

Initial X-ray absorption spectroscopy measurements in LIGO mirror coatings*

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- **Motivation**
- **Experiment**
- **Data & Analysis**
- **Results**
- **Summary & Future Work**

Objective:

Obtain physical correlations between chemical impurities and/or dopants (Ti, Cr, Fe, Co, etc.) and optical absorption characteristics of materials under consideration for use as test masses and **optical coatings** in advanced LIGO.

X-ray Fluorescence (XRF)

CAMD, SSRL, ALS

**Extended X-ray absorption
fine structure (EXAFS)**

CAMD



**X-ray absorption near edge
spectroscopy (XANES)**

CAMD, SSRL



Louisiana State University
CENTER FOR ADVANCED MICROSTRUCTURES & DEVICES



Neutron Activation Analysis (INAA)

NIST

Prompt Gamma

Neutron Activation Analysis (PGNAA)

NIST



Neutron Depth Profiling (NDP)

NIST

Electron Spin Resonance (ESR)

NIST

Four (4) samples received June 23, 2005.

Sample #1

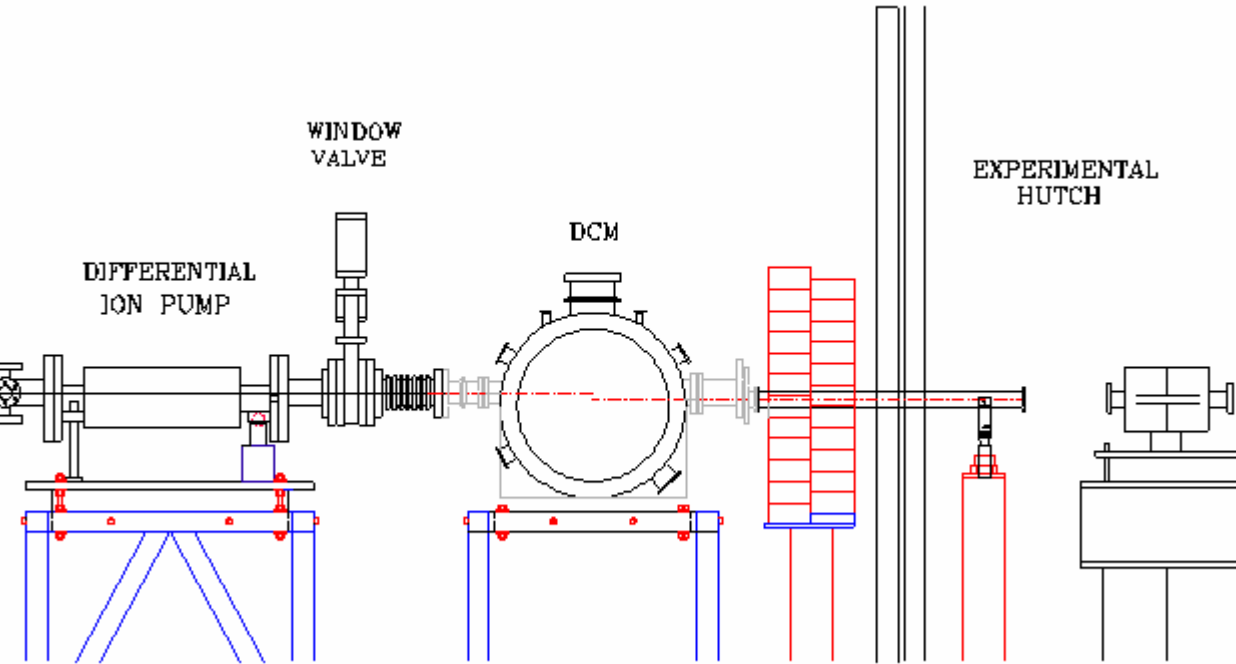
300°C MLD

Sample #3

Sample #4

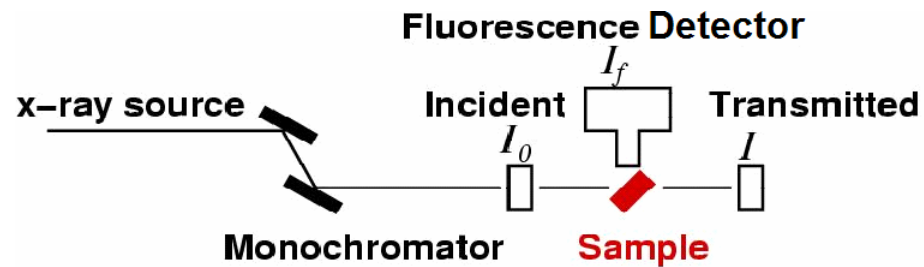
Samples are ~9.5 mm SiO₂ wedges with coatings.

Double Crystal Monochromator Beamline



CAMD Experimental Hall

13 element Ge photon detector array



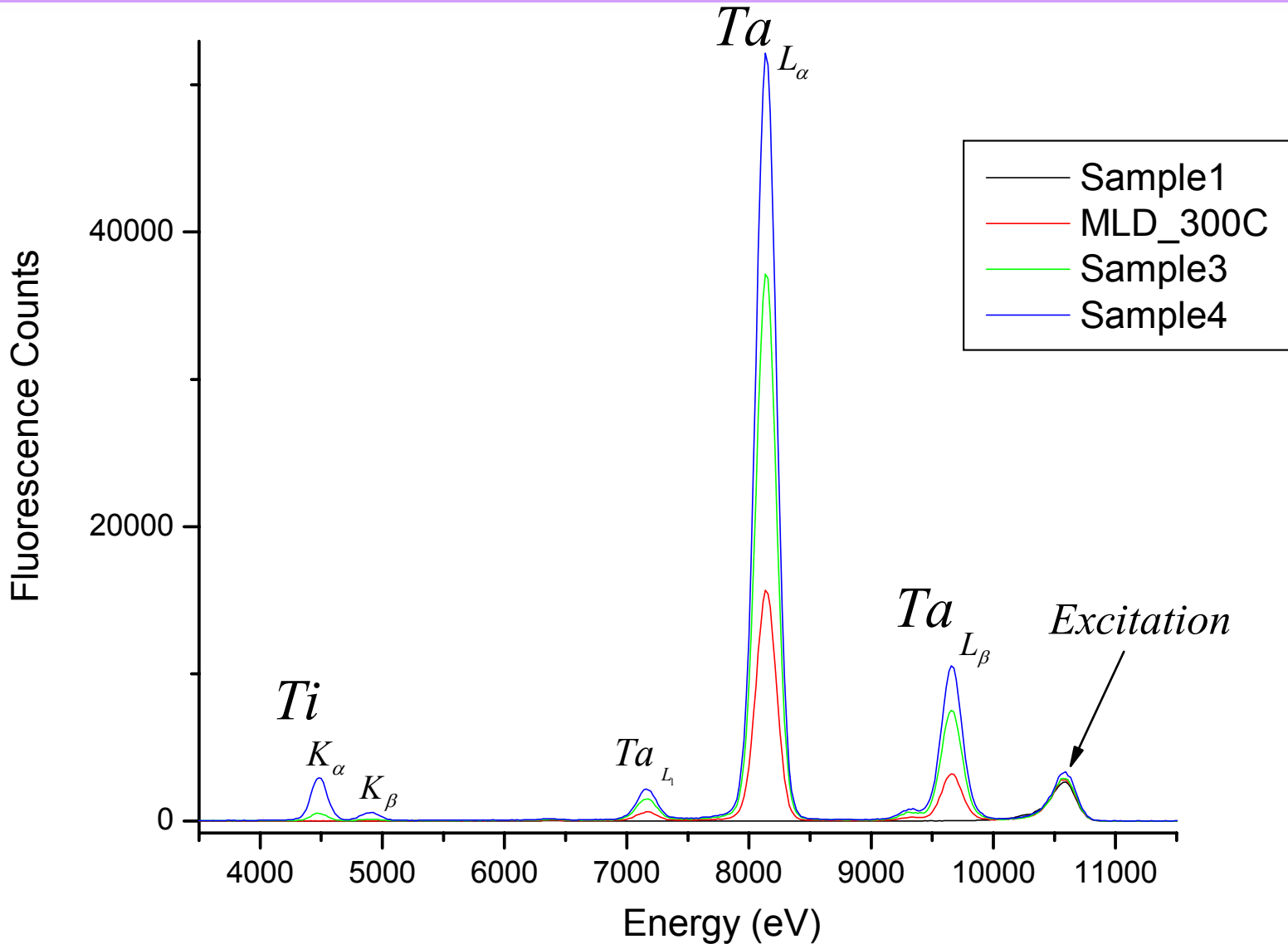
$$N_{X\text{-rays}} = N_{atoms} \int \sigma(E) \phi(E) dE \cdot [\omega(\exp(-\sum_i \mu_i \delta x_i)) \varepsilon \Delta T] \quad (1)$$

For a **monoenergetic** beam of energy, E_0 ,

$$N_{X\text{-rays}} = N_{atoms} \sigma(E_0) \phi(E_0) \omega(\exp(-\sum_i \mu_i \delta x_i)) \varepsilon \Delta T \quad (2)$$

where

$N_{X\text{-rays}}$	=	number of background-subtracted X-rays	
N_{atoms}	=	number of target atoms seen by the beam	
σ	=	photoelectric cross section	(cm ²)
ω	=	fluorescence yield	
ϕ	=	flux of incident X-rays	(#/cm ² ·s)
μ_i	=	linear attenuation coefficient(s)	(/cm)
δx_i	=	Secondary X-ray pathlength(s)	(cm)
ε	=	photopeak detection efficiency	
ΔT	=	detector live time	(s)



X-ray Absorption Spectroscopy

Excitation of a specific core electron with monochromatic X-ray source

Events prior to and subsequent to ejection of electron provide useful information about the system

XANES

(X-ray Absorption Near-Edge Spectroscopy)

Information Provided

Presence or absence of specific bonds
 Oxidation state
 Orientation

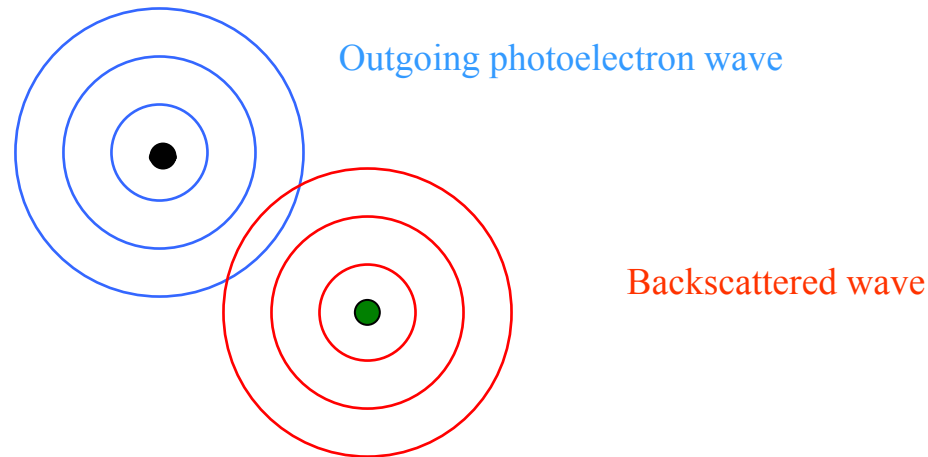
EXAFS

(Extended X-ray Absorption Fine Structure)

Information Provided

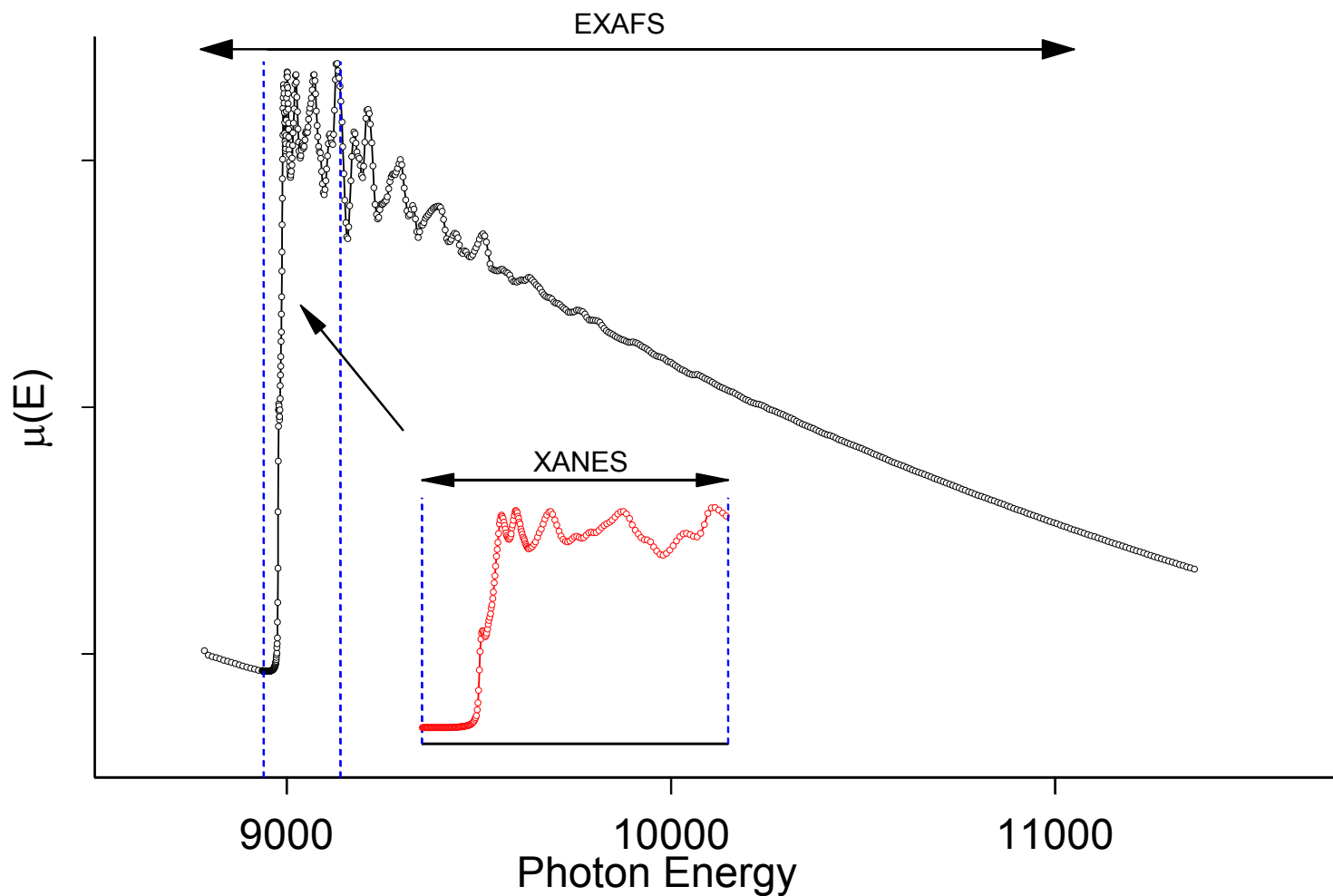
Identity of neighbors
 Neighbor coordination numbers
 Interatomic distances
 Thermal or static disorder

A visual representation

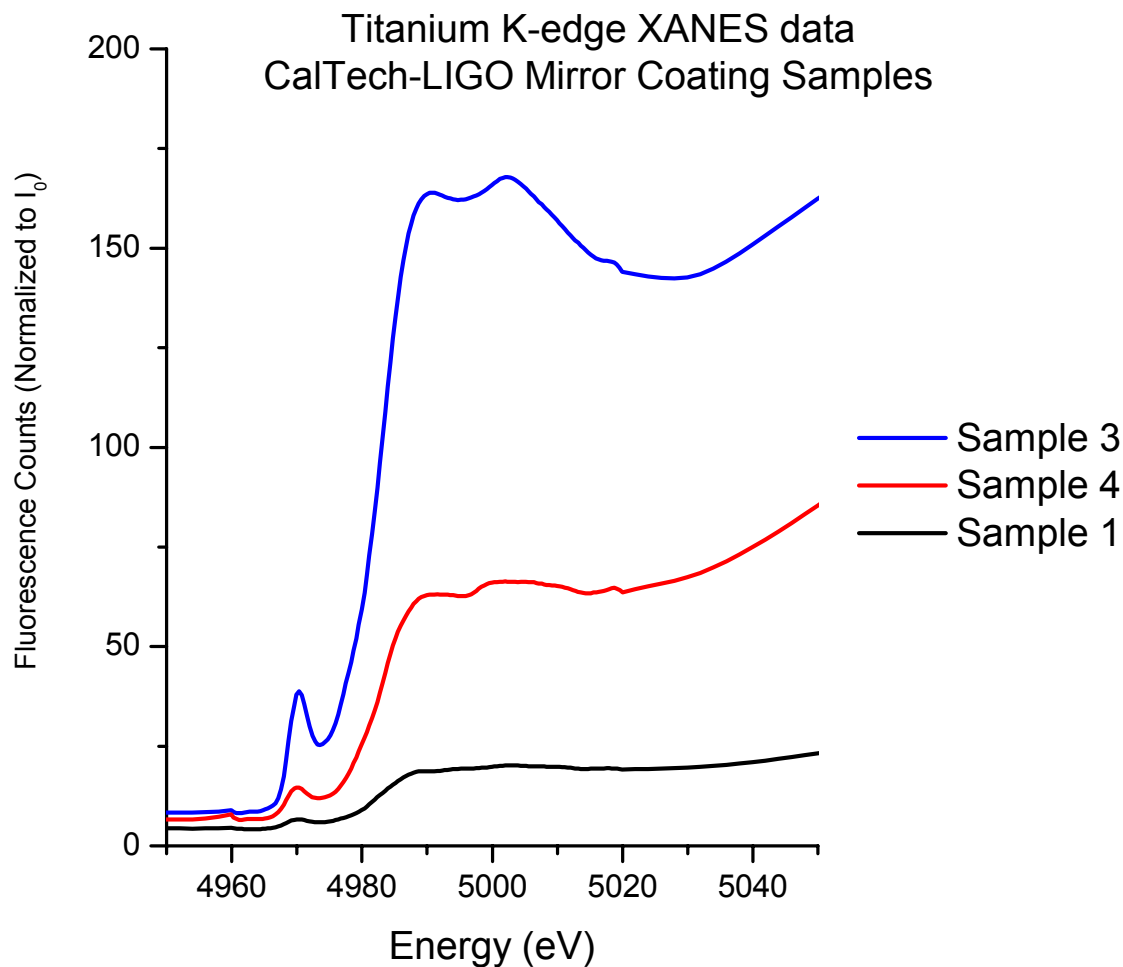


- Interaction of waves from absorber and backscattering neighbor yield information on the system.

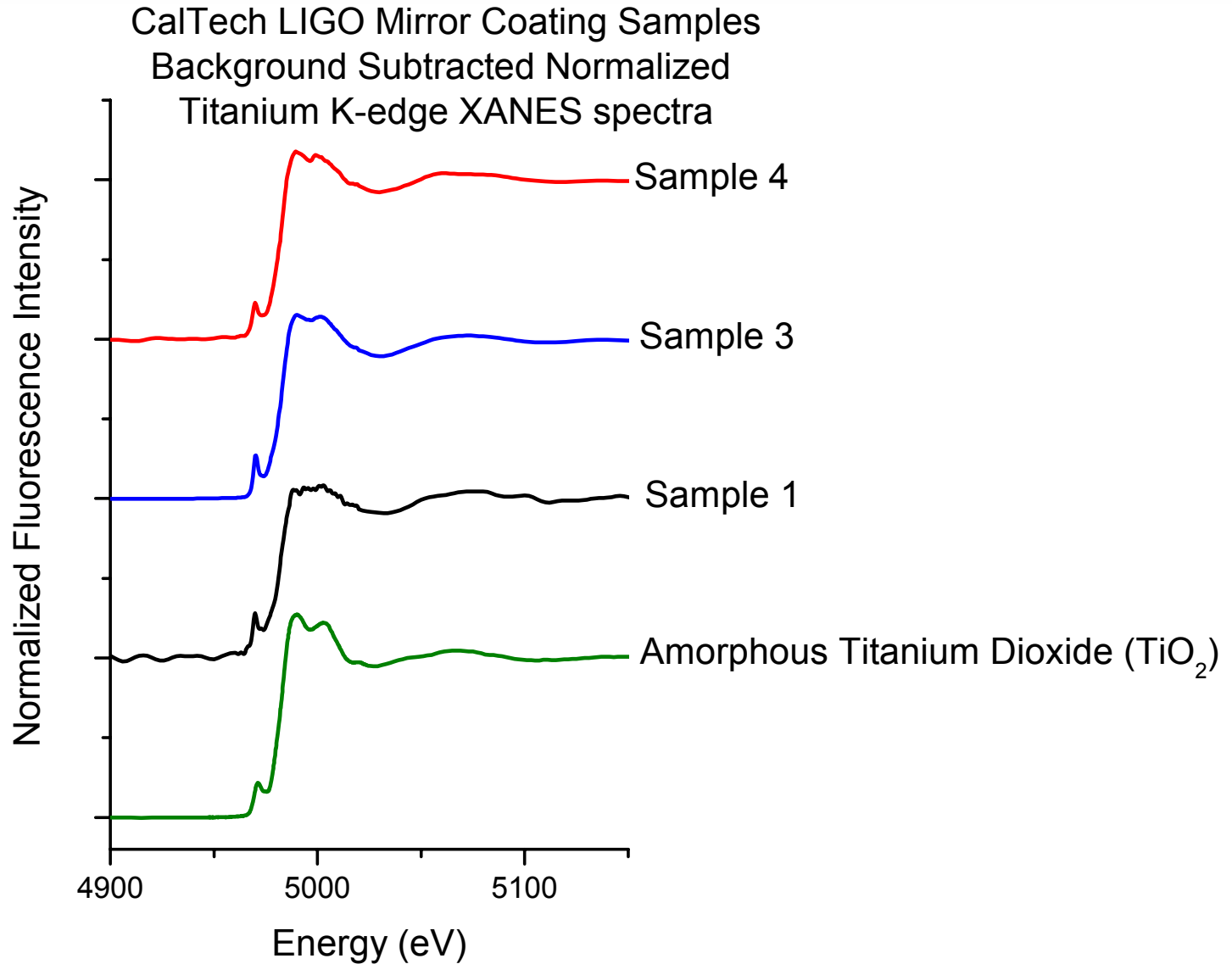
X-ray Absorption Spectrum



- The near edge region is dominated by multiple-scattering of the low kinetic energy photoelectron by neighboring atoms
- Edge position is a function of oxidation state
- Coordination and molecular orbitals influence peak shapes
- Comparison with known spectra allows for ‘fingerprinting’ of samples



- The intensity of each uncorrected spectrum in the previous figure is an indicator of the amount of titanium present (fluorescence intensity is roughly proportional to # of titanium atoms present)
- The shape of the spectrum is a function of the chemical state of the titanium (titanium oxide)



In the experiments conducted thus far:

- **Clear signals for elements of primary interest (Ti and Ta) have been observed for samples 3 and 4.**
- **Relative concentration of titanium in samples 3 & 4 has been estimated: $Ti(S4)/Ti(S3) = 6.1 \pm 2.01$ % (counting statistical uncertainty only).**
- **These initial results are encourage a long-term program of measurements.**
- **CAMD beamtime proposal is in preparation.**
- **Measurements are applicable to the investigation of other dopant/substrate systems.**

Summary & Future Work

- Identified major coating components and some trace elements on surfaces.
- Titanium is deposited as an amorphous oxide.
- Future grazing incidence experiments will allow for selective examination of coating surface with reduced scatter from substrate.
- Complementary X-ray diffraction studies are planned.
- Simulations of spectra with FEFF code are ongoing.





Questions? and/or Discussion

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