

Multiphoton processes in two-level atoms in two intense pump beams

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The dynamical behavior of a two-level atom under the influence of two intense pump beams is examined. Exact equations for the nonlinear susceptibilities to all orders in the field strengths are derived. Numerical results for the nonlinear susceptibilities giving the absorption of the radiation from either of the pump beams and the generation of the four-wave mixing signal are given. The susceptibility for four-wave mixing, in addition to showing various saturation peaks, shows a peak at $\omega_1 = \omega_2$ even in the case of radiative relaxation. Higher-order multiphoton peaks in the absorption spectra are found to be extremely sensitive to the temporal fluctuations in the pumps.

1. INTRODUCTION

The dynamical behavior of a two-level atom in the presence of a single-mode field is well understood.¹ Such a dynamical behavior can be probed²⁻⁸ by using another external beam of a different frequency, assuming that the probe field is weak. Such probe-field absorption spectra are known² to show regions of the parametric gain provided that the pumping field is strong enough to saturate the two-level transition. The predictions on the weak-probe-field absorption spectra have been experimentally verified^{3,4} using both rf and optical fields. Some years ago Bonch-Bruевич *et al.* also investigated the effect of the temporal fluctuations⁴ of the pump field on the weak-field absorption spectra. Their results are in good agreement with the theoretical models of the pump-field fluctuations.^{9,10} Bonch-Bruевич *et al.* later carried out a series of experiments in which even the probe field was taken to be an intense one.⁸ The new spectra, when both pump and probe were intense, exhibited a number of remarkable features, for example, the appearance of the subradiative structures. Toptygina and Fradkin¹¹ have recently presented a theoretical explanation for the appearance of subradiative structure and various multiphoton absorption peaks. Since fluctuations of the pump beams were an important factor in the weak-field spectra, the question arises: What happens to the multiphoton absorption peaks if the temporal fluctuations of the two intense fields pump and probe¹² are taken into account? This is the object of our present theoretical study. Another problem that has attracted considerable attention is four-wave mixing¹³ in two-level atoms. If the probe and the pump fields are weak, then the four-wave mixing, the generation of a coherent signal at $2\omega_2 - \omega_1$, is described by the nonlinear susceptibility $\chi^{(3)}(-2\omega_2 + \omega_1, \omega_2, \omega_2, -\omega_1)$; such a nonlinear susceptibility in the usual perturbation treatment is independent of the field strengths. However, if the fields are strong enough to saturate the transition, then $\chi^{(3)}$ becomes intensity dependent. It is then desirable to know the variation of $\chi^{(3)}$ as a function of the intensities, relaxation parameters, etc. Our analysis also yields this information along with a host of other higher-order¹⁴ χ 's.

In Section 2 we formulate the basic problem. In order to keep the system of equations tractable, we adopt the phase diffusion model to describe the temporal fluctuations in each pump. Previous experience has shown that this model is quite adequate. Exact equations for the ensemble averages of various atomic operators are given. In Section 3 the results of the numerical calculations are given. The rate of the energy absorption from each field is calculated for arbitrary values of the intensities of the two fields. The effect of the field temporal fluctuations on the multiphoton absorption peaks is discussed in detail. In Section 4 we investigate numerically the behavior of $\chi^{(3)}$ and discuss the various saturation peaks that $\chi^{(3)}$ can show. We also discuss the effect of relaxation parameters on $\chi^{(3)}$ and show that, because of saturation, the type of extra resonance discussed by Bloembergen and co-workers¹⁵ can also arise even in the case of radiative relaxation.

2. BASIC DYNAMICAL EQUATIONS IN THE FIELD OF TWO FLUCTUATING BEAMS

In this section we formulate dynamical equations giving the behavior of a two-level atom with states $|1\rangle$ and $|2\rangle$ separated by $\hbar\omega_0$, in the field of two fluctuating fields. The fluctuations in each field are described by the phase diffusion model, i.e., we write the fields in the form

$$\begin{aligned} \mathbf{E}(t) = & \epsilon_1 \exp[-i\omega_1 t - i\Phi_1(t)] \\ & + \epsilon_2 \exp[-i\omega_2 t - i\Phi_2(t)] + \text{c.c.}, \quad (2.1) \\ = & \epsilon(t) + \text{c.c.}, \end{aligned}$$

with

$$\dot{\Phi}_i(t) = \mu_i(t), \quad i = 1, 2. \quad (2.2)$$

The μ_i 's are delta-correlated Gaussian random processes with zero mean:

$$\langle \mu_i(t) \rangle = 0, \quad \langle \mu_i(t) \mu_j(t') \rangle = 2\Gamma_{ij} \delta(t - t'). \quad (2.3)$$

The optical Bloch equations in the rotating wave approximation are given by

$$\frac{d}{dt} \begin{bmatrix} \rho_{21} \\ \rho_{12} \\ \rho_{11} - \rho_{12} \end{bmatrix} = \begin{bmatrix} i\omega_0 - \frac{1}{T_2} & 0 & +i\mathbf{d} \cdot \boldsymbol{\epsilon}^*(t) & \rho_{21} \\ 0 & -i\omega_0 - \frac{1}{T_2} & -i\mathbf{d} \cdot \boldsymbol{\epsilon}(t) & \rho_{12} \\ 2i\mathbf{d} \cdot \boldsymbol{\epsilon}(t) & -2i\mathbf{d} \cdot \boldsymbol{\epsilon}^*(t) & -\frac{1}{T_1} & \rho_{11} - \rho_{22} \end{bmatrix} + \begin{bmatrix} 0 \\ 0 \\ \frac{\rho_{11}^{(0)} - \rho_{22}^{(0)}}{T_1} \end{bmatrix}, \quad (2.4)$$

where \mathbf{d} is the dipole matrix element. On making the transformation to the rotating frame

$$\begin{aligned} \rho_{21} &= \exp[i\omega_2 t + i\Phi_2(t)]\psi_1, \\ \rho_{12} &= \exp[-i\omega_2 t - i\Phi_2(t)]\psi_2, \\ \rho_{11} - \rho_{22} &= 2\psi_3, \end{aligned} \quad (2.5)$$

we can write Eq. (2.4) in the following matrix form:

$$\begin{aligned} \dot{\psi} &= A\psi + i\mu_2(t)f\psi + I + B_+ \exp[i\{\Phi(t) + \Omega t\}]\psi \\ &\quad + B_- \exp[-i\{\Phi(t) + \Omega t\}]\psi, \\ \Omega &= \omega_1 - \omega_2, \quad \Phi = \Phi_1 - \Phi_2, \end{aligned} \quad (2.6)$$

where the nonvanishing elements of f , I , and B_{\pm} are

$$\begin{aligned} f_{11} &= -1, \quad f_{22} = +1, \quad I_3 = \eta \equiv (\rho_{11}^{(0)} - \rho_{22}^{(0)})/T_1, \\ (B_+)_{13} &= 2i\mathbf{d} \cdot \boldsymbol{\epsilon}_1^*, \quad (B_+)_{32} = -i\mathbf{d} \cdot \boldsymbol{\epsilon}_1^*, \\ (B_-)_{23} &= -2i\mathbf{d} \cdot \boldsymbol{\epsilon}_1, \quad (B_-)_{31} = i\mathbf{d} \cdot \boldsymbol{\epsilon}_1, \end{aligned} \quad (2.7)$$

and

$$A = \begin{bmatrix} i\Delta - \frac{1}{T_2} & 0 & 2i\mathbf{d} \cdot \boldsymbol{\epsilon}_2^* \\ 0 & -i\Delta - \frac{1}{T_2} & -2i\mathbf{d} \cdot \boldsymbol{\epsilon}_2 \\ i\mathbf{d} \cdot \boldsymbol{\epsilon}_2 & -i\mathbf{d} \cdot \boldsymbol{\epsilon}_2^* & -\frac{1}{T_1} \end{bmatrix}, \quad \Delta = \omega_0 - \omega_2, \quad (2.8)$$

We next perform the ensemble average over the fluctuations of the fields, i.e., over the phases Φ_1 and Φ_2 . For this purpose we define

$$\psi^{[n]} = \exp\{i[\Omega t + \Phi(t)]n\}\psi, \quad (2.9)$$

the equation of motion for which is given by

$$\dot{\psi}^{[n]} = A\psi^{[n]} + i\mu_2(t)f\psi^{[n]} + (i\Omega n + i\mu_1 n - i\mu_2 n)\psi^{[n]} + B_+\psi^{[n+1]} + B_-\psi^{[n-1]} + I \exp\{in[\Omega t + \Phi(t)]\}. \quad (2.10)$$

Ensemble averaging can now be done by using the theory of multiplicative stochastic processes as explained in Ref. 9, with the result that

$$\begin{aligned} \langle \dot{\psi}^{[n]} \rangle &= A\langle \psi^{[n]} \rangle + B_+\langle \psi^{[n+1]} \rangle + B_-\langle \psi^{[n-1]} \rangle + \delta_{n0}I \\ &\quad + [in\Omega - \Gamma_1 n^2 - \Gamma_2(f-n)^2 \\ &\quad - 2\Gamma_{12}n(f-n)]\langle \psi^{[n]} \rangle. \end{aligned} \quad (2.11)$$

where $\Gamma_1 = \Gamma_{11}$ and $\Gamma_2 = \Gamma_{22}$ represent the bandwidths of the beams at frequencies ω_1 and ω_2 , respectively, and Γ_{12} , represents the correlation between the two beams [Eqs. (2.3)]. The steady-state situation is simple as $\langle \dot{\psi}^{[n]} \rangle = 0$. These equations are strictly valid for $\Phi(t) \neq 0$, so that the well-defined steady states exist. Equation (2.11) is a recursion

relation involving $\langle \psi^{[n]} \rangle$ and $\langle \psi^{[n\pm 1]} \rangle$. We now reduce Eq. (2.11) to be a single equation for $\langle \psi_3^{[n]} \rangle$, whose solution is written in the form of a continued fraction.

A calculation using the components of Eq. (2.11) shows that, in the steady state,

$$\langle \psi_1^{[n]} \rangle = (2ig_2\langle \psi_3^{[n]} \rangle + 2ig_1\langle \psi_3^{[n+1]} \rangle)/D_n, \quad (2.12)$$

$$\langle \psi_2^{[n]} \rangle = (-2ig_2\langle \psi_3^{[n]} \rangle - 2ig_1\langle \psi_3^{[n-1]} \rangle)/D_{-n}^*, \quad (2.13)$$

with $g_i (\equiv \mathbf{d} \cdot \boldsymbol{\epsilon})$ representing coupling of the field at ω_i with the atom end

$$\begin{aligned} D_n &= \frac{1}{T_2} - i\Delta_2 - in\Omega + \Gamma_1 n^2 \\ &\quad + \Gamma_2(n+1)^2 + 2\Gamma_{12}n(n+1). \end{aligned} \quad (2.14)$$

The equation for $\langle \psi_3^{[n]} \rangle \equiv X_n$ has the form

$$a_n X_n + b_n X_{n+1} + c_n X_{n-1} = \eta \delta_{n0}, \quad (2.15)$$

where

$$\begin{aligned} a_n &= \frac{1}{T_1} + n^2(\Gamma_1 + \Gamma_2 - 2\Gamma_{12}) - in\Omega \\ &\quad + \frac{2g_2^2}{D_{-n}^*} + \frac{2g_1^2}{D_{n-1}} + \frac{2g_1^2}{D_{-n-1}^*} + \frac{2g_2^2}{D_n}, \\ b_n &= 2g_1 g_2 \left[\frac{1}{D_n} + \frac{1}{D_{-n-1}^*} \right], \\ c_n &= 2g_1 g_2 \left[\frac{1}{D_{n-1}} + \frac{1}{D_{-n}^*} \right]. \end{aligned} \quad (2.16)$$

The recursion relation [Eq. (2.15)] for X_n can be solved in terms of the continued fraction, with the result that

$$X_0 = \eta[a_0 + 2 \operatorname{Re}(b_0 Y_1)]^{-1}, \quad (2.17)$$

$$Y_n = -c_n [a_n + b_n Y_{n+1}]^{-1}, \quad Y_n \equiv X_n/X_{n-1}, \quad n \neq 0. \quad (2.18)$$

The system of equations is thus completely solved once the solution of Eqs. (2.17) and (2.18) is known. The off-diagonal elements of the density matrix are then obtained from Eqs. (2.12) and (2.13).

The rate of absorption of the energy W_i from the i th field is obtained by using

$$W_1 \cong 2\omega_1 g_1 \operatorname{Im} \langle \psi_1^{[-1]} \rangle, \quad (2.19)$$

$$W_2 \cong 2\omega_2 g_2 \operatorname{Im} \langle \psi_1^{[0]} \rangle. \quad (2.20)$$

We close this section by commenting on the case $\Phi = 0$, i.e., the case when both the lasers are completely correlated. In such a case Eq. (2.6) can be directly averaged as

$$\begin{aligned} \langle \dot{\psi} \rangle &= A\langle \psi \rangle - \Gamma_2 f^2 \langle \psi \rangle + I \\ &\quad + B_+ \exp(i\Omega t) \langle \psi \rangle + B_- \exp(-i\Omega t) \langle \psi \rangle, \end{aligned} \quad (2.21)$$

whose steady-state solution is

$$\langle \psi \rangle = \sum \langle \psi^{[n]} \rangle \exp(-in\Omega t), \quad (2.22)$$

with

$$A \langle \psi^{[n]} \rangle + B_+ \langle \psi^{[n+1]} \rangle + B_- \langle \psi^{[n-1]} \rangle + \delta_{n0} I + (in\Omega - \Gamma_2 f^2) \langle \psi^{[n]} \rangle = 0. \quad (2.23)$$

The set [Eq. (2.23)] is the same as Eq. (2.11) with $\Gamma_1 = \Gamma_2 = \Gamma_{12}$.

3. EFFECT OF PUMP TEMPORAL FLUCTUATIONS ON MULTIPHOTON PEAKS IN THE ABSORPTION SPECTRA

We now use the dynamical equations of Section 2 to study how the temporal fluctuations of the two pump fields affect the multiphoton absorption peaks. It is clear that $\langle \psi^{[n]} \rangle$ depends on the intensities of the two fields to all orders. $\langle \psi^{[n]} \rangle$ is expected to have the structure

$$\langle \psi^{[n]} \rangle = \sum_{n=0}^{\infty} g_1^{2n+1} \alpha_{2n+1}, \quad (3.1)$$

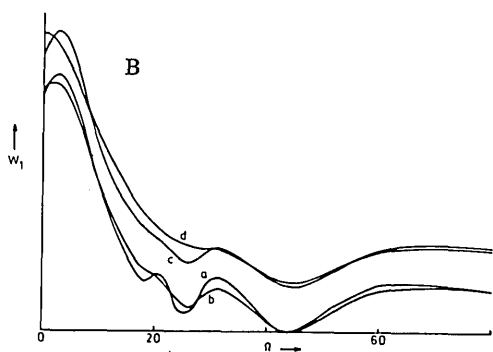
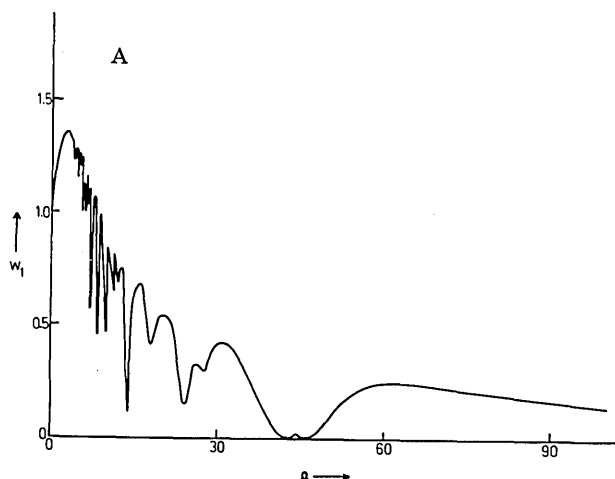


Fig. 1. A, rate of absorption $W_1 [\propto \langle \psi^{[n]} \rangle]$ as a function of Ω for $g_1 = 30, g_2 = 20, \Delta = 0, \Gamma_1 = \Gamma_2 = 0$, and $T_1 = T_2 = 1$. B, W_1 for various values of Γ_1 and Γ_2 as a function of Ω for $g_1 = 30, g_2 = 20, \Delta = 0$, and $T_1 = T_2 = 1$. Curve a, $\Gamma_1 = 0.4, \Gamma_2 = 0.0$. Curve b, $\Gamma_1 = \Gamma_2 = 0.4$. Curve c, $\Gamma_1 = 0.0, \Gamma_2 = 0.8$. Curve d, $\Gamma_1 = \Gamma_2 = 0.8$. Note that the spectrum for $\Gamma_1 = 0.0$ and $\Gamma_2 = 0.4$ is similar to that for curve a. For convenience of illustration, curves c and d have been shifted.

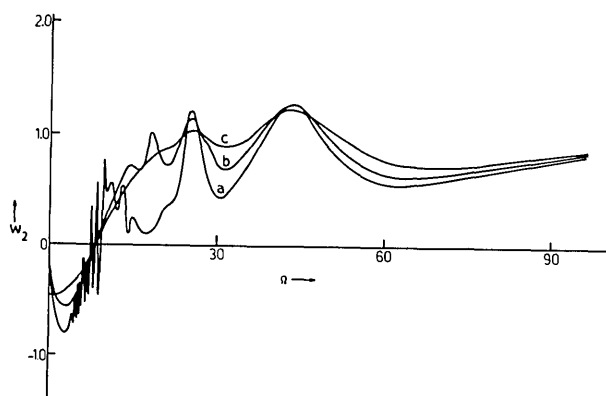


Fig. 2. Absorption spectrum $W_2 [\propto \langle \psi^{[0]} \rangle]$ as a function of Ω for $g_1 = 30, g_2 = 20, \Delta = 0$, and $T_1 = T_2 = 1$. Curve a, $\Gamma_1 = \Gamma_2 = 0.0$. Curve b, $\Gamma_1 = 0.2, \Gamma_2 = 0.0$. Curve c, $\Gamma_1 = 0.8, \Gamma_2 = 0.0$.

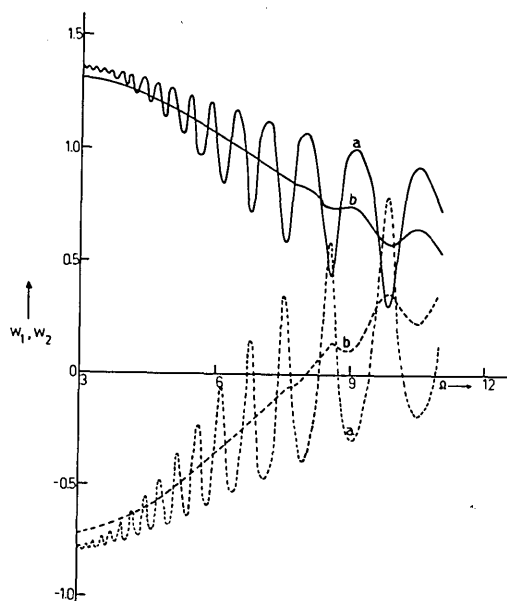


Fig. 3. Multiphoton absorption region of the spectrum W_1 (solid line) and W_2 (dashed line) for $g_1 = 30, g_2 = 20, \Delta = 0$, and $T_1 = T_2 = 1$. Curve a, $\Gamma_1 = \Gamma_2 = 0.0$. Curve b, $\Gamma_1 = 0.05, \Gamma_2 = 0.0$.

where α 's depend on g_2^2 to all orders. α_1 represents the absorption of a single photon from the beam at ω_1 in the presence of another field of arbitrary strength and this is the quantity measured in the experiments of Wu *et al.*³ The field ϵ_2 can be imagined to dress the eigenstates of the atom. Thus α_{2n+1} will represent the absorption of $(n + 1)$ photons of frequency ω_1 by the atom dressed by the field ϵ_2 . The effect of the temporary fluctuations on α_1 is well known both experimentally and theoretically.

In the case when both the pumps are completely correlated, $\Gamma_1 = \Gamma_2 = \Gamma_{12} = \gamma$, then the multiphoton absorption spectra in the presence of fluctuations can be obtained from those in the absence of fluctuations by using the simple substitution rule¹⁶

$$\frac{1}{T_2} \rightarrow \frac{1}{T_2} + \Gamma. \quad (3.2)$$

The general case is extremely involved, and it does not appear to be possible to obtain a simple result even for the line center $\omega_1 = \omega_2 = \omega_0$. Hence we have numerically investigated the

solution of Eq. (2.18), and the results are shown in a number of figures. The parameters (all in frequency units) used in our numerical computations are based on the ones used in the experimental investigations of the Russian workers.^{8,11} We also assume that the two fields are uncorrelated, $\Gamma_{12} = 0$. Figure 1 gives the behavior of the absorption spectra for the absorption of the energy from field 1 for $g_1 > g_2$ and when both the fields are strong. The figure shows that the multiphoton absorption peaks are wiped out with increase in Γ_1 . The effect of Γ_2 is similar. Only the peak corresponding to the frequency $2(g_1^2 + g_2^2)^{1/2}$ and the first few multiphoton peaks survive. The behavior of the energy absorption from field 2 is similar, as is shown in Fig. 2. Figure 3 shows in detail the effect of the laser fluctuations on the multiphoton absorption peaks. In order to see these absorption peaks, the laser bandwidths must not be more than few percent of the natural line width. Figure 4 is for much larger values of the bandwidths, whence only the structure at the line center survives. The case when $g_1 < g_2$ is illustrated in Fig. 5.

The extreme sensitivity of the multiphoton absorption peaks on the temporal fluctuations of pump beams can be understood from the structure of the coefficients D_n [defined

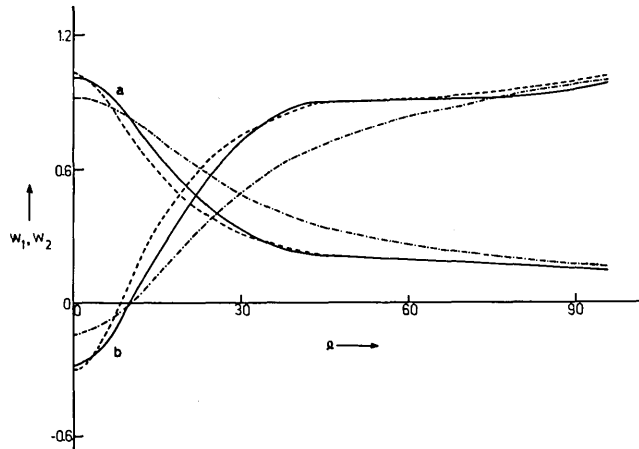


Fig. 4. Absorption spectrum W_1 (curves a) and W_2 (curves b) for high values of Γ_1 and Γ_2 . $g_1 = 30$, $g_2 = 20$, $\Delta = 0$, and $T_1 = T_2 = 1$. (Solid line, $\Gamma_1 = 10.0$, $\Gamma_2 = 0.0$; dashed line, $\Gamma_1 = 0.0$, $\Gamma_2 = 10.0$; and dashed-dotted line, $\Gamma_1 = \Gamma_2 = 10.0$.)

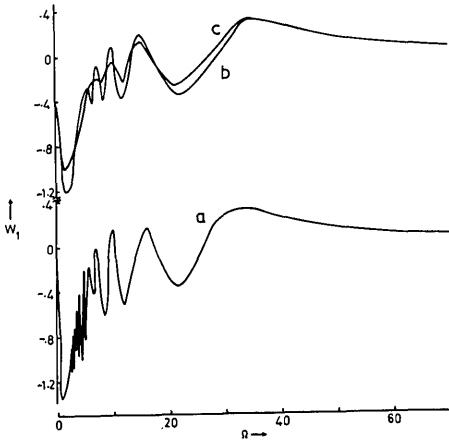


Fig. 5. Rate of absorption W_1 for $g_2 > g_1$. The numerical values are $g_1 = 9.52$, $g_2 = 11.9$, $\Delta = 0$, and $T_1 = T_2 = 1$. The various values for Γ_1 and Γ_2 are curve a, $\Gamma_1 = \Gamma_2 = 0.0$; curve b, $\Gamma_1 = 0.0$, $\Gamma_2 = 0.05$; curve c, $\Gamma_1 = 0.0$, $\Gamma_2 = 0.2$.

by Eq. (21.4)]. The position and the width of the multiphoton absorption peaks are approximately determined from the zeros of a_n . For large n and in the absence of any temporal fluctuations, these are extremely narrow—for $T_1 = T_2$ these are located at $\pm(g_1^2 + g_2^2)^{1/2}/n$ with widths $\sim(1/nT_2)$. Note that the laser temporal fluctuations (bandwidths) enter as $n^2\Gamma$ in the coefficients D_n and a_n . Hence these multiphoton absorption peaks acquire an additional width of the order of $n\Gamma$. Since the relative separation of such peaks is of the order of gT_2/n , it is clear that a small fluctuation will wash away such multiphoton structures.

4. EFFECT OF SATURATION ON THE NONLINEAR SUSCEPTIBILITY $\chi^{(3)}(-2\omega_2 + \omega_1, \omega_2, \omega_2, -\omega_1)$

As was mentioned in the introduction, $\chi^{(3)}$ yields the four-wave mixing signal in the direction $2\mathbf{k}_2 - \mathbf{k}_1$. The theory described in Section 2 is fairly general and should also enable us to calculate $\chi^{(3)}$. We show that $\chi^{(3)}$ has an interesting saturation behavior. In order to keep the analysis simple, we ignore the complications that arise because of finite temporal fluctuations¹⁷ of the fields. From Eqs. (2.22) and (2.5) it is clear that

$$\rho_{12}(t) = \sum_n \psi_2^{[n]} \exp\{-it[\omega_2(1-n) + n\omega_1]\} \quad (4.1)$$

$$= \psi_2^{[-1]} \exp[-it(2\omega_2 - \omega_1)] + \text{other terms.} \quad (4.2)$$

The four-wave mixing signal is related to the induced polarization at $2\omega_2 - \omega_1$. The induced polarization is proportional to ρ_{12} , and thus the component responsible for four-wave mixing is $\psi_2^{[-1]} = \psi_1^{[1]*}$. Note that, in the limit of weak fields (Rabi frequencies \ll detunings), the induced polarization at $2\omega_2 - \omega_1$ is known¹³ to be proportional to $\epsilon_2^2 \epsilon_1^*$, and hence $|\psi_1^{[1]}|^2 \propto |\epsilon_2|^4 |\epsilon_1|^2$. In general, $\psi_1^{[1]}$ will depend on all powers of ϵ . The component $\langle \psi_1^{[1]} \rangle$ is given by Eq. (2.12) with $n = 1$, which when Eq. (2.18) is used, becomes

$$\langle \psi_1^{[1]} \rangle = 2i(g_2 + g_1 Y_2) Y_1 X_0 D_1^{-1}. \quad (4.3)$$

Thus the continued fraction of Section 2 will give the four-wave mixing signal if Eq. (4.3) is used. The continued fraction can be summed up exactly for $\omega_1 = \omega_2$ leading to

$$Y_1 = -\frac{a}{2b} \left[1 - \left(1 - \frac{4cb}{a^2} \right)^{1/2} \right], \quad (4.4)$$

where

$$a = \frac{1}{T_1} + \frac{4(g_1^2 + g_2^2)}{T_2 \left[\Delta_2^2 + \left(\frac{1}{T_2} \right)^2 \right]}, \quad b = c = \frac{4g_1 g_2}{T_2 \left[\left(\frac{1}{T_2} \right)^2 + \Delta_2^2 \right]}. \quad (4.5)$$

Bloembergen *et al.*¹⁵ have shown that S , in the limit of weak fields, has an interesting resonance at $\omega_1 = \omega_2$ with a width T_1^{-1} provided that the system has collisions; otherwise, for the case of radiative relaxation, S has no structure. This is illustrated in Fig. 6, curve a. We also show in Fig. 6 (curves b and c) the effect of the saturation, which leads to the peak at $\omega_1 = \omega_2$, even in the case of the radiative relaxation,¹⁸ in

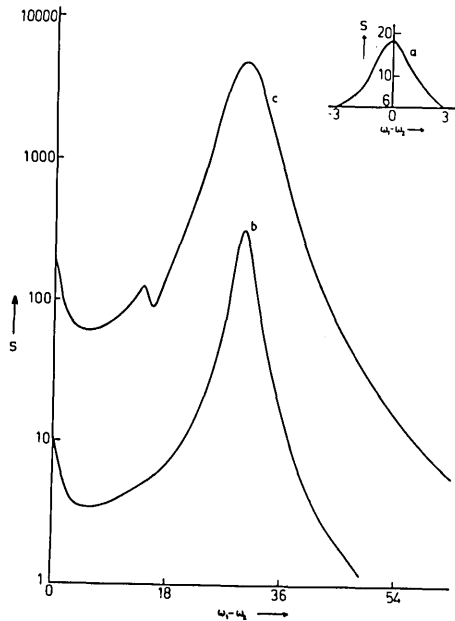


Fig. 6. Four-wave mixing signal $S(\alpha|\psi_1^{(1)}|^2)$ as a function of $(\omega_1 - \omega_2)$ for curve a, inset: $\Delta = 200$, $T_1 = T_2 = 1$, $g_1 = 4$, and $g_2 = 15$ and for $T_1 = 0.5$, $T_2 = 1.0$, $\Delta = 0$; curve b, $g_1 = 1$, $g_2 = 15$; curve c, $g_1 = 4$, $g_2 = 15$.

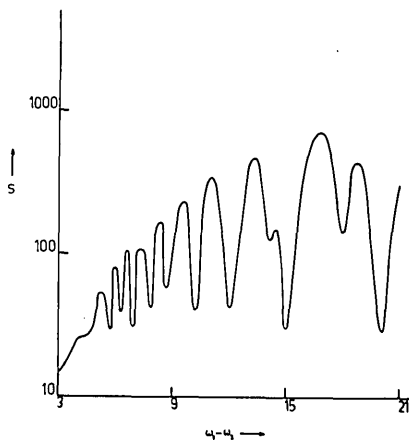


Fig. 7. Multiphoton absorption region of S as a function of $(\omega_1 - \omega_2)$ for $g_1 = 30$, $g_2 = 20$, $T_1 = T_2 = 1$, and $\Delta = 0$.

addition to the multiphoton (saturation) peaks.¹⁹ The effect of collisions is not dramatic if the saturation effects are important. Note that, as the field intensities are increased

further, more and more multiphoton absorption peaks start appearing as shown in Fig. 7.

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