

CO₂ laser operating at 10.6 μ with optical pumping at 9.6 μ

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We observed for the first time lasing of the CO₂ molecule in the region of the 10.6 μ with optical pumping of the composite 02⁰-00¹ vibrational band by a pulsed CO₂ laser in the 9.6- μ band.

We report here the first successful experimental realization of lasing of the CO₂ molecule in the region of 10.6 μ with pumping by a pulse from a CO₂:N₂He laser

at $\lambda = 9.6 \mu$ with a quantum efficiency 90%. We obtained amplification and lasing in pure CO₂ gas up to atmospheric pressure. Observation of lasing of the CO₂ mol-

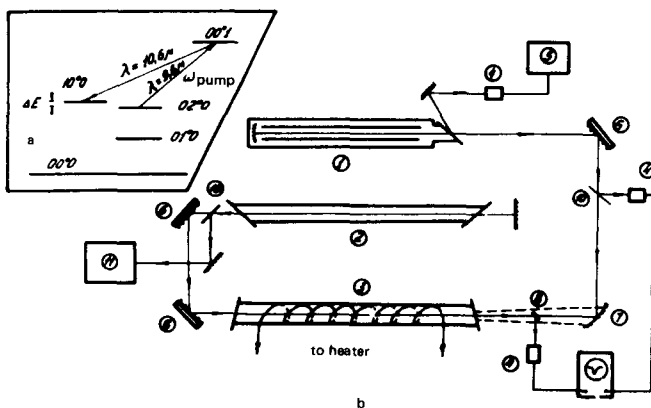


FIG. 1. a) Lower energy levels of the CO₂ molecule. b) Block diagram of setup: 1 – pulsed CO₂ laser, 2 – cw CO₂ laser, 3 – cell, 4 – radiation receiver, 5 – F118 nanovoltmeter, 6 – diffraction gratings, 7 – focusing mirror, 8 – small-aperture mirror, 9 – oscilloscope, 10 – NaCl plates, 11 – IKM-1 monochromator.

ecule with optical pumping is of great interest for the development of simple and effective tunable sealed CO₂ lasers.

The idea of a CO₂ laser optically pumped at 9.6 μ is based on the singularities of the structure of two composite vibrational bands of the CO₂ molecule (Fig. 1), which have a common upper level.

Owing to the closeness of the vibrational levels 10⁰ and 02⁰ (Fermi resonance) and the small energy defect in the vibrational exchange between the levels 00⁰, 01⁰, and 10⁰, the time needed to establish an equilibrium population ratio of these levels in pure CO₂ is of the order of $\tau_{rel} = 10^{-9}$ sec-atm⁻¹.^[1,2] Thus, under the influence of resonant optical pumping in the 02⁰–00¹ band, with a pulse duration $\tau_{pul} \gg \tau_{rel}$, and equalization of the populations of the levels 02⁰ and 00¹ will take place in the absorbing CO₂ gas, whereas the rapid relaxation processes will maintain an equilibrium population of the levels 02⁰ and 10⁰. Then, at a sufficient degree of saturation of the transition 02¹–00¹ there will occur on the transition 00¹–10⁰ a population inversion with a maximum gain $\kappa_g = \kappa_{abs} [\exp(\Delta E/kT) - 1]$, where κ_{abs} is the absorption coefficient of the weak signal in the 10⁰–00⁰ transition and ΔE is the energy difference between the levels 10⁰ and 02⁰. With increasing temperature of the absorbing CO₂ gas, κ_{abs} and κ_g increase. However, with increasing temperature the populations of the levels 10⁰ and 02⁰ become equalized $[\exp(\Delta E/kT) - 1 \rightarrow 0]$, and in addition, the decrease in the pump radiation over the length of the cell with the absorbing CO₂ gas becomes appreciable, as does also the decrease of the time of vibrational relaxation of the 00¹ level, which amounts to $\sim 3.8 \times 10^{-6}$ sec-atm⁻¹ at $T = 296$ K.^[3] These processes limit the growth of the gain with increasing temperature.

A block diagram of the experimental setup for the measurement of the gain produced by pulsed pumping on the 10⁰–00¹ transition is shown in Fig. 1. Previously purified CO₂ at various pressures, 50–760 Torr, is admitted into a cell (3) whose temperature could be varied in the range 23–240 °C and kept constant within

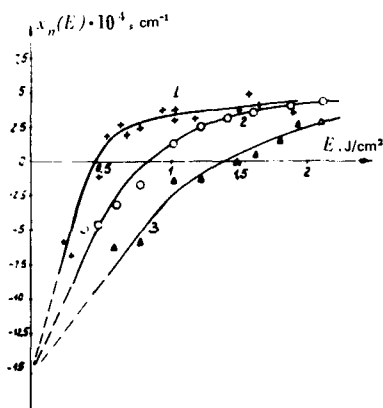


FIG. 2. Gain vs the pump energy density: a – $t = 23$ °C, 1 – $P_1 = 330$ Torr, 2 – $P_2 = 500$ Torr, 3 – $P_3 = 1$ atm. b – $P = 330$ Torr, 1 – $t_1 = 320$ °C, 2 – $t_2 = 343$ °C, 3 – $t_3 = 400$ °C.

± 1 °C. The radiation from the two CO₂ lasers, with selection of the vibrational-rotational transitions, was directed to the same cell. Pump radiation with a pulse energy 10 J and durations 3 μsec, and a pump radiation wavelength 9.6 μ (P branch of the 00¹–02⁰ transition) was applied from one side. From the other side, direction of a probing cw laser at 10.6 μ [the $P(20)$ line of the 00¹–10⁰ transition] was directed to the same cell. To prevent the pulsed CO₂ laser from affected the operation of the cw probing laser, the two lasers were optically decoupled by diffraction gratings (6) and a system of diaphragms.

The lasing pulse of the CO₂ pump laser, the radiation of which saturated the 02¹–10⁰ transition of the molecule CO₂ was fed to the screen of a two-beam oscillo-

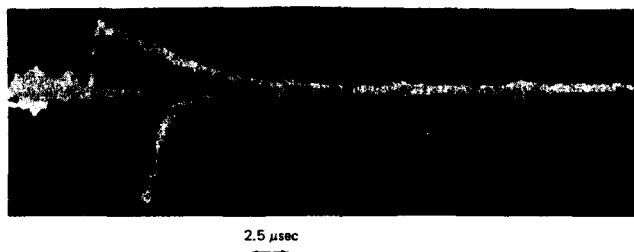


FIG. 3. Oscillogram of pump pulse and generation pulse.

scope together with the signal representing the change of absorption on the $P(20)$ line of the $10^{\circ}0-00^{\circ}1$ transition.

Figure 2 shows plots of the gains on the $P(20)$ line of the $10^{\circ}0-00^{\circ}1$ transition against the pump energy density (J/cm^2) for different pressures and temperatures. The gain at room temperature reached $4 \times 10^{-4} \text{ cm}^{-1}$. The maximum value was $\kappa_g = 2.1 \times 10^{-3} \text{ cm}^{-1}$ at a pressure 100 Torr and a temperature 130°C , corresponding to a 23% gain per pass in a cell 10 cm long.

To observe the generation, the following modifications were made to the experimental setup. The mirror (8) together with a total-reflection mirror with $R = 2 \text{ m}$ formed a semiconfocal resonator. Part of the radiation was extracted from this resonator by reflection from the Brewster window of the cell, was incident on a system of dispersion filters cutting off the 9.6μ radiation. Thus, we were able to eliminate almost completely the radiation of the pulsed pump laser. The generation signal was observed simultaneously with the pump pulse. An oscillogram of the pump pulse at a wavelength 9.6μ and of the generation pulse at the wavelength 10.6μ is shown in Fig. 3. The generation was effected in a wide

range of pressures and temperatures of the molecular CO_2 gas (80–600 Torr, $40-240^{\circ}\text{C}$). We measured the value of $E_{\text{threshold}}$ for the pump energy density.

The developed CO_2 laser with optical pumping can be used to produce a sealed CO_2 laser using a mixture of the isotopic molecules $^{12}\text{C}^{16}\text{O}_2$, $^{13}\text{C}^{16}\text{O}_2$, $^{12}\text{C}^{18}\text{O}_2$, etc. This will permit overlap of the lines a smooth tuning of the frequency at atmospheric pressure.

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