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TEA LASER EFFECTIVE OSCILLATION ON $01^1 1-11^1 0$ HOT
BAND LINES OF ISOTOPIC SPECIES CARBON DIOXIDE

KEY WORDS: TEA CO_2 -laser; $01^1 1-11^1 0$ hot band;
 $^{12}\text{CO}_2$, $^{13}\text{CO}_2$ and $^{14}\text{CO}_2$ isotope modifi-
cations; uranium isotope separation.

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The increase of frequency number generated by conventional CO_2 laser systems is undoubtedly of interest for various applications associated with selective action on substance.

In this paper we report on TEA CO_2 laser oscillation spectrum expanding owing to $01^1 1-11^1 0$ hot band lines of $^{12}\text{CO}_2$ ($10.9-11.4 \mu\text{m}$) and $^{13}\text{CO}_2$ ($11.4-11.9 \mu\text{m}$) molecules at high laser system efficiency.* Besides

* For the first time the weak $^{12}\text{CO}_2$ hot band os-
cillation was registered by C.Frapard et al. in longi-
tudinal discharge system.¹

the possibility of effective powerful radiation source development with $\lambda = 12.15 \mu\text{m}$ that may be used for commercial laser separation of uranium isotopes is also discussed.

To obtain considerable hot band gain K_h of CO_2 molecule it is necessary to heat up ν_2 mode, characterized by vibrational temperature T_2 , along with the excitation of ν_3 mode (temperature T_3). For the conventional values of $T_3 \sim 1600 - 2200 \text{ K}$ realized in an electrical discharge K_h gain is shown to achieve its maximum if $T_2 \sim \frac{1}{3} T_3^2$. Such a relationship between T_2 and T_3 can be reached in gas mixtures with greater CO_2 and lesser He contents, as compared to those optimal for $10.6 \mu\text{m}$ regular band oscillation³. In addition, an increased specific energy input is also required.

The hot band gain investigations have shown the $\text{CO}_2:\text{N}_2:\text{He} = 1:0.8:1$ mixture to be optimal for the UV-preionization TEA CO_2 laser described in details in⁴. The maximum gain of 0.7 m^{-1} for that mixture was achieved at total pressure of $\sim 220 \text{ Torr}$, $0.25 \mu\text{F}$ capacitor bank voltage of $\sim 30 \text{ kV}$ and operational volume of $4 \times 2 \times 70 \text{ cm}^3$. In this case the vibrational temperatures T_3 and T_2 are $\sim 1850 \text{ K}$ and $\sim 630 \text{ K}$, respectively, according to the technique described in⁴.

The hot band line oscillation spectrum studies have been performed for the above optimal active medium parameters. The laser cavity was formed by a full reflective 5-m radius concave AL-coated mirror and

100-line/mm Al-coated grating used as an output coupler. The cavity length was ~ 1 m. Grating reflectivity could be changed depending on the plane of incident radiation polarization from $\sim 60\%$ to 90% . In this case output coupling changes from $\sim 37\%$ to 7% , respectively. Polarization changing was obtained by NaCl Brewster window rotation around the cavity axis. Such a grating mode enables the optimal output coupling to be chosen for rotation-vibration lines with variable J , that is essential for effective oscillation on lines with strongly different gains.

Figure 1 shows the output energy distribution on hot band P-branch for $^{12}\text{CO}_2$ and $^{13}\text{CO}_2$ isotope modifications. Figure 1 a) also shows the output energy value for $00^0 1-10^0 0$ regular band P(42) line obtained under similar conditions. Oscillation spectrum identification was accomplished with Carl Zeiss Jena mirror monochromator SPM-2 and the spectroscopic data.^{5,6} Oscillation was achieved on 36 lines over the range of $10.9-11.4 \mu\text{m}$ with pulse energy exceeding 200 mJ for $^{12}\text{CO}_2$ molecule. The strongest line oscillation energy was about 2.3 J , that is approximately 35% when compared to the regular band strong line output energy for the laser system in optimal mode. In this case the hot band line oscillation efficiency was $\sim 2.3\%$, and specific output energy was $\sim 21 \text{ J L}^{-1} \text{ atm}^{-1}$.

In case of $^{13}\text{CO}_2$ isotope molecule oscillation was obtained on 33 hot band lines over the range 11.45 to

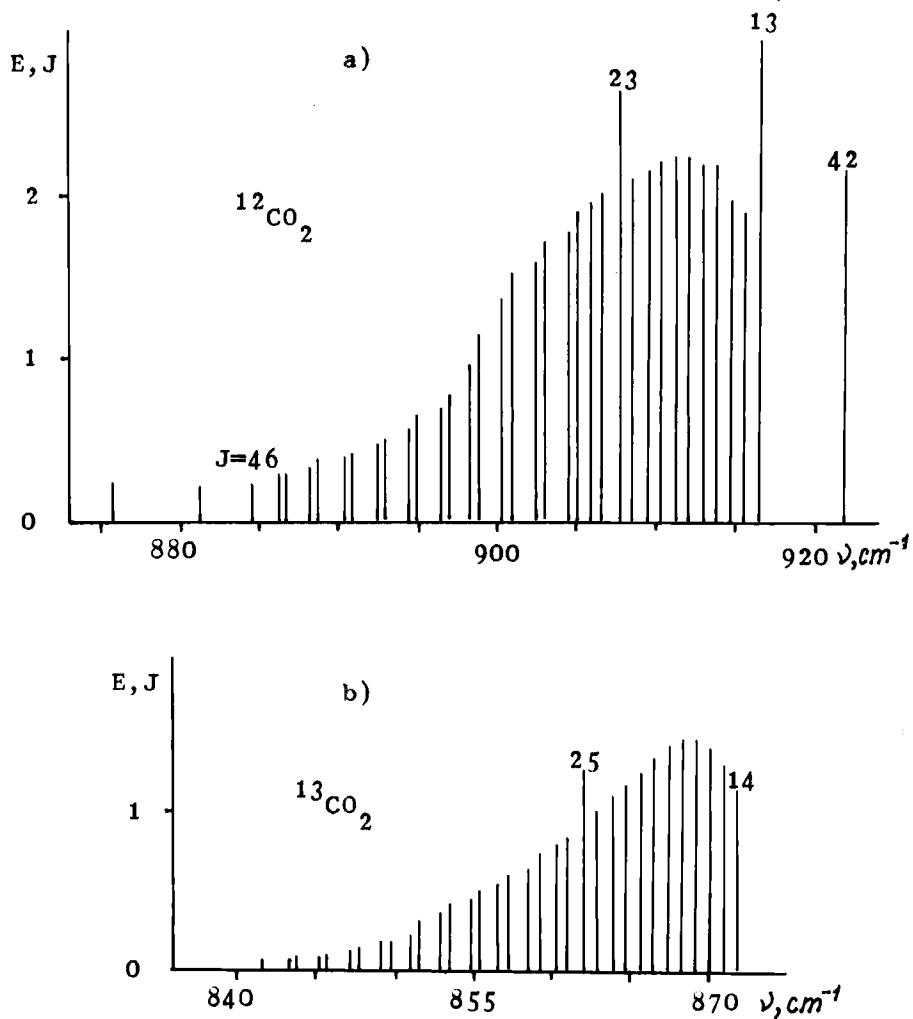


Fig. 1. The output energy distribution on the hot band P-branch for $^{12}\text{CO}_2$ and $^{13}\text{CO}_2$ molecules

11.9 μm . The strongest line oscillation energy was about 1.5 J. Slightly lower efficiency as compared to that of $^{12}\text{CO}_2$ molecule was due to gain reduction because of carbon dioxide mixture use with $^{13}\text{CO}_2 \sim 69\%$ and $^{12}\text{CO}_2 \sim 31\%$.

Note, that abnormally high output energy values for a number of hot band lines for both isotopic species may be explained by their overlapping with the adjacent lines of $00^0 1-10^0 0$, $10^0 1-20^0 0$... bands.

The use of isotopic $^{13}\text{CO}_2$ molecule enables megawatt radiation pulses with frequencies spanning the promising spectral range over 11.4 to 11.9 μm at $\sim 1 \text{ cm}^{-1}$ interval to be received for the first time. The radiation with such parameters may be used for the development of ND₃-optical pump laser with oscillation wavelength about 16 μm range.

The spectral range of 12-12.5 μm which is covered by $^{14}\text{CO}_2$ molecule hot band lines is of special interest for some applications. In this paper we have not dealt with $^{14}\text{CO}_2$ molecule. However, the presented results and analysis of some papers on $^{14}\text{CO}_2$ molecule regular band oscillation⁶ reveal no principal physical difficulties in obtaining the effective $^{14}\text{CO}_2$ molecule hot band oscillation.

The most promising hot band $^{14}\text{CO}_2$ laser application may be its radiation use on one of the strong hot band lines ($\nu = 823 \pm 0.5 \text{ cm}^{-1}$) for uranium isotope separation by multiphoton dissociation of UF_6 .⁷ To

date TEA CO₂ power lasers are well developed and widespread, therefore the proposed source can be easily scaled for commercial uranium isotope separation.

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