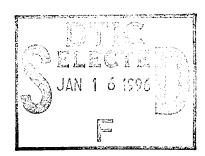
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THEORY AND NUMERICAL VALUE CALCULATIONS ASSOCIATED WITH STIMULATED RAMAN SCATTERING IN THE ATMOSPHERE

by

Lin Dewen





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ABSTRACT Starting out from the basic principles of quantum mechanics, the dynamics equations for stimulated Raman scattering (SRS) under semi-classical theoretical conditions are obtained. Theoretical verification is done of the fact that classical descriptions are only appropriate provided the distance between upper and lower energy levels is relatively great. Numerical simulations are done of several types of effects associated with stimulated rotational Raman scattering (SRRS) when strong lasers pass through the atmosphere. In these are included influences associated with Raman rotational properties, SRRS threshold values, and molecular "memory" on pulse series transmission.

KEY WORDS Stimulated rotational Raman scattering Semiclassical theory Transforming threshold Pulse series

I INTRODUCTION

In research on stimulated Raman scattering, particle systems are normally described by the use of classical resonance sub-equations [1]

$$\frac{\partial Q^{\bullet}}{\partial t} + \Gamma Q^{\bullet} = \frac{i\alpha'}{4\omega, M} \widetilde{E}_{L}^{\bullet} \widetilde{E}_{s}$$
 (1)

As far as stimulated vibratory Raman scattering (SVRS), which occurs in molecules under outside field effects, is concerned, opting for the use of classical descriptions discussed above can be understood. It has already been clearly demonstrated that, when strong laser light passes through the atmopshere, SRRS is primarily involved. With respect to this type of molecular transition between rotation states, there is no corresponding classical model. There are also problems awaiting investigation as to whether or not it is possible to make use of classical resonance sub-equation (1) to apply descriptions. From the fundamental principles of quantum mechanics $\partial \hat{\rho} / \partial t = -i[\hat{H}, \hat{\rho}]/\hbar$, this article sets out to solve $p = Tr(\hat{\rho}\hat{\mu})$ and particle field Bloch equations and field polarization p. Moreover, fields themselves still obey classical wave equations.

2 DYNAMICS EQUATION SETS ASSOCIATED WITH SEMI-CLASSICAL THEORIES OF

STIMULATED RAMAN SCATTERING

2.1 Particle System Bloch Equations

Assume that particle systems possess the three energy levels shown in Fig.1. As far as lasers in TEMoo are concerned,

Fig. 1
$$\frac{1}{\Omega_{L}} = \varepsilon_{L} \cos \psi_{L} = \frac{\varepsilon_{L}}{2} \left(e^{i\psi_{L}} + e^{-i\psi_{L}} \right) \\
\psi_{L} = \omega_{L} t - k_{L} z + \Phi_{L}$$
(2)

Under these effects, Raman scattering will occur. The scattered light is

$$E_{s} = \varepsilon_{s} \cos \psi_{s} = \frac{\varepsilon_{s}}{2} \left(e^{i\psi_{s}} + e^{-i\psi_{s}} \right)$$

$$\psi_{s} = \omega_{s} t - k_{s} z + \Phi_{s}$$
(3)

(Illegible) system Hamilton quantities are

$$\hat{H} = \hat{H}_0 + \hat{H}_1$$

$$\hat{H}_{0} = \begin{bmatrix} \hbar \omega_{1} & 0 & 0 \\ 0 & \hbar \omega_{2} & 0 \\ 0 & 0 & \hbar \omega_{3} \end{bmatrix} \qquad \hat{H}_{1} = \begin{bmatrix} 0 & 0 & V_{L} \\ 0 & 0 & V_{S} \\ V_{L}^{*} & V_{S}^{*} & 0 \end{bmatrix}$$

Dipole transition $V_{\alpha} = -\mu_{x}E_{\alpha}$, $(\alpha = L, S)$, Dipole

transition matrix elements $\mu_L = -\langle 1|q\,r|3\rangle$, $\mu_s = -\langle 2|q\,r|3\rangle$. The atmosphere is primarily composed of dual homonuclear molecules such as N2, O2, H2, and so on. Dual homonuclear molecules, in and of themselves, have no permanent dipole moments. Transitions associated with molecular vibrating electrons still maintaining base states invariablly do not produce induction dipole moments either. As far as the production of dipole moments is concerned, it is necessary that they be accompanied by stimulated electron states -- under the effects of electric field incidence, electron charge distributions give rise to relative shifts. This is nothing else than to say that |3> states belong to the class of excited electron states. With regard to SRRS, |1> states and |2> states should belong to two rotation states in the same vibratory state.

By solving equation $\partial \hat{\rho}/\partial t \equiv \hat{\rho} = -i[\hat{H},\hat{\rho}]/\hbar$ satisfying density calculations $\hat{\rho} = |\psi\rangle < \psi|$

$$\dot{\rho}_{12} = -i\omega_{12}\rho_{12} + i[d_{L}\varepsilon_{L}\rho_{32} (e^{i\psi_{L}} + e^{-i\psi_{L}}) - d_{\tau}^{*}\varepsilon_{5}\rho_{13} (e^{i\psi})$$
(4)

$$\dot{\rho}_{22} - \dot{\rho}_{11} = i[\,\varepsilon_S\,(d_S\rho_{23}^{\bullet} - d_S^{\bullet}\rho_{23}^{})\,(e^{i\psi_S} + e^{-i\psi_S})$$

$$-\varepsilon_{L}\left(d_{L}\rho_{13}^{*}-d_{L}^{*}\rho_{13}\right)\left(e^{i\psi_{L}}+e^{-i\psi_{L}}\right)]\tag{5}$$

$$\dot{\rho}_{33} = i[(d_L^* \varepsilon_L \rho_{13} - d_L \varepsilon_L \rho_{13}^*)(e^{i\psi_L} + e^{-i\psi_L})]$$

$$+ \left(d_s^* \varepsilon_s \rho_{23} - d_s \varepsilon_s \rho_{23}^* \right) \left(e^{i \psi_s} + e^{-i \psi_s} \right) \right] \tag{6}$$

 $d_z = \mu_z/2\hbar$, $(\alpha = L, S)$, In this, . Use δ i to stand for normal δi phases. Moreover, note $\delta L = \delta 3 - \delta 1 = \delta 13$, $\delta S =$ $\delta 3 - \delta 2 = \delta 23$, and $\delta lambda = \delta 2 - \delta 1 = \delta 12$.

From similar equations associated with $\dot{\rho}_{13}$. $\dot{\rho}_{23}$ solutions are done for |13 and |23 (approximate solutions). Again, substituting into equations (4), (5), and (6), high frequency terms are eliminated, and one obtains [2]

$$\dot{\sigma}_{12} = i\{ (\Omega_{\lambda} + \beta \Omega_{sf}) \sigma_{12} + \Omega_{e} (\rho_{22} - \rho_{11}) [e^{i\psi_{1}} + (\beta - 1)e^{-i\psi_{2}}] \}$$
 (7)

$$\dot{\rho}_{22} - \dot{\rho}_{11} = i \left[\tilde{d}_{s} \varepsilon_{s} (e^{i\psi_{s}} + e^{-i\psi_{s}}) \left(\sigma_{23}^{\bullet} - \sigma_{23} \right) - \tilde{d}_{L} \varepsilon_{L} (e^{i\psi_{L}} + e^{-i\psi_{L}}) \left(\sigma_{13}^{\bullet} - \sigma_{13} \right) \right]$$

$$4$$
(8)

$$\dot{\rho}_{33} = 0$$

(9)

Here are introduced the signs $d_{\alpha} = \widetilde{d}_{\alpha} e^{i\delta_{\alpha}} \; (\alpha = L \;, S \;, \lambda) \;, \; \sigma_{nm} = \rho_{nm} e^{i\delta_{nm}} \;, \; \Omega_{\lambda} = \Omega_{21} = - \omega_{12} \;, \; \Omega_{\epsilon} = 2\widetilde{d}_{L}\widetilde{d}_{S}\varepsilon_{L}\varepsilon_{S}/\Delta_{L} \;, \; \Omega_{sj} = (\widetilde{d}_{L}^{2}\varepsilon_{L}^{2} - \widetilde{d}_{S}^{2}\varepsilon_{S}^{2})/\Delta_{L} \;\; \text{called drift frequencies).}$

$$\begin{array}{l})=1+\frac{1}{1+(2\omega_{s}/\Delta_{L})}\;\;,\;\;\psi_{\lambda}=(\omega_{L}-\omega_{s})t\;\;\;-(k_{L}-k_{s})z+(\Phi_{L}-\Phi_{s})=\omega_{\lambda}t-\Delta kz+\Delta\Phi\;\;.\\ \\ \Leftrightarrow\;\;R_{1}=\sigma_{12}+\sigma_{12}^{\bullet}\;\;,\;\;R_{2}=-i\;\;(\sigma_{12}-\sigma_{12}^{\bullet})\;\;,\;\;R_{3}=\\ \\ \rho_{22}-\rho_{11}\;\;;\;\;\omega_{1}=-\beta\Omega_{e}\mathrm{cos}\psi_{\lambda}\;\;,\;\;\omega_{2}=(\bar{\beta}-2)\Omega_{e}\mathrm{sin}\psi_{\lambda}\;\;,\;\;\omega_{3}=\Omega_{\lambda}+\beta\Omega_{sf}\;\;. \end{array}$$

Using (R1, R2, R3) to be the three components constituting vector R, and using (R1, R2, R3) to be the three components constituting vector $\vec{\omega}$, one then has

$$\dot{R} = \dot{\overline{\omega}} \times R \tag{10}$$

That is, during stimulated Raman scattering, particle systems satisfy Bloch equations. Normally, $\Omega e << \Omega lambda$. R is primarily rotational around 3 axses.

Doing transforms

$$\begin{bmatrix} R_1 \\ R_2 \\ R_3 \end{bmatrix} = \begin{bmatrix} \cos\psi_{\lambda} & -\sin\psi_{\lambda} & 0 \\ \sin\psi_{\lambda} & \cos\psi_{\lambda} & 0 \\ 0 & 0 & 1 \end{bmatrix} \begin{bmatrix} u_{\lambda} \\ v_{\lambda} \\ w_{\lambda} \end{bmatrix}$$

/195 Leaving out high frequency terms, one obtains

The three components of vector R_i are (u_i, v_i, w_i) , . The three components of the vector 2.5 are $(-\Omega_e, 0, \beta\Omega_{if} - \Delta^{\Phi})$. The rate of time change associated with Rlambda will be much slower than R.

This article adopts $\beta=1$. At this time, it is equivalent to adopting $E_z=\frac{1}{2}\,\varepsilon_\alpha e^{i\phi_\alpha}$, $(\alpha=L\,,S)$. This is nothing else

than saying that rotational wave approximations in resonance optics are also appropriate for use with regard to SRRS processes.

Taking stimulated particle attenuations and phenomenologically introducing them into motion equations, one has

$$\dot{u}_{\lambda} = -\left(\Delta_{\epsilon} - \Delta^{\bullet} \Phi\right) v_{\lambda} - \Gamma u_{\lambda} \tag{12}$$

$$\dot{v}_{\lambda} = (\Delta_{e} - \Delta^{\bullet} \Phi) u_{\lambda} + \Omega_{e} w_{\lambda} - \Gamma v_{\lambda}$$
(13)

$$\dot{w}_{\lambda} = -\Omega_{\epsilon} v_{\lambda} - \Gamma_{1} \left(w_{\lambda} - w_{\epsilon} \right) \tag{14}$$

In these, $1/\Gamma 1=t1$ is longitudinal relaxation times. $1/\Gamma=t2$ is transverse relaxation times. we is an equilibrium value for particle number reversal when there is no field. We = Ω sf + Δ T. Δ T is Fig.1's detuning amount [3]. 2.2 Field Equations

Electric field Ea ($\alpha = L,S$) satisfies wave equation

$$\left(\frac{\partial^2}{\partial t^2} - \frac{1}{c^2} \frac{\partial^2}{\partial t^2}\right) E_{\alpha} = \frac{4\pi N_0}{c_{\alpha}^2} \frac{\partial^2 p_{\alpha}}{\partial t^2}$$
 (15)

No is particle number density. $p = \Sigma pa$ is a particle polarization. No p is macropolarization strength. c = co/n is the speed of light in media. n is index of refraction (Here, chromatic dispersion is ignored). co is the speed of light in a vaccuum.

According to the basic principles of quantum mechanics, $p = Tr(\hat{\rho}\hat{\mu})$. Solving, one gets

$$p_L = \widetilde{\mu}_L \ (\sigma_{13} + \sigma_{13}^{\bullet})$$
$$p_S = \widetilde{\mu}_S \ (\sigma_{23} + \sigma_{23}^{\bullet})$$

In this, $\widetilde{\mu}_z = |\mu_z| = 2 \hbar \widetilde{d}_z$, , that is, $\mu_z = \widetilde{\mu}_z e^{i\delta_z}$.

Taking polarization and dividing it into two parts--isophase and nonisophase

(16)

Let

$$R_{zi} = \sigma_z + \sigma_z^*$$
, $R_{z2} = -i (\sigma_z - \sigma_z^*)$, In this

$$\sigma_{z} = \begin{cases} \sigma_{13} , & \exists \alpha = L \text{ 时} \\ \sigma_{23} , & \exists \alpha = S \text{ H} \end{cases}$$

Between $(R_{z1},\,R_{z2})$ and (u_{x},v_{x}) , the transforms below are satisfied

$$\begin{bmatrix} R_{z1} \\ R_{z2} \end{bmatrix} = \begin{bmatrix} \cos\psi_{\alpha} & -\sin\psi_{\alpha} \\ \sin\psi_{\alpha} & \cos\psi_{\alpha} \end{bmatrix} \begin{bmatrix} u_{\alpha} \\ v_{\alpha} \end{bmatrix}$$

Making use of σ 13 and σ 23 as already solved for, one obtains

(17)
$$u_{L} = \frac{1}{\Delta_{L}} \left[\widetilde{d}_{s} \varepsilon_{s} u_{\lambda} - 2 \widetilde{d}_{L} \varepsilon_{L} \left(\rho_{33} - \rho_{11} \right) \right]$$

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(18)
$$v_L = \frac{1}{\Delta_L} \tilde{d}_s \varepsilon_s v_{\lambda}$$

(19)
$$u_{s} = \frac{1}{\Delta_{L}} \left[\widetilde{d}_{L} \varepsilon_{L} u_{\lambda} - 2 \widetilde{d}_{s} \varepsilon_{s} (\rho_{33} - \rho_{22}) \right]$$

(20)
$$v_s = \frac{1}{\Delta_L} \tilde{d}_L \varepsilon_L v_\lambda$$

Particle number conservation requires $\rho 33 + \rho 22 + \rho 11 = 1$. From equation (9), one obtains $\rho 33 = C1$. Therefore, $\rho 22 + \rho 11 = C2$. C1 and C2 are constants. It is possible to see that, during

stimulated Raman scattering processes, transitions only arise between state $\rho 1>$ and state $\rho 2>$. The arrangement of state $\rho 3>$ particles is maintained invariable throughout. $\rho 3>$ is designated as intermediate state. C1 and C2 are determined by initial conditions.

It has been pointed out before that, with regard to dipole transitions, $\rho 3>$ is an electron excitation state. The energy difference between it and a base state is approximately 10eV. Therefore, C1 = 0. C2 = 1. Because of this, one solves to get

$$\rho_{33} - \rho_{22} = -\frac{1}{2} (w_1 + 1) \tag{21}$$

$$\rho_{33} - \rho_{11} = \frac{1}{2} (w_{\lambda} - 1)$$
 (22)

Taking equations (17)-(22) and substituting into equation (16), one gets

$$p_{L} = \widetilde{\mu}_{L} \left\{ \frac{1}{\Delta_{L}} \left[\widetilde{d}_{S} \varepsilon_{S} u_{\lambda} - \widetilde{d}_{L} \varepsilon_{L} \left(w_{\lambda} - 1 \right) \right] \cos \psi_{\lambda} - \frac{1}{\Delta_{L}} \widetilde{d}_{S} \varepsilon_{S} v_{\lambda} \sin \psi_{L} \right\}$$
 (23)

$$p_{s} = \widetilde{\mu}_{L} \left\{ \frac{1}{\Delta_{L}} \left[\widetilde{d}_{L} \varepsilon_{L} u_{\lambda} + \widetilde{d}_{s} \varepsilon_{s} \left(w_{L} + 1 \right) \right] \cos \psi_{L} + \frac{1}{\Delta_{L}} \widetilde{d}_{L} \varepsilon_{L} v_{\lambda} \sin \psi_{s} \right\}$$
 (24)

$$\frac{\partial^2 p_{\alpha}}{\partial t^2} = -(\omega_{\alpha} + \dot{\Phi}_{\alpha})^2 p_{\alpha} \approx -\omega_{\alpha}^2 p_{\alpha}$$
 (25)

Taking equations (23)-(25) and substituting into equation (15), comparisons of $\cos\psi_a$ and $\sin\psi_a$ coefficients arrive at equations associated with slow change amplitudes ϵ_a and slow change phases Φ_a of fields (consider $\omega_a\gg\dot{\Phi}_a$, $k_z\gg\frac{\partial\Phi_a}{\partial z}$

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \varepsilon_L = \beta_L \varepsilon_s v_\lambda \tag{26}$$

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \Phi_L = -\beta_L \left[\frac{\varepsilon_S}{\varepsilon_L} u_\lambda - \frac{\widetilde{\mu}_L}{\widetilde{\mu}_S} (w_\lambda - 1)\right]$$
 (27)

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \varepsilon_{s} = -\beta_{s} \varepsilon_{L} v_{\lambda} \tag{28}$$

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \Phi_{s} = -\beta_{s} \left[\frac{\varepsilon_{L}}{\varepsilon_{s}} + \frac{\widetilde{\mu}_{s}}{\widetilde{\mu}_{L}} \left(w_{\lambda} + 1\right)\right]$$
(29)

In this,

$$\beta_{\alpha} = \frac{\pi N_0 \omega_{\alpha} \widetilde{\mu}_L \widetilde{\mu}_S}{c_0^2 \hbar \Delta_s} , \quad (\alpha = L, S)$$

3 CLASSICAL STIMULATED RAMAN SCATTERING DESCRIPTIONS AS WELL AS THE APPROPRIATE CONDITIONS FOR USE

Let $\ \ \ \stackrel{\diamondsuit}{=} \ \widetilde{E}_{a} = \varepsilon_{a} e^{i\Phi_{a}}$ (30)

$$Q = (u_{\lambda} + iv_{\lambda}) e^{i\Delta\Phi} \qquad (\Delta\Phi = \Phi_L - \Phi_S)$$
 (31)

If one adopts $w_{\lambda} = w_{\epsilon} (\dot{w}_{\lambda} = 0)$, then the fixed solution equation set becomes

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \widetilde{E}_L = -i\beta_L \left[\widetilde{E}_S Q - \frac{\widetilde{\mu}_L}{\widetilde{\mu}_S} \left(w_{\epsilon} - 1\right) \widetilde{E}_L\right]$$
 (32)

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \widetilde{E}_{s} = -i\beta_{s} \left[\widetilde{E}_{L} Q^{*} + \frac{\widetilde{\mu}_{s}}{\widetilde{\mu}_{L}} \left(w_{\epsilon} + 1\right) \widetilde{E}_{s}\right]$$
 (33)

$$\left[\frac{\partial}{\partial t} + (\Gamma + i\Delta_{\epsilon})\right] Q^{\bullet} = -ik_{1}w_{\epsilon}\tilde{E}_{L}\tilde{E}_{S}$$
(34)

In this, $k_1 = \frac{\mu_L \mu_S}{2\hbar \Delta_L}$. Normally, one has $\Delta e << \Gamma$. If we = -1, then equations (33) and (34) change to become

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) \widetilde{E}_s = -ik_2 \widetilde{E}_L Q^*, (k_2 = \beta_s)$$
 (35)

$$\left(\frac{\partial}{\partial t} + \Gamma\right) Q^{\bullet} = ik_1 \tilde{E}_L^{\bullet} E_S \tag{36}$$

When pump light E_L does not attenuate, equations (35) and (36) form a closed equation set. It is nothing else than molecular Raman scattering (Stokes wave and photon phonon wave) coupling equations. Q is simple positive molecular vibration coordinates, also designated as photon phonon waves. In reality, from equation (31), it is possible to see that Q reflects changes associated with particle systems (u_λ, v_λ) . At the same time, it also reflects changes in field phase ($|\Phi\rangle$). So long as field phase differences $|\Phi\rangle$ are maintained constant, Q then, and

only then, belongs purely to particle "coordinates".

Key conditions for transitioning to classical descriptions are $w_1 = w_1 = -1$, , that is, particle lay out numbers associated with transition upper energy levels are zero (ρ 22 = 0). The ratio associated with first molecular vibration excitation states and base state lay outs are [4] ρ 22 / ρ 11 = 1.4x10-5. Therefore, classical descriptions are capable of giving good results with regard to molecular vibration Raman scattering. With regard to arrangement differences associated with two rotational energy levels on vibration base states, they are [5] we = -0.035. It is possible to see that classical equations will have relatively large errors with regard to describing SRRS. However, provided corrections are made to |N=No|we|, classical equations are still capable of giving good results.

4 CALCULATION RESULTS

Fixed solution equations are simultaneous equation sets composed of 7 equations—(12)—(14) and (26)—(29).

4.1 Parameter Determination

When material reaches a stable state $(\dot{u}_{\lambda} = \dot{v}_{\lambda} = \dot{w}_{\lambda} = 0)$, it is possible to solve and obtain

$$v_{\lambda} = \frac{\Omega_{\epsilon} w_{\epsilon} \Gamma}{\Gamma^{2} + (\Gamma/\Gamma_{1}) \Omega_{\epsilon}^{2} + (\Delta_{\epsilon} - \Delta \Phi)^{2}}$$
 (37)

(Illegible) the equations above and substituting into equation (28), it is possible to obtain

$$\left(\frac{\partial}{\partial z} + \frac{1}{c} \frac{\partial}{\partial t}\right) I_s = \gamma I_L I_s$$

In this $I_x = \frac{c_0 \varepsilon_x^2}{4\pi}$, $(\alpha = L, S)$;

$$\gamma = \frac{\delta \pi^2 N_0 \left(- w_e \right) \Gamma}{c \hbar \lambda_s \left[\Gamma^2 + \left(\Gamma / \Gamma_1 \right) \Omega_e^2 + \left(\Delta_e - \Delta^2 \Phi \right)^2 \right]} \approx \frac{8 \pi^2 N_0 \left(- w_e \right)}{c \Gamma \hbar \lambda_s} \left(\frac{\widetilde{\mu}_L \widetilde{\mu}_S}{\hbar \Delta_L} \right)^2$$

(Illegible) stable state gain coefficient. The setting up of the second equality in the equations above is because $\Omega e << \Gamma$ and $(\Delta - \Delta \Phi)$ << Γ .

kererence (5) gives

$$\gamma = \frac{64\pi^{3}N_{0}(-w_{e})}{15\hbar M_{S}\Delta v_{R}c^{2}} \frac{(j+1)(j+2)}{(2j+1)(2j+3)} \gamma_{00}^{2}$$

Therefore,

$$\left(\frac{\widetilde{\mu_L}\widetilde{\mu_S}}{\hbar\Delta_L}\right)^2 = \frac{4}{15} \frac{(j+1)(j+2)}{(2j+1)(j+3)} \gamma_{00}^2$$

 $\Gamma = \Gamma_1 \ , \ \Delta_T = 0 \ , \ \lambda_L = 10^{-4}$ (photon phonon wave number). Adopting $\widetilde{\mu}_L = \widetilde{\mu}_S$. cm , $\lambda_S = \lambda_L - \lambda_{ph} \ , \lambda_{ph}$ is photon phonon wave length. The values are solved for from photon phonon wave numbers $\widetilde{\nu}_{ph}$. Materials parameters are seen in Table 1 [5].

Table 1 The parameters of materials

	$N_{0}(0)$	$N_{0}(0)$ γ_{∞} B		10,		β (j)/.10-3	ns -1	
	$/10^{19} \text{cm}^{-3}$	/ 10 ⁻²⁴ cm ³	/ cm ⁻¹		6	8	10	12	14
N ₂	2.237	0.693	2.001	- 0.035	3560	3270	3060	2870	2660
Ο,		1.099	1.4378	- 0.039					

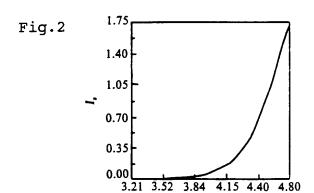
4.2 Fixed Solution Conditions

Initial conditions: when t=0, $u_{\lambda}(0)=v_{\lambda}(0)=0$, $w_{\lambda}(0)=w_{\epsilon}$. Boundary conditions: when t=0, at ground locations (z = 0), add a planar wave propogating along the z direction. These are Gaussian distributions following along with time changes. Wave lengths are |L . Half widths are τ . Peak value strengths are IL (0). Seed illumination IS (0) is adopted as IS (0) = IL (0) x 10-7.

4.3 Calculation Results

4.3.1 SRRS Transformation Characteristics and Threshold Values When strong lasers pass through the atmosphere, there is an intimate relationship between scattered light strengths IS associated with transformations and initial incident light strengths IL (0). Concrete calculation results are as shown in Fig.2. When IL (0) > 3.9MW/cm2, increases in IS following along with IL (0) are relatively slow. When IL (0) = 3.9 MW/cm2, IS increases rapidly following along with increases in IL (0). When IL (0) = 3.9 MW/cm2, IS/IL (0) \approx 1%. This pattern during strong laser atmospheric transmission possesses universality. The normal definition

 $\frac{I_s}{I_r(0)} = 1\%$



 $I_L/MW \cdot cm^{-2}$

Fig. 2 Stokes intensity I_s as a function of the incident intensity $I_L(0)$ 11

of IL (0) is SRRS threshold value It. When IL (0) > It , emited laser light IL will be scattered. Therefore, during research on strong laser light passing through the atmosphere, study of It for various types of wave lengths |L| and pulse widths τ is significant.

Table 2 gives threshold values solved for for different $(\tau, | L)$.

Table 2

Table 2 Threshold of different (τ, λ_L)

τ/ns	0.01	0.01	0.02	0.1	1.0	2.0	2.0
$\lambda_L/\mu\mathrm{m}$	0.35	1.0	1.0	1.0	1.0	1.0	0.4
$I_{t}/MW \cdot cm^{-2}$	322	928	464	98.5	14.8	9.8	3.9

4.3.2 Pulse Series Transmission Characteristics
Pulse series are composed of multiple pulses. The width of each pulse is τ . The interval between various pulses is t. Therefore, what is different from single pulses about the transmission of pulse series is due to molecules possessing "memory" effects. Speaking in concrete terms, this is nothing else than a previous pulse later being forced by stimulated molecules into exponential attenuations as a function of time using $e^{-\Gamma t}$ rules. See equations (12) and (13). In this way, with t < 1/ Γ , when the second pulse arrives, the molecular stimulation which the first pulse creates is still partially retained. The second pulse is scattered by this type of stimulated molecule. If t >> 1/ Γ , obviously, pulse series are no different from the transmission of single pulses.

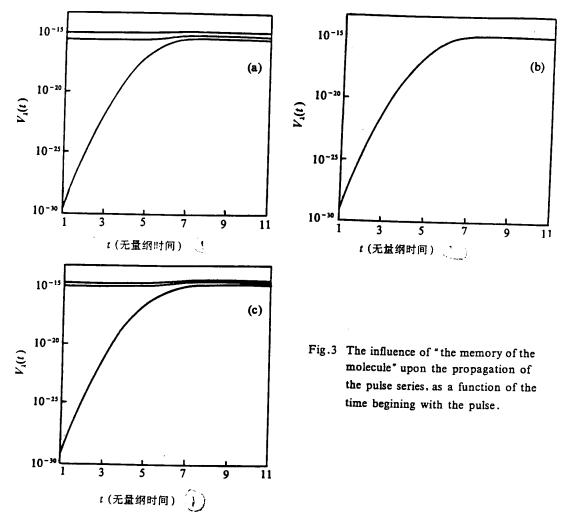


Fig.3 The influence of "the memory of the molecule" upon the propagation of the pulse series, as a function of the time beginning with the pulse. (1) Dimensionless Time

What Fig.3 graphs out is curves (v_a) associated with material stimulation created by pulse series pulses with serial numbers 1, 2, and 10. τ is 0.02ns. IL (0) = 0.085MW/cm2 . Fig.3(a) is t = 2ns. This clearly shows that molecules have "memories". Stimulations are cumulative. Fig.3(b) is t = 20ns. The three lines converge together presenting single pulse characteristics.

Due to collision attenuation coefficient $\Gamma_a \propto e^{-bz}$, the higher the atmospheric level is, therefore, the stronger is molecular "memory", and the clearer accumulative effects are. Fig.3(b) is a case where the altitude is 10km. Fig.3(c) is a case where the altitude is 50km (t=20ns). Molecules have "memories", and stimulations are cumulative.

5 ANALYSIS AND DISCUSSION

The driving term Ω e associated with w_{λ} in equation (14) is normally very small. Therefore, w_{λ} deviation equilibrium values we are very small. This is nothing else than the reason improved classical descriptons [6] give good results.

Due to the fact that the driving term Ω e associated with v in equation (12) is far larger than the driving term $(\Delta, -\Delta\Phi)v_{\lambda}$ associated with u_{λ} in equation (13), material polarization, therefore, is primarily characterized by nonisophasal component

Seen from the view point of equations (26) - (28), v_{λ} directly influences changes in field amplitudes ε_{z} . Then, u_{λ} , and ε_{z} directly influence changes in field phases Φ_{z} . Field amplitudes ε_{z} and field phases Φ_{z} do not directly give rise to coupling. By contrast, coupling is given rise to indirectly through materials systems.

When setting up computational plans, use is made of this type of coupling property to make program efficiencies relatively high. In 5 million operations a second computers, program cycles each only require 100 μs .

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MEASUREMENT OF ATMOSPHERIC COHERENCE LENGTH AND ISOPLANATIC ANGLE

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Translation of "Da Qi Xiang Gan Chang Du Yu Deng Yun Jiao De Ce Liang"; HIGH POWER LASER AND PARTICLE BEAMS, Vol.6, No.2, May 1994, pp 215-220

ABSTRACT This article introduces measurement principles associated with atmospheric coherence length and isoplanatic angle measuring instruments as well as the basic structures of instruments. It reports the results of measurements carried out at the Yunnan Observatory on Kunming's Phoenix Hill. The average value measured for ro is 12.4cm. Average value for $|{\rm o}$ is 10.6 $\mu{\rm rad}$. Compared to data associated with temperature pulsation probe instruments, the results obtained were demonstrated to be reliable. Finally, analyses were carried out with regard to a number of factors influencing measurement errors which can possibly exist in instruments.

KEY WORDS Atmospheric coherence length Isoplanatic angle Atmospheric seeing Adaptive optics

I INTRODUCTION

In situations where use is made of adaptive optics systems, it is necessary to understand atmospheric seeing at the places and times concerned. The selection of sites for observatories and atmospheric seeing associated with daily astronomical observations are also one of the indispensible myriad parameters. Quantitative descriptions of atmospheric seeing parameters are nothing else than the so called Fried length [6] --- atmospheric coherence length ro. This characterizes atmospheric coherence properties along certain transverse distances in a certain specially designated light path. In the same way, people are concerned with coherence characteristics between different light courses within certain ranges of distances or certain angles. This then requires making use of isoplanatic zones or isoplanatic angles in order to make descriptions. In actuality, what are made relatively more use of are isoplanatic angles, normally recorded as Θo . In the middle 1980's, we began to make efforts in research work relating to atmospheric coherence [2]. conjunction with this, an instrumentation unit to measure ro and Θ o has been recently developed and manufactured successfully. This article is nothing else than the carrying out of a simple introduction to its measurement principles, instrument structure, and preliminary measurement results. Finally, analysis is done of error factors which can possibly exist.

2 MEASUREMENT PRINCIPLES

2.1 Atmospheric Coherence Length The definition of atmospheric coherence length is

$$D(r) = 6.88 (r/r_0)^{5/3}$$
 (rad²)

That is, when receiving optical system radius r is equal to coherence length ro, the value of wave structure funcion D (r) is 6.88 rad2. At this time, the properties of the systems in question reach optimum limits. Due to random propogation theory associated with waves in media, in air-ground light courses, planar waves ro, are capable of being expressed as

$$r_0 = [0.422 k^2 \sec \psi \int_{h'}^{\infty} C_n^2(h) dh]^{-3/5}$$
 (2)

In the equation, $k=2\pi/\lambda$. λ is wave length. ψ is the zenith angle. C_{π}^2 (h) is height distributions associated with turbulent flow structure constants. h' is the altitude of measuring stations.

If it is possible to measure C_n^2 (h), it is then possible to calculate ro from equation (2). Barletti and others [3] used this method to obtain height distributions associated with ro.

The results were the same as Γ distribution models [4] we use/216

$$C_n^2(h) = C_0 h^{-2/3} \exp(-h/H_0)$$
 (3)

Calculation results agree. In the equation, Co and Ho are two set up parameters. Ho is analogous to the equivalent atmospheric altitude at the place and time in question. Co corresponds to turbulence strengths associated with measurement station positions. However, normally, making use of this type of method in order to precisely determine ro is very difficult. Moreover, it is also not possible to obtain results in real time. As a result, it is necessary to opt for the use of a type of indirect measurement method. In it, use is made of the earliest and most wide spread methods—which make use of arrival angles in order to do conversions. The theoretical expression associated with arrival angle fluctuation variences is

$$\sigma_a^2 = 2.905 D^{-1/3} \sec \psi \int_{h'}^{\infty} C_n^2(h) dh$$
 (4)

In the equation, D is the diameter of the receiving telescope. From equation (2) and equation (4), it is possible to obtain

$$r_0 = 3.18k^{-6/5}D^{-1/5}\sigma_{\alpha}^{-6/5} \tag{5}$$

In early periods, measurements associated with arrival angles made use of photographic time exposure methods in order to record star fluctuations. From one dimensional directional time exposures, it is possible to obtain angular varience C_{up}^2 . In order to make σ_{up}^2 , and σ_{up}^2 equal to each other, an empirical constant exists between the two, that is,

$$\sigma_s^2 = b \, \sigma_{Hp}^2 \tag{6}$$

As far as adopted b values are concerned, there exists very great dispersion, making equation (6) become very imprecise. Besides this, due to such causes as data processing troubles, vibrations in instruments themselves, and so on, they can easily bring with them very great errors. Single light path arrival angle methods have gradually become known as "dual point difference arrival angle methods" or "difference imagery movement methods" [6] instead. This type of method takes light coming from a certain celestial body and, through certain measures, forms two celestial images on the receiving plane. Respective solutions are made for the differences $\Delta \rho$ ci of the two celestial images from their central position. Through the formula

$$r_0 = \left\{ \frac{2f^2 \left[0.36 \left(\lambda/D \right)^{1/2} - 0.242 \left(\lambda/d \right)^{1/3} \right] \lambda^{5/3}}{\langle \Delta \rho_{ci}^2 \rangle - \langle \Delta \rho_{ci} \rangle^2} \right\}^{3/5}$$
 (7)

ro is calculated. In the equation, f is telescope focal length. d is the central interval associated with the two light paths (that is also the theoretical interval associated with two images when there is no fluctuation). This type of method is not sensitive to such factors as perturbations in the instruments themselves, optical system quality, temperature effects on telescope focal length, fluctuations in star image brightness, and so on. As a result, it is possible to hope to obtain very good measurement precisions.

2.2 Isoplanatic Angles

The theoretical expression associated with isoplanatic angels is

 $\theta_0 = [2.905 \, k^2 \sec \psi \int_{h'}^{\infty} C_n^2 (h) h^{5/3} dh]^{-3/5}$ (8)

In the same way as ro, Θ o daily observations must also go through observations of another physical quantity in order to carry out conversions. The most convenient solution is to use star light scintillation methods. This is due to the fact that the scintillation method σ_i^2 also possesses altitude weighting functions similar to Θ o. Theoretical calculations clearly show that, when D = 11cm, it is then possible to make the weighting functions associated with the two equal to each other [7]. In conjunction with this, the relationship set out below is established

$$\theta_0 = C \left[\log \left(1 + \sigma_s^2 / < S > ^2 \right) \right]^{-3/5}$$
 (9)

<S> is average values associated with star light signals. C is a
constant. Through large amounts of experimentation, Eaton and
others [8] indicate that C = 0.9676. Equation (9), during weak
turbulence and medium strength turbulence, the precisions
possessed are very good. It is only under strong turbulence
conditions that errors can reach approximately 20%. In general
cases, turbulence strengths throughout the entire atmosphere will
not be in a strongly turbulent state. As a result, equation (9)
is normally effective. /217

3 INSTRUMENT CONSTRUCTION

In our coherence length and isoplanatic angle measurement instruments, as far as measurements associated with ro are concerned, use is made of dual point difference arrival angle methods. With regard to measurements associated with Θ o, use is made of star light scintillation methods. The schematic diagram is as shown in Fig.1. Principal parameters are seen in Table 1. The instruments in question break down into two large parts—optical receiving systems and data processing systems. The former includes two recieving telescopes connected together with completely uniform properties, image merging light paths and two detectors (4720 model CCD pickup cameras and photoelectric

multiplier tubes). The entire system is installed on an equatorial type base. It is possible to use computers to control step electrical equipment to make searches and track in declination and right ascension directions. The latter includes video frequency cards, A/D cards, and 386 microcomputer systems.

Fig.1

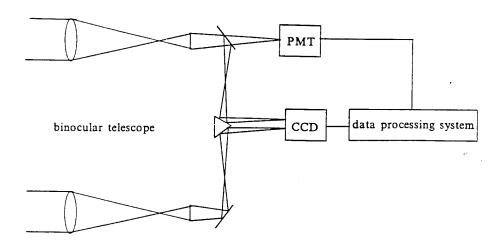


Fig. 1 Schematic diagram of the r_0 and θ_0 measurer

Table 1

Table 1 The main parameters of the r_0 and θ_0 measurer

Optical system	Receiving aperture	D/cm	12	
	Effective focal length	f/m	5	
	Separaton of centers	d/cm	15	
	Visual angle of telescope	(')	3	
	Visual angle of guiding telescope	(°)	0.5	
Image signal	Minimum illumination	$E_{min}/1x$	0.02	
	Signal - to - noise tatio	SNR/dB	52	
	Sampling rate	R/Hz	25	
	Sampling time	t/s	2.5	
Intensity signal	Dark current	I_0/nA	50	
	Sampling rate	R/Hz	1000	
	Sampling time	t/s	5	
Mechanical	Туре	Equatorial style		
system	Right ascension	± 100 °		
	Declination	±45°		
	Tracking precision	≼4 ″		
	Weight 约200kg			

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The height of the entire instrument is approximately 1.2m. The length is approximately 1.6m. Right ascension and declination coordinates are picked up from 21 place photoelectric coding disks. At the same time as numeric tubes display right ascension and declination coordinate values, they are transferred to microcomputer to be recorded in order to act as a basis during normalized data processing.

CCD pick up cameras possess characteristics of low illumination and high signal to noise ratios. In order to be advantageous to the reception of celestial images with relatively weak optical powers, the signals go through video frequency collection cards, and are, first of all, entered into internal computer storage. The collection rate is 25 frames/s. Each iteration of observations is 2.5 s. After that, solutions are done for the two central positions $\rho 1i$ and $\rho 2i$, thereby obtaining $\Delta \rho$ ci = $\rho 2i$ - $\rho 1i$. Finally, ro is calculated from equation (7).

When measuring Θ o , computer processing inputs celestial image strength signals by A/D cards. Collection speed is lkHz. The sampling period for each set of data is 5s. Reckoning up

signal fluctuation varience σ_s^2 and average value <S> , by using fomula (9), it is then possible to calculate Θo .

In order to resolve such problems as inadequate computer internal storage capacity and excessively long calculation times, when processing celestial image signals, option is made for the use of "window" methods. The size of windows must pay attention to all of imagery fluctuation ranges, mechanical adjustment precisions, and observation times. Considering imagery fluctuations, they are generally not larger than 2". Mechanical precisions are approximately 4". As a result, window height should not be less than 10". This corresponds to the 64 lines of a 512x512 imagery card. Window widths should also be at least 10". The total of the two imagery paths is 20". That is, window size is 64 lines x 128 columns. During observations, first of all, adjusting telescope right ascension and declination coordinates makes celestial imagery enter into the window placed on the top of the screen. After it is appropriately placed, the base then stops rotating. After the center of gravity of images distinguished by computer are placed at suitable locations, number gathering begins. Windows follow celestial images as they shift downward, synchronously tracking. After completing the gathering of predetermined amounts of data, calculations are then From the beginning of collection to the presenting carried out. of results only requires 4 seconds. Utilization and operation of measurement instruments is very simple. Automation levels are very high.

4 MEASUREMENT RESULTS

From November to December, 1992, we carried out measurements to be completed in 40 days at the Yunnan Observatory on Phoenix Hill in Kunming. This observatory is located 25°N at 2000m above sea level. Due to the fact that the weather is sometimes rainy or cloudy, only 20 segments of useful data were acquired. time period for each iteration of observation was between 18:40 -Through analysis of measurement results, it was discovered that the average value of ro is 12.4cm (Unitized to the zenith location. The same below.). The range of changes was The average value associated with Θ o is 10.6 μ rad. range of changes is 3-27 μ rad. Fig.2 and Fig.3 are, respectively, observation results for ro and Θ o on the two days, 4 and 5 December. From these Fig.'s, it is possible to see that changes associated with ro are relatively smooth. However, Θ o changes are still relatively large. This clearly shows that changes associated with low atmosphere turbulence are not large. However, atmospheric turbulence flows associated with relatively high layers (h > 3 - 5 km) are very unstable. Looking at this overall, the atmospheric seeing for Kunming in this period was very good. It can be compared to excellent observatory sites on a world scale, such as the U.S. Mauna Kea and Chile's La Silla.

During periods when use was made of coherence length and isoplanatic angle measurement instruments to carry out

observations of ro and Θ o, Zeng Zongyong and others fired temperature fluctuation space probe instruments at the same time to carry out measurements of C^2 . Fig.4 is an actual example measured at 23:00 on 29 November 1993. In the Fig., curve b is the Hufnagel model [9]. Curve c is given by equation (3). In this, Co = 3.5×10^{-14} . Ho = 5400 m. Their standard deviations relative to experimentally measured data are, respectively, 4.6 and 0.026. Coefficients of correlation are, respectively, 0.70 and 0.73. From this, it is possible to know that Γ models agree relatively well.

We made use of C_s^2 probe data and ro and Θ o observation data for the same periods to carry out comparisions. In accordance with equation (2) and equation (8), respectively, calculations were gotten of ro = 14.4cm and Θ o = 8.3 μ rad. These two are in basic agreement. However, there are some relatively small differences. The causes for the differences that exist will be discussed in the sections that follow.

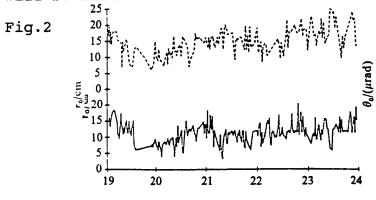
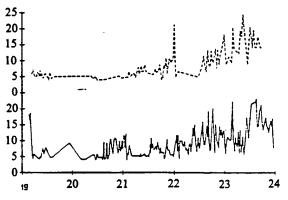


Fig.3

Beijing time/h
Fig. 2 The atmospheric coherence length vs time
——: 1993. 12.4 ---: 1993. 12.5



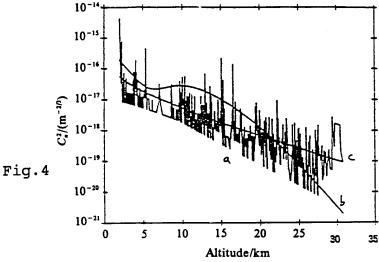


Fig. 4 An example of measured C_n^2 (23:00, Nov.29, 1993) a. measured data; b. Hufnagel's model; c. Eq. (3)

Beijing time/h
Fig. 3 The isoplanatic angle vs time
----: 1993. 12. 4 ---: 1993. 12. 5

5 CONCLUSIONS

The fact that there exist differences associated with measurement values and theoretical calculation values is also seen in the work of other scholars. For instance, Sydney and others [10] discovered ro observed values to be 4.4cm. However, calculated values were 3.35cm. The two differed by 24%. The differences associated with Θo were even larger. This is entirely consistent with our results. In fact, it is reasonable that certain differences should exist. The reason is the measurement instruments themselves. No matter whether it is measurements of ro and Θo or measurements of such parameters as C_n^2 , there always exists a certain measurement error. As random quantity statistics, differences of around 20% are quite small. At the same time, theoretically, it is also possible for

certain flaws to exist.

Speaking in terms of measurements, our imagery gathering speeds are not high enough. Sampling times are not long enough. This then creates frequency spectra in signals not being capable of complete coverage. Seen on the basis of frequency spectrum fluctuations, high frequency components (f210Hz) are already extremely weak. Very large errors are not necessarily created. However, paying inadequate attention to low frequency components (f≤0.2Hz) will give rise to relatively large errors. As a /220 result, sampling times should be increased later. Secondarily, there is only a 15cm distance between the centers of the two telescopes associated with measuring instruments. It may be somewhat smaller. On the basis of Roddier's opinions, when d/D ≥ 2, the accuracy of formula (7) is somewhat higher. However, this is not important. The shortening of interval distances will, no doubt, influence low frequency components. However, on the basis of the Taylor principles of frozen turbulence flows, if option is made for the use of multistage averaging, it will reduce in the same way the effects of low frequency components. One possible cause for the existence of relatively large differences in isoplanatic angles is that, according to theory, D should be However, what we actually used was 12cm. Due to relatively large apertures possessing relatively large aperture smoothing effects, this leads to signal fluctuations becoming relatively weak. Thus, relatively large isoplanatic angles are obtained. Another possible cause which is more important is that, when solving formulae (2) and (8), consideration was not given to the influences of turbulent flow outside scales. As far as problems in this area are concerned, we have already made detailed analyses [11] in another report. It is not necessary to repeat them here. We only wish to point out this. considering finite outside scales, based on a certain outside scale height distribution model, it is possible--with regard to the turbulent flow data discussed above--to do calculations and obtain ro = 20.9cm and Θ o 12 μ rad. These numerical values and

observed values are then in quite good agreement.

In summary, we are able to reach the following conclusions. Results obtained from coherence length and isoplanatic angle measurement instruments are reliable. After making certain improvements, it will then be possible to obtain even higher accuracies.

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A NEW APPROACH TO LASER CONDITIONING TECHNOLOGIES AND MECHANISMS ASSOCIATED WITH RAISING HIGH POWER LASER REFLECTOR MIRROR COATING DAMAGE THRESHOLD VALUES

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ABSTRACT This article introduces a type of laser conditioning technology associated with raising high power laser reflector mirror coating damage threshold values. In conjunction with this, it stresses using experimental and theoretical research on the mechanisms of raising laser damage threshold values to draw conclusions and make inquiries.

KEY WORDS Optical coating Laser damage Laser conditioning Mechanism

INTRODUCTION

Following along with the wide spread application of high power lasers in fields such as tactical laser weapons as well as laser thermonuclear fusion, and so on, reserach associated with optically coated reflector mirrors possessing high laser damage threshold values has received more and more serious attention from people. The reason is that laser damage threshold values associated with optical coatings limit laser output powers. the last more than ten years, a new type of technology has been developed--laser conditioning technology. It is capable of making laser damage threshold values associated with certain highly reflective mirrors go up two to three fold. Inside China and abroad, a number of hypotheses have been put forward in order to explain this phenomenon. What is regretable is that there is still not one type of theory capable of completely explaining it.

CHARACTERISTICS OF LASER CONDITIONING

Laser Conditioning Technology

As far as laser conditioning is concerned, it refers to a new technology which is capable of raising 2-3 fold laser damage threshold values after taking optical coatings or large pieces of optical materials and putting them through laser irradiation lower than laser damage threshold values. There are two types of laser irradiation methods associated with laser conditioning. One is using multiple iterations of laser irradiation of the same strength (s-on-1). Another type is using multiple iterations of laser irradiation with optical strengths which gradually go up over time (r-on-1). Using Hf02/Si02 and Zr02/Si02 reflective film systems as examples, laser damage threshold values before laser conditioning are 7.4J/cm2 and 7.1J/cm2. Going through laser conditioning, they rise respectively to 19.8J/cm2 and Moreover, the latter type of laser conditioning 18.8J/cm2.method raises laser damage threshold values somewhat higher.

Relationships to Laser Parameters

Laser system parameters are different. Not only are damage threshold values different, but damage mechanisms also differ. Relationships between damage threshold values and wave lengths have roughly the relationship $D_{\pm} \propto \lambda^{x}$ coefficient which changes along with film properties. laser wave lengths are, the higher damage threshold values then The relationship between damage threshold values and pulse are. $D_{\pm} \propto \tau^{0.35}$. When pulse widths are long, damage is widths is primarily caused by absorption of focused heat. When laser pulses are short, dialectric breakdown created by electric fields is the primary factor in damage. Besides this, there are also relationships to such things as repetition frequencies of lasers, modes, and so on. Due to the fact that the relationships of laser damage threshold values and widths to wave lengths are the same before and after laser conditioning, as a result, damage mechanisms should also be the same before and after laser This article limits itself to neodymium glass conditioning.

laser spectral lines associated with irradiation laser light of 1.06 microns. Pulse widths are a few nanoseconds.

2.3 Relationships to Methods of Coating

Three types of methods are used to carry out tests: (1) commonly used electron beam evaporation; (2) ion beam spattering (IBS); and, (3) plasma plating (PP). Making use /298 of the latter two types of methods, the plated coating apertures are few (densities are high). Mechanical properties and environmental characteristics are good. However, experiments clearly show that their laser damage threshold values are still lower than films associated with electron beam evaporation. As far as carrying out laser conditioning on them is concerned, PP film laser damage threshold values are not raised much. Moreover, IBS laser conditioning is not necessarily effective. Generally speaking, laser conditioning is only effective with regard to electron beam evaporation coatings.

2.4 Relationships with Thin Film Materials

Thin film materials associated with high reflection mirrors in the visible and near infrared often use pairs of HfO2/SiO2, TiO2/SiO2, and ZrO2/SiO2 to form reflection film systems. Because, after TiO2/SiO2 plating, there is a requirement for drying in oxygen in order to reach complete oxydation so as to reduce absorption, this is not appropriate for reflection mirrors with large apertures. Besides that, in plating processes, TiO2/SiO2 reflection mirrors cannot avoid producing intense absorption of low valance oxides of titanium. As a result, we only select ZrO2/SiO2 and HfO2/SiO2. Making use of laser conditioning associated with pulse widths of lns, laser damage threshold values associated with the latter two are all raised roughly 3 fold. Replicability of laser damage threshold values associated with HfO2/SiO2 is very good. Moreover, levels of increase with regard to various types of pulses are the same. Laser damage threshold values associated with ZrO2/SiO2 follow along with different positions on reflector surfaces and change. Laser conditioning results are sometimes good and sometimes bad. However, making use of Nomarski microscopic techniques for observations, the flaw densities for the two are the same. Relationships with Coating System Designs

With regard to three types of samples—lambda/4 HfO2, rich SiO2 samples (includes 13 layers of lambda/4 SiO2 between 10nm HfO2), and rich HfO2 samples (includes 17 layers of lambda/4 HfO2 between 10nm SiO2)—after laser conditioning, laser damage threshold values all go up approximately 2 fold. Because of this, there is no relationship between conditioning results and relative thicknesses of film systems. However, there is a type of exceptional special case. That is nothing else than that when the SiO2 thickness is O, that is, there is single layer HfO2, there are no conditioning effects.

3 INQUIRIES INTO LASER CONDITIONING MECHANISMS

3.1 Early Explanations of Laser Conditioning The first type of hypothesis is "laser cleanness", that is, laser conditioning takes volatile impurities (in particular, water in the atmosphere) and gets rid of them from surfaces and the interior of coating materials. These impurities lead to laser damage. If this is the case, damage threshold value increases should be temporary. We took HfO2 reflection mirrors which had not gone through laser conditioning and put them separately into environments at 20°C with relative humidities of approximately 50% for a period of time. Laser damage threshold values of 1064nm are measured. After that, use is then made of anhydrous P2O5 and measurements are made again in 10 weeks. Observations are done of the shortening of wave lengths after the removal of water. It was discovered that, comparing laser damage threshold values before and after the removal of water, there were no alterations at all. However, The explanation of this is reflectivity drift is reversible. that laser damage threshold value changes are permanent.

Another type of hypothesis is that lasers take materials and heat them right up until their crystalinities In order to check this hypothesis, use was made of TEM and X ray diffraction in order to determine crystal structures associated with single and multiple layer films of HfO2 and SiO2. Results clearly showed that SiO2 in single and multiple layer films of 1 micron and 266 nm was no different at all. cases, it was in amorphous states. In contrast to that, HfO2 crystal structures were related to thickness--smaller than 20nm, states were indefinite. As growth continued, by contrast, they were cylindrical crystals. They were primarily monoclinic crystals. Small numbers of cubes were also included. changes in relative thickness certainly did not influence conditioning effects. As a result, there was no relationship between laser conditioning and crystal structures. Making use of diffraction meters to carry out scanning with regard to thin films before and after laser conditioning, no differences were discovered.

3.2 Electron Defect Model

Research personnel of the Lawrence Livermore National Laboratories (LLNL) put forward the existence [1] of a relationship between laser conditioning and electron defects inherent in films. Because electron beam evaporation processes are high speed quenching from high temperature steam to low temperature substrate, as a result, a porous film is produced with defects in its interior. There is a relationship between nonequilibrium states and laser conditioning characteristics. They believe that laser damage associated with thin films goes through the four steps below:

(1) lasers take valence carrying electrons from restricted zones and excite them into conducting zones;

(2) through photoelectric field or free current carrying particle absorption effects, free current carrying particles are

stimulated to achieve high energies;

- (3) through ionization avalanche or photoelectric effects (crystal lattice heating), excessive amounts of energy are transmited into crystal lattices; /299
- (4) film materials are heated to critical damage temperatures—for instance, media material melting points.

The occurence of laser conditioning phenomena is due to the elimination of the sources of conducting zone electrons in the first step, that is, as far as media material are concerned, under low flux laser irradiation, electrons in defects are excited into conduction zones. However, at this time, photoelectric fields are still not adequate to give rise to damage (steps 2-4). Electrons then decay into relatively deep energy levels. After that, it is not easy for them to be stimulated into conduction zones. When laser conditioning media make use of laser light below laser damage threshold values for irradiation, the number of electrons it is possible to use in order to transfer to conduction zones is already very small. As a result, the amounts of net energy transmited to crystal lattices is also small and does not go so far as to cause damage. In this way, conditioning effects are then produced.

Above is the basic thinking associated with the model in question. However, LLNL personnel also discovered a few experiments [2] that do not agree with the results that this

model predicts.

Using ND.YAG lasers with wave lengths of 1064nm, 532nm, 355nm, and pulse widths of 10ns, respectively, to carry out laser conditioning with regard to 1 micron thick SiO2 and HfO2 single layer and multiple layer films, it was discovered that, in three wave lengths, SiO2 single layers increased roughly 2 fold in all cases. However, HfO2 single layer film laser damage threshold values still showed no increase at all. At the same time, LLNL personnel determined 1.06 micron, lambda/4 single layer HfO2 and SiO2 films. In them, HfO2 films and 1 micron thick ones are the same. There are still no conditioning effects. However, the levels of increase associated with SiO2 laser conditioning laser damage threshold values are still not higher than 30%.

If, with respect to laser damage threshold value $D_T^{\infty\lambda^x}$, , x = approximately 1.0 for SiO2 single films and x = approximately 1.9 for HfO2, laser damage threshold values are intensely influenced by wave length. In another area, damage forms are different. SiO2 damage is a few small points (smaller than 5 microns), and they do not get any bigger with continued laser irradiation. Yet typical HfO2 damage has diameters or 0.1-1 mm. This clearly shows that the methods of laser coupling to HfO2 and coupling to SiO2 are different.

With regard to multiple layer Hf02/Si0 films, after 1064nm laser conditioning, laser damage threshold values increase 2 fold. 355nm only have 40%. 533nm is then between the two. Laser damage threshold values at 1064nm are controlled by SiO2. However, at very short wave lengths, by contrast, the control is by HfO2. Due to HfO2 absorption fringes associated with electron

beam evaporation at around 375nm, narrow band linear absorption inside HfO2 may possibly be the cause controling 355nm multiple film layer laser damage the shold values.

As a result, it is certainly not a single electron defect inside HfO2 or SiO2 which is then capable of determining conditioning effects. In that case, are electron defect models right in the final analysis or not? Up to now it has still not been possible to reach a conclusion. Perhaps a part of them is correct.

3.3 Recent Progress

Edwards and others carried out analyses [3] with regard to 1m ZrO2/SiO2 Nova laser reflection mirrors plated by LLNL (The surface of the mirrors in question is unusually smooth. They discovered that the production of Roughnesses are 2.5-5nm). damage is begun from microscopic defects with diameters of approximately 30 microns. These defects form damage of approximately 250 microns, sometimes covering over 20% of the mirror surface. These defects lie inside thin films. conjunction with this, they are of a two dimensional "disc" type. What is peculiar is that the damaged film surface is enveloped by a layer of "fog". They extracted these "fog" granules and discovered that they were precisely ZrO2 and SiO2 film materials. The explanation for this lies in material being splattered out in the damage process. In conjunction with this, it piles up on the film surface. Making use of SEM to observe these granules, diameters did not reach 1 micron. There was both surface smoothness and surface roughness. The former may possibly be formed from gas states condensing. However, the latter may possibly be solid state remnants thrown out.

Edward's explanation is that it is due to the defects in ZrO2 material being heated, after laser irradiation, to temperatures sufficient to make the surrounding ZrO2 gasify, that In reference [4], Chase confirms that is, in excess of 3000K. the cause which leads to this process is linear absorption. this type of so called defect a pure substance granule or not? Schildbach made use of laser mass analysis systems (LIMA), and, in conjunction with this, discovered no impurities at all [5]. In another area, due to defect densities being on the order of 100mm-2 (with regard to the Nova reflection mirror), it is only when a vaccuum chamber is severely polluted that it is possible to create this kind of outcome. In that case, may they not be low valence oxide materials -- for instance, SiO2? Due to the fact that the Nova mirror laser damage threshold value is 7.8J/cm2, in this way, absorption coefficients associated with low valence oxides should be greater than 1000cm-1. Unfortunately, SiO2 absorption coefficients (1 micron) still do not reach 100cm-1. What defects in SiO2 are in the final analysis remains a puzzle even now.

With regard to HfO2 films, LLNL personnel, when studying electron paramagnetic defects, discovered that, contained in rich HfO2 samples, there were special signals which were not in other samples. This type of signal clearly shows that HfO2 may contain

in it Fe3+ impurities (Commercially used HfO2 film materials contain less than 0.05% Fe). This cannot be eliminated under laser irradiation below damage threshold values, leading to there being no conditioning effects.

Schildbach discovered that, when laser irradiation lower than the laser damage threshold (16-18J/cm2) is used on HfO2/SiO2 reflective films, there are neutral particles of Hf, Si, and oxygen (including O and O2) sent out [5]. During initial emission, laser optical strength is 10J/cm2. Moreover, the critical optical strength required for conditioning is also just 10J/cm2. As a result, the emission of neutral particles is very possibly produced by laser conditioning. Neutral particle kinetic energies reach as high as a few score eV. High energy neutral particle emission corresponds to heat induced fractomission. Even if there is no obvious damage, this may still be the source of neutral particles. After heating of thin film defects, heat induced cracking is produced, leading to the elimination of absorbed material or the improvement of thermal stress fields (These are all causes giving rise to damage).

At the same time, Schildbach analyzed 8 points at different locations on mirror surfaces. He discovered that the conditioning results associated with them were certainly not the same. With regard to the 2 points which had not gone through processing, laser damage threshold values were 18-20J/cm2-approaching laser damage threshold values associated with other points after processing. He believed that the cause for this was due to these two points not having any defects.

Staggs and others used atomic force microscopes (AFM) to observe reflection mirrors after plating. They discovered that most SiO2 surfaces contain small mounds 200nm wide and 10nm high Moreover, Hf02/Si02 reflective film build ups are primarily irregular and are shallow pits or cores. Longitudinal dimensions Transverse dimensions are approximately 1/3 of are 1-10 microns. longitudinal dimensions. Densities are on the order of 10mm-2. After damage, the largest changes are the emission of cores. A round tunnel remains. Staggs paid attention to their being cracking on the edges of the tunnels. The cores are very sensitive to damage. When cores grow on thin films, due to shadow effects, moreover, the peripheries of cores are made to produce gaps, and they become flexible. Another characteristic of damage is the enlargement around 8 fold of small mounds in areas without defects in SiO2 surfaces, that is, their growth or agglomeration. This type of surface agglomeration temperature is high enough to reach an ability to make components transition within time periods on the order of nanoseconds, causing the production of structures with large volumes and small surface Structures clearly show that emission of cores occurs areas. when using laser irradiation lower than laser optical strengths associated with making small mounds agglomerate.

Staggs carried out irradiation on defect areas. During the process of optical strengths increasing from 7.8J/cm2 to 54.6J/cm2, AFM was used to analyze the appearance of defects.

There were still no obvious changes to the original outlines. However, after increasing to 19.8J/cm2, most 10nmx100nm cores disappeared from surfaces. After increases continued to 54.6J/cm2, there were no further changes. That is nothing else than to say that, as far as the production of surface smoothness phenomena in optical strengths emited by cores is concerned, even though optical strengths continue to increase—exceeding laser damage threshold values—at the locations associated with these cores, still no damage occured.

4. CONCLUSIONS

To summarize what was discussed above, relfection mirror thin film damage is capable of definitely not resembling that type of complete cause in microscopic nonstable states considered by electron defect models, and, at the same time, is also not capable of being explained in all phenomena from a macro viewpoint. Combining electron defect models, we believe that the damage may go through the several steps below:

(1) Defects Inside Thin Films--Under Laser Irradiation--Stimulate the Production of Large Amounts of Free Electrons Method A: Electron Ionization Avalanche

If free electrons which have broken out of their bonding already exist inside thin films, after laser irradiation, they go through electron ionization avalanche and form plasma bodies. The avalanche speed is represented as

$$\frac{dN_c}{d\tau} = \gamma(E) \times N_c$$

In the equation, $\gamma(E)$ is ionization avalanche coefficient, Nc is conductance belt electron density, and E is field strength.

After that, energy absorption proceeded. Assuming that $\eta(\xi,\tau)$ is the number of electrons with energies between $\xi+\delta\xi$. It is expressed as

$$\eta(\xi, \tau) = \eta(\xi) \exp(\xi \times \tau)$$

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Moreover, electron flux through isoenergetic planes is

$$J(\xi,\tau) = v(\xi) \times \eta(\xi,\tau) - D(\xi) \times \delta \eta(\xi,\tau) / \delta \xi$$

In this, $\nu(\xi)$, and $D(\xi)$ are, respectively, effective speeds and coefficients of diffusion associated with electron flows. Then, the energy distribution equation is

$$\frac{dJ}{d\xi} = \gamma(E) \times \eta(\xi)$$

Attached boundary conditions are

$$\eta(E_g, \tau) = 0$$

$$J(0, \tau) = 2 \times J(E_g, \tau)$$

In equations, Eg is the electrical gap of materials.

Method B: Going Through Valence Electron Transitions Given
Rise to by Multiple Photons

At this time, the free electron densities produced by

stimulated valence electron transitions are

$$\frac{dN_c}{d\tau} = N \times \sigma^m \times F^m$$

In the equation, N is particle density, αm is multiple photon absorption, Fm is incident photon flow density.

(2) Free Current Carrying Particle Absorpton Inside Defects
If crystal lattice temperature is T. On the basis of Drude
models

$$c \rho \frac{dT}{d\tau} = \frac{N_c e^2 \tau E^2}{m(1 + \tau^2 \omega^2)}$$

In the equation, c is specific heat, ρ is crystal density, τ is photon and electron collision time (approximately on the order of 10-16 s), E is electric field strengths (effective value), m is electron effective mass, and ω is laser angular frequency.

Combining the energy distribution equations above, the carrying out of integration over the entire energy space give laser energy absorption as

$$c \rho \frac{dT}{d\tau} = \delta \left\{ \int_{0}^{+\infty} \xi \cdot \eta(\xi, \tau) d\xi \right\} / \delta \tau = 1.09 \left(m^{\circ} kT / 2\pi \right)^{1/2} \cdot \frac{N_c}{L_{ac} v_s} \cdot \left(\frac{eE\omega}{m^{\circ}} \right)^3$$

In the equation, Lac is the average free travel associated with electron and photon collisions, vs is vertical wave speed, and m* is electron belt mass.

(3) After Heating, High Density Defects Produce Agglomeration. In Conjunction with This Cracks which Are Made to Cover Thin Films Lead to Damage.

Laser conditioning is then the production of cracking in thin films under low flux laser irradiation, and, in conjunction with that, the causing of neutral particle emission. The results lead to laser damage cores being ejected. Conditioning results depend only on defect properties. Moreover, there is no relationship to such factors as thin film density, layer number, and so on.

The significance of laser conditioning research lies not only in being able to make our advances and improvements in laser conditioning technology, which is not yet mature. It is also being able to make us advance our understanding of the basic nature of laser damage.

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