

5-17-1979

A Feasibility Analysis of a Directly Sun-Pumped Carbon Dioxide Laser in Space

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<https://doi.org/10.15760/etd.2794>

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AN ABSTRACT OF THE THESIS OF Seiichi Morimoto for the Master of Science in Applied Science presented May 17, 1979.

Title: A Feasibility Analysis of a Directly Sun-Pumped Carbon Dioxide Laser in Space.

APPROVED BY MEMBERS OF THE THESIS COMMITTEE:



George A. Tsongas, Chairman



James M. Heneghan



Selmo Tauber

The possibility of using sunlight to pump a CW carbon dioxide laser has been analyzed. Such a laser could be of interest for such applications as space communication and power transmission. In order to optically pump CO₂ using sunlight, the intense vibrational-rotational absorption bands of CO₂ in the 4.3 micron spectral region would have to be utilized. The total pumping power from sunlight can be calculated

from the known data of the solar spectral irradiance outside the atmosphere and the infrared absorption by carbon dioxide at 4.3 microns. The pumping power is proportional to the collector area of the sunlight and is also dependent on the characteristics of the absorbing gas mixture, such as the gas composition, the gas temperature, the total pressure of the mixture, the partial pressure of CO_2 , and the absorption path length of the sunlight in the gas.

To analyze the carbon dioxide laser system, a thermodynamic approach was used with a simplified CO_2 chemical kinetic model. The gain and saturation intensity were obtained by solving a set of energy balance equations which describe the processes among the various vibrational modes. From those results, along with the estimated cavity losses, the output power was found.

Initially, the computation was carried out for a CO_2 -He mixture for the total pressure range 0.01-0.05 atm with various concentrations of CO_2 at temperatures of 200°K and 300°K , and with the following system parameters specified: a tube of 100 cm length and 1.7 cm diameter, pumping from one end, 1% residual losses, and a 10 m^2 collector. All the collected sunlight was assumed to enter the cavity without losses. Then the parameters were varied to examine their influence on the results. The 10 m^2 collector and a 2 m absorption path length were found to be necessary to achieve sufficient gain and high output power. While the performance was found to be quite sensitive to a number of the individual parameters, the net effect was that the optimum output was obtained under conditions which are close to those initially set. The optimum performance at 200°K was obtained

at 0.02 atm total pressure with 50% CO₂; then the gain was about 5%/m and the output power was about 250 milliwatts. The corresponding efficiency of converting absorbed pump light into laser output was about 4%. At 300°K the gain was found to be unacceptably low because the lower laser level was more highly populated. Thus, the gas must be cooled to achieve sufficient gain. At total pressures above about 0.05 atm the small signal gain was too low, while at pressures below 0.01 atm the gain and output power decreased on account of the low input power and diffusion effects. Therefore, this particular type of CO₂ laser would not appear to be capable of operation at pressures above atmospheric where continuous tunability can be achieved.

This preliminary analysis indicates that it should be feasible to directly sun-pump a carbon dioxide laser in space. The laser would be a modest power, low gain system compared with the demonstrated sun-pumped Nd:YAG laser. If CO₂ output is required or desirable for any of a number of reasons, then this study indicates that direct sun-pumping is possible.

A FEASIBILITY ANALYSIS OF A DIRECTLY
SUN-PUMPED CARBON DIOXIDE LASER IN SPACE

by
SEIICHI MORIMOTO

A thesis submitted in partial fulfillment of the
requirements for the degree of

MASTER OF SCIENCE
in
APPLIED SCIENCE

Portland State University
1979

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ACKNOWLEDGEMENTS

The author wishes to express his sincere appreciation to Dr. George Tsongas for his invaluable guidance and unending assistance throughout the course of this study. In addition, his kind and patient contribution of so much time and effort to improve my writing is gratefully appreciated.

It is also a pleasure to acknowledge Dr. Gail Massey of the Oregon Graduate Center and Dr. Walter Christiansen and Dr. Oktay Yesil of the University of Washington for fruitful discussions and helpful suggestions.

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NOMENCLATURE

English:

| | |
|-------------|--|
| A | Einstein A coefficient, sec^{-1} |
| A_m | Cross-sectional area of the beam, cm^2 |
| A_s | Area of the sun collector, cm^2 |
| B | Rotational constant, cm^{-1} |
| c | Velocity of light, $\text{cm}\cdot\text{sec}^{-1}$ |
| c_{c-k} | The mean relative velocity of CO_2 and the species k (C for CO_2 and H for He), $\text{m}\cdot\text{sec}^{-1}$ |
| D | Diffusion constant, $\text{cm}^2\cdot\text{torr}\cdot\text{sec}^{-1}$ |
| E_v | Vibrational energy per unit volume, $\text{J}\cdot\text{cm}^{-3}$ |
| \bar{E}_v | Vibrational energy in equilibrium, $\text{J}\cdot\text{cm}^{-3}$ |
| E_1 | Vibrational energy of symmetric mode, $\text{J}\cdot\text{cm}^{-3}$ |
| E_2 | Vibrational energy of bending mode, $\text{J}\cdot\text{cm}^{-3}$ |
| E_3 | Vibrational energy of asymmetric mode, $\text{J}\cdot\text{cm}^{-3}$ |
| E_{12} | Combined vibrational energy of E_1 and E_2 , $\text{J}\cdot\text{cm}^{-3}$ |
| $g(\nu)$ | Lineshape function, sec |
| g_J | Degeneracy of J-th rotational level |
| g_1 | Degeneracy of lower vibrational level |
| g_3 | Degeneracy of upper vibrational level |
| h | Planck's constant = 6.626×10^{-34} J·sec |
| I | Intensity of light, $\text{watts}\cdot\text{cm}^{-2}$ |
| I_0 | Intensity of incident light, $\text{watts}\cdot\text{cm}^{-2}$ |

NOMENCLATURE (Cont'd)

English:

| | |
|-------|---|
| I_s | Saturation intensity, $\text{watts} \cdot \text{cm}^{-2}$ |
| i | Vibrational quantum number |
| J | Rotational quantum number of the upper level |
| J' | Rotational quantum number of the lower level |
| k | Boltzmann constant = $1.38 \times 10^{-23} \text{ J} \cdot \text{K}^{-1}$ |
| L | Total intensity loss per pass, % |
| L_i | Residual intensity loss per pass, % |
| l | Length of the laser tube, cm |
| l_a | Absorption path length of the sunlight, cm |
| m_c | The mass of a CO_2 molecule, kg |
| m_k | The mass of a molecule of species k , kg |
| N | Population density of CO_2 , $\text{molecules} \cdot \text{cm}^{-3}$ |
| N_k | Population density of species k (C for CO_2 , H for He), $\text{molecules} \cdot \text{cm}^{-3}$ |
| N_i | Population density of the i -th vibrational level, $\text{molecules} \cdot \text{cm}^{-3}$ |
| N_1 | Population density of the lower vibrational level, $\text{molecules} \cdot \text{cm}^{-3}$ |
| N_3 | Population density of the upper vibrational level, $\text{molecules} \cdot \text{cm}^{-3}$ |
| n_j | Population density of the j -th rotational level, $\text{molecules} \cdot \text{cm}^{-3}$ |
| n_1 | Population density of the vibrational-rotational lower level, $\text{molecules} \cdot \text{cm}^{-3}$ |

English:

| | |
|-------------|--|
| n_3 | Population density of the vibrational-rotational upper level, molecules \cdot cm ⁻³ |
| P_{\max} | Maximum available power, watts |
| P_{\min} | Minimum pumping power, watts |
| P_e | Cavity power (net power generated in the cavity), watts |
| P_o | Output power, watts |
| P_T | Total pumping power, watts |
| P_λ | Solar spectral irradiance at 4.3 microns, watts \cdot cm ⁻² \cdot μ ⁻¹ |
| P | Total pressure of the mixture, atm |
| P_c | Partial pressure of CO ₂ , atm |
| Q_v | Vibrational partition function |
| Q_1 | Vibrational partition function for symmetric mode |
| Q_2 | Vibrational partition function for bending mode |
| Q_3 | Vibrational partition function for asymmetric mode |
| Q_R | Rotational partition function |
| R | Pumping power density, watts \cdot cm ⁻³ |
| r | Radius of the laser tube, cm |
| r_m | Beam radius, cm |
| T | Translational temperature (gas temperature), ^o K |
| T | Useful mirror transmission, % |
| T_v | Vibrational temperature, ^o K |
| T_1 | Vibrational temperature of symmetric mode, ^o K |
| T_2 | Vibrational temperature of bending mode, ^o K |
| T_3 | Vibrational temperature of asymmetric mode, ^o K |

NOMENCLATURE (Cont'd)

English:

| | |
|-------------------|--|
| T_R | Rotational temperature, $^{\circ}\text{K}$ |
| t_{sp} | Spontaneous emission lifetime, sec |
| V | Volume of the laser tube, cm^3 |
| V_m | Volume of the beam, cm^3 |
| v_1 | Symmetric stretching mode |
| v_2 | Bending mode |
| v_3 | Asymmetric stretching mode |
| W | Induced emission rate, sec^{-1} |
| W_{CO_2} | The amount of CO_2 (product of CO_2 partial pressure and absorption path length), $\text{atm}\cdot\text{cm}$ |
| Z | The number of collisions required for deactivation |
| z | Distance, cm |

Greek:

| | |
|--------------|--|
| α | Spectral absorptivity |
| α_i | Spectral absorptivity at a specific wavelength |
| γ | Gain (gain coefficient), cm^{-1} |
| γ_o | Small signal gain, cm^{-1} |
| γ_t | Threshold gain, cm^{-1} |
| ϵ_o | Zero-point energy, J |
| ϵ_i | Quantized vibrational energy of i-th state, J |
| ϵ_j | Quantized rotational energy of j-th level, J |
| η | Energy conversion efficiency |
| θ_v | Characteristic temperature for vibration, $^{\circ}\text{K}$ |

NOMENCLATURE (Cont'd)

Greek:

| | |
|------------------------|--|
| θ_1 | Characteristic temperature for symmetric mode (1920°K) |
| θ_2 | Characteristic temperature for bending mode (960°K) |
| θ_3 | Characteristic temperature for asymmetric mode (3380°K) |
| θ_R | Characteristic temperature for rotation ($.565^{\circ}\text{K}$) |
| λ | Laser wavelength ($\sim 10.6\mu$), μ |
| λ_P | Pumping wavelength ($\sim 4.3\mu$), μ |
| $\Delta\lambda$ | Spectral interval in wavelength, μ |
| ν | Frequency, Hz |
| $\bar{\nu}$ | Wavenumber, cm^{-1} |
| ν_c | Collision frequency, sec^{-1} |
| ν_0 | Center frequency, Hz |
| ν_P | Pumping frequency, Hz |
| $\Delta\bar{\nu}$ | Spectral interval, cm^{-1} |
| $\Delta\nu_D$ | Doppler linewidth, Hz |
| $\Delta\nu_L$ | Lorentzian linewidth, Hz |
| σ | Optical cross section, cm^2 |
| σ_{c-k} | Collision cross section of each species with CO_2 (C for CO_2 and H for He), cm^2 |
| τ | Relaxation lifetime, sec |
| τ_c | Time interval between collisions, sec |
| τ_{C-C} | Relaxation lifetime by collision with CO_2 , sec |
| τ_{C-H} | Relaxation lifetime by collision with He, sec |
| $\tau_{\text{coll},3}$ | Collisional relaxation lifetime of upper laser level, sec |

NOMENCLATURE (Cont'd)

Greek:

| | |
|--------------|---|
| τ_d | Diffusion lifetime, sec |
| τ_{rad} | Radiative lifetime, sec |
| τ_1 | Relaxation lifetime of lower laser level, sec |
| τ_3 | Relaxation lifetime of upper laser level, sec |
| ψ_C | Fractional concentration of CO_2 |
| ψ_H | Fractional concentration of He |

1. INTRODUCTION

At the present rate of growth, the worldwide space communication demand for new channels will be serious in the near future. The capacity of a communication channel is proportional to the width of its band of frequencies. Thus, a laser emitting in the visible or infrared region of the spectrum, where enormously wide bands of frequencies are available, has the potential to carry many more times the amount of information than radio waves or even microwaves. This is the main reason for wanting to use a laser, but optical communication has another advantageous feature. The spatial coherence of a laser beam makes possible highly directional transmission unattainable by conventional radio techniques, which allows lower transmission losses and at the same time maintains secrecy by preventing easy interception of the signals. Moreover, in space use laser communication is not limited at all by atmospheric scattering, absorption, and turbulence. For many of the same reasons and others, lasers are also of interest for power transmission in space.

In view of the interest in laser communication or power transmission in space, it is natural to consider the possibility of sun-pumping a laser since the external pumping source could possibly be eliminated. A long lifetime and lightweight system might be possible by this pumping approach compared with other conventional pumping methods such as electrical pumping using solar cells or other types of power supplies. One of the most appropriate applications of a space-qualified sun-pumped

laser would be on a synchronous orbiting satellite which is occluded from the sun during only a small fraction of its orbit.

To date mostly solid state lasers have been studied for sun-pumping: Both theoretical and experimental investigations on sun-pumping Nd:YAG, Nd:Glass, Nd:CaWO₄, and ruby lasers have been reported.¹⁻⁴ Among these candidates, the Nd:YAG laser seems most promising since it is most highly excited above the threshold value for a given sun-pumping level and hence has the highest output power. In fact, GTE Sylvania has developed and successfully demonstrated a sun-pumped Nd:YAG laser obtaining about two watts CW output power with a 0.3 m² collector using an end-pumping method and conductive cooling.⁵

However, for both space communication and power transmission other types of lasers including CO₂ lasers have been considered. For example, carbon dioxide lasers have several advantages for both applications. First, they are considered to be relatively reliable and long-lived. In addition, high temporal and spatial coherence and outstanding frequency stability compared with solid state lasers is expected. Moreover, CO₂ lasers offer a signal-to-noise advantage over Nd:YAG lasers.⁶ Since infrared lasers have a greater beam divergence than visible lasers, their tracking systems need not be as precise. But this fact also means CO₂ lasers need more power output. The critical technology problem for a CO₂ laser system in space is that of providing Doppler compensation in heterodyne detection systems.⁶

Regarding the possibility of sun-pumping carbon dioxide, the CO₂ molecule has intense and wide vibration-rotational absorption bands in the 4.3 micron region, among others. It is those bands that would have

to be utilized to optically pump CO_2 . Actually the feasibility of achieving considerable power by optical excitation of a CO_2 laser by its own radiation and also by blackbody radiation was hypothesized by Bokhan.^{7,8} In fact, he experimentally proved optical pumping to be practical by obtaining one watt of laser power for a CO_2 (50 Torr) - N_2 (1 atm) mixture with radiation from the hot combustion process of CO in oxygen providing the pump light.⁹ Similar work was done by Wieder.¹⁰ More recently Yesil and Christiansen¹¹ and others¹² have measured gain in a CO_2 gas mixture optically pumped by blackbody radiation in an effort to demonstrate the feasibility of solar pumping. In addition, Chang and Wood optically pumped a CO_2 laser with a pulsed HBr laser and produced a peak power of a few kilowatts for pulses of about 2.2 nsec using extremely high pressure (33 atm) pure CO_2 .

However, the possibility of using the sun to directly pump CO_2 has not previously been pursued. Yet it would be extremely beneficial to know whether such pumping is feasible, and if so to what degree. How might its performance compare with that of the Nd:YAG sun-pumped laser? Thus, the objective of this study is to analyze the feasibility of a directly sun-pumped CO_2 laser for space application. To do that, a simplified theoretical model of such a CO_2 laser is developed, as described in the chapters to follow.

2. FEASIBILITY ANALYSIS

2.1 SOLAR POWER OUTSIDE THE ATMOSPHERE AND ITS ABSORPTION BY CO₂ MOLECULES

In order to determine whether a carbon dioxide laser can be pumped using sunlight, two questions must be answered:

1. In what spectral region does carbon dioxide absorb light for pumping?
2. Is enough sunlight available in that region to achieve sufficient gain and power output, given the absorption characteristics of CO₂?

As noted earlier, carbon dioxide absorbs incident light strongly in the 4.3 micron region. Thus, the amount of solar power available at that wavelength must be determined. Although a great deal of solar spectrum data is available for the visible region, beyond 2.4 microns no complete spectrum has been obtained outside (or inside) the atmosphere. However, it is similar to that of a 6000°K gray-body and can be extrapolated from the known data for shorter wavelengths. From a table of solar spectral irradiance data¹⁵ so determined, the spectral irradiance (defined as the incident radiant energy flux per unit surface area per unit spectral interval) outside the atmosphere for the 4.3 micron region is given as $P_{\lambda} = 0.00073 \text{ watts cm}^{-2} \mu^{-1}$.

With the available solar power known, the absorption of the sunlight by carbon dioxide must be considered. The infrared absorptivity (the fraction of the incident radiation absorbed) of carbon dioxide

at 4.3 microns has been calculated for a wide range of values of absorption path length, pressure, and temperature.^{16,17 *}

Here some typical values of the spectral absorptivity α are plotted in Figure 1 for various total pressures p and partial pressures of carbon dioxide p_c ; the absorption path length of the sunlight ℓ_a is taken to be 2 meters (round trip path in a 1 m tube is assumed).^{**} Clearly, there is more absorption at high pressures where more absorbing molecules exist. Pressures higher than those given are not presented because they will be shown to correspond to non-optimum laser performance.

Assuming no losses transmitting the incident sunlight into the cavity (in reality some loss factor would exist), the total solar pumping power P_T into a given cavity can be calculated from the following expression (see Figure 2):

$$P_T = P_\lambda A_s \sum_i \alpha_i \Delta\lambda \quad (1)$$

where

P_λ : solar power at 4.3 μ per unit area per unit spectral intervals (watts $\text{cm}^{-2} \mu^{-1}$)

A_s : sun collector area (cm^2)

α_i : spectral absorptivity of CO_2 at λ_i , where λ_i is the center wavelength of the i -th spectral segment

* In references 16 and 17 absorption data are for CO_2 in air (i.e. almost a CO_2 - N_2 gas mixture), but it can be shown that these data can be used for a CO_2 -He gas mixture as well with an error of less than a few percent. The reason is that broadening caused by N_2 is not very much different from that caused by He when either is in a mixture with CO_2 . In addition, variation of total pumping power with broadening is even much smaller than that of broadening itself.

** The data from references 16 and 17 were actually tabulated for various p and W_{CO_2} , where $W_{\text{CO}_2} = P_c \ell_a$.

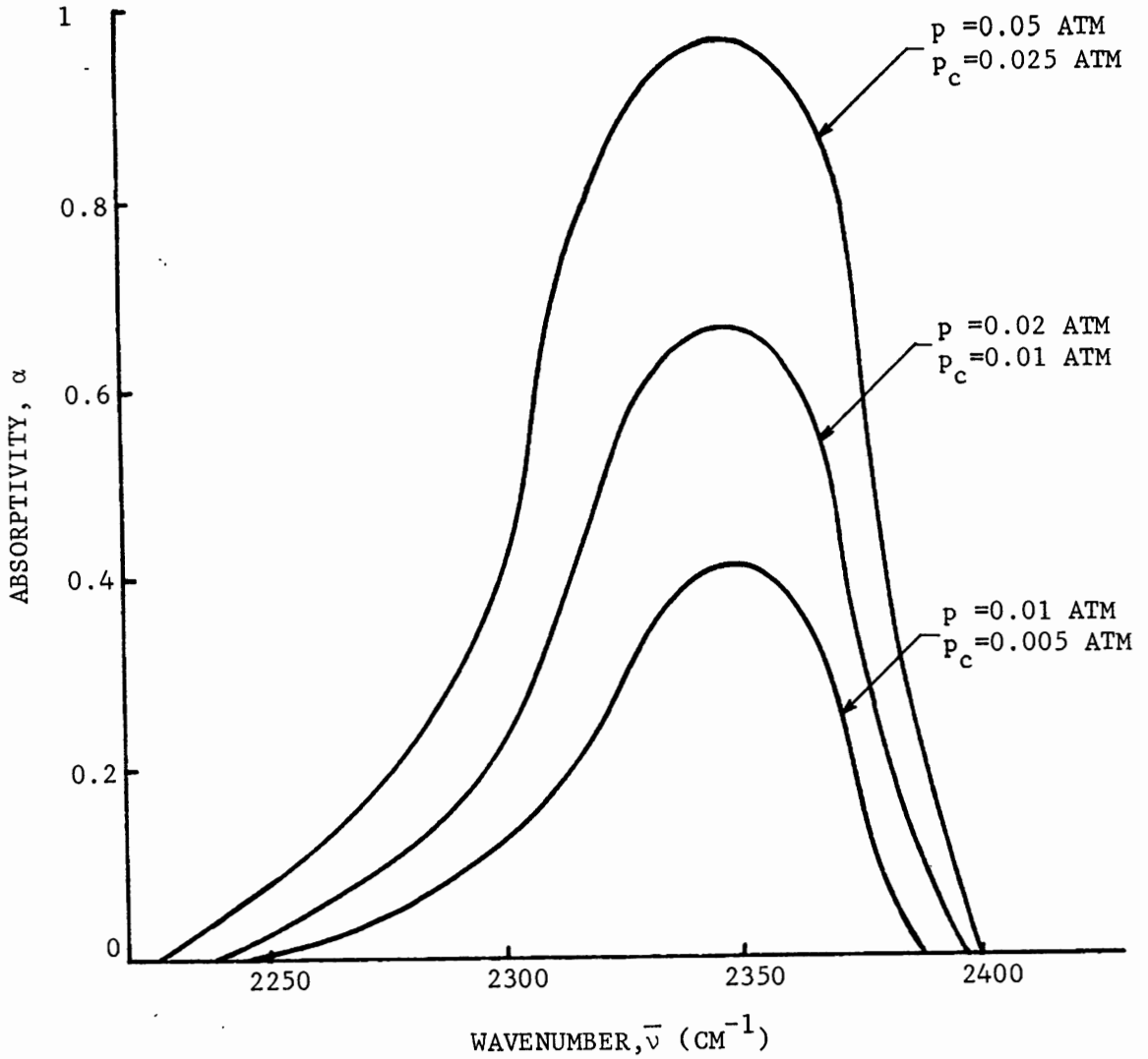


Figure 1. The infrared absorptivity of CO₂ in the 4.3 micron region at 200°K for various total pressures p and CO₂ partial pressures p_c for a 2 meter absorption path length.

$\Delta\lambda$: spectral interval of the given absorptivity data * (μ)

It should be noted that this procedure for the determination of P_T is used because the absorptivity data is provided in tabular form at discrete wavelength intervals.

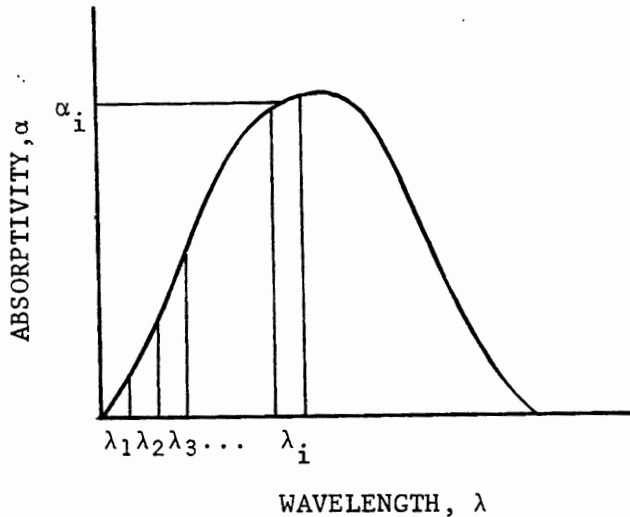


Figure 2. Segmentation of the Absorptivity Curve for Calculation of P_T .

Using equation (1) and the absorptivity data, P_T was calculated. The results are plotted in Figures 3 and 4 versus $\log p_c$ (where $\lambda_a = 2$ m) for each p for the temperatures 200°K and 300°K , respectively.** These temperatures will later be shown to be in the range of prime importance for laser action.

It can be seen from the graphs that P_T increases with temperature (15-20% increase in P_T from 200°K to 300°K), and the variation of P_T with p_c is fairly logarithmic.

* $\Delta\lambda = 0.018\mu$, which corresponds to $\Delta\bar{\nu} = 10 \text{ cm}^{-1}$.

** The horizontal scale $\log p_c$ is actually $\log W_{\text{CO}_2}$ with fixed value of $\lambda_a = 2$ m. P_T for values of λ_a other than 2 m can be found easily from the relation $W_{\text{CO}_2} = \lambda_a P_c$.

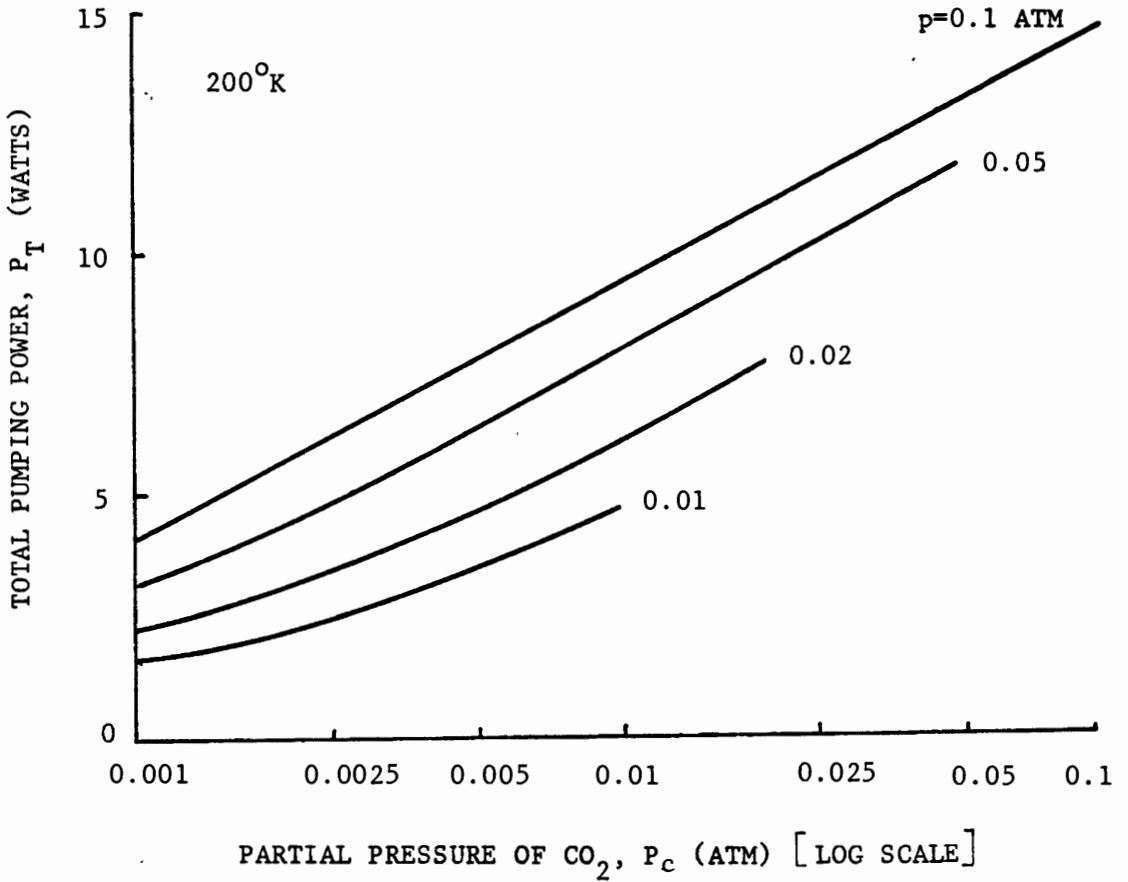


Figure 3. Total pumping power, P_T , at 200°K for a 10 m² collector and a 2 meter absorption path length.

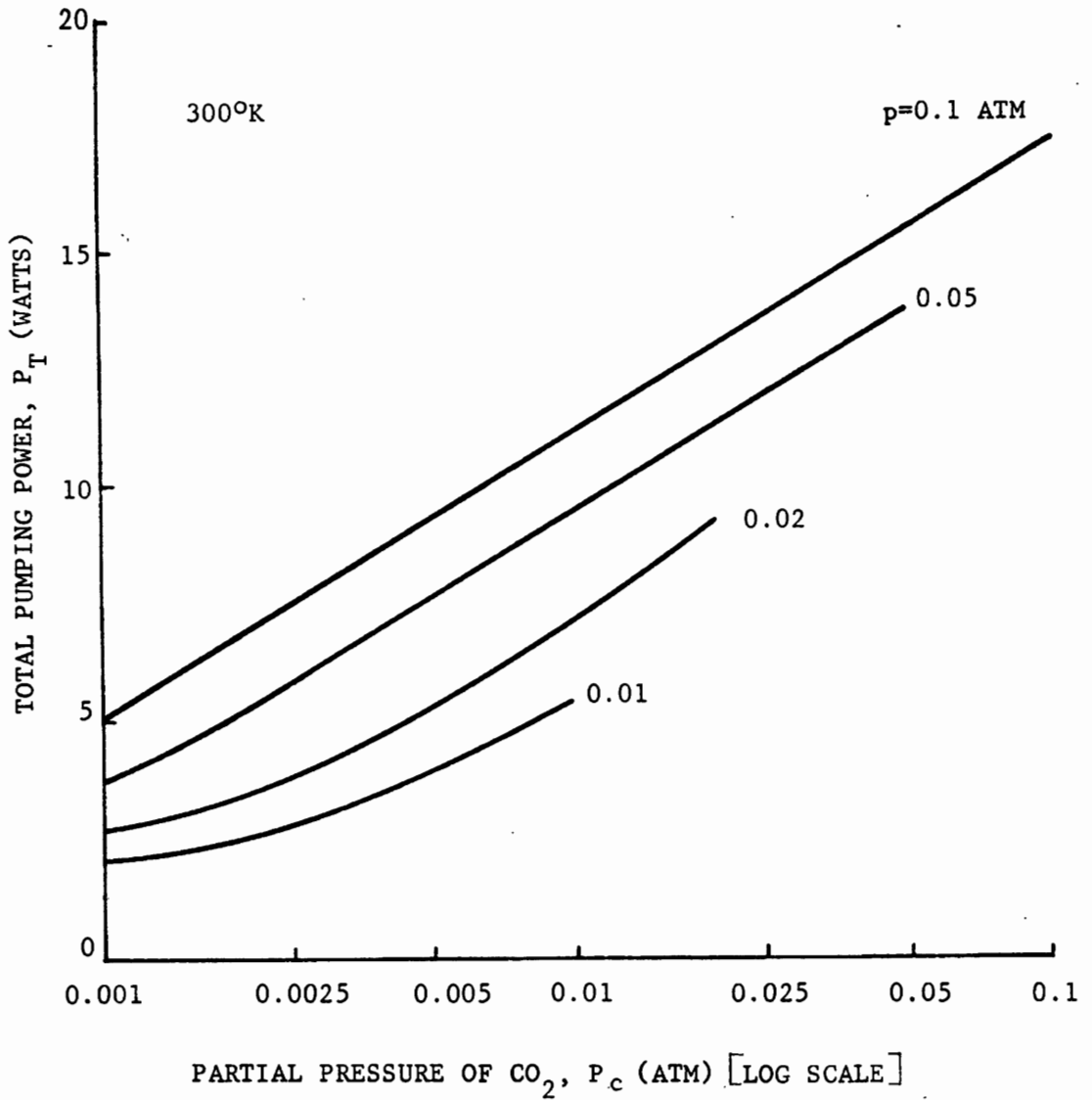


Figure 4. Total pumping power, P_T , at 300°K for a 10 m² collector and a 2 meter absorption path length.

2.2 THEORETICAL MODEL AND CHEMICAL KINETICS OF CARBON DIOXIDE

Since a thermodynamic approach will be used later to analyze the carbon dioxide laser system, it will be appropriate to present here some fundamental facts about vibrational and rotational levels of CO_2 and the relaxation processes among the vibrational levels. This includes specific equations for the energy of the vibrational modes, for the population of vibrational and rotational levels, and for the relaxation lifetimes of vibrational states. These equations will be later used to directly calculate the gain and saturation intensity of the sun-pumped CO_2 gas mixture, as well as the laser output power.

Carbon dioxide is a linear symmetrical molecule which has three normal modes of vibration:

| | | |
|------------------------------------|-------|-----------------|
| the symmetric stretching mode | v_1 | $(i_1 \ 0 \ 0)$ |
| the doubly degenerate bending mode | v_2 | $(0 \ i_2 \ 0)$ |
| the asymmetric stretching mode | v_3 | $(0 \ 0 \ i_3)$ |

where $i_1, i_2, i_3 = 0, 1, 2, 3, \dots$ specify the vibrational states. A partial energy level diagram for the carbon dioxide molecule is illustrated in Figure 5. Shown are the pertinent vibrational energy levels together with a set of rotational levels for the $(1 \ 0 \ 0)$ and the $(0 \ 0 \ 1)$ states on a much expanded scale. The 10.6 micron laser transition and some principal relaxation processes are shown in the diagram.

2.2.1 Vibrational Energy Levels

The molecules may be regarded to a good approximation as harmonic oscillators of frequency ν . According to quantum mechanics, the

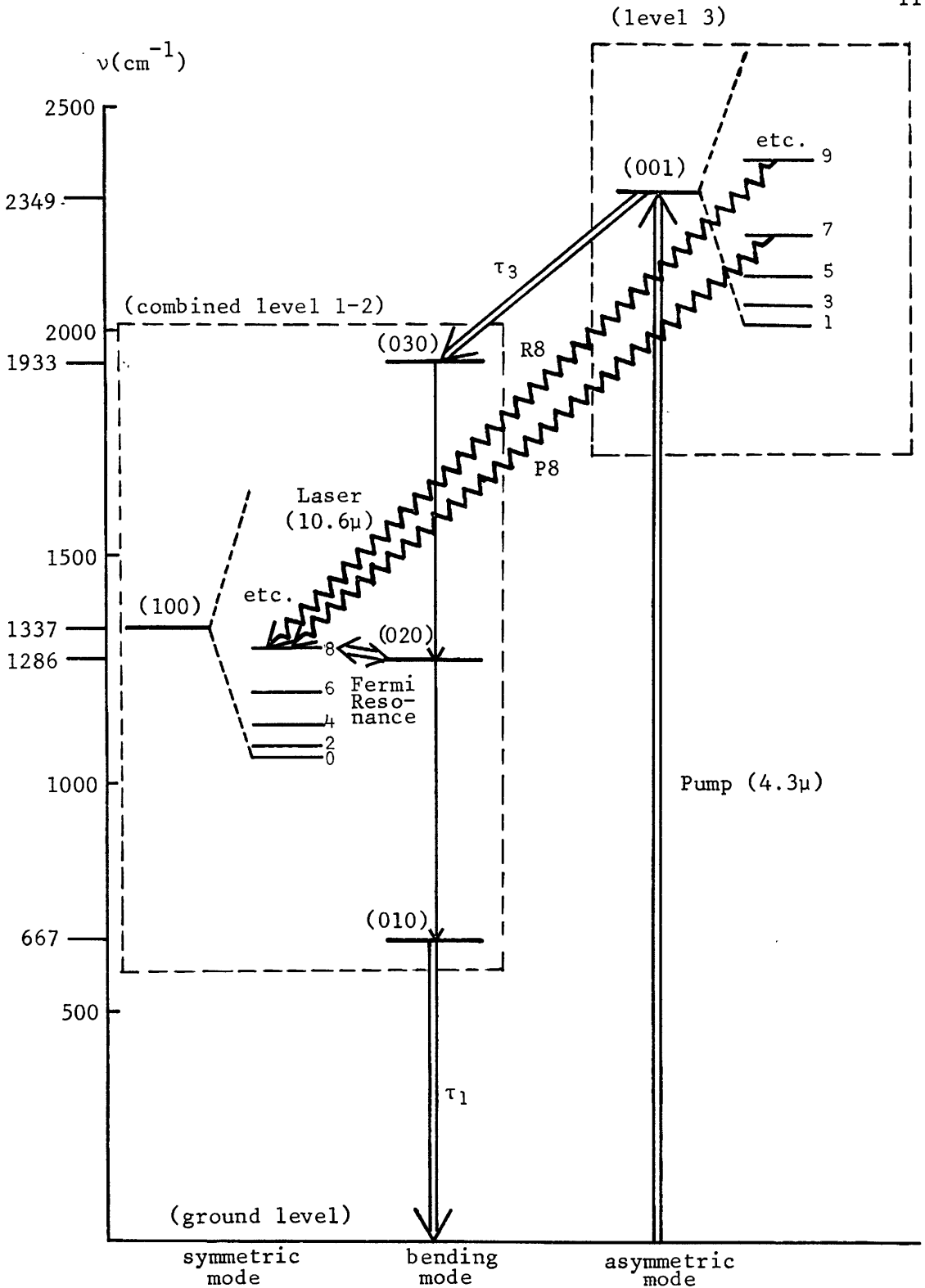


Figure 5. Partial energy level diagram for CO₂.

degrees of vibrational excitation are quantized and the permissible energy states of a molecule are given by

$$\epsilon_i = (i + \frac{1}{2})h\nu \quad \text{where } i = 0,1,2,3, \dots \quad (2)$$

Omitting the zero-point energy $\epsilon_0 = \frac{1}{2}h\nu$ of a vibrational ground state (i.e. vibrational energies are measured from the ground state),

$$\epsilon_i = ih\nu \quad \text{where } i = 0,1,2,3, \dots \quad (3)$$

Now the vibrational partition function is introduced (i.e. how the particles are partitioned or divided up among the energy states), as defined by

$$Q_v = \sum_{i=0}^{\infty} \exp(-\epsilon_i/kT_v) = \sum_{i=0}^{\infty} \exp(-ih\nu/kT_v) \quad (4)$$

where T_v is the vibrational temperature. That temperature is, in effect, defined in the Einstein function of equation (7) below.

When $T_v = T$ (gas temperature), a thermodynamic non-equilibrium condition exists. Note that T_v (and E_v shown later) indicates the degree of population of a certain vibrational state; the higher the vibrational temperature, the larger the population of the vibrational state.

The sum in equation (4) is recognized as the geometrical series

$$Q_v = 1/[1 - \exp(-\theta_v/T_v)] \quad (5)$$

where

$$\theta_v = h\nu/k : \text{characteristic temperature for vibration}$$

Using equation (5) and the relation¹⁸

$$E_v = NkT_v^2 \frac{\partial \ln Q_v}{\partial T_v} \quad (6)$$

where N is the total population density of CO_2 , the total vibrational energy of the oscillators per unit volume is

$$E_v = Nk\theta_v / [\exp(\theta_v/T_v) - 1] \quad (7)$$

This is the well known Einstein function (in terms of vibrational temperature T_v) for a simple harmonic oscillator with a Boltzmann distribution of the states.

When T_v is in equilibrium with the translational temperature T (i.e. $T_v = T$),

$$\bar{E}_v = Nk\theta_v / [\exp(\theta_v/T) - 1] \quad (8)$$

The population distribution for the i -th vibrational level is given by the formula¹⁸

$$N_i/N = \exp(-i\theta_v/T_v)/Q_v \quad (9)$$

where

N_i : population density of the i -th vibrational level

N : total population density

Now, from the ideal gas law

$$N(\text{molecules/cm}^3) = 7.37 \times 10^2 p_c(\text{atm})/T(^{\circ}\text{K}) \quad (10)$$

So far we have been referring to one vibrational mode. However, since carbon dioxide has three modes, the vibrational partition function Q_v for CO_2 is (taking the double degeneracy of the bending mode into account)

$$Q_v = Q_1 Q_2^2 Q_3 \quad (11)$$

$$= \{ [1 - \exp(-\theta_1/T_1)] [1 - \exp(-\theta_2/T_2)]^2 [1 - \exp(-\theta_3/T_3)] \}^{-1} \quad (12)$$

where Q_1 , Q_2 , and Q_3 correspond to the v_1 , v_2 , and v_3 modes, respectively.

Then, the energies of the three modes are, respectively,

$$E_1 = NkT_1^2 (\partial \ln Q_v / \partial T_1) = Nk\theta_1 / [\exp(\theta_1/T_1) - 1] \quad \text{for } v_1 \text{ mode} \quad (13)$$

$$E_2 = NkT_2^2 (\partial \ln Q_v / \partial T_2) = 2Nk\theta_2 / [\exp(\theta_2/T_2) - 1] \quad \text{for } v_2 \text{ mode} \quad (14)$$

$$E_3 = NkT_3^2 (\partial \ln Q_v / \partial T_3) = Nk\theta_3 / [\exp(\theta_3/T_3) - 1] \quad \text{for } v_3 \text{ mode} \quad (15)$$

where T_1, T_2 , and T_3 are the vibrational temperatures of the v_1, v_2 , and v_3 modes, respectively, and θ_1, θ_2 , and θ_3 are the characteristic temperatures of vibration of the v_1, v_2 , and v_3 modes, respectively.

2.2.2 Rotational Energy Levels

Each vibrational level is subdivided into a number of rotational levels denoted by J's. J can only be odd or even numbers for a given vibrational level, and the selection rule for rotational transitions is $J = \pm 1$ ($J = +1$ is a P-transition; $J = -1$ is an R-transition).

The quantized rotational energy is given by

$$\epsilon_J = hcBJ(J+1) \quad (16)$$

where

$$J = 0, 1, 2, 3, \dots$$

B = rotational constant for each vibrational state

Taking the degeneracy $g_J (=2J+1)$ into account, the rotational partition function is

$$Q_R = \sum_{\substack{J=0 \\ J=\text{even or odd}}}^{\infty} g_J \exp(-\epsilon_J/kT_R) \quad (17)$$

$$= \sum_{\substack{J=0 \\ J=\text{even or odd}}}^{\infty} (2J+1) \exp[-J(J+1)hcB/kT_R] \quad (18)$$

By approximating the series by an integral for a linear and symmetric molecule like CO_2 ,

$$Q_R = \frac{1}{2} \int_0^{\infty} dJ (2J+1) \exp[-J(J+1)hcB/kT_R] \quad (19)$$

which can be shown to be ¹⁸

$$Q_R = T_R/2\theta_R \quad (20)$$

where

$\theta_R = hcB/k$: characteristic temperature for rotation

$T_R =$ rotational temperature

Among the various rotational levels of a vibrational level a Boltzmann distribution in terms of the gas temperature $T(=T_R)$ is established, which results from the fact that the spacing of the rotational energy levels is comparable to the kinetic energy (0.025eV) of a molecule and these energies are easily exchanged in collisions (thermalization).

Hence, at gas temperature $T (=T_R)$ the population of the J -th rotational level is determined by¹⁸

$$n_J = g_J N_i \{ \exp[-J(J+1)\theta_R/T] \} / Q_R \quad (21)$$

where

$n_J =$ population density of the J -th vibrational-rotational level

By differentiating this equation with respect to J , the value of J for which n_J is maximum is given by

$$J_{\max} = \sqrt{kT/2hcB} - \frac{1}{2} \approx 0.95 \sqrt{T} - \frac{1}{2} \quad (22)$$

where $B \approx 0.39 \text{ cm}^{-1}$ for (100) and (001).¹⁹

The maximum gain occurs when using this expression for J_{\max} ; the value of J_{\max} at those temperatures where the gain will be evaluated is as follows.

At $T = 200^\circ\text{K}$

$$J_{\max} = 13 \quad \text{i.e. } J = 13, J' = 14 \quad \text{for P(14)}$$

At $T = 300^\circ\text{K}$

$$J_{\max} = 16 \quad \text{i.e. } J = 15, J' = 16 \quad \text{for P(16)}$$

2.2.3 Vibrational Relaxation

Excited molecules in vibrational levels will decay in three different fashions: by radiative decay (spontaneous emission), collisional deactivation and spatial diffusion. Over the pressure range of interest in this study, the latter two processes are dominant since the radiative decay lifetimes are considerably larger than the corresponding collision or diffusion lifetimes.*

One means of effectively raising the relaxation rate at which molecules start or cease to interact with the lasing field is by diffusions of molecules into and out of the laser beam (i.e. the region in the gas where amplification exists). Diffusion can play an important role in laser beams with both small radii and low total pressure. An expression for the diffusion lifetime τ_d is found to be²¹⁻²³

$$\frac{1}{\tau_d(\text{sec})} = \frac{8D}{p(\text{torr}) r_m^2(\text{cm}) [1+4\ln(r/r_m)]} \frac{T(^{\circ}\text{K})}{300} \quad (23)$$

For a CO_2 -He mixture where $D = 400 \text{ cm torr/sec}$ ²³, the expression becomes

$$\tau_d(\text{sec}) = \frac{71 p(\text{atm}) r_m^2(\text{cm}) [1 + 4 \ln(r/r_m)]}{T(^{\circ}\text{K})} \quad (24)$$

where

D : diffusion constant

r_m : beam radius

r : radius of the tube

* The radiative lifetime τ_{rad} is about 2.5 msec for the (001) - (100) transition (all P-transitions and R transitions), while for other transitions it is much larger.²⁰

The second important deactivation process is collisional relaxation. Since this is the dominant process for the pressure range of interest in this study (i.e. $p = 0.01-0.05$ atm), the rest of our discussion will be concerned with the detailed mechanism of collisional relaxation.

Transfer of energy between two vibrational states or one vibrational and one translational degree of freedom occurs through molecular collisions. The collisional relaxation lifetime τ (which is the average time for a molecule to exist in an excited state before deactivation) is defined by

$$\tau = \tau_c Z \quad (25)$$

where

τ_c : the time interval between collisions, which is inversely proportional to pressure

Z : the average number of collisions to deactivate one vibrational quantum to its e^{-1} value (i.e. $1/Z$ is the probability of deactivation of vibrational energy in one collision)

In a mixture involving CO_2 and the foreign gas He (as noted later, this mixture is chosen for our laser system), relaxation of a vibrationally excited CO_2 molecule can take place through binary collision processes with either ground state CO_2 molecule or with a He atom. The relaxation rate for binary collisions is given by ^{24,25}

$$\frac{1}{p\tau} = \frac{\psi_C}{p\tau_{C-C}} + \frac{\psi_H}{p\tau_{C-H}} \quad (26)$$

where

τ_{C-C} : relaxation lifetime for collisions with CO_2

τ_{C-H} : relaxation lifetime for collisions with He

$1/p\tau_{C-C}$: relaxation rate for pure CO_2 at p atm pressure

$1/p\tau_{C-H}$: relaxation rate for collisions with He at p atm pressure

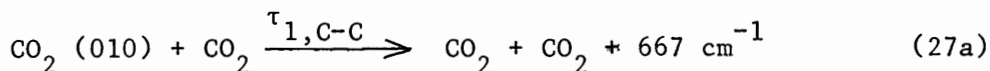
ψ_C and ψ_H : mole fractions of CO_2 and He, respectively

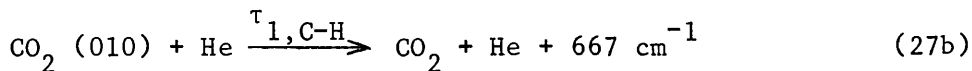
$$(\psi_C + \psi_H = 1)$$

In what follows the kinetic reactions and the values of the lifetimes τ_{C-C} and τ_{C-H} for the pertinent levels will be examined. In modeling the important kinetic processes involved in a CO_2 laser, certain simplifications have been made as shown in the diagram of Figure 5. It has been assumed that V - V (vibrational - vibrational) transitions are very fast in each mode (i.e. in the vertical direction, such as (030) \rightarrow (020), (020) \rightarrow (010), but not from one mode to any other (i.e. in the horizontal direction). It should, however, be noted that process (100) \rightleftharpoons (020) is very rapid because of the known Fermi resonance (F.R.). Therefore, with this simplified model, mode v_1 and v_2 are considered to be one combined level 1-2. Then, the rate-controlling processes among the groups of vibrational levels (i.e. level 3, combined level 1-2, and the ground level) are those from (001) to level 1-2 (i.e. lifetime τ_3) and from level (010) to ground level (i.e. lifetime τ_1).

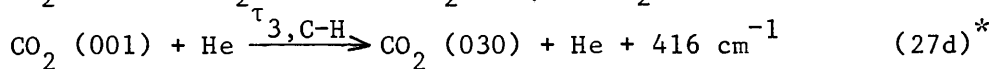
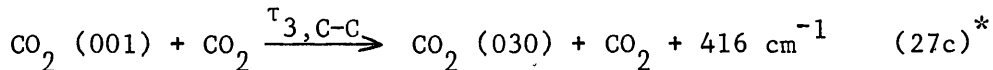
The pertinent processes for energy transfer between the noted vibrational states can be expressed as kinetic reactions as follows:

T - V (translational - vibrational) processes:





V - V (vibrational - vibrational) processes:



Crucial to laser oscillation is the fact that the lifetime of the upper level τ_3 must be larger than the lifetime of the lower level τ_1 . Only then can a population inversion be achieved. Furthermore, the population inversion could be increased by enhancing the lower level depopulation rate with the addition of helium.

The values of the collisional relaxation lifetimes $\tau_{3,C-C}$, $\tau_{3,C-H}$, $\tau_{1,C-C}$ and $\tau_{1,C-H}$ for the kinetic processes noted above have previously been measured experimentally and are plotted versus temperature for convenience in Figures 6 and 7.^{24,26-29} The lifetimes vary inversely with pressure.

Now, since the diffusion effect on the upper level is considerable and radiative decay is not negligible in the lowest pressure range of this study, a combined lifetime τ_3 which incorporates all of these effects must be used; it is given as:

$$\frac{1}{\tau_3} = \frac{1}{\tau_{d,3}} + \frac{1}{\tau_{\text{coll},3}} + \frac{1}{\tau_{\text{rad},3}} \quad (28)$$

Note that the τ_3 referred to earlier was really $\tau_{\text{coll},3}$. The values of $\tau_{d,3}$ are calculated using equation (24).

* Reactions (27c) and (27d) are considered to be dominant for the relaxation process from (001) to the 1-2 level. However, since those from (001) to any other levels in the 1-2 level could be responsible, $\tau_{3,C-C}$ and $\tau_{3,C-H}$ should be understood to correspond to the process from (001) to the whole 1-2 level.

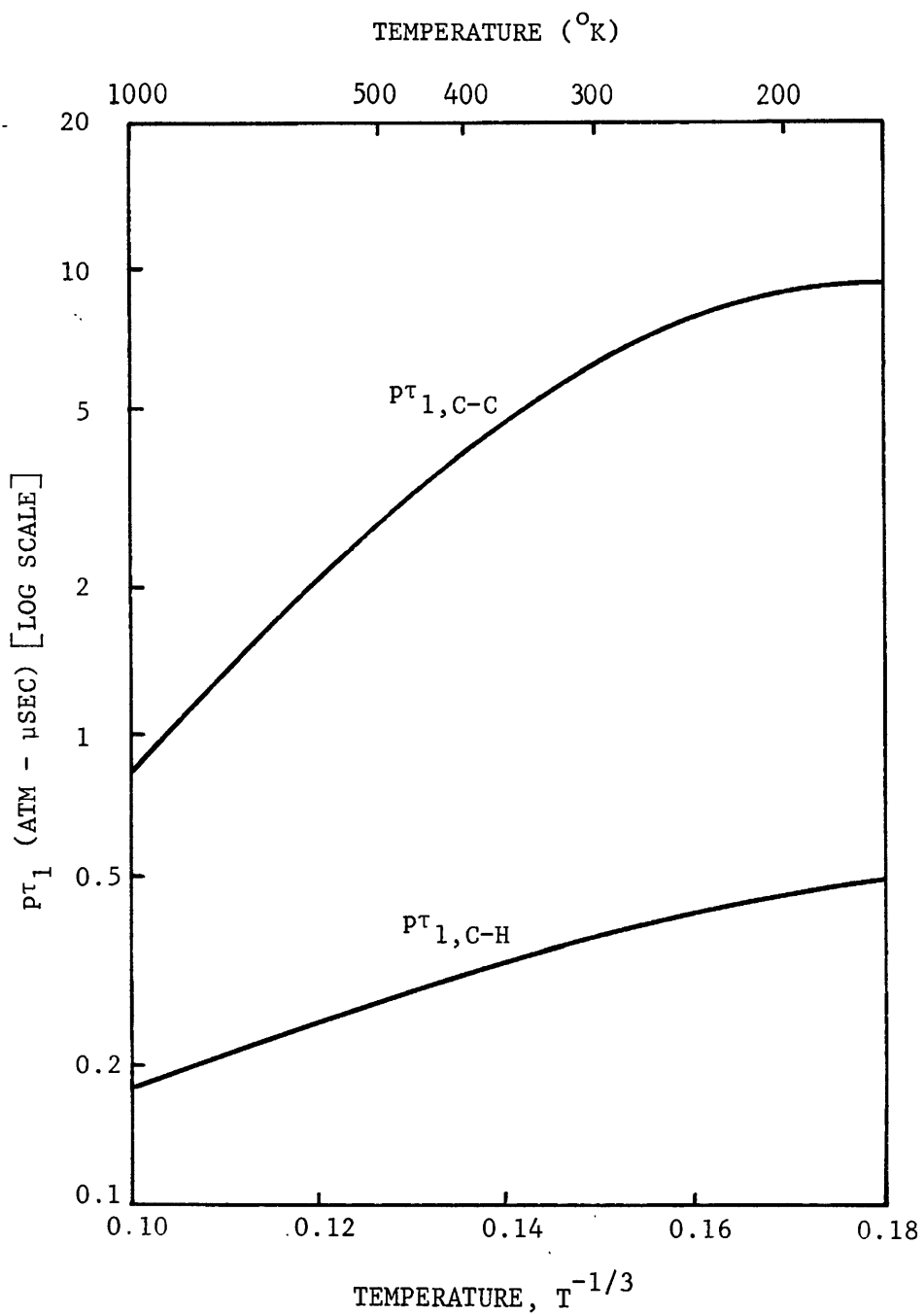


Figure 6. Temperature dependence of the relaxation lifetime of the (010) level, τ_1 .^{24,26-29}

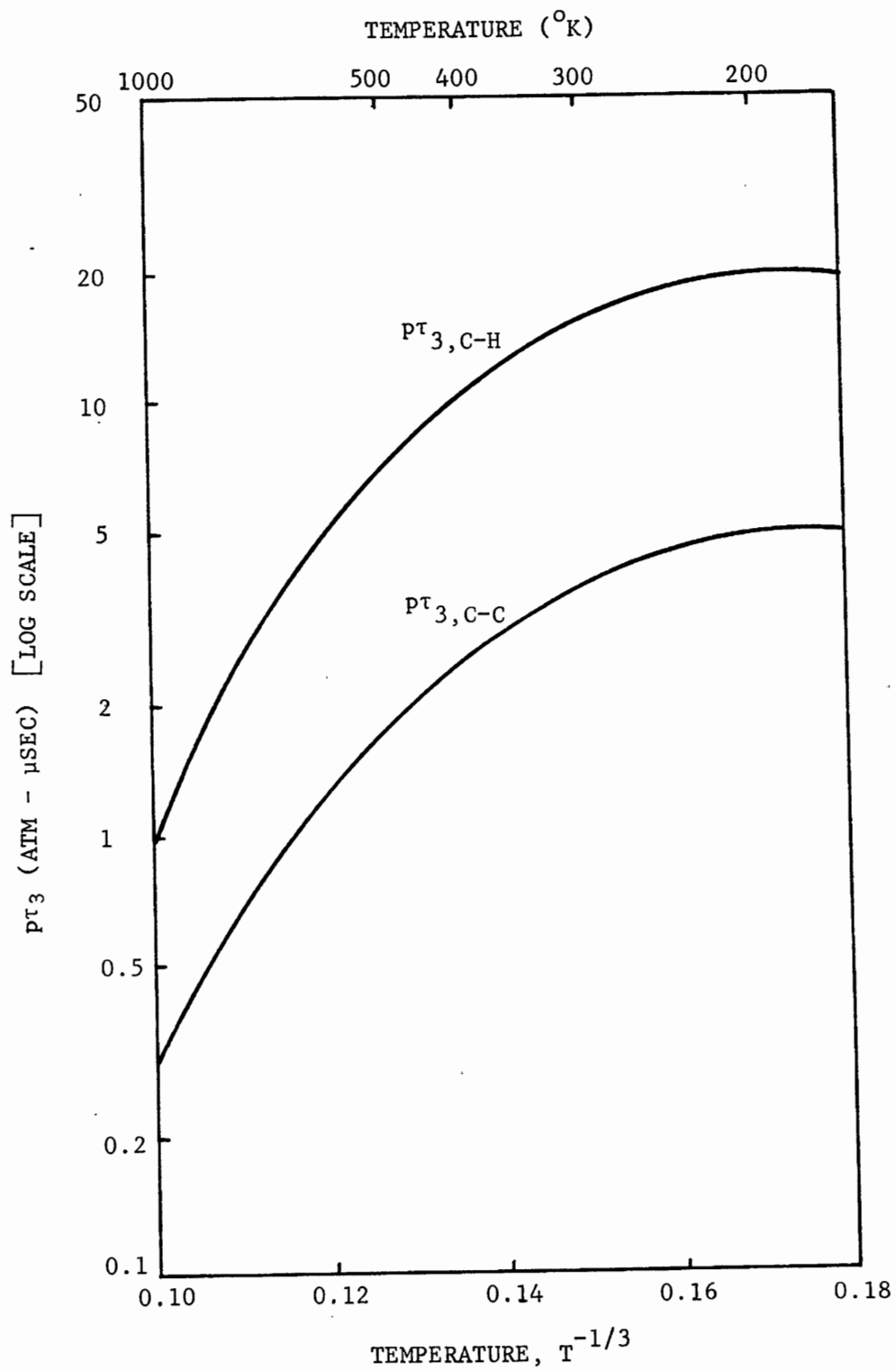


Figure 7. Temperature dependence of the relaxation lifetime of the (001) level, τ_3 .^{24,26-29}

Therefore, combining the equations (24) and (26)

$$\frac{1}{\tau_3} = \frac{T (^{\circ}\text{K})}{71 p (\text{atm}) r_m^2 (\text{cm}) [1 + 4 \ln (r/r_m)] + \left[\frac{\psi_C}{p\tau_{3,C-C}} + \frac{\psi_H}{p\tau_{3,C-H}} \right] p (\text{atm}) + \frac{1}{2.5 \times 10^{-3}}} \quad (29)$$

where

$$p\tau_{3,C-C} = 3.76 \text{ atm } \mu\text{sec at } 300^{\circ}\text{K}$$

$$p\tau_{3,C-H} = 15.5 \text{ atm } \mu\text{sec at } 300^{\circ}\text{K}$$

The values of $p\tau_{3,C-C}$ and $p\tau_{3,C-H}$ for 200°K are 1.2 times those of 300°K .

Thus, in summary the pertinent mode energies, level populations, and relaxation rates are functionally available to predict the actual laser gain and the power output for a given set of conditions. The necessary gain and output power relations which require the expressions presented in this section are discussed in the following section.

2.3 PREDICTION OF GAIN AND POWER OUTPUT

2.3.1 Minimum Pumping Power Required for Population Inversion

When an electromagnetic field is applied to atoms (or molecules), the interaction between the field and the atoms results in absorption or stimulated emission depending upon the relative population in the upper and lower levels. For any collection of atoms in thermal equilibrium, there are always more atoms in a lower energy level than in a higher energy level, and the transition is absorptive.

The essential feature of laser action is that under proper circumstances, by pumping or exciting the atoms in appropriate fashion, a non-equilibrium condition can be created which produces a temporary "population inversion" such that Boltzmann's law in terms of gas temperature does not apply. In order to achieve a population inversion the available total pumping power, P_T , must be greater than some minimum pumping power, P_{min} , where P_{min} is given by ⁷

$$P_{min} = \frac{N_1 V h \nu_p}{\tau_3} \quad (30)$$

where

$$\nu_p = c/\lambda_p = c/4.3\mu = \text{pumping frequency}$$

Thus,

$$R > \frac{N_1 h \nu_p}{\tau_3} \quad (31)$$

where

$$R = P_T/V = \text{pumping power density}$$

In terms of vibrational temperatures, the following relation has to be satisfied for population inversion of the vibrational levels,

$$\frac{N_3}{N_1} = \frac{N_3 e^{-\theta_3/T_3}/Q_v}{N_1 e^{-\theta_1/T_1}/Q_v} > 1 \quad (32)$$

so that

$$\frac{T_3}{T_1} > \frac{\theta_3}{\theta_1} = 1.76 \quad (33)$$

Relations (31) and (33) will be useful to roughly check the possibility of achieving lasing action.

2.3.2 Gain Coefficient

Electromagnetic radiation of intensity I propagating through an

amplifying medium (where a population inversion exists) with a gain coefficient γ will be amplified with an increase per unit length

$$\frac{dI}{dz} = \gamma I \quad (34)$$

Or integrating

$$I(z) = I_0 e^{\gamma z} \quad (35)$$

That is, the intensity grows exponentially.

From the principle of energy conservation, the increase in intensity per unit length, dI/dz , is equal to the net power generated per unit volume of length dz , i.e.,

$$\gamma I = g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) W h\nu \quad (36)$$

where W , the induced emission rate, is given by

$$W = \frac{\lambda^2 I g(\nu)}{8\pi h\nu t_{sp}} = \frac{\sigma I}{h\nu} \quad (37)$$

with

σ : optical cross section

$g(\nu)$: line shape

g_3, g_1 : degeneracy of rotational levels of upper and lower vibrational levels, respectively

$$g_3 = 2J + 1, \quad g_1 = 2J' + 1$$

$J' = J + 1$ for P-transitions (strongest gain)

n_3, n_1 : vibrational-rotational populations of upper and lower levels, respectively (cm^{-3})

Therefore, the expression of gain is obtained as

$$\gamma = \frac{\lambda^2 g(\nu)}{8\pi t_{sp}} g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) = \sigma g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) \quad (38)$$

where

t_{sp} : spontaneous emission lifetime (t_{sp} for the P-laser transition at 10.6μ is 5.38 sec)³⁰

$g(\nu)$: lineshape function which describes the distribution of emitted intensity versus the frequency ν ; it is normalized according to $\int_{-\infty}^{+\infty} g(\nu) d\nu = 1$

For the pressure broadening case (which applies to most of the pressure range in this study),

$$g(\nu_o) = \frac{\Delta\nu_L}{2\pi [(\nu-\nu_o)^2 + (\Delta\nu_L/2)^2]} = \frac{2}{\pi\Delta\nu_L} \quad (\text{since } \nu = \nu_o) \quad (39)$$

where $\Delta\nu_L$ is the Lorentzian linewidth which is given by the equation

$$\Delta\nu_L = \frac{1}{\pi \tau_c} = \frac{\nu_c}{\pi} \quad (40)$$

The collision frequency ν_c for CO_2 is in general given by

$$\nu_c = \sum_k N_k \sigma_{c-k} \bar{c}_{c-k} \quad (41)$$

where

N_k : the population of each species

σ_{c-k} : the collision cross section of each species with CO_2

\bar{c}_{c-k} : the mean relative velocity of the species

$$\bar{c}_{c-k} = \sqrt{\frac{8kT}{\pi m}}, \quad \frac{1}{m} = \frac{1}{m_c} + \frac{1}{m_k}$$

m_k : the mass of a molecule of each species

m_c : the mass of a CO_2 molecule

In the case of a CO_2 -He mixture

$$\nu_c = N_C \sigma_{C-C} \bar{c}_{C-C} + N_H \sigma_{C-H} \bar{c}_{C-H} \quad (42)$$

While the above can in general be used to calculate $\Delta\nu_L$, experimental values of σ_{C-H} are required. Recent experiments have resulted in the following formulation³¹

$$\Delta\nu_L = (9.96 \times 10^{10}) (\psi_C + 0.64 \psi_H) \quad (43)$$

The results obtained by using equation (43) are in good accord with the results obtained by using the more general equations noted above.

For the combined Doppler and pressure broadening case (Voigt Profile),^{32,33} which applies to the lowest pressure range of this study,

$$g(\nu_o) = \frac{2\sqrt{\ln 2}}{\sqrt{\pi} \Delta\nu_D} \exp(x^2) \operatorname{erfc}(x) \quad (44)$$

where

$$\Delta\nu_D : \text{Doppler Linewidth} = (3.05 \times 10^6) \sqrt{T} \text{ Hz for a CO}_2 \text{ line} \quad (45)$$

$$x = \sqrt{\ln 2} \Delta\nu_L / \Delta\nu_D \quad (46)$$

$\operatorname{erfc}(x)$: complimentary error function, whose values are

tabulated or given by

$$\operatorname{erfc}(x) = \frac{2}{\sqrt{\pi}} \int_x^\infty du e^{-u^2} \quad (47)$$

It is easily verified that the lineshape function reduces to the results for pure Doppler or pressure broadening, if the limit $x \rightarrow 0$ or $x \rightarrow \infty$ is taken in equation (44):

- 1) Doppler broadened limit: $x \rightarrow 0$

$$g(\nu_o) \rightarrow \frac{2\sqrt{\ln 2}}{\sqrt{\pi} \Delta\nu} \quad (48)$$

- 2) Pressure broadened limit: $x \rightarrow \infty$

$$g(\nu_o) \rightarrow \frac{2}{\pi \Delta\nu_L} \quad (49)$$

where the asymptotic relation $\exp(x^2) \cdot \operatorname{erfc}(x) \approx 1/(x\sqrt{\pi})$ was used.

Now recalling the relations for rotational and vibrational populations, as given by equations (21) and (9), the rotational and vibrational populations of the upper and lower levels are given by

$$\frac{n_3}{g_3 N_3} = \frac{e^{-J(J+1)\theta_R/T}}{Q_R} \quad (50)$$

$$\frac{n_1}{g_1 N_1} = \frac{e^{-J'(J'+1)\theta_R/T}}{Q_R} \quad (51)$$

$$N_3 = \frac{N e^{-\theta_3/T_3}}{Q_v} = \frac{\xi_3}{Q_v} \frac{E_3}{k\theta_3} \quad (52)$$

$$N_1 = \frac{N e^{-\theta_1/T_1}}{Q_v} = \frac{\xi_1}{Q_v} \frac{E_{12}}{k\theta_1} \quad (53)$$

where

N : total CO_2 population

N_3, N_1 : population densities of the upper and lower vibrational levels, respectively

n_3, n_1 : population densities of the upper and lower rotational-vibrational levels, respectively

T_3, T_1 : vibrational temperatures of the upper and lower levels, respectively

and where

$$E_3 = Nk\theta_3 / [e^{\theta_3/T_3} - 1] \quad (15)$$

$$E_{12} = E_1 + E_2 = Nk\theta_1 / \{1/[e^{\theta_1/T_1-1}] + 1/[e^{\theta_2/T_1-1}]\} \quad (54)$$

$$\xi_3 = [1 - e^{-\theta_3/T_3}] \quad (55)$$

$$\xi_1 = e^{-\theta_1/T_1} / [(e^{\theta_1/T_1-1})^{-1} + (e^{\theta_2/T_1-1})^{-1}] \quad (56)$$

$$Q_v = [(1 - e^{-\theta_1/T_1}) (1 - e^{-\theta_2/T_1})^2 (1 - e^{-\theta_3/T_3})]^{-1} \quad (12)$$

T_2 is assumed equal to T_1 (by Fermi resonance).

Hence, the gain can be expressed in terms of the energies of the modes characterized by the vibrational temperatures as follows

$$\gamma = \frac{\lambda^2 g(\nu)}{8\pi t_{sp}} g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) \quad (57)$$

$$= \frac{\lambda^2 g(\nu)}{8\pi t_{sp}} g_3 \left(\frac{n_3 N_3}{g_3 N_3} - \frac{n_1 N_1}{g_1 N_1} \right) \quad (58)$$

$$\approx \left(\frac{G_0}{N} \right) (N_3 - N_1) \quad \left(\text{since } \frac{n_3}{g_3 N_3} \approx \frac{n_1}{g_1 N_1} \right) \quad (59)$$

$$\gamma = \left(\frac{G_0}{N} \right) \frac{1}{kQ_v} \left[\frac{\xi_3}{\theta_3} E_3 - \frac{\xi_1}{\theta_1} E_{12} \right] \quad (60)$$

where

$$\left(\frac{G_0}{N} \right) = \frac{\lambda^2 g(\nu)}{8\pi t_{sp}} g_3 \frac{e^{-J(J+1)\theta R/T}}{Q_R} \quad (61)$$

2.3.3 Energy Balance Equations

In order to find E_3 and E_{12} and to get the final form of the gain expression, a set of energy balance equations among the vibrational modes will be introduced. Figure 8 below is the simplified energy level diagram for the energy equations.

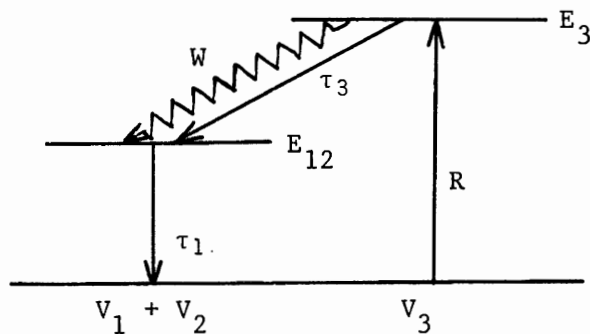


Figure 8. Simplified Energy Level Diagram.

Knowing that vibrational energy will always tend toward the equilibrium value and that the energy relaxation rate is expressed as $\frac{dE_v}{dt} = \frac{E_v - \bar{E}_v}{\tau}$, energy equations for the three modes in steady state are ³⁰

$$\frac{dE_3}{dt} = R - \frac{1}{\tau_3} [E_3 - \bar{E}_3] - \frac{\theta_3}{\theta_3 - \theta_1} g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) Wh\nu = 0 \quad (62)$$

$$\begin{aligned} \frac{dE_{12}}{dt} &= \frac{1}{\tau_3} \frac{3\theta_2}{\theta_3} [E_3 - \bar{E}_3] - \frac{1}{\tau_1} [E_{12} - \bar{E}_{12}] \\ &+ \frac{\theta_1}{\theta_3 - \theta_1} g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) Wh\nu = 0 \end{aligned} \quad (63)$$

where

$$g_3 \left(\frac{n_3}{g_3} - \frac{n_1}{g_1} \right) Wh\nu = \gamma I \quad (34)$$

However, $\bar{E}_3 \approx 0$ since the thermal population of the upper level is negligible compared with the pumped population. Then the equations for the energy balance and the gain are

$$R - \frac{E_3}{\tau_3} - \frac{\theta_3}{\theta_3 - \theta_1} \gamma I = 0 \quad (64)$$

$$\frac{1}{\tau_3} \frac{3\theta_2}{\theta_3} E_3 - \frac{1}{\tau_1} [E_{12} - \bar{E}_{12}] + \frac{\theta_1}{\theta_3 - \theta_1} \gamma I = 0 \quad (65)$$

$$\gamma = \left(\frac{G_0}{N}\right) \frac{1}{kQ_V} \left[\frac{\xi_3}{\theta_3} E_3 - \frac{\xi_1}{\theta_1} E_{12} \right] \quad (60)$$

Thus, there are the three unknowns E_3 , E_{12} , γ in the three equations above; solving for γ ,

$$\gamma = \frac{\left(\frac{G_0}{N}\right) \frac{1}{kQ_V} \left[\left(\frac{\xi_3}{\theta_3} \tau_3 - \frac{3}{2} \frac{\xi_1}{\theta_3} \tau_1\right) R - \frac{\xi_1}{\theta_1} \bar{E}_{12} \right]}{1 + \left(\frac{G_0}{N}\right) \frac{1}{kQ_V} I \left[\frac{\xi_3}{\theta_3 - \theta_1} \tau_3 - \frac{\xi_1}{2(\theta_3 - \theta_1)} \tau_1 \right]} \quad (66)$$

This is the general expression for the gain in terms of T_3 , T_1 , and I . For a small signal (where $I \approx 0$) in the energy equations, (64), (65), and (60)

$$E_3 = \tau_3 R = \frac{Nk\theta_3}{e^{\theta_3/T_3} - 1} \quad (67)$$

Therefore

$$T_3 = \frac{\theta_3}{\ln\left(\frac{Nk\theta_3}{\tau_3 R} + 1\right)} \quad (68)$$

Also,

$$\begin{aligned} [E_{12} - \bar{E}_{12}] &= \frac{3\theta_2}{\theta_3} \tau_1 R \\ &= \frac{kN\theta_1}{\xi_1} e^{-\theta_1/T_1} - \frac{kN\theta_1}{\xi_1'} e^{-\theta_1/T_1} \end{aligned} \quad (69)$$

where

$$\xi_1' = e^{-\theta_1/T} / \left[(e^{\theta_1/T} - 1)^{-1} + (e^{\theta_2/T} - 1)^{-1} \right] \quad (70)$$

therefore

$$T_1 = \frac{\theta_2}{\ln\left(\frac{1 + \sqrt{4\beta^2 + 8\beta + 1}}{2\beta}\right)} \quad (71)$$

where

$$\beta = \frac{1}{kN} \left(\frac{3}{2\theta_3} \tau_1 R \right) + (e^{\theta_1/T} - 1)^{-1} + (e^{\theta_2/T} - 1)^{-1} \quad (72)$$

The small signal gain γ_0 is

$$\gamma_0 = \left(\frac{G_0}{N}\right) \frac{1}{kQ_V} \left[\frac{\xi_3}{\theta_3} E_3 - \frac{\xi_1}{\theta_1} E_{12} \right] \quad (60)$$

$$= \left(\frac{G_0}{N}\right) \frac{1}{kQ_V} \left[\left(\frac{\xi_3}{\theta_3} \tau_3 - \frac{3}{2} \frac{\xi_1}{\theta_3} \tau_1\right) R - \frac{\xi_1}{\theta_1} \bar{E}_{12} \right] \quad (73)$$

Thus, for a small signal ($I \approx 0$), the values of the vibrational temperatures T_3 , T_1 and the gain (small signal gain) γ_0 can be calculated from equations (68), (71), and (73).

2.3.4 Saturation Intensity and the Maximum Available Power

When a population inversion is created in a laser such that an electromagnetic field (laser oscillation) builds up, the presence of that field reduces the population inversion and hence the gain coefficient. This effect is called gain saturation and is the mechanism which reduces the gain inside laser oscillators to a point where it just balances the losses so that steady oscillation can result. The maximum power of any laser is limited by saturation of the gain.

In a homogeneously broadened (pressure broadened) case,* the gain profile is uniformly saturated and the coefficient decreases with radiation intensity I according to the relation

$$\gamma = \frac{\gamma_0}{1 + \frac{I}{I_s}} \quad (74)$$

where I_s , the saturation intensity, defined as the intensity level that causes the inversion (and therefore the gain) to drop to one-half of its unsaturated value, will be given by comparing the gain expression obtained previously (equation (66)) and equation (74), i.e.

$$I_s = \frac{kQ_V}{\left(\frac{G_0}{N}\right)} \frac{1}{\left[\frac{\xi_3}{\theta_3 - \theta_1} \tau_3 - \frac{\xi_1}{2(\theta_3 - \theta_1)} \tau_1 \right]} \quad (75)$$

Now multiplying both sides of equation (74) by I , the net generated power per unit volume is

* Saturation behavior in the pressure range of this study is represented by the homogeneous broadening limit since $\sqrt{1 + I/I_s} \Delta\nu_L \geq \Delta\nu_D$.³² This is because broadening actually increases as intensity increases (intensity broadening).

$$IY = \frac{I\gamma_0}{1 + \frac{I}{I_s}} = I_s\gamma_0 \left(1 - \frac{I}{I + I_s}\right) \quad (74)$$

and its profile is plotted in Figure 9 below.

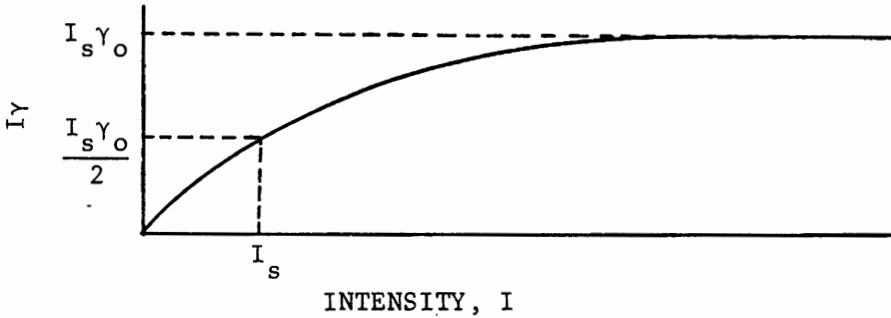


Figure 9. Net generated power per unit volume versus intensity.

As can be seen from Figure 9, the maximum available power per unit volume is $I_s\gamma_0$ (where $I \gg I_s$), the value of which is given by the expression below as obtained from equations (73) and (75).

$$\frac{P_{\max}}{V_m} = I_s\gamma_0 = \frac{\left(\frac{\xi_3}{\theta_3} \tau_3 - \frac{3}{2} \frac{\xi_1}{\theta_3} \tau_1\right)R - \frac{\xi_1}{\theta_1} \bar{E}_{12}}{\frac{\xi_3}{\theta_3 - \theta_1} \tau_3 - \frac{\xi_1}{2(\theta_3 - \theta_1)} \tau_1} \quad (77)$$

2.3.5 Cavity Losses and Output Coupling

So far we have not considered cavity losses such as those due to scattering, absorption, diffraction (these are called residual losses), and useful output transmission. But in order to figure the actual output power, we have to determine those losses.

Scattering and absorption by the mirrors will generally be small and is usually less than a percent (mainly by absorption). Diffraction is dependent on many parameters such as cavity length ℓ , radius of the mirrors, curvature of the mirrors, and the transverse mode configuration

of the laser beam. For the cavity dimensions found necessary in this study the diffraction loss can be held to less than a percent (see Section 3.1.1).

Regarding the output loss, it should be noted that there is a trade-off between the two extremes; i.e. if output mirror loss is too large, internal cavity losses are too large for laser oscillation, and on the other hand if the mirror transmission is too small, almost no useful power is coupled out.

The fraction of the intensity lost per single pass, L , is

$$L = L_i + T \quad (78)$$

where

L_i : residual loss per single pass

T : useful mirror transmission (average of both mirrors)

The optimum T to maximize the power output is given by³⁴

$$T_{\text{opt}} = -L_i + \sqrt{\gamma_o \ell L_i} \quad (79)$$

2.3.6 Threshold Gain and Output Power

Once the oscillation starts, the gain is reduced by gain saturation and clamped at the threshold value regardless of the pumping. At this point the gain per pass is equal to the loss per pass, i.e.

$$\gamma_t \ell = L \quad (80)$$

This is valid for small losses, i.e. $L \ll 1$.

From equation (80) and (74) at threshold, the intensity I can be written as

$$I = I_s \left(\frac{\gamma_o}{\gamma_t} - 1 \right) = I_s \left(\frac{\gamma_o \ell}{L} - 1 \right) \quad (81)$$

The cavity power considering losses is consequently given by

$$P_e = (IY_t)V_m = I_{s_m} A_m (\gamma_o \ell - L) \quad (82)$$

where $A_m = V_m/\ell$ is the cross-sectional area of the beam.

The fraction of the cavity power that is coupled out of the laser cavity as a useful output is T/L .

The power output is thus,

$$P_o = \frac{T}{L} P_e \quad (83)$$

Combining equations (73), (75), (82), and (83), P_o is finally written

as

$$P_o = \left\{ \frac{kQ_V}{\left(\frac{G_o}{N}\right)(\theta_3 - \theta_1)} \frac{1}{(\xi_3 \tau_3 - \frac{1}{2} \xi_1 \tau_1)} \right\} \frac{V_m}{\ell} T$$

$$\times \left\{ \frac{\left(\frac{G_o}{N}\right) \ell}{kQ_V \theta_3 L} \left[(\xi_3 \tau_3 - \frac{3}{2} \xi_1 \tau_1) \frac{PT}{V} - \frac{\theta_3}{\theta_1} \xi_1 \bar{E}_{12} \right] - 1 \right\} \quad (84)$$

3. RESULTS AND DISCUSSION

A computer program was written to facilitate the calculation of small signal gain γ_0 , maximum available power P_{\max} , cavity power P_e , output power P_o , optimum mirror transmission T_{opt} , efficiency ($\eta = P_o/P_T$) and vibrational temperatures T_3 and T_1 of the upper and lower levels for any set of parameters for the CO_2 -He laser system. A CO_2 -He mixture was chosen as the laser medium; addition of helium effectively depopulates the lower level while populating the upper level only by a slight degree. Nitrogen was not used because it provides no advantage to a sun-pumped CO_2 laser. The parameters that were varied included the gas temperature T , the total gas pressure p , the partial pressure of carbon dioxide p_c , the beam radius r_m , the radius of the tube r , the length of the tube ℓ , the absorption path length of the sunlight ℓ_a , the residual loss per pass L_1 , and the area of the solar energy collector A_s . Total pressure was varied in the range of $p = 0.01 - 0.05$ atm where acceptable gain and output power were found to exist. Gas temperatures of 200°K and 300°K were taken since at lower temperatures (about 170°K) carbon dioxide solidifies, and at temperatures above 300°K the gain was found to be too low for laser oscillation because of thermal population of the lower level.

Computed results are given first for a specific system model and then the effects on the laser performance of varying selected parameters are examined. The proposed system model is described in the following section.

3.1 PROPOSED LASER SYSTEM MODEL

3.1.1 Description of the System

End pumping was chosen because it provides the necessary long sunlight absorption path length in comparison to side pumping. Side pumping is of course possible as another alternative approach that could be studied. Although the length of the tube should be long to allow enough absorption, it cannot be too long because of practical size limitations and a higher diffraction loss. A tube of 1 m length was arbitrarily chosen with a 2 m absorption path length for the sunlight (i.e. one round-trip of the incident sunlight in the tube is assumed).

A collector size of 10 m^2 was also arbitrarily chosen as the largest reasonable collector size physically and economically practical. The minimum possible spot size concentrated from a 10 m^2 collector is limited to a 1.66 cm diameter area by the Abbe sine condition; therefore, the radius of the tube was determined to be 0.85 mm.

The radius of the beam should be large so that as many of the excited molecules as possible participate in lasing action. This is especially important for a sun-pumped CO_2 laser since the pump light is limited by collector size and should not be wasted. On the other hand, beam size for a given tube radius is limited by diffraction; the diffraction loss for a single mode beam is 1% for $r_m = \frac{2}{3}r$ and even larger for a multimode beam.³⁶ The beam radius is therefore taken to be 5 mm in the initial case.

The parameters for the laser tube and the collector area were thus selected as follows:

Area of the sun collector, $A_s = 10 \text{ m}^2$

Residual loss per pass, $L_i = 1 \%$

Beam radius, $r_m = 0.5 \text{ cm}$

Radius of the tube, $r = 0.85 \text{ cm}$

Length of the tube, $l = 1 \text{ m}$

Absorption path length of the sunlight, $l_a = 2 \text{ m}$

3.1.2 Computed Results for Proposed Laser

Computed results for small signal gain and output power are presented here as a function of the gas temperature and the total and partial pressures. The gain and power generally decrease as the gas temperature goes up because of the increase of the lower level population which is almost in equilibrium with the gas temperature. Thus, for the range of gas temperatures considered, the lowest possible temperature is optimum for this system.

3.1.2.1 Small signal gain: The variation of the gain with CO_2 partial pressure p_c is plotted in Figure 10 for 200°K with total pressure as a parameter. A total pressure of greater than 0.05 atm yields too low a gain and is thus not considered further. Gain is a function of many parameters such as linewidth, CO_2 concentration, pumping power, and upper and lower level lifetimes. Among these parameters, pumping power and upper level lifetime have the dominant effects on gain at 200°K , and these vary with the CO_2 and total pressures.

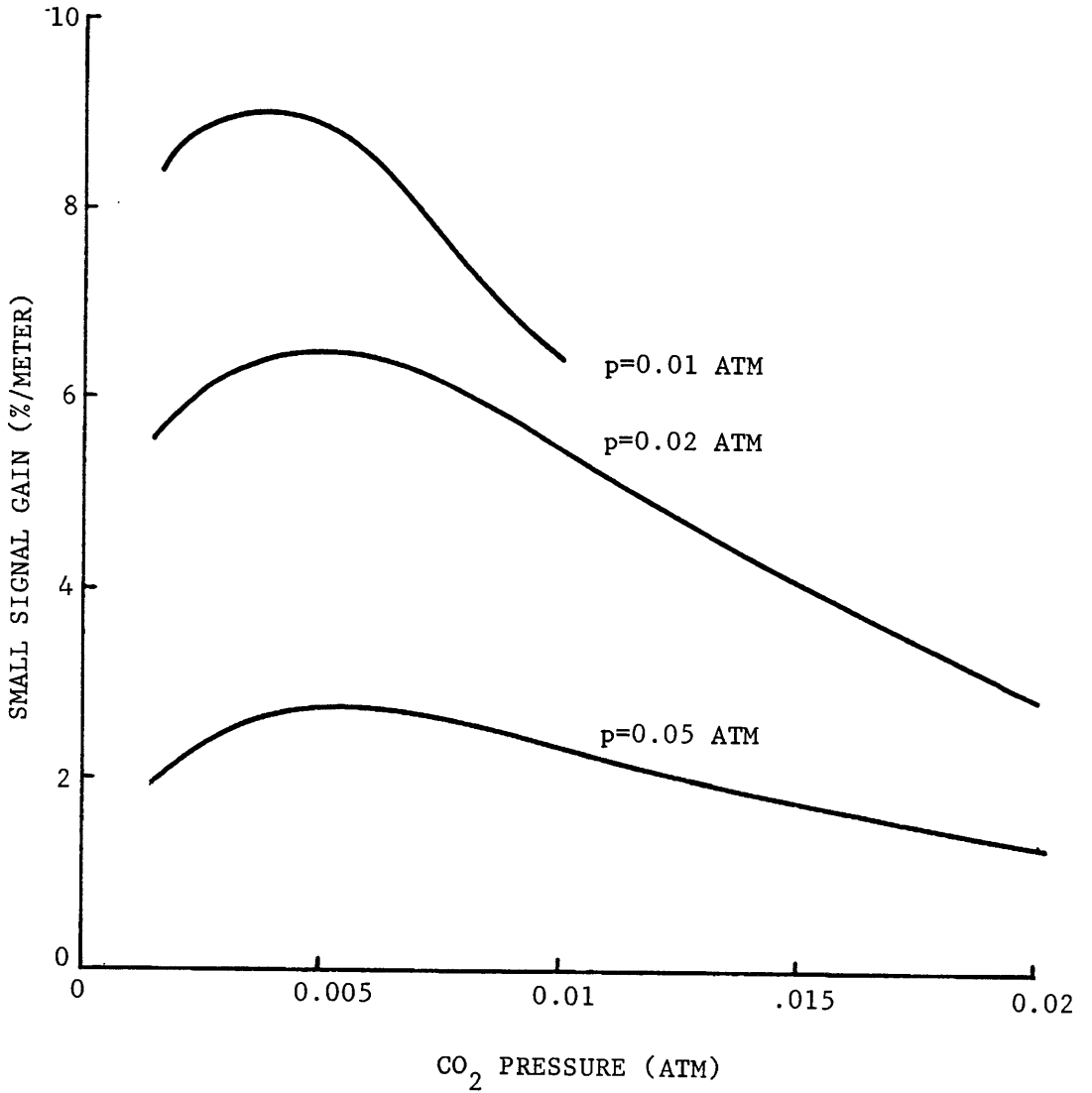


Figure 10. Variation of small signal gain with total and partial pressures.

As the CO_2 and total pressures increase, pumping power increases, but the upper level lifetime decreases. The maximum gain, then, occurs when the product of pumping power and upper level lifetime is a maximum. Because pumping is relatively weak such that the effect of the relaxation of the upper level predominates, maximum gain occurs at low partial and total pressures. At these low pressures thermal population of the lower level (100) is almost negligible compared with that of the pumped population of the upper level (001). Although the maximum gain is given at $p = 0.01$ atm and $p_c = 0.005$ atm, maximum output power is achieved at higher pressures (with high enough gain) as will be seen in the next section.

At 300°K thermalization of the lower level reduces the gain considerably to a value which is generally too low to be able to enable lasing action. A maximum gain of only 2.5%/meter was obtained in the 300°K case. Hence, the gas temperature was held at 200°K .

3.1.2.2 Output Power: Shown in Figure 11 is the plot of output power at 200°K . Output power is the power which comes out of the laser cavity through the partially transmitting mirror whose optimum transmission is calculated from equation (79). As one can see from expressions (82) and (83), output power varies with saturation intensity, small signal gain and residual losses as well as beam volume. Saturation intensity is a function of linewidth, pumping power, and upper and lower level lifetimes.

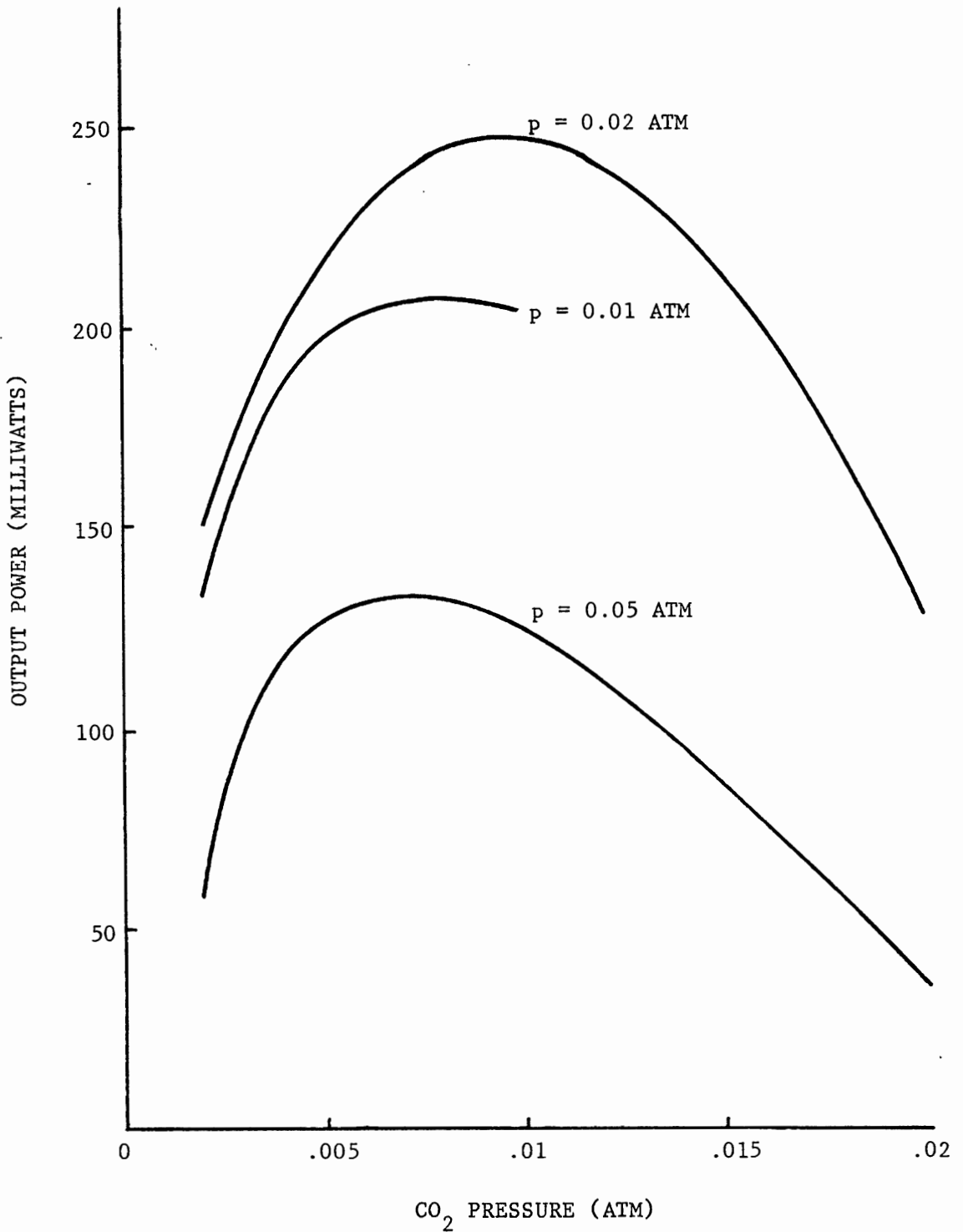


Figure 11. Variation of output power with total and partial pressures.

At 200°K a maximum output power of about 250 milliwatts was obtained at $p = 0.02$ atm and $p_c = 0.01$ atm with 1.3% mirror transmission. These pressures correspond to a rather low, non-optimum gain of 5.4%/meter, which would, however, still be sufficient to sustain laser oscillation.

3.1.2.3 Overall Performance: Enough gain and high output power are the requirements for good laser performance. Since the sun-pumped CO_2 laser is found to be a modest power, low gain system, it must operate at conditions where the gain is sufficient and the power is highest. The optimum performance of the assumed laser system is summarized as follows:

| | |
|------------------------------|-------------------------|
| Temperature: | 200°K |
| Total pressure: | 0.02 atm |
| CO_2 pressure: | 0.01 atm |
| Small signal gain: | 5.4%/meter or 5.4%/pass |
| Output power: | 250 milliwatts |
| Efficiency: | 4.1% |
| Optimum mirror transmission: | 1.3% |

3.2 EFFECTS OF PARAMETER VARIATION ON THE SMALL SIGNAL GAIN AND THE OUTPUT POWER

For the selected temperature of 200°K , various parameters that were preselected to initially determine the gain and the output power

are now varied to note the sensitivity of the performance to variations of the individual variables.

Each of the Figures 12-15 shows the variation of the gain and the output power due to the change of one parameter selected from the proposed system model; the six parameters changed are residual loss L_i , collector area A_s , radii of the beam r_m and the tube r , and the lengths of the absorption path ℓ_a and the tube ℓ .

The effect of varying residual losses is noted in Figure 12. The losses are taken to increase from 1% to 2% and then to 3%. The gain remains the same since it is independent of losses, but the output power decreases to 1/2 for $L_i = 2\%$, and to 1/3 for $L_i = 3\%$. Achieving losses of less than 1% would be difficult, but would result in a large power increase. Thus, the magnitude of residual loss is an important parameter.

The collector size directly affects the output power and gain as shown in Figure 13. As collector size varies, tube radius and beam radius are also adjusted accordingly. The tube radius was matched with the minimum spot size (by the Abbe Sine condition) concentrated from each collector area. The beam radius was kept slightly smaller than 2/3 of the tube radius for minimum diffraction loss. Since the pumping power density stays the same with varying collector size and, hence, tube radius (i.e. the maximum intensity of concentrated sunlight is limited by the Abbe Sine condition), the gain stays essentially constant. On the other hand, output power increases proportionally with collector size and, hence, with beam volume. For example, by reducing collector area by one-half (from 10 m^2 to 5 m^2) so that the beam volume

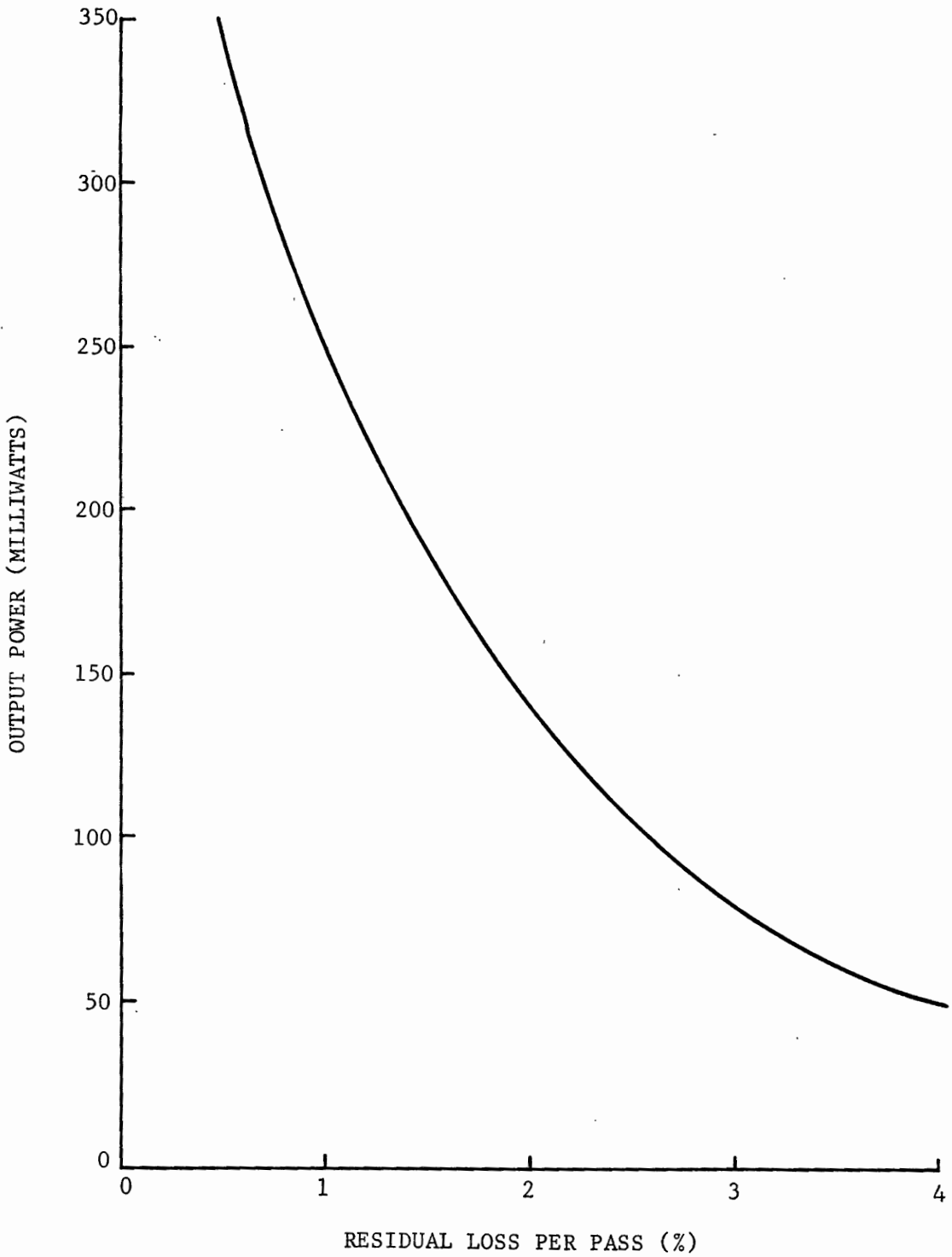


Figure 12. Output power versus residual loss per pass at 200°K.

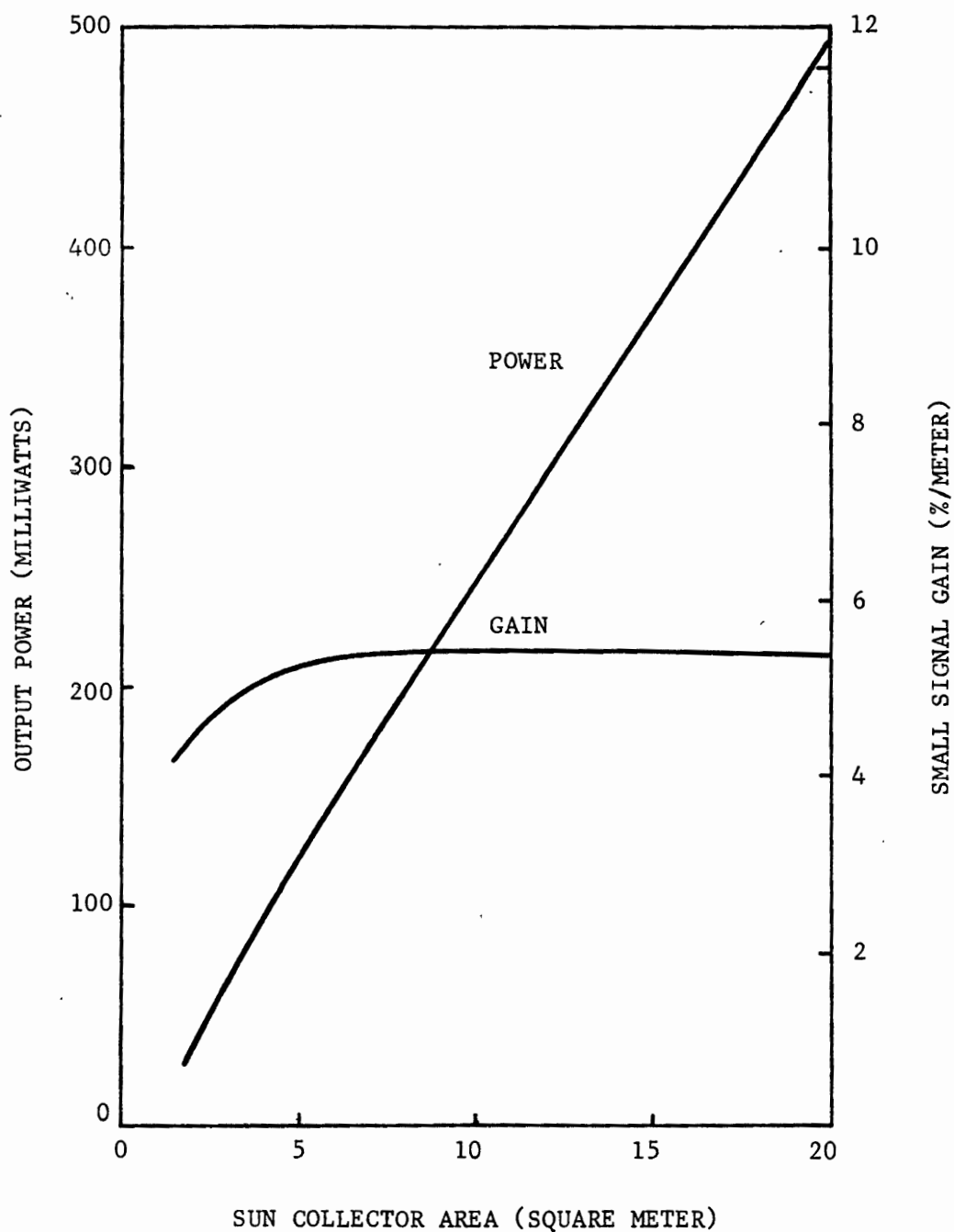


Figure 13. Output power and small signal gain versus sun collector area at 200°K.

(or beam spot size) is decreased by one-half, the output power goes down by one-half, while the pumping power density and, therefore, the gain stays constant. Although power increases with collector area, considering physical constraints, a collector much larger than 10 m^2 would not be very practical. Besides, a larger beam size is also required for a larger collector. That would also be hard to achieve.

For the constant collector (10 m^2) and the tube size (0.85 cm radius), only the beam size is varied in Figure 14. A smaller beam causes the pumping power to be wasted outside the beam volume, reducing the output power drastically. A beam larger than about 0.5 cm, on the other hand, will result in too high a diffraction loss which would lower the output. The results of Figure 14 do not include the effect of such increased losses.

As shown in Figure 15, both the gain and the output power are fairly strong functions of the absorption path length of sunlight. This result comes from the fact that in this low pressure range ($p = 0.02 \text{ atm}$) an absorption path length longer than 2 m would be necessary to completely absorb the incident sunlight. The output power approaches a maximum for an absorption length of about 8 m. An absorption length longer than 2 m would appear from Figure 15 to result in increased output power. However, the longer tube results in increased diffraction loss or a reduced mode volume that does not fully utilize the pump light. In fact, the output power is a maximum with about 2 m absorption length. It should be noted that gain per pass is plotted in Figure 15 rather than gain per meter. This is because comparison of gains should be made for single pass gain, not

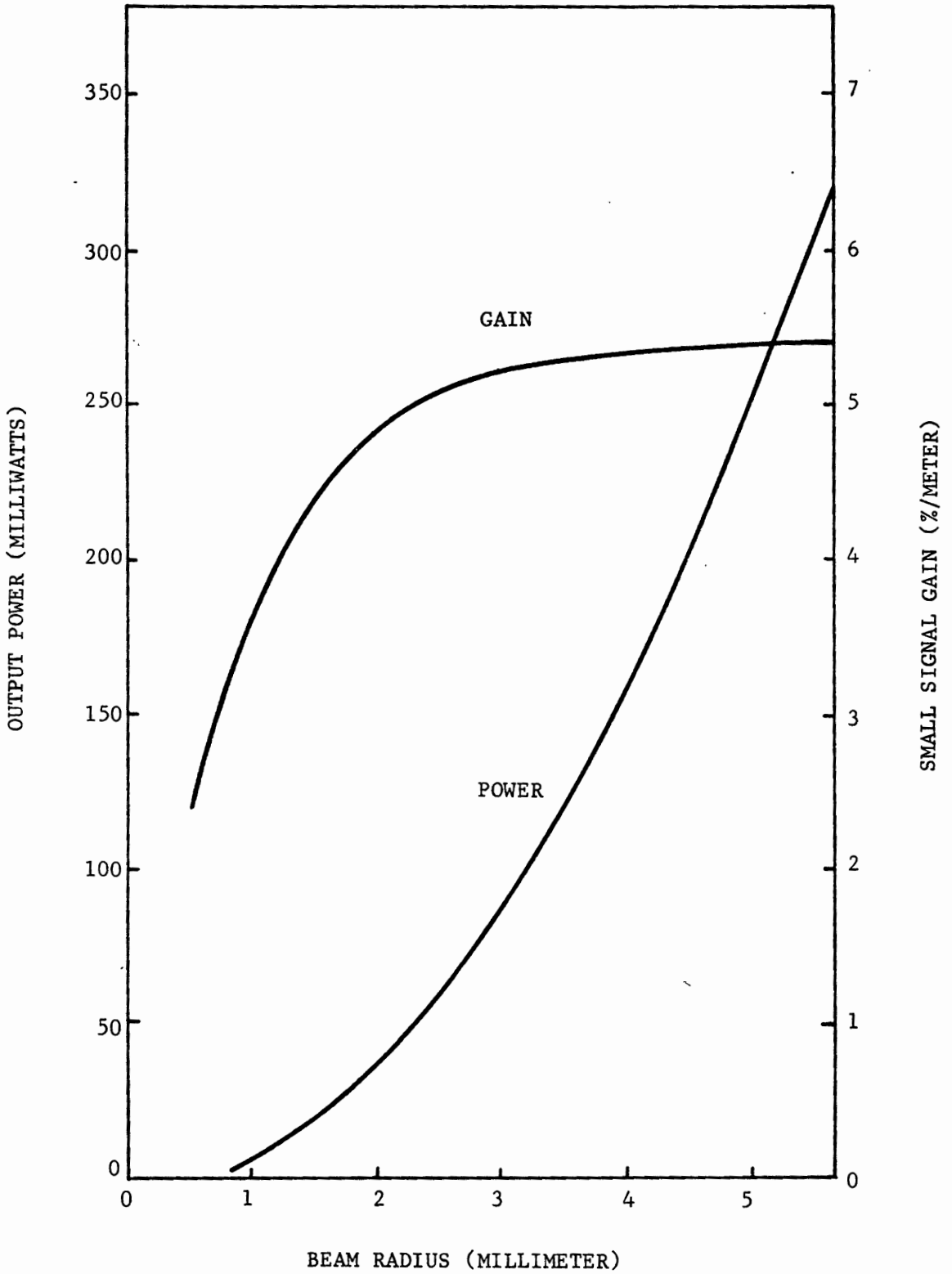


Figure 14. Output power and small signal gain versus beam radius.

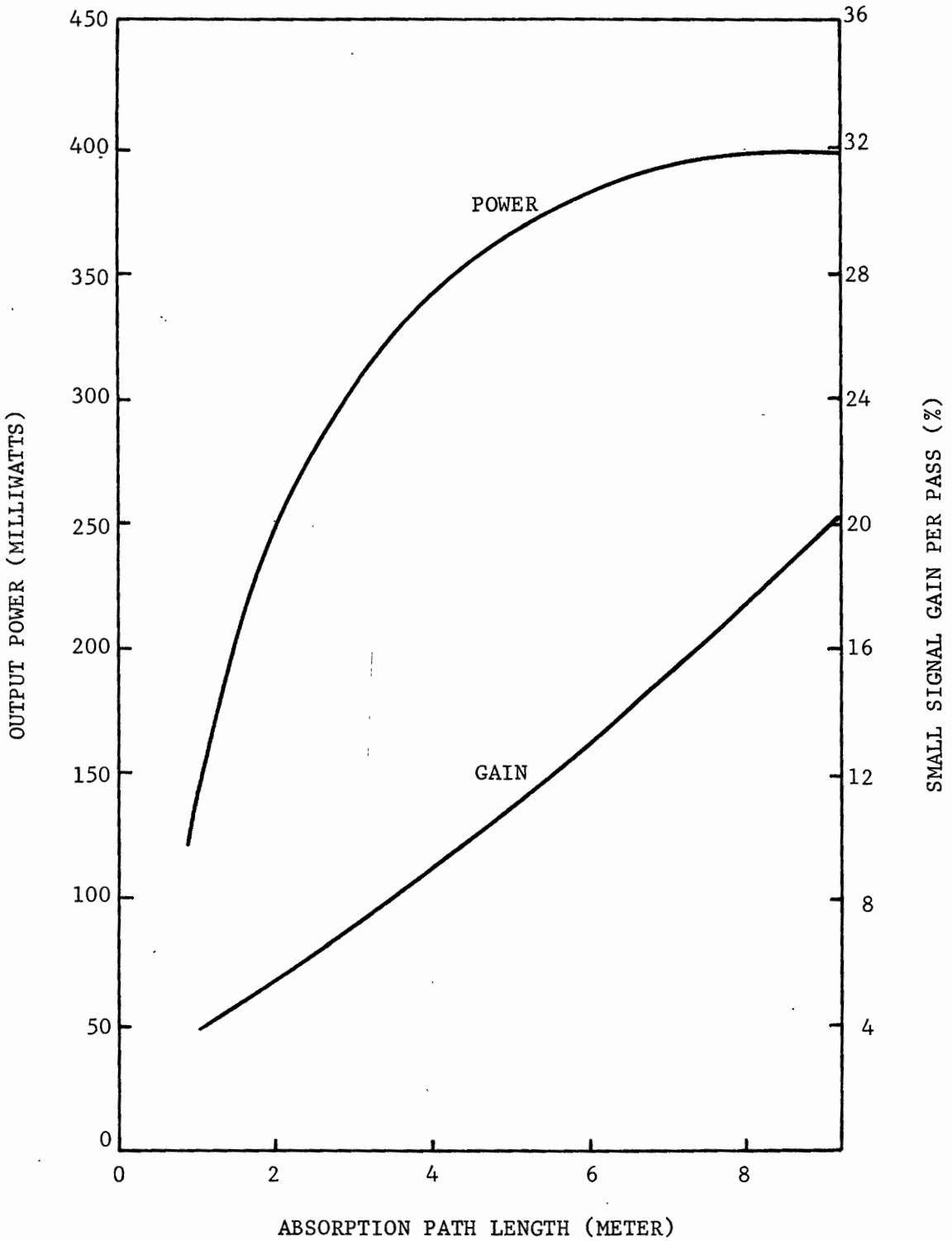


Figure 15. Output power and small signal gain per pass versus absorption path length at 200°K.

gain per unit length for different tube lengths.

In summary, when the various parameters are varied as above, the output power varies more than the gain. The initially proposed laser system in fact has the optimum power and reasonable gain. However, it is not likely that more than 500 milliwatts could be obtained from the sun-pumped CO₂ laser design presented here.

3.3 PRACTICAL DESIGN CONSIDERATIONS

In the foregoing discussions it has been assumed that the sunlight could appropriately be collected, concentrated, and then absorbed in the CO₂ gas mixtures. A schematic diagram of the end-pumping method assumed is shown in Figure 16 below.

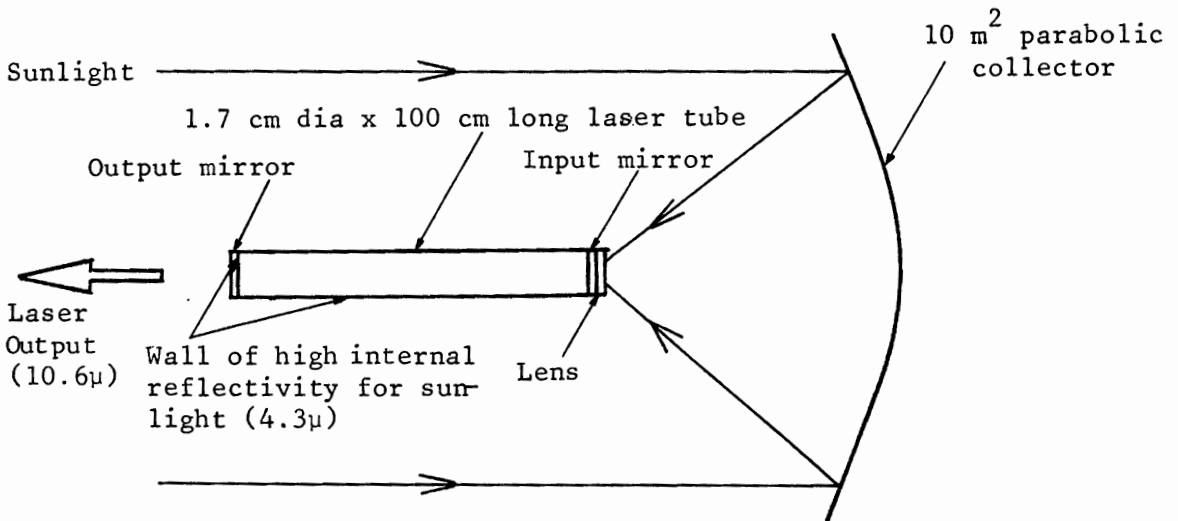


Figure 16. Schematic diagram of a sun-pumped CO₂ laser with end-pumping.

In this scheme sunlight collected by a large parabolic collector is focused into a lens that produces essentially parallel pump light which is transmitted into the laser tube. Other possible means of collecting and leading sunlight into the laser tube could be developed and used. For example, side-pumping might be another alternative method if long sunlight absorption path lengths could be achieved.

The laser output mirror and sunlight input mirror must be appropriately coated so that they transmit light selectively according to wavelength; the output mirror is partially transmitting for 10.6 micron laser light and reflective for 4.3 micron pumping light. The input mirror is transparent in the 4.3 micron region and reflective at 10.6 microns.

The other main practical consideration for a sun-pumped space-qualified CO₂ laser is cooling the laser medium. Conductive cooling of the laser tube and use of heat pipes to remove the heat to a radiative cooler is considered to be the preferred design technique for a space qualified laser because of its low weight and high reliability.^{1,5}

4. CONCLUSIONS

The possibility of directly sun-pumping a CW CO₂ laser in space has been analyzed for the first time, and the gas kinetic analysis indicates that end-pumping such a laser is feasible. However, since this laser system is pump deficient, it is essential to keep the residual losses such as diffraction, absorption, and scattering to a minimum, and also to concentrate the sunlight down to an almost theoretical minimum spot size in order to obtain the highest pumping power density. The gas temperature must also be held low (but higher than about 170°K) to achieve the highest possible gain and output power.

Since pumping must take place at low gas pressures to achieve sufficient gain, the power output is modest. In principle the output can be increased by utilizing a larger sunlight collector area, but practical difficulties with a larger collector and the necessarily larger laser beam diameter limit this approach. Long absorption paths could also be used to achieve higher powers, but the amount of improvement is again practically limited. It is possible that addition of other gases to the CO₂-He mixture could increase the output power by increasing the absorption of sunlight without adversely affecting the upper level lifetime. This possibility for increasing the output should be further examined. Nonetheless, at this time it would appear that the maximum achievable continuous power obtainable with the laser design presented here is about 500

milliwatts. Therefore, it does not look like high powers can be achieved using this approach. In addition, it should be noted that this particular type of CO_2 laser would not appear to be capable of operation at pressures above atmospheric where continuous tunability can be achieved.

It should be noted that output powers of greater than one watt have already been achieved with a sun-pumped Nd:YAG laser. However, if it is desirable or necessary to utilize a modest power carbon dioxide laser or if its output wavelength is preferred, then this study shows that such a laser is feasible. To examine its performance in more detail, laboratory measurements would have to be made to determine how much sunlight can actually be absorbed and how much gain can be achieved under the conditions predicted to give optimum performance.

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