

MASTER

CONF-761204-1

LA-UR -76-2301

TITLE: OBSERVATION OF COHERENT RESONANCE FLUORESCENCE IN HOT CO₂

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SUBMITTED TO: American Physical Society Meeting at Stanford,
California

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INTRODUCTION

Hot CO₂ has often been considered as an isolator for large, short-pulse CO₂ laser systems because of the hot CO₂ line-to-line absorption coincidences with the CO₂ laser transitions. Schappert¹ first showed that in the absence of beam diameter reduction, hot CO₂ was unsuitable because of its large saturation flux. Feldman, et.al.,² showed that hot CO₂ is a potentially attractive baseline precursor suppressor because of the difference in the bandwidth of the undesirable baseline and the desirable short pulse. Feldman subsequently showed³ that by reducing the beam diameter in the hot CO₂ (and therefore raising the intensity) the effective saturation parameter of hot CO₂ could be reduced so that it would simultaneously behave as a practical saturable absorber and as a pulse bandwidth-discriminating baseline suppressor.

ONE NANOSECOND PULSE RESULTS

Although hot CO₂ was originally considered for quarter-nanosecond systems^{2,3}, we have elected to first check our theoretical understanding by using the recently available one-nanosecond pulses from the Gigawatt Test Facility (GWTF) in TSL-85. The results reported here are in relatively good agreement with theory. The experimental setup (as shown in Fig. 1) is as follows: a one nanosecond duration 1/2 millijoule switched-out pulse from the smoothing-tube stabilized GWTF oscillator was directed through two heated pyrex cells filled with CO₂ and fitted with NaCl windows at Brewster's angle. The cells were 190 cm long of

which 160 cm were heated to $300 \pm 10^\circ$ C for a total heated path of 320 cm. Temperatures were measured with three chromel-alumel thermocouples on each cell. The 1 ns pulse clipper consisted of a CdTe crystal (8 mm x 8 mm x 4 cm) switched by a 13 - 16 kv pulse from a laser-triggered spark gap. The resulting pulses were detected by a Molectron P5-00 pyroelectric detector, and signals were recorded on a Tektronix 7904 oscilloscope.

Figure 2 shows typical results. Note the growth of the second peak as the CO₂ pressure is increased. Even at zero torr hot CO₂ there is a second peak; this is merely ringing in the 7904. This ringing was subtracted from each picture before evaluating the ringing due to the hot CO₂. Figure 3 shows the measured corrected ratio of the second peak to the first peak as a function of hot CO₂ pressure. This figure indicates what has already been known; that in the linear regime one must use low CO₂ pressures to reduce the undesired secondary ringing. Unfortunately, at these reduced pressures the baseline rejection will be only slight since $\alpha_0 l$, the absorption coefficient times cell length, is small (0.554/torr up to 5 torr). The calculated baseline reduction factor ($e^{-\alpha_0 l}$) is shown in Fig. 4. Note that adequate baseline reduction is not available at low pressures. This is why hot CO₂ operating as a linear filter was not proposed for one nanosecond systems; good baseline rejection and lack of ringing are incompatible for one nanosecond pulse durations.

Saturation of either the hot CO₂ or the subsequent amplifier chain will reduce the secondary ringing. Although the pulse did not have enough energy to saturate the entire sample length of CO₂, we telescoped the beam diameter down to see if any evidence of hot CO₂ saturation could be obtained; our results showed only a slight reduction of the second peak. The testing of saturated operation of hot CO₂ at one nanosecond will be addressed in a future experiment.

170 PICOSECOND RESULTS - PULSE COMPRESSION IN SF₆

To test the effects of hot CO₂ on pulses shorter than one nanosecond, we were first tempted to use the ultrashort pulses generated

by optical Free-Induction Decay⁴ (an effect also obtained in hot CO₂), but these FID pulses are characteristically different from the electro-optically switched-out pulses. The FID pulses have a narrow spectral notch missing from an otherwise broad spectrum, and there is post-pulse temporal ringing out of phase with the main lobe. In our attempt to obtain a shorter pulse we have tried to compress a one nanosecond switched-out pulse by saturating SF₆ in a short cell. The beam was telescoped down to a 3 mm diameter (corresponding to ~ 5 MW/cm² in the one nanosecond pulse) and was passed through a 3.3 cm-long cell of SF₆ heated to 120°. Unfortunately, at this time, all Molectron detectors had been unable to meet the sensitivity originally specified, and all had been returned to the factory for repair. A Rofin photon-drag detector with slightly different ringing characteristics had to be used. After passage through the SF₆; the pulse passed through the hot CO₂ cells and then through amplifiers 1 and 2. The amplifiers were needed because of the attenuation in the SF₆ ($\sim 90\%$). Fig. 5 shows typical results obtained with 10 torr of hot CO₂ in the cells. As the pressure of the SF₆ was increased, the hot CO₂-induced ringing was reduced. Several data points were taken at each of a set of SF₆ pressures; the averaged results are graphed in Fig. 6. Again, detector and scope ringing was subtracted prior to obtaining this figure.

Although this evidence is preliminary, it is consistent with the explanation that at 18 torr, the SF₆ cell reduced the pulse duration by a factor of 6; this would have reduced the hot CO₂-induced ringing by just that same factor. In our numerical simulations in 3.5 meters of hot CO₂, the ratio of the second peak to the first is approximately $0.61 T_p/T_2$, where T_p is the $\text{FW}_{\frac{1}{e}}$ M duration of the pulse and T_2 is the homogeneous lifetime of the absorbing transition. There are, of course, other explanations which do not require SF₆ induced pulse shortening; these other possibilities must be eliminated before we can be assured that shortening does occur. We intend to carefully measure the shorten-

ing effect with the new 5 GHz scope and fast detectors.

All results obtained thus far have been free of any conflicts with our theoretical models of short pulse propagation in resonant absorbers.

REFERENCES:

1. G. T. Schappert, LASL Memorandum: L-1/75-1393, September 1975.
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3. B. J. Feldman, LASL Memorandum: L-2/76-45, January 1976.
4. B. J. Feldman, R. A. Fisher, E. J. McLellan, and S. J. Thomas, LA-UR 75-2321, Presented at IXth International Quantum Electronics Conference, Amsterdam, The Netherlands, June (1976), also Opt. Commun. 18, 72 (1976).

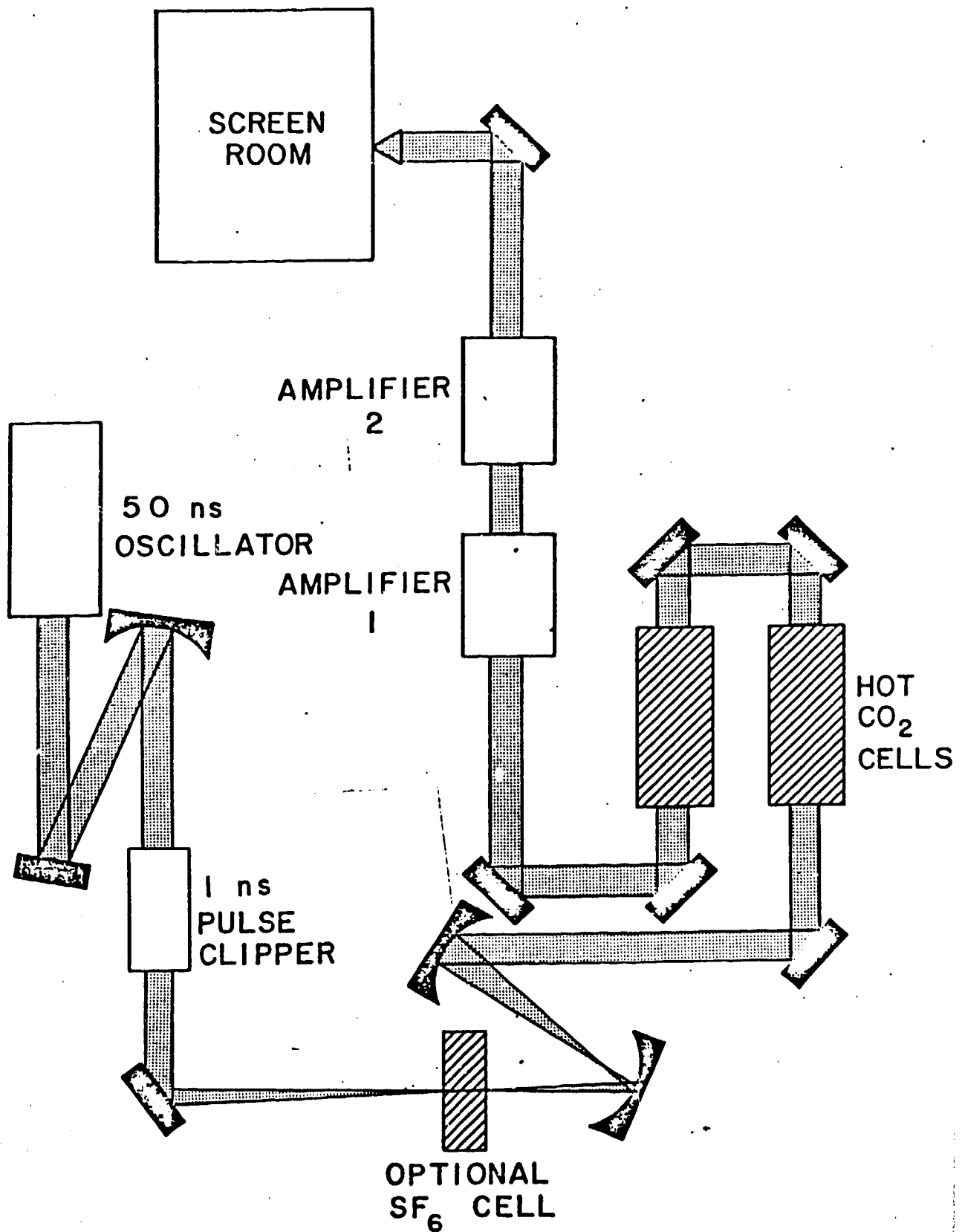


Fig. 1

Adaption of the Gigawatt Test Facility (GWF) for Experiments - Normally the SF₆ Cell and Hot CO₂ Cells are Missing.

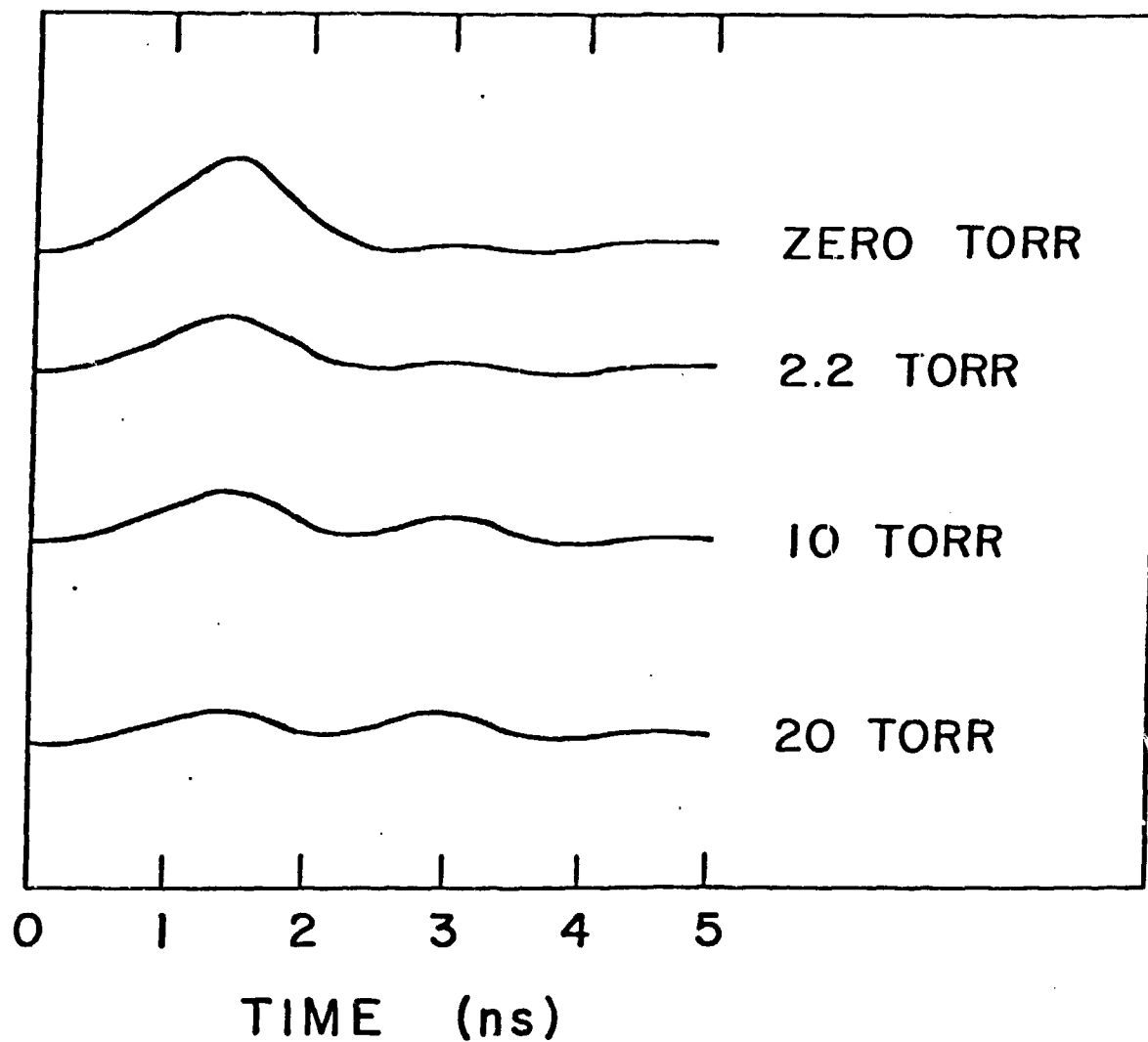


Fig. 2

Artists' Reconstruction of Oscilloscrams Taken to Study Ringing Introduced by Short Pulse Transmission in Hot CO_2 . The Hot CO_2 Pressures Corresponding to Each Shot are Indicated in the Figure. The Cell Temperature is 2000 K.

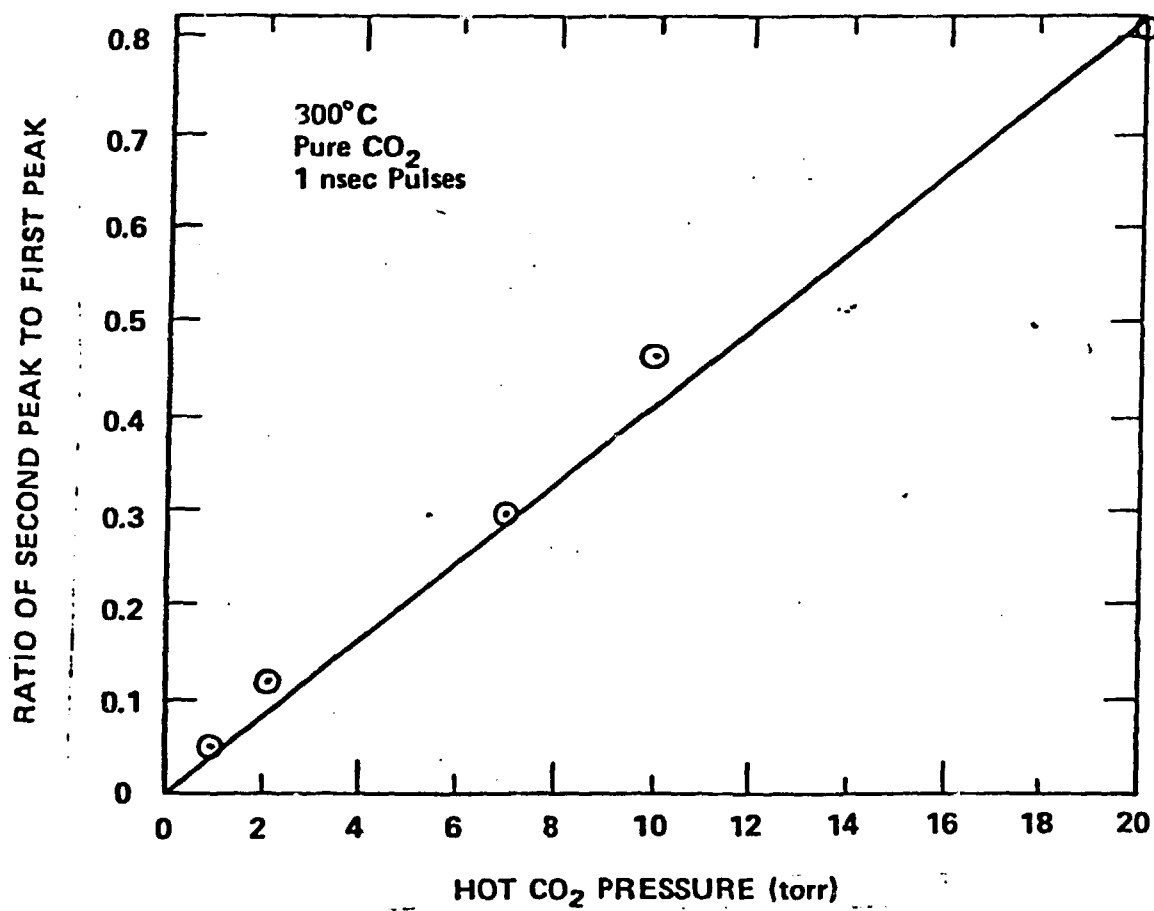


Fig. 3

Measured Fractional Height of Second Peak After Propagation Through the Hot CO₂ Cells. The Ringing in the Oscilloscope (as in the zero torr trace in Fig. 2) Has Been Subtracted

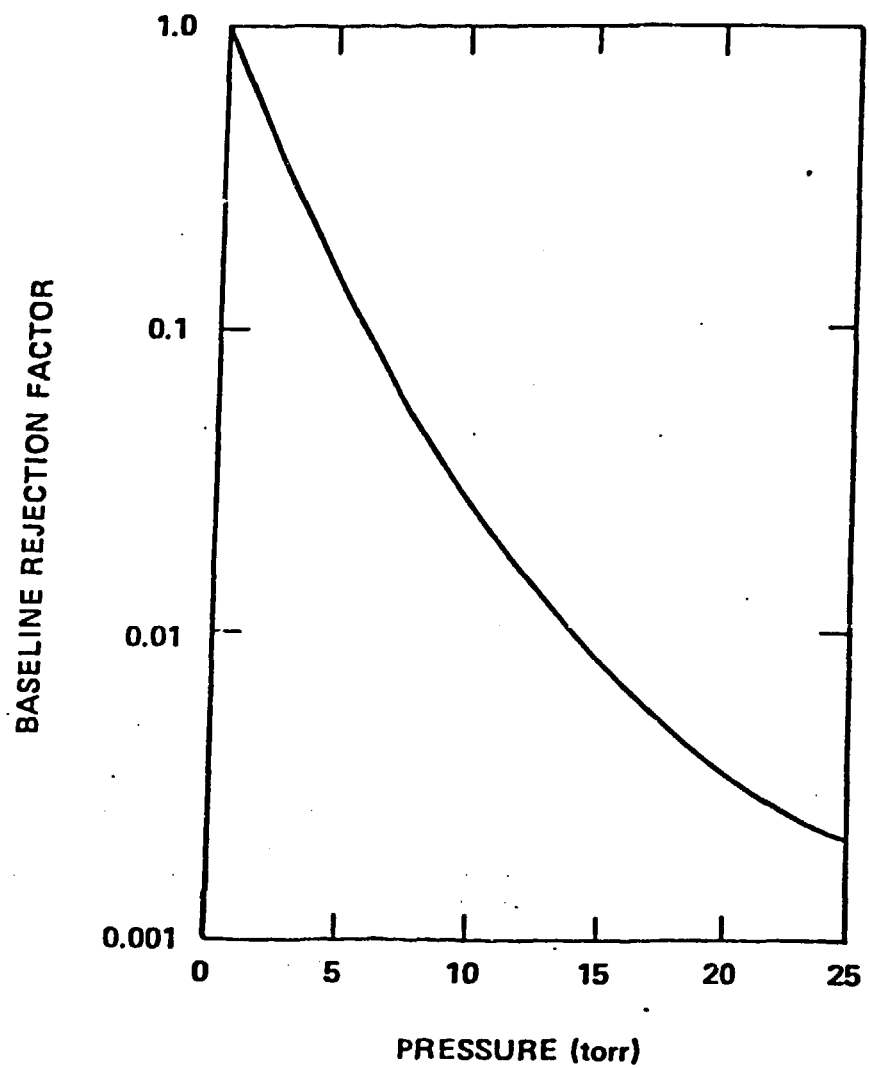


Fig. 4
Calculated Baseline Reduction Factor ($e^{-\alpha_0 l}$) Versus Pressure
in the Hot CO₂ Cells

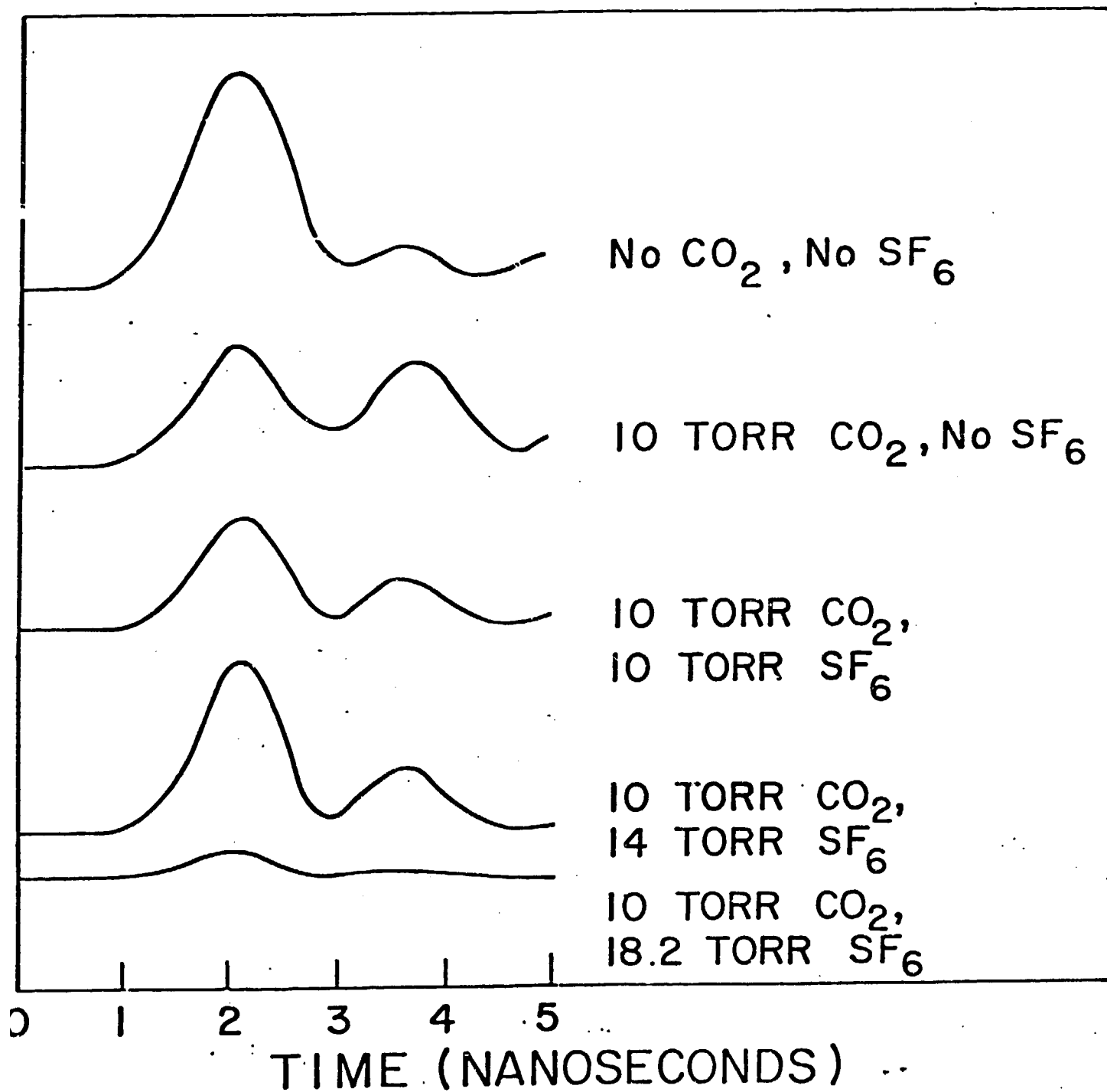


Fig. 5

Artists' Reconstruction of Oscillograms Showing the Reduced Ringing in the Hot CO_2 When the Pressure in the Preceding 3.3 cm SF_6 Cell is Increased - The Vertical Scales are Different on Each of These Traces

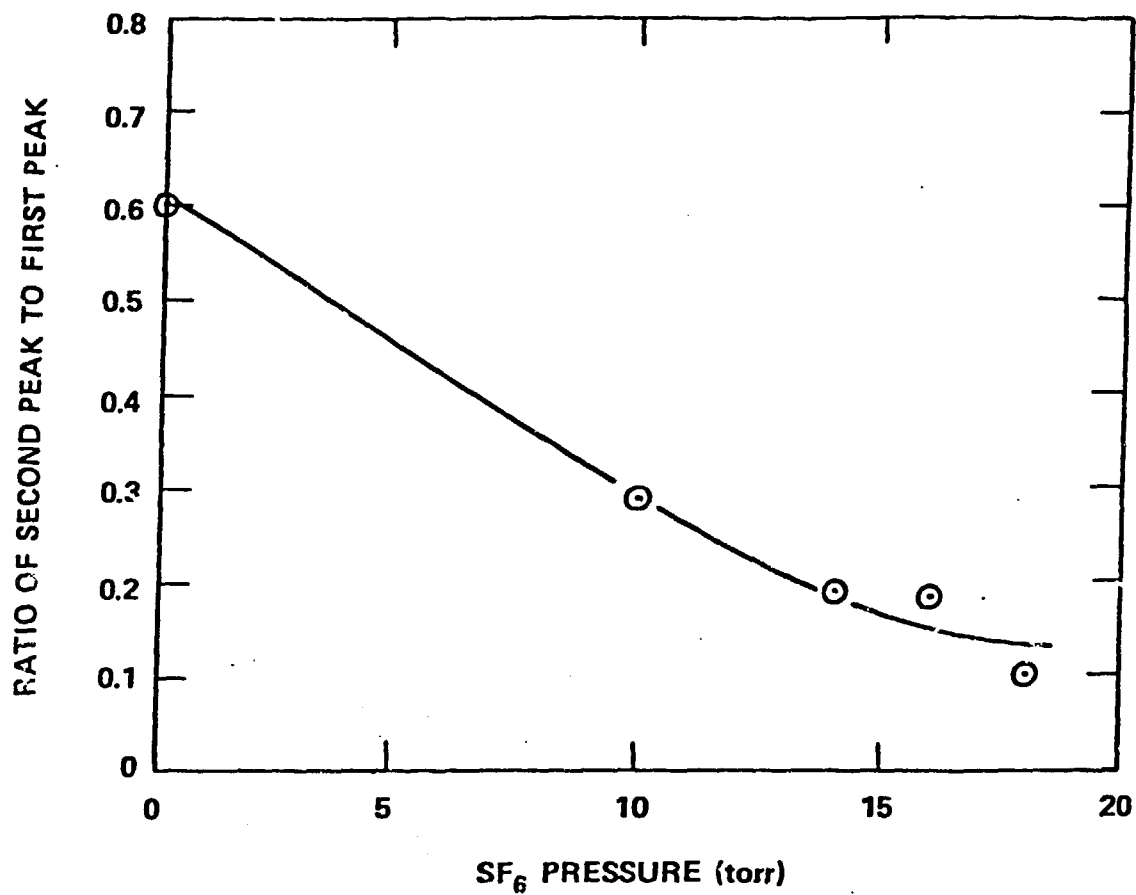


Fig. 6

Measured Ratio of Hot CO₂-Induced Secondary Peak as a Function of Pressure in the 3.3-cm SF₆ Cell (T = 49° C)