

be thought of as using interference between left and right circularly polarized waves. It could also be used for labeling a particular lower level, so that all of its absorption lines appear as bright lines in the probe output. Moreover, it can be equipped with a servo adjustment to keep the phase difference between the beams always zero. Then the probe intensity is proportional to the square of the difference in absorption of the two beams, and so the linewidth is reduced.

A particularly simple interferometer velocimeter for continuous wave lasers was described by Kowalski and Hawkins. It uses two-beam interferometer in which beams from the unknown laser and a standard laser traverse identical paths in opposite directions. The path length is changed by moving a corner cube reflector, and the wavelength is determined from the ratio of the number of fringes of the standard and unknown laser for a given displacement of the reflector. A higher accuracy, close to one part in a hundred million, is obtained by an improved method in which the reflector velocity is held constant. Then frequency counting techniques can be used, including fringe-frequency multiplication by a phase-locked oscillator. We are using this interferometer to study pressure shifts in molecular spectra at pressures low enough so that individual lines are clearly resolved. Also, in collaboration with J. E. M. Goldsmith, T. W. Schuch, and E. W. Eeber it is being used to improve the accuracy of the wavelength of the Lyman line of atomic hydrogen, which is the basis for determining the Rydberg con-

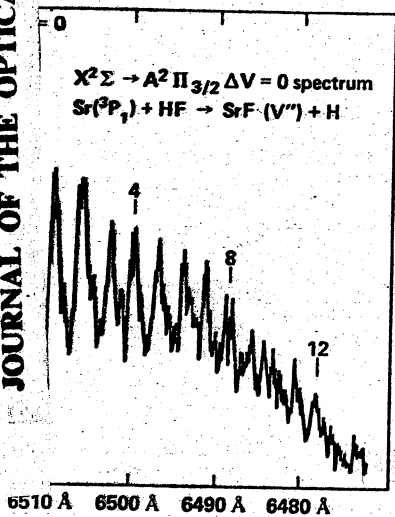


FIG. 1. Laser-induced fluorescence spectrum for SrF.

The polarization labeling technique provides a method of recognizing all absorption lines from a chosen lower level. It has been used with both pulsed and continuous-wave lasers. It has recently been applied to simplify the very complex spectrum of nitrogen dioxide and to identify sodium molecular lines in the A band where perturbations displace some of the lines from their expected positions.

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A.4. A Stable cw Waveguide CO₂ Laser for High-Resolution Saturation Spectroscopy. Application to the ν₃ Band of SF₆.* A. VANLERBERGHE, S. AVRILLIER, AND C. J. BORDE, *Laboratoire de Physique des Lasers, Université Paris-Nord, 93430 Villetaneuse, France*, AND C. D. CANTRELL, *Los Alamos Scientific Laboratory, University of California, Los Alamos, N. M. 87545*. (15 min.)

We have constructed a very stable waveguide CO₂ laser for high-resolution saturation spectroscopy. The square bore waveguide is made from BeO plates epoxied together and has ZnSe Brewster windows. The optical resonator consists of a spherical mirror at one end and a diffraction grating illuminated by a ZnSe lens at the other end. These elements are held in a rigid Invar mechanical structure. The tuning bandwidth of this laser is limited by the 600 MHz free spectral range of the resonator reduced to around 520 MHz by the anomalous index of refraction of the medium. The short-term frequency jitter estimated by

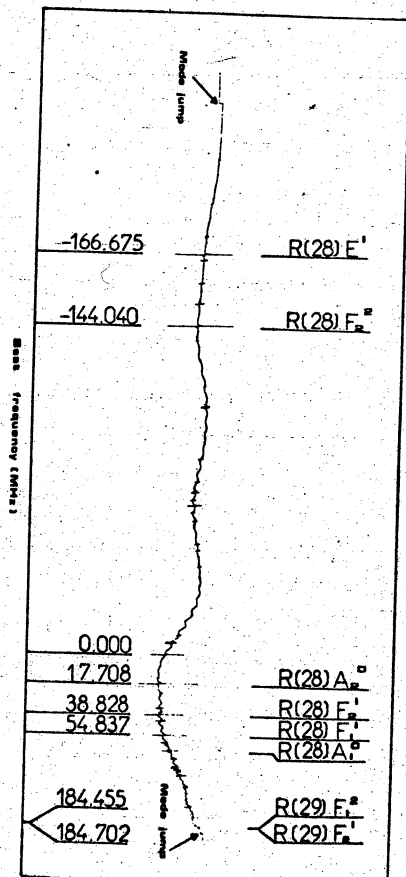


FIG. 1. Saturation spectrum of SF₆ for the P(14) line of the CO₂ laser at 949.4793 cm⁻¹.

beating this laser with a very stable conventional laser is less than 10 kHz. Many saturation spectra of various molecules have already been obtained with this new laser used as one of the sources in our saturation spectrometer. A conventional laser locked to a

saturation peak provides the reference frequency. The beam from both lasers are expanded by 30-cm-diam telescopes to reduce transit-time broadening and curvature-induced shifts. As an example, Fig. 1 shows the saturation spectrum of SF₆ obtained with the free-running laser for the P(14) transition of CO₂ centered at 949.4793 cm⁻¹. The observed peak-to-peak width is of the order of 20 kHz and will be discussed in detail. The frequencies of the main resonances have been measured by locking the waveguide laser at the corresponding line center and measuring the beat frequency with the reference laser locked on the SF₆ peak closest to the CO₂ line center. Similar measurements were performed for most SF₆ resonances that are within ±250 MHz of the P(12) to P(20) CO₂ line centers of the 10.4 μm band. A very good signal-to-noise ratio has enabled us to lock the waveguide laser on weak lines such as the P(4) Coriolis components of SF₆. A detailed interpretation of the spectra and the new set of constants for the ν₃ band of SF₆ derived from them will be given.

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A.5. Precision Laser Photodetachment Spectroscopy in Magnetic Fields.* W. A. M. BLUMBERG, R. M. JOYSON, AND D. J. LARSON, *Lyman Laboratory of Physics, Harvard University, Cambridge, Mass. 02138*. (15 min.)

We have studied the effect of a strong magnetic field on the photodetachment cross section for negative sulfur ions near the threshold for detachment from the ²P_{3/2} state. The magnetic field produces structure in the cross section which is periodic in the light frequency. This structure represents a dramatic departure from the Wigner law for threshold behavior which has been found to be valid at zero magnetic field.¹ The departure is found to be a result of the excitation of the detached electrons to discrete cyclotron levels.

The S⁻ ions are confined in a Penning² trap in magnetic fields ranging from 6 to 16 kG. The ions are illuminated by light of approximately 597 nm from a single-mode dye laser. The number of ions present after a pulse of light is measured by observing the currents induced on the trap electrodes while driving the axial motion of the ions. The data consists of measurements of the number of ions remaining as a function of laser frequency.

An example of data obtained at 15.7 kG is shown in Fig. 1 together with a predicted curve which has three variable parameters adjusted to give reasonable agreement with the data. These parameters are the frequency of the first threshold, an overall cross section scale factor, and the average velocity of the ions.

The basic element of the prediction is the assumption that the detached electrons are excited to discrete cyclotron levels. Since the transverse motion of the electron is quantized, it is left with only one degree of freedom and thus the density of final states is proportional to 1/k (where k is the relative momentum of the electron) rather than proportional to k as it is in the case of zero mag-