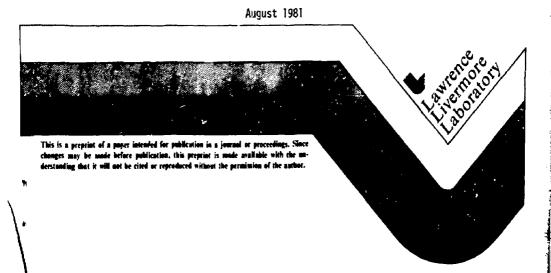
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A Streaked X-Ray-Transmission-Grating Spectrometer

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#### A Streakeu, X-Ray-Transmission-Grating Spectrometer\*

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#### Abstract

A free standing x-ray transmission grating has been coupled with a soft x-ray streak camera to produce a time resolved x-ray spectrometer. The instrument has a temporal resolution of  $\sim 20$  psec, is capable of covering a broad spectral range, 2-120 Å, has high sensitivity, and is simple to use requiring no complex alignment procedure. In recent laser fusion experiments the spectrometer successfully recorded time resolved spectra over the range 10-120 Å with a spectral resolving power,  $\lambda/\Delta\lambda$  of 4-50, limited primarily by source size and collimation effects.

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A free standing x-ray transmission grating is a linear, periodic grid of gold wires or bars with sub-micron spatial period. The grating structure is supported by a coarse grid orthogonal to the grating lines. Because of its simple geometry, the TG is a uniquely versatile spectroscopic dispersion element. It accepts radiation at all angles of incidence, thereby requiring no detailed angular alignment in its operation. It can, therefore, be easily coupled to instruments of high temporal resolution. In addition the TG's have high efficiency, and provide continuous spectra with moderate resolution over a broad spectral range.

Fig. 1 shows a simple x-ray transmission grating spectrometer viewing a small laboratory source. In such a configuration the spectral resolution (limited by source size and degree of collimation) is given by

$$\Delta \lambda = \frac{d}{L} (S + A) + \frac{d}{D} (A)$$
(1)

where d is the grating period, D is the grating to detector distance, L is the grating to source distance, S is the source size, and A is the collimation aperture opening. When

$$D \gg \frac{A}{(S+A)} L \tag{2}$$

the second term in Eq. (1) may be neglected. However, in many applications involving limited detector size (such as the slit of a streak camera) Eq. (2) cannot easily be satisfied. The spectral range of the spectrometer is typically limited by detector characteristics<sup>1</sup>. The maximum wavelength that can be recorded is

 $\lambda = \frac{d}{D}T$  (3)

where T is the linear size of the detector<sup>2</sup> (e.g. for a streak camera T would be the slit length). The minimum wavelength that can be distinctly recorded will be limited either by spatial resolution at the detector, or by source size and collimation (leading to overlap of the D.C. and first order spectral line). When detector spatial resolution

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limits  $\lambda_{\min}$ , then its value is given by Eq. (3) using the detector spatial resolution as the value for T. When source size and collimation limit  $\lambda_{\min}$  then  $\lambda_{\min} = \Delta \lambda$  given by Eq. (1).

A time integrated x-ray spectrum from a laser illuminated,  $SiO_2$  disk target is shown in Fig. 2. The spectrometer configuration of Fig. 1 was used with photographic x-ray film as the detector. In this configuration the spectrometer characteristics were:

d = 3000 m	S ≃ 100 µm
D = 63 cm	A = 100 µm
L = 74 cm	film resolution ≃ 10 µm

yielding a spectral resolution (Eq. (1)):

 $\Delta\lambda \approx 1.3 \text{ A}$ 

The spectral range was limited in this case at low energy by the carbon absorption of the polymer overcoat on the x-ray film, and at high energy by source size and collimation effects.

A free standing x-ray transmission grating has been coupled to a suft x-ray streak camera, and used to record the first time-resolved ( $\sim 20$  psec resolution), continuous, soft x-ray spectrum from a laser produced plasma. The experimental arrangement used is the same as that shown in Fig. 1 with the slit of the x-ray streak camera serving as the detector. Fig. 3 is a schematic representation of the experiment along with the data record from a gold disk target. In this time-resolved configuration the spectrometer characteristics were:

d =	3000 A	S ≃ 100 µm
0 =	31 cm	A = 100 µm
L =	76.6 cm	Streak camera resol. $\simeq$ 150 $\mu$ m

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The spectral resolution, limited primarily by collimation, was

## $\Delta \lambda \approx 2$ Å

The long wavelength limits to the spectral range were the finite slit length (1.3 cm), and absorption due to the 50  $\mu$ gm/cm<sup>2</sup> carbon window at the streak camera slit. In each case the long wavelength limit was around 120 A. The short wavelength limit, around 2 A, was established by collimation limitations, and the limited spatial resolution of the streak camera.

The time resolved TG spectrometer has been used in a number of different geometrical configurations (varying resolution and spectral range) to address the important issues of radiation production and plasma transport in laser fusion experiments. Fig. 4 shows time resolved spectral data recorded from a laser irradiated Al-Ti disk target. In this case the directly irradiated aluminimum disk was surrounded by an unirradiated annular ring of titanium. The difference in temporal characteristics (e.g. turn-on time) between the Al-K-lines and Ti-L-band provides important insight into radial plasma transport phenomena.

#### CONCLUSION

A marriage of two powerful technologies has provided a new spectroscopic capability. An x-ray transmission grating has been coupled with an x-ray streak camera to provide time resolved ( $\infty$  20 psec), continuous x-ray spectra over a broad spectral range. This new spectroscopic tool offers insight to the key issues of radiation production and plasma transport in laser fusion experiments.

#### ACKNOWLEDGEMENTS

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and a number of groups at Lawrence Livermore National Laboratory. The authors wish to acknowledge the skilled contributions of their colleagues both at MIT and LLNL: G. Howe, G.F. Stone, E.M. Campbell, R. Turner, D. Ciarlo, H.I. Smith, J. Melngailis, J. Carter, H. Medecki, and E. Pierce.

- 1. The other possible limit to spectral range is the transparency of the grating material at high x-ray energy. In this case, using gold gratings greater than 0.5 µm thick, grating transparency is not the limit to spectral range. Streak camera characteristics: finite slit length, finite spatial resolution at the slit, reduced sensitivity at high x-ray energy, and low energy x-ray absorption by the carbon window, provide the limiting factors in time resolved applications.
- 2. We have in practice not found spectral overlap with higher orders to ue a limit to  $\lambda_{max}$  in our laser fusion experiments. This is in large part due to the reduction in diffraction efficiency in higher order.

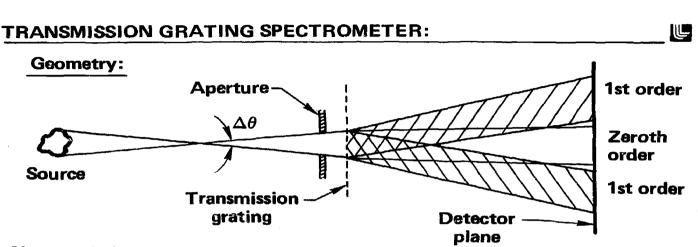
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## Figure Captions

- Fig. 1 A simple x-ray transmission grating spectrometer design.
- Fig. 2 A time integrated x-ray spectrum from a laser produced plasma, recorded by a transmission grating spectrometer.
- Fig. 3 A time-resolved transmission grating spectrometer.
- Fig. 4 Time resolved x-ray spectrum from a "plasma transport" laser target.



Characteristics:

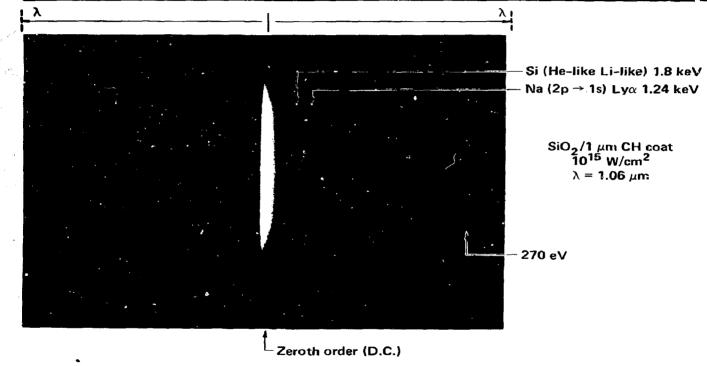
• Spectral resolution: limited by source size and collimation

$$\frac{\Delta\lambda}{\lambda} = \Delta\theta \frac{\mathbf{d}}{\lambda}$$

• Spectral range: limited by transparency of grating material at high energy. (e.g.)  $R\lambda \sim (2-3000)$  Å

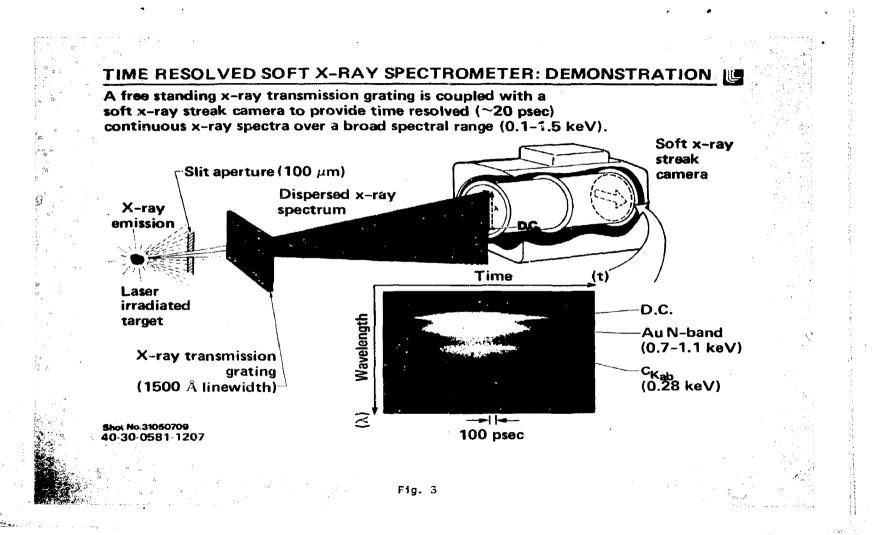
• Sensitivity: direct trade-off between  $d\Omega$  and  $\Delta\lambda$ 40-90-0681-1579

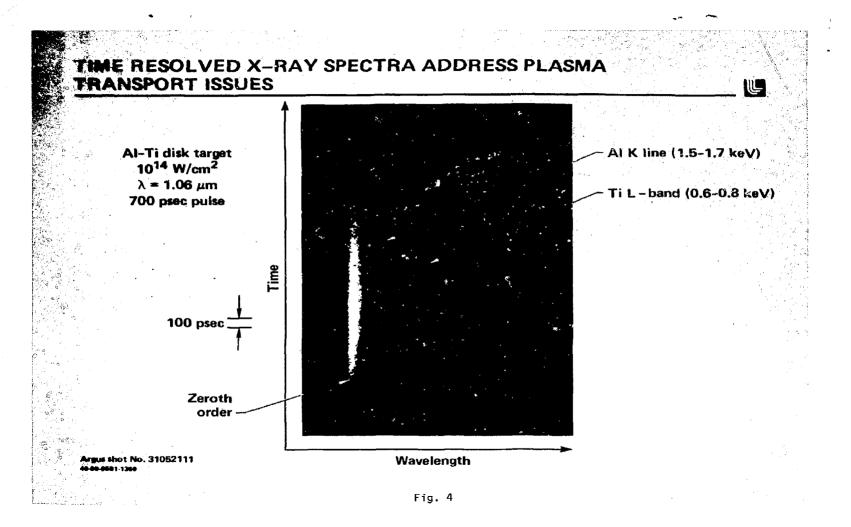
# X-RAY TRANSMISSION GRATING SPECTRUM FROM LASER PRODUCED PLASMA



Arg#a shot No. 31040606 40-90-0481-1132

Fig. 2





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