

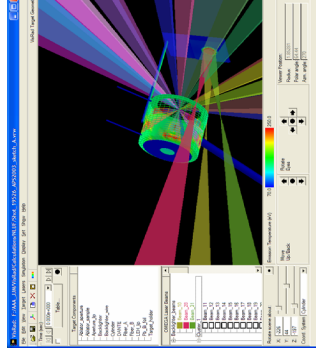
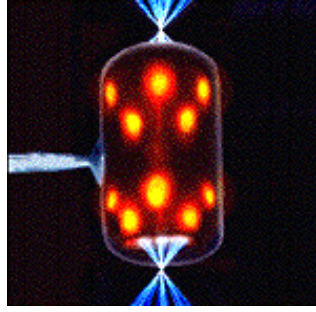
Tracer Spectroscopy Diagnostics of Doped Ablators in Inertial Confinement Fusion Experiments on OMEGA

David Cohen
Department of Physics and Astronomy
Swarthmore College

with

Joseph MacFarlane, Prism Computational Sciences
Paul Jaanimagi, University of Rochester/LLE
Otto Landen, Lawrence Livermore National Laboratory
Donald Haynes, Los Alamos National Laboratory

and
David Conners ('03), Katherine Penrose ('04), and Nathan Shupe ('05) Swarthmore College



OUTLINE

1. Scientific Context
 - ablator dopants
 - tracer spectral diagnostics
2. Experiment Design
3. Targets
4. Data and Overview of Results
5. Modeling
6. Conclusions

Context

Ablator dopants are used to control fuel pre-heat, but they also affect the radiation hydrodynamics of the interaction between the hohlraum radiation field and the capsule.

Effects of variable x-ray preheat shielding in indirectly driven implosions*

O. L. Landen,[†] C. J. Keane, B. A. Hammel, W. K. Levedahl, P. A. Amendt, J. D. Colvin, M. D. Cable, R. Cook, T. R. Dittrich, S. W. Haan, S. P. Hatchett, R. G. Hay, R. A. Lerche, R. McEachern, T. J. Murphy, M. B. Nelson, L. Suter, and R. J. Wallace
Lawrence Livermore National Laboratory, P.O. Box 5508 Livermore, California 94550

(Received 7 November 1995; accepted 25 January 1996)

The performance of indirectly driven fusion capsules has been improved by mid Z doping of the plastic capsule ablator. The doping increases x-ray preheat shielding leading to a more isentropic compression, higher convergence, and higher neutron yield. A 4× increase in neutron yield is both calculated and observed as the Ge doping level is increased from 0% to 3% by atomic fraction. A predicted 40% decrease in x-ray image core size with increasing Ge content is confirmed. © 1996 American Institute of Physics. [S1070-664X(96)93105-5]

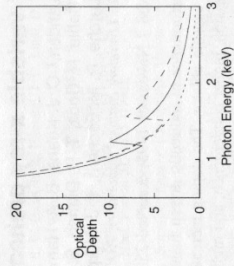
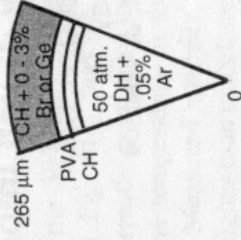


FIG. 3. Initial capsule ablator opacity vs photon energy for 58 μm thick undoped plastic ablators (short dashed curve), for 45 μm thick 1.3% Ge-doped plastic ablators (solid curve), and for 45 μm thick 1.9% Br-doped plastic ablator (long dashed curve).

(G. 1. Cross section of typical deuterated fuel capsule design.



(G. 1. Cross section of typical deuterated fuel capsule design.

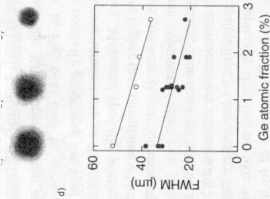
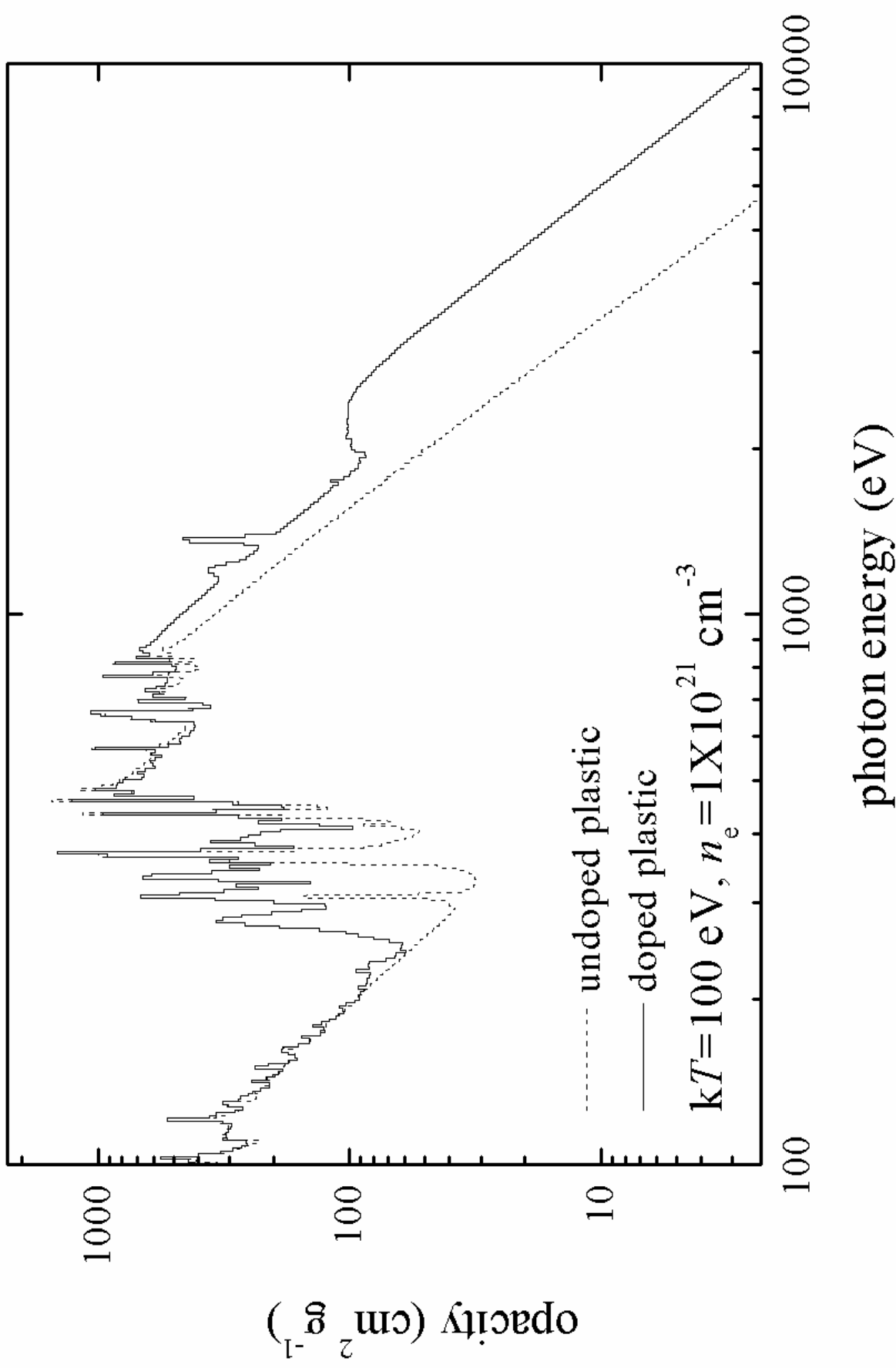


FIG. 7. X-ray images (4 keV) of imploded cores from smooth capsules at emission time for various Ge-dopant levels: (a) no Ge, (b) 1.3% Ge, (c) 2.7% Ge. Measured (open circles) and calculated (open circles) azimuthally averaged diameter (50% contours vs dopant level). Solid lines are linear fits to data and simulations.

A small amount of dopant - here 1.8% by atom of germanium - can significantly increase the opacity of a low-Z ablator



Ablator dopants affect the opacity and density, changing the manner in which energy is absorbed by the ablator.

Controlling the process requires a means of **diagnosing** the properties of doped and undoped ablators in the hohlraum environment.

Burnthrough and shock breakout experiments (ex. shown at right) measure the properties of ablators and their response to hohlraum radiation fields *integrated* over the duration of the experiment.

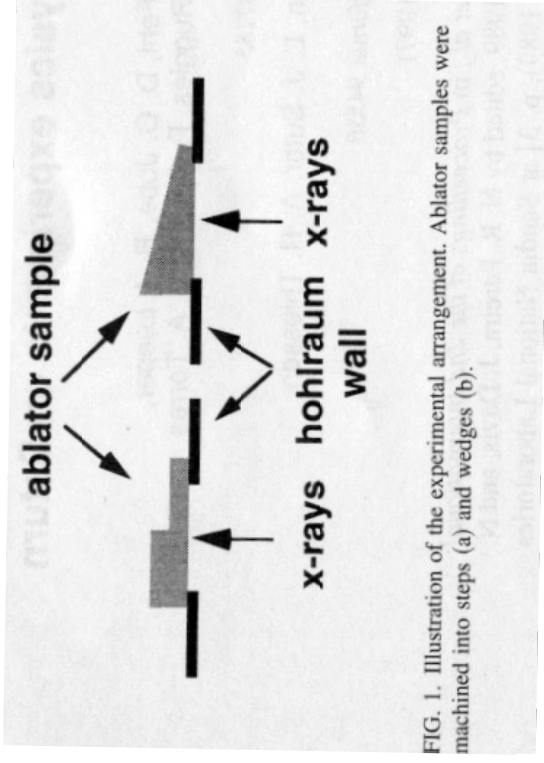


FIG. 1. Illustration of the experimental arrangement. Ablator samples were machined into steps (a) and wedges (b).

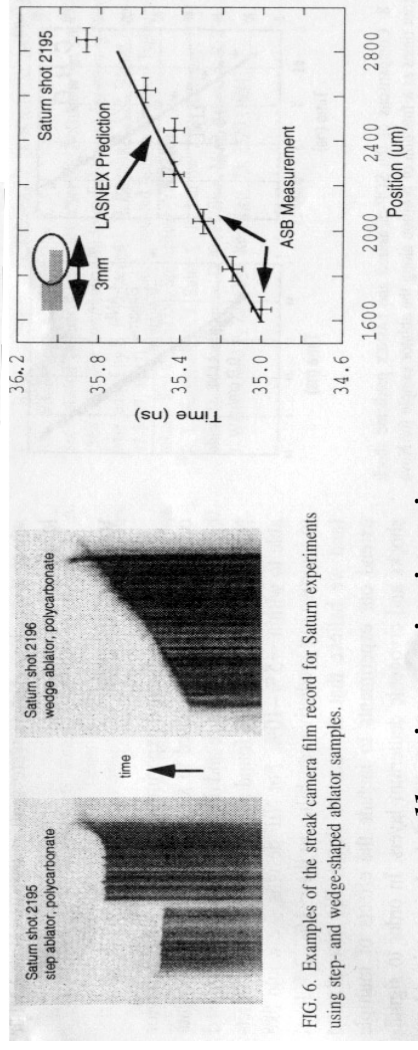


FIG. 6. Examples of the streak camera film record for Saturn experiments using step- and wedge-shaped ablator samples.

Also, ablators have been evaluated *spectroscopically* via emission in gas-filled capsule implosions.

In a different context, Perry *et al.* showed that absorption spectroscopy in multi-layered targets could diagnose radiation transport.

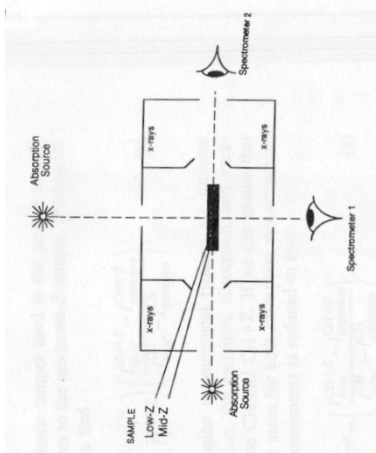


FIG. 1. (a) Schematic of an experiment designed to allow quantitative analysis of the absorption spectrum. The key elements are the backlight, the sandwich target, and a recording medium that allows the various sources to be measured at one time. (b) Schematic showing two spectrometers as used in the PPS technique.

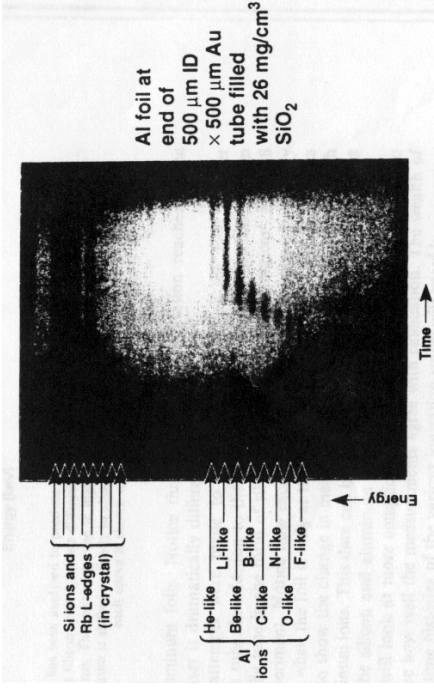


Fig. 8. The x-ray crystal streak camera record for a filled tube is shown. Filling the tube with 26 mg/cm^3 of silicon aerogel only slightly slows the transfer of heat but provides more information about the transfer of heat throughout the entire length of the tube. The Rb *L*-edges that overlay some of the silicon lines can be eliminated on future experiments by using a KAP crystal.

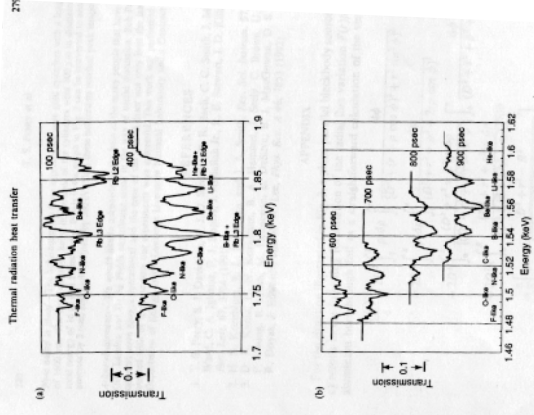


Fig. 9. The data in Fig. 8 has been analysed to produce spectra in transmission as a function of time. (a) The upper graph shows the transmission spectra at four different times. The shift with time to higher ionization states as time changes is clearly seen. (b) Note that each spectrum is shifted vertically to allow easy comparison. The transmission scale for

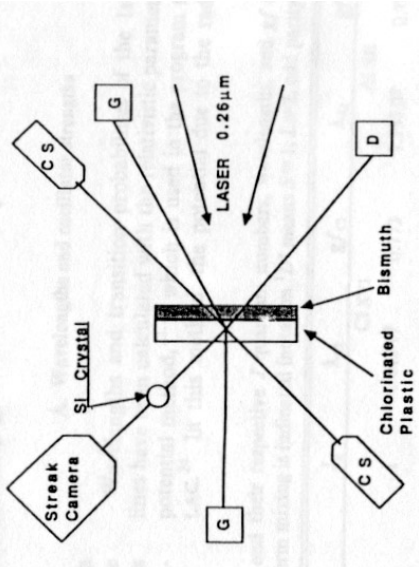


FIG. 2. Set up of the experiment. CS; crystal spectrographs with PET or ADP crystal; G; transmission grating spectrograph; D; filtered diodes array.

And Chenais-Popovics *et al.* showed that Cl K_{α} absorption spectroscopy could diagnose material properties. Laser-produced Bi plasma provided the backlighter continuum source.

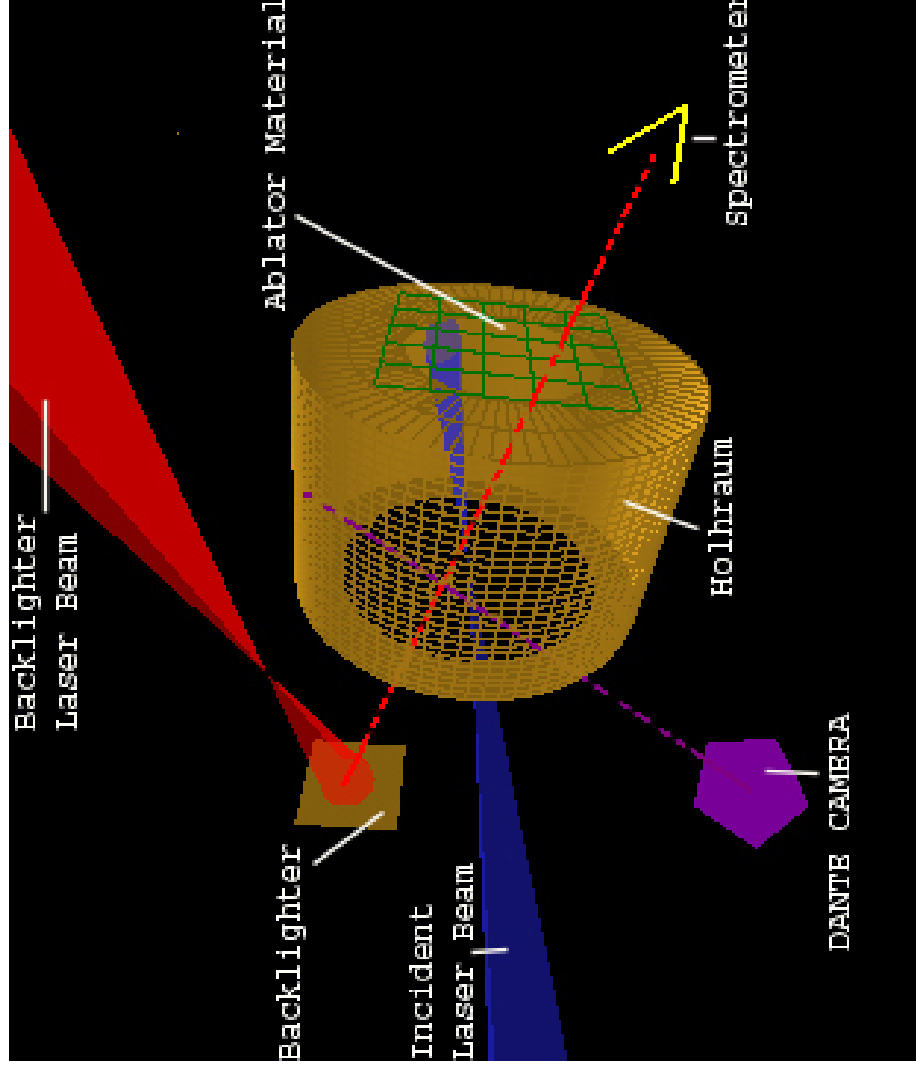
We have combined these ideas, and building on a previous effort to measure tracer emission spectra from aluminum witness plates with Tina Back, proposed an experimental campaign to use backlit Cl K $_{\alpha}$ absorption spectroscopy to diagnose radiation physics in the *interior* of ablator samples.

We proposed to do this by placing thin tracer layers at specified depths in the interiors of ablator samples mounted on hohlraums. The spectroscopy monitors the ionization conditions in that layer, effectively diagnosing the time-dependent plasma properties at a specific location inside the sample.

A time-delay in the turn-on time of the tracer signal between doped and undoped samples should allow us to determine the effects of dopants on the Marshak wave propagation.

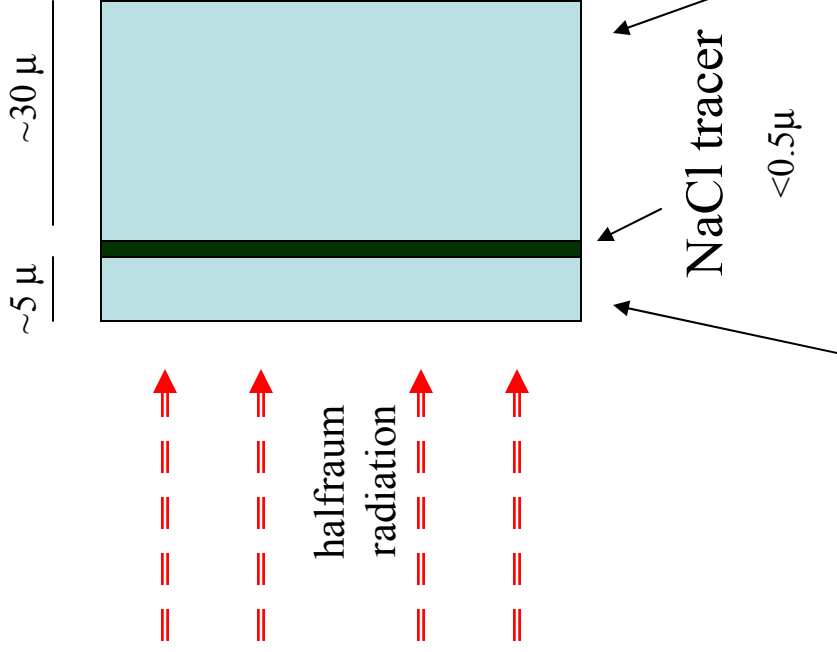
Experimental Set-up

Including schematics of diagnostic lines-of-sight



Note: only one (blue) beam into the halfraum is shown here, for simplicity.
All shots were carried out with 15 beams.

Thin tracers embedded at known depths in planar ablator samples provide a spectroscopic signal when the Marshak wave reaches a specific location in the ablator.



We will be comparing data from two shots in our OMEGA campaign:

19526: undoped, tracer depth 6.3μ

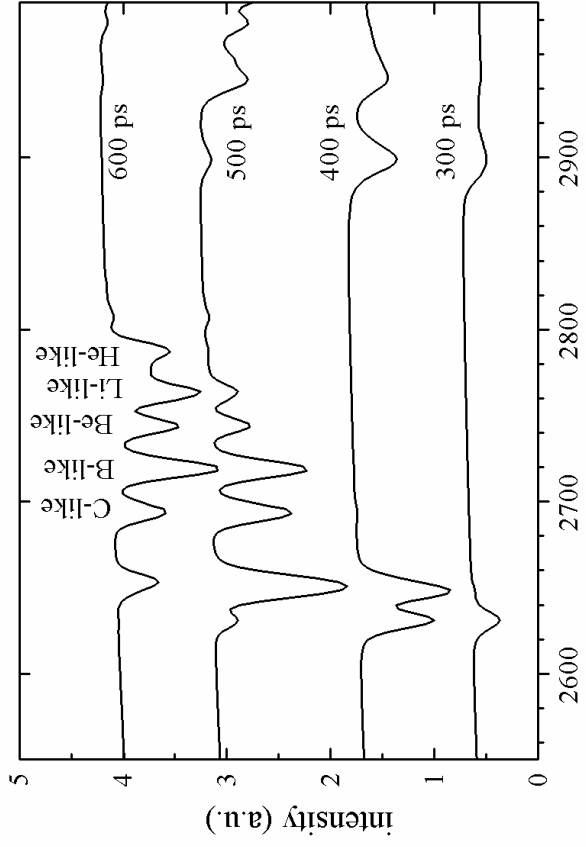
19528: doped, tracer depth 4.1μ

The time-dependent drive spectrum is *modeled* and constrained by DANTE

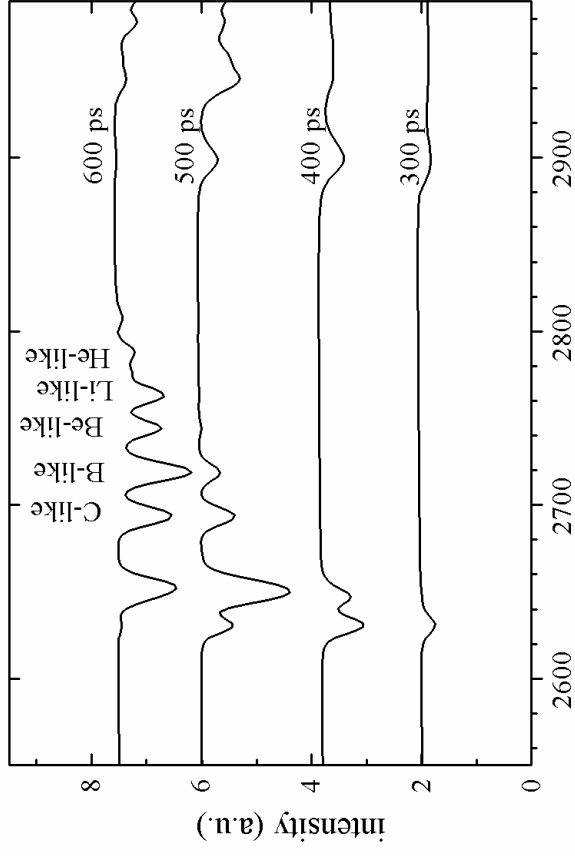
Ge-doped or undoped plastic

What do we expect to see?

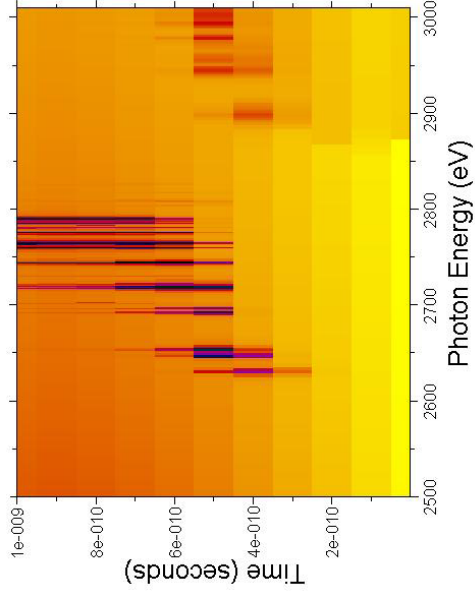
undoped



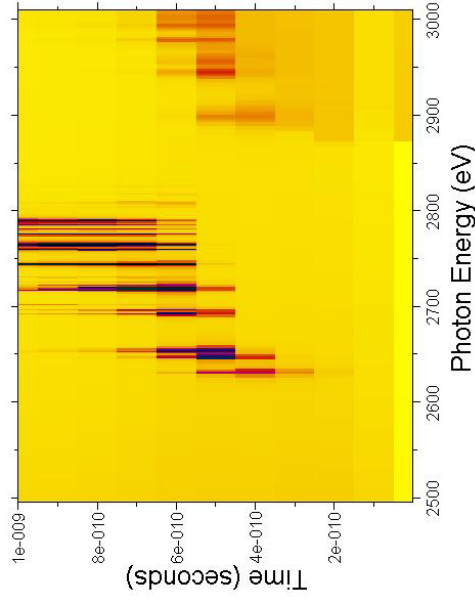
doped



photon energy (eV)



photon energy (eV)



Tracer signals turn on earlier in the undoped sample

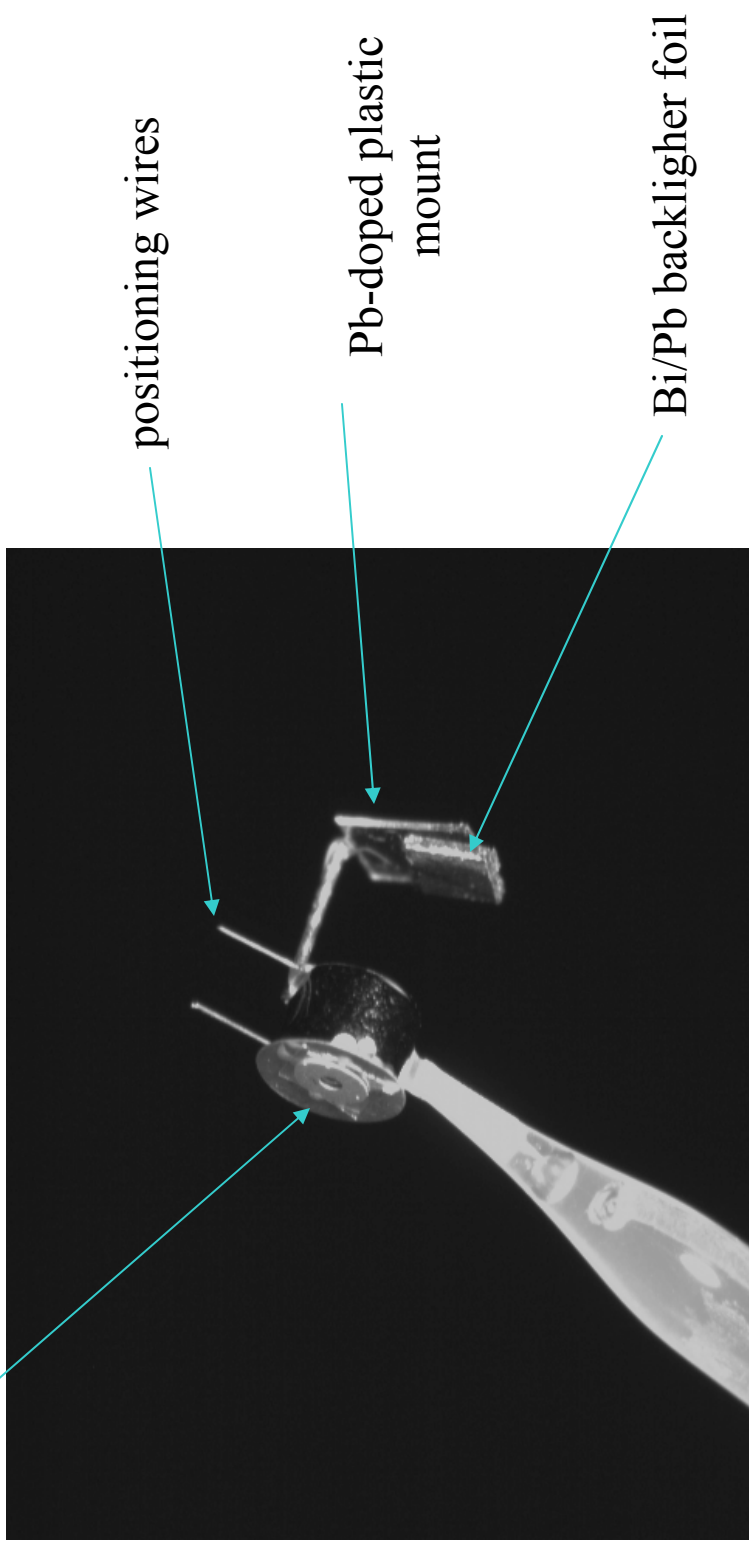
Targets: Fabrication and Assembly

Experimental Configuration

LEH facing P-7 (LXS in P-6)

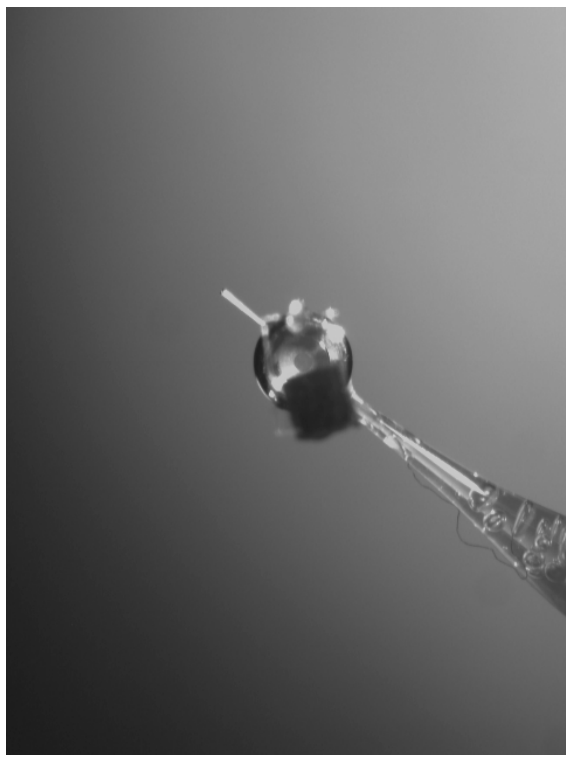
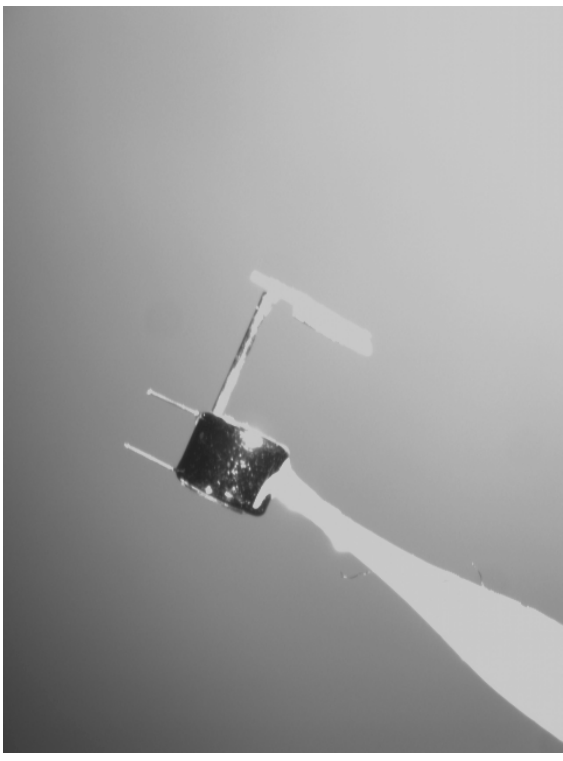
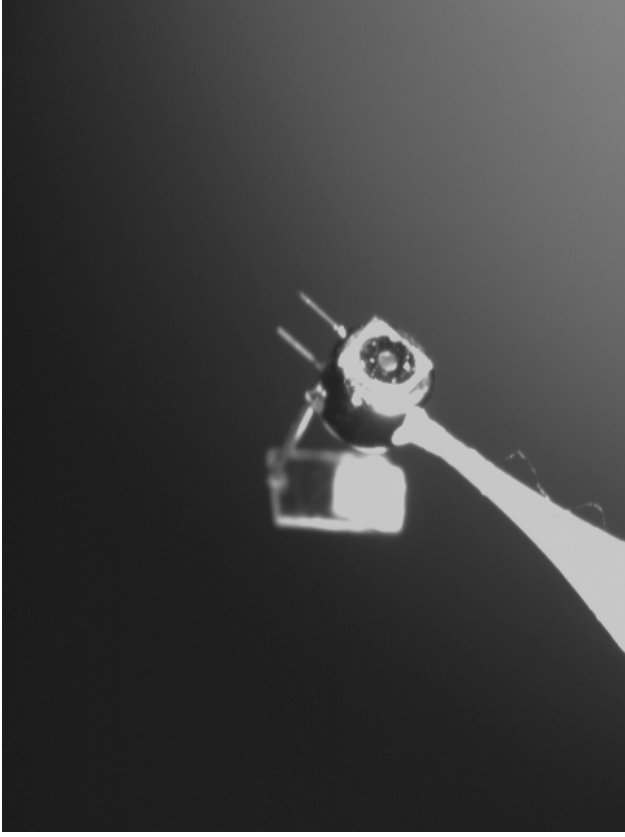
Gold Halfraums: $L=1200\mu$, $R=800\mu$

washer/aperture



TVS-X view

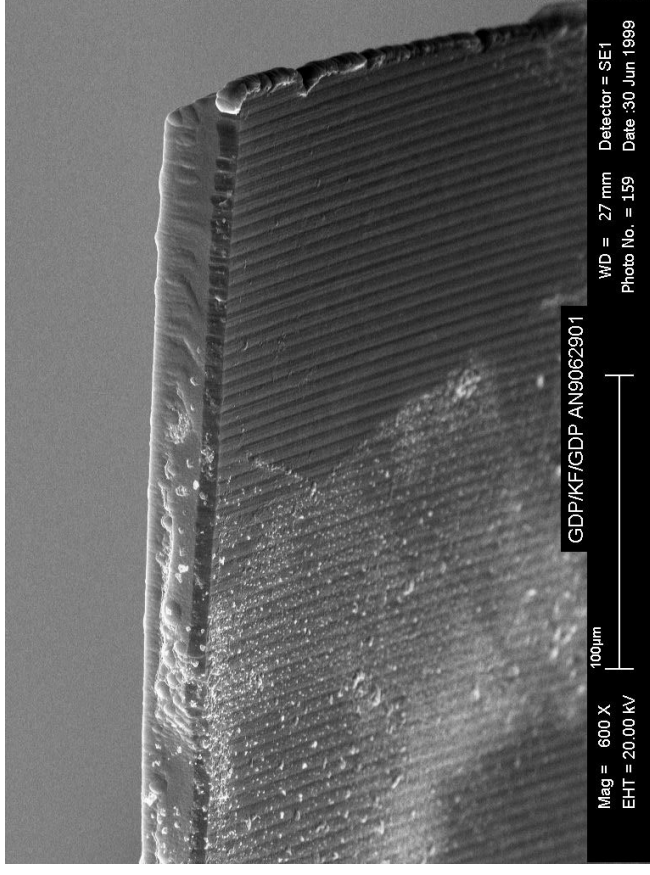
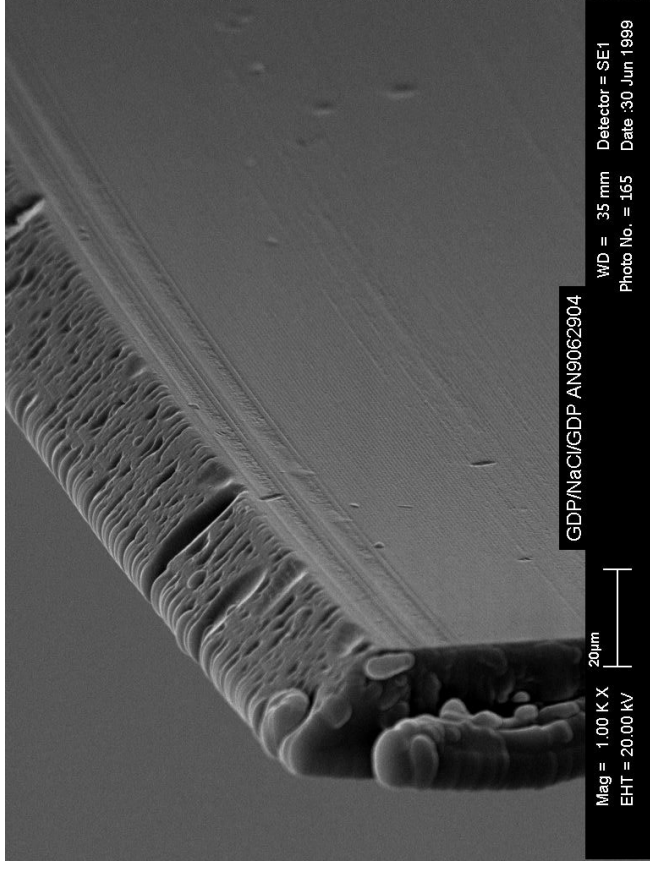
Ablator samples were mounted on the ends of halfraums; backlighter foils hung $\sim 1.5\text{mm}$ from LEH



Several views of a target: taken at the LLE metrology lab

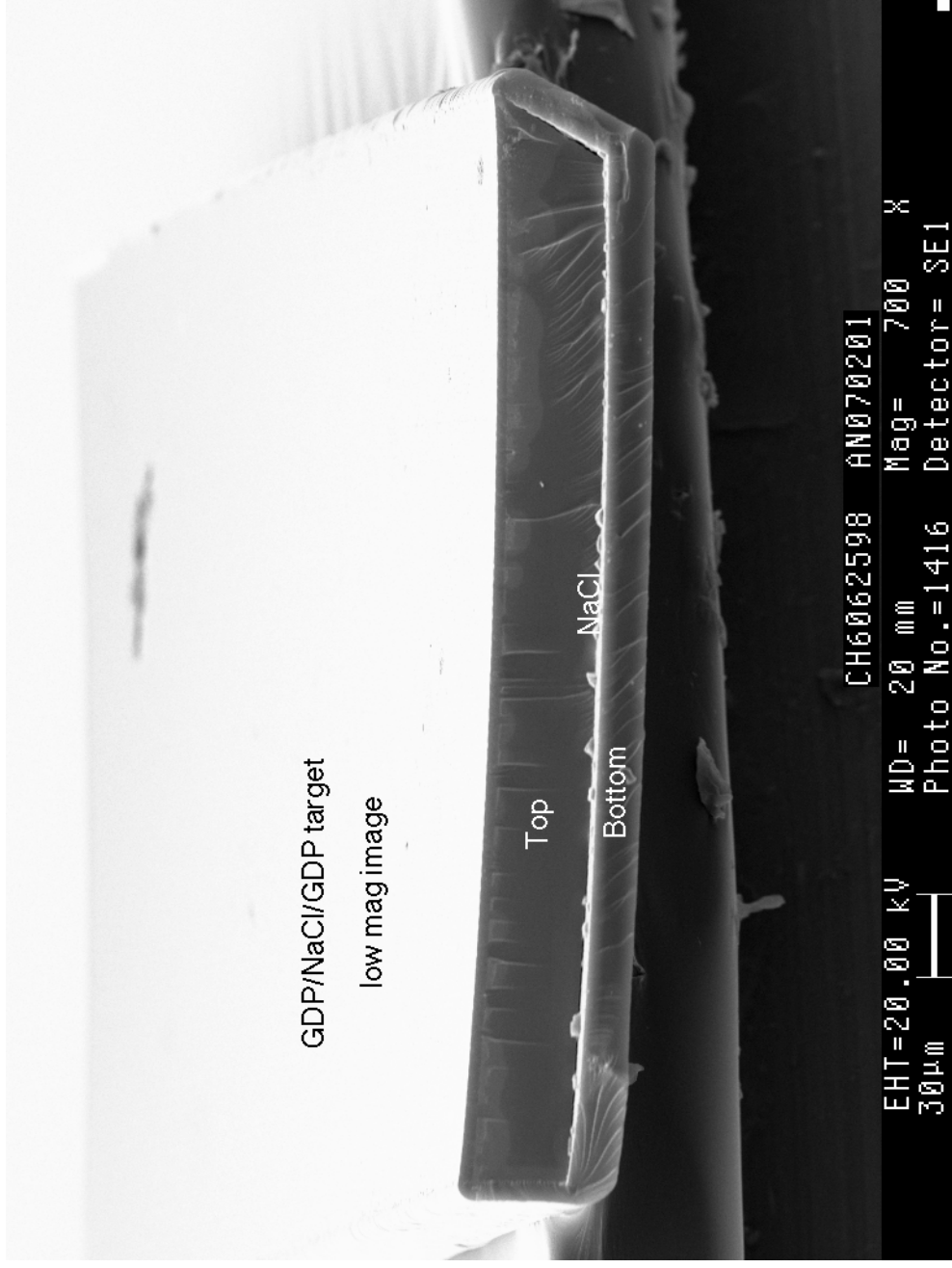
SEM images of finished ablator samples

Leakage of KF tracer onto front of plate (right), but no similar problem in samples with NaCl tracer (left); we used the ablator samples with NaCl tracers.



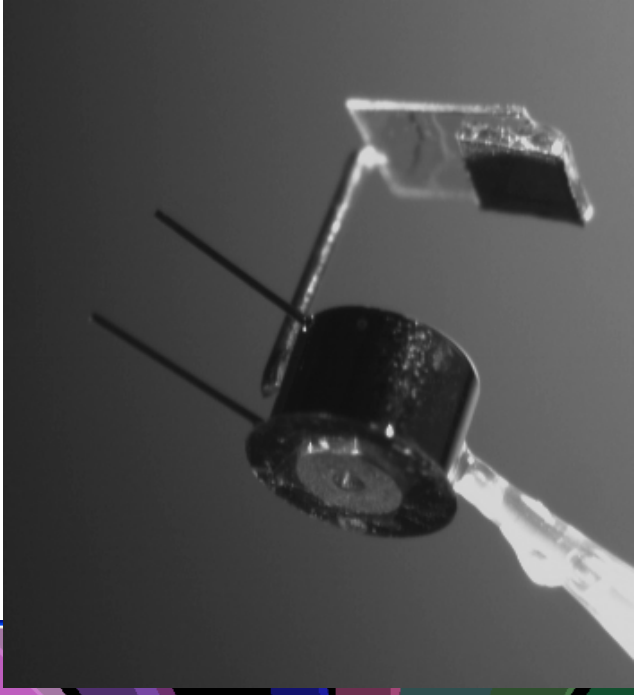
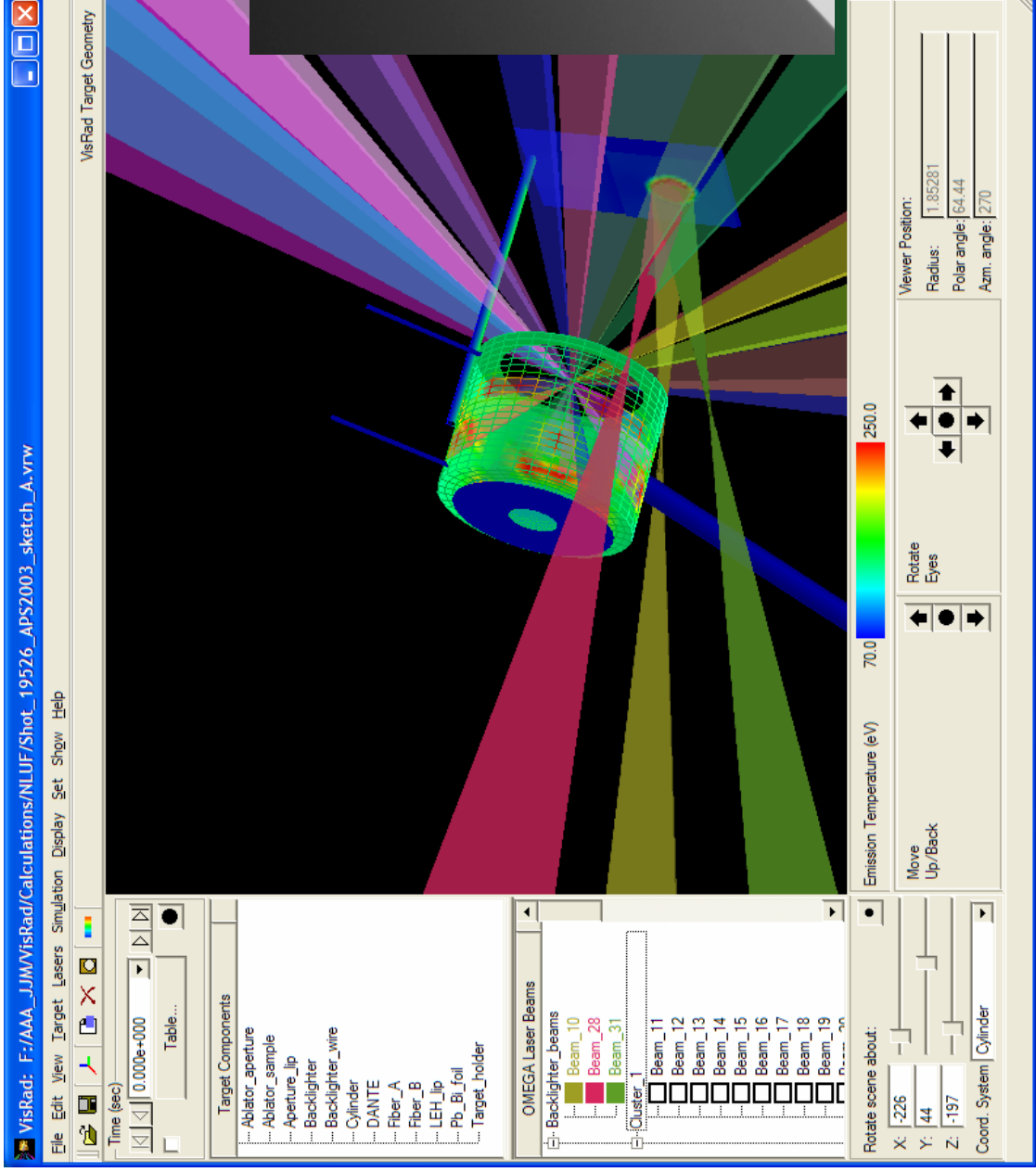
General Atomics produced these witness plates by first making the thicker plastic layer via glow-discharge polymerization (GDP); The salt layer was deposited on this plastic, and then the whole assembly was put back in the GDP chamber and an additional $\sim 5\mu$ of plastic was deposited on top.

SEM image of cross-section of target



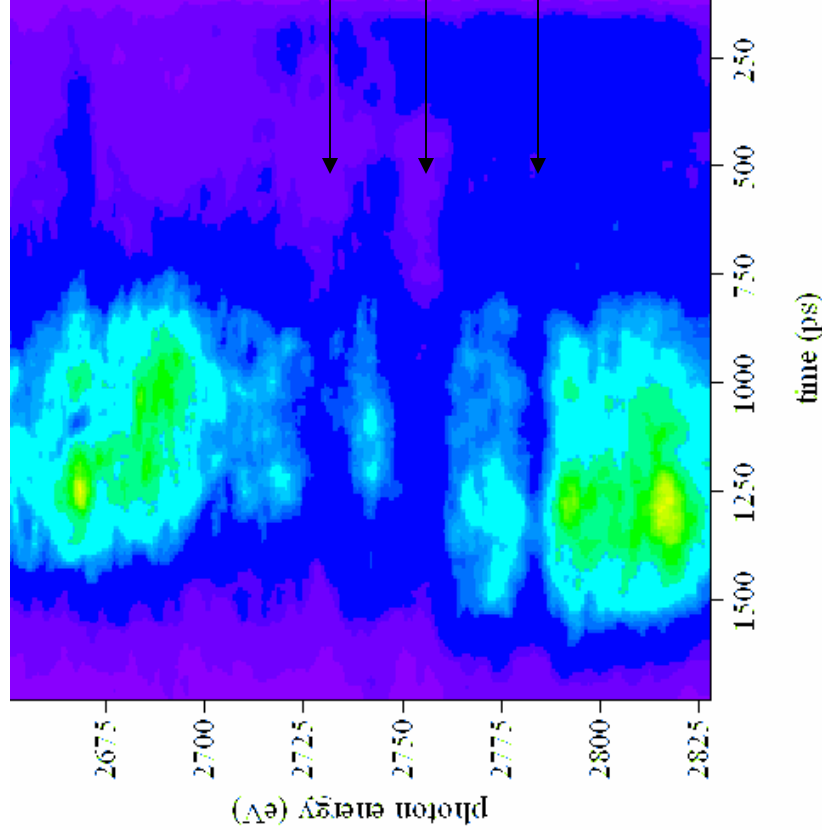
Note the salt crystals...perhaps an artifact of SEM sample preparation, but would weaken the spectral signal if representative of tracer properties in the samples we shot.

So, we've got our assembled targets, and now we're going to shoot the halfraums

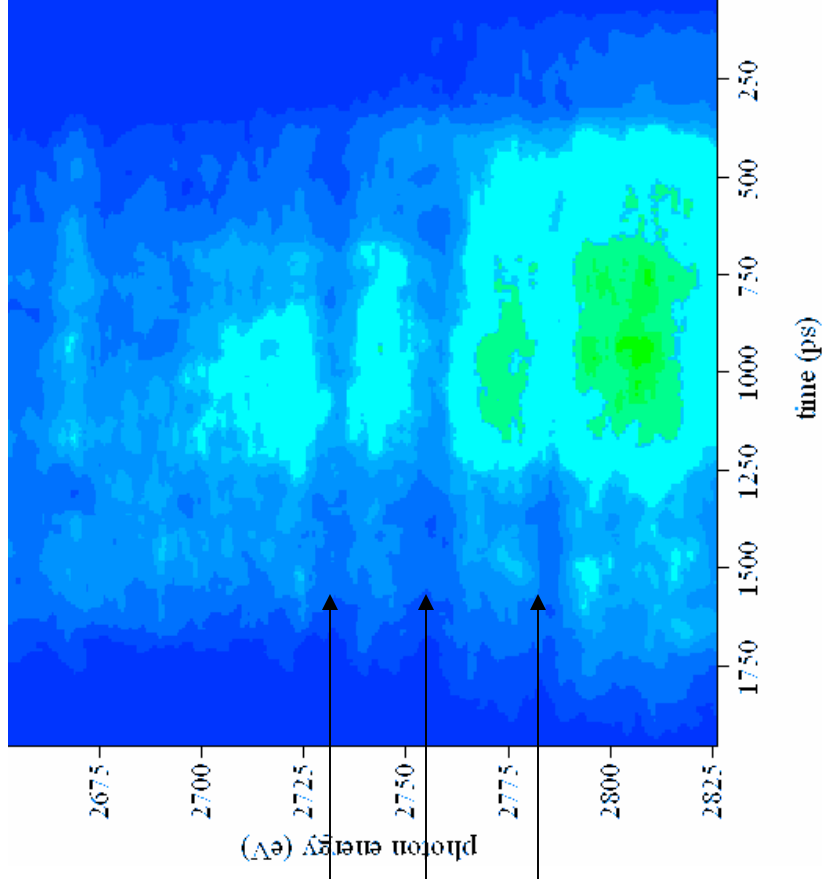


The Spectroscopic Data

undoped



doped



Digitized section of film from LXS/streak-camera:

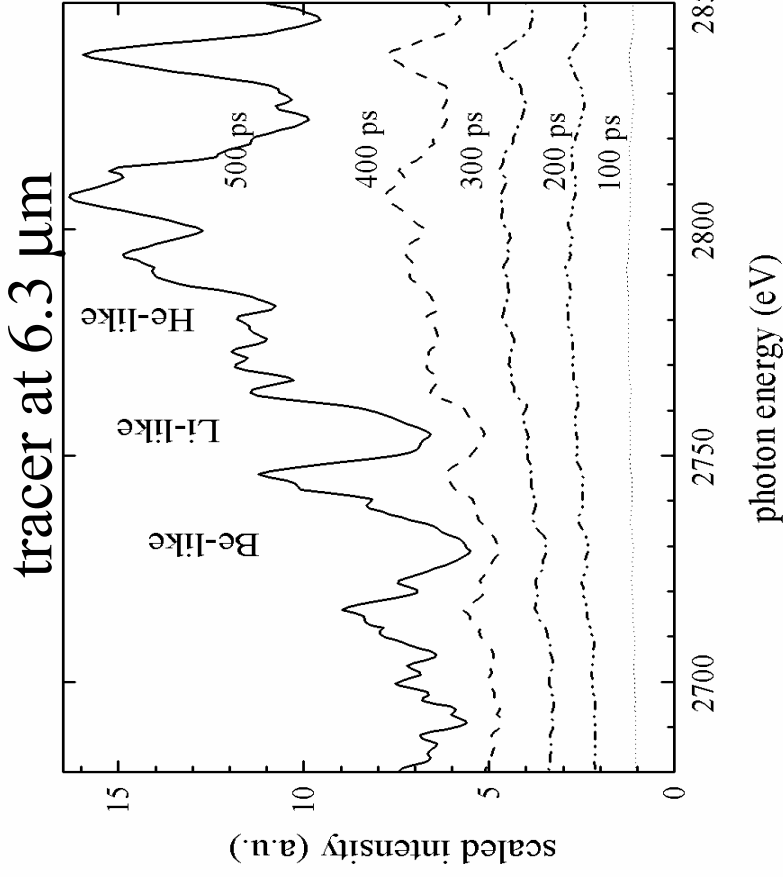
Features from three ion stages of chlorine are indicated;

Note the temporal and spectral structure in the backlighter emission; this makes it difficult to bring out the tracer features effectively at all times in this color-scale representation. Note also that the backlighter signal is weak at early times (we staggered the backlighter beams in time).

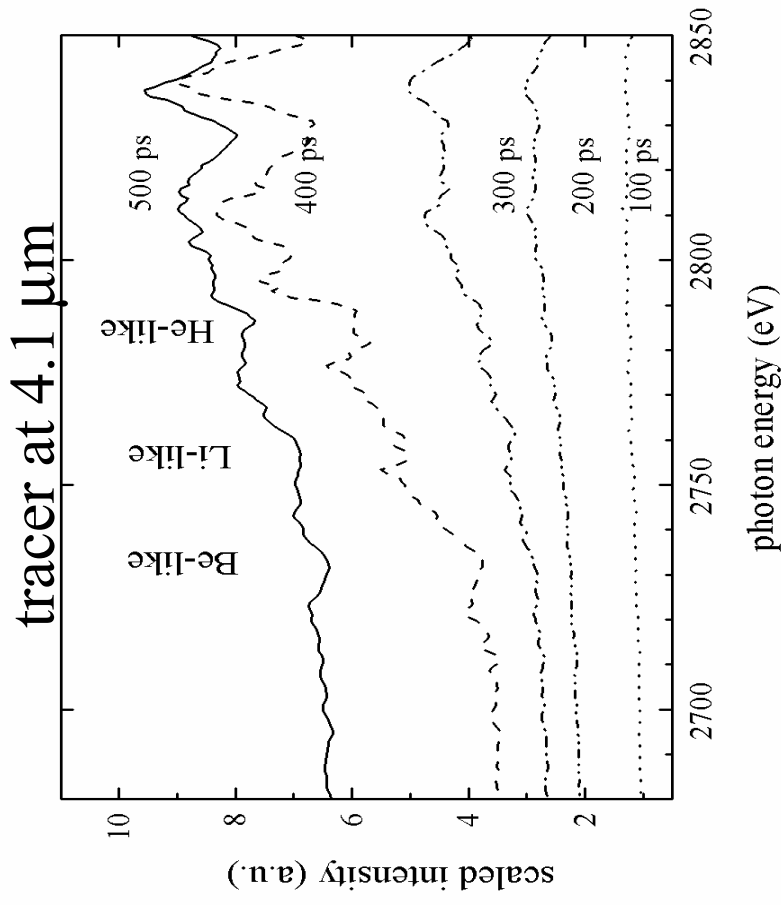
Data: time-resolved tracer spectra:

lineouts from film shown on previous slide

shot 19526: undoped



shot 19528: doped



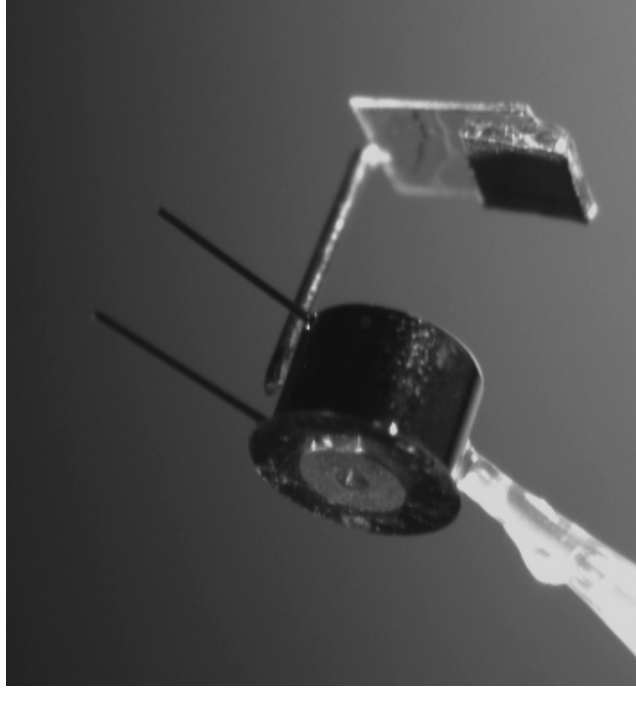
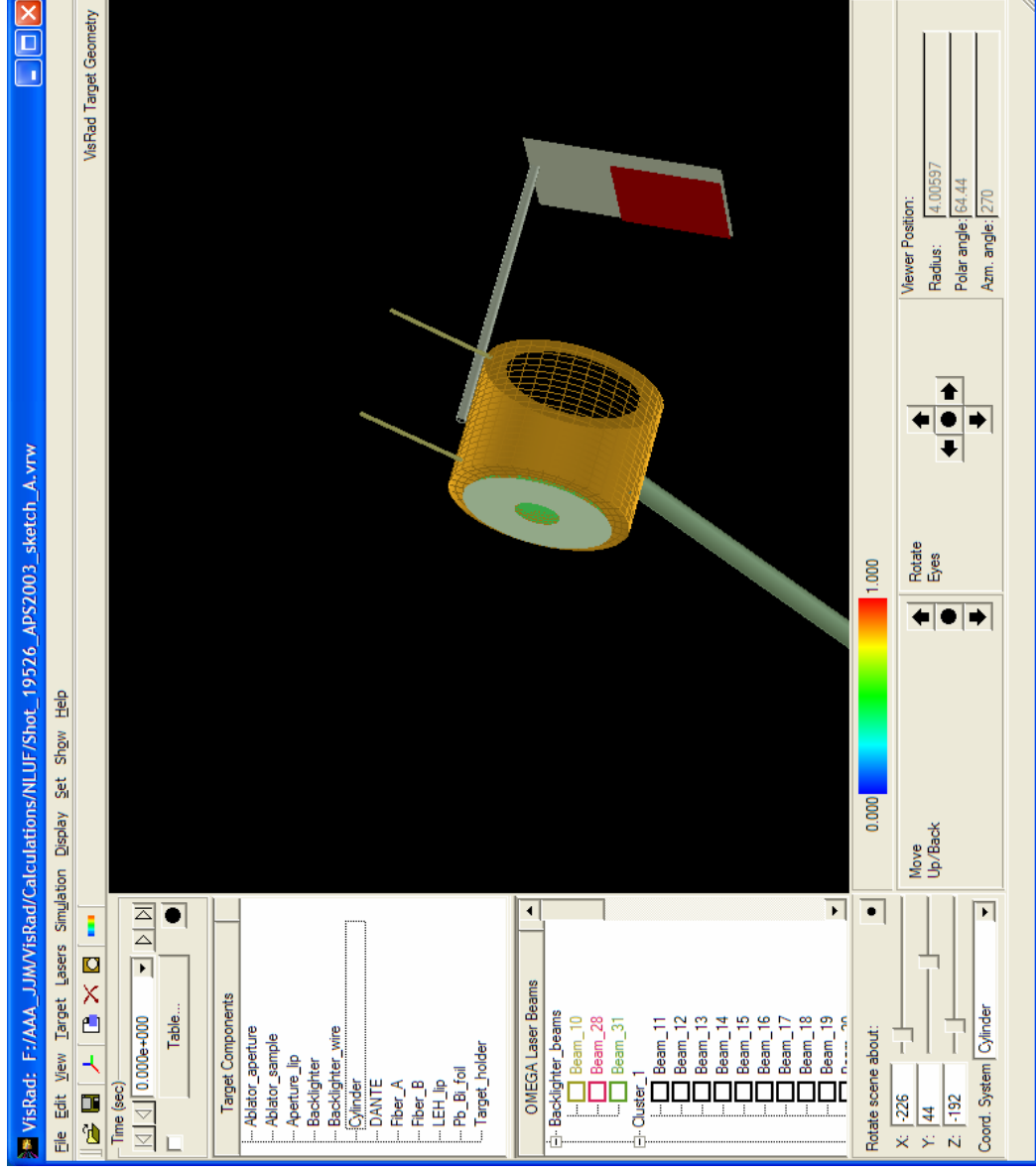
Undoped: Be-like @ 300 ps, Li-like @ 400 ps;

Doped: up to He-like @ 400 ps

Taken at face value, the data seem to indicate the slower Marshak wave propagation in the doped sample, but before we can draw any conclusions...

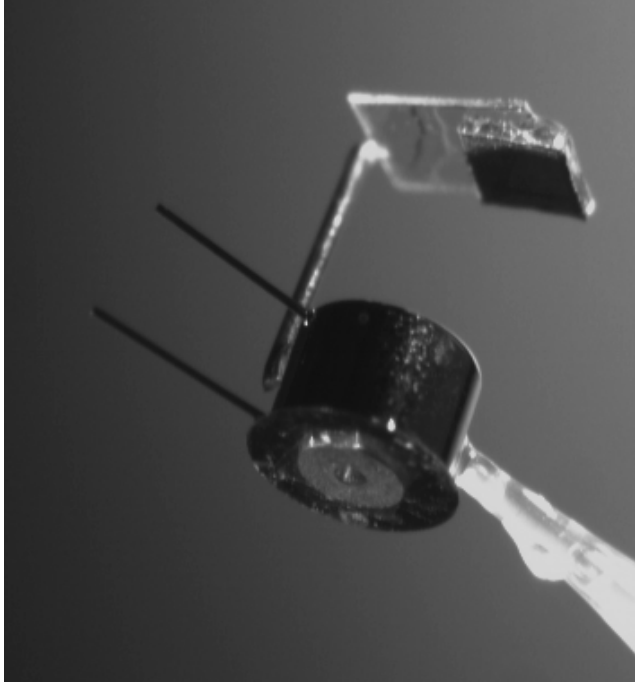
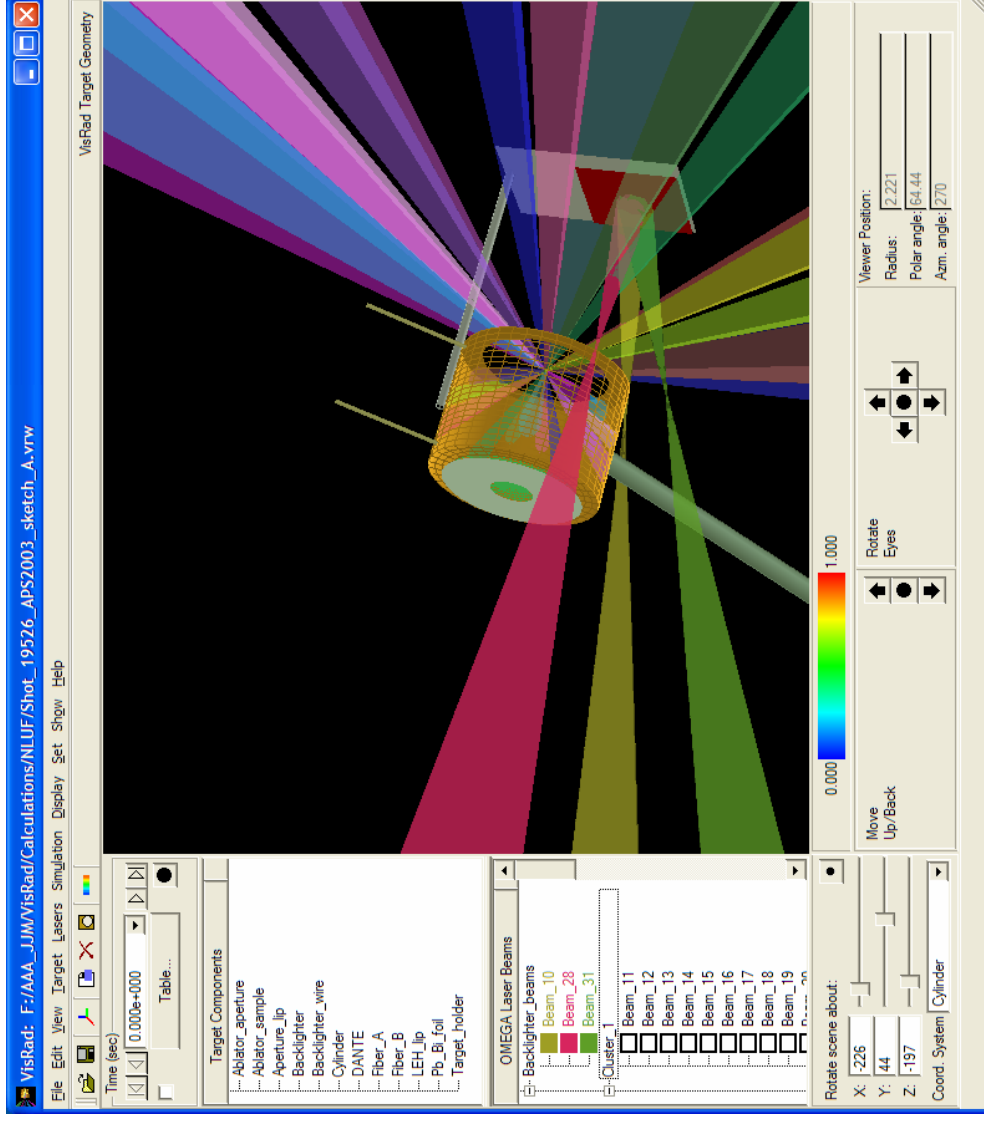
Modeling

Set up a halfraum target model in the *VisRad* view factor code



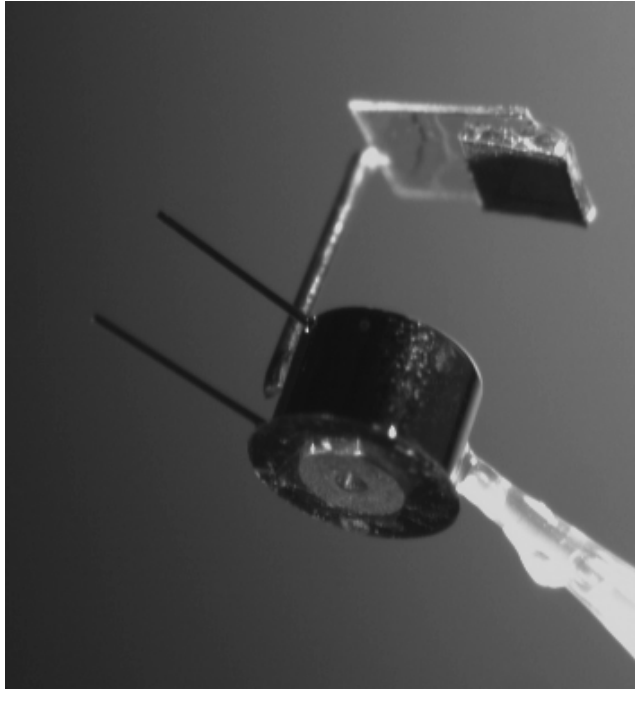
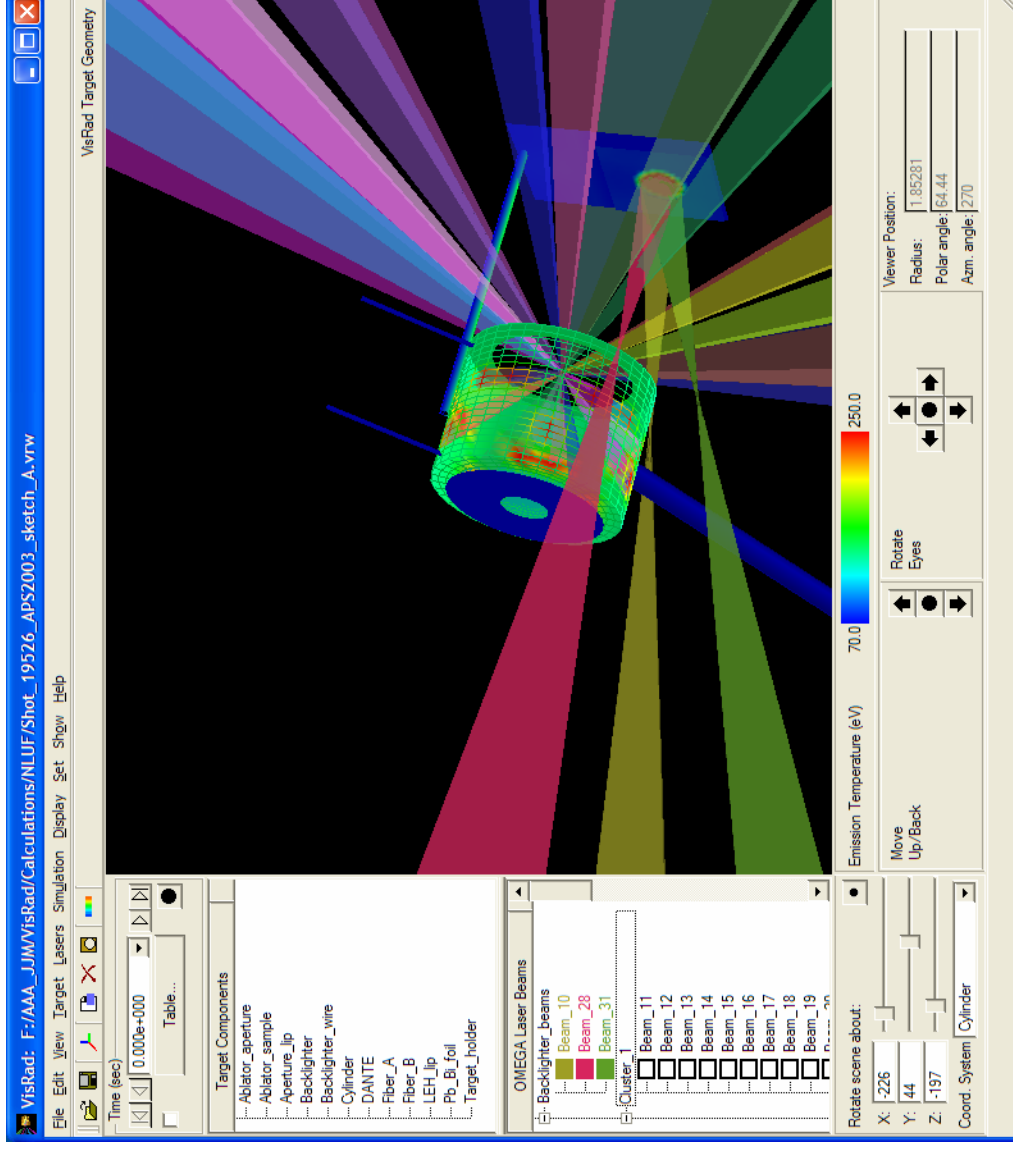
Specify material properties (albedos, laser reflectivities)

Put actual beam powers onto target (backlighter too)



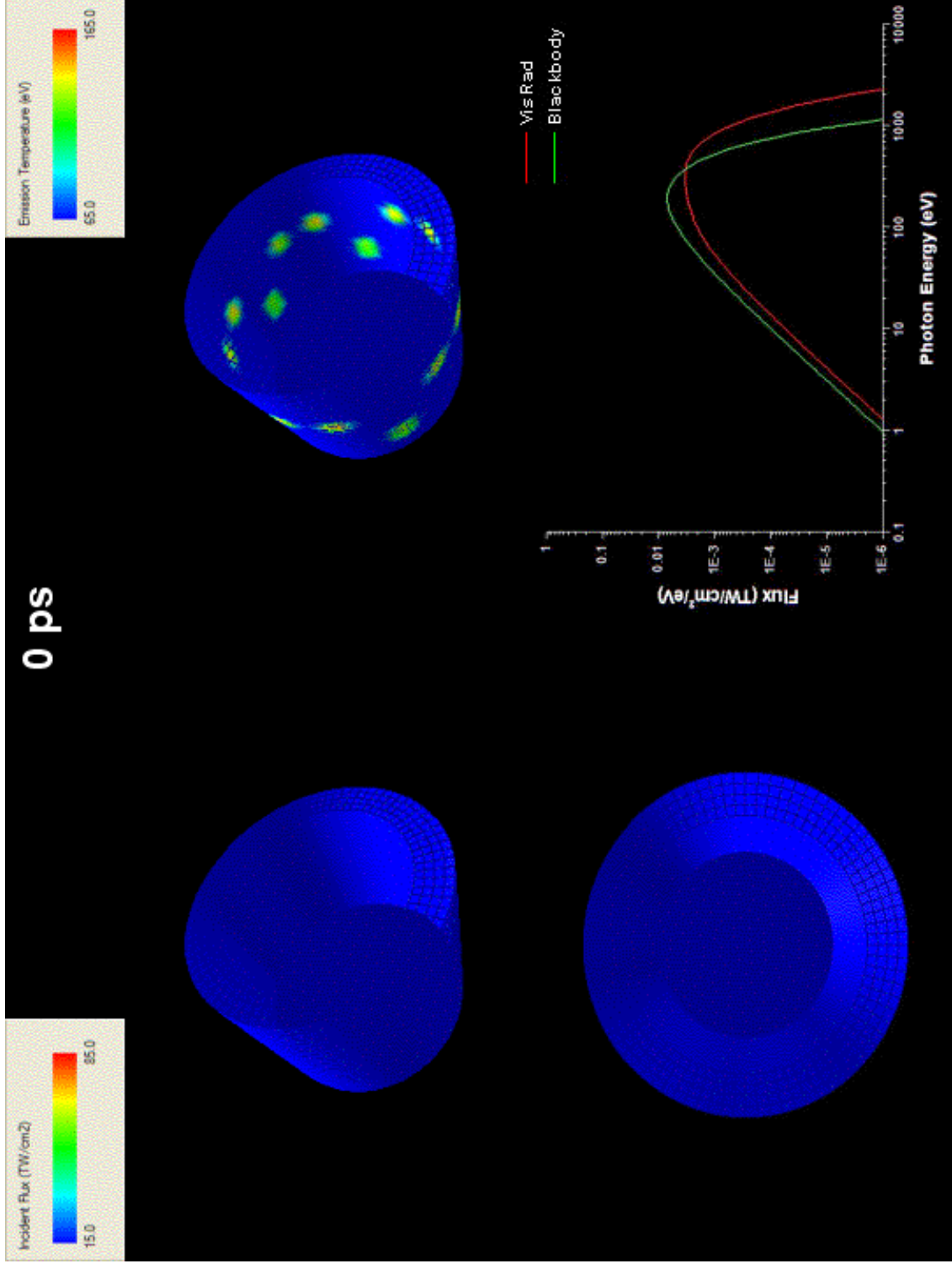
A series of time-independent calculations, with beam powers, albedos, XCEs varying at each time step

Calculate radiation field and material properties at every surface



Emission temperature shown here (note laser hot spots), but also calculate incident flux on ablator sample

VisRad simulations

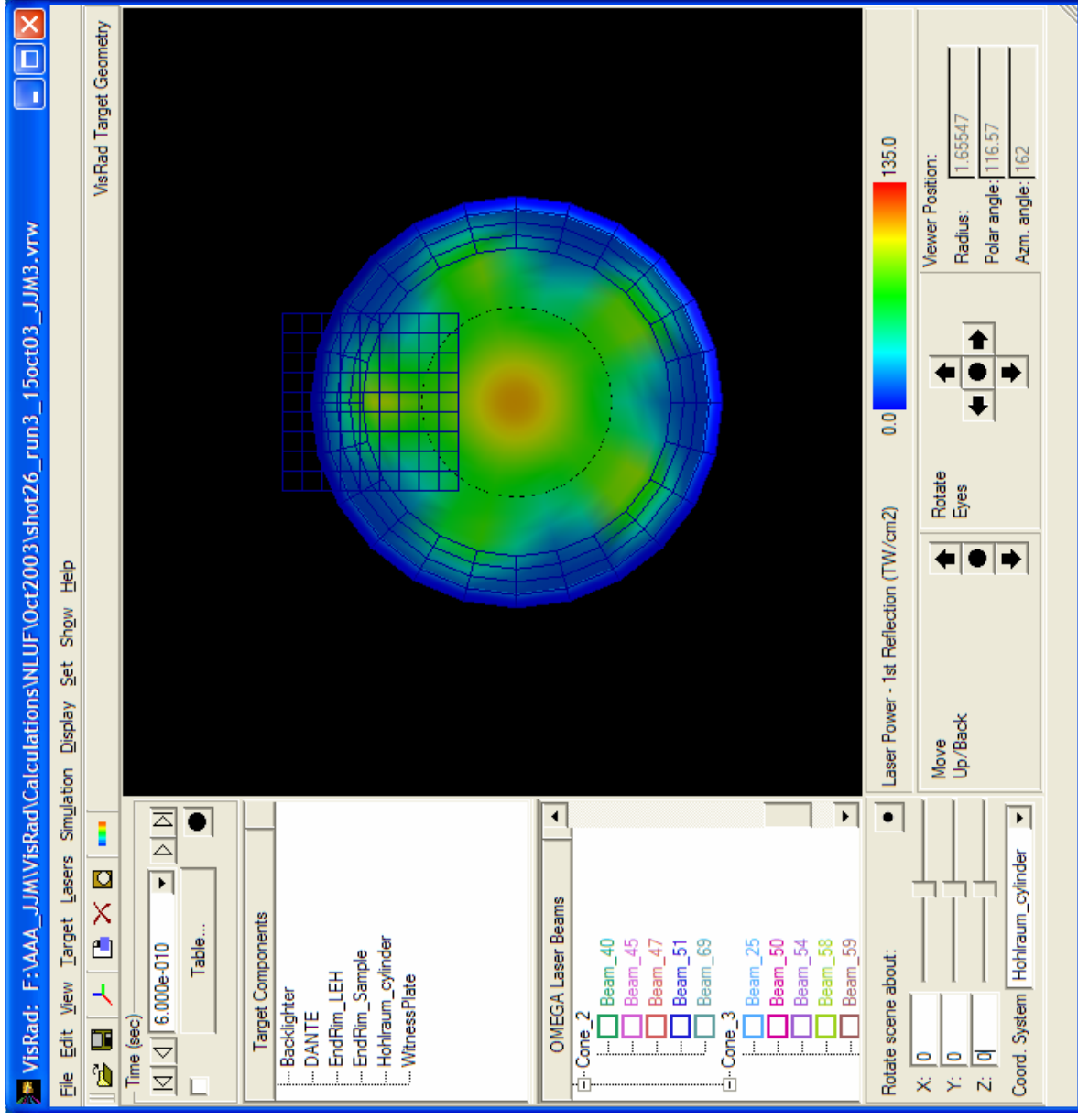


DANTE view on top, incident flux on left; spectrum
incident on ablator sample in lower right

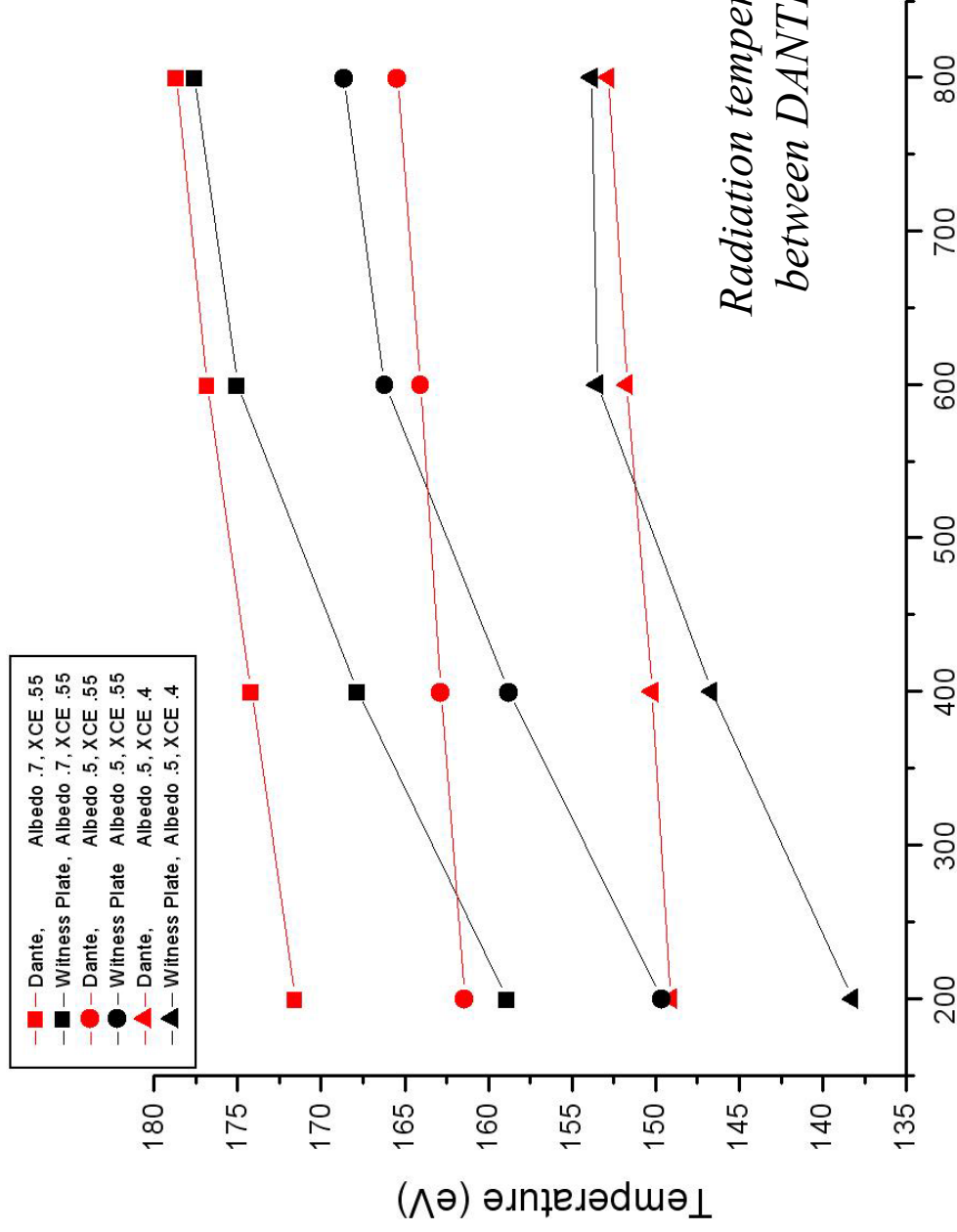
View-factor modeling is useful for
addressing various issues related to
hohlraum radiation conditions

Glint

Up to 135 TW/cm² of reflected laser light on to ablator sample.



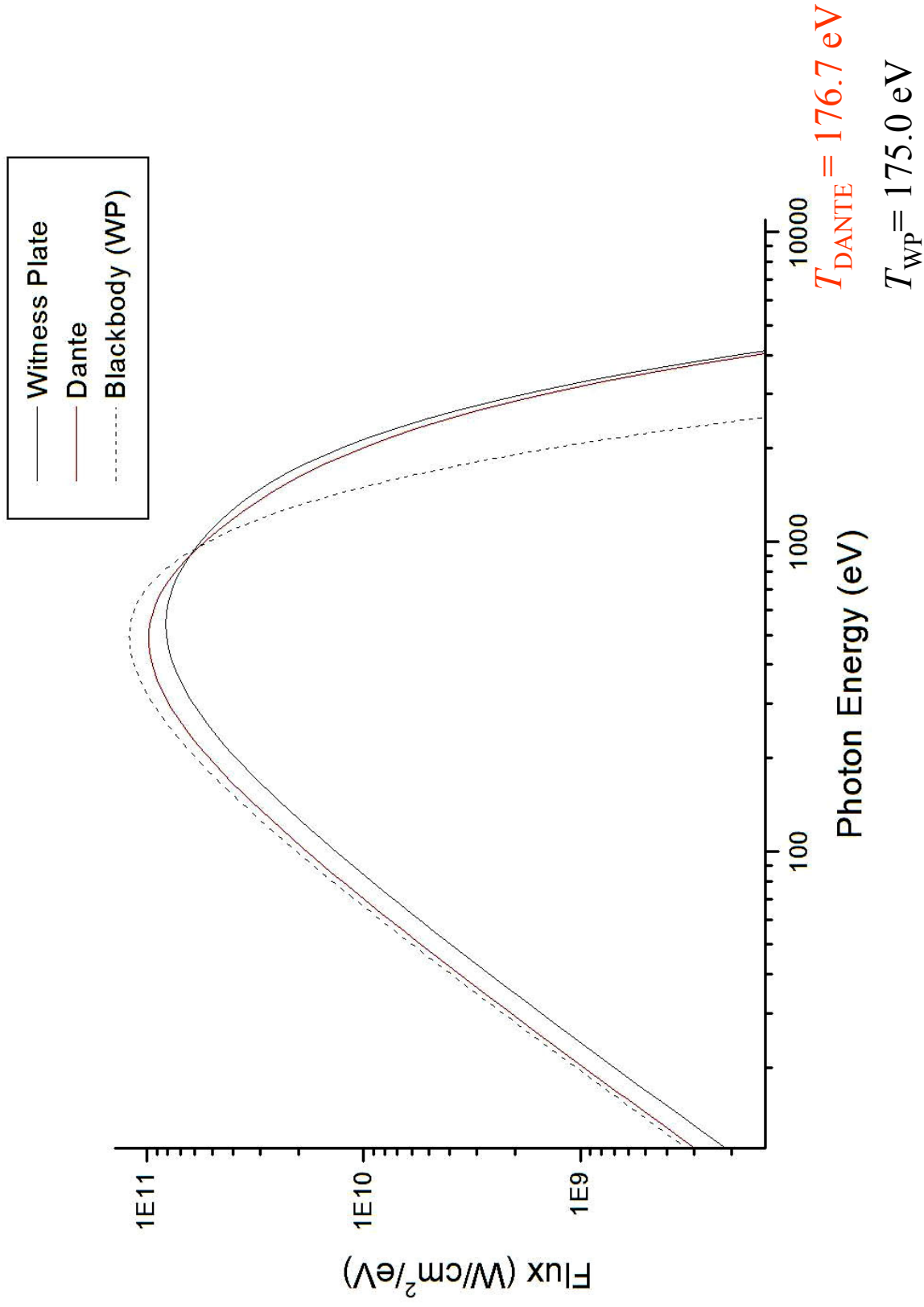
Radiation temperature as a function of beam pointing for three different albedo/XCE combinations, comparing DANTE (red) and WP (black) values.



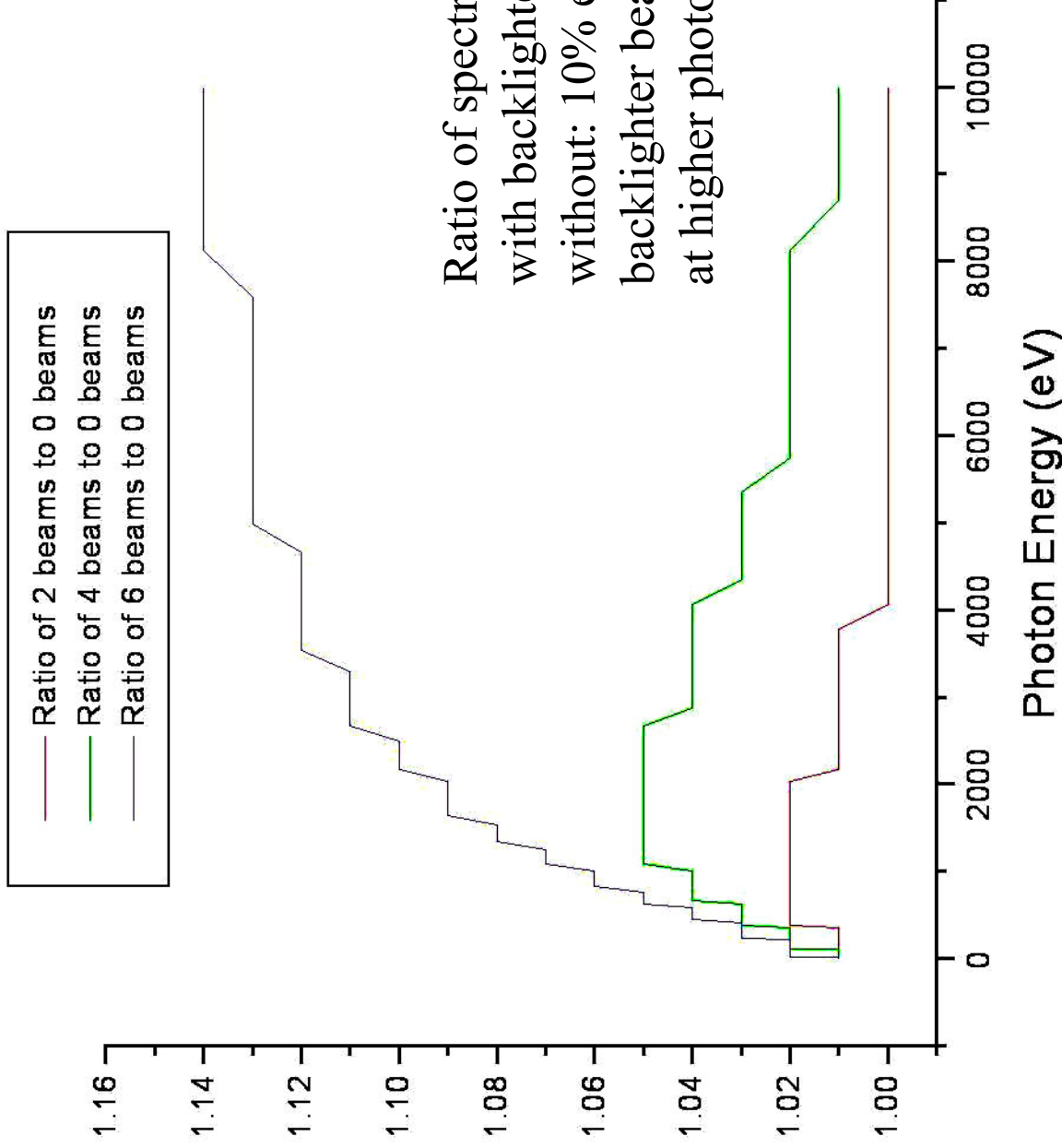
Radiation temperature differences between DANTE and sample can be significant.



Modeled spectra onto the sample and DANTE-view: even when the radiation temperatures are very similar, the spectral energy distributions are different (from each other and compared to a blackbody).

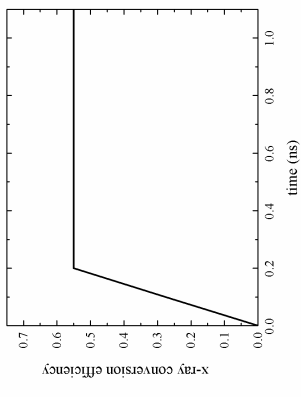
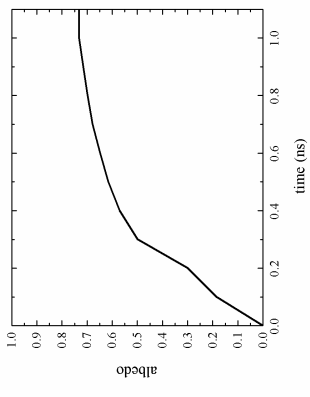
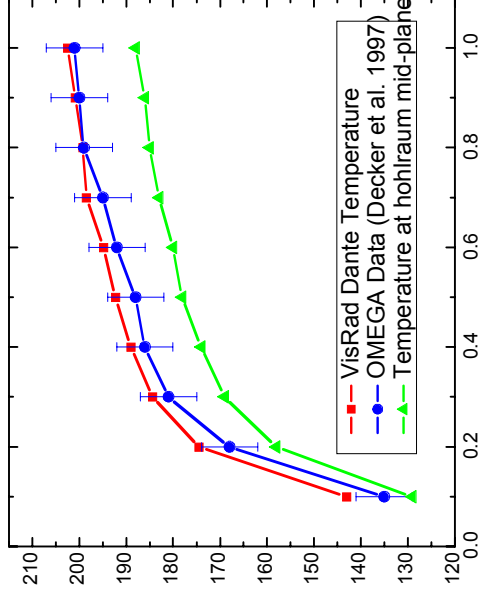
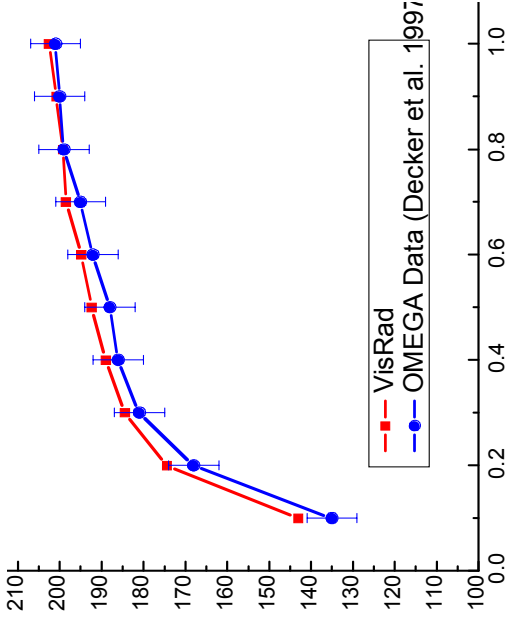
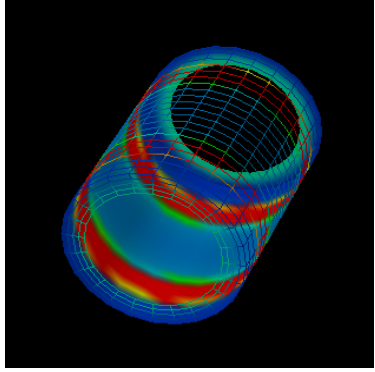
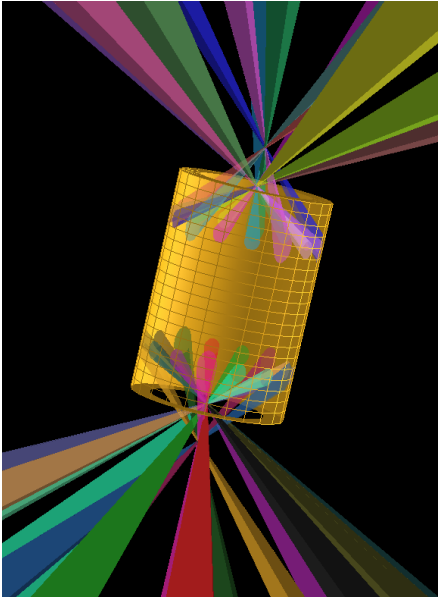


Effect of backlighter on radiation incident on ablator sample



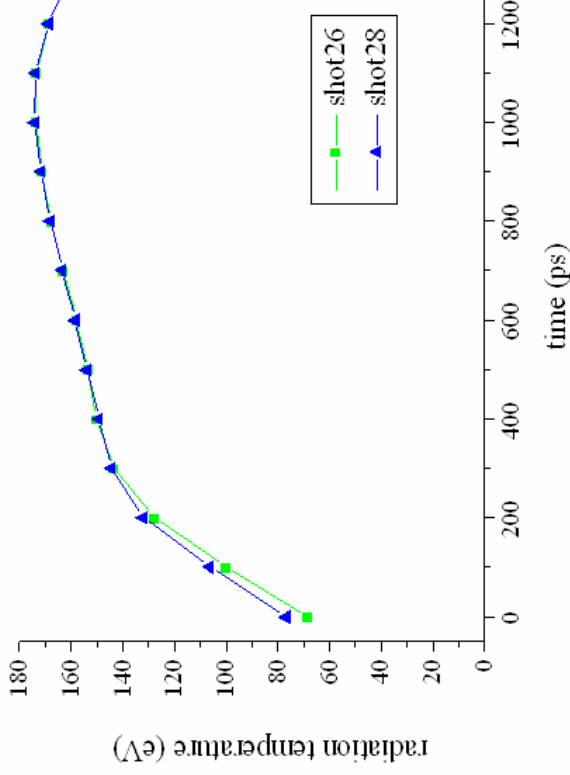
Ratio of spectrum onto ablator with backlighter included to that without: 10% effect with six backlighter beams; bigger effect at higher photon energies.

VisRad successfully modeled OMEGA hohlraum shots: DANTE data (Decker et al., PRL, 1997)



Assumed albedo (top) and XCE (bottom) vs. time

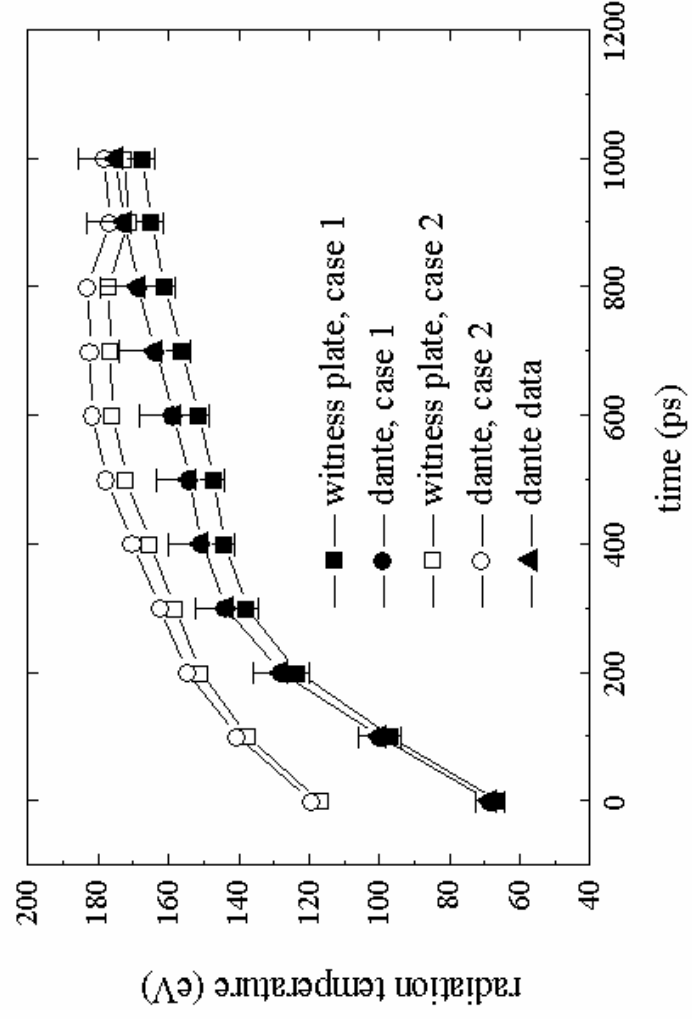
T_R at the midplane (green triangles) is different than wall re-emission seen by DANTE



But it is low (filled triangles, right) compared to our modeling (open symbols, right)

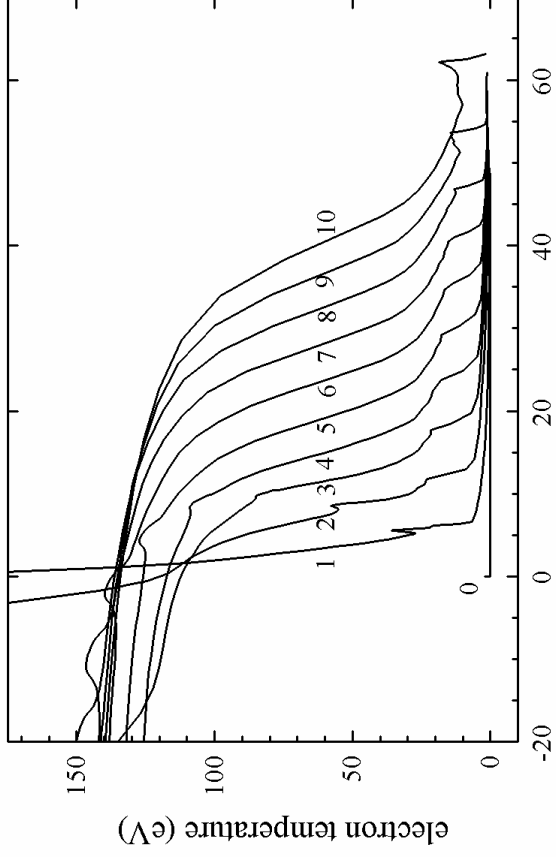
Note also: different T_R on ablator sample (squares) and seen by DANTE (circles)

DANTE data was almost identical for our two shots (left)



We are in the process of correcting for the reduced DANTE sensitivity at the time of our shots

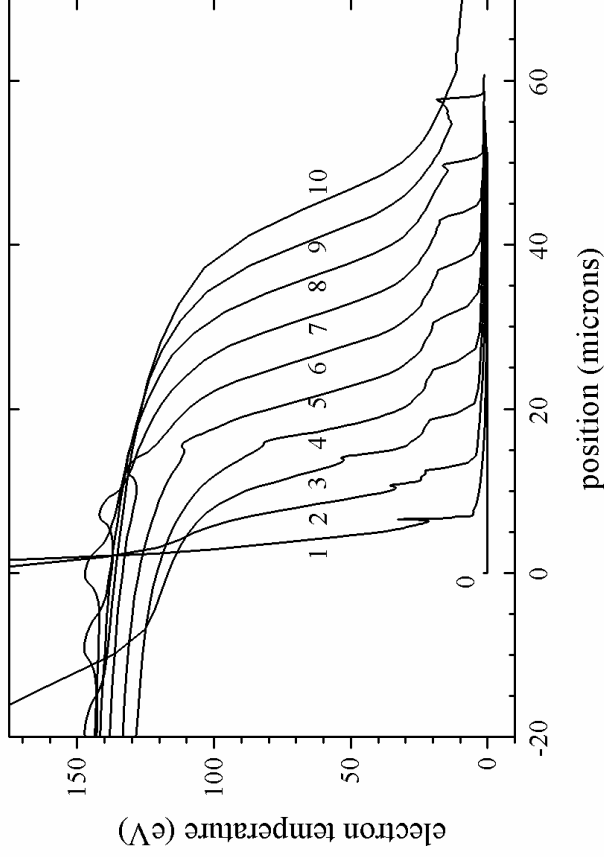
Times indicated in units of 100 ps



Undoped: faster
hydro evolution



HELIOS hydro modeling,
using incident flux from
view-factor modeling

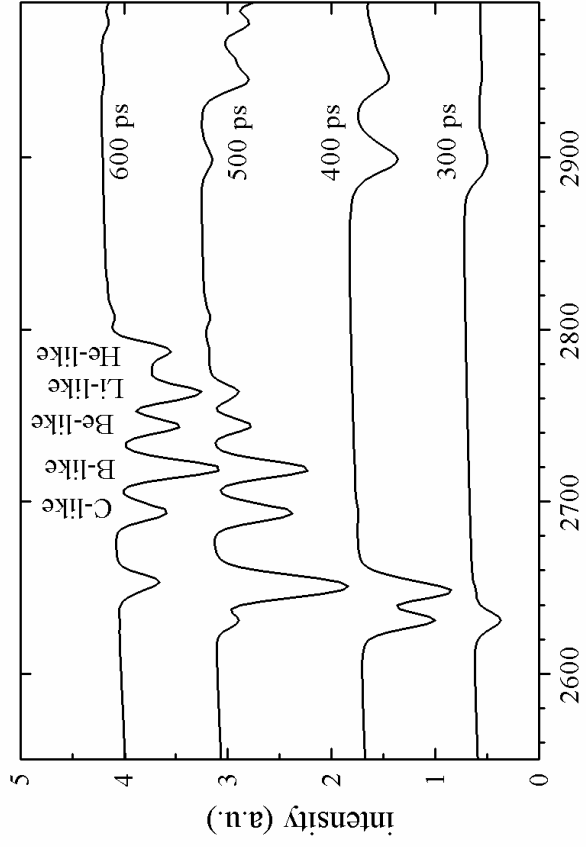


Doped: slower hydro
evolution

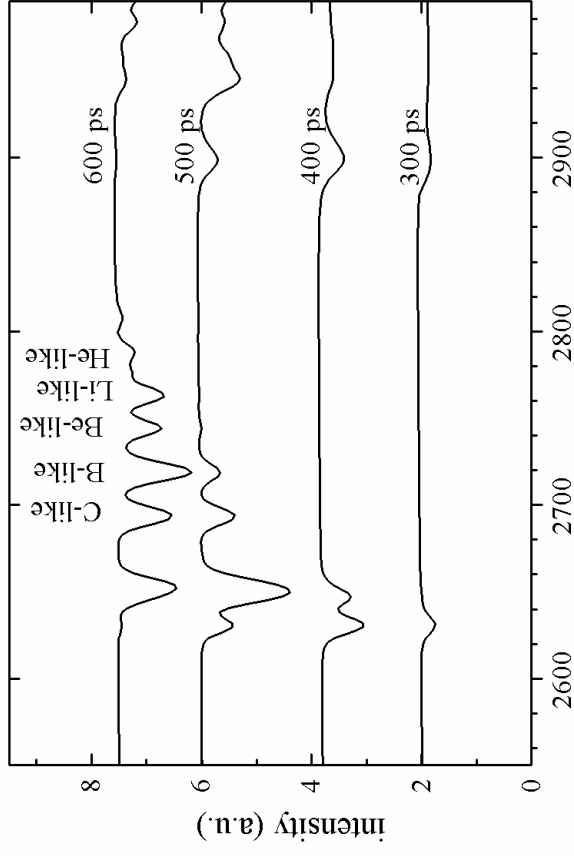


The hydro output is then post-processed with
Spect3d: statistical equilibrium and spectral
synthesis

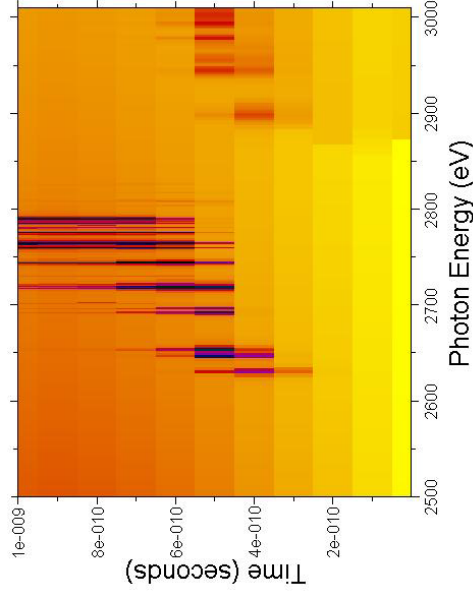
undoped



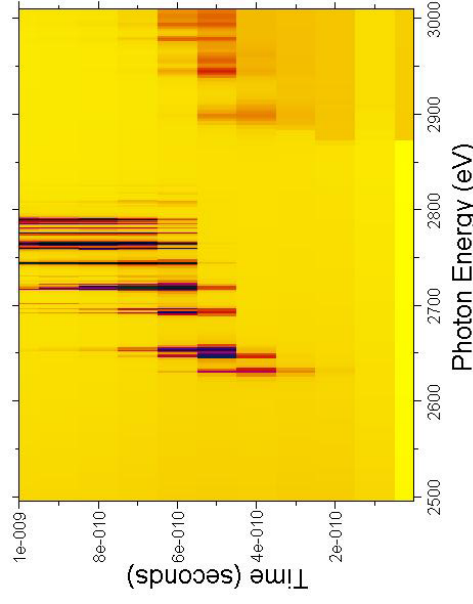
doped



photon energy (eV)

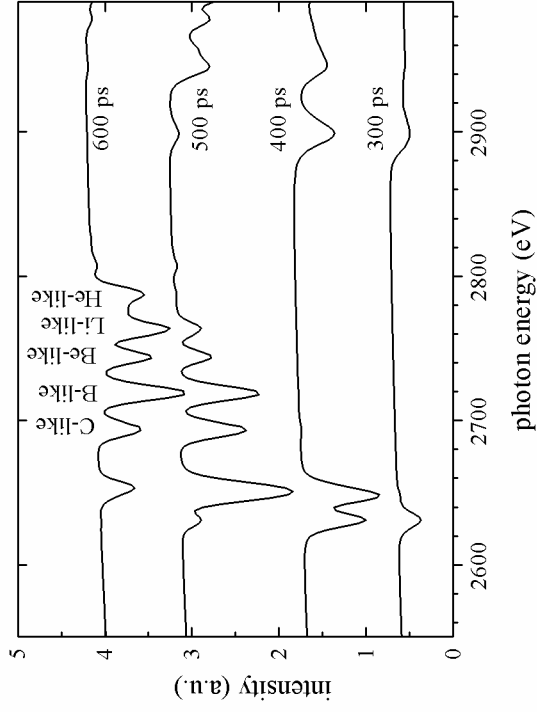


photon energy (eV)

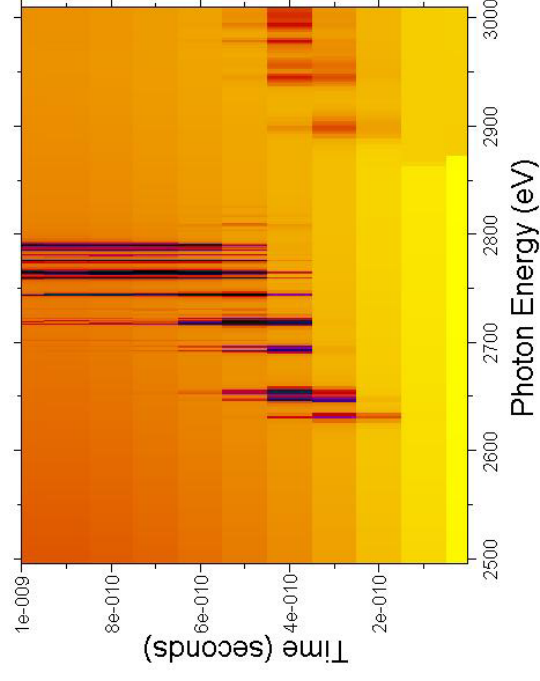
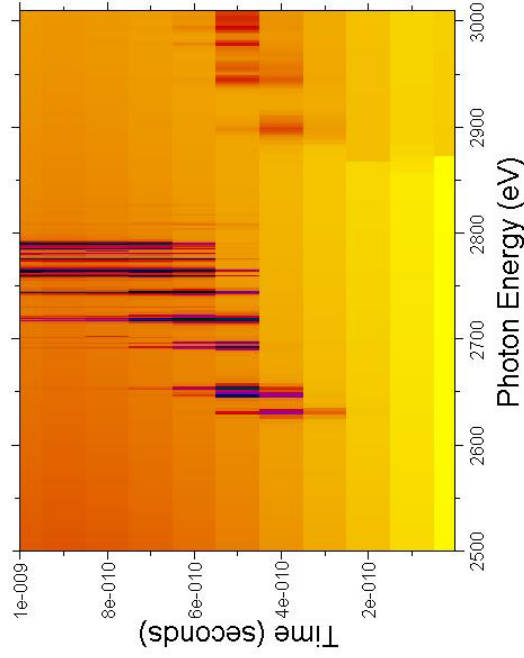
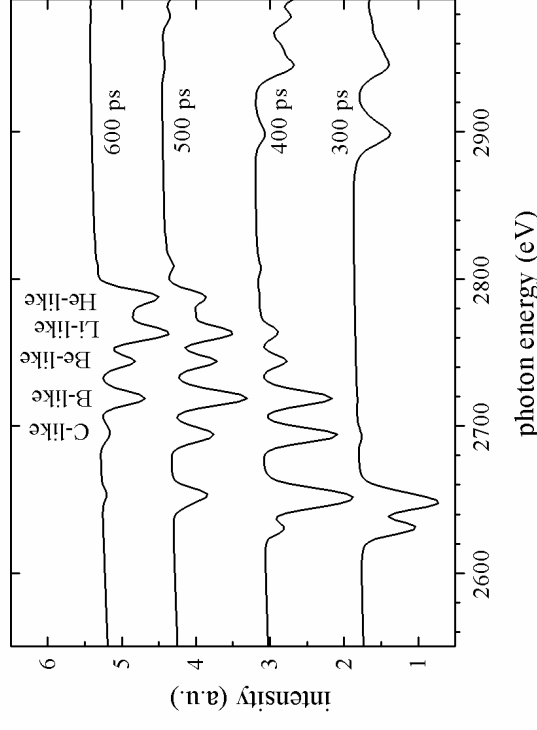


Tracer signals turn on earlier in the undoped sample
(as we saw before; when tracers in two samples are at the same depth)

Undoped: tracer at 6.3 μm

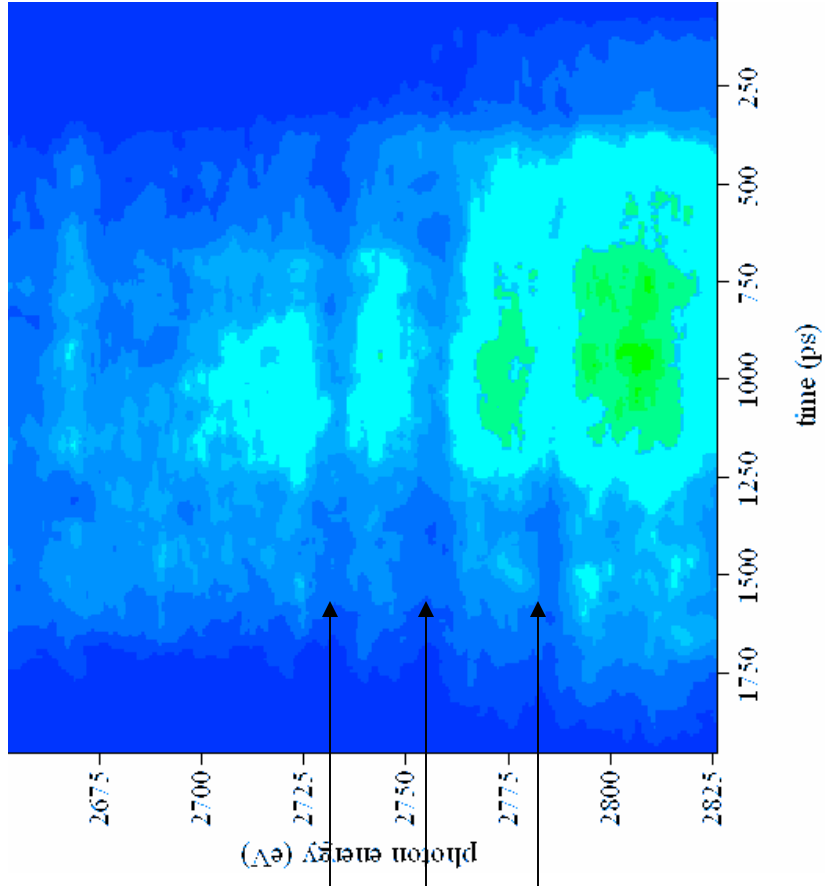
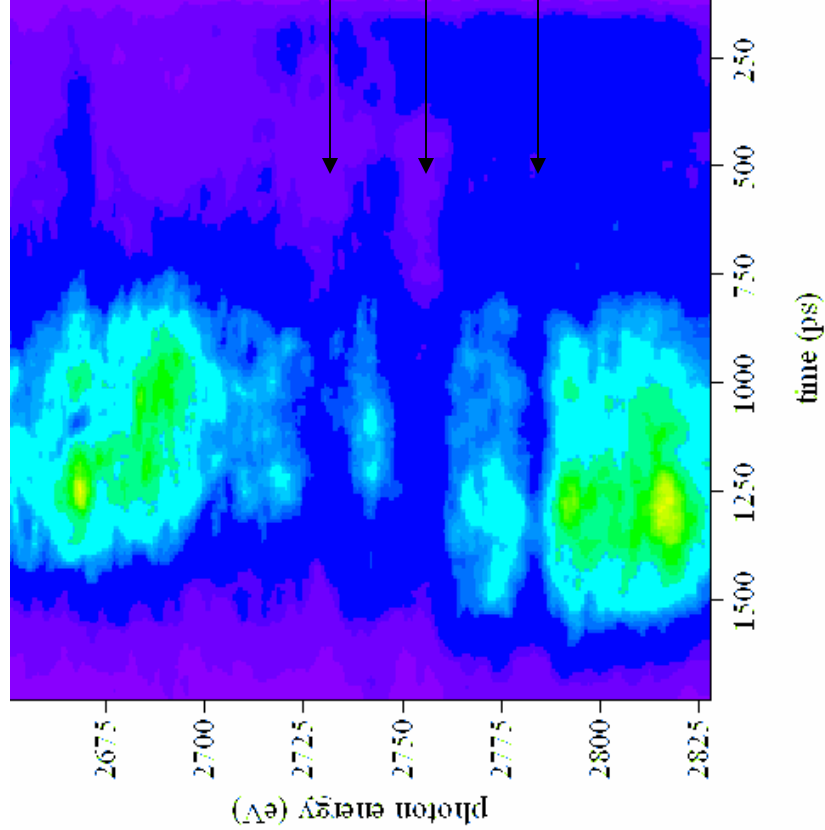


Doped: tracer at 4.1 μm



Shallower tracer in doped samples actually causes *earlier* turn on in this specific doped target

undoped



doped

Be

Li

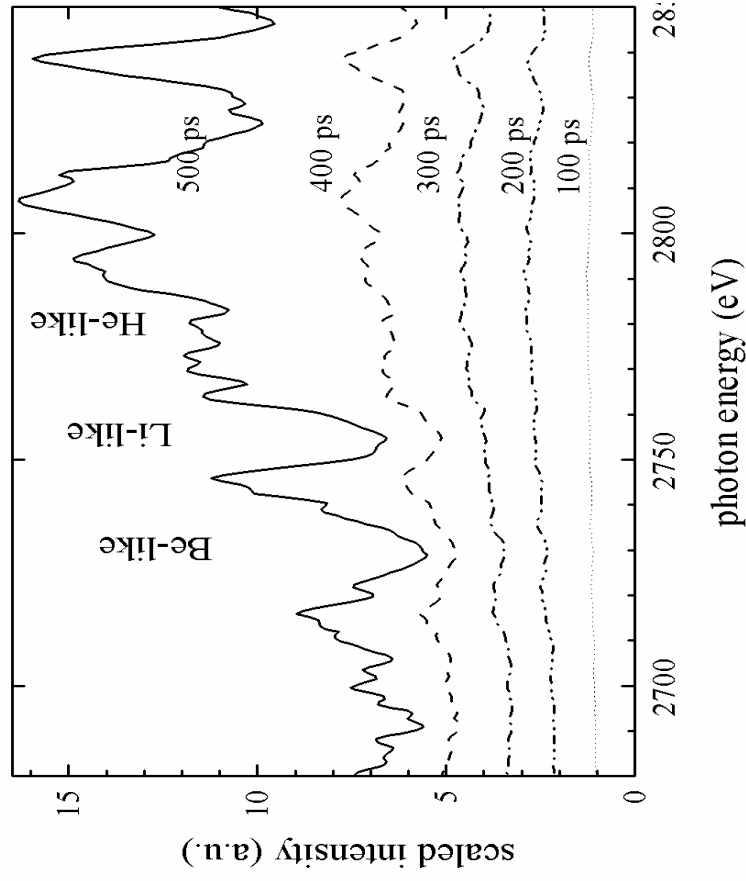
He

Back to the Data...

Data: time-resolved tracer spectra

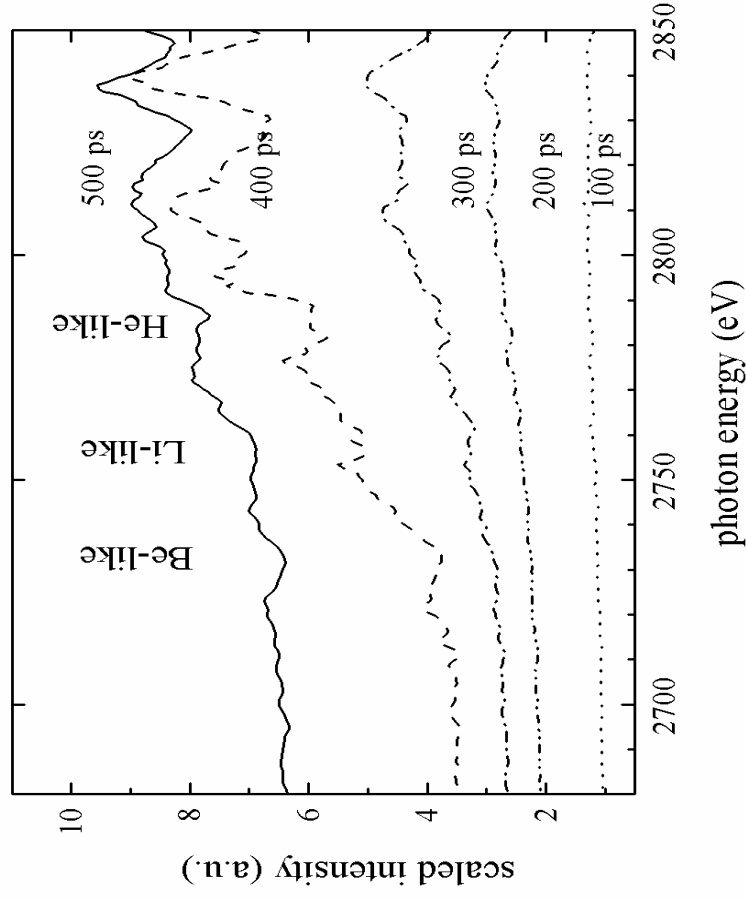
shot 19526: **undoped**

tracer at 6.3 μm

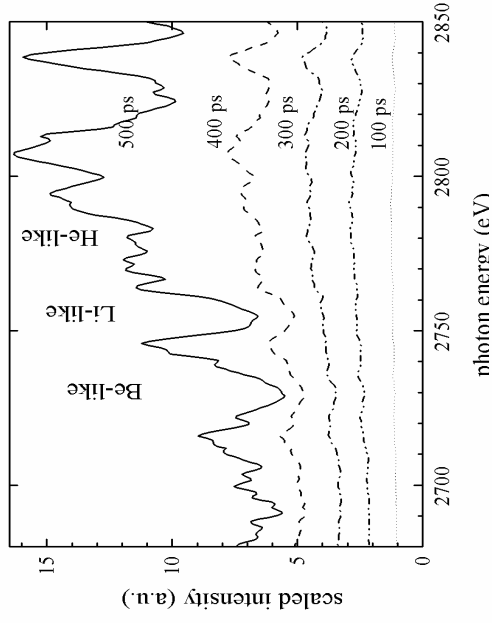


shot 19528: **doped**

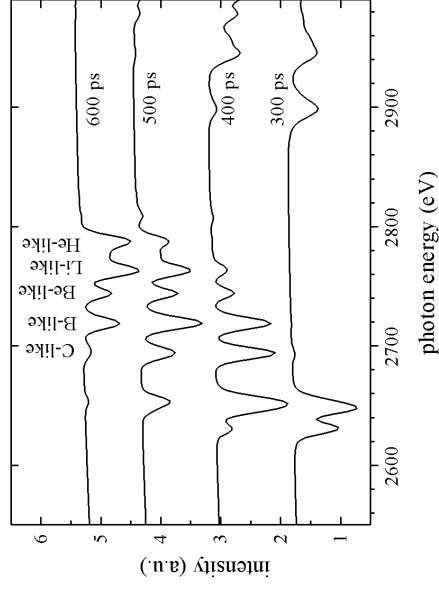
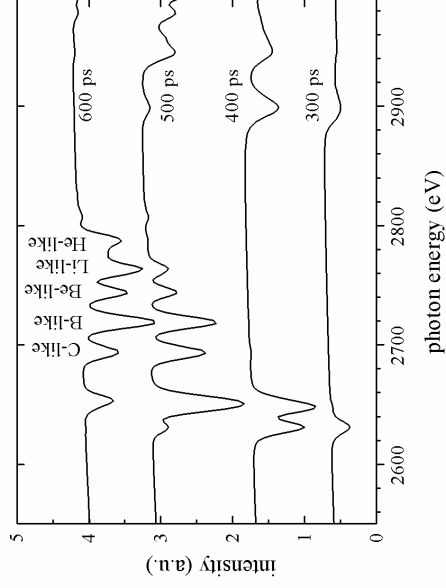
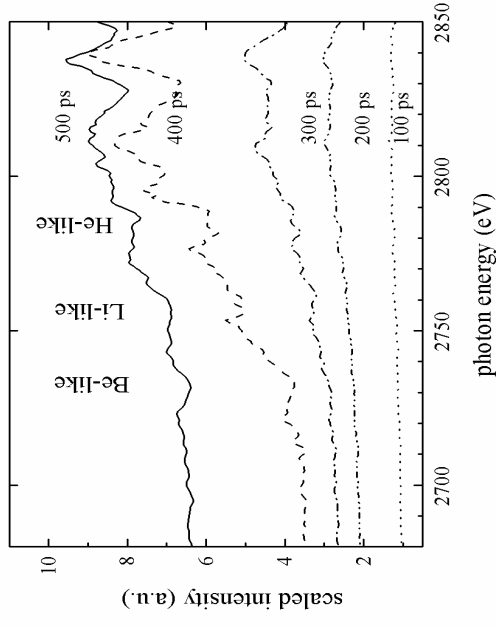
tracer at 4.1 μm



undoped



doped



Turn-on of Be-like: earlier in data
than model (300 vs 500 ps)

Turn-on time in better
agreement: 400 ps

Taking the turn-on times from the data, and the tracer depths:

The Marshak wave propagation:

2.5 mg cm⁻² ns⁻¹ for the undoped ablator

1.3 mg cm⁻² ns⁻¹ for the doped ablator

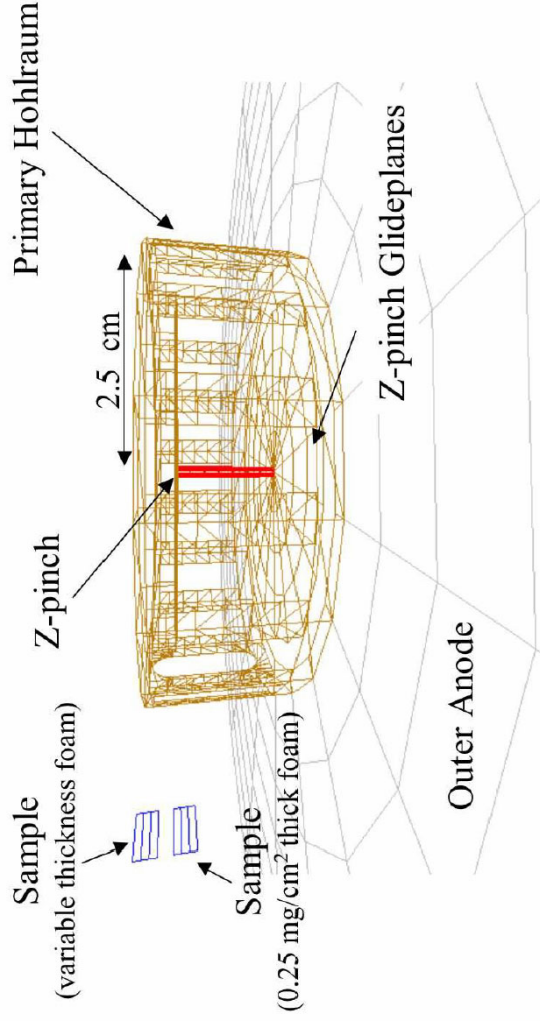
The dopant has a significant effect on the rad hydro

But, early turn-on time in undoped sample, relatively weak signal, inability to measure ion stages in the Cl tracer lower than Be-like...

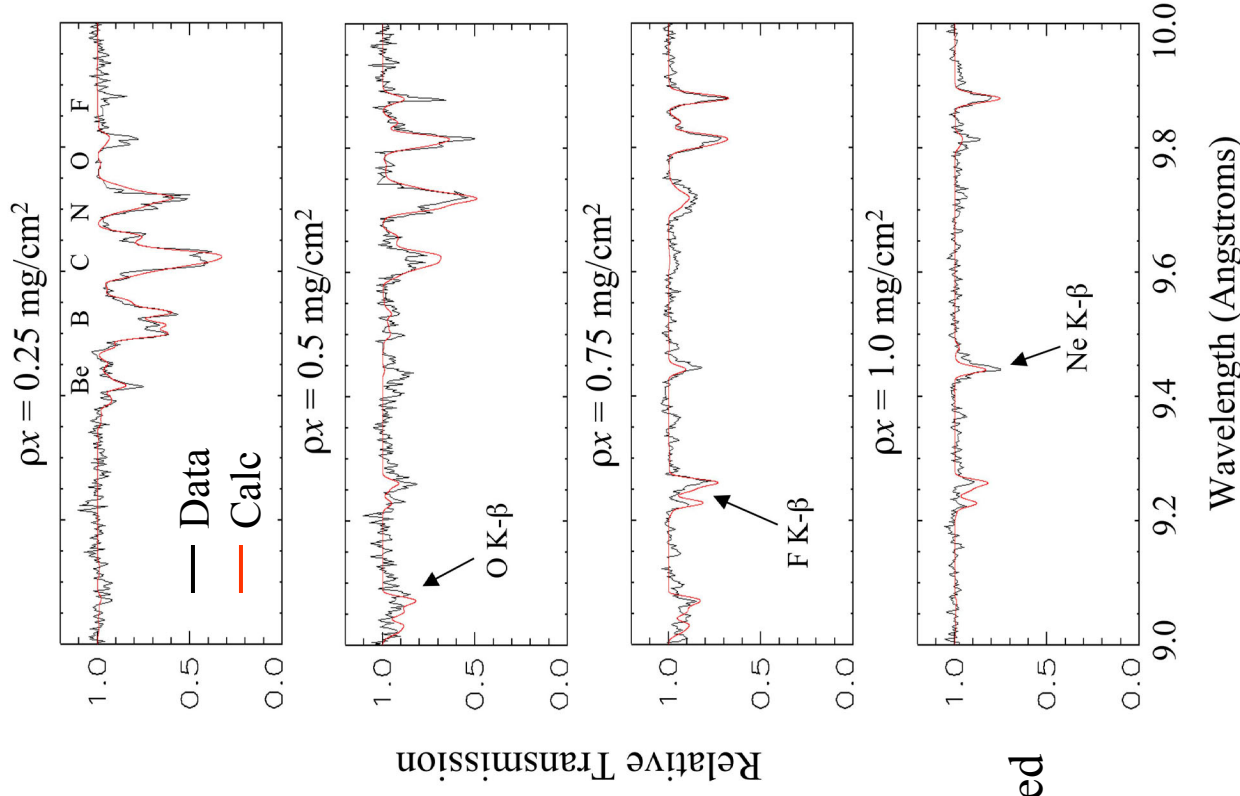
Similar tracer spectral diagnostics have been shown to be quite useful on Z-machine experiments involving foams

And the *Spect3d* spectral modeling has shown very good agreement with those data

Mg absorption spectra



- 5 mg/cc CH₂ foam samples
- Z-pinch heats and backlights samples
- 3 samples on 3 lines-of-sight
- MgF₂ tracers placed at 4 different depths
- Mg absorption spectrum recorded on time-integrated convex crystal spectrometer (R ~ 800)



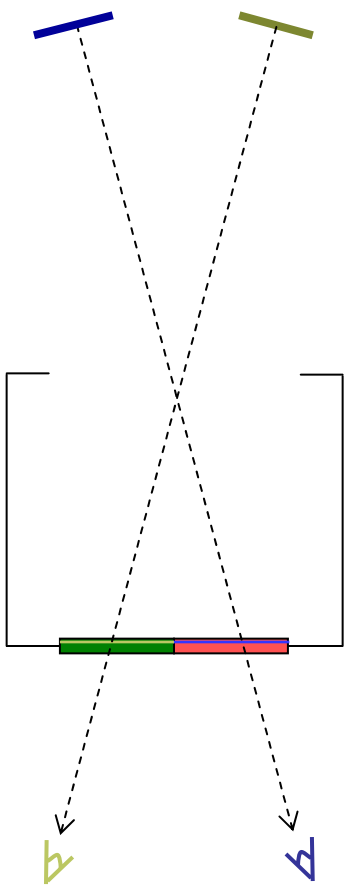
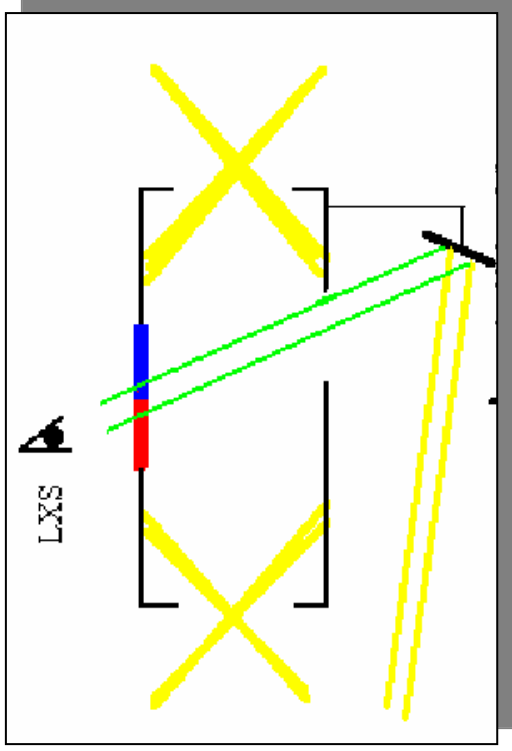
CONCLUSIONS

- Backlit tracer absorption spectroscopy is hard to accomplish cleanly in a hohlraum environment
- But, the tracer spectral data seem to indicate qualitatively the expected effect of the ablator dopant
- Modeling the hohlraum environment is important
- Future experiments might employ chlorinated plastic rather than salt as a tracer and a more evenly powered backlighter

Supplemental Slides

Our original plan was to make side-by-side measurements on doped and undoped ablaters mounted on the same hohlraum

Due to the complexity of these targets, we were not able to get good data



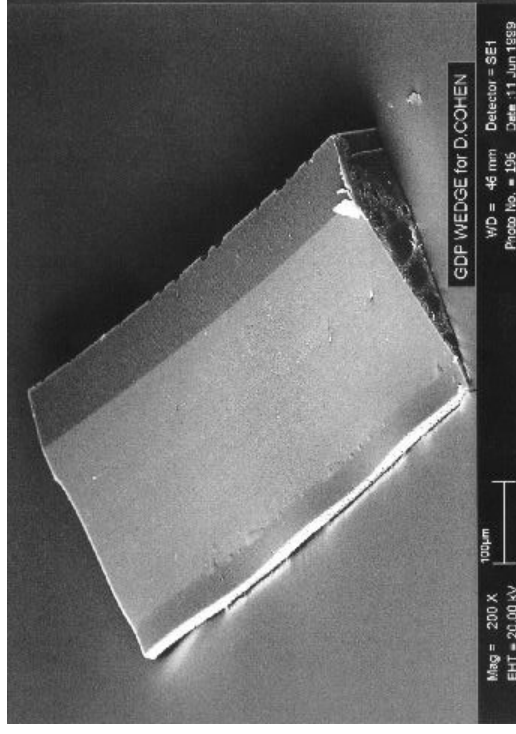
We mounted them first on the outside of hohlraums, near the midplane; one spectrometer with two separate crystals was used (K tracer in one side, Cl in the other)

A hohlraum with two samples (two tracers, two backlighters) and two spectrometers

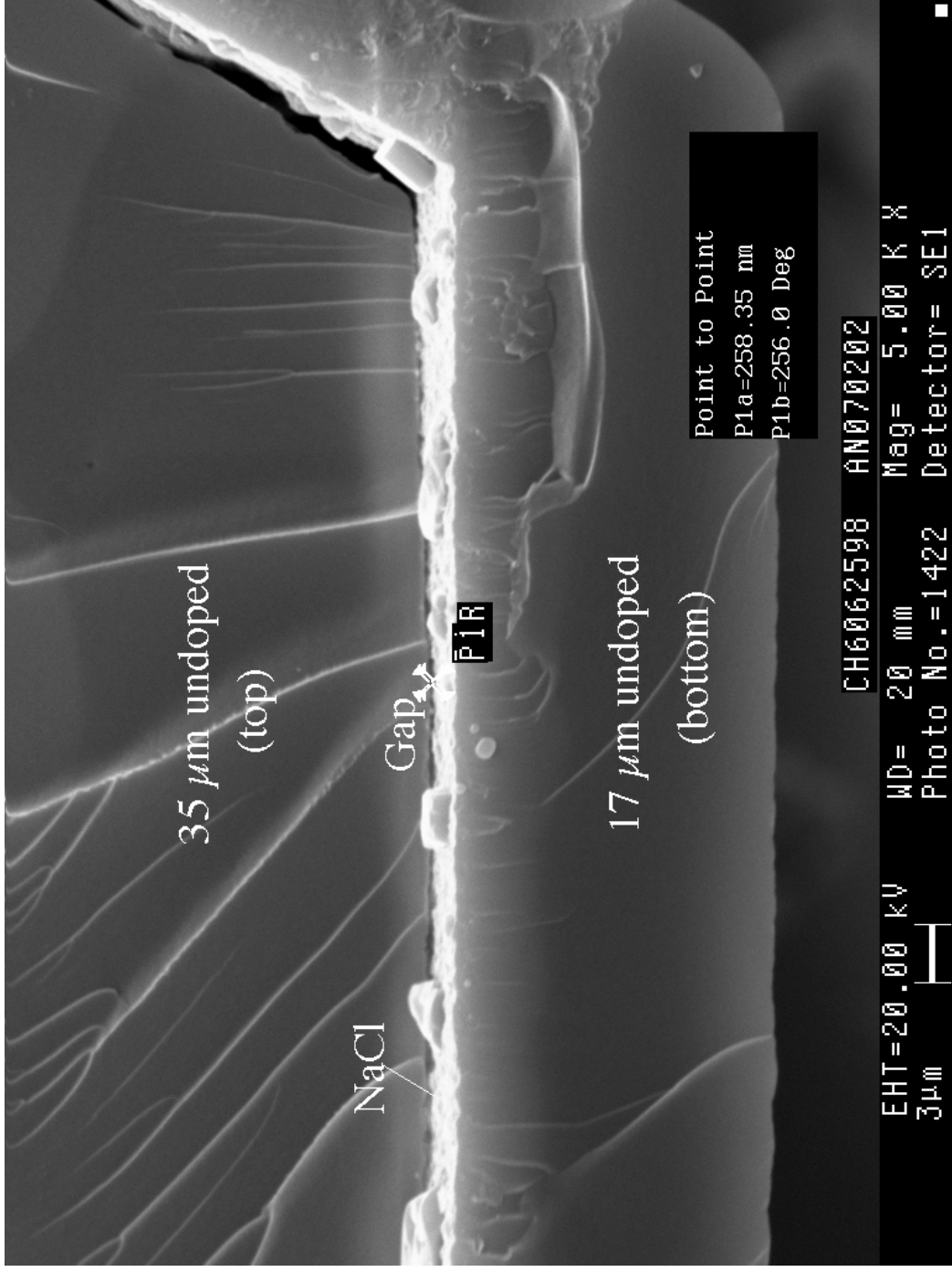
We never were able to successfully measure a tracer spectral signal on these experiments:

- Lower-than expected drive temperatures (tracers deeper than they ought to have been)
- “Cross-talk” between samples; emission seen by spectrometers not coming from line-of-sight through samples?
- Problems with one spectrometer

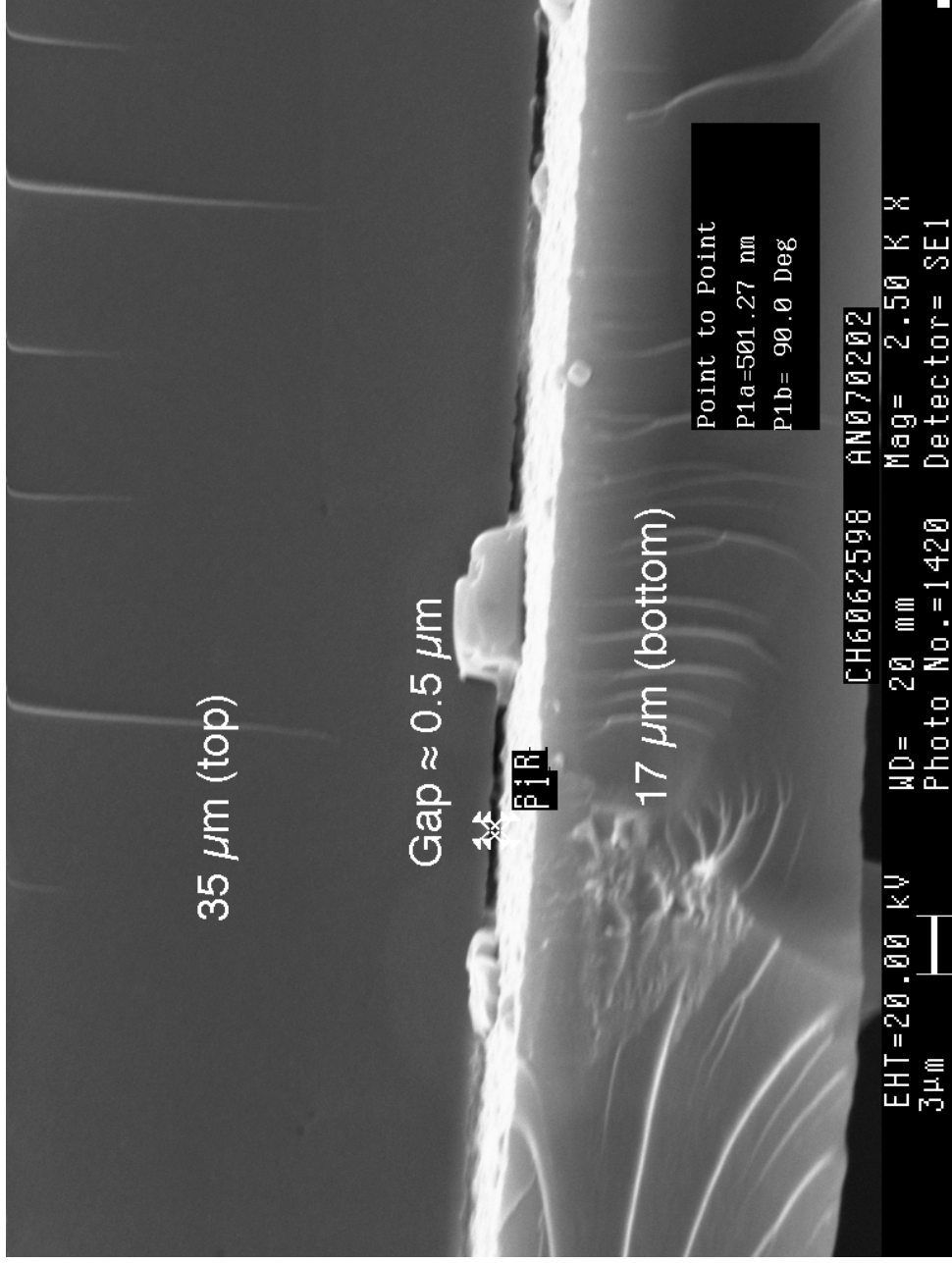
Other ambitious plans included use of wedge witness plates to make passive shock breakout measurements (*VISAR* with J. Oertel) simultaneously with tracer spectroscopy



higher magnification SEM image of ablator sample



And higher still...



If there are large gaps in the tracer layer, no amount of average areal mass will provide a strong signal.