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on  $\Delta\rho$  according to a simple power law over the full density range of Fig. 3, although the slope of  $\rho^2\kappa_T$  appears to approach 2 asymptotically near  $\rho_c$ . Along the vapor side of the coexistence curve,  $\rho^2\kappa_T$  obeys a power law (exponent  $1.8 \pm 0.3$ ) over a wider range of densities than does  $\kappa_T$  alone. More extensive experimental data are needed to investigate the existence of a natural thermodynamic variable for the compressibility.

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## Transient Self-Focusing in a Nematic Liquid Crystal in the Isotropic Phase

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We have made the first quantitative experimental study of transient self-focusing in a Kerr liquid by propagating Q-switched laser pulses in an isotropic nematic substance which has a long orientational relaxation time. Our results show good qualitative, and occasionally quantitative, agreement with the existing theoretical calculations.

Self-focusing of light in a Kerr liquid has been a subject of extensive investigation.<sup>1</sup> It gives rise to the observed intense streaks and many other related phenomena. Quasi-steady-state self-focusing is now well understood. Most of the observed results can be explained satisfactorily by the picture of moving foci.<sup>2,3</sup> Transient self-focusing is however still in a state of confusion. There already exist a number of detailed numerical calculations on the subject,<sup>4-6</sup> but no quantitative experimental results are yet available to check these calculations. The reason is obvious. Transient self-focusing occurs when the laser pulse width  $W$  is shorter than or comparable to the relaxation time  $\tau$  of the field-induced refractive index  $\Delta n$ . Since in ordinary Kerr liquids  $\tau$  is in the picosecond range, picosecond pulses must be used in the transient studies.<sup>7</sup> Unfortunately, picosecond pulse technology is still in such a primitive stage that quantitative measurements are extremely difficult and expensive. Experiments on transient self-focusing would be so much easier if nanosecond pulses could be used, but then the medium should have a relaxation time  $\tau$  in the 50-nsec range. This happens to be the case for liquid crystalline material in the isotropic phase.<sup>8</sup> In this paper, we would like to report the preliminary results of our ex-

perimental investigation on transient self-focusing in such a medium. Our results are in good agreement with the existing theoretical calculations.<sup>5,6</sup>

Let us first have a brief review on the theory of transient self-focusing. The phenomenon is presumably governed by the equations

$$[\nabla^2 - (\partial^2/c^2 \partial t^2)(n_0 + \Delta n)^2]E = 0, \quad (1)$$

$$(\tau \partial/\partial t + 1)\Delta n = \Delta n_0, \quad (2)$$

where  $n_0$  is the linear refractive index and  $\Delta n_0 \cong n_2|E|^2$ ,  $n_2$  being a constant. The solution of Eqs. (1) and (2) has been obtained numerically.<sup>4-6</sup> It can be described qualitatively from the following physical reasoning. If the input pulse is short, then  $\Delta n$  can never reach its steady-state value. We can write

$$\Delta n(z, \zeta) = (1/\tau) \int_{-\infty}^{\zeta} n_2 |E(z, \eta)|^2 \exp[-(\zeta - \eta)/\tau] d\eta,$$

where  $\zeta = t - zn_0/c$  is the local time. For an input pulse width comparable to  $\tau$ , Eq. (3) shows that the later part of the pulse may see a larger  $\Delta n$ . Consequently, different parts of the pulse propagate differently in the medium as shown in Fig. 1.<sup>6</sup> The front part ( $a, b$ ) of the pulse sees little induced  $\Delta n$  and diffracts accordingly. It however leaves a sufficiently large  $\Delta n$  in the

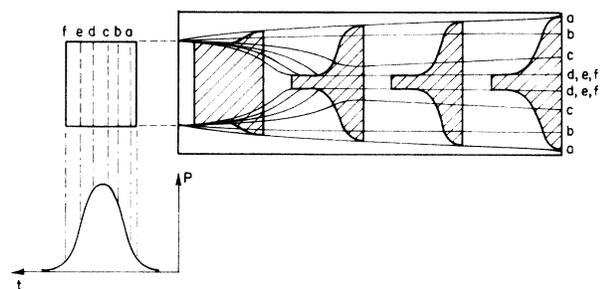


FIG. 1. Schematic drawing showing how an input pulse gets deformed through self-focusing into a horn-shaped pulse which then propagates on with little change in shape.

first section of the medium to cause the lagging part ( $c-f$ ) first to self-focus and then to diffract. Because of the transient response, both self-focusing and diffraction are expected to be gradual.

Figure 1 shows that the spatial distribution of the input pulse first deforms into a horn shape and then propagates on without much further change.<sup>6,9</sup> If one probed the pulse at a point sufficiently deep in the medium, one would see a horn-shaped pulse pass through; the pulse describing the variation of the on-axis intensity with time appears distorted as compared to the input pulse. The minimum diameter of the self-focused beam should be a sensitive function of the input power since it depends strongly on the magnitude of the induced  $\Delta n$ . However, it turns out that other nonlinear processes may set in to limit the diameter.<sup>10-12</sup> When this happens, we may find that the minimum diameter of the self-focused beam remains nearly unchanged for an appreciable section of the pulse or, in other words, the horn-shaped pulse propagates over a finite distance with little change in its neck diameter. The numerical results of transient self-focusing, expressed in terms of the normalized parameters such as  $T/\tau$ , etc., are given in Refs. 5 and 6.

In our experimental study, we used the liquid crystalline material N-[*p*-methoxybenzylidene]-*p*-butylaniline (MBBA) in its isotropic phase (above  $T_K = 42.5^\circ\text{C}$ ) as the nonlinear medium. This material has a large steady-state optical Kerr constant (about 70 times that of  $\text{CS}_2$  at  $50^\circ\text{C}$ ) and an orientational relaxation time which varies with temperature from  $\sim 40$  to  $> 800$  nsec.<sup>8</sup> It is therefore ideal for study of transient self-focusing with  $Q$ -switched laser pulses. Our experi-

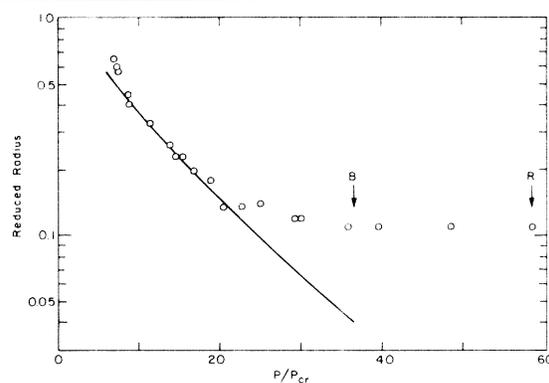


FIG. 2. Radius of the self-focused beam at the end of a 9.5-cm MBBA cell at  $50^\circ\text{C}$  as a function of the incoming peak power. The reduced radius is normalized against the radius of the incoming beam, and  $P_{cr} = 0.12$  kW. The solid curve is a theoretical curve extrapolated from the calculations in Ref. 6. The arrows *B* and *R* indicate where stimulated Brillouin and Raman scattering set in.

mental setup was the same as the one described earlier.<sup>2,3</sup> The input single-mode ruby laser pulse had a diameter of  $210 \mu\text{m}$ , a pulse width  $W$  of 10 nsec, and a maximum peak power of 50 kW. The cell containing MBBA was sealed under  $\text{N}_2$  atmosphere, and was thermally controlled to  $\pm 0.03^\circ\text{C}$  in the range between 40 and  $53^\circ\text{C}$ . Self-focusing was readily observed in such a medium.

We first photographed the self-focused beam at the end of a 9.5-cm cell and used the densitometer traces to measure the reduced beam diameter at different input power levels. The results for MBBA at  $50^\circ\text{C}$  are shown in Fig. 2 on a semilog plot. When the input peak power increases, the beam diameter first decreases almost exponentially and then approaches a limiting diameter. As seen in Fig. 2, the nearly exponential decrease agrees very well with the theoretical curve extrapolated from the curves calculated by Shimizu.<sup>6</sup> At  $P/P_{cr} = 20$  ( $P_{cr} = 0.12$  kW), some other nonlinear process presumably sets in to limit the beam diameter. At this power level, the peak intensity at the focus is several orders of magnitude below the avalanche breakdown threshold<sup>12</sup> and therefore breakdown cannot be the limiting mechanism. We also found that stimulated Brillouin scattering and stimulated Raman scattering do not appear until the input power reaches appreciably higher values, as indicated in Fig. 2, and therefore they also cannot be responsible for the limiting diameter.<sup>11</sup>

We suspect that two-photon absorption is probably the limiting mechanism in this case because of the near-uv absorption band of MBBA. Anyway, this suggests that different nonlinear processes must be responsible for the limiting diameter of the self-focused beam in different cases.

The results at different temperatures are nearly the same after the correction due to scattering loss. The reason is as follows. In our case,  $W \ll \tau$ , and hence, from Eq. (2),  $\Delta n(t) \cong \int_{-\infty}^t \Delta n_0(t') \times dt'/\tau$  in the first-order approximation. Both  $\Delta n_0$  and  $\tau$  due to molecular reorientation are proportional to  $(T - T_c)^{-1}$ , where  $T_c$  is the temperature at which a second-order isotropic-nematic transition would occur.<sup>8</sup> As a result,  $\Delta n(t)$  is nearly independent of temperature except for the change of scattering loss with temperature. At 50°C or higher, the effect of scattering loss on self-focusing in our case is negligible.

We also measured the on-axis intensity variation of the self-focused beam with time at the end of the cell. In Fig. 3, we show the results. As the input power increases, the peak of the output intensity pulse first appears (case *b*) appreciably delayed from the peak of the input pulse (case *a*) and then gradually moves (cases *c*–*e*) towards it. At sufficiently high input power, weak oscillation starts to develop on the lagging

part of the pulse (case *d*) and gets gradually more pronounced (case *e*). At even higher input power, stimulated Brillouin scattering begins to show up. The above results agree qualitatively very well with the predictions derived physically from Fig. 1, and also with the numerical calculations of Refs. 5 and 6 (which also show oscillation in the output pulses at high input powers). Quantitative comparison between theory and experiment is presently not possible since no numerical calculation corresponding to the experimental parameters we used is available.

We also calculated from Fig. 3 the temporal variation of the self-focused beam diameter at the end of the cell by assuming that the shape of the intensity profile remains nearly unchanged. The results are shown in Fig. 4, cases *a*–*d*, which correspond to the intensity pulses of Fig. 3, cases *b*–*e*, respectively. It is seen that the incoming pulse does gradually develop into a horn shape as described qualitatively in Fig. 1. In particular, the neck diameter of the horn remains almost constant over a fairly long section, although at high input power, it shows some weak oscillation which correlates with the oscillation on the output intensity pulse.

We have also made measurements at different cell lengths. The results are similar to what we

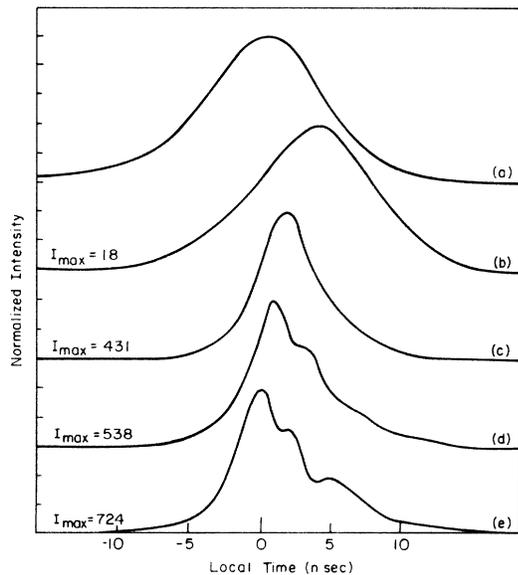


FIG. 3. On-axis intensity variation with time at the end of a 9.5-cm MBBA cell at 50°C with different input peak powers: case *a*, the input pulse; *b*,  $P = 1.6$  kW; *c*,  $P = 3$  kW; *d*,  $P = 3.2$  kW; *e*,  $P = 3.5$  kW.  $I_{\max}$  is in  $\text{MW}/\text{cm}^2$ . Note that the baseline is shifted for each pulse.

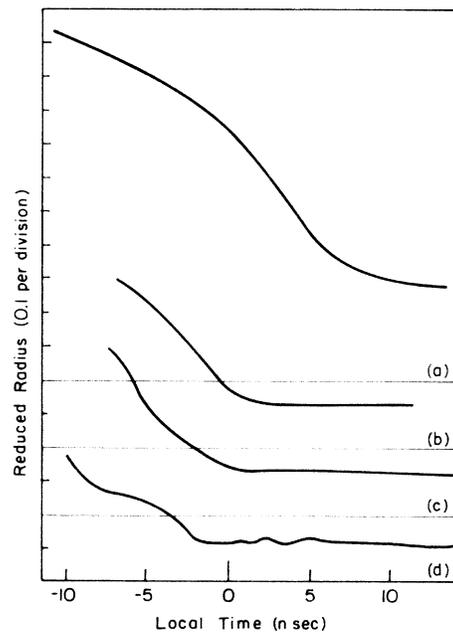


FIG. 4. Temporal variation of the self-focused beam radius at the end of the cell. Cases *a*–*d* correspond respectively to the cases *b*–*e* in Fig. 3. Note the shift of the baseline for different cases.

have already described, indicating that after the initial period of deformation through self-focusing, the deformed horn-shaped pulse does propagate on with little further change in shape. The self-focused beam showed practically no spectral broadening since the phase modulation rate here is expected to be small.

Finally, we should briefly comment on the recent work of Rao and Jayaraman on self-focusing in isotropic MBBA. In analyzing their data, they have neglected the facts that self-focusing is transient and that the scattering loss is non-negligible in their 30-cm cell, especially at temperatures close to the transition temperature. It is also difficult to understand how they could obtain a self-focusing threshold  $P_{th}$  (defined as the input power at which the self-focused beam at the end of the cell reaches a limiting diameter) which decreases as  $(T - T_c)^{0.16}$  with temperature. Our results show that  $P_{th}$  increases as  $(T - T_c)$  decreases. This is what one should expect because the scattering loss increases as<sup>14</sup>  $\exp[A/(T - T_c)]$ .

In conclusion, we have presented the results of the first quantitative experimental study on transient self-focusing. They agree qualitatively, and also quantitatively when comparison can be made, with the earlier theoretical predictions.<sup>5,6</sup> The limiting diameter in the present case is due to neither stimulated Raman and Brillouin scattering nor avalanche breakdown.

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## Restraining of Runaway Electrons in High-Temperature Plasmas\*

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A theoretical model is proposed in relation with the observed lack of electron runaway processes in magnetically confined, high-electron-temperature plasmas on which electric fields of the order of or larger than the critical runaway field have been applied. The excitation of standing modes locked in with the periodic inhomogeneities of the magnetic field is considered in order to provide, via mode-particle resonance processes, a momentum exchange between the (accelerated) current-carrying electrons and the magnetic field itself.

One of the motivations for this Letter is the experimental observation of strongly anomalous electrical resistivity<sup>1,2</sup> in high-electron-temperature plasmas on which electric fields of the order of or larger than the runaway critical field

have been applied. The relevant experiments<sup>1</sup> involved magnetically confined toroidal plasmas for which the Langmuir frequency was of the order of or smaller than the electron gyrofrequency.<sup>1,2</sup> In particular, since no large anomalous