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INFRARED EMISSION SPECTROSCOPY OF LOW PRESSURE GASEOUS DISCHARGES

NO.

Peter Hansen Hajime Sakai John Strong

Astronomy Research Facility University of Massachusetts Amherst, MA 01003

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central electrode. It is powered by an AC supply. Power dissipation ranges from 1 to 4 kilowatts. Due to the discharge geometry, pressure is limited to the range of 0.1 to 0.6 torr; however, broad adjustments in current density are possible over this pressure range.

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This contract is (1) to create a discharge condition similar to those existing in the upper atmosphere, and (2) to study the infrared emission spectra of atomic and molecular nature which are generated from it. The work described in this scientific report represents our effort made during the first year of a multi-year program. Our principal scheme of experiment is to utilize the technique of Fourier spectroscopy for observation of the infrared emission generated in low pressure, long pathlength flow discharge. Since an insignificant amount of information is available in literature for the laboratory study of the infrared discharge emission, our main effort has been concentrated to develop the technique most adaptable to our need.

During this phase of work, we have succeeded to produce a very stable discharge condition which is essential for the spectrometry using the Fourier technique. We believe that the problem of generating the infrared emission spectra has been solved at least to our satisfaction. The recovered spectra show a good signal-to-noise ratio with a spectral resolution of 1.0 cm⁻¹, a figure currently achieved. The interferogram recording scheme used in the measurement is antiquated and preventing us from extending the maximum path difference much beyond the presently achieved figure of 1.0 cm⁻¹. Within the next few weeks, we will introduce a new recording scheme, expecting to improve the resolution to a figure of 0.1 cm⁻¹, the limit imposed by the interferometer currently used.

The scientific report which is contained in the following pages was presented at the 1977 Fall Meeting of the Optical Society of America held at Toronto, Canada.

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INTRODUCTION

This report covers work done in the first year of a multi-year contract to determine the infrared spectra of atoms and molecules under conditions similar to those existing in the upper atmosphere. Low pressure, long pathlength electronic flow discharges are generated in various gases and the resulting emission spectra are obtained by means of Fourier spectroscopy. The discharge facility and data handling system are described.

As evidence of the capability of the overall system, sample spectra are presented of some of the species (CO, CO₂, N₂, NI, OI) radiating in the 1800 to 7900 cm⁻¹ spectral region. Optimum resolution to date is $\Delta v = .96 \text{ cm}^{-1}$. This effort continues with the current objective of increasing the resolving power of the spectra, and a substantial increase of the flow rate of the discharge to decrease the effects of contaminants.

EXPERIMENTAL

The discharge column is contained in a steel vessel (Fig. 1) one meter in diameter and 33 meters long. A 12-meter-long polished aluminum electrode is centrally suspended in the midsection of this tank. This central electrode and the tank walls form the discharge configuration.

The discharge is powered by a 10 kw variable 60 Hz supply (Fig. 2). Due to the high power dissipation, the central electrode is water cooled. Typical operating conditions are 2.5 kw (V $\stackrel{\triangle}{=}$ 1000 VAC, I = 2.5 A) supplied to the discharge and 1 kw dissipated in the load. The discharge may be operated in the pressure range of 200 to 600 millitorr, depending on the gas under consideration. By variation of the pressure and load, stable operating conditions can be achieved over prolonged periods of time. As the pressure range is limited, current density is essentially the only independent variable affecting discharge conditions.

The optical system consists of two 1-meter diameter, 33-meter focal length spherical mirrors. The mirrors are focused on one another, resulting in three passes through the discharge column, for an equivalent optical path of 36 meters. The resulting f/33 beam is then collimated by

a KBr lens and enters the interferometer, an Idealab IF6 that has been modified for use in the 1 to 6 micron region. The interferometer uses a CaF beamsplitter and InSb and PbS cooled detectors. Its ultimate resolving power is near $\Delta\nu$ % 0.1 cm⁻¹. The path difference is monitored with a He-Ne laser.

There are two basic data handling systems presently available or near completion. For the spectra presented in this report, the interferograms were punched on paper tape and subsequently the data was placed on a CDC 6600 computer for the Fourier transformation and subsequent plot. This method, while workable, is extremely time-consuming (slow punch rate) and has some inherent limitations as to the number of data points that may be taken (16K) and the resolution of the A/D conversion (10 BITS). This system limits resolution to about Av 2.6 cm-1. Near completion now is the system outlined in Fig. 3. Here the radiation from the discharge column (already chopped at 120 Hz) enters the interferometer. The detector signal (interferogram) is synchronously amplified and passed to a 12 BIT A/D convertor locked to the interferometer's laser signal. The A/D output is passed through a parallel interface to a DEC LSI 11/03 minicomputer and stored on a floppy disk. The interferogram is then transmitted via a serial interface and a timesharing system to the CDC 6600 for processing.

RESULTS

Some earlier (1974-75) spectra of the discharge column were taken with an interferometer and data reduction system provided by Dr. Randall Murphy of AFGL. These are low resolution spectra (10 cm⁻¹ < $\Delta \nu$ < 50 cm⁻¹), as the operating time of the discharge column was limited due to the fact that the central electrode at the time was not water cooled, and the overall spectrometric system was set up for low resolution. Fig. 4 shows a nitrogen flow discharge spectrum in the PbS region. The molecular electronic band structure of the B³II \rightarrow A³E⁺ system is indicated according to the scheme of Saum and Benesch. Fig. 5 indicates the spectrum of an

oxygen flow discharge. The significant features here are atomic oxygen lines at 5549 cm⁻¹, 5485 cm⁻¹, 3773 cm⁻¹, 3617 cm⁻¹, and 3456 cm⁻¹. These have previously been identified by Saum and Benesch.² Both the nitrogen and oxygen spectra were compared to a black body spectrum at 2300° K. The spectral radiance values 10^{-3} to 10^{-5} W/cm²-SR-wavenumber appear to be typical of the signal strengths that are encountered.

Fig. 6 shows the spectrum of a nitrogen flow discharge. The conditions under which this spectrum was obtained were the following: The power supplied to the discharge was % 2 kw, i.e., 800 VAC at 2.5 amperes, and the pressure was 250 millitorr. It took 3 hours to take the spectrum; however, here the limiting factor was the speed of the paper tape punch. The flow rate was 1 l/min. The discharge acts as its own chopper (120 Hz) and thus the powerline frequency is used as a reference for synchronous rectification. The spectral range covered is $1800 \rightarrow 7900 \text{ cm}^{-1}$, the lower limit being the cutoff frequency of the InSb detector, while the upper limit is due to the sampling interval. Spectral resolution is $\Delta v = 1 \text{ cm}^{-1}$. Figs. 7 through 30 show this spectrum in more detail in increments of 250 cm⁻¹. It should be noted that the vertical gain is not the same for all the spectra.

As perhaps could be expected, the strongest spectral features are caused by impurities CO, CO₂, and possibly OH and NH. Electronic transitions in molecular and atomic nitrogen are also observed. From $1850~\rm cm^{-1}$ to $2150~\rm cm^{-1}$ the CO vibration rotation bands predominate. From $2290~\rm cm^{-1}$ to $2365~\rm cm^{-1}$ the $\rm V_3$ fundamental band of CO₂ is clearly defined (Fig. 9). In the $2500~\rm cm^{-1}$ to $3500~\rm cm^{-1}$ region several band systems have been schematically indicated. The wide spacing of the emission lines in these spectra (high rotational constant ${\rm \mathfrak{A}}$ 15 cm $^{-1}$) indicates vibration rotation bands of OH and NH. We feel that higher resolving power is desirable for complete analysis of this spectral region. Another CO₂ feature appears at $3700~\rm cm^{-1}$. From $3800~\rm cm^{-1}$ to $6600~\rm cm^{-1}$ there are some small features as yet not identified. Between $6600~\rm and$ $7700~\rm cm^{-1}$ we again have electronic transitions in molecular nitrogen. Also atomic nitrogen is observed at $7360~\rm cm^{-1}$ and $7444~\rm cm^{-1}$.

CONCLUSION

We have shown that during the first year of this contract we have established a low pressure, long pathlength gaseous discharge facility effectively simulating conditions in the upper atmosphere. The infrared radiant flux from the discharge column is sufficient to produce spectra with good S/N. With the new data handling system, and a faster pumping system, both under construction, we will be able to produce higher resolving power ($\Delta v \% .1 \text{ cm}^{-1}$) emission spectra of the atmospheric gases with lower impurity content.

REFERENCES

- 1. K.A. Saum and W.M. Benesch, Appl. Opt. 9, 195 (1970).
- 2. K.A. Saum and W.M. Benesch, Appl. Opt. 9, 1419 (1970).

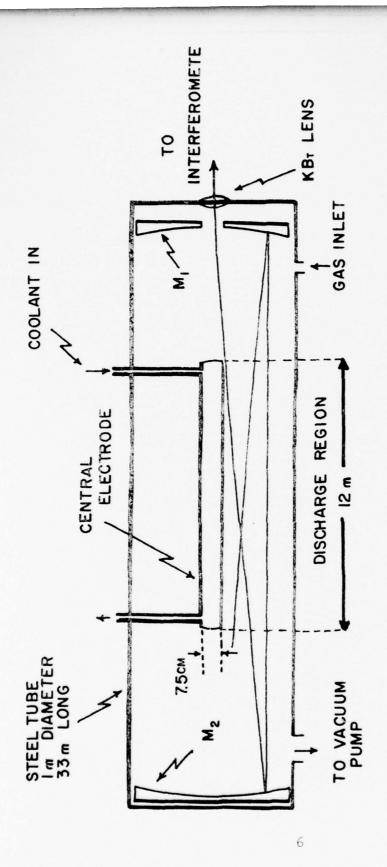


FIG. 1 SCHEMATIC REPRESENTATION OF THE DISCHARGE FOCUSED ON EACH OTHER RESULTING IN THREE COLUMN (not to scale). MIRRORS M, AND M2 ARE PASSES THROUGH THE DISCHARGE COLUMN.

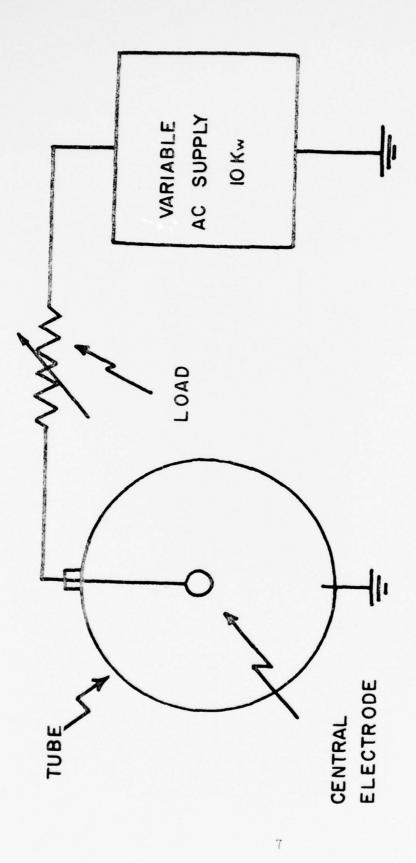


FIG. 2 END ON SCHEMATIC REPRESENTATION OF TUBE INDICATING BASIC ELECTRICAL CONNECTIONS.

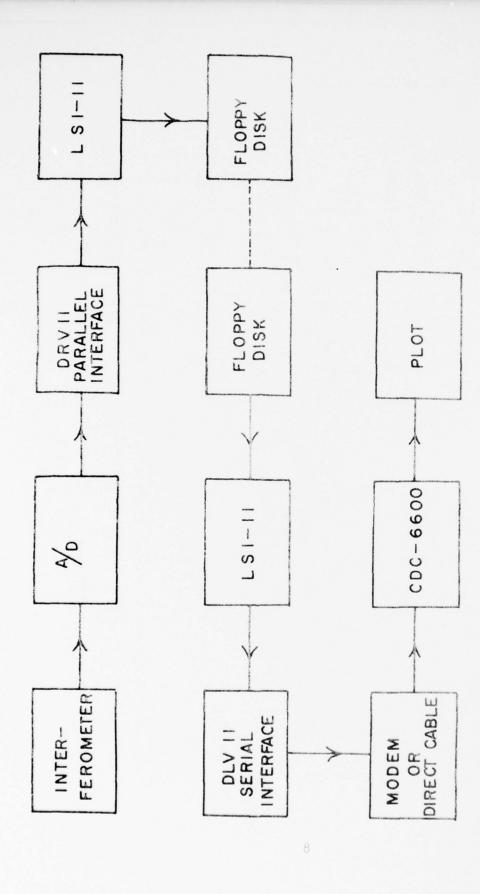


FIG. 3 DATA COLLECTION AND PROCESSING SCHEME

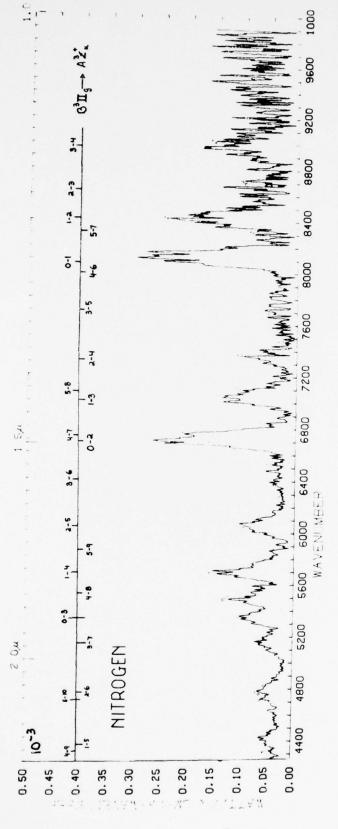
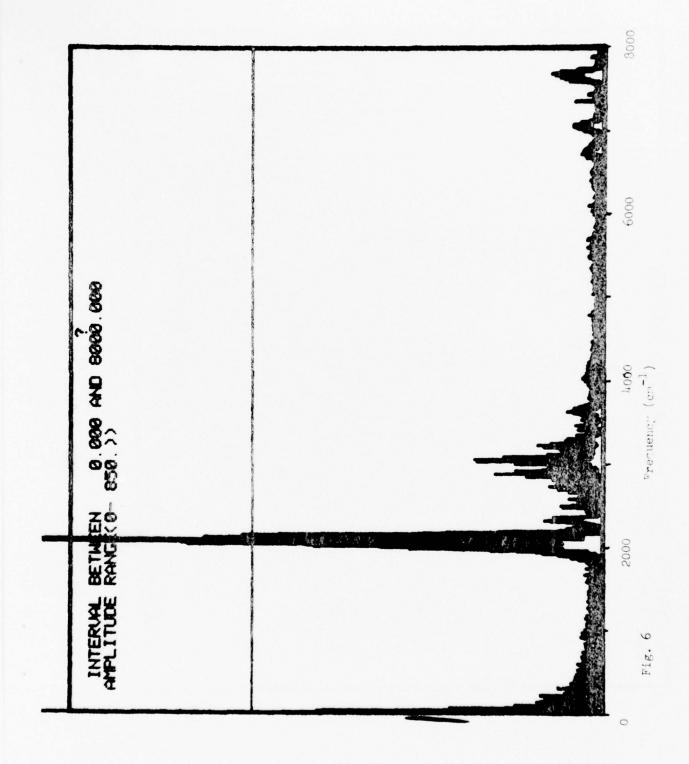
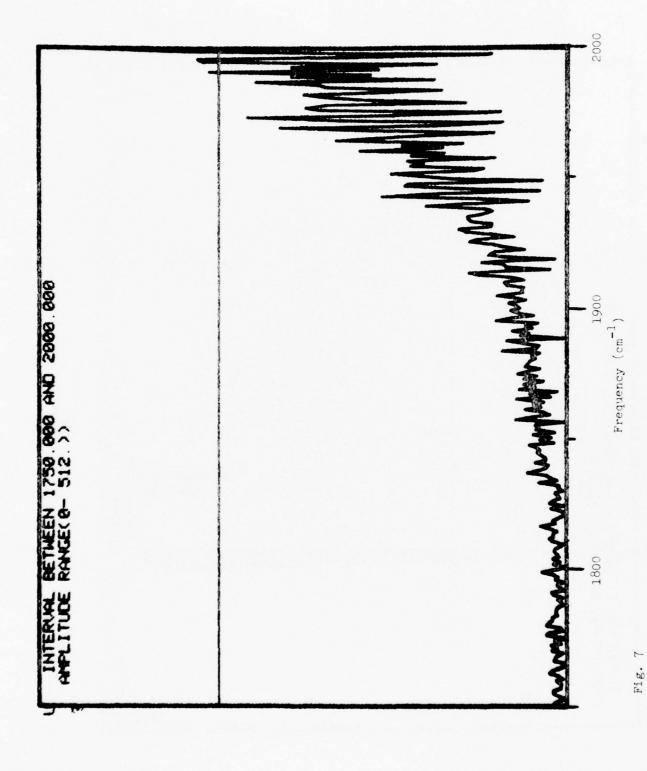


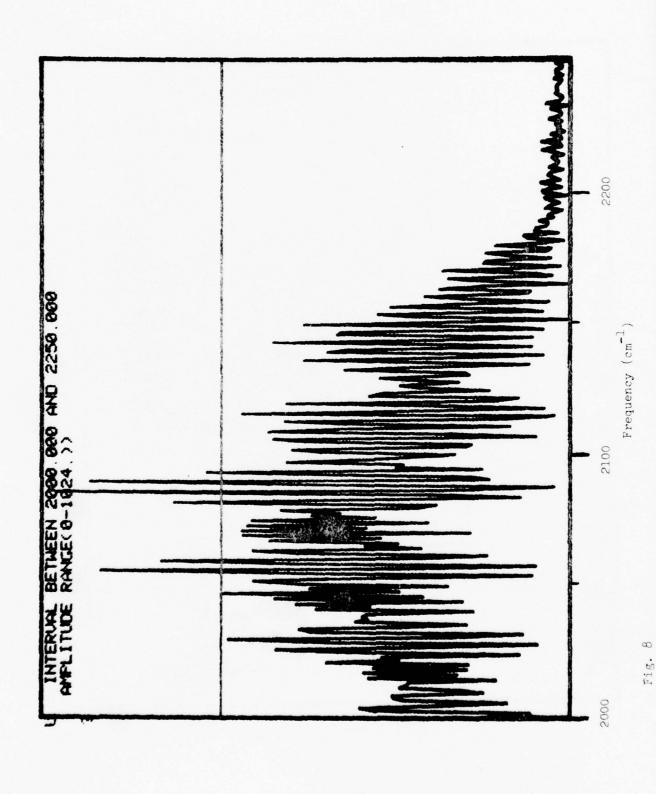
FIG. 4 EMISSION SPECTRUM OF A NITROGEN DISCHARGE

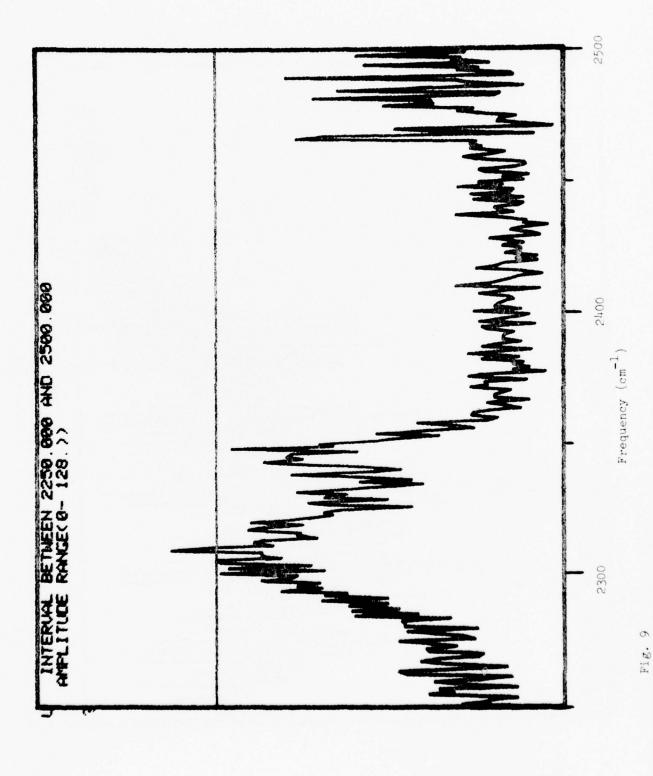


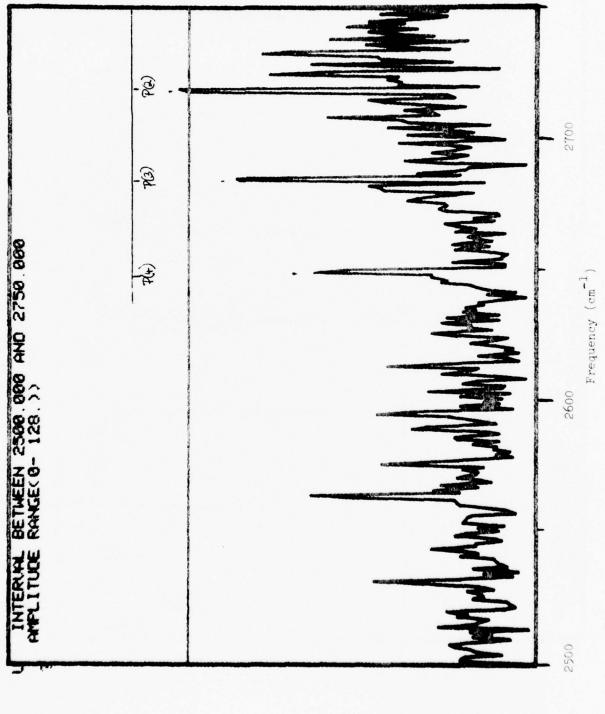
EMISSION SPECTRUM OF AN OXYGEN DISCHARGE FIG. 5











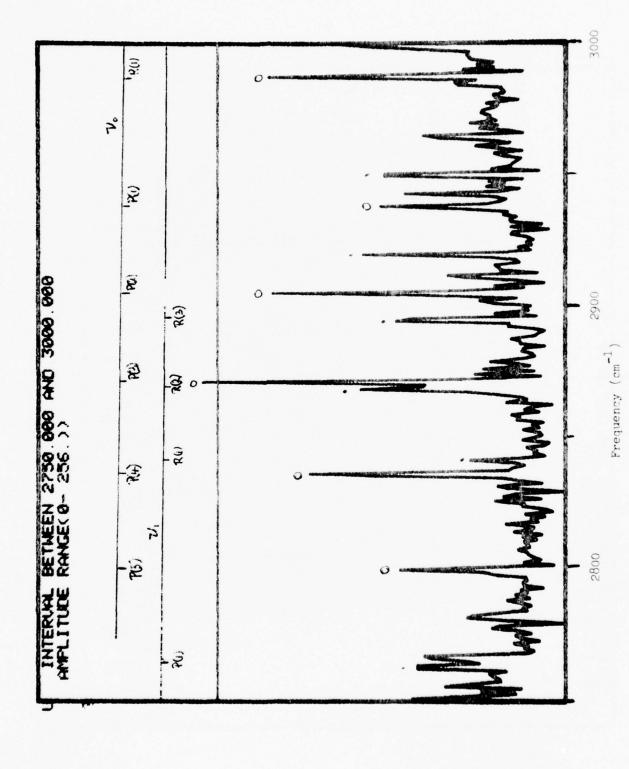


Fig. 11

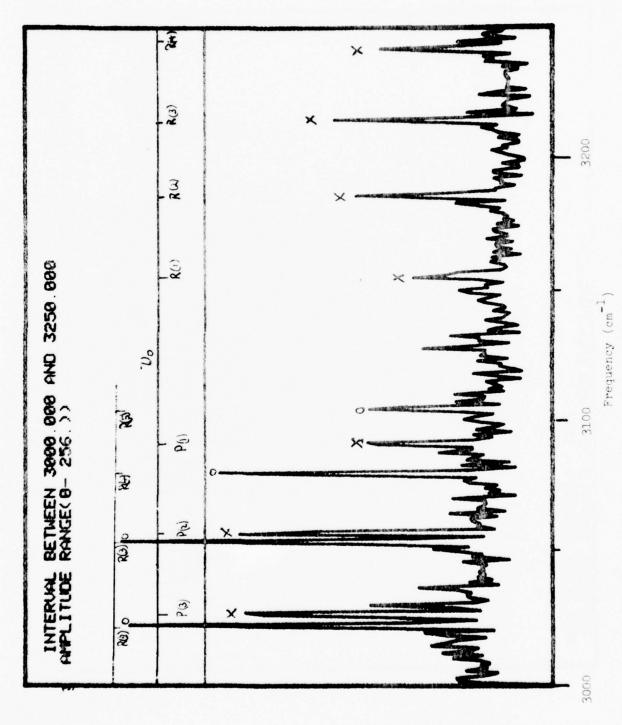


Fig. 12

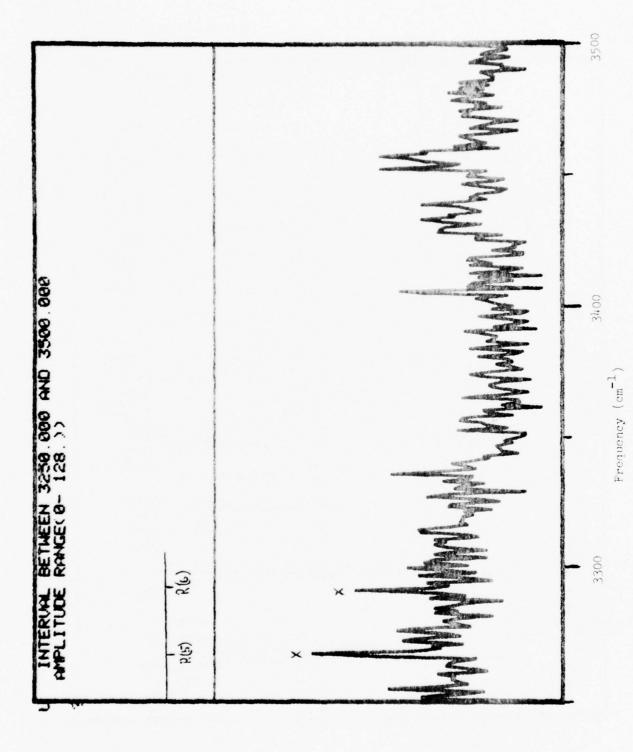


Fig. 13

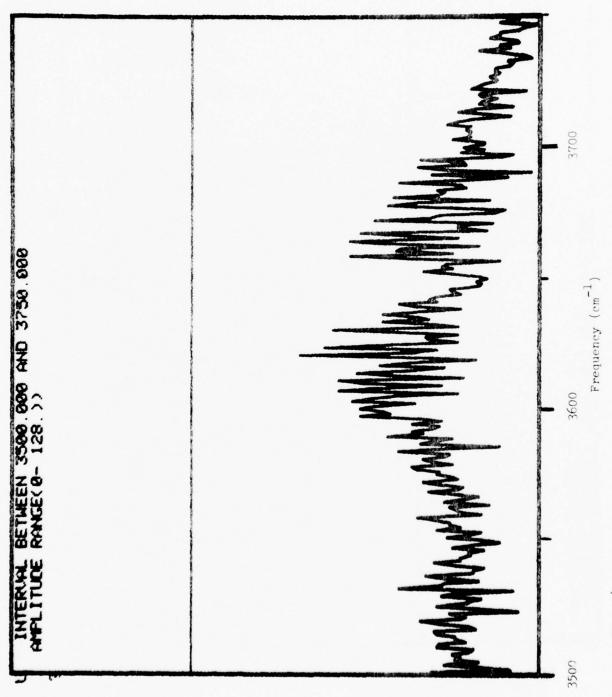
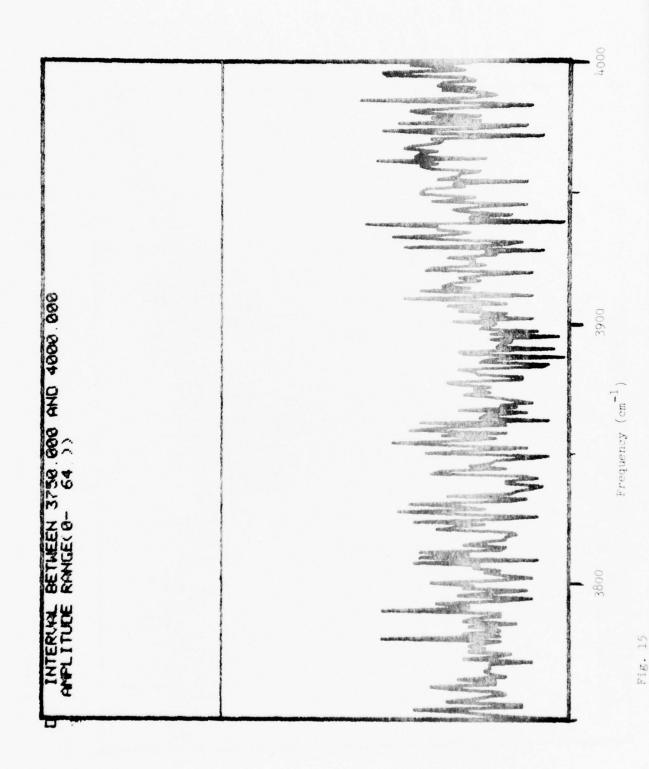
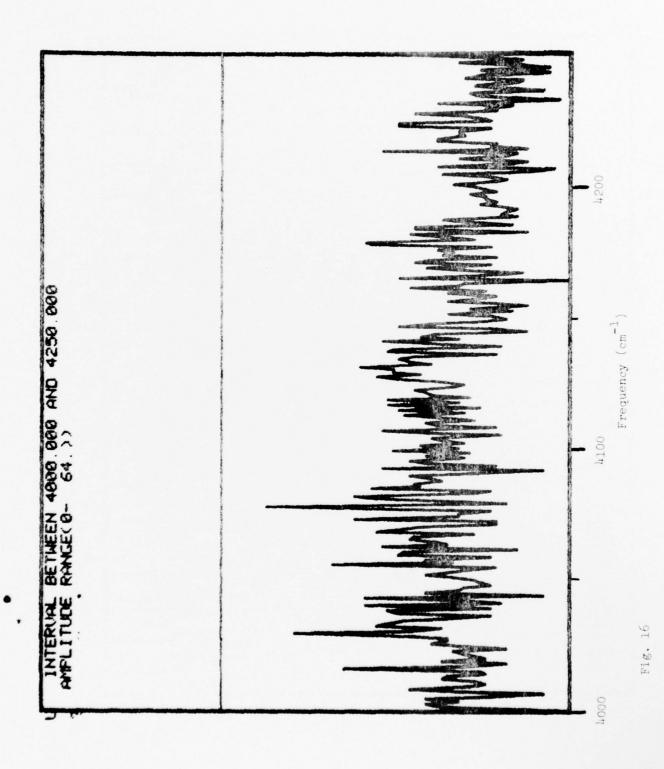


Fig. 14





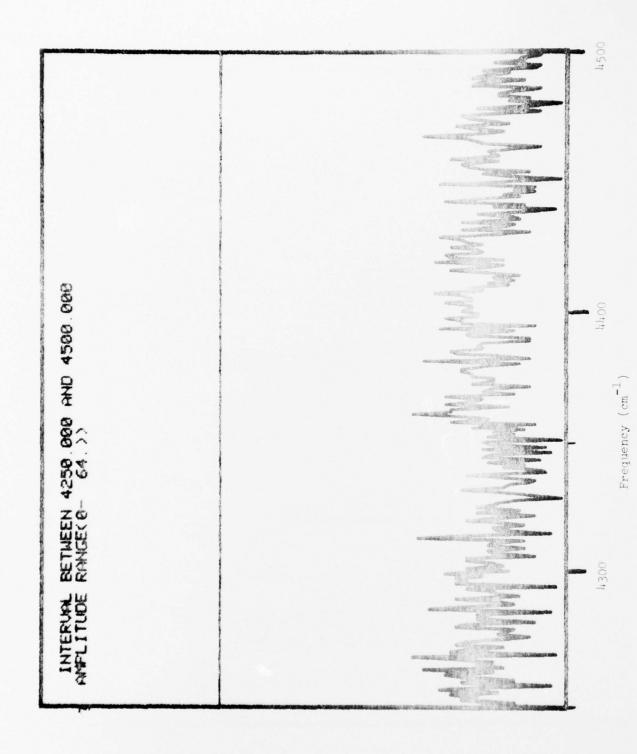
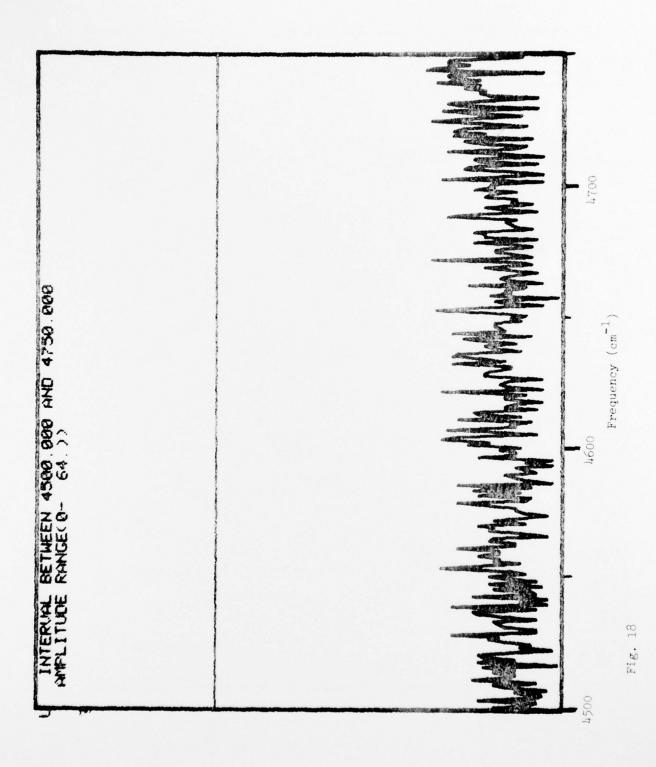


Fig. 17



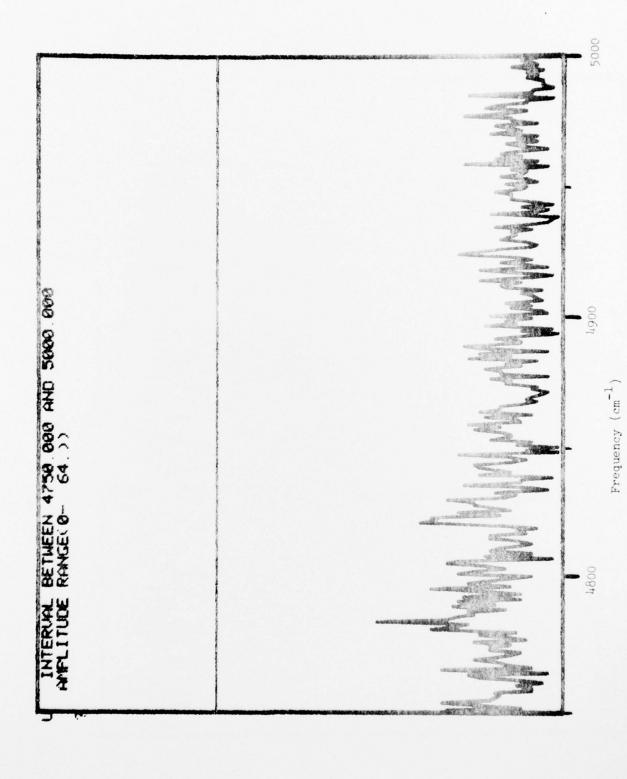


Fig. 19

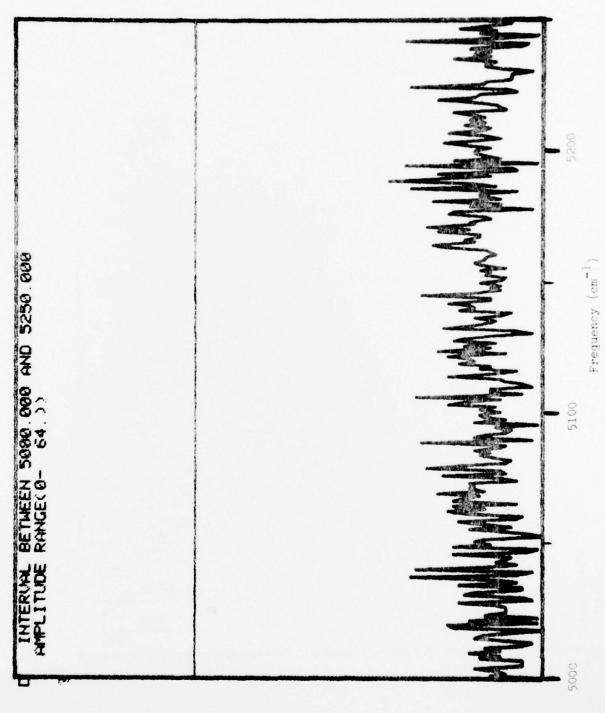


Fig. 20

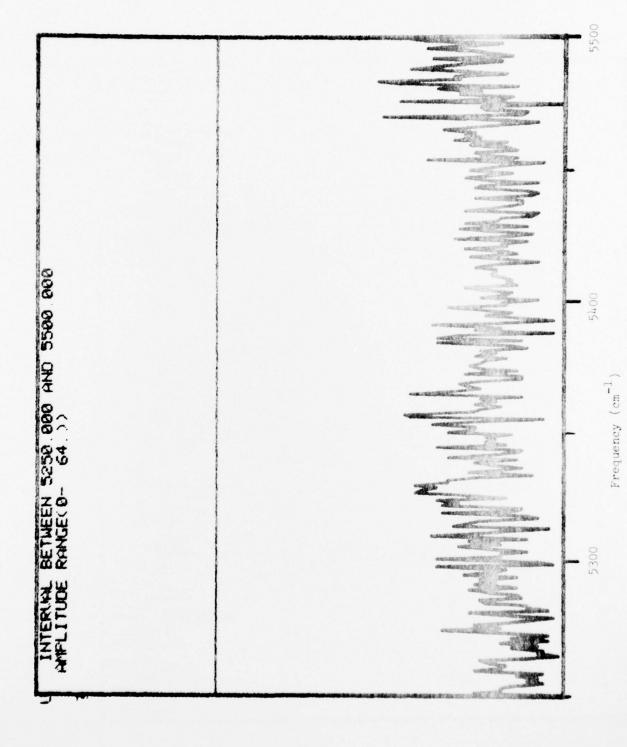


Fig. 2

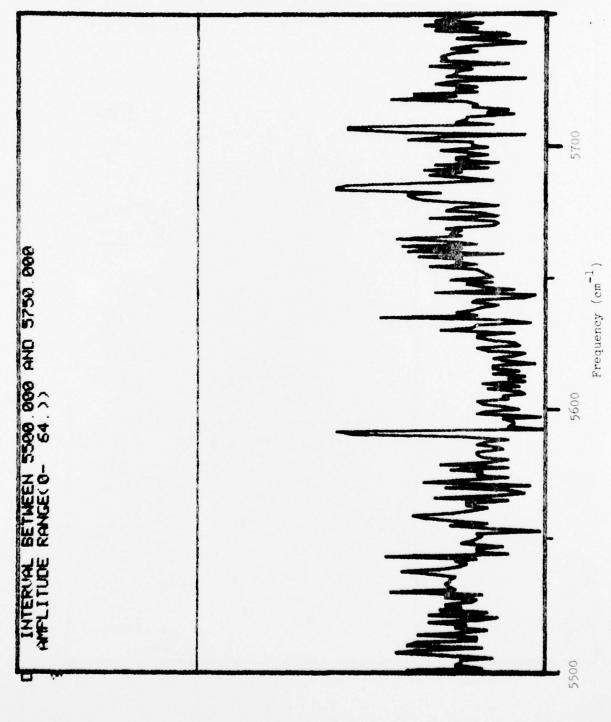


Fig. 22

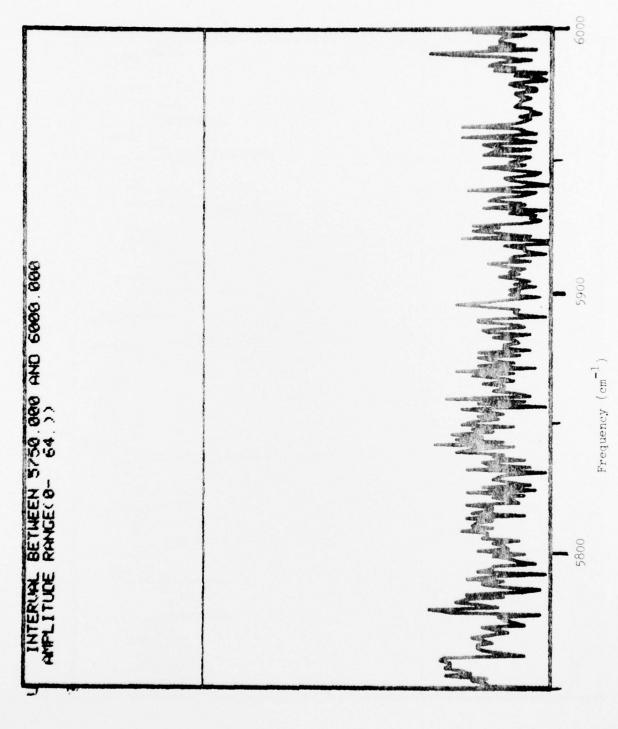


Fig. 23

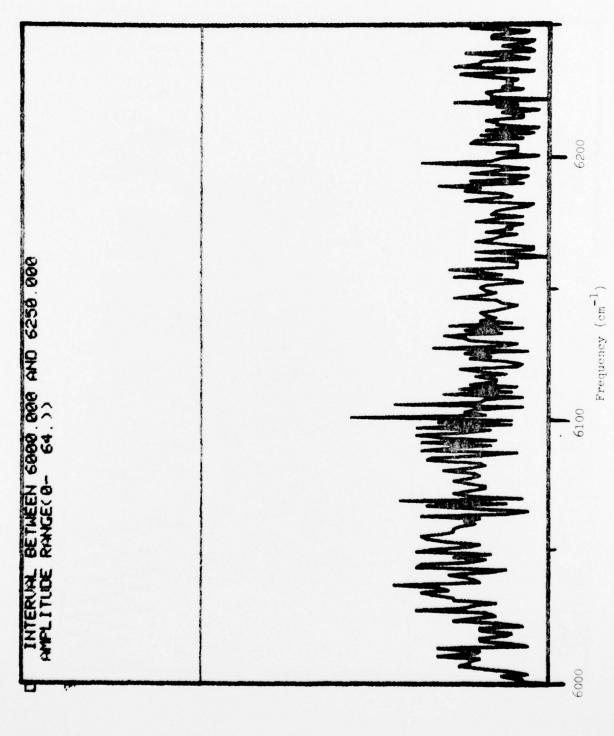


Fig. 24

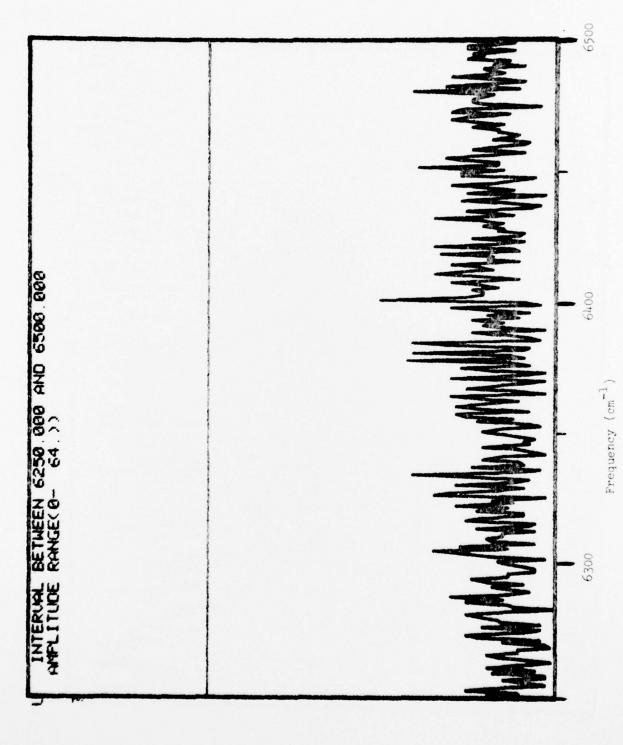


Fig. 25

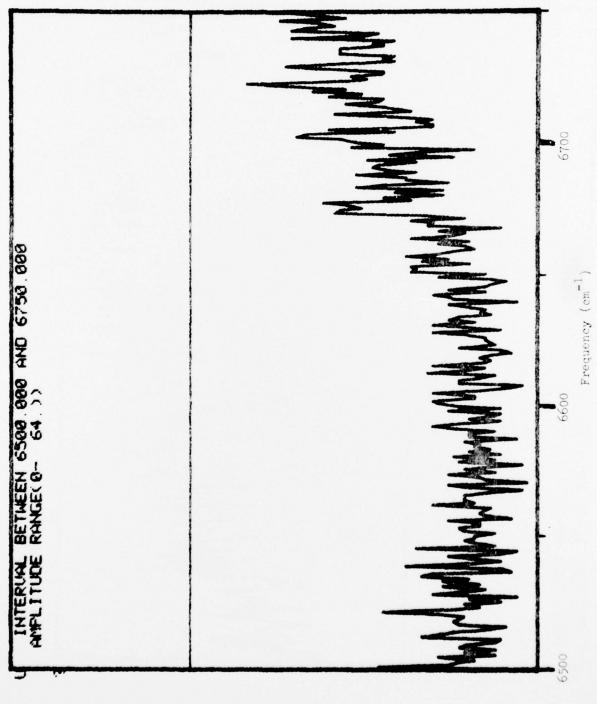


Fig. 26

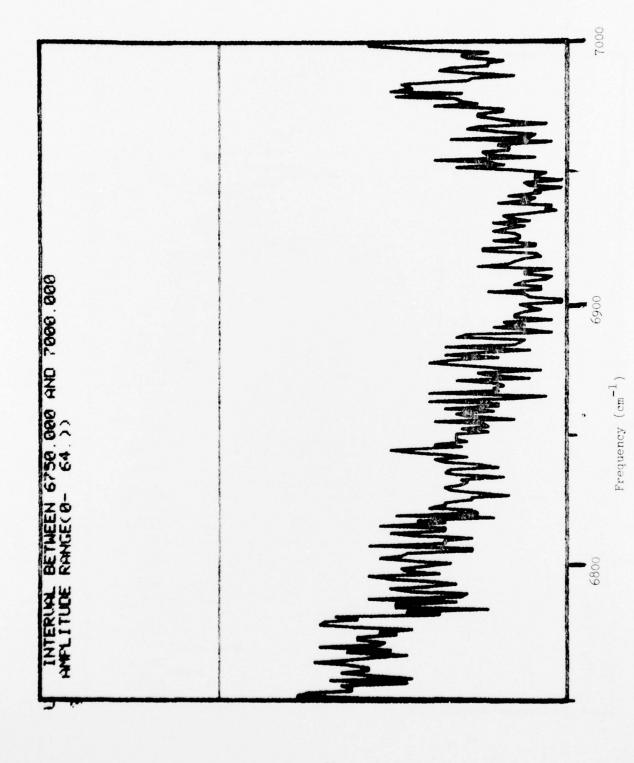
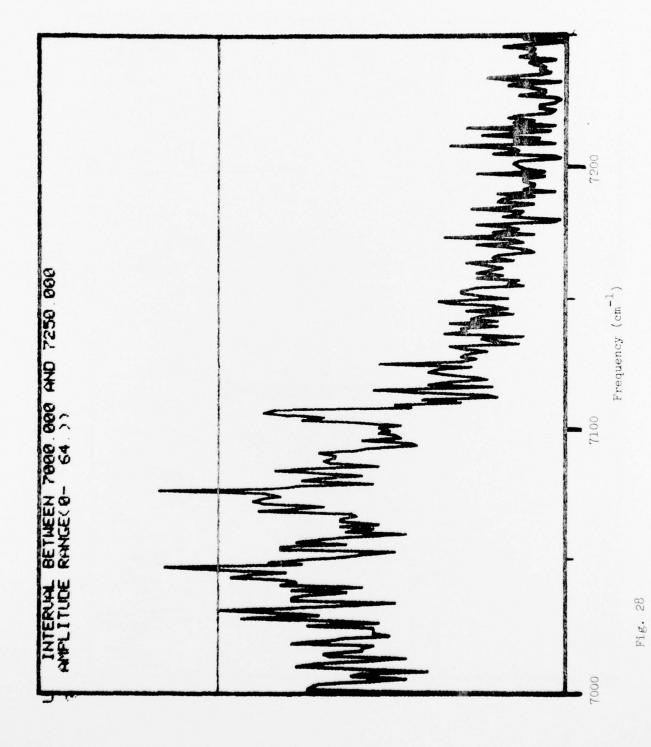


Fig. 27



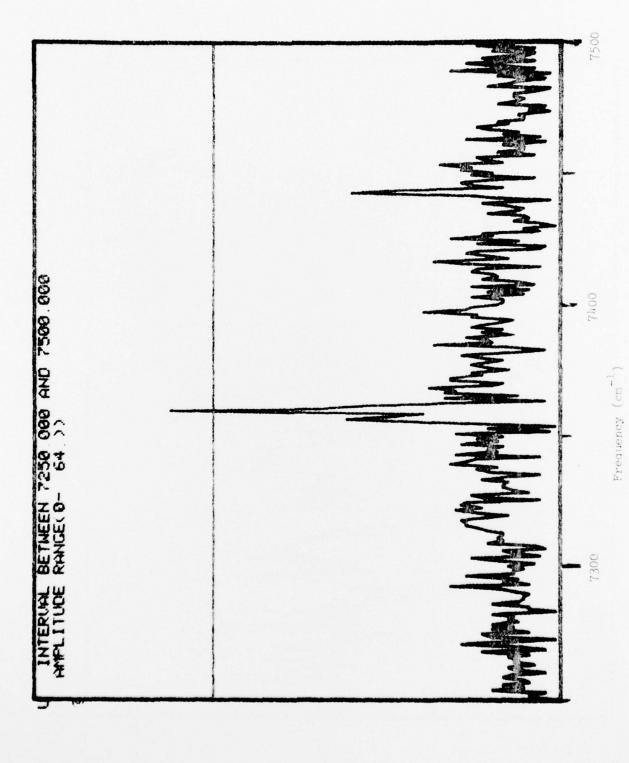


Fig. 29

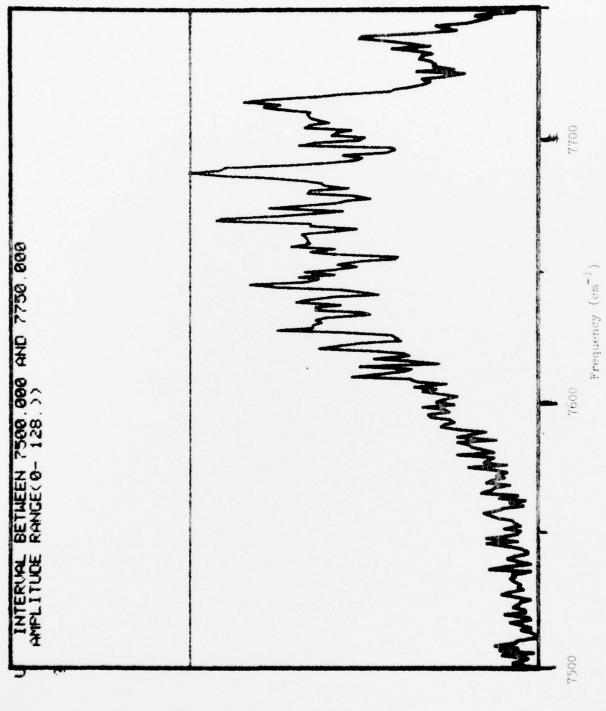


Fig. 30