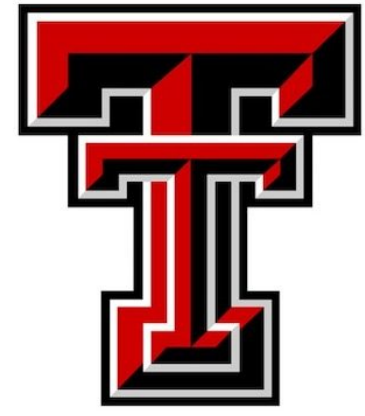




ETH Institute for
Particle Physics



Istituto Nazionale di Fisica Nucleare
SEZIONE DI TORINO

Cerium-doped fused-silica fibers for particle physics detectors

IPRD19

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N. Akchurin¹, N. Bartosik⁴, J. Damgov¹, F. De Guio¹, G. Dissertori³, E. Kendir², S. Kunori¹, T. Mengke¹,
F. Nessi-Tedaldi³, N. Pastrone⁴, S. Pigazzini³, S. Yaltkaya²

¹Texas Tech University, Department of Physics and Astronomy, USA

²Akdeniz University, Department of Physics, Turkey

³ETH, Zurich, Switzerland

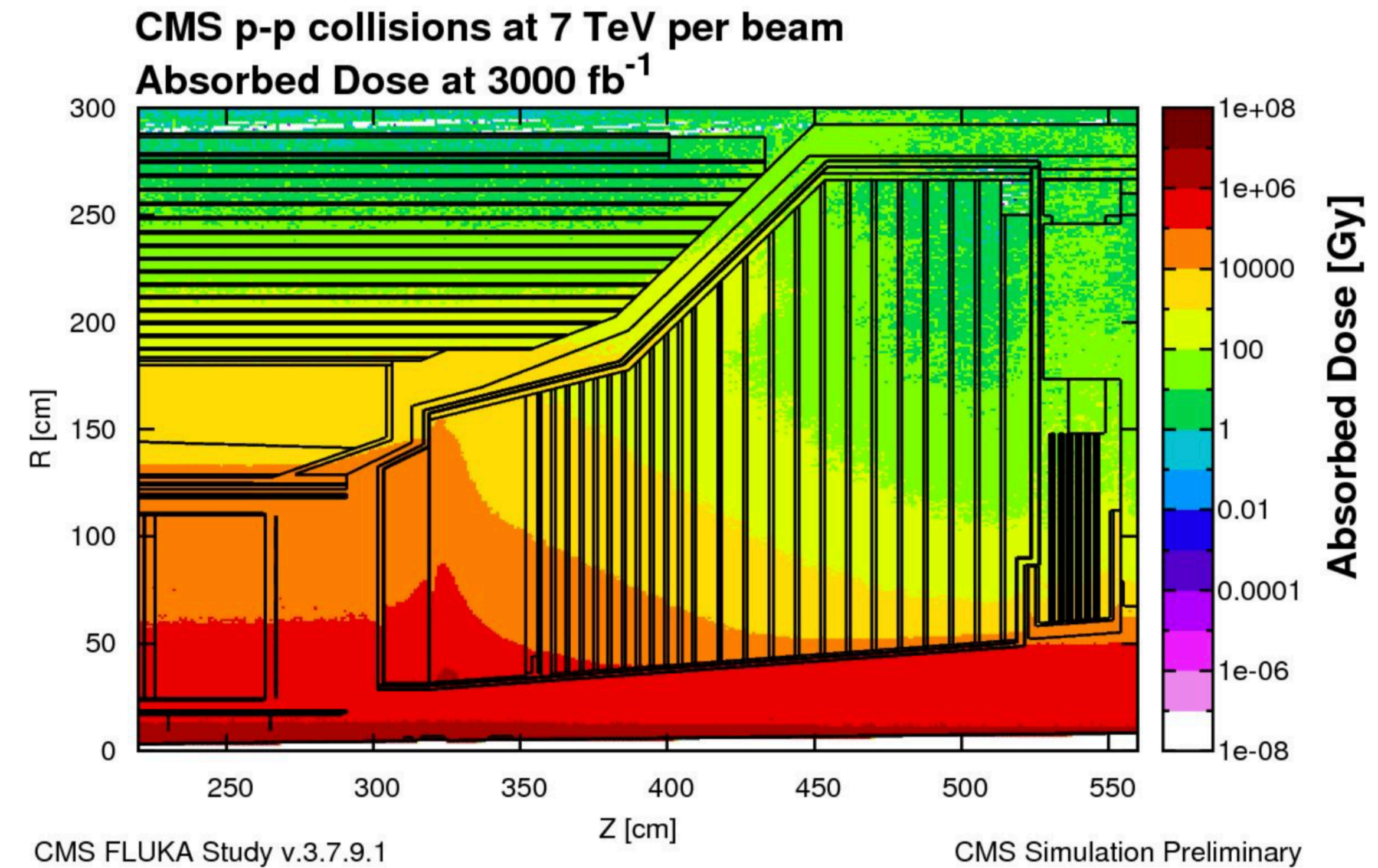
⁴INFN-Torino, Italy

Outline

- Context and motivation
- Fibers layout, composition and characteristics
- Irradiation tests: attenuation and recovery measurements
- Different applications: scintillating fibers vs WLS fibers
- Summary and future development

Motivation

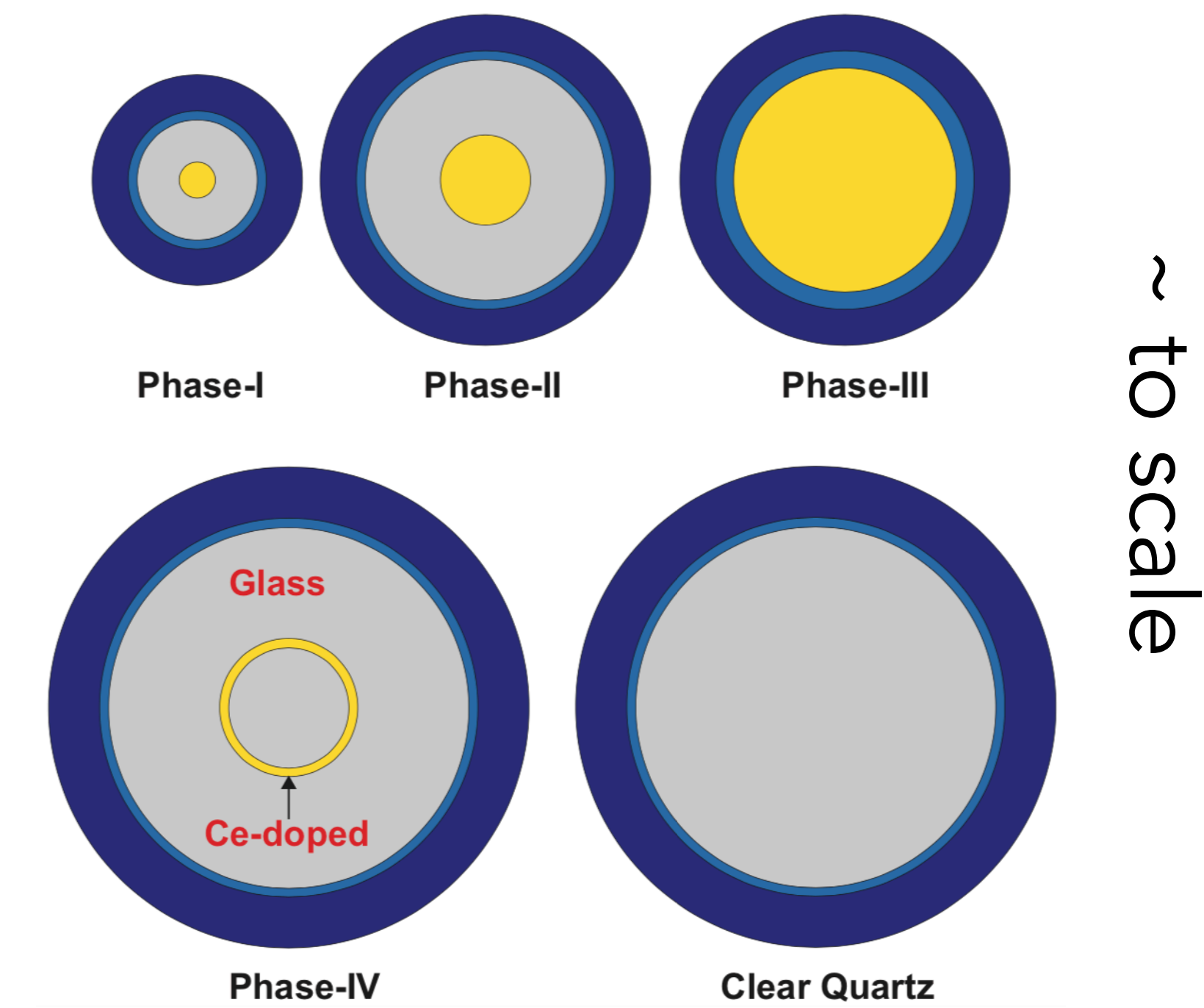
- Need for **radiation-hard** scintillators for applications at colliders
 - **Fused-silica** is a good base material (see CMS HF)
 - **Cerium** as a dopant gives several advantages
 - relatively large LY
 - good overlap between excitation of $\text{SiO}_2\text{:Ce}^{+3}$ and the emission band of the CeF_3
 - peak emission wavelength appropriate for bialkali PMTs



- Doped fibers can be applied either as **scintillators radiators** or as **wavelength shifters** for the collection and transport of light
- The combination of clear and doped fibers opens the door to a **multiple-readout approach**
 - **"Hybrid" fibers: Scintillation and Cherenkov signals in the same fiber**
- Fibers enable the design of detectors with high spatial resolution, good timing properties and their production is cost effective

Fibers layout

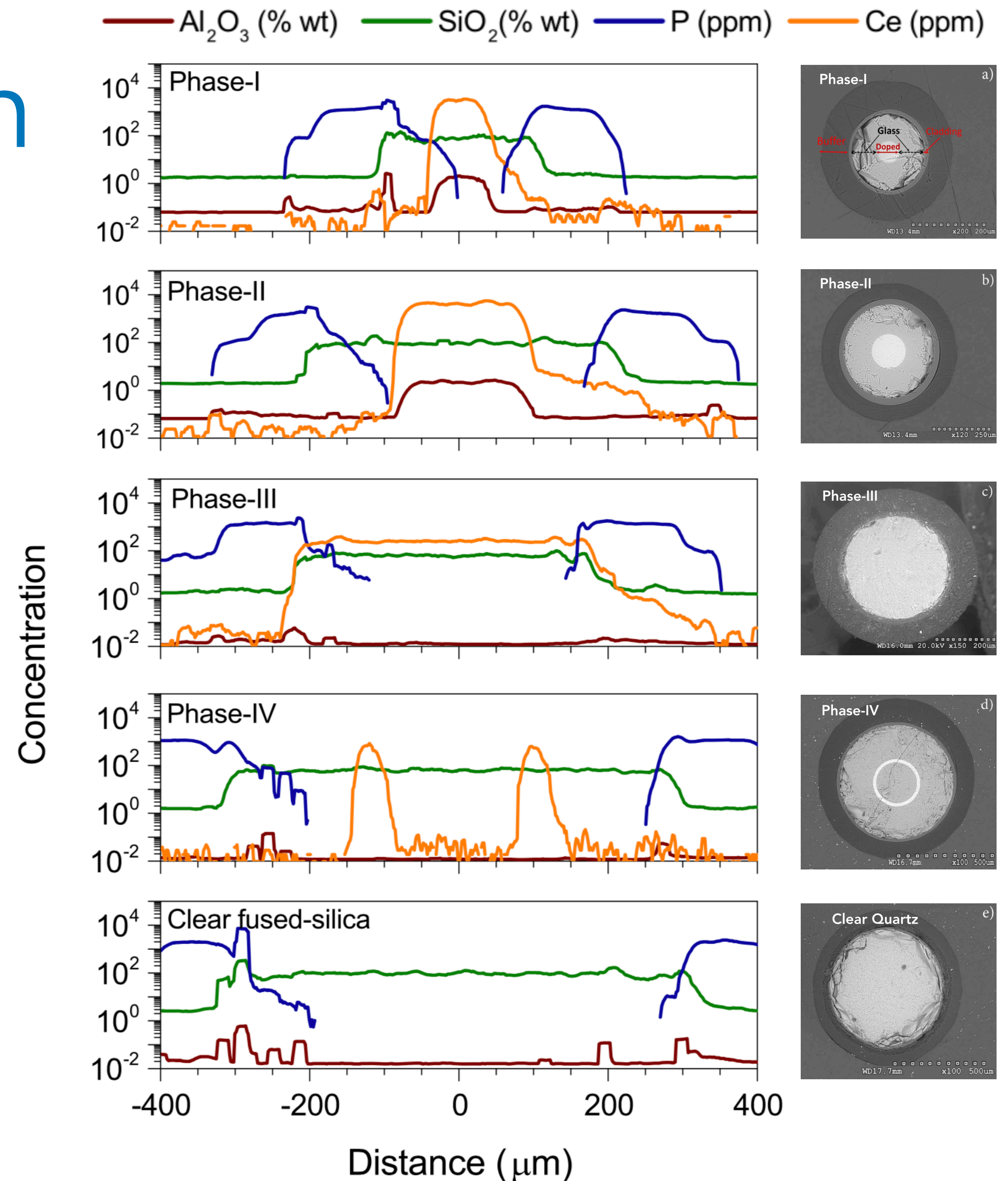
- Composition:
 - Hard-polymer clad + acrylate buffer
 - High OH- fused synthetic silica
- Different layouts explored to improve radiation hardness and light transmission



Fiber Name	Core OD [μm]	Glass OD [μm]	Clad OD [μm]	Buffer OD [μm]
Phase-I	60±7	200±6	230 ⁺⁵ ₋₁₀	350±15
Phase-II	150±20	400±10	430±10	550±30
Phase-III	370±8	—	400 ±8	550±15
Phase-IV	200±8(clear) 230±8(doped)	600±6	630±10	800±30
Clear fused-silica	600±10	—	630 ⁺⁵ ₋₁₀	800±30
T14	150±5	400±5	450±5	500±15

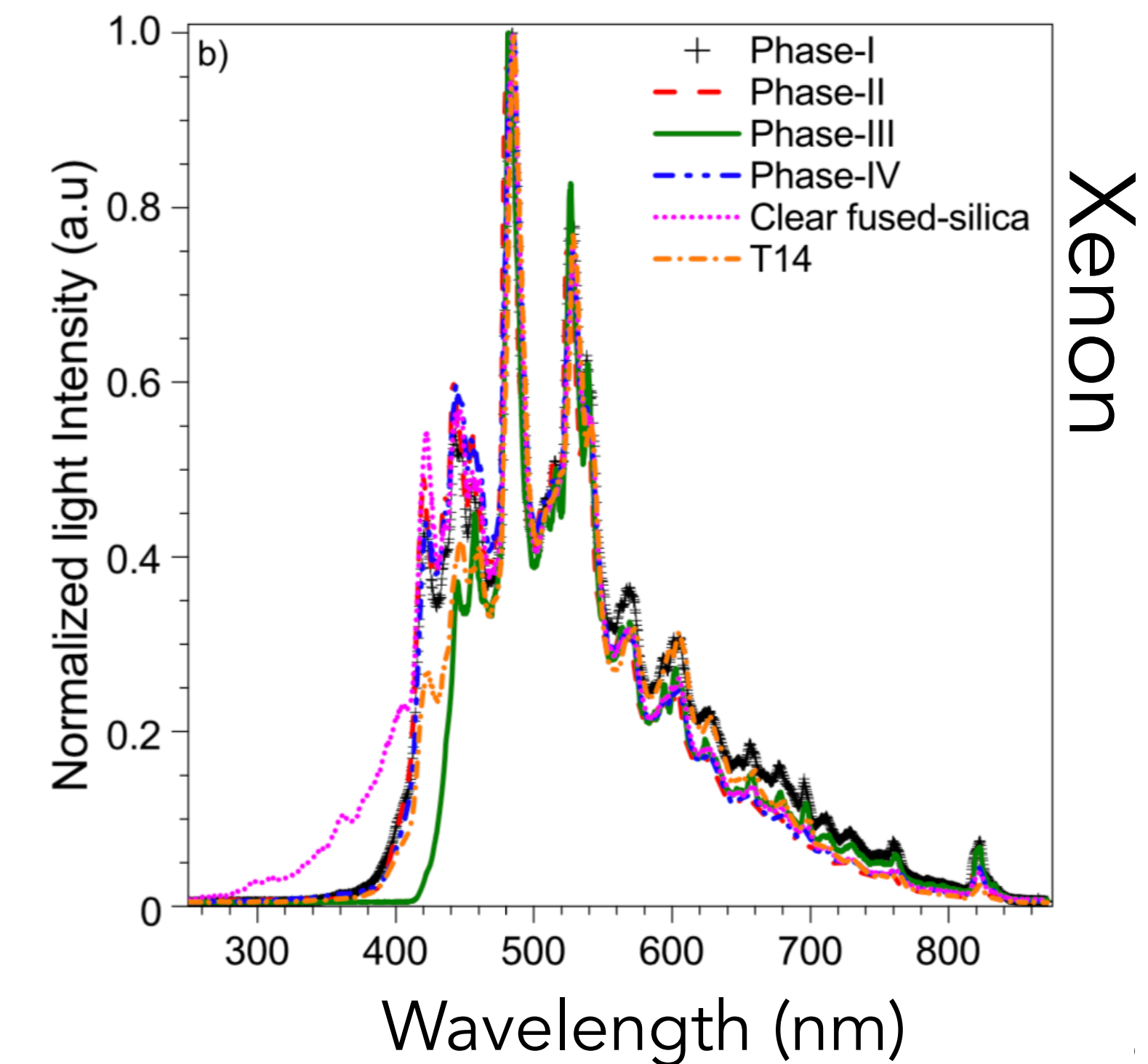
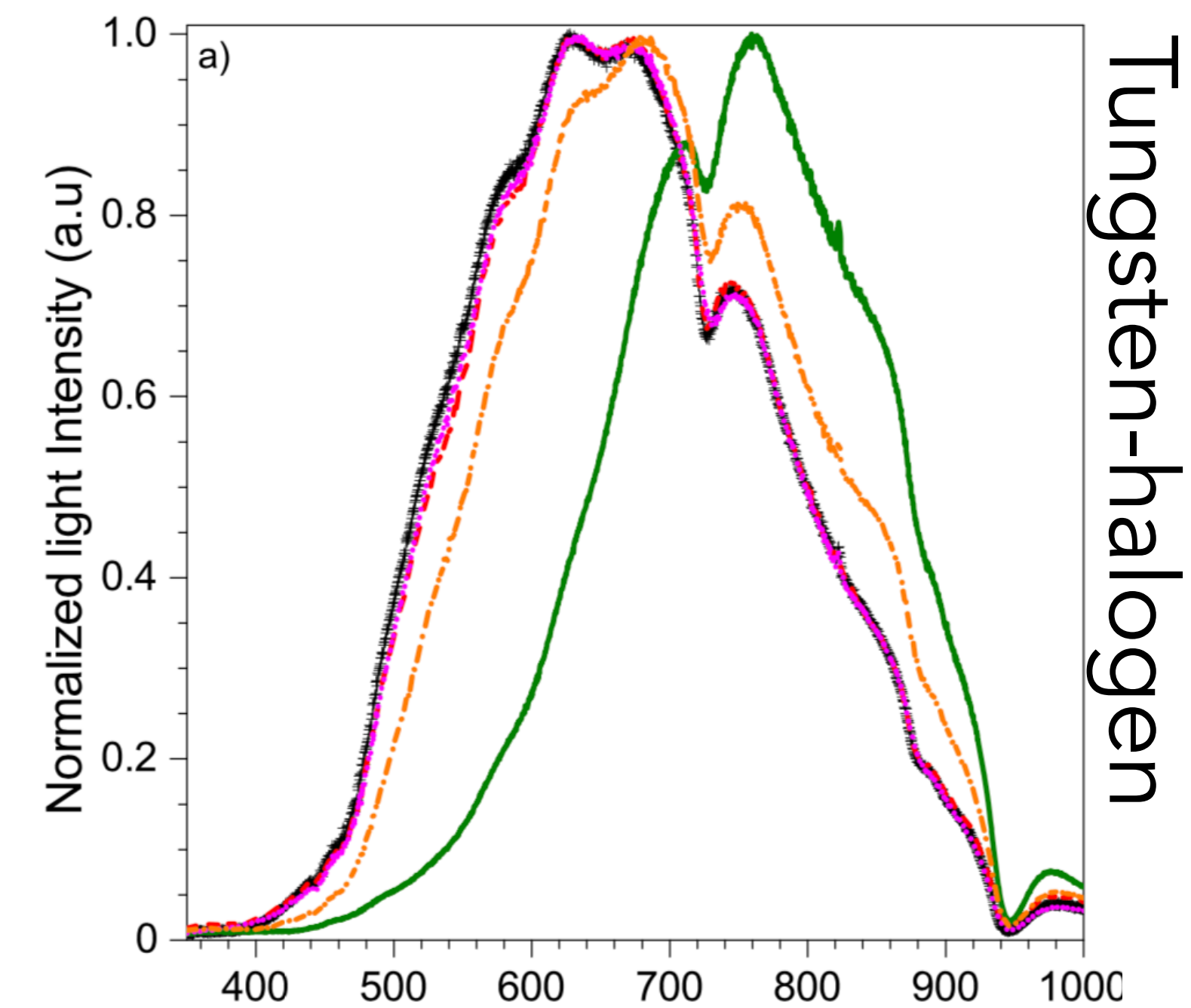
Fibers composition

- Mass spectrometer (Agilent 7500cs) used to determine the chemical composition of all fibers
 - Scan rate of 5 $\mu\text{m/s}$
 - 40- μm diameter laser spot size
- Small concentrations of Ga, La and Gd tend to accompany Ce in glasses
- Significant concentrations of Al in Phase-I, -II
 - Reduced in latest prototypes



Spectral characteristics

- Ce:SiO₂ absorption peak: 329 nm
- Compare transmission of light through 2.5 m of un-irradiated fibers using tungsten-halogen and xenon lamps
 - Reference: **Clear Fiber**
- Observations:
 - Effective light transmission when clear fused-silica component is present (**Phase-I**, **Phase-II**, **Phase-IV**)
 - Absorption of short wavelength particularly evident for **Phase-III**



Radiation hardness studies

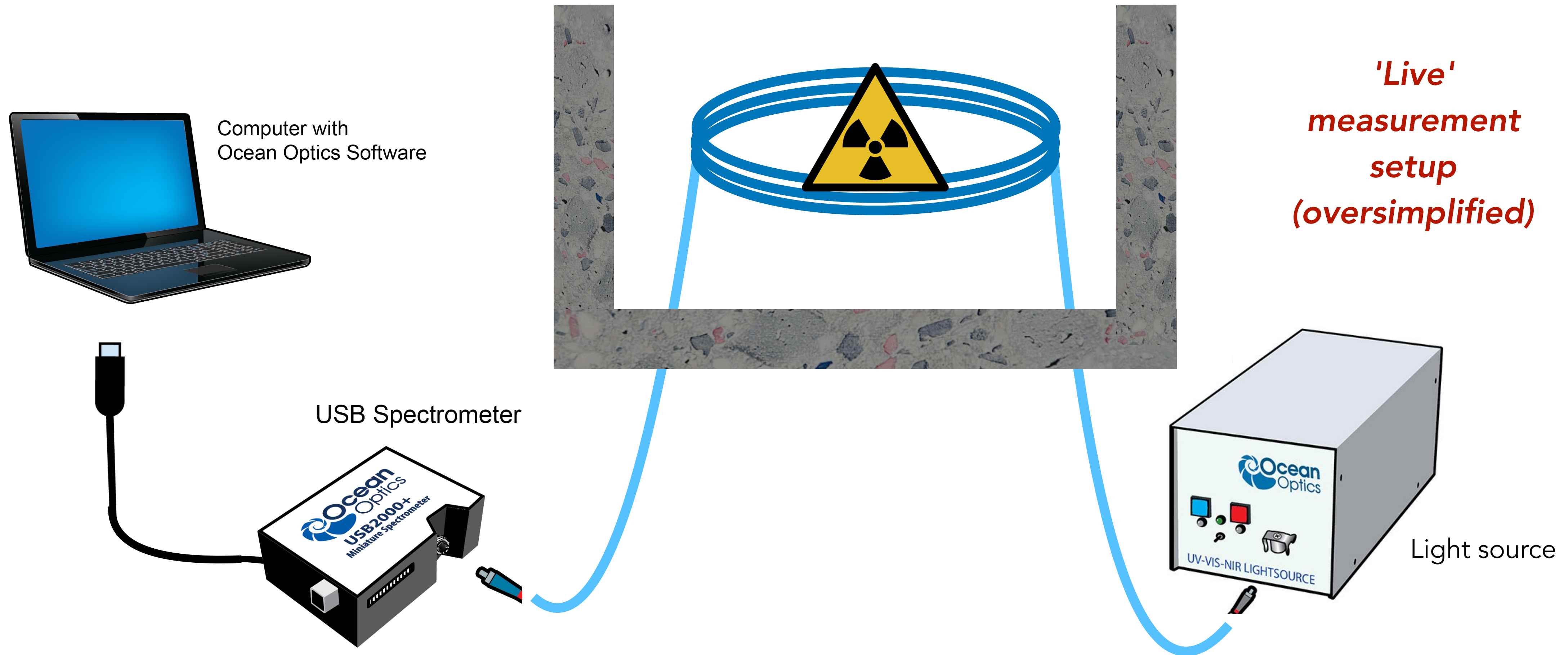
GOAL: study the response after irradiation

- evaluate degradation in photo-luminescence and optical transmission
- model recovery phenomena

Setup

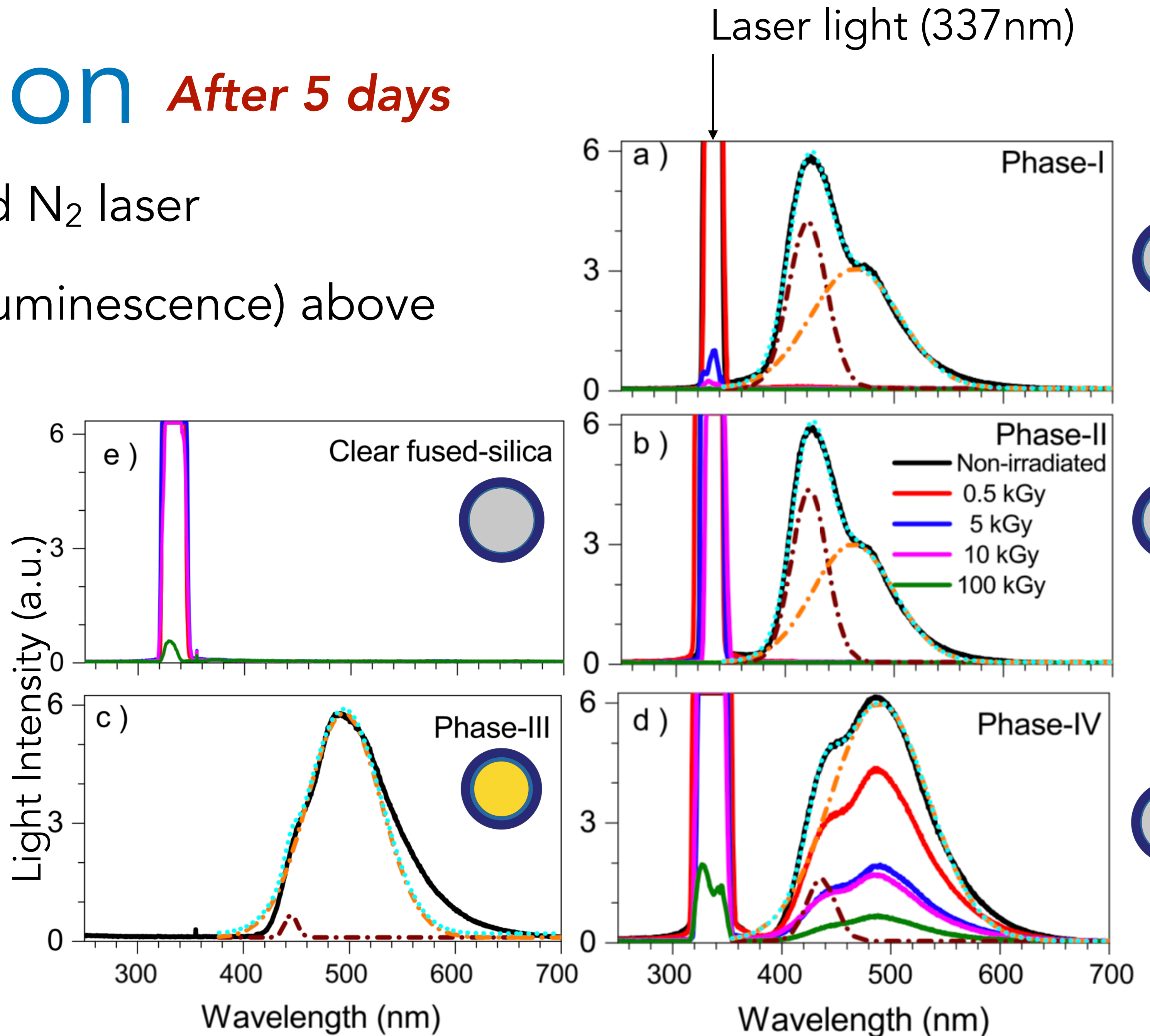
Two irradiation sessions are performed:

- 0.5, 5, 10, 100 kGy at rates of 0.05, 0.5, 1, 10 kGy/h
measure **after 5 days**
- Lower doses, 'live' measurements

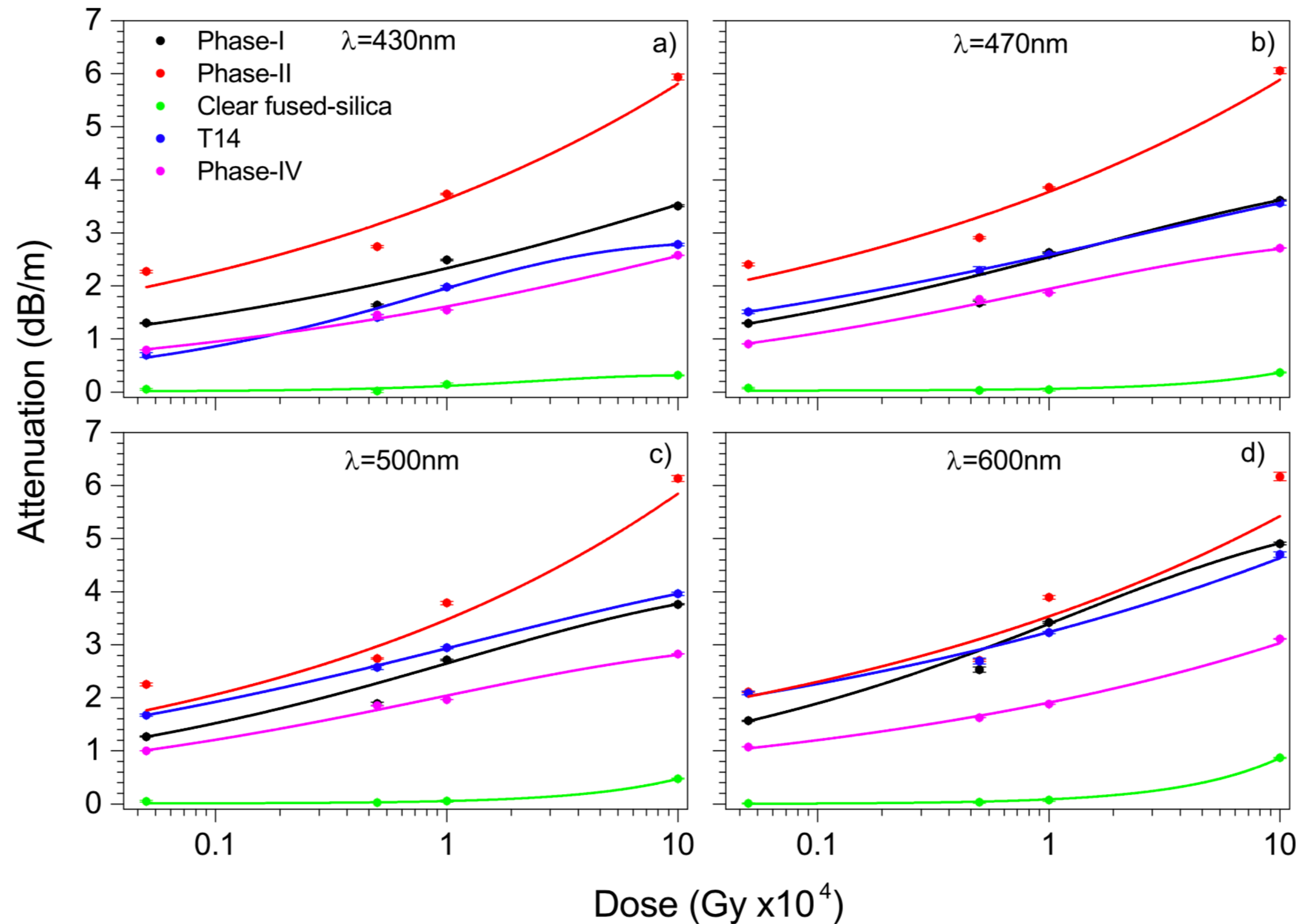


Light emission *After 5 days*

- Fibers excited by a pulsed N₂ laser
- Photon emission (photo-luminescence) above 440 nm with a long tail
- Good UV light transmission when clear glass is present
- **Emission still visible for Phase-IV up to 100 kGy**



Light transmission *After 5 days*



- Inject Xenon light axially and measure spectra
- Four wavelength
- Among the doped fibers, **Phase-IV performs the best**

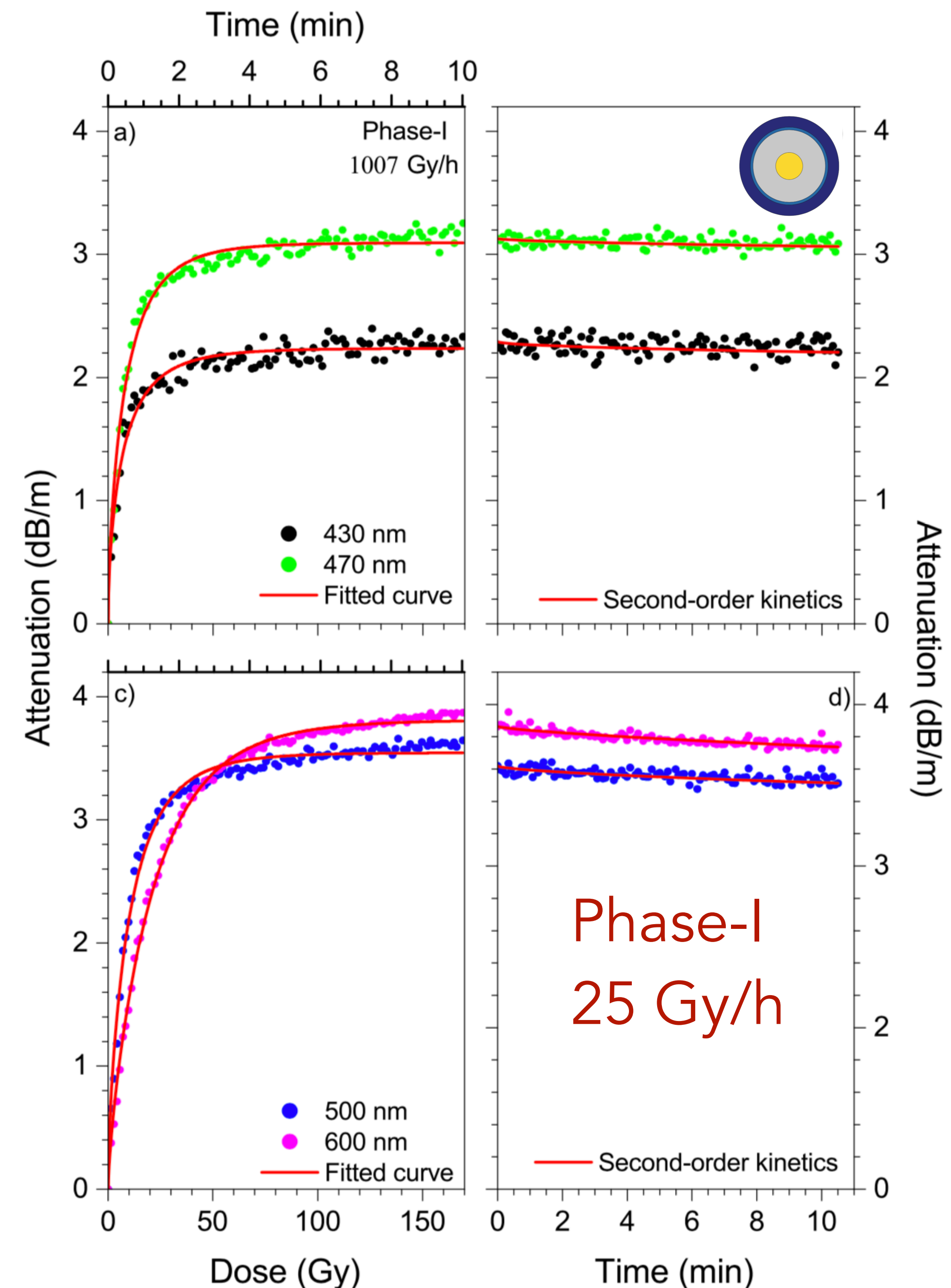
$$A(\lambda, D) = -\frac{10}{L} \log \left(\frac{I(\lambda, D)}{I_0(\lambda)} \right)$$

Response evolution Phase-I *'Live' meas.*

- Attempt to **model rad-induced damage** and optical **recovery**
 - Follow Griscom's classical rate equation with solution

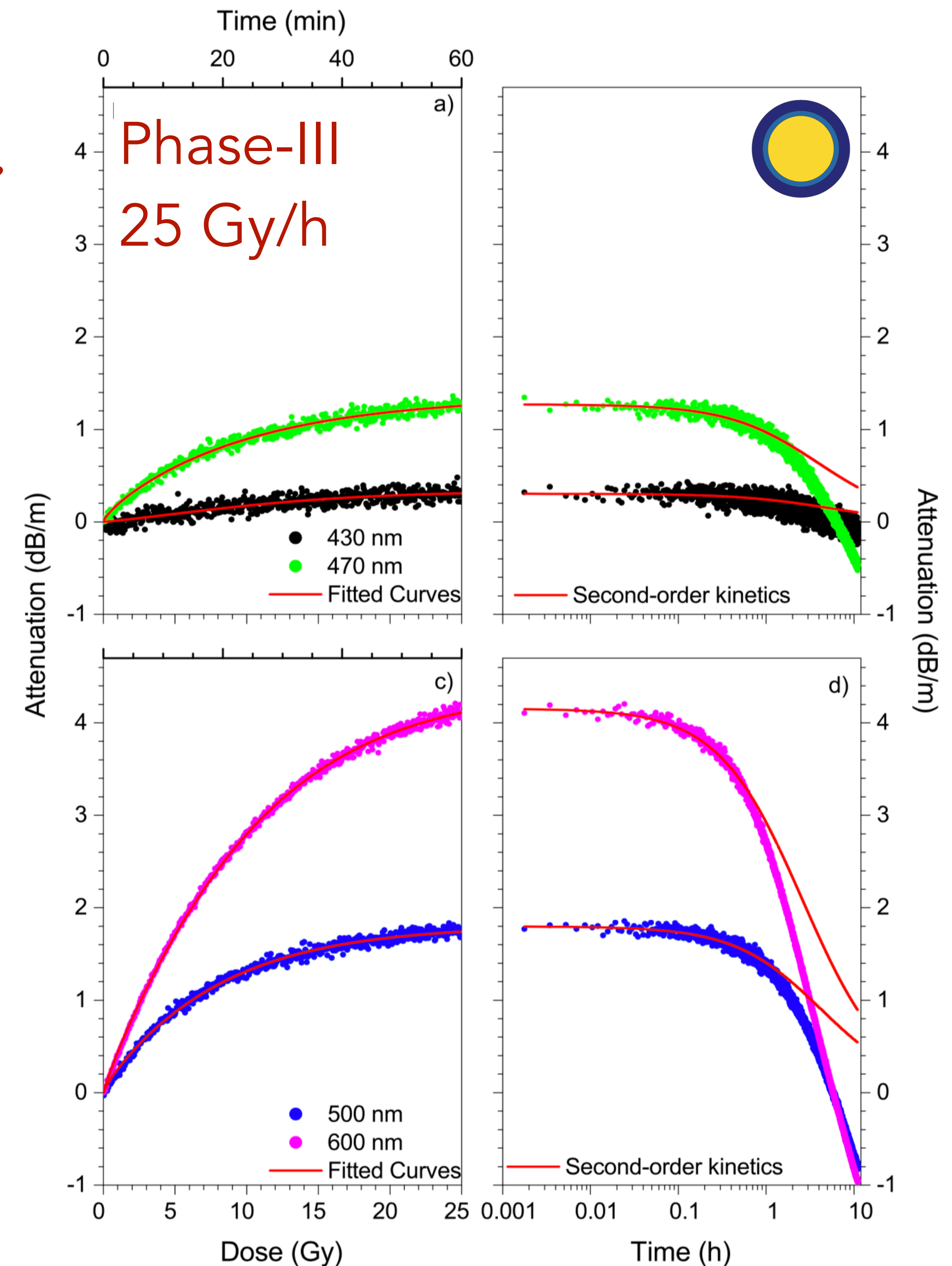
$$N[(kt)^\beta] = N_{\text{sat}} \tanh[(kt)^\beta]$$

- Fit N_{sat} , k and β during irradiation and extrapolate during recovery
- Main features of damage and recovery are **well described by the fitted model**
 - **Al and P contamination** prevents fast recovery making color centers stable

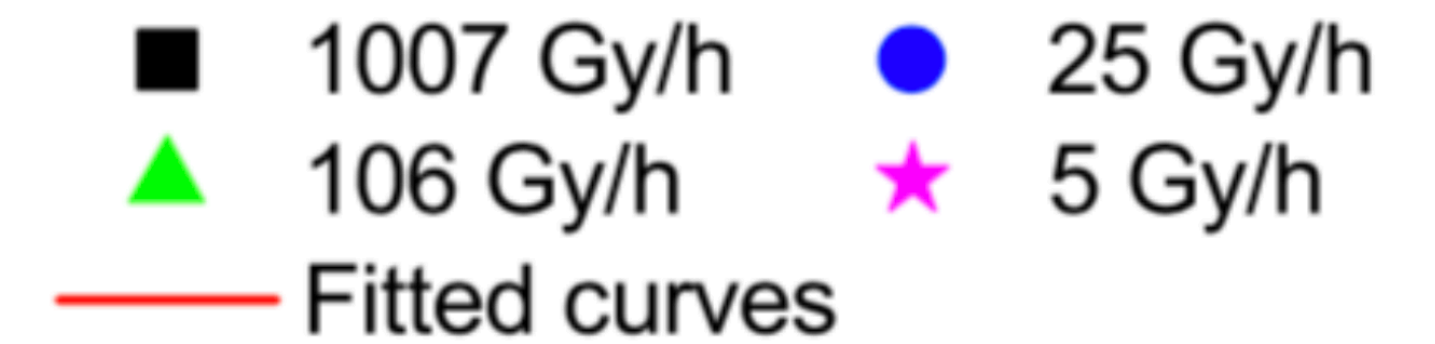


Response evolution **Phase-III** *'Live' meas.*

- For Phase-III fibers, the recovery prediction fails to represent the data after an hour of recovery
- **Over-recovery** is observed
 - absence of AI and P is likely the responsible of the different behavior wrt Phase-I fibers

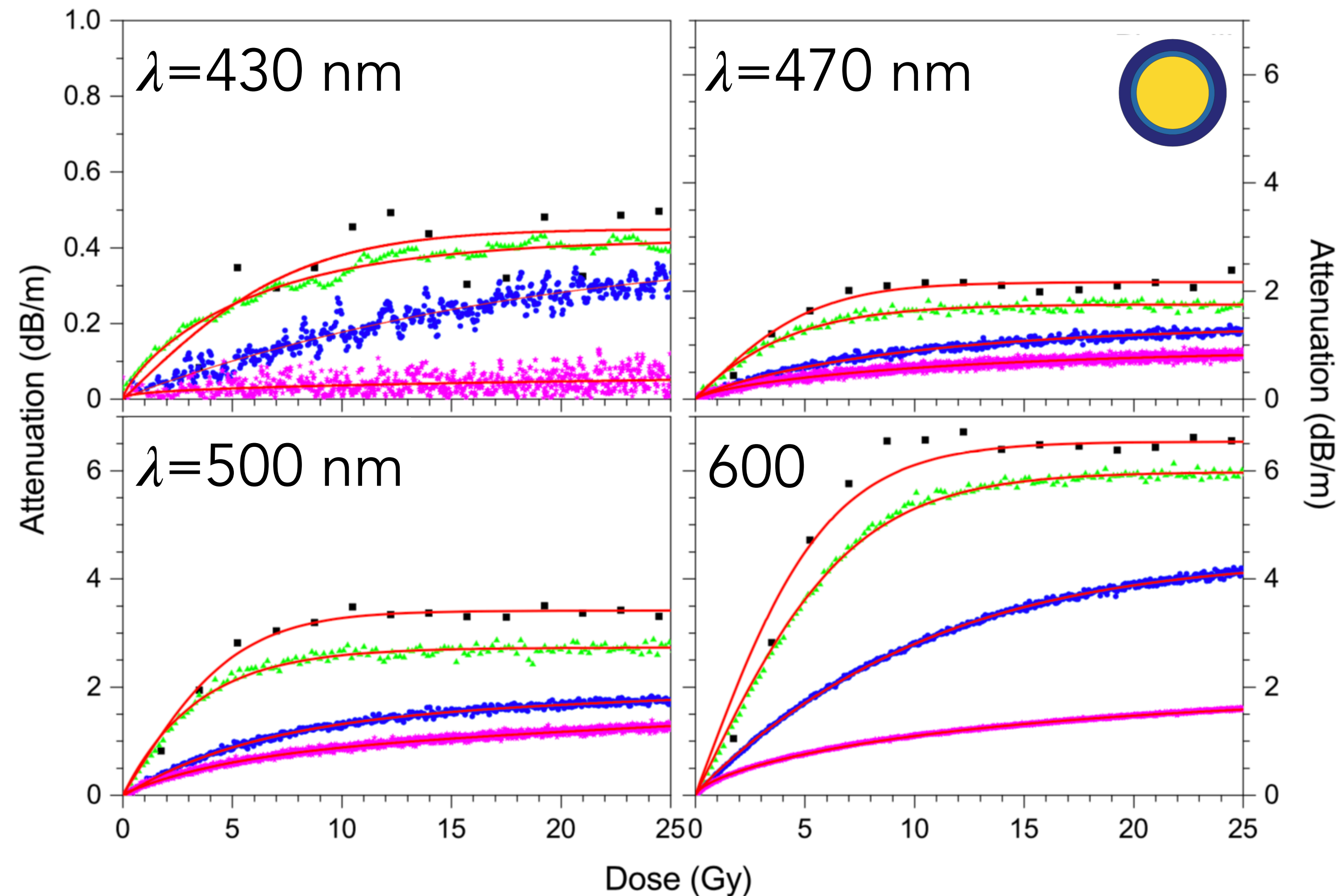


Dose-rate effects



Phase-III

- At all wavelengths, the **attenuation increases as a function of dose** and reaches a plateau
- The degree of attenuation at the plateau depends on the wavelength



Response to electron beam

GOAL: characterize the response of the fibers

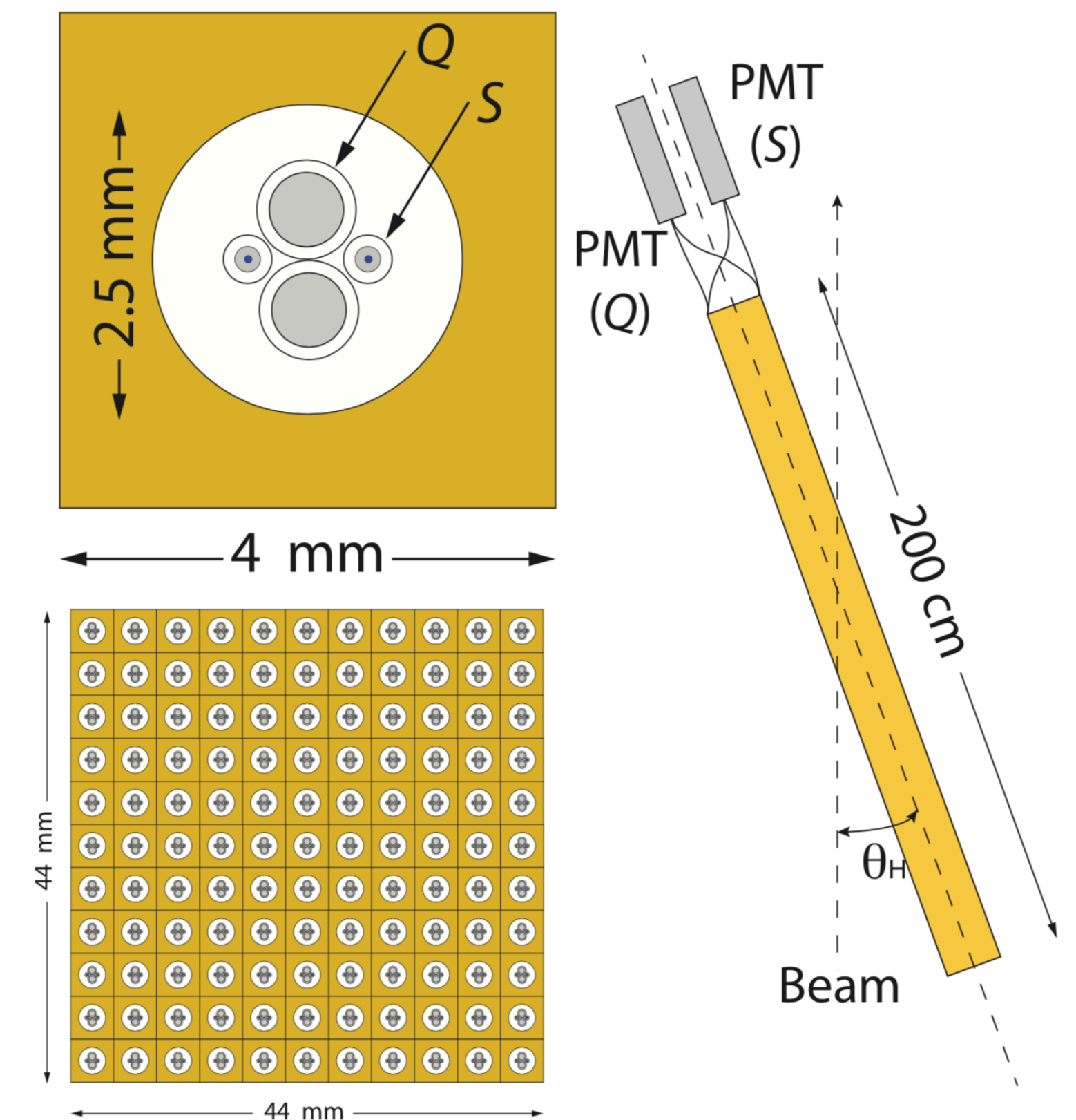
- Pulse shape
- Timing characteristics
- Scintillation light yield

Setup

- A very crude "calorimeter" using Phase-I and clear fibers
 - Phase-I and clear fibers were embedded in a **copper matrix**
 - Transverse size of the matrix: $4.4 \times 4.4 \text{ cm}^2$
 - Length: 200 cm
 - Scint (S) and clear (Q) fibers are **readout separately** by PMT (Hamamatsu R6427)
- This setup was exposed to 4 to 32 GeV positrons at the MTest beamline at Fermilab

By volume:

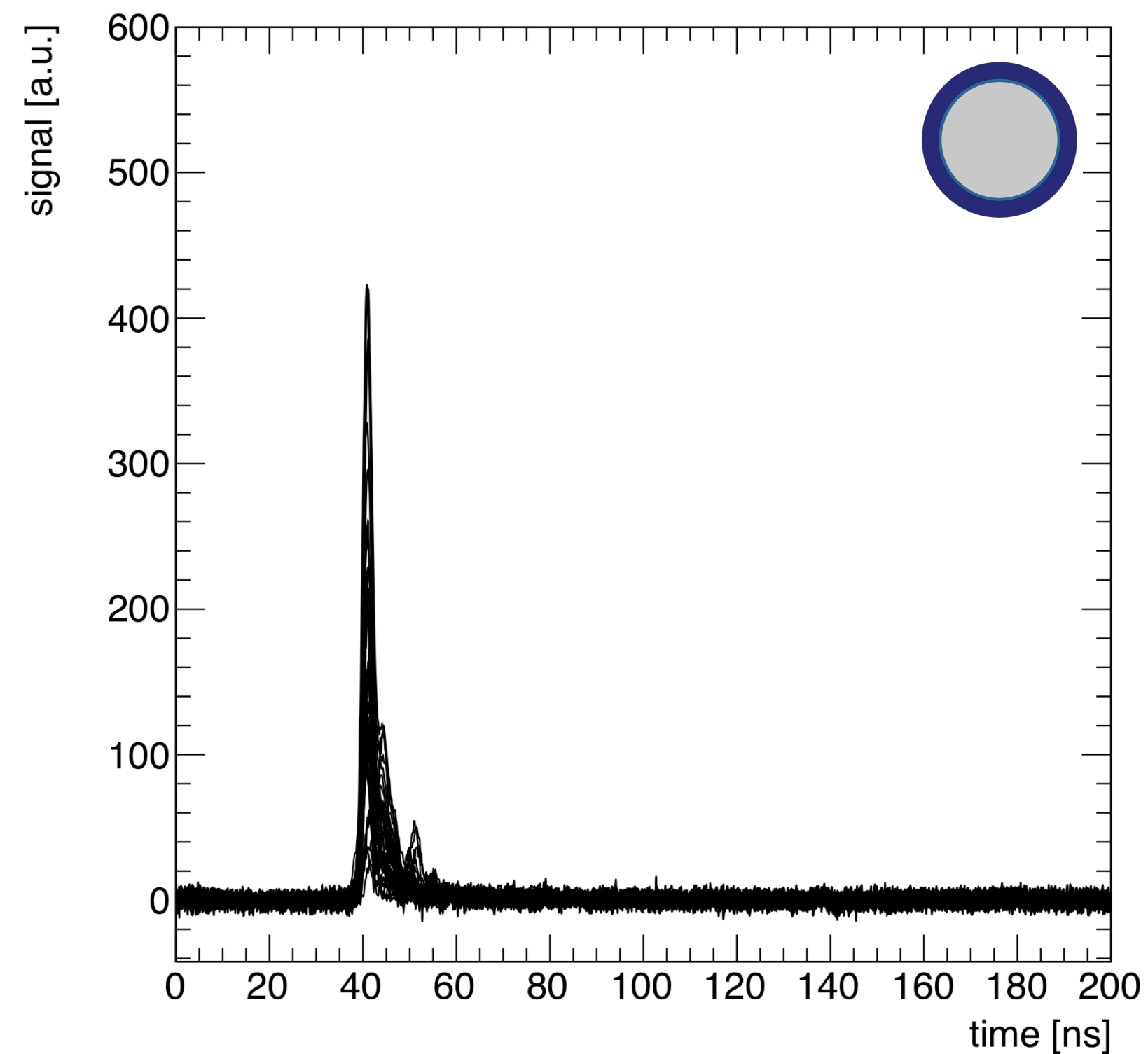
- Cu 71.7%
- Air 20.2%
- Clear quartz 3.5%
- Ce-doped core 0.035%
- Glass in Phase-I fiber 0.35%



Signals from 16 GeV positrons

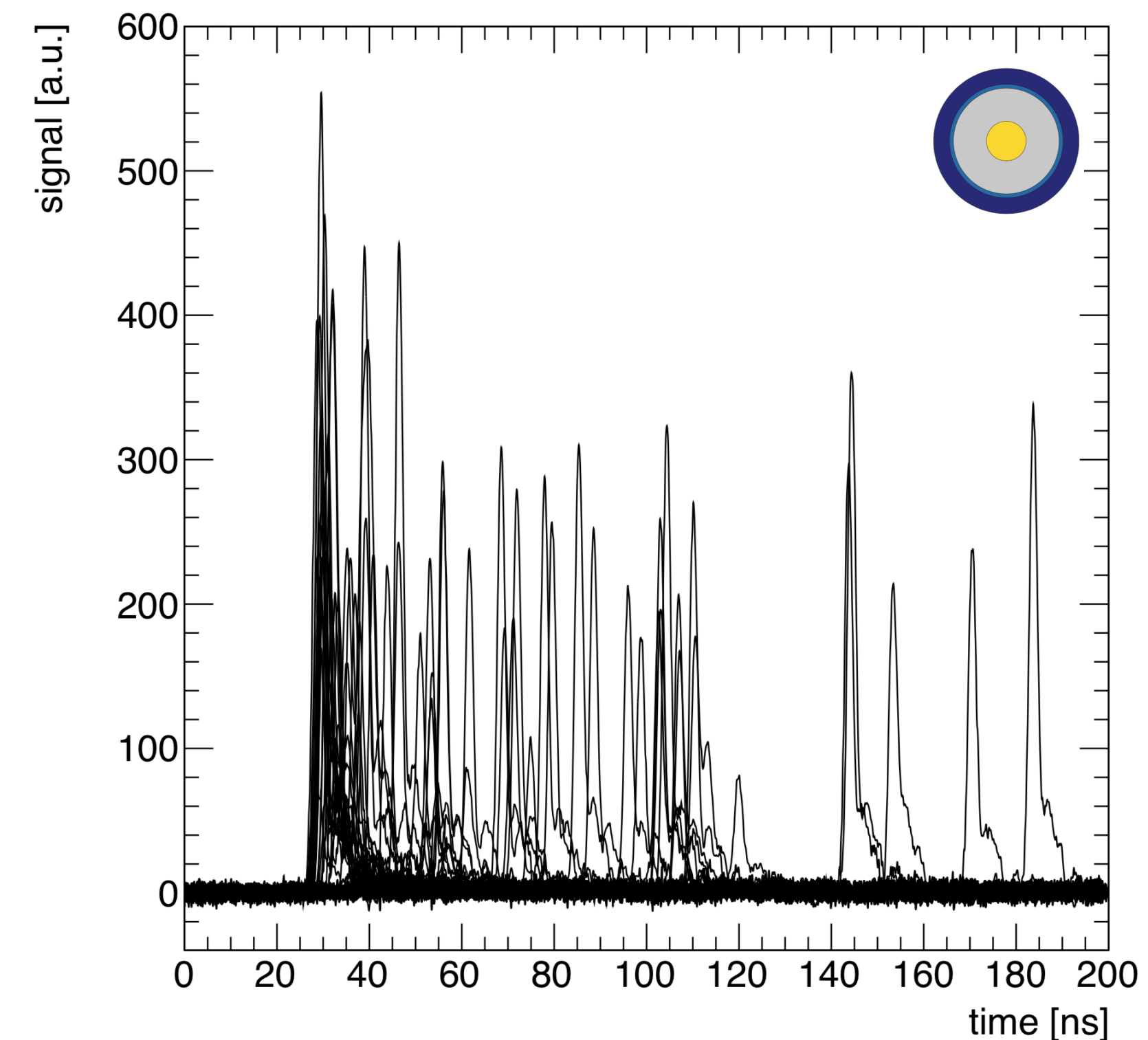
Clear quartz fiber:

- Prompt and fast pulses from Cherenkov



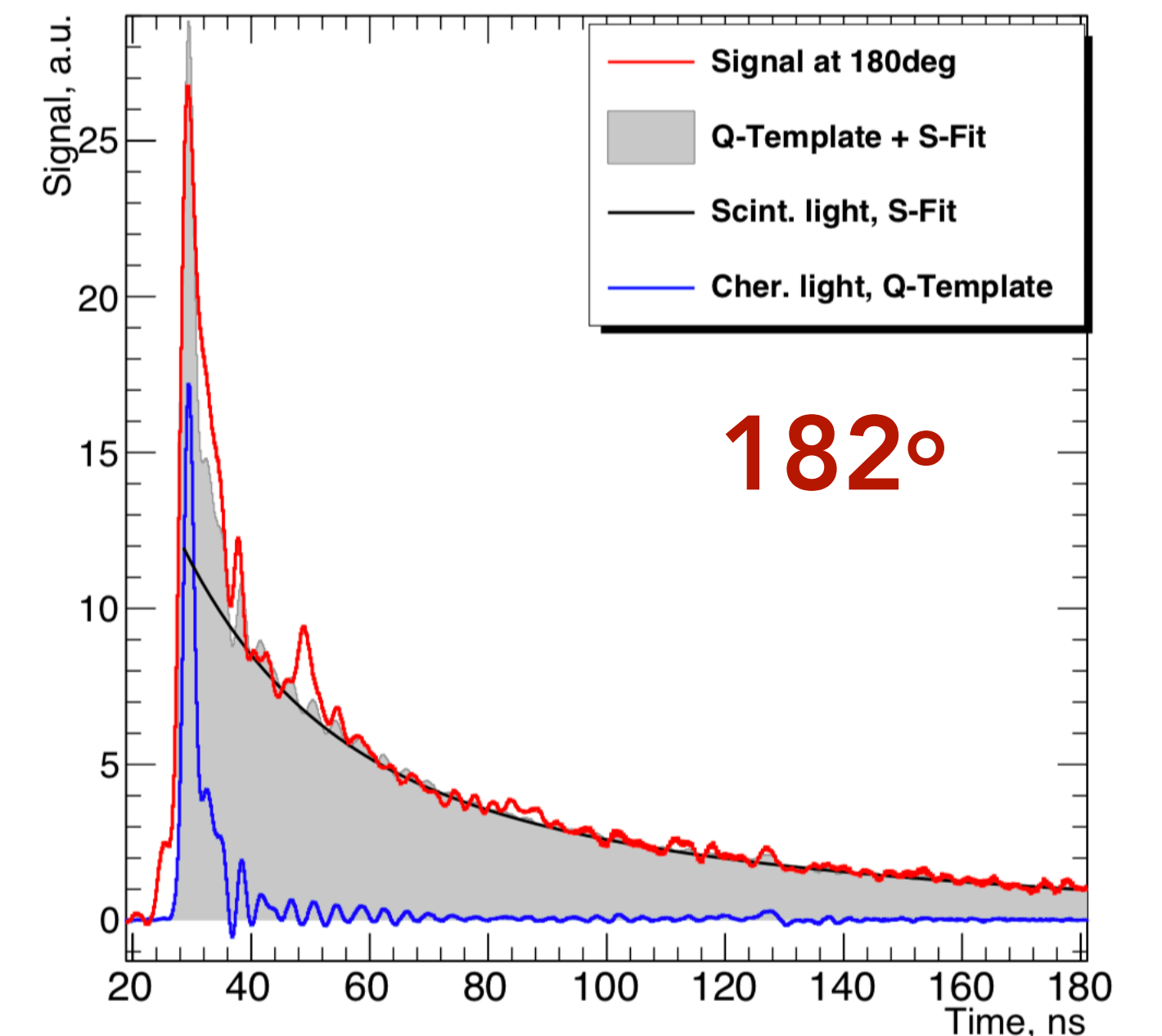
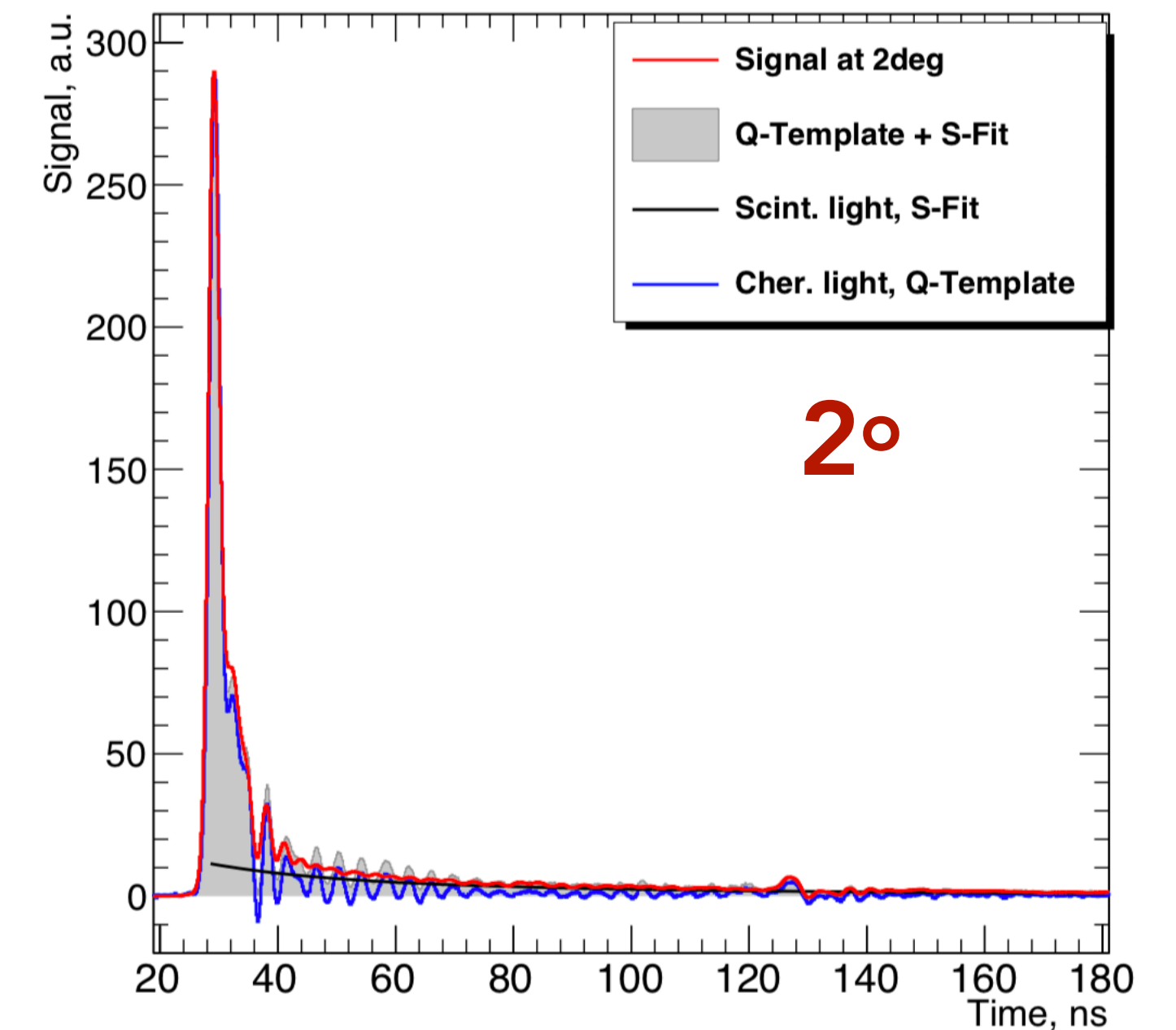
Scintillating Phase-I fiber:

- Both prompt and delayed signals
- Most pulses correspond to single p.e.



Pulse shape

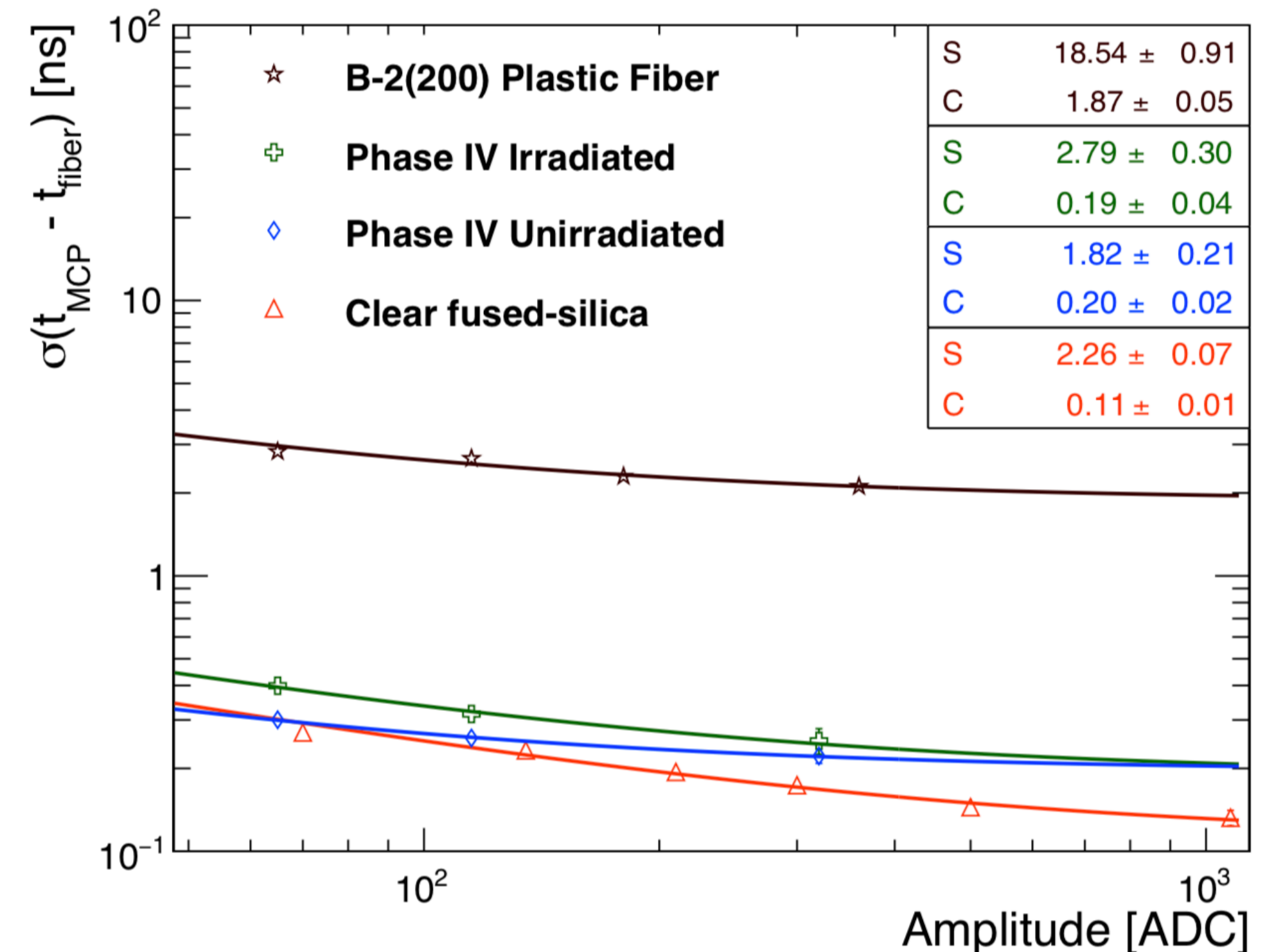
- Can control the abundance of Cherenkov light by changing the orientation of the fibers wrt beam
 - Cherenkov: **directional**
 - Scintillation: **isotropic**
- The scintillation contribution has **two components**:
$$S(t) = a_1 e^{-\frac{t}{\tau_1}} + a_2 e^{-\frac{t}{\tau_2}}$$
 - Fast: $\tau_1 = 20.8 \pm 5.4$ ns (23.9±5.3%)
 - Slow: $\tau_2 = 93.0 \pm 12.6$ ns (76.1±5.3%)
- **Absolute Light yield** was measured to be between 500 and 700 γ /MeV depending on the fiber's type



Timing measurements

- The prompt Cherenkov signal **could be exploited for timing applications**
- Using a fast MCP as time reference it is possible to measure and compare the time resolution for different fibers
 - Better performance when comparing to scintillation signal (expected)
 - Degraded performance for the irradiated sample due to a worsening of the light transmission
- Precise measurement should be performed with a dedicated setup

$$\sigma_t^2 = \left(\frac{s}{\sqrt{A}} \right)^2 + c^2$$



Fibers as wavelength shifters

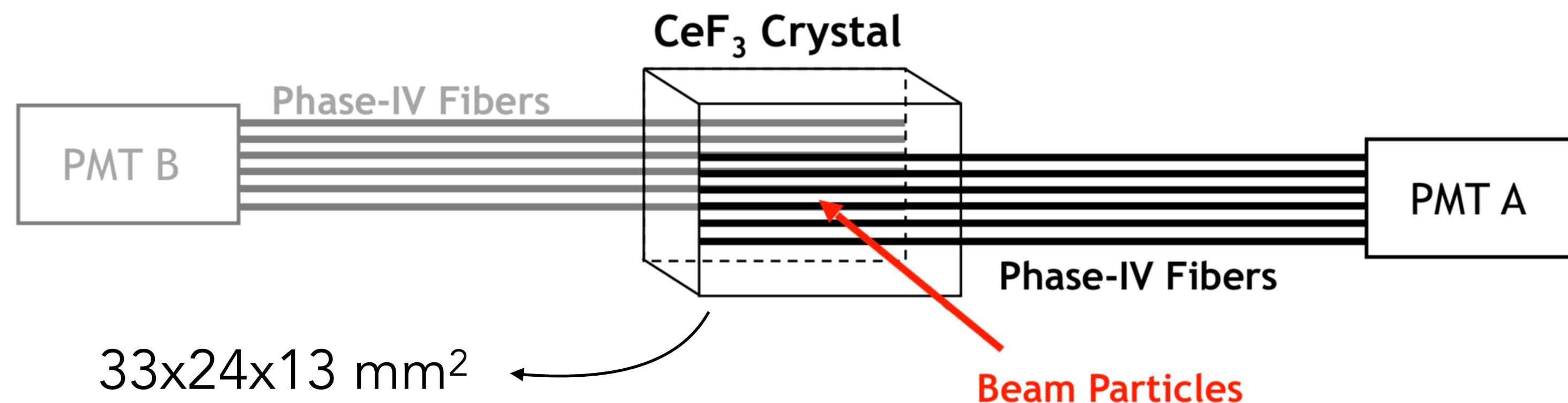
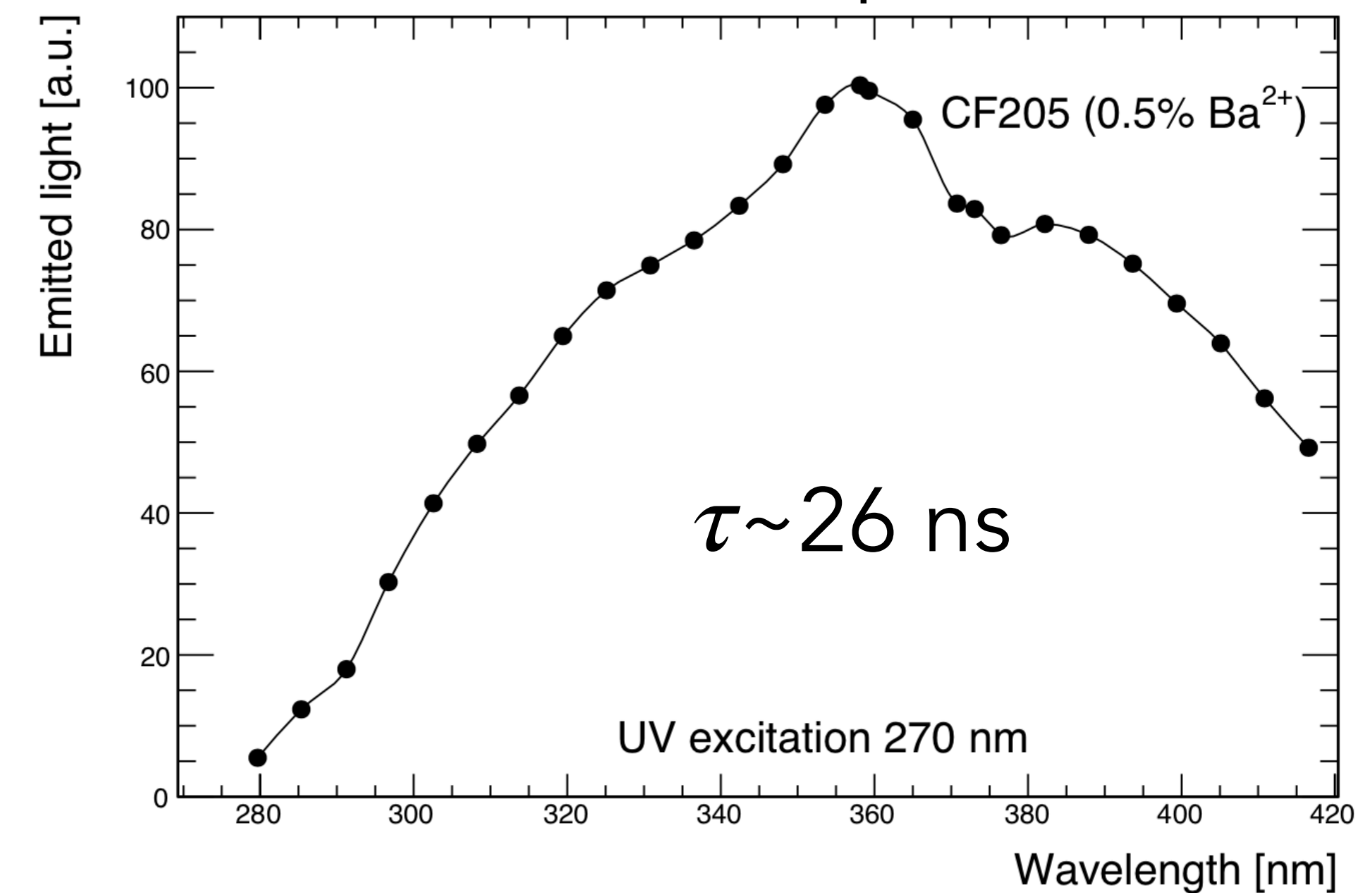
GOAL: evaluate fiber's performance when coupled to a scintillating crystal

- Pulse shape
- Collection efficiency
- Evolution of response after irradiation

Couple fibers with CeF_3 crystal

- Two sets of Phase-IV fibers are coupled to a CeF_3 crystal
 - New vs irradiated (100 kGy) bundle
- Substantial overlap between the excitation of the $\text{SiO}_2:\text{Ce}^{+3}$ and the emission band of the CeF_3

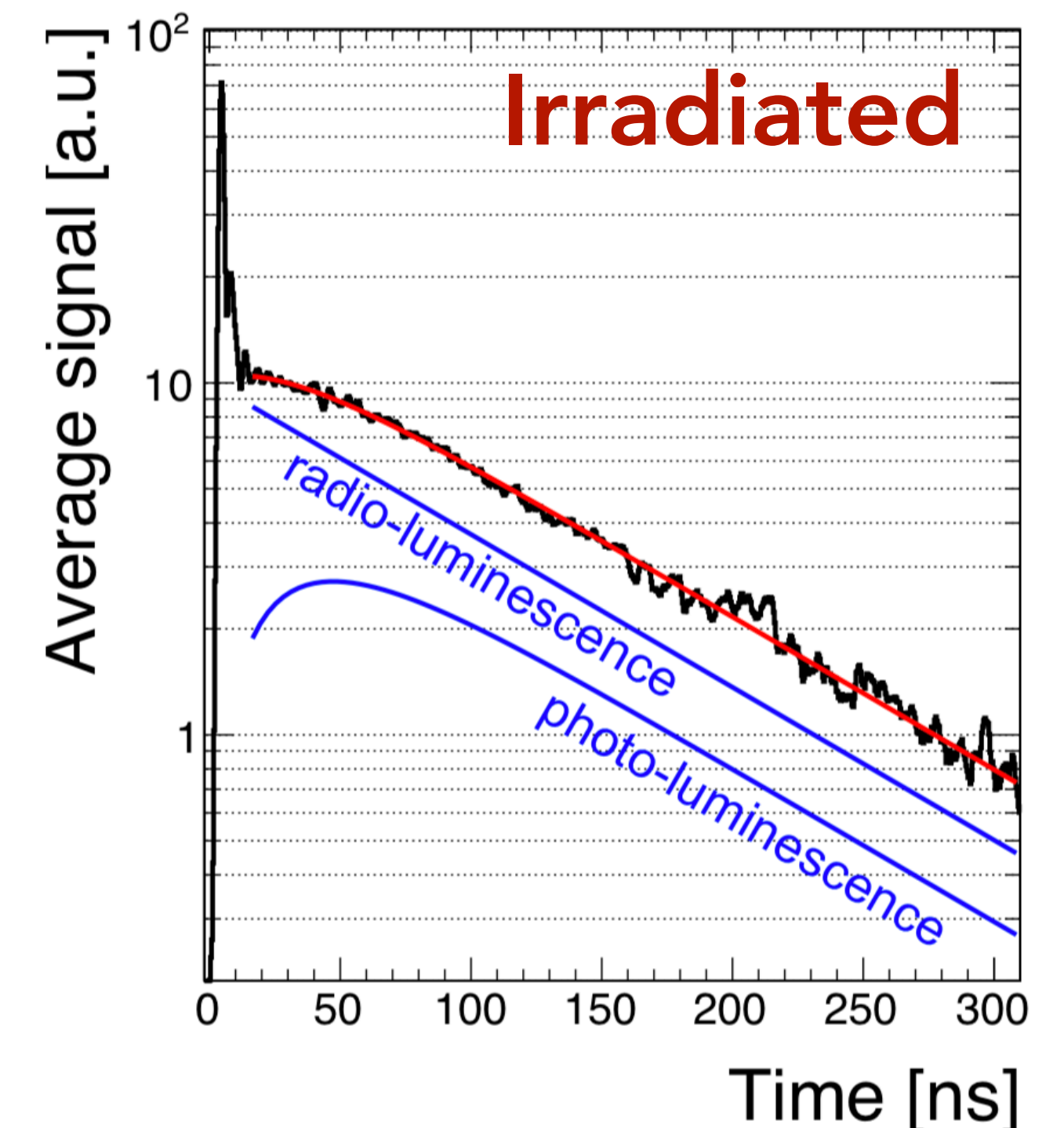
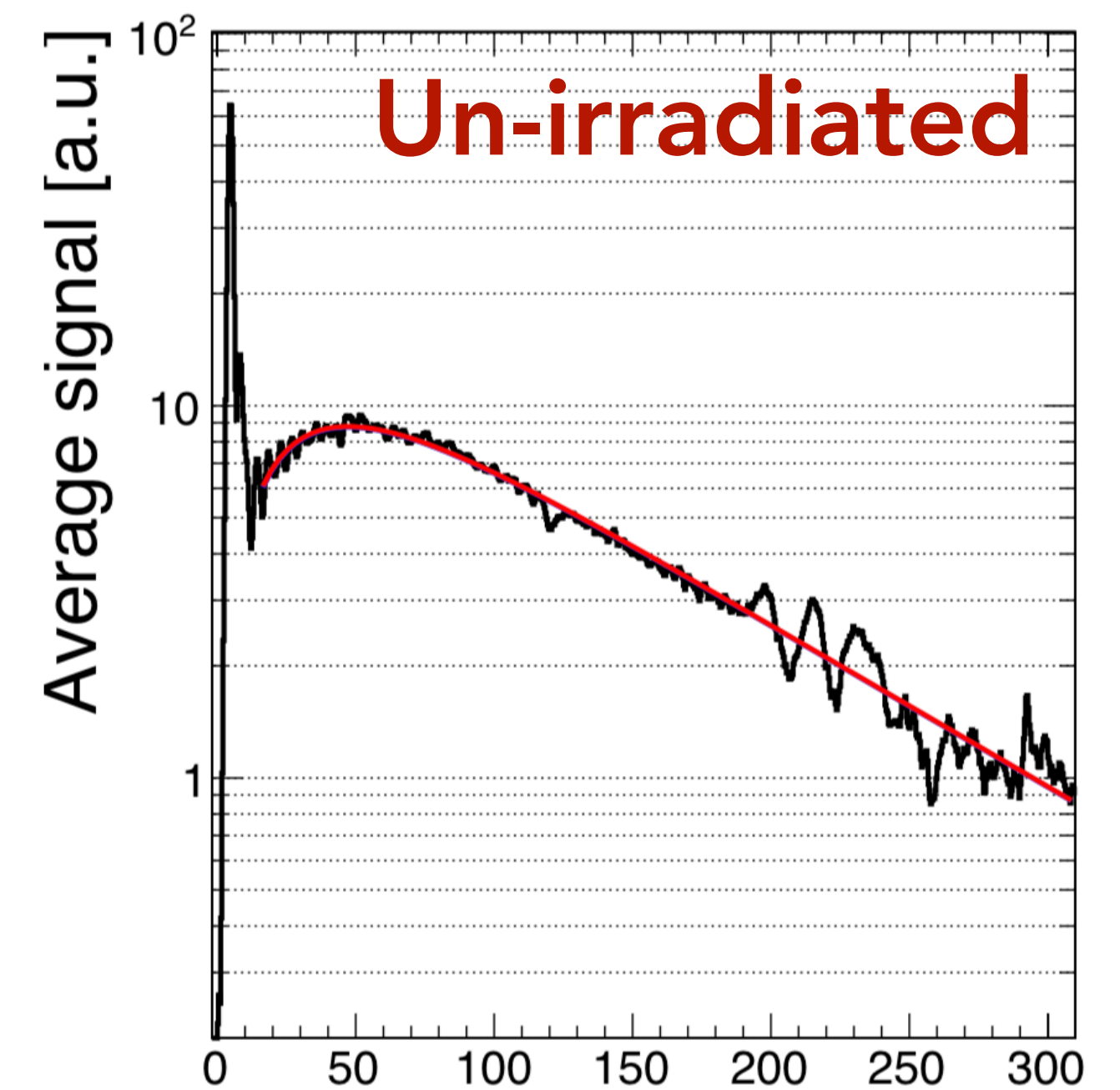
CeF_3 emission spectrum



Abs peak: 329 nm	RM = 2.6 cm
$\rho = 6.16 \text{ g/cm}^3$	$\lambda_l = 25.9 \text{ cm}$
$X_o = 1.68 \text{ cm}$	$n = 1.68$

Compare irradiated vs un-irradiated

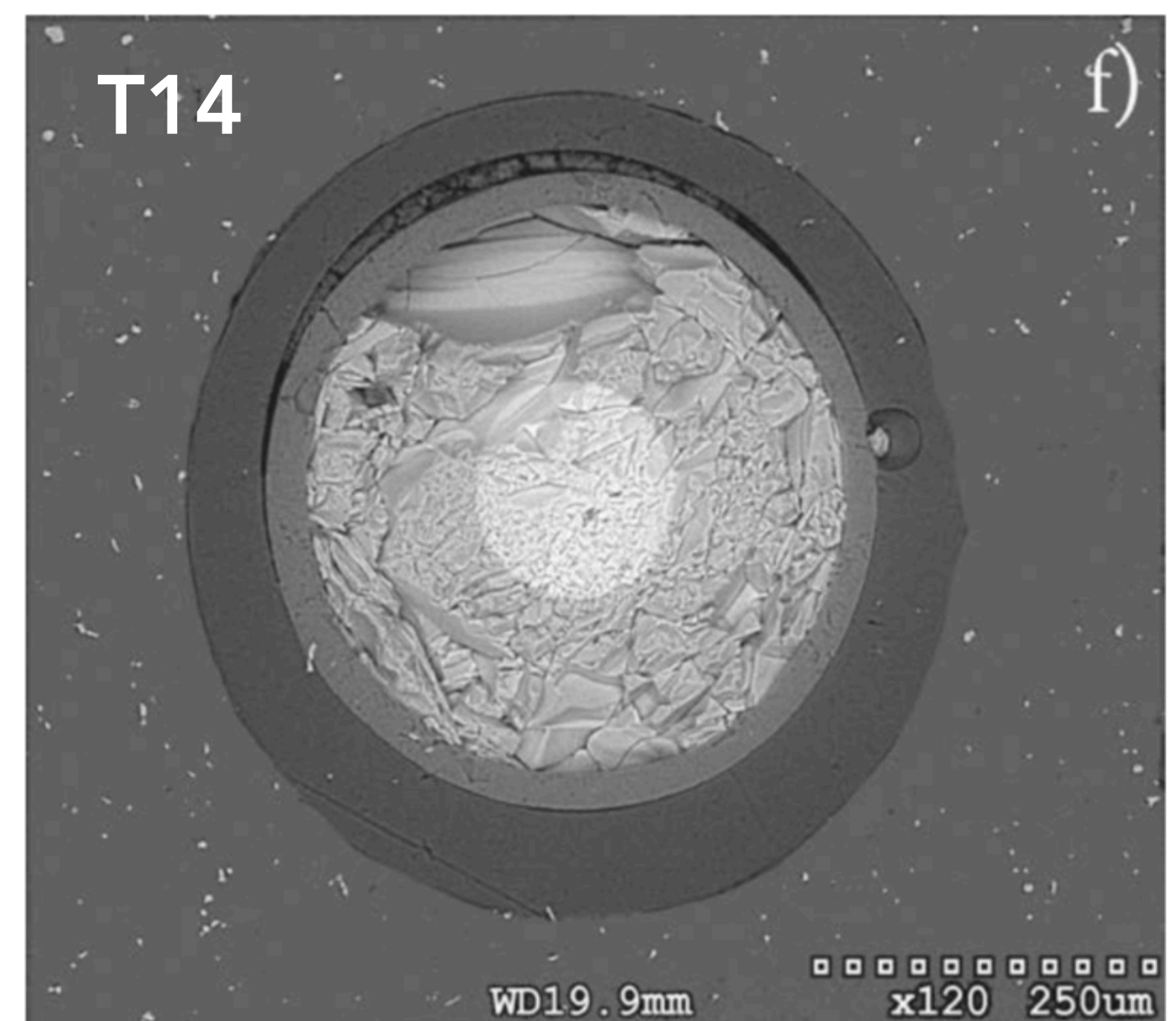
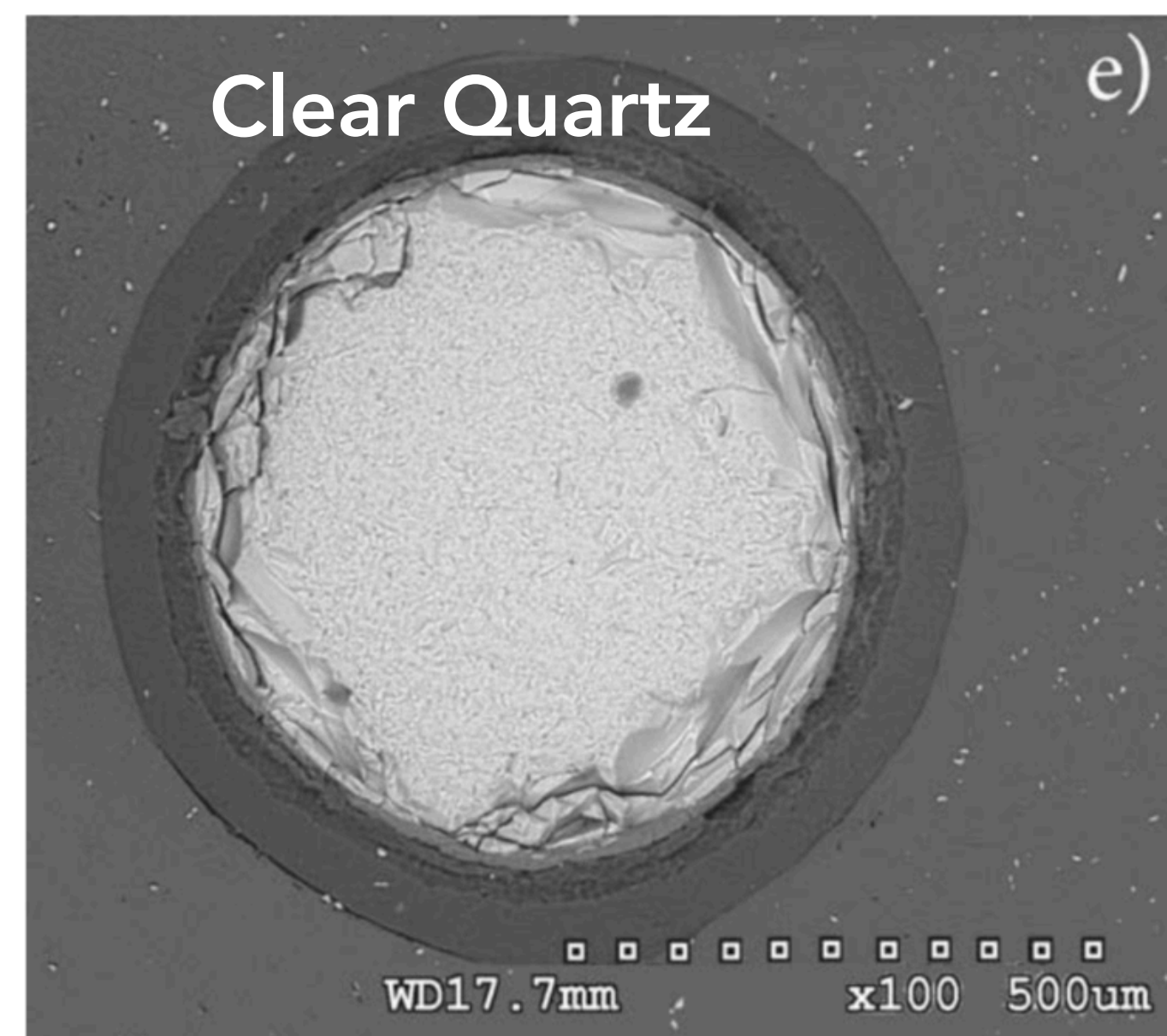
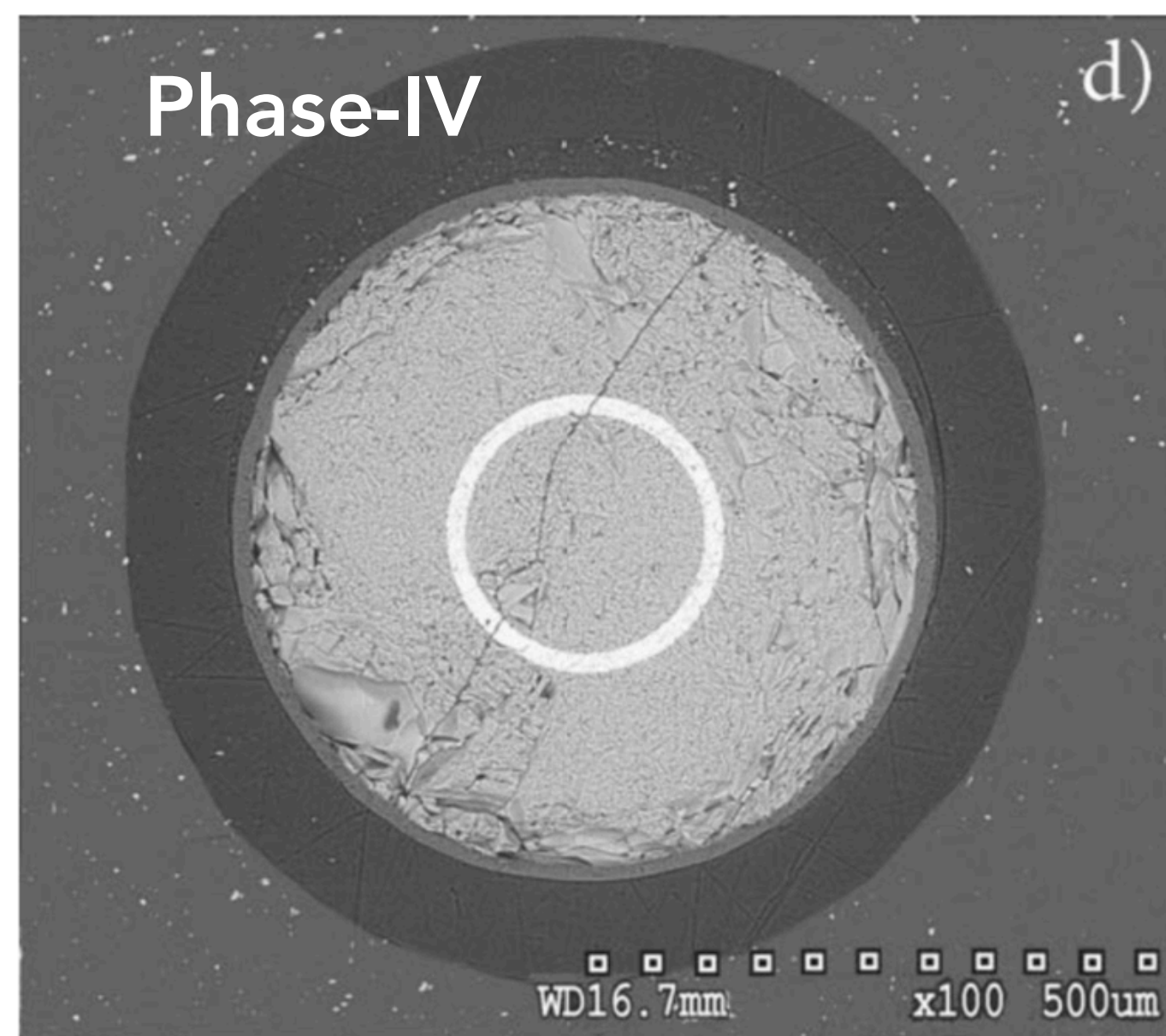
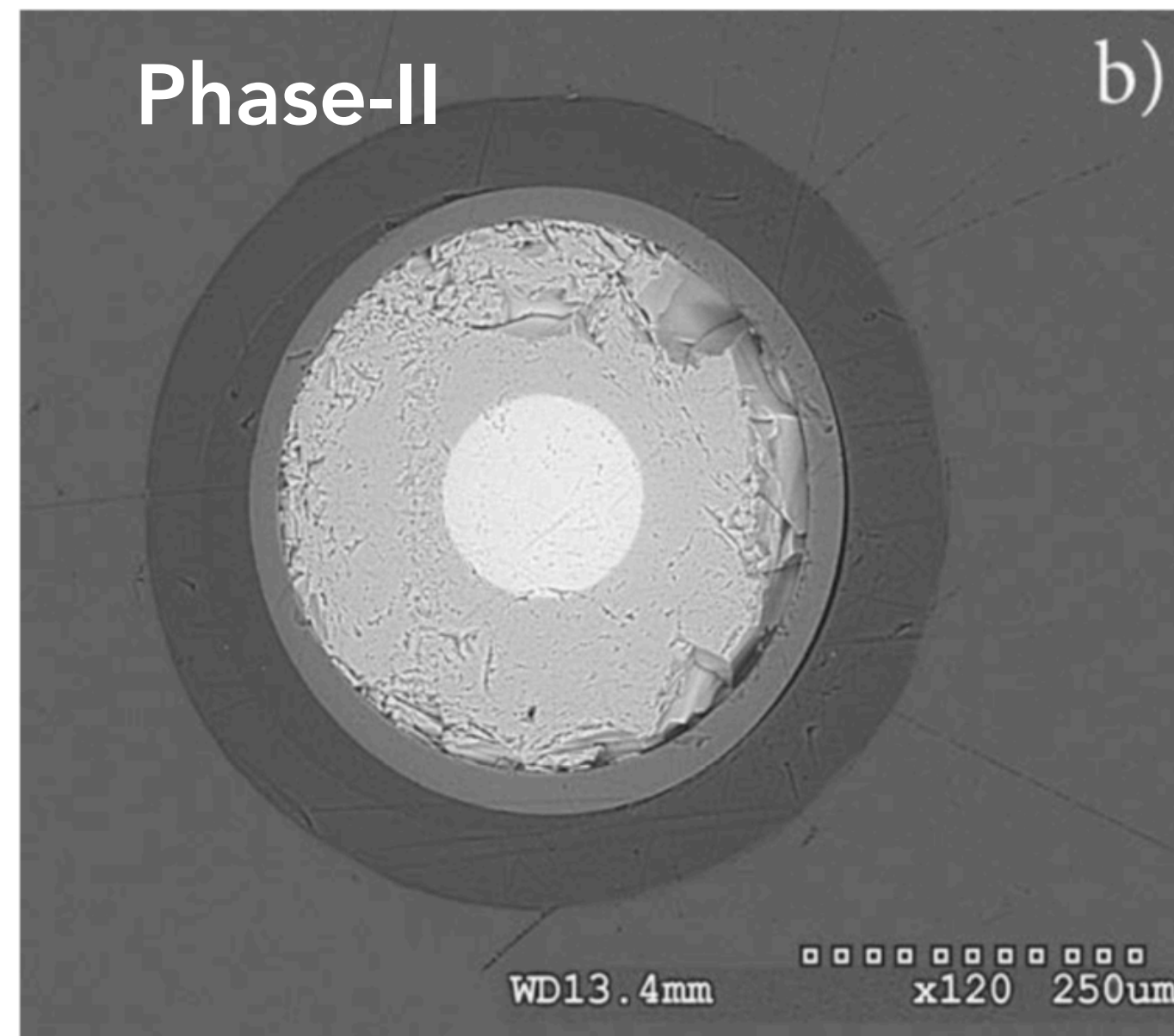
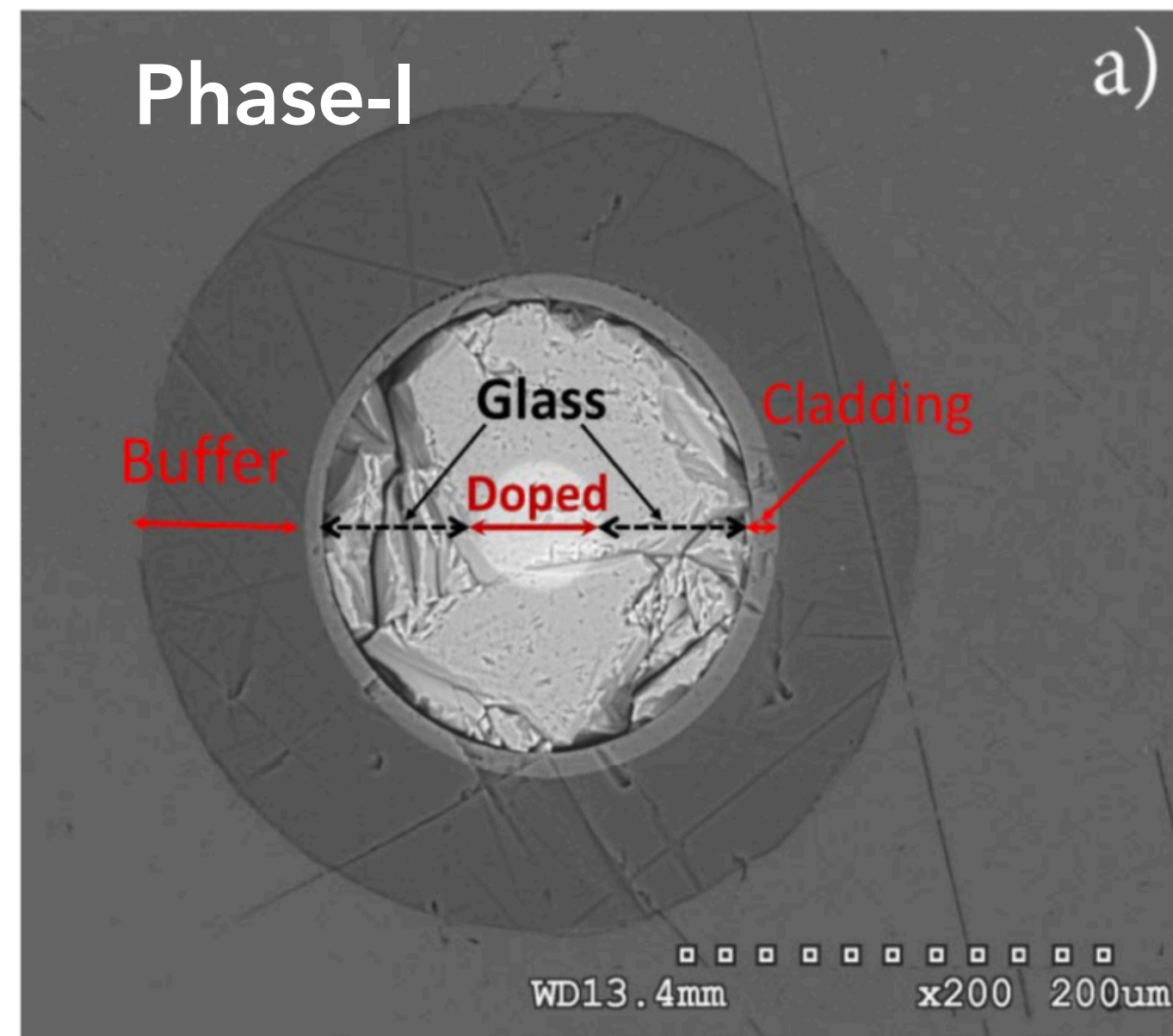
- **Two components** of the signal:
 - direct excitation of the fibers (**radio-luminescence**)
 - light from the crystal that excites the fibers (**photo-luminescence**)
- Model the signal shape analytically (**different time charact.**)
 - Un-irradiated fiber is best described with a ~100% contribution from the photo-luminescence
 - Irradiated fiber suggests that only 1/4 of the total signal comes from photo-luminescence
- Possible explanations:
 - loss of performance for both emission and transmission
 - reduced transmission of the scintillation light from cerium fluoride through the cladding of the fiber



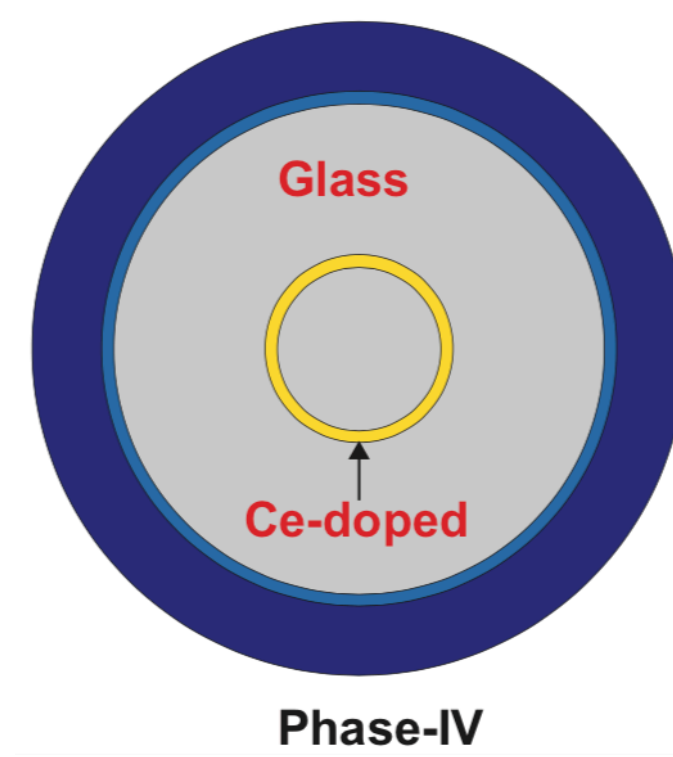
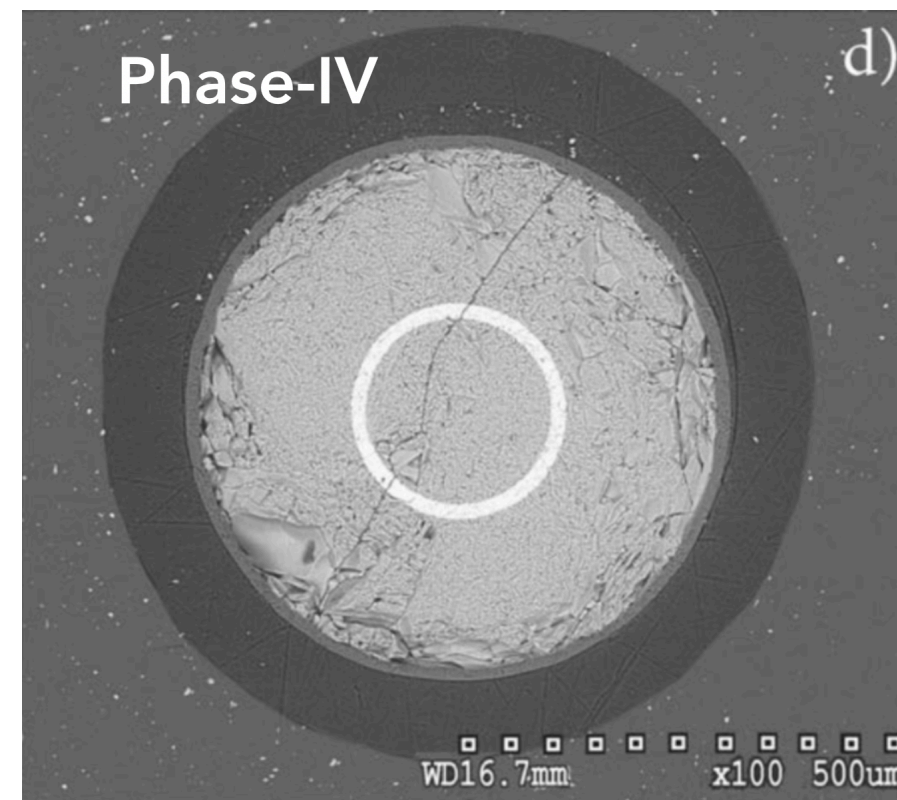
Summary and next steps

- We continue studying the light emission and transmission characteristics of cerium-doped fused-silica fibers as a potential active element in detectors
 - pulse shape, decay time, light yield, propagation speed, WLS, attenuation length, etc
- Hybrid fiber structure ("clear" + "doped") may prove useful in some detector applications in HEP
- Attention is paid to reproducibly manufacture fibers in an industrial setting
- Phase-V fibers are currently under development

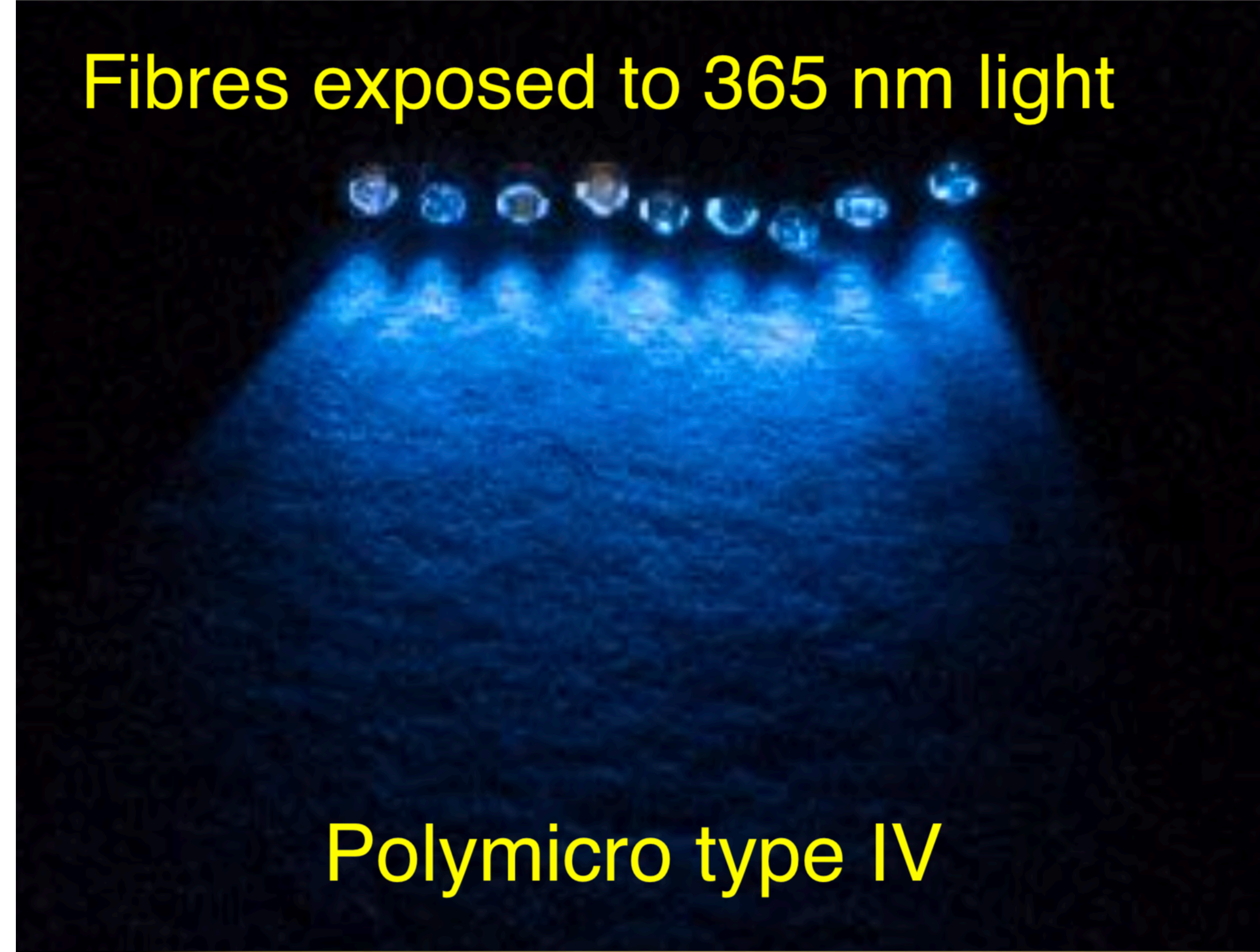
Backup



Phase-IV



Fibres exposed to 365 nm light



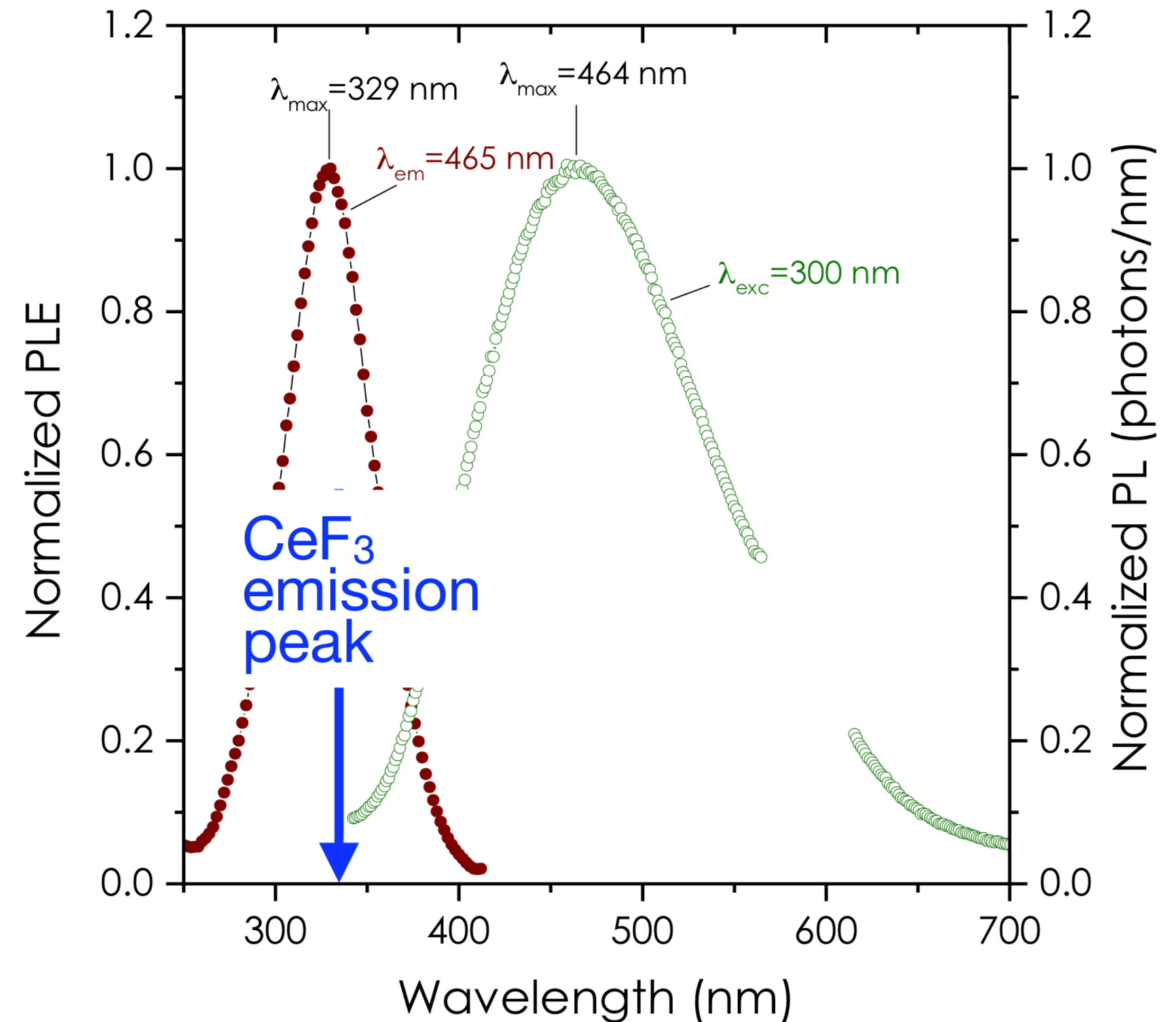
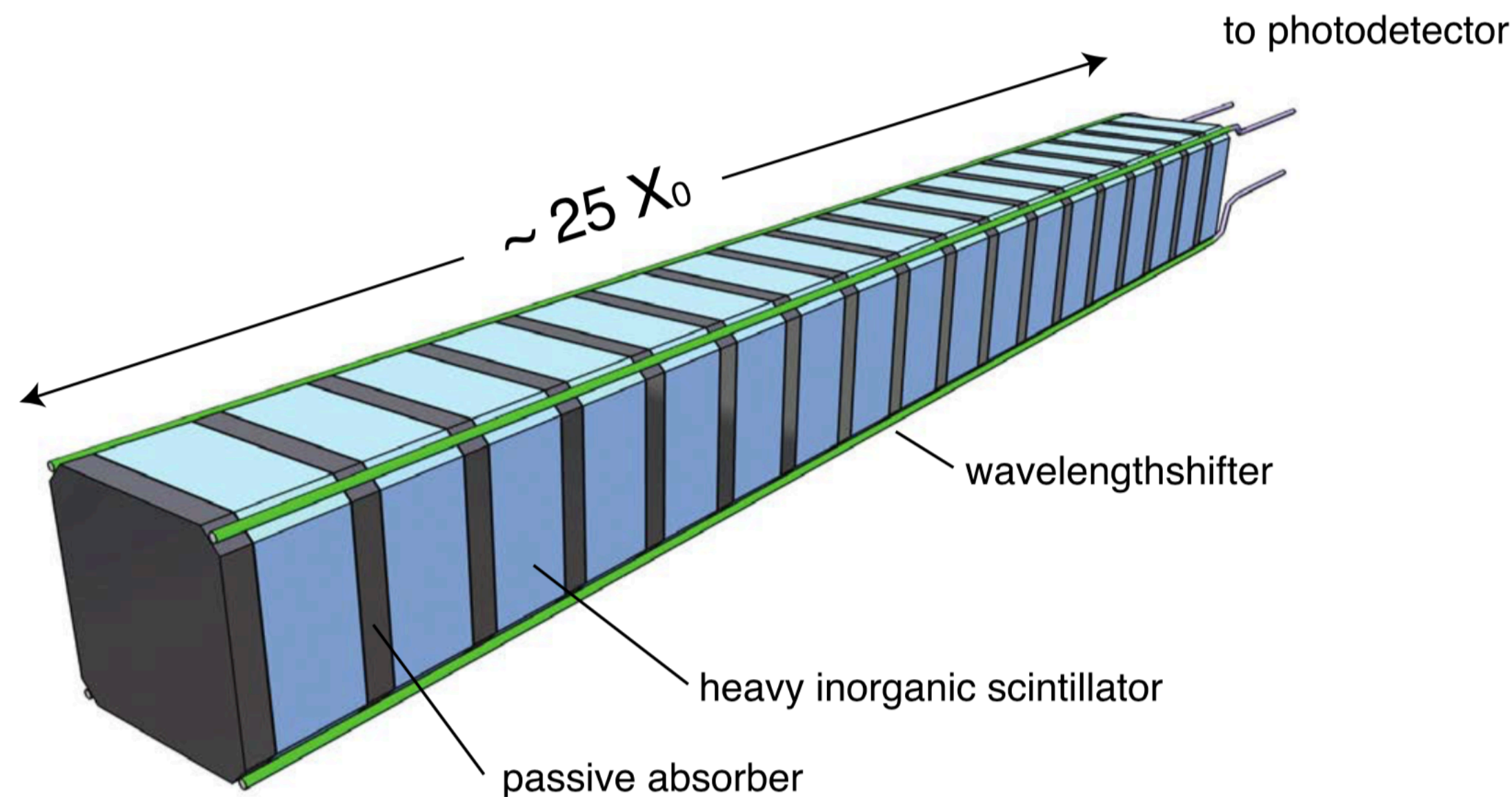
Polymicro type IV



Visible light through fibers

Tungsten CeF_3 array prototype

Francesca Nessi et al
TIPP2017



Rad-damage and optical recovery modeling

We follow Griscom's classical rate equation for second-order growth with thermally activated decay approach [34]:

$$dN(t)/dt = K\dot{D}N^* - RN^2/N^* \quad (4.4)$$

with its solution

$$N(t) = N_{\text{sat}} \tanh(kt) \quad (4.5)$$

where $N_{\text{sat}} = (K\dot{D}/R)^{1/2}N^*$ and $k = (K\dot{D}R)^{1/2}$. N stands for the number density of color centers, $R = 1/\tau$ is the decay rate constant, K governs the creation of new color centers, N^* has the dimensionality of the color center number density. The “stretched” version of the second-order growth kinetics is accomplished by the $kt \rightarrow (kt)^\beta$ substitution where kt is dimensionless.

$$dN[(kt)^\beta]/d(kt)^\beta = (K\dot{D}/R)^{\beta/2}N^* - (R/K\dot{D})^{\beta/2}N^2/N^* \quad (4.6)$$

with $0 < \beta < 1$ and $k = (K\dot{D}R)^{1/2}$. The solution is

$$N[(kt)^\beta] = N_{\text{sat}} \tanh[(kt)^\beta]. \quad (4.7)$$

Here, $N_{\text{sat}} = (K\dot{D}/R)N^*$ and $k = (K\dot{D}R)^{1/2}$. The classical time-independent rate constant K and β are determined by fitting the experimental data. The recovery in this system of equations is

$$dN[(Rt)^\beta]/d(Rt)^\beta = -N^2/N^* \quad (4.8)$$

with solution

$$N[(Rt)^\beta] = N(0)\{1 + [N(0)/N^*](Rt)^\beta\}^{-1}. \quad (4.9)$$

Experimental Setup at MTest at Fermilab



The module was installed on a movable table

The angle between the beam direction and the fiber axis was set at 2, 90 and 182 degrees

Trigger counters 1 cm x 1 cm

Cherenkov and Phase-I fiber bundles were separately readout by two PMTs

Signals were digitized at 5 GHz by a CAEN DRS unit (V1742)