Secondary modulation instability in partially coherent beams

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Abstract

Due to modulation instability partially incoherent optical beams break up into stripe filaments in noninstantaneous media at a first threshold. We numerically and experimentally report on the formation of two-dimensional filaments at a secondary threshold, if increasing the nonlinearity further, which is due to material anisotropy. Particularly, we investigate the dependence of this secondary modulation instability on the coherence properties of the beam, using a photorefractive nonlinearity. From the measurement of modulation contrast in two dimensions and additionally using a Fourier method we quantitatively derive experimental thresholds for first and secondary onset of modulation instability and study how both onsets relate to the coherence properties of the beam.

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1. Introduction

Modulation instability (MI) is a very general phenomenon in nonlinear science [1]. In presence of a nonlinearity a noisy uniform solution becomes unstable against perturbations of a specific length or time scale. These specific perturbations become self-amplified and finally a self-organized solution with a scaling typical for the system evolves. Examples for such behaviour may be found in many fields and also play an important role in optics. In the context of optical systems at first regarded as a nuisance, later on the potential importance of these processes for the development of concepts in optical information processing have been recognized and consequently intensive

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research activity was triggered. Especially the for-
mation of temporally and spatially localized struc-
tures such as solitons has attracted considerable
attention for this reason [2,3]. A two-dimensional
transversal modulation instability of the optical
beam is to be considered as a crucial pre-condition
for the existence of such spatial optical solitons,
because a fundamental requirement for a spatial
optical soliton, namely the balance between dif-
fraction-induced beam broadening and a self-
focussing nonlinearity, is found at the same
parameter regions as MI. If MI is present, a broad
optical beam will break up into small transversal
filaments while propagating within a self-focussing
nonlinear medium [4–8]. Assuming a spatially
isotropic nonlinear response of the material, the
filamentation transversal to the propagation direc-
tion, would also be expected to occur isotropically.
However, in case of anisotropic media, such as
photorefractive crystals, an anisotropic nonlinear
response must be expected. And indeed, in photo-
refractive materials it is found that dependent on
both the strength of the nonlinearity and the prop-
agation distance the uniform beam first breaks up
into (1 + 1) dimensional (D) stripes and only later
on at either higher propagation distances or stron-
ger nonlinearities (2 + 1)D beam filaments develop
[5].

In principle one expects MI to occur only with
coherent beams. If the nonlinear material however
exhibits a noninstantaneous response, the observa-
tion of MI with incoherent beams is possible as
long as the phase fluctuations of the beam are far-
ter than the response time of the material [8–14].
MI of such partially incoherent beams has gained
attraction, because a change in interaction charac-
teristic of incoherent solitons has been observed
and therefore the prospect for more closely packed
arrays of non-interacting solitons is opened
[15,16]. Up to now MI of incoherent beams has
been mainly studied with an approach regarding
(1 + 1)D solutions, which may be extended to
(2 + 1)D structures if assuming a spatially isotro-
pic material response [11]. As a first approach this
approximation is sufficient to explain the major ef-
fects. The delayed onset of (2 + 1)D filamentation
via (1 + 1)D stripe solution as it has been observed
in the coherent case [5] cannot be covered if the
material anisotropy is excluded. Although the be-
haviour of the in experiment most commonly
used photorefractive nonlinearity is determined
by anisotropy, up to now the threshold behaviour
of MI in noninstantaneous media and partially
incoherent beams has not yet been studied includ-
ing anisotropy-induced phenomena. In our con-
tribution we especially address the aspect of
anisotropy induced phenomena of MI when using
incoherent light and a photorefractive nonlinear-
ity. Particularly, we are interested in the onset of
the secondary (2 + 1)D MI in dependence on the
beam’s coherence properties, which cannot be
covered in an isotropic approach. At first we will
discuss MI of incoherent beams, using a numerical
approach. Later on we present experimental
results, which we analyse with different methods
in respect to the onset of the secondary fully
two-dimensional modulation instability.

2. Numerical model for incoherent modulation
instability in anisotropic media

In order to numerically study modulation insta-
bility of incoherent beams we model the propaga-
tion of incoherent light in a photorefractive
medium. The model explicitly includes the full
characteristics of the photorefractive nonlinearity
and particularly the anisotropic material response
is covered.

A light beam propagating through a photore-
fractive medium in z-direction can be described
in paraxial approximation by the equation

$$2ik_0n_e \frac{\partial}{\partial z} A + \nabla_\perp^2 A = -k_0^2n_e^2r_{33}\frac{\partial \varphi}{\partial x} A,$$  (1)

where \(\nabla_\perp = \hat{\partial}_x + \hat{\partial}_y\), \(k_0\) is the vacuum wavenumber
of the light beam, \(n_e\) is the refractive index of the
(unperturbed) medium and \(r_{33}\) is the effective
element of the electro-optical tensor. \(\varphi\) is the elec-
tric space charge potential inside the medium. For
our simulations we used, in agreement with the
experimental configuration, \(k_0 = 2\pi/532\) nm, \(n_e =
2.3\) and \(r_{33} = 180\) pm/V. \(A\) is the envelope of the
electric field of the light beam. In order to cover
for the incoherence of the light beam, the coherent
density model is used, which states that incoherent

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density model is used, which states that incoherent
light can be described as a superposition of many coherent, yet mutually incoherent components, propagating in slightly different directions. We thus express the light beam at the input face of the medium as:

\[
A(x, y, 0) = \sum_{k_x, k_y} P(x, y) \exp(i(k_xx + k_yy)) \frac{l_c}{\sqrt{\pi}} \times \sqrt{\exp(-(k_x^2 + k_y^2)l_c^2/2)} \exp(i\gamma_{k_x, k_y}(t)),
\]

where \([P(x, y)]^2\) is the intensity profile of the beam at the input face, \(k_x\) and \(k_y\) stand for the slight tilting of the different components, \(l_c\) is the coherence length of the light and \(\gamma_{k_x, k_y}(t)\) are random phase factors that are uncorrelated to each other, thus making the single components mutually incoherent. The factor \(\exp(-(k_x^2 + k_y^2)l_c^2/2)\) stands for a Gaussian distribution of the mutually incoherent \(k\)-vector components. We chose the maximum intensity of our incoherent beam to be around unity (measured in units of the dark intensity \(I_d\)). To excite the instability, we add around 1% of noise to the initial input beam.

Since plane waves with truly infinite spatial extent cannot be realized in numerical simulations, we chose to use a large grid and periodic boundary conditions to minimize the effects of the edges on the instability.

Photorefractive crystals are known to be well-described by the Kukhtarev equations, which yield, using a few well justified approximations, the following differential equation for the space charge potential [17]:

\[
\nabla^2 \varphi + \nabla \cdot \ln(1 + I) \varphi = E_0 \frac{\partial}{\partial X} \ln(1 + I),
\]

where \(E_0\) is the external field applied parallel to the crystal’s \(\hat{c}\)-axis (in the following we will choose \(E_0 = 2.5 \text{kV/cm}\)) and \(I\) is the intensity of the light beam, scaled to the dark intensity \(I_d\), describing the excitation of charge carriers into the conduction band by thermal excitation or background illumination of the crystal. Remember that we are dealing with incoherent light. Hence, the intensity \(I\) is given by the sum of the intensities of the single components introduced in Eq. (2) and propagated using Eq. (1).

\(L_c = \infty\) 16\(\mu\)m 13\(\mu\)m 11\(\mu\)m

Fig. 1 shows the results of the numerical propagation of light with different degrees of coherence. Each column shows light with a different value of \(l_c\). The leftmost column shows the fully coherent case, i.e., \(l_c = \infty\), while for the other columns \(l_c = 16, 13\) and 11 \(\mu\)m from left to right. The light beam is shown at the input face, after \(z = 6, 8, 10\) and 12 mm of propagation. The pictures are oriented such that the \(\hat{c}\)-axis of the photorefractive crystal is horizontal. The pictures show that for all degrees of coherence, MI first breaks the plane wave into stripes oriented perpendicular to the \(\hat{c}\)-axis. In the coherent case, these stripes then break into filaments, as is known for quite a while [5]. If the degree of coherence of the light is decreased, however, the breaking of the stripes can be suppressed. For \(l_c = 16\) \(\mu\)m one can see that extent of the filaments in the vertical direction is much longer than in the case of the fully coherent light. Decreasing of the degree of coherence further then leads to a complete stabilization of the stripes. Reducing the degree of coherence even further
would lead to a complete suppression of MI. The plane wave then would propagate through the medium without changing its shape.

3. Experimental observation of incoherent modulation instability in photorefractive crystals

The light source for our experiment is a Nd:YAG cw-laser at \( \lambda = 532 \text{ nm} \). To create a partially incoherent beam, a rotating diffuser is placed between two lenses in a 4f arrangement. The degree of spatial coherence is controlled with the diffuser position on the z-axis. According to the beam diameter at the diffuser the speckle size and thus the degree of spatial coherence length is modified. The collimated beam then propagates through a Cerium doped Strontium Barium Niobate crystal SBN:61 (5 mm \( \times \) 5 mm \( \times \) 23 mm, in width, height and length) to which an external electric field is applied along the crystal’s \( \hat{c} \)-axis. The beam is extraordinarily polarized and thus experiences a nonlinear change of refractive index depending on the external field and on the total light intensity. The dark current density of the crystal is controlled with a white light background illumination. The intensity distribution at the crystal’s backplane is recorded with a CCD-camera, while the input intensity is monitored with a photodiode. As an equivalent to the spatial coherence length \( l_c \) the average full half width of the speckles at the crystal’s frontplane is taken, when the diffuser was stopped. It is made sure that the diffuser rotates much faster than the material’s response time \( \tau \), which is in the order of a second for the crystal in use [18].

The intensity output was monitored at a fixed coherence length \( l_c \), while the nonlinearity of the SBN crystal was increased by raising the externally applied voltage \( U_{\text{ext}} \) in increments of 100 V. For analysis, the images of the beam profile were recorded at every voltage step after approximately 2 min, permitting for transients to disappear. The applied external electric field was restricted to maximum values of 3 kV/cm to avoid damage of the crystal. Experiments were performed in the same manner for different coherence lengths varying between \( l_c = 280 \) and 9 \( \mu \text{m} \). During all measurements input intensity \( (I = 480 \mu \text{W/cm}^2) \) as well as background illumination \( (I_{\text{white light}} \approx 700 \mu \text{W/cm}^2) \) were kept constant. Only approximate levels for the intensity of the background illumination at the crystal position can only be given due to the divergence and the spectral width of the white light. In Fig. 2, images of developing MI are shown. Rows depict system states at equal nonlinearity, while columns show pictures at equal coherence properties. At a fixed coherence length \( l_c \) (e.g., the \( l_c = 280 \mu \text{m} \) column) starting from a nearly uniform state the system develops first a stripe pattern, perpendicularly to the \( \hat{c} \)-axis, followed by the formation of spot like filaments at higher voltages. Small modulations found in the uniform state (0 V) most likely result from striations of the crystal and probably act as a trigger for the formation of stripes [14]. From the images it becomes also

![Fig. 2. Experimental images of modulation instability. Images in a row: the nonlinearity is equal. Images in a column: the degree of coherence is constant. With decreased coherence modulation instability sets in later.](image-url)
obvious that as well formation of (1 + 1)D stripes as the decay of stripes in (2 + 1)D filaments occurs later, if the coherence length is decreased. Other groups observed tilting of the stripes up to an angle of 45° [11, 12], in our experiment however only a very small tilt of approximately 1–5° compared to the initial striations in the crystal has been observed. The resulting (1 + 1)D MI stripes are in good approximation oriented perpendicularly to the crystal's $c$-axis. At very small coherence lengths (e.g., 9 $\mu$m) (2 + 1)D MI is strongly suppressed and we observe zig-zag-kind of patterns instead of pronounced stripes. With decreasing coherence also the contrast of the pattern decreases.

4. Experimental thresholds for secondary modulation instability

To gain a quantitative measure for the experimental thresholds of MI the mean contrast $C(x_i)$ of image profiles has been determined along the images’ horizontal $x$- and the vertical $y$-axis, respectively, with the following equation:

$$C(x_i) = \frac{(I_{\text{max}} - I_{\text{min}})}{(I_{\text{max}} + I_{\text{min}})},$$

where $x_i$ denotes either the $x$- or $y$-direction. We determined the mean contrast by first calculating the average contrast of a single image profile line. The total mean contrast $C(x)$ plotted over the external bias in Fig. 3 was derived from 600 lines. The curves in the graph are guides for the eye, which connect the measurement points at fixed coherence lengths $l_c$. The legend in the insertion of Fig. 3 denotes the coherence lengths at which measurements were performed. Fig. 3(a) shows the averaged contrast $C(x)$ (parallel to the crystal's $c$-axis). Typically (e.g., at $l_c = 284$ $\mu$m) at first a rise in contrast is observed, which is generated by the formation of the (1 + 1)D stripe pattern. The rise is used to determine an experimental threshold for (1 + 1)D MI, as which we define the rise above the level of $(1 - 1/e) = 63\%$ times the maximum of contrast (plotted as $+$ in Fig. 7). As in our experiment the observed tilt angle between $x$-axis and stripe pattern is very small, this method to us seems justified. However for comparison later on also thresholds are derived from the angular modes in Fourier space are derived [13, 14].

If only the first (1 + 1)D MI was involved one would expect the contrast $C(x)$ to remain at a constant saturation level, but typically the contrast decreases from its maximum value onwards (e.g., at 1 kV/cm at $l_c = 284$ $\mu$m). Note, that the decline in contrast is less pronounced for measurements at smaller coherence lengths $l_c$ and also sets in later. In the case of coherence lengths of $l_c = 19$ and 9 $\mu$m even no significant decrease of the contrast is observed below 3 kV/cm. We conclude that the drop in contrast is caused by the onset of the secondary (2 + 1)D MI. Due to the break up of the stripe solution, now also the $y$-profiles become modulated, which reaffects $C(x)$. To check this assumption also the mean contrasts of the $y$-profiles
C(y) are plotted in Fig. 3(b). Along the y-profiles the rise of contrast occurs at higher nonlinearities (e.g., at l_c = 280 μm the maximum of C(y) is reached at 1.4 kV/cm, instead of at 0.8 kV/cm for C(x).) We define an experimental threshold for (2 + 1)D MI as before as the rise above the level of (1 − 1/e) times the maximum of C(y) (* in Fig. 7). However, at the coherence lengths l_c = 9 and 19 μm C(y) does not yet reach a well pronounced maximum below 3 kV/cm, therefore the (2 + 1)D thresholds for these coherence length are to be neglected. Comparison with images of MI at l_c = 9 μm (Fig. 2) indicates that at these low coherence length potentially the formation of (1 + 1) D MI is suppressed in favour of a kind of zig-zag pattern, shown in Fig. 2, which seems to smears out the contrast measurement.

Similar results can be obtained from the numerical simulations. The plots of the average contrasts C(x) and C(y) can be found in Fig. 4. Qualitatively good agreement in the progression of experimental- and numerical contrast functions is observed. Quantitatively however comparison to the experimental results is difficult because the propagation length was used as parameter in numerics, while in the experiments the nonlinearity of the material is varied. Note, that typically the threshold for the onset of nonlinear behaviour is derived theoretically using a more rigid stability analysis. Here, we therefore refrain from the determination of numerical thresholds and restrict ourselves to a qualitative comparison.

To validate our previous results we also derived experimental thresholds from a Fourier analysis of the light field, as also used by Chen et al. [13,14]. From the Fourier transform we calculate the angular distribution of spatial frequencies:

\[ f(\theta_0) = \int_{\theta_0 - \Delta \theta}^{\theta_0 + \Delta \theta} \int_0^{k_{\text{max}}} dk \, d\theta \mathcal{F}(k, \theta), \]

where \( \mathcal{F}(k, \theta) = \mathcal{F}(\mathcal{F}(I(x, y))) \) is the Fourier transform of the intensity distribution, and with \( \Delta \theta = 1^\circ \) and \( \theta_0 \) varying from 0° to 360° in increments of 2°. In presence of the stripe (1 + 1)D MI perpendicularly to the stripe orientation pronounced peaks appear in the angular distribution \( f(\theta) \), while as well without MI present as at fully developed (2 + 1)D MI, \( f(\theta) \) shows no peaking (see Fig. 5(a)). Thus, now we determined the averaged contrast \( C(f(\theta)) = (\max\{f(\theta)\} - \min\{f(\theta)\})/ (\max\{f(\theta)\} + \min\{f(\theta)\}) \) of the angular distribution profiles, which is plotted in Fig. 5(b). From the contrasts we define experimental thresholds as follows: As the threshold of (1 + 1)D MI we take the rise above 1/e*max(f(θ)) (□ in Fig. 7) and for the (2 + 1)D MI we define the descend below (1 − 1/e)*max(f(θ)) as threshold (○ in Fig. 7). As before similar results are obtained for the contrast function \( C(f(\theta)) \) derived with the Fourier method from numerical simulation as comparison with the plot in Fig. 6 shows.

In Fig. 7 experimental thresholds for (1 + 1)D (+, □) and (2 + 1)D MI (*, ○), defined above are plotted. Comparison of the thresholds derived from C(x) and C(y) with the ones derived from C(f(θ)) shows that for (1 + 1)D MI the C(f(θ)) thresholds (□) remain below the C(x) thresholds (+), while the C(f(θ)) (2 + 1)D MI thresholds (○) stay above the C(y) thresholds (*). This differences are due to
the choice of values for the definition of thresholds. Fine tuning of these values probably would gain better coincidence, but the difference can also be used as an estimation for the error bars of the measurement. The difference for the (2 + 1)D MI thresholds however becomes especially pronounced for very small coherence lengths. Here, the difference can be regarded as an indicator for the formation of the zig-zag like pattern. The onset of full filamentation is in this parameter region covered better by the Fourier method $C(f(\theta))$. Regarding the general progression of (1 + 1)D experimental thresholds in relation to (2 + 1)D MI thresholds at first these values do not change with decreased coherence length. In this parameter region also both threshold approximatively have a fixed ratio to each other, which can be viewed as indication for the anisotropy of the SBN crystal’s nonlinear response. Only at a coherence length of approximately $l_c = 45 \mu m$ both experimental thresholds start to alter and a steep increase of both is observed. Considering only the behaviour of the experimental thresholds of MI, the coherence length $l_c = 45 \mu m$ can be interpreted as the nonlinear coherence length of MI. Up to this point the nonlinear response to the partially coherent beam is equivalent to the coherent behaviour, from then on the influences of incoherent fluctuations push the experimental thresholds for MI upwards and also the fixed relation between thresholds ceases to exist.
5. Conclusion

We have presented experimental and numerical results on two-dimensional modulation instability of incoherent beams in anisotropic media. The onset of $(1 + 1)D$ and $(2 + 1)D$ MI was investigated experimentally and numerically in dependence of the spatial coherence length. At moderate incoherence the experimental thresholds of MI are not affected by alteration of coherence properties. If the beam becomes more strongly incoherent the experimental thresholds of MI increase steeply. At very low coherence lengths numerics and experiment indicates to the development of zig-zag kind of patterns in favour of $(1 + 1)D$ filamentation, which was analysed by help of two different methods in determining experimental threshold of MI. A good qualitative agreement of numerical and experimental results can be stated.

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