Pyrometric Measurement of the Temperature of Shocked Molybdenum

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Measurements of the temperature of Mo shocked to \( \sim 60 \) GPa and then released to \( \sim 28 \) GPa were previously attempted using high explosive driven flyer plates and pyrometry. Analysis of the radiance traces at different wavelengths indicates that the temporal evolution of the radiance can be explained by a contribution from the LiF window to the measured thermal radiation.

Fitting the radiance traces with a simple model, supported by continuum dynamics studies which were able to relate structures in the radiance history to hydrodynamic events in the experiment, the contribution of the window was obtained and hence the temperature of the Mo sample. The shock- and release temperature obtained in the Mo was 762\( \pm \)40K which is consistent with calculations taking the contribution of plastic work to the heating into account. The radiance obtained for the LiF window shows a non thermal distribution which can be described by a bulk temperature of 624\( \pm \)112K and hot spots (less than 0.5% in total volume) within the window at a temperature of about 2000K.

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I. INTRODUCTION

The measurement of temperature in shock physics experiments is of paramount importance since temperature is required as constraint (additional to pressure and volume) for the development and credibility of robust multiphase equation of states (EOS). The application of modern material strength models requires these EOS with accurate temperatures and phase boundaries, particularly melt curves. Temperature is an important parameter in geophysics and planetary sciences for an accurate description of planetary structure and astrophysical impacts. Many measurements of the EOS of planetary materials, and the pressure calibration on which diamond anvil cell studies rely, are made using shock experiments, and rely on temperature corrections to infer states off the shock Hugoniot. General research and model development for the response of materials to extreme dynamic conditions is focusing on multi-scale approaches, in which thermally-activated processes including plastic flow, phase changes, and chemical reactions requires an accurate knowledge of the temperature. As a specific example, the melting curve and kinetics of Beryllium and Carbon (Diamond) at elevated pressures are important to understand the behavior of ablator capsules for the national ignition facility (NIF) to succeed in harvesting fusion energy from laser driven capsule implosions.

Pyrometry is a form of thermal emission spectrometry in which the emission spectrum is collected in a small number of spectral bands of significant bandwidth, compared with energy-dispersive spectrometers such as prisms and gratings. Energy-dispersive spectrometers are not practical for most shock experiments at temperatures below a few thousand Kelvin because the thermal emission is too weak to take advantage of the relatively fine spectral resolution. Pyrometry is thus the most promising technique applicable to many materials and experimental configurations (besides Neutron Resonance Spectroscopy and Raman Spectroscopy, both with their advantages and limitations) to achieve the goals stated above. Nevertheless, although pyrometry has been fielded on dynamic loading experiments for more than four decades, it still suffers from problems such as background light, thermal and non-thermal emission from the window material and sample/window interface effects which are hard to take into account.

Neutron Resonance Spectroscopy (NRS) has been investigated as a possible bulk-temperature diagnostic for shock physics experiments, capable of measuring the temperature within opaque samples. Initial experiments on shock loaded Mo showed significant discrepancy with theoretical predictions. To investigate this discrepancy, microsecond-duration pyrometric measurements were performed on Mo samples through LiF windows. These experiments used a shock generation method that was the same as used in the NRS experiments, and also included Doppler velocimetry measurements of the surface velocity history of the sample to verify the loading conditions. Previous analysis of the pyrometry experiments did not succeed in extracting temperatures. Here we report a more rigorous study of the effects of shock and release waves on pyrometry measurements with a release window, identifying two-temperature population from pyrometry data. We are able to correlate features of the radiance history at a level...
usually dismissed as too complicated for further analysis and provide valuable insight into dynamic processes occurring in shock experiments using a window. This allows us to extract temperatures more accurately and with greater confidence. The results presented here are the first simultaneous extraction of sample and window temperature to our knowledge which enhances the ability to reliably interpret pyrometry data. This is a first step and key to future research and physics-based understanding of response of condensed matter subject to dynamic loading and heating.

II. EXPERIMENTAL SETUP

As in the NRS experiments, a shock was induced in the Mo sample by the impact of an Al disk accelerated to a speed of about 3.5 km/s by means of high explosive (HE) gases. A LiF window was glued to the backside of the Mo disk (where the measurement was made) in order to maintain high pressure over an area accessible for contactless temperature measurements. Emitted thermal radiation was focused onto a 1-mm core-diameter near-infrared (NIR) fiber as well as onto a 1-mm core-diameter visible-glass, low OH fiber. Two different multichannel pyrometers were used to infer thermal radiance over a wide range of wavelengths: 5 channels in the visible and NIR wavelength region, using photomultiplier tubes (PMTs), and 4 channels in the NIR, using InSb detectors. The velocity history at the Mo/LiF interface was measured by laser Doppler velocimetry, using a Velocity Interferometer System for Any Reflector (VISAR) in order to infer the pressure history applied to the sample.

In the case of low-temperature pyrometry measurements, special precautions have to be taken in order to avoid background light, which can easily overwhelm especially the short wavelength channels. These backgrounds can be generated either by bright HE gases blowing by the target, air lighting up due to shock luminescence from nitrogen, ejecta or jets generated due to improper surface preparation, or sharp edges. In order to minimize the background, the experiments were performed in a vacuum (10-3 Torr). Two CaF lenses were used to focus the thermally emitted light onto a 1-mm diameter NIR fiber (centered 2.5 mm off axis at the fiber bundle) and a 1-mm diameter visible low OH fiber (centered 2.5 mm diametrically opposed). The center fiber was a VISAR probe used to determine the velocity of the sample after impact. The LiF window, 30 mm diameter by 20 mm thick, was included to maintain an elevated pressure in the Mo when the shock reached its surface, avoiding a release to atmospheric pressure. The LiF was attached to the Mo coupon using Loctite 326 glue, which has been previously found not to cause considerable amounts of thermal emission. This simplifies relating the surface temperature to the bulk temperature of the shocked sample. However thermal light emission from both the glue layer and the LiF is a concern.

The NIR pyrometer collimates and divides the incoming light from a single NIR fiber into four spectral ranges, using three custom dichroic beamsplitters. These four collimated beams are spectrally narrowed by bandpass filters centered at 1.8 µm, 2.3 µm, 3.5 µm and 4.8 µm and then refocused onto the 1 mm2 active area of the 50 MHz bandwidth InSb detectors using ZnSe lenses. The lower temperature limit of the two longer wavelength channels is about 340 K (assuming an emittance of 1). For more information on the NIR instrument and on data analysis see Ref. 11.

The 5 channel visible/NIR pyrometer also uses dichroic beamsplitters to spectrally divide the incoming light into five beams, which are then refocused onto the active area of photomultiplier tubes (PMT). Also using bandpass filters, these five channels are centered at 505 nm, 725 nm, 850 nm, 1.23 µm and 1.59 µm. A holographic notch filter is used to suppress the bright 532 nm laser light used for the VISAR measurements. More information on this instrument can be found in Ref. 12.

For further details on the HE driven flyer system see references 7 and 13.

III. RESULTS

Eight experiments have been performed at the "Chamber-8" high explosive experimental facility at the Los Alamos National Laboratory in September 2004; these experiments are summarized in Table I. Figure 2 shows the radiance as function of time as well as the sample/window interface velocity for experiment #06. The time of shock breakout is set to t = 0 µs. If no unwanted background light occurs the radiances are...
TABLE I: Experiments performed. Surface condition (free surface or window), diagnostics (V, VISAR, IR, IR-pyrometer; vis, visible-pyrometer)

| Exp # | Surface     | Diagnostics | Comments            |
|-------|-------------|-------------|---------------------|
| 01    | LiF window  | V, IR       | Data clipped        |
| 02    | LiF window  | V, IR       | Data lost           |
| 03    | LiF window  | V, IR       | Good data           |
| 04    | free surface| V, IR       | HE problems, no data|
| 05    | free surface| V, IR       | Good data           |
| 06    | LiF window  | V, IR, vis  | Good data           |
| 07    | LiF window  | V, IR, vis  | Good data           |
| 08    | free surface| V, IR, vis  | Good data           |

FIG. 2: Measured radiances for seven wavelengths and sample/window interface velocity for experiment #06.

expected to be constant from the time of shock breakout until the release wave from the back of the flyer reaches the sample/window interface (about 0.7 μs after breakout). At this time the sample/window interface is decelerating to a lower velocity as can be seen from the particle velocity trace.

Continuum dynamics simulations were used to investigate the origin of the varying signals in the shocked state by comparing them with the various hydrodynamic events occurring in the experiments. One and two dimensional (1D and 2D) simulations were performed, using general-purpose multi-physics hydrocodes. Shock dynamics calculations were also performed using the same material properties, solving the Rankine-Hugoniot equations to obtain shock states and hence wave speeds more precisely than by derivation from the discretized hydrocode solutions\textsuperscript{14}. The 1D and 2D hydrocodes both used a finite difference representation of spatial fields, a second-order predictor-corrector algorithm for time integration, and artificial viscosity to stabilize shocks. The 1D simulations, along the axis of symmetry, were Lagrangian, avoiding inaccuracies caused by numerical advection, and used the LAGC1D hydrocode\textsuperscript{15}. The 2D simulations, axisymmetric in the axial-radial plane, were Eulerian for robustness in treating highly distorted flow from sharp corners, with an operator-split third order van Leer flux limited method for advection, and used the EUL2D hydrocode\textsuperscript{16}.

The simulations included the HE-driven flyer, the Mo target, and the LiF window. The HE acceleration system itself was not modeled: the flyer was treated as flat and moving at a constant speed, starting at the instant of impact. In reality, the flyers were slightly dished, still accelerating slightly, and reverberating, but these details should not make a significant difference to the pyrometry data. The equations of state (EOS) used the cubic Grüneisen form with published parameters for each material\textsuperscript{17}. Some simulations were repeated using EOS from the SESAME tabular compendium\textsuperscript{18}, the difference was negligible. Strength was treated using the Steinberg-Guinan model\textsuperscript{19}, with published parameters for each material\textsuperscript{17}. In the 1D simulations, the cell size in the flyer, sample, and window was 0.1, 0.05, and 0.2 mm respectively. In the 2D simulations, the cell size was 0.1 mm.

A position-time diagram for the key mechanical waves from the simulations is shown in figure 3 (lower part) together with the measured spectral radiance at 2.3 μm for experiment #06.

The radius increased from the time of shock breakout until about 2.1 μs. Then the radiation decreased at a rate slightly lower than the increase. The kink in the radiance at about 3.1 μs occurs at about the time when the release from the edge of the window reaches the center of the sample-window interface, cooling it by adiabatic expansion. The shock wave reaches the rear surface of the window at about 3.75 μs, at this time the signal of all wavelengths goes into saturation because of a bright

FIG. 3: x-t-diagram (lower part: black lines, boundaries between flyer (F), sample (S) and window (W); grey lines, shock front; dashed lines, head and tail of flyer release waves), and measured spectral radiance at 2.3 μm for experiment 06 (upper part: the time has been shifted so that the breakout at the sample-window interface occurs at the same time as at the simulated x-t-diagram).
The recorded pyrometer output signals for experiment #01 have been clipped because the settings of the digitizer have been too sensitive (the increase of radiance with time was not expected), the measured radiances for the other successful experiments look very similar to the ones shown in figure 2 (for experiment #06) and are not shown here.

The radiance traces for the free surface experiments are not analyzed and discussed here because, additionally to problems with background light, the surface temperature was not homogeneous. (This can be inferred from the response of different wavelengths shortly after the time of shock breakout. For more details on spatial temperature non-uniformities see reference 20.) The background light at the free surface experiments was most likely caused by thermal radiation from APIEZON Q®, a soft, black, putty-like substance that was intended to absorb thermal radiation from hot jets formed at the edge of the sample/window interface.

IV. DATA ANALYSIS

Figure 3 shows the parts involved in the process of emission of thermal light for the sample/window combination used in these experiments, including the effect of the interface between shocked and unshocked window material.

Thermal emitted radiance from the sample \(L_{\text{sample}}\) is multiply reflected between the interfaces \(R_1\), \(R_2\) and \(R_3\). These reflectivities can be obtained from the optical properties of the sample, the shocked and the unshocked window. The index of refraction of the shocked window can be estimated from the Gladstone-Dale relation, from this it can be seen that \(R_2\) is very small and can be neglected. The index of refraction and extinction coefficient of the unshocked window can be found in Ref. 22, the absorption in this part can be neglected as well. The shocked part of the window can emit thermal radiation \(L_{\text{window}}\) as well as absorb thermal radiation emitted by the sample. The emissive power of translucent materials (equivalent to emissivity of opaque materials) can be expressed by \(1 - e^{-\alpha x}\), and the absorption can be expressed by \(e^{-\alpha x}\), where \(x\) is the thickness of the shocked layer and \(\alpha\) is the absorption coefficient. The absorption coefficient is related to the extinction coefficient by the following relation:

\[
\alpha = \frac{4 \cdot \pi \cdot k}{\lambda_0}
\]

where \(k\) is the extinction coefficient and \(\lambda_0\) is the wavelength in vacuum. Thus the radiance from this assembly can be expressed as:

\[
L = \frac{(1 - R_3)}{(1 - R_1 \cdot R_3 \cdot e^{-2\alpha x})} \cdot [L_{\text{sample}} \cdot \epsilon \cdot e^{-\alpha x} + L_{\text{window}} \cdot (1 + R_1 \cdot e^{-\alpha x}) \cdot (1 - e^{-\alpha x})]
\]

where \(\epsilon\) is the effective emissivity of the sample into the window, taking the interface effects and absorption of the glue layer into account. The emissivity is mainly determined by the optical properties (\(n\) and \(k\)) of the sample and the window, but can increase if the interface roughness increases following shock breakout, for instance if the interface is Richtmyer-Meshkov unstable or because of flow or jetting from surface texture.

The first term in equation 2 \((1 - R_3)/(1 - R_1 \cdot R_3 \cdot e^{-2\alpha x})\) is very close to 1 (between 0.98 and 1.0 in all practical cases) and can be set to 1 without compromising the accuracy of the analysis. The term \((1 + R_1 \cdot e^{-\alpha x}) \cdot (1 - e^{-\alpha x})\) can be simplified to \((1 - e^{-2\alpha x})\) without compromising the fitting results for the sample radiance, introducing an error in the window radiation and absorption coefficient of less than 10%, which is far below the uncertainty of the fitting parameters. Thus equation 2 can be approximated as:

\[
L = L_{\text{sample}} \cdot \epsilon \cdot e^{-\alpha x} + L_{\text{window}} \cdot (1 - e^{-2\alpha x})
\]

The thickness \(x\) of the shocked layer is given by: \(x = (u_s - u_p) \cdot t\), where \(u_s\) is the shock velocity in LiF (7.10 mm/μs at a pressure of 27GPa) and \(u_p\) is the particle velocity at the sample/window interface (1.45 mm/μs on release into LiF from a Hugoniot pressure of 64GPa in Molybdenum) and \(t\) is time after shock breakout at the Mo/LiF interface. Equation 3 was used to fit the radiance traces of experiments #01, 03, 06 and 07. The results are shown in table 4.

While the emissive power of the window material was obtained from the fitting parameters, and hence the temperature of the shocked window can be obtained directly, the inferred temperature of the sample depends strongly on the effective emissivity \(\epsilon\) of the sample-window combination. Since this property is very hard to determine
TABLE II: Fitting results for the experiments #01, 03, 06 and 07. Where there are no data available (due to low signal level or poor quality of the fit) the cells have been left blank.

| Exp # | $\lambda$ (µm) | $\epsilon \cdot L_{M0}$ (W/m$^2$·nm·sr) | $L_{LiF}$ (W/m$^2$·nm·sr) | $a_{LiF}$ (mm$^{-1}$) |
|-------|----------------|--------------------------------------|-----------------------------|------------------------|
| 01    | 3.5            | 0.117                                | 2.27                        | 0.0109                 |
|       | 4.8            | 0.051                                | 0.229                       | 0.0988                 |
| 03    | 3.5            | 0.110                                |                             |                        |
|       | 4.8            | 0.062                                |                             |                        |
| 06    | 0.85           |                                      |                             |                        |
|       | 1.23           |                                      |                             |                        |
|       | 1.59           |                                      |                             |                        |
|       | 1.8            | 0.072                                | 4.732733                    | 0.0472                 |
|       | 2.3            | 0.088                                | 3.2168                      | 0.0239                 |
|       | 3.5            | 0.06401                              | 0.85708                     | 0.03692                |
|       | 4.8            | 0.02808                              | 0.45                        | 0.011792               |
| 07    | 1.23           | 0.004                                |                             |                        |
|       | 1.59           | 0.025                                |                             |                        |
|       | 1.8            | 0.102                                |                             |                        |
|       | 2.3            | 0.108                                |                             |                        |
|       | 3.5            | 0.132                                |                             |                        |
|       | 4.8            | 0.069                                |                             | 0.1234                 |

experimentally, it is common practice to assume lower and upper emissivity bounds and use these bounds to calculate upper and lower bounds for the temperature. This has been done for experiments #06 and 07 (where a sample radiance $L_{sample}$ was obtained) with the following lower $\epsilon_l$ and upper $\epsilon_u$ emissivity bounds:

$\lambda=1.23$ µm: $\epsilon_l=0.2$, $\epsilon_u=1.0$; $\lambda=1.59$ µm: $\epsilon_l=0.15$, $\epsilon_u=0.8$; $\lambda=1.80$ µm: $\epsilon_l=0.12$, $\epsilon_u=0.8$; $\lambda=2.30$ µm: $\epsilon_l=0.1$, $\epsilon_u=0.6$; $\lambda=3.50$ µm: $\epsilon_l=0.08$, $\epsilon_u=0.4$; $\lambda=4.80$ µm: $\epsilon_l=0.05, \epsilon_u=0.3$. These estimates are based on the room temperature values (obtained from the optical properties) for polished surfaces and the melting temperature values at ambient pressures. The calculated true temperature bounds for experiment #06 and 07 as function of wavelength are shown in figure 5 (for readability reasons the wavelengths for experiment #06 have been shifted by 0.1 µm to the red).

It can be seen that the uncertainty due to unknown emissivity is far higher for longer wavelengths, as expected. (For an explanation see reference 21.) The temperature ranges for all wavelengths overlap between 722 K and 802 K; this is the range where the temperature of the sample shocked to a Hugoniot pressure of 63.9±2.4 GPa and released into LiF (at an interface pressure of 27.1±1.0 GPa) is determined. The Hugoniot pressures in the sample have been obtained from the measured particle velocity at the sample window interface, the pressures in the LiF window have been obtained from jump conditions and known shock properties of the window material, through the published EOS and strength model.

The temperature of the LiF window as function of wavelength from experiment #06 is shown in figure 5. If the window is at a homogenous temperature and no background light occurs, the measured window temperature should be independent of wavelength. The increase of the measured window temperature (apparent temperature) with decreasing wavelength indicates a non-homogenous window temperature. Fitting the results with a simple two temperature model (for details see reference 20) gives a temperature of the window of 624±112 K with hot cells at a temperature of 1945±210 K. The area fraction (as seen from the pyrometer) of these hot cells was determined to be 3.0±1.5%, corresponding to a
volume fraction of less than 0.5%. The absorption coefficient as function of wavelength for the shocked LiF window is shown in figure 7. The solid line shows data from reference 22 at room temperature and ambient pressure. It can be seen that the absorption coefficient increases by roughly two orders of magnitude compared to the literature data independent of wavelength.

It can be seen from figure 3 that the radiance increases as long as the thickness of the shocked layer in LiF is increasing (from shock breakout at the Mo-LiF interface at 1.0 μs until the release wave from the back of the aluminum flyer reaches this interface at about 2.1 μs). The thickness of the shocked LiF layer decreases after t=2.1 μs because the release wave propagates at a higher velocity than the shock wave and hence the measured radiation decreases between t=2.1 μs to about 3.7 μs. At 3.7 μs the shock front in the LiF reaches the free surface of the window, and the measured radiance increases abruptly, which is assumed to be due to non thermal light emission when the window fractures on release from the shocked state and possibly the air shock.

From fitting the spectral radiance traces at different wavelengths and assuming lower and upper bounds for the effective emissivity of the sample/window assembly, a sample temperature (shocked to 63.9±2.4 GPa and released to a pressure of 27.1±1.0 GPa into LiF) at the interface of 762±40 K could be obtained. The release temperature into LiF was calculated to be 670±25 K using the Steinberg-Guinan strength model to predict the contribution to heating from plastic work 25. The resulting temperatures are in relatively good agreement 22 with experiments performed on a gas gun at a slightly lower pressure (T=683±41 K at a Hugoniot pressure of 58.7 GPa, with a release pressure of 24.8 GPa into LiF). A summary of all experimental and calculated results can be seen in figure 8.

The measured temperature in the LiF window is 624±112 K with hot spots (3.0±1.5% in area) with a temperature of 1945±210 K. These hot spots are believed to occur due to the slightly non uniaxial loading of the sample and window due to the dishing of the flyer plate which most likely causes local heating by shear strain localization. This localized heating in LiF windows was also observed in infrared imaging experiments 26. The temperature of the LiF window was calculated to be 569±15 K 25 which is in excellent agreement with the temperature inferred from the experiment. An analysis of the free surface experiments was not possible because of temperature non uniformities combined with background light caused by the hot APIEZON Qc⃝ (used to block thermal emission from hot jets from the edges, as discussed above).

V. CONCLUSIONS

Reliable pyrometric temperatures were obtained at shocked and released surfaces despite an ϑ(100%) background from shocked window material. This improvement was achieved by using the emission spectrum to infer a volume fraction of hotspots in the window, and was supported by relating changes in the emitted radiance to predictions of hydrodynamic events in the window. The measured temperature for Mo at a Hugoniot pressure of 63.9±2.4 GPa released into LiF to an interface pressure of 27.1±1.0 GPa of 762±40 K is in good agreement with temperatures measured using a powder gun at a slightly lower pressure (58.7 GPa released to 24.8 GPa) of 683±41 K. The measured temperatures are slightly higher than
calculations using the Steinberg-Guinan model \[19\] to take strength effects into account. It is not clear whether the remaining discrepancy is caused by additional sources of thermal emission not taken into account, or inaccuracy in the models used in the simulations. However, this level of agreement is unusually good for thermal emission from shocked metals.

Besides the release temperature of Mo, the shock temperature of LiF at a pressure of 27.1±1.0 GPa was also obtained and is in good agreement with calculations. The shock temperature in the window was much less sensitive to the strength model.

Because of problems controlling the thermal background, free surface temperatures could not be extracted from the experiments without a window.

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29. One injection fiber in the center surrounded by seven receiving fibers each 100 μm in diameter
30. At a pressure 200kbar in the anvil the reflectivity R2 is less than 0.1%, it increases to 0.25% at 500kbar and to 0.5% at 1Mbar.
31. The normal spectral emissivity for a flat surface is obtained by: \( \epsilon_\perp = \frac{4n_1n_2}{(n_1+n_2)^2} + k_2 \) with n1 and n2 index of refraction of the anvil and the sample and k2 the extinction coefficient of the sample
32. For both calculations (with and without strength) the temperature at 24.8 GPa is about 55 K lower than the one at 27.1 GPa. The measured data point at 24.8 GPa is 79±35 K lower than the 27.1 GPa one.