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SOFT X-RAY SPECTROSCOPIC MEASUREMENTS
OF PLASMA CONDITIONS AT EARLY TIMES
IN ICF EXPERIMENTS ON OMEGA

Final Technical Report
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Final Technical Report

SOFT X-RAY SPECTROSCOPIC MEASUREMENTS OF PLASMA CONDITIONS AT EARLY TIMES IN ICF EXPERIMENTS ON OMEGA

During this reporting period, we first analyzed the data obtained during our final campaign under this grant on the OMEGA laser facility, which took place on April 27, 1999 (one day). As explained in our previous semi-annual report, for the first time we fielded our Ten Inch Manipulator- (TIM-) mounted flat-field, grazing-incidence extreme-ultraviolet (euv) spectrograph [1] with a four-channel gated-stripline microchannel plate (MCP) detector. This spectrograph covers the spectral range of 30-250 Å ($h\nu=50-400$ eV). This instrument reached closer to the target than did the previous version mounted on the chamber wall; such that the sensitivity increased by at least a factor-of-10 for viewing weak spectral features.

During this April 1999 campaign all except one of the 60 beams were used, with a total on-target energy of nominally 20 kJ in a 1 ns pulse. The beams were focused onto a spherical target of overall diameter 940 μm , for an irradiance of $\sim 2 \times 10^{14}$ W/cm². Beam smoothing was provided by a combination of spectral dispersion (SSD) as well as distributed phase plates (DPP's). The plastic microballoon targets of 20- μm wall thickness were filled with neon to a pressure of 10 atm, including a 10% admixture of argon. A sealant coating of aluminum of thickness 0.0125 μm (125 Å) was followed by an outer coating of Mg of thickness 2 μm . Also, in this campaign, some outer coatings of LiF were used in place of Mg.

In the April 1999 campaign, we were able to track F VII, VIII and IX as well as, at higher temperatures, Mg X, XI and XII spectral lines from $n=3$ to $n=2$ transitions in Li-, He- and H-like ions, beginning at very early times in the event (prepulse) and continuing through the heating cycle, after which continuum emission dominated the spectra during compression. Also, Li III lines were measured.

During this final-report period, we also completed the analysis of our prepulse measurements which began with the May 1998 campaign. The results were presented at the November 1999 meeting of the American Physical Society Division of Plasma Physics in Seattle [2] and have been submitted for publication [3] (a recent preprint is included as Attachment A here). The prepulse measurements agree with modeling carried out at LLE by Dr. Jacques Delettrez, and by Dr. J. Dahlburg and colleagues at the Naval Research Laboratory and represent a prime example of code verification by experimentation. We were not able to continue to monitor the prepulse after

April 1999 since our grant was not renewed for FY-2000. This is unfortunate, since it is an event that should be monitored regularly, after it appeared suddenly during the May 1998 campaign and varied in significance later, never disappearing.

Particularly interesting in the prepulse data obtained in April 1999 were aluminum absorption spectra in the 40-120 Å wavelength region, as shown in Fig. 5 of Attachment A. The strongest and broadest absorption feature (at a wavelength centered near 110 Å) matches a blend of the energies [4] for photoionization from the 2s innershell levels in neutral aluminum as well as in Al^{1+} and Al^{2+} ions. Also, photoionization edges [5] associated with valence electrons in Al^{3+} to at least Al^{5+} match other narrower absorption features in the spectra, as indicated. One possible source for the continuum emission in which the absorption occurs is the surface of the CH microballoon (the balloon wall as well as the gaseous fill are opaque in this region, ruling out the possibility of a rear-target-surface backlighting). The 0.0125-μm thick solid aluminum sealant layer is semi-transparent (~25%) to the 351-nm wavelength laser light [6], while also being semi-opaque (~40% absorption) at the x-ray edges [7]. As such, laser preheater light can be transmitted through the thin aluminum layer to the outer surface of the polymer target, generating a continuum, and forming the 2s absorption feature in the cold aluminum layer. This holds similarly for n=2 absorption in an aluminum plasma that might be formed, such that the product of density and thickness remains constant over a limited range. Also, the absorption features from multiple ionization stages could be arising from different points on the target, and perhaps from varying thicknesses of the aluminum layer, if it is not completely uniform. An alternative explanation for the absorption observed is that it takes place along chords in the direction of view, tangential to the surface of the microballoon, i.e., in a "limb" absorption process. This provides a longer path length for euv emission, perhaps up to 100's of μm's instead of 20 μm for direct view. The backlighting could arise from beams focused to the rear of the target which may have prepulses included. This latter explanation would be consistent with later carbon spectra to be described next, both of which require space-resolved spectra to definitively interpret. Clearly, an identification of the local region of absorption and the backlighter is vital.

Another observation which may bear on such limb absorption is observed at approximately 8 to 10 ns after the onset of the main laser, i.e., late in the event. Time-gated, spatially-integrated (along the axis of the slit of the spectrograph) recombination spectra show a rich series of spectral lines from n→2 "Balmer" series transitions in C VI in the 106-182 Å region, and similarly in C V and O VIII. The series extends to n=9 in some cases before merging at n=10 into a distinct continuum. For C VI this indicates an electron density of $\sim 10^{18} \text{ cm}^{-3}$ using the Inglis-Teller relation [8,9:p. 148] for the merging of high-lying lines

with the continuum. Another observation is that the $n=3-2$ and $n=4-2$ transitions are unusually intense compared to those for $n>5$.

Fortunately we also were able to record the $n\rightarrow 1$ series of C VI (and C V) on the same spectra but in the 27-34 Å region, i.e., at the same time. However, we noticed that the series are much shorter there, ending with $n=4$ followed by a continuum step at $n=5$. This corresponds for to an electron density of $\sim 10^{21} \text{ cm}^{-3}$, i.e. much higher than that deduced from the $n\rightarrow 2$ series. None of the data we obtained with the same spectrograph using slab targets displayed such a difference in series limits and, hence, electron densities.

We can begin to understand this apparent discrepancy if we imagine that we are viewing two concentric spherical plasmas, one at the core emitting short-series (both) spectra at high density and the other in an expanding shell emitting a superimposed longer-series spectra at lower density. The $n\rightarrow 1$ x-ray lines viewed through an extended plasma chord of a spherical shell could be readily absorbed, leaving only the core spectra ending at $n=4\rightarrow 1$, while the $n=2$ core spectra enhances the low- n lines from the shell. This is not unlike the phenomena observed at early times with aluminum line absorption, likely along chords, as discussed above. Again, space-resolved spectra using our sensitive TIM-mounted spectrograph could map these features in detail and explain the phenomenology involved.

During this period, we also published in J. Quant. Spectrosc. and Radiative Transfer [10] (reprint included as Attachment B) the analysis of data obtained on the TRIDENT laser facility at the Los Alamos National Laboratory in September 1998, experiments partially supported by NSF. In that campaign, the results of which are quite relevant to the OMEGA experiments, we used two opposing beams (at 0.53 μm wavelength) onto slab targets, and recorded, using a time-gated x-ray crystal spectrograph, spectra from Mg, Al and Si. Digital recording from a CCD camera was used. We also were able to achieve spatial resolution along a direction normal to the target surface. With 170 J per beam in a 1 ns Gaussian pulse, focused to a diameter of 500 μm , we obtained a typical target irradiance of $9 \times 10^{13} \text{ W/cm}^2$; and hence we could simulate the OMEGA irradiance by using small focal spot sizes on TRIDENT. We observed line radiation from resonance transitions in hydrogenic and helium-like ionic species, as well as innershell satellite lines in the next lower species. We also observed $K\alpha$ type transitions at the target surface, which arose from the first few ionization species. Thomson scattering diagnostics provided us with temperature and density values useful in numerical modeling of the data and for future "calibration" of our temperature measurements on OMEGA.

Typical results are indicated in the spectral scan presented in Fig. 2 of Attachment B for silicon. Notable features include unusually intense (relative to

nearby lines) well-displaced satellite lines arising from $2s^2$ - $1s2p$ and $1s2s^2$ - $1s^22p$ transitions in He-like and Li-like ions, respectively, which involve a two-electron process with one photon being emitted. They arise from interactions between the $2s^2$ and $2p^2$ configurations for the two cases. Various explanations for the apparent anomalous satellite intensities have been offered. One candidate is opacity in the radiative transfer of the more intense dipole satellite lines. This has been confirmed tentatively for the lithium-like satellites by numerical modeling carried out by Dr. Ehud Behar of Hebrew University and for the helium-like satellites by Prof. Roberto Mancini of the University of Nevada at Reno. Another explanation under consideration at present is ion-ion charge transfer pumping, two electron included, as suggested by Dr. Frank Rosmej during a six-weeks collaboration with us in August 1999 (that may be repeated during the Summer of 2000).

A very useful aspect of the (mostly) magnesium targets on TRIDENT was our ability to reproduce irradiances as low as a few $\times 10^{12}$ W/cm² on two-sided layered planar targets consisting of 20- μ m thick CH covered with 300-Å of Al and finally a 2- μ m thick Mg coating, simulating the surfaces of the spherical targets used at LLE. We were able to demonstrate that Mg XI and XII x-ray spectral lines are produced at measureable intensities at such an irradiance. This was a point in question in our prepulse measurements at that level of irradiance carried out at LLE and described above. It is also consistent with modeling mentioned above.

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Spectroscopic Measurements of Target Preheating on OMEGA

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ABSTRACT

The preheating of laser-heated microballoon targets has been measured by time-resolved x-ray and extreme ultraviolet (euv) spectroscopy on the 30 kJ, 351 nm, 60-beam laser-fusion system at the University of Rochester Laboratory for Laser Energetics. Thin coatings of aluminum overcoated with magnesium served as indicators. Both the sequence of the x-ray line emission and the intensity of euv radiation were used to determine a preheating peaking at ~ 10 ns prior to onset of the main laser pulse, with a power density $\lesssim 1\%$ of the main pulse. The measurements are supported by numerical modeling. Further information is provided by absorption spectra from the aluminum coating, backlighting by continuum from the heated surface. The exact source of the preheating energy remains unknown at present, but most likely arrives from early laser leakage through the system. The present target diagnostic is particularly useful when all beams cannot be monitored directly at all laser wavelengths.

PAC's: 52.50.Jm; 52.25.Nr; 52.40.Nk; 52.70.La

I. INTRODUCTION

Two related key issues in inertial confinement fusion (ICF) research are the uniformity and integrity of the irradiated spherical targets during surface heating and collapse. This is particularly important for the direct-drive approach, where the surface ablation and acceleration is achieved directly from the focused driving beams. Even small mass or density nonuniformities between layers undergoing mutual acceleration can be greatly amplified according to Rayleigh-Taylor¹ hydrodynamic instabilities or, when shock-accelerated, by Richtmyer-Meshkov² instabilities. Either can lead to pellet breakup prior to ignition. Great care is normally applied to forming targets of smooth surfaces. To avoid the imprinting of non-uniformities from the driving beam, considerable efforts are invested towards creating beams with irradiance uniformity within 1% variation or better on the target surface.

Design and numerical modeling usually are based upon the sudden heating of the target by the driving beam. Hence, any premature surface ablation that takes place prior to the onset of the main driving beam due to significant undesirable energy deposition could lead to a premature slow expansion of the ablator with an enlargement of surface area and decrease in density of the absorber, i.e., to a loss of target integrity. (This is separate from an early "foot" sometimes added prior to the beginning of the laser pulse, with a smoothness comparable to that of the main beam.) If not completely uniform, such a preheating could lead to the imprinting of auxiliary mass uniformities and hence instabilities. In the case of cryogenic targets (e.g., frozen deuterium), it could result in premature sublimation and interior cracking. Therefore, a large effort in laser design involves an enhancement of contrast, i.e., a ratio of laser-to-

prepulse irradiance of perhaps five to six orders of magnitude. With laser drivers operating at a fundamental wavelength in the infrared and frequency upconverted into the ultraviolet, such contrasts are enhanced by a strong dependence of the conversion efficiency upon irradiance.

Such contrasts are routinely checked in laser beam design and development. However, when tens-to-hundreds of beams are involved in irradiating spherical ICF targets, it becomes impractical to routinely monitor the contrast for each beam for all wavelengths present. Instead, it becomes more prudent and meaningful to monitor emission arising from the target prior to the onset of the laser pulse as well as during the pulse to measure the contrast in deposited energy. A simple monitor of reflected light from the target surface at the laser wavelengths involved during the preheat period is a first approach; and indeed such preliminary results provide support for our interpretations. More definitive are time-resolved spectroscopic measurements which relate the emission to a specifically-identified target material (or materials in the case of layered coatings of varying substances). In this paper we describe such extreme-ultraviolet (euv) spectroscopic measurements³ of both emission and absorption from the preheating of multilayered coated spherical targets on the OMEGA laser-fusion system⁴ at the University of Rochester Laboratory for Laser Energetics (LLE). The techniques and interpretations described can be applied to other existing and future facilities.

II. EXPERIMENT

The experiments were carried out between May 1998 and April 1999 in three campaigns. The experimental layout is shown in Fig. 1. Fifty-nine beams from

the "main" Omega laser system constituted the main-laser pulse at a wavelength of 351 nm, with a combined energy of 20 kJ on target in a 1 ns Gaussian pulse. The beams were directed into the vacuum chamber of 3.3-m diameter and focused onto a spherical target of nominal overall diameter 940 μm , resulting in a peak irradiance of $\sim 2 \times 10^{14} \text{ W/cm}^2$. Beam uniformity was provided by a combination of smoothing by spectral dispersion as well as by the use of distributed phase plates. Polymer microballoon targets of 20- μm wall thickness were filled with neon to a pressure of 10 atm, including a 10% admixture of argon as a tracer. A thin coating of aluminum (0.03- μm thick in May 1998 and earlier, reduced to 0.0125- μm thereafter) was applied to the surface to help retain the gas. An additional laser shinethrough barrier (typically 0.1 μm of aluminum) was not applied. Instead, a thicker (2 or 4 μm) outer coating of magnesium was added over the thin aluminum layer to form a tamper layer, also useful for spectroscopic diagnostics.

The euv diagnostics consisted of two 1-m grazing-incidence flat-field spectrographs⁵ covering the 30-300 \AA range with a spectral resolution of 0.1 \AA . Both were equipped with a spherical collecting mirror near the slit. One of these was always equipped with a gated ($\leq 1 \text{ ns}$) four-stripline microchannel plate (MCP) for time-resolution, followed by photographic recording. Cables from these strips (of varying lengths to provide time delays) were properly matched and terminated at the MCP output to minimize pulse reflections. One of the spectrographs was mounted on the outside of the vacuum chamber with the collecting mirror at a distance of 210 cm from the target. The other spectrograph was mounted on a standard re-entrant platform ("TIM" in Fig. 1), with the collecting mirror 28 cm from the target for enhanced sensitivity. The use of such highly-selective (in wavelength) spectrographs was essential to

discriminate against extraneous radiation when measuring large intensity ratios between the preheat and main phases of the event. In contrast, simpler thin-metal bandpass-filtered euv detectors alone can transmit x-rays freely above a certain photon energy at the high temperatures encountered in the main laser-plasma event; while only measuring euv emission in a cooler preheater phase. This can result in a large underestimate of the relative preheat euv emission.

We also had available at LLE a soft x-ray, 100-cm radius, convex-curved crystal (KAP) spectrograph set to measure some of the magnesium and aluminum hydrogenic (Mg XII, Al XIII) and helium-like (Mg XI, Al XII) resonance lines in the 6.4-8.6 Å (1.4-1.9 keV) range. This instrument was equipped with a streak camera and photographic readout for temporal resolution. Since this spectrograph is slitless, the spectral resolution is source-size limited; this feature provided a measure of the spatial extent of the plasma, as determined from the spectral line widths. Also fielded were various LLE-provided x-ray imaging cameras with varying degrees of filtration, and operating in both the time-resolved and time-integrated modes.

III. OBSERVATIONS

The first evidence of significant preheating of the target arose following the first two shots taken in May 1998, i.e., in a rather sudden change of events that continued throughout that campaign. On those first two shots (both at ~14 % lower energy than for the shots that followed), the temporal evolution shown in the x-ray streak spectral data was typical of those obtained in previous campaigns, in that intense and narrow magnesium resonance spectral lines arising from the heating of the outer layer preceded in time those from the thin

aluminum inner layer. Such a time-resolved spectrum is shown in Fig. 2 and is considered "normal", such as we had observed during several previous campaigns at LLE. The magnesium spectral lines are more intense than those from aluminum because of the much thicker coating of the former. Resonance series lines up to $n=5$ (transitioning to $n=1$) are distinct for both helium-like and hydrogenic magnesium. The narrowest aluminum and (initial) magnesium lines corresponds to ~ 1 mm at the target, i.e., approximately the original target size.

In stark contrast to the first two shots, data obtained similarly on the succeeding 19 shots in this campaign showed broadened aluminum lines already at the onset of the laser pulse, accompanied or followed by quite diffuse (broadened) magnesium lines of less intensity. An example of such an "alternate" spectrum is shown in Fig. 3. In such spectra, the magnesium lines are expanded to the extent that series members above $n=2$ are barely distinguishable. The predominance of the aluminum lines indicates that most of the magnesium has been vaporized and that the laser directly heats the aluminum undercoating. Hence, the aluminum lines also become broadened due to expansion.

The Mg XII $n=2$ to $n=1$ Lyman- α line appears to be composed of two lines, which is associated with an already expanded shell of ~ 4 mm diameter at the onset of the main laser pulse, i.e., having experienced an ~ 1.5 -mm radial expansion of the surface during a preheat period of 10 ns (from the euv data discussed below). This results in a radial expansion velocity of $\sim 1.5 \times 10^7$ cm/sec, which is not unreasonable. This measured velocity agrees with numerical modeling of the initial blowoff towards the laser, using the FAST radiation transport hydrocode⁶ for spherical geometry, and assuming an overall

preheating irradiance on target of $3 \times 10^{12} \text{ W/cm}^2$ (as inferred below from the euv data).⁷ There is also experimental evidence of a narrow central line, presumably from the target surface, which indicates that some of the coatings continue to be relatively intact. This is also consistent both with late-time data following compression and with modeling of the target surface area, as described below.

Also present (and not observed previously at all on more "normal" shots) at a time of $\sim 0.7 \text{ ns}$ after onset of the main pulse and persisting for $\sim 0.2 \text{ ns}$ was the K_{α} line formed following innershell ionization in neutral (and slightly-ionized) aluminum at a wavelength of 8.34 \AA . This suggests the presence of fast electrons often associated with instabilities, and perhaps target breakup in the form of both hot spots and cold aluminum, since the K_{α} line is observed concurrently with resonance lines of Al XII and Al XIII. (A much weaker K_{β} line at 7.96 \AA was also detected on one shot, adding confirmation to the identification of the K_{α} line.)

Coincident with the x-ray observations of an apparent preheating and vaporization of the target coatings was the corresponding observation of early euv spectra that included continuum as well as line emissions arising from $n=3$ to $n=2$ transitions, mostly of lithium-like Mg X spectra, but also of less intense helium-like and hydrogenic Mg XI and XII spectra, respectively. The latter emissions from the higher ionization stages peaked at $\sim 10 \text{ ns}$ before the onset of the main laser pulse, giving way to Mg X (and lower) spectra in the last 5 ns. (Similar emissions from the 70-times thinner aluminum layer are expected to be much less intense, with some unfortunate wavelength overlaps, and are difficult to distinguish.) Also observed prior to the main laser pulse onset were carbon lines, presumably arising from: (a) the microballoon surface following localized penetration, (b) the target surface as contaminants, or (c) the $10\text{-}\mu\text{m}$ diameter

supporting stalk. They showed a dominance of C VI over C V $n=2$ to $n=1$ resonance lines.

Such preheating euv spectral features are illustrated in the temporal plot of overall emission in the 100-200 Å wavelength region, shown in Fig. 4 from multiple shots during the May 1998 campaign, using four MCP gated striplines for temporal resolution. This indicates a preheating euv emission of $\sim 0.7\%$ of that of the corresponding continuum spectrum observed during the main event. This suggests a power flux deposited at the target of $\sim 3 \times 10^{12}$ W/cm² in the preheat period, which was used as input for the numerical modeling⁶ discussed elsewhere here.

That an irradiance of this magnitude could produce some Mg XI and XII euv emission is evidenced experimentally by a distinct x-ray streak spectrum obtained at the lowest irradiance used at LLE, namely 7×10^{12} W/cm². Additionally, recent experiments at Darmstadt⁸ with a laser-irradiated magnesium slab target produced an x-ray spectrum that included Mg XI and XII at an irradiance as low as 1×10^{12} W/cm². (Again, one cannot rule out hot spots as a source of radiation from these species, particularly when K_α lines are observed, as described above.) As with the x-ray streak spectra, the euv preheating began on the third shot in the May 1998 campaign, i.e., no euv spectral emission was measured on the two striplines set for early timing on the first two shots (the remaining two striplines timed later did record the main event, which verified proper operation of the detector). This again suggests some sudden event after the first two shots, resulting in target preheating.

The C VI and Mg X spectral lines observed at the peak of the preheating period shown in Fig. 4 signify an electron temperature of $kT_e \sim 100\text{-}150$ eV, which again is consistent with numerical computations⁶ for the same pulse shape⁹ as shown in Fig. 4 and using an overall target irradiance of 3×10^{12} W/cm², as inferred above. Also, the calculations predict an expanding (towards the laser) plasma consisting of ions dominated by charge states of Mg⁹⁺ (Li-like) and Mg¹⁰⁺ (He-like) at the peak of the preheating pulse. This accounts for the Li-like nd-2p emission from recombination (as well as excitation). Excitation of line emission in the He-like and H-like ions during this period most likely occurs in hot spots (as supported again by K α emission). A measured decline to a predominantly Li-like Mg X spectral feature in the 4-5 ns prior to main laser pulse onset is consistent with numerical predictions of a temperature drop to a few 10's of eV and charge states ranging from Mg⁶⁺ to Li-like Mg⁹⁺ during that period. This is followed by a reheating of residual magnesium at main laser pulse onset, once again producing Mg XI and XII line emissions in a collapsing plasma.

The numerical modeling of density, using the pulse in Fig. 4, also predicts at least a partial breakdown in target integrity beginning at $\sim 3\text{-}4$ ns after onset of the preheat, followed by a heat wave flowing through the 20- μm thick microballoon wall into the gas filling to a depth of ~ 100 μm at the time of onset of the main laser pulse. This breakup could be contributing to the decrease in euv emission shown in Fig. 4 beginning at ~ 8 ns prior to main laser onset. However, a boundary between the CH and the magnesium also is predicted to be somewhat sustained, along with some residual magnesium available for reheating at main laser pulse onset, as observed. All modeling^{6,7} predicts a critical-density layer traveling inward during the preheating phase. Since this

layer is at a density much less than that for a solid, it indicates once again that surface material such as magnesium has expanded away.

That the weaker euv spectrum observed in the preheat period is not simply later-time emission recorded by a weak secondary pulse to the MCP striplines has been verified experimentally by the vast difference in the simple, few-line, few-species spectra obtained at early times compared to a multi-line spectrum from recombination to many species at the later times, including Ne VIII (Li-like) 2p-3d emission at 98 Å from the core filling, not present in the preheated plasma. Another distinct aspect of the late-time recombination spectra is a ratio of C VI to C V resonance lines of order unity, signifying a temperature of $kT_e \sim 75$ eV, lower than at the peak of the preheating phase as reported above. Hence, this alternative explanation for the preheater spectra observed is first ruled out on the basis of non-time-correlated spectral features. Furthermore, a measured ratio of peak voltages between the secondary and primary pulses of $\sim 1/5$ results in a MCP gain ratio of 10^{-8} , using a power-law gain dependence^{10,11} on applied stripline voltage V as strong as V^{11} , thereby resulting in a negligible gain on the secondary pulse.

A cause for this sudden onset of preheating could not be established at the time of the May 1998 campaign. During the October 1998 campaign, which included 21 shots, the anomalies in the x-ray streak spectral data were not readily apparent. The euv preheat emission, while still present with the same ionic species, was $\sim 1/4$ of that measured in May. These data indicated that the preheating effects had diminished somewhat, perhaps associated with improved binding of the magnesium with the use of aluminum layers of less than one-half the thickness.

A further understanding of the preheating mechanism at the target arose from the third campaign, carried out in April 1999. For this limited series of 10 shots, the closer-in and hence more sensitive TIM-mounted euv spectrograph was equipped with the time-gated MCP detector. Again, the aluminum inner layer was thin ($0.0125\text{-}\mu\text{m}$). For this campaign, the $2\text{-}\mu\text{m}$ thick magnesium outer layer was overcoated with $2\text{ }\mu\text{m}$ of CH to provide additional tamping. Also, some $2\text{-}\mu\text{m}$ thick LiF-coated targets (without CH overcoatings) were used. The results to be described next were similar for all of these target designs.

The euv data in this campaign showed overall continuum emission during the preheat phase equivalent to that from the October 1998 campaign. Particularly obvious in this series of experiments were a number of absorption features in the euv spectra during the preheating phase which were not present during the main event. These are shown in the spectrum in Fig. 5. This attests to the "purity" of the euv continuum, i.e., not being an accumulation of higher orders and scattered light of other wavelengths which would not have been absorbed in these lines. The strongest and broadest absorption feature (at a wavelength of $116\text{ }\text{\AA}$) matches a blend of the energies¹² for photoionization from the 2s innershell levels in neutral aluminum as well as in Al^{1+} and Al^{2+} ions. In addition, photoionization edges¹³ associated with valence electrons in Al^{3+} to at least Al^{5+} match other narrower absorption features in the spectra, included in Fig. 5. (That 2s-innershell absorption at $156\text{ }\text{\AA}$ did not appear for magnesium is another indication that the magnesium coating vaporizes and expands at early times in the preheat phase.)

One interpretation of such absorption features is that they arise from a continuum background formed at the front surface of the CH microballoon (the balloon wall as well as the gaseous fill are opaque in this region, ruling out the possibility of a rear-target-surface backlighting, unless one invokes major target breakup). The 0.0125- μm thick solid aluminum sealant layer is semi-transparent ($\sim 25\%$) to the 351-nm wavelength laser light,¹⁴ while also being semi-opaque ($\sim 40\%$ absorption) at the x-ray edges.¹⁵ As such, laser preheater light can be transmitted through the thin aluminum layer to the outer surface of the polymer target, generating a continuum backlighter, and forming the 2s absorption feature in the aluminum layer. This holds similarly for n=2 absorption in an aluminum plasma that might be formed such that the product of density and thickness remains constant over a limited range. In addition, the absorption features from multiple ionization stages could be arising from different points on the target, and perhaps from varying thicknesses of the aluminum layer, if it is not completely uniform in the case of hot spot formation.

Alternatively, or additionally, it is possible that the absorption is forming along chords in the "limb" of the spherical shell of aluminum, rather than through the layer itself. This would increase the absorption length by perhaps a factor-of-10 compared to that for absorption perpendicular to the layer. In this scenario, the continuum backlighting could be arising from a heated coating, as well as the CH surface.

In campaigns that took place earlier than that of April 1999, only a close examination of the October 1998 data taken with the more remote spectrograph showed a weak presence of the 2s-innershell absorption band. This marked difference may have been a result of a different mounting position on the

spherical target chamber for the April 1999 campaign, being almost opposite to the earlier ones. This could be interpreted as an effect of certain selective beams being involved in the preheating, rather than all 59 uniformly. This could also have contributed to the backlighting and absorption along a chord.

Turning to the x-ray streak spectra obtained during this April 1999 campaign, a very sensitive fresh CsI photocathode was in place on the x-ray streak spectrograph for two shots. As such, the late-time spectra were very overexposed. However, a close inspection of the films obtained indicate some evidence of a low-intensity Mg XII Ly- α line for approximately 1 ns prior to the main pulse. The maximum preheater emission at -10 ns, expected from the euv data, could not be detected on these two shots because of a limited sweep range, and remains a topic for future experiments.

IV. CONCLUSIONS

The early-time spectrographic data presented here are consistent with a numerical model of low-energy preheating in regions of sufficient energy deposition in the target to produce Mg X (and XI and XII in spots) spectra and premature evaporation and subsequent expansion of the outer coating, along with a loss of target integrity. Transmission of this beam through the thin aluminum sealant layer leads to a preheating of the target surface, thereby producing C V and VI emission along with a continuum that backlights the aluminum and produces observed absorption features, either perpendicular to the layer surface or through chords in the shell, along the axis of view, or perhaps both.

An obvious explanation for the target preheating is that there is a relatively weak laser prepulse occurring at early times, if not from the ultraviolet (0.351 μm wavelength) beam, perhaps of a fundamental- (1.053 μm) or doubled- (0.527 μm) equivalent frequency not completely converted. Other experiments as well as numerical modeling now show that a projected irradiance of $\sim 10^{12} \text{ W/cm}^2$ is sufficient to produce the observed features. If indeed the preheating arises from laser beams, such heated zones would represent severe non-uniformities in the beam profiles and could result in laser imprinting at the beginning of the main pulse, thereby generating instabilities as mentioned earlier here and breakdown of target integrity throughout. Also, excessive preheating of the low mass CH surface adjacent to the aluminum layer could produce pressure differences to further drive instabilities. It will be very interesting to obtain additional measurements of the preheating pulse shape for a more exact modeling of the effects.

The source of the energy for the observed target preheating remains under investigation, as does the wavelength (i.e., fundamental, frequency doubled or tripled), should it be associated with the OMEGA laser. The power of this target technique in diagnosing a prepulse and the overall contrast with the main beam when multiple beams are present is demonstrated. The fact that this preheating began as a sudden event which may have subsided somewhat over time indicates the importance of a thorough understanding and continuing monitoring of the energy reaching the target, lest it reoccur randomly during future experiments.

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FIGURE CAPTIONS

Fig. 1. Layout of the experiments. "TIM" refers to the ten-inch manipulator re-entrant platform upon which one of the euv spectrographs was mounted.

Fig. 2. X-ray streak spectrum typical of those obtained for the first two shots in the May 1998 and in previous campaigns, showing the expected sequence of Mg XI (He-like) and Mg XII (H-like, Lyman series) resonance lines followed by similar Al XII and Al XIII lines, from outer and inner layers, respectively.

Fig. 3. X-ray streak spectrum typical of those obtained after the first two shots in the May 1998 campaign, showing the early onset of aluminum lines from the inner layer. This shows the dominance of Al XII (He-like) and Al XIII (H-like Lyman series) lines from the inner layer at early times.

Fig. 4. Logarithmic time history of euv emission in the 100-200 Å wavelength range. The data points shown and connected by a smooth curve were obtained from various shots and striplines used in the May 1998 campaign, with normalization between strips obtained from pointing shots taken with gold targets. Corrections have been applied for film saturation. The ~4 ns duration of emission during the main pulse agrees with that of continuum emission in the x-ray streak spectra (not readily apparent in the copy in Fig. 3), possibly even including here (but not curve-fitted) a decrease over an interval of ~0.5 ns at 3 ns from the onset of the main laser pulse.

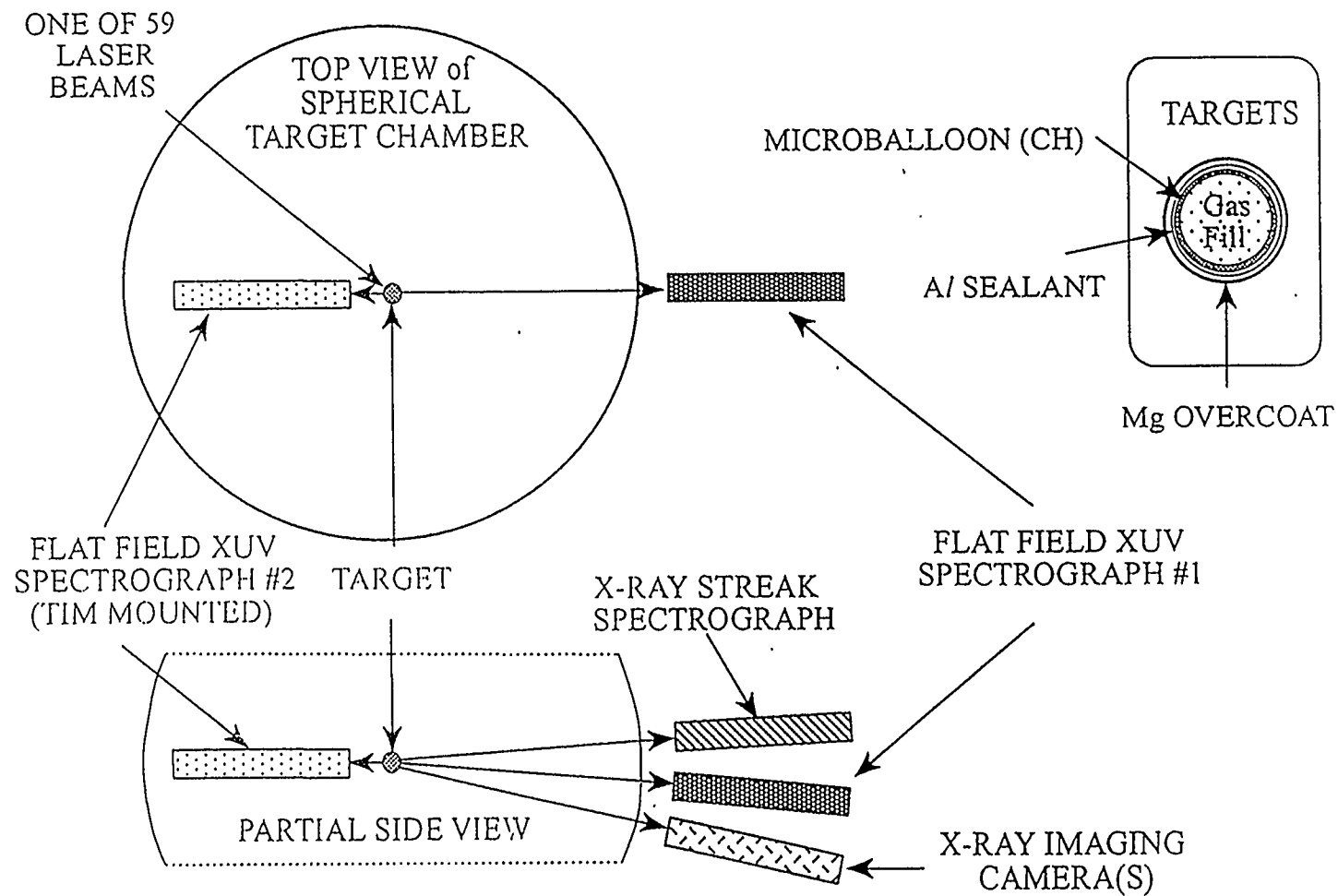


Fig. 1. Layout of the experiments. "TIM" refers to the ten-inch manipulator re-entrant platform upon which one of the euv spectrographs was mounted.

Fig. 5. Euv spectra obtained from two adjacent MCP striplines during the April 1999 campaign, indicating some main emission as well as absorption features. The carbon emission is most likely associated with target surface vaporization.

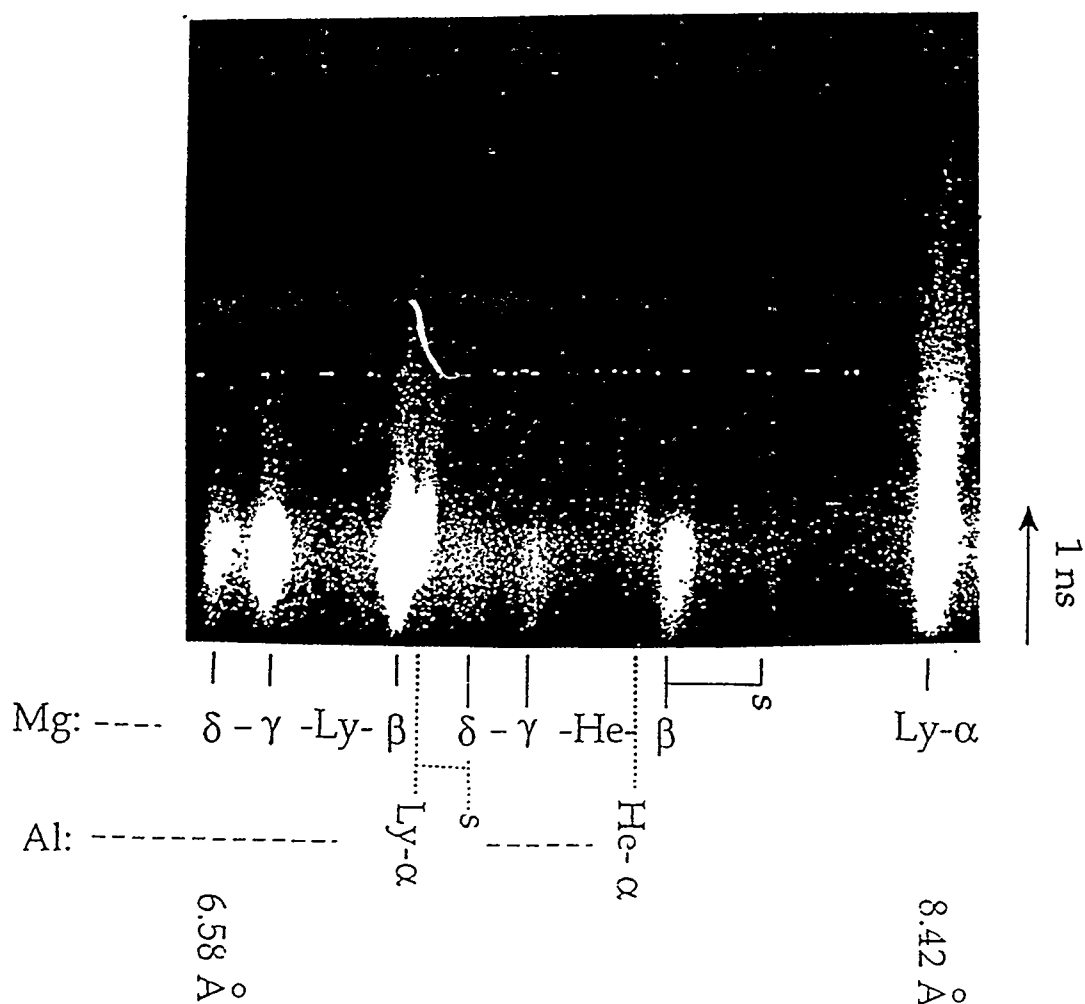


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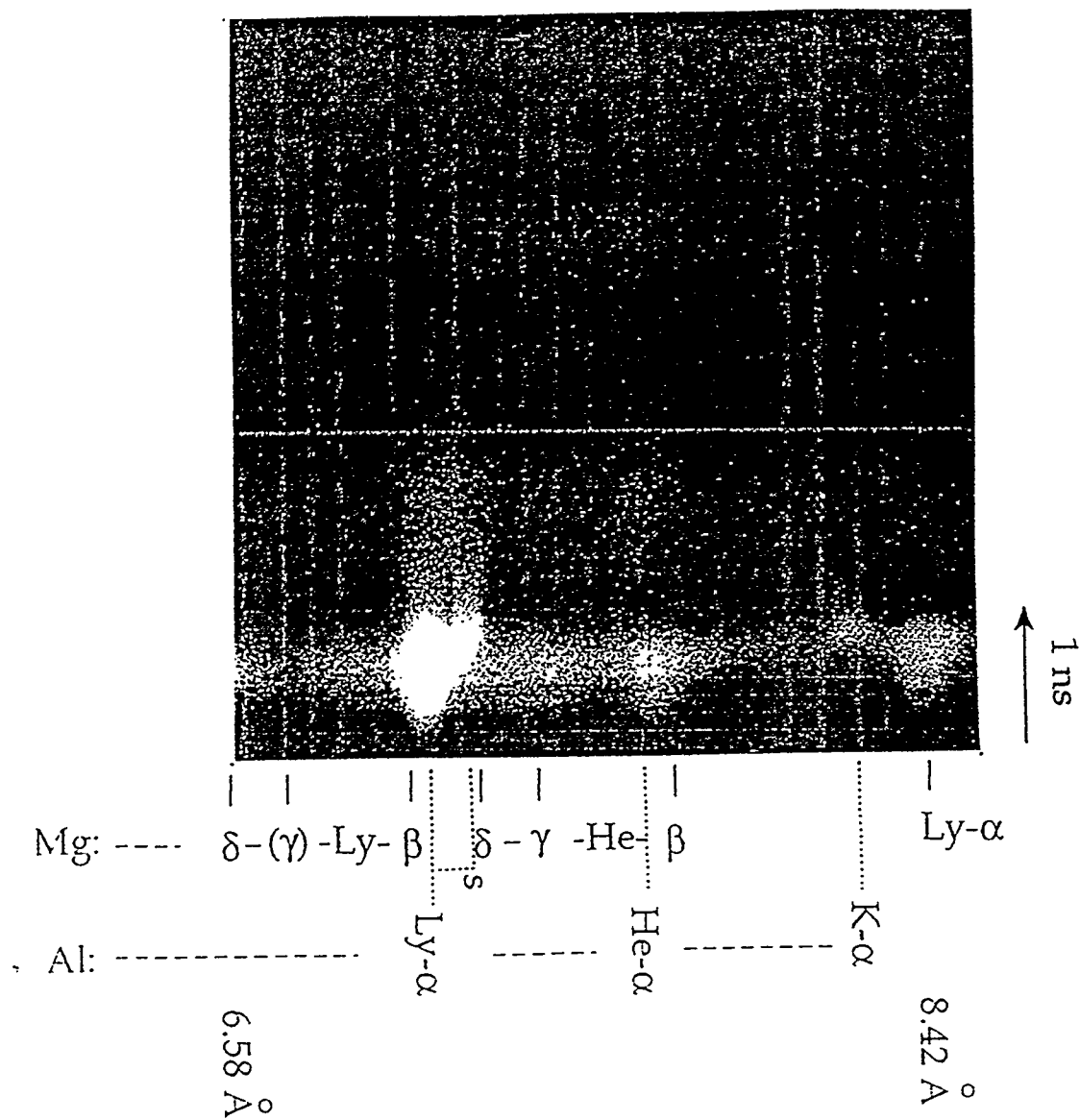


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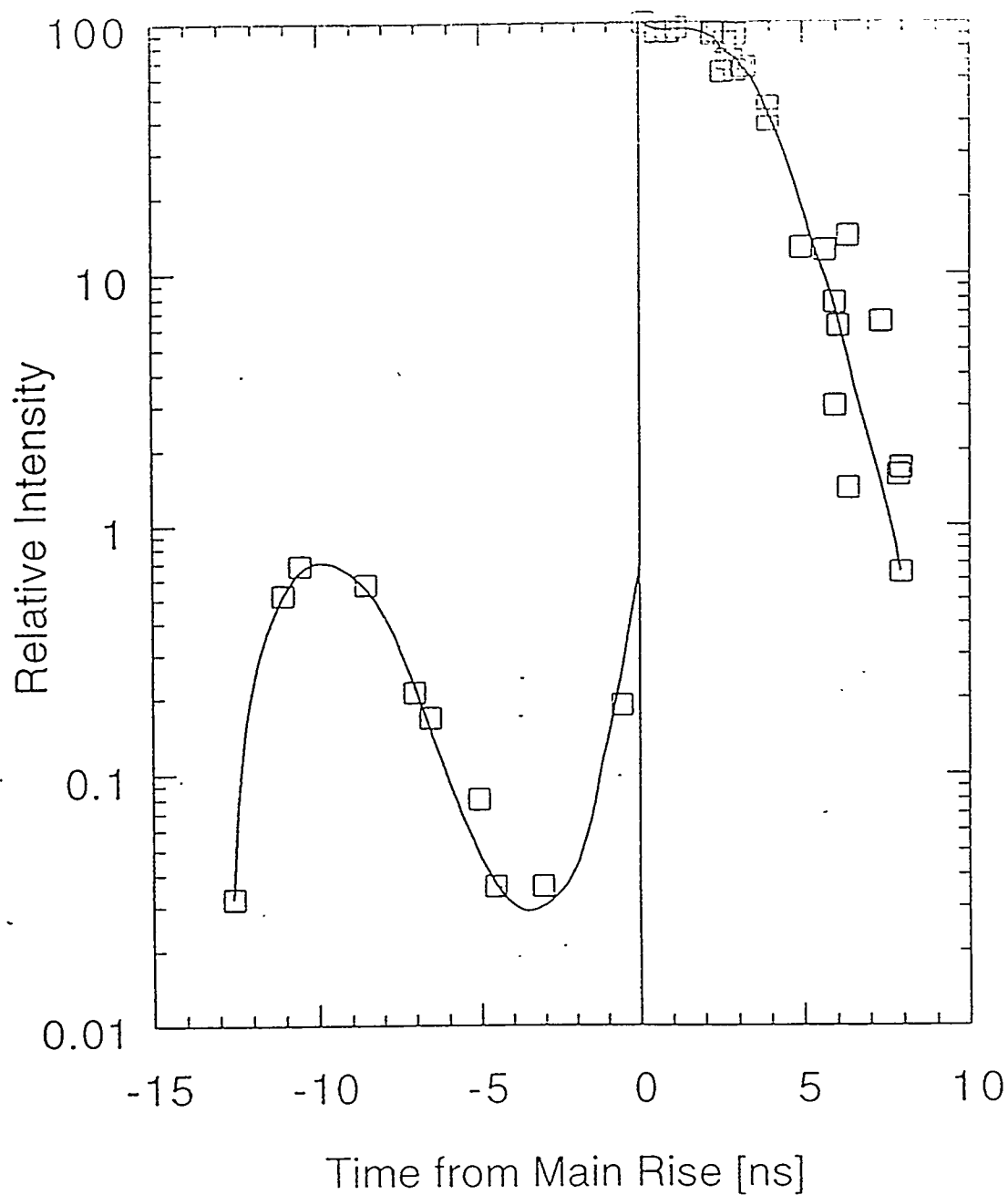


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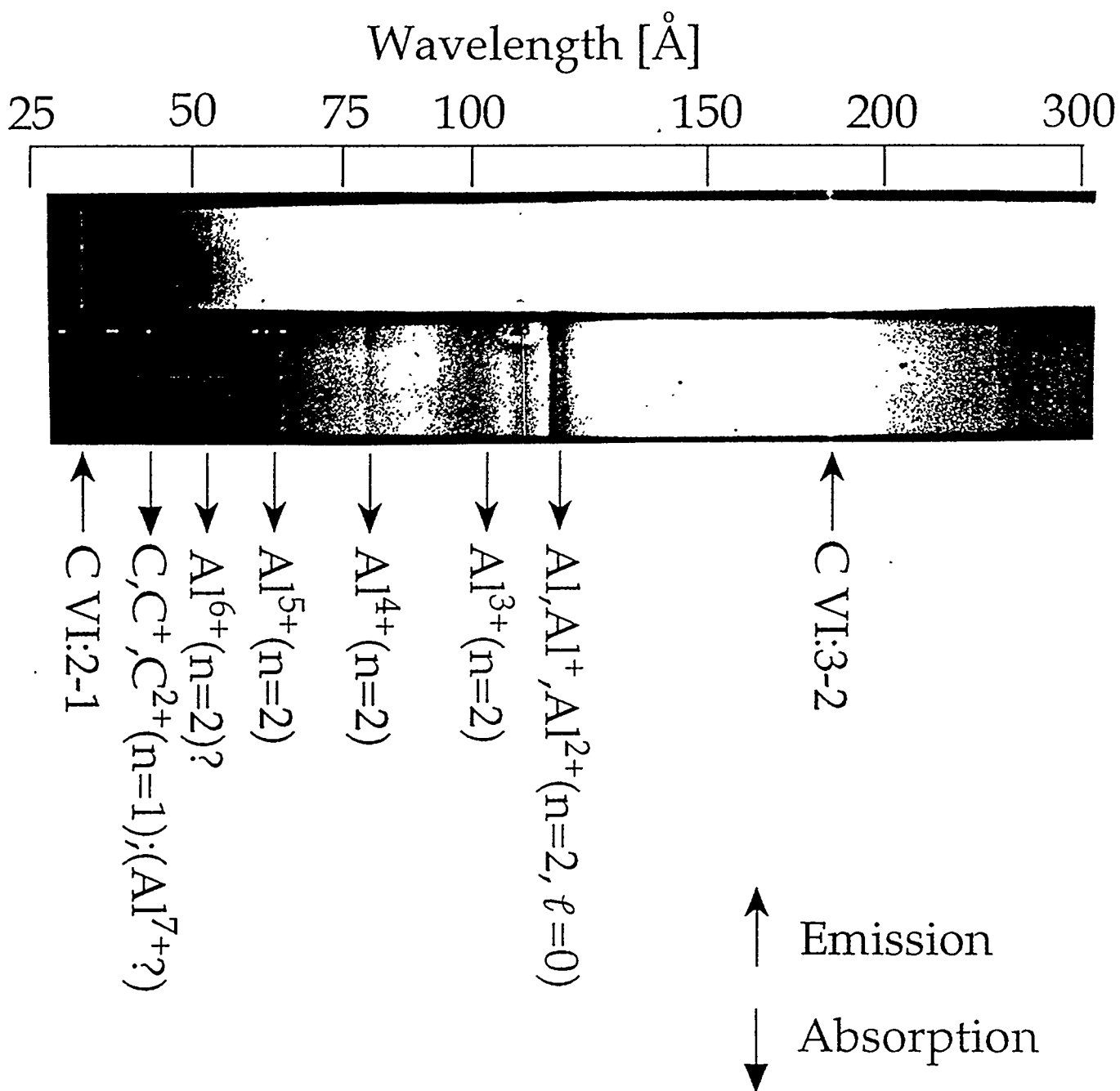


Fig. 5. EUV spectra obtained from two adjacent MCP striplines during the April 1999 campaign, indicating some main emission as well as absorption features. The carbon emission is most likely associated with target surface vaporization.

ABSTRACT

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Further Evidence for a Prepulse on OMEGA* R. Elton, H. Griem, and E. Iglesias, *Univ. of Maryland Institute for Plasma Research*. Our previously-reported [Bull. Am. Phys. Soc. 43, 1646 (1998)] observation of a disruptive prepulse on OMEGA, possibly as large as 1% of the extreme ultraviolet (euv) radiation measured from the main pulse, has recently been substantiated by the measurement of euv absorption spectra, prior to the main pulse. The absorption features have been identified with $n=2$ photoionization in aluminum atoms and ions up to Al^{5+} . Cold aluminum is originally present as a 0.0125- μm thick sealant coating applied to a neon-filled (10 atm) CH microballoon, with an euv transmission at the L-absorption edge of $\sim 50\%$. The aluminum in turn is overcoated with 2 μm of Mg. The spectra which show the absorptions also include continua as well as line emissions from Mg^{9+} to Mg^{11+} ions. These occur prior to the onset of laser target irradiation by at least 10 ns, and imply a prepulse irradiance of about $10^{12} W/cm^2$. Since the neon and CH are opaque to euv radiation from the rear, a likely scenario is early (prepulse) vaporization of the outer Mg layer, perhaps in hot spots, followed by laser radiation transmitted through the thin Al layer, thereby heating the CH surface. This could provide an euv continuum backlighter for the aluminum that leads to the euv absorption features in various ionic species, perhaps from different points on the target surface.

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