Characterization of the x-ray emission from spherical shells for x-ray absorption spectroscopy experiments on OMEGA-60



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University of Rochester

D. A. Chin

We have characterized the performance of CH shell implosions, a common source for x-ray absorption spectroscopy (XAS) experiments*

- XAS provides the temperature, density and complex chemistry of the probed material
- Implosion core emission is an ideal x-ray source for XAS, because it is bright, broadband, short duration and small
- The corona and afterglow emission stages can account for 25% of the total x-ray emission and can impact the spectral resolution due to source size broadening
- Improved illumination strategies and inner metal layers increase x-ray
 emission from CH shell implosions



Collaborators



P.M. Nilson, D.T. Bishel, E. Smith, R.S. Craxton J.R. Rygg, G.W. Collins University of Rochester Laboratory for Laser Energetics

> J.J. Ruby, F. Coppari, A. Coleman and Y. Ping Lawrence Livermore National Laboratory

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Spectral features near an x-ray absorption edge can be used to deduce the electronic density of states and atomic structure of matter



E29799



International X-ray Absorption Society, Fe2O3 Data, https://xaslib.xrayabsorption.org M. Newville, Rev. Min. Geo. <u>78</u>, 33 (2014).

Precision XAS requires bright, broadband, short duration and small x-ray sources



- Multiple different implosion shells have been studied to determine the optimal target design^{*,**,†}
- Recent XAS experiments used CH shells with a vacuum fill[‡]



CH Shell

This work will focus on 9 μ m thick CH shells with an 865 μ m outer diameter and a vacuum fill



The corona and afterglow emission stages can be sizeable contributors (~25%) to the x-ray emission from an implosion





Time resolved imaging and spectroscopy measured the x-ray emission stages





Lilac simulations were used to construct models for each of the three x-ray emission stages





Models were applied to each stage and fit to the data to characterize the fraction of the signal in each stage



- An 8 parameter model was developed to characterize the fraction of the signal in each stage
- The model was verified using the hydrodynamic simulation and then fit to the data

	Corona	Core	Afterglow
Signal Fraction	$12^{+2}_{-2}\%$	76 ⁺⁷ ₋₉ %	$12^{+7}_{-6}\%$





The source broadening from each stage was calculated and used to simulate the absorption spectrum of iron

A point source spectrum was degraded using the spectral resolution corresponding to each stage[†]



The additional broadening terms prevent the ability to use XANES to determine the iron melting*

> The additional broadening terms decrease the Debye-Waller Factor^{**} by 5 – 10%



By repointing the drive beams to more symmetrically illuminate the shell, we increased the symmetry of the core stagnation

- In XAS experiments, five beams are used to drive the XAS target while the remaining beams illuminate the CH shell
- The drive beams were repointed in configuration 2 resulting in a 4x decrease in the variation of the absorbed laser intensity





An improved target illumination strategy resulted in a factor of two increase in the x-ray emission

The target illumination strategy was improved by changing the laser phase plates and repointing the target drive beams





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Preliminary results indicate metal layers can further increase the total x-ray emission





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Future experiments are planned to characterize the spatial and temporal x-ray emission profile of the metal layered shells for XAS experimental use



Backup





An 8 parameter model was used to fit the x-ray emission and constrain the three emission stages

Corona Emission:

$$J_{corona}(t; A_1, B_1) = \frac{A_1}{\sqrt{\tau}} e^{-\frac{1}{\tau}} \text{ and } \tau = B_1 \left(\frac{p(t)}{r_{crit}(t)}\right)^{\frac{2}{3}}$$

• Core Emission:

$$J_{core}(t; A_2, \mu_2, \sigma_2) = A_2 e^{-\frac{1}{2} \left(\frac{(t-\mu_2)}{\sigma_2}\right)^2}$$

Afterglow Emission:

$$J_{afterglow}(t; A_3, v_3, t_3) = \begin{cases} 0 & t < t_3 \\ A_3 \eta_e^2 & t \ge t_3 \end{cases} \text{ and } \eta_e = \frac{1}{\left(\frac{50 + v_3(t - t_3)^2}{2}\right)^3}$$

• Total Emission:

 $J_{model} = J_{corona} + J_{core} + J_{afterglow}$



