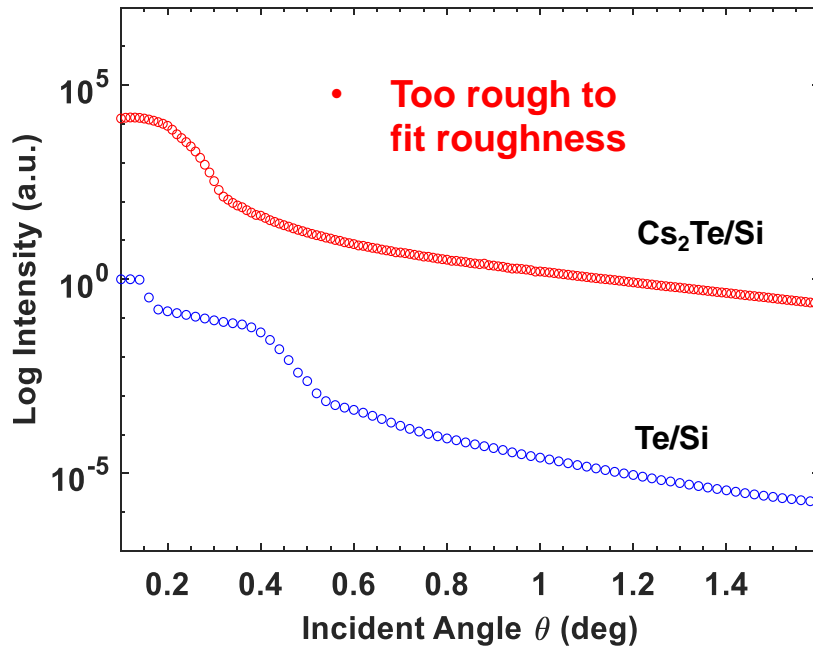
# CsTe cathode Quantum efficiency



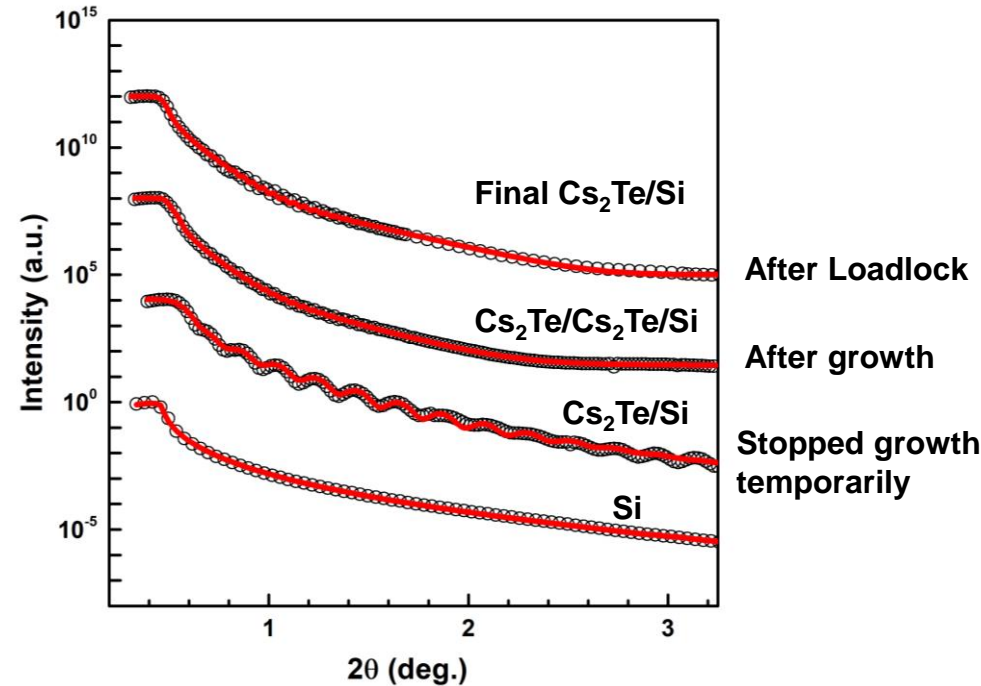
- **Co-dep can result in much higher QE than sequential**
- **over cesiation might lead to Cs build-up on the cathode surface and lower the QE**
- **QE can be recovered by removing the access Cs**

# CsTe cathode surface roughness: XRR analysis

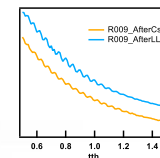
R008 Sequential - CsTe on Si



R009 Co-dep - CsTe on Si



	Thickness ( $\text{\AA}$ )	Roughness ( $\text{\AA}$ )
FINAL $\text{Cs}_2\text{Te/Cs}_2\text{Te}$	$968.3 \pm 2.9$ (total $\text{Cs}_2\text{Te}$ )	$19.1 \pm 0.2$
$\text{Cs}_2\text{Te/Cs}_2\text{Te}$	$1026.1 \pm 1.6$ (total $\text{Cs}_2\text{Te}$ )	$19.10 \pm 0.07$
$\text{Cs}_2\text{Te}$	$245.5 \pm 1.7$	$9.55 \pm 0.14$
Si Substrate	-	$3.75 \pm 0.02$





# Summary

We presented results of real time structural analysis and in situ XRF, XRR and spectral response measurements.

The cathode from our deposition procedure yields stoichiometry that is less than Cs: Te = 2:1, The comparison between sequential and co-dep shows that the co-dep method incorporates more Cs than sequential method

CsTe photocathodes grown by pure alkali effusion cells and co-deposition method have stable and highly textured crystal structure and ultra smooth surface

QE of Co-dep cathode is higher in both high energy and low energy region. Excess Cs might result in lower QE.

# Thanks for your attention!

- Thanks to all the contributors!
- Staff @ CHESS and beamline G3
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