Unconditional generation of bright coherent non-Gaussian light from exciton-polariton condensates

Tim Byrnes, ¹ Yoshihisa Yamamoto, ^{1,2} and Peter van Loock ³

¹National Institute of Informatics, 2-1-2 Hitotsubashi, Chiyoda-ku, Tokyo 101-8430, Japan

²E. L. Ginzton Laboratory, Stanford University, Stanford, California 94305, USA

³Institute of Physics, Johannes Gutenberg-Universität Mainz, 55099 Mainz, Germany

(Received 2 November 2012; published 13 May 2013)

Exciton-polariton condensates are considered as a deterministic source of bright, coherent non-Gaussian light. Exciton-polariton condensates emit coherent light via photoluminescence through the microcavity mirrors due to the spontaneous formation of coherence. Unlike conventional lasers which emit coherent Gaussian light, polaritons possess a natural nonlinearity due to the interaction of the excitonic component. This produces light with a negative component to the Wigner function at steady-state operation when the phase is stabilized via a suitable method such as injection locking. In contrast to many other proposals for sources of non-Gaussian light, in our case, the light typically has an average photon number exceeding one and emerges as a continuous wave. Such a source may have uses in continuous-variable quantum information and communication.

DOI: 10.1103/PhysRevB.87.201301 PACS number(s): 71.36.+c, 42.50.Ex, 42.50.Gy

The optical coherent state is one of the most fundamental states of quantum optics, and has been studied from theoretical and application perspectives for half a century.^{1,2} Coherent states, and related optical states such as squeezed coherent states, are characterized by a Wigner quasiprobability distribution of a Gaussian form, containing no negative regions.³ While such Gaussian states are fairly easily produced and manipulated in the laboratory,⁴ the creation of general non-Gaussian states is a more difficult task, due to the lack of materials possessing strong nonlinearities. Current experimental approaches to creating states with a negative Wigner function are highly probabilistic.^{5,6} Even theoretically deterministic single photon production schemes in practice suffer from less than perfect efficiencies, and would still not be capable of producing bright states with negative Wigner functions. A more desirable way of generating non-Gaussian light is a more "plug-and-play" style device, where the device is simply switched on and bright non-Gaussian light emerges deterministically.

One of the most promising applications of such non-Gaussian light is for continuous-variable (CV) quantum information tasks. CV quantum optics has been shown to be a complementary approach to traditional discrete-variable methods using qubits. §-12 While elementary quantum protocols such as teleportation¹³ can be realized using Gaussian CV states and Gaussian operations, a more advanced application such as universal quantum computation is known to require non-Gaussian elements. 14-16 However, even those states that are non-Gaussian but positive may be insufficient as resources for quantum computation^{17,18} (for the discrete analog, see Ref. 19). As a consequence, the occurrence of a negative Wigner function can be seen as a prerequisite for a potential quantum mechanical speedup²⁰ (for related results on the discrete case, see Ref. 21). While such negativities are necessary, only small instances of it are sufficient for universality. ^{20,22,23}

In this Rapid Communication we describe a deterministic method of producing non-Gaussian quantum states of light with a negative Wigner function using exciton-polariton condensates. Exciton-polaritons are bosonic quasiparticles in semiconductor microcavities, corresponding to a coherent superposition of an exciton and a cavity photon.²⁴ Exciton-polaritons have been observed to undergo a dynamical form of Bose-Einstein condensation.^{25,26} One of the attractive features of this system is that the state of the polaritons inside the semiconductor can be directly imaged using the leakage of photons through the imperfect microcavity mirrors. Thus the spontaneous buildup of coherence, one of the signals of Bose-Einstein condensation, can be directly measured simply by analyzing the properties of the emerging light.

One of the principal differences between an excitonpolariton condensate and a laser is the presence of, or lack of, respectively, strong coupling between the excitons and photons. Strong coupling leads to polariton-polariton interactions, giving much richer physics than its noninteracting counterpart.²⁷ These interaction effects also lead to phenomena such as superfluidity of the polaritons.²⁸ In this Rapid Communication, we take advantage of the polariton-polariton interactions to propose a device for producing non-Gaussian light. The key technological advance that has allowed for this is the availability of high-Q cavities where polariton lifetimes of the order of ~ 100 ps are now possible.²⁹ In addition, exciton-polaritons may be confined spatially using trapping techniques such as metal deposition on the surface of the microcavity, 30 increasing the self-interaction energy U. This gives the possibility of energy scales of the self-interaction U and the cavity decay $\hbar \gamma_0$ to be of the same order, leading to the creation of non-Gaussian light in the steady state. One of the attractive features of this method is that the light is produced completely deterministically and continuously while it emerges from the microcavity. This is in contrast to most other existing methods which work either conditionally, or with an efficiency that is typically less than 1 for deterministic schemes.⁷ Moreover, the created quantum states have an average photon number much greater than one, thus are much brighter than those from other sources of non-Gaussian light such as single photon emitters.

In the work of Schwendimann *et al.*, a master equation for exciton-polariton condensates was derived, where the "system" is considered to be the k = 0 condensate, and the "reservoir" is all the modes with $k \neq 0$.^{31,32} The equations

are essentially a variation of standard lasing equations, where there are a combination of one and two polariton loss and gain terms, which under suitable conditions gives rise to a condensate population. In Ref. 33 the authors obtain the Fokker-Planck equations for the Wigner functions, and solve the equations numerically and analytically for the approximate case. The analytical form of the radial dependence of the Wigner function showed no trace of negativity, thus based on this alone it appears that exciton-polaritons cannot be used to generate coherent non-Gaussian light that would be useful for continuous variables.

There is, however, an aspect which was not discussed in Ref. 33, which generally applies to all lasing master equations. For simplicity let us first consider the standard lasing equations, with a hypothetical built-in nonlinearity which will be clear that it is analogous to the polariton case. The Scully-Lamb lasing equation³⁴ with such a nonlinearity reads

$$\frac{d\rho}{dt} = -\frac{i}{\hbar} [H_{\text{int}}, \rho] - \frac{A}{2} \mathcal{L}[a^{\dagger}, \rho] - \frac{\gamma}{2} \mathcal{L}[a, \rho]
+ \frac{B}{8} [\rho (aa^{\dagger})^2 + 3aa^{\dagger} \rho aa^{\dagger} - 4a^{\dagger} \rho aa^{\dagger} a + \text{H.c.}], \quad (1)$$

where

$$H_{\rm int} = \frac{U}{2} a^{\dagger} a^{\dagger} a a \tag{2}$$

is the nonlinear interaction term, and

$$\mathcal{L}[O,\rho] \equiv \rho O^{\dagger} O + O^{\dagger} O \rho - 2O\rho O^{\dagger} \tag{3}$$

is a Lindblad loss or gain term, with O an arbitrary operator. Here A is the gain coefficient, B is the self-saturation coefficient, γ is the cavity loss rate, and U is the nonlinear Kerr interaction (self-interaction for condensates). A similar system for a $\chi^{(2)}$, as opposed to the $\chi^{(3)}$ nonlinearity as we consider here, was analyzed in Refs. 35–37.

The steady-state solution of these equations can be obtained by setting $\frac{d\rho}{dt}=0$. From the structure of Eq. (1) it can be deduced that in the steady-state limit only the diagonal components are nonzero.² This is most easily seen by calculating the behavior of $\frac{dP_k}{dt}$, where $P_k=\sum_n \rho_{n,n+k}$. For diagonal terms $\frac{dP_0}{dt}=0$, so that $P_0=1$ for all times. In contrast, the off-diagonal terms decay away, and in the steady-state limit tend to zero. Physically, this can be interpreted as the effect of phase diffusion, where any initial coherence becomes randomized in the long-time limit. Far above threshold, the density matrix of such a state is

$$\rho = \frac{1}{2\pi} \int d\theta |re^{i\theta}\rangle \langle re^{i\theta}| = e^{-r^2} \sum_{n=0}^{\infty} \frac{r^{2n}}{n!} |n\rangle \langle n|, \qquad (4)$$

where $|re^{i\theta}\rangle$ is a coherent state and $|n\rangle$ are Fock states.³⁸ The state is diagonal in the density matrix, and the Wigner function has a "doughnut" shape with equal amplitude but a mixture of all phases [see Fig. 1(a)]. The point is then that for the diagonal components of the density matrix, the nonlinear term H_{int} is completely ineffective, no matter how large U may be. This can be seen by looking at the matrix elements

$$\langle n|[H_{\rm int},\rho]|m\rangle = \frac{U}{2} [n(n-1) - m(m-1)] \rho_{nm},$$
 (5)

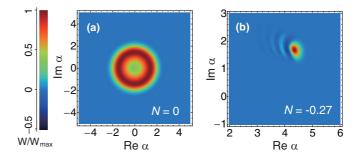


FIG. 1. (Color online) The steady-state Wigner function for (a) a laser with no phase fixing K=0 and (b) an injection locked laser with nonlinear interaction U=0.1 and K=50. Parameters used are A=3, B=1, $\gamma=1$. The negativities $N=\int d^2\alpha [W(\alpha)-|W(\alpha)|]/2$ are marked on the plots.

which disappear for diagonal components. Since diagonal components only couple to other diagonal components in (1), the distribution is independent of U.

How then is it possible to make use of the nonlinearity? Clearly we must recover the off-diagonal terms in the density matrix, instead of the completely phase-diffused state (4). This can be achieved by employing a suitable phase fixing method such as injection locking³⁹ or feedback phase stabilization. 40,41 The effect of such phase fixing is to take a completely mixed state such as (4), which is immune to the interaction H_{int} , and make it approach a pure state depending upon the strength of the phase fixing. For a pure state it is well known that the interaction H_{int} produces highly non-Gaussian states such as Schrödinger cat states, 42 and hence we may expect that the interaction will produce non-Gaussian features to the state of the light.

For simplicity, in this Rapid Communication we consider injection locking to be the phase fixing mechanism. This corresponds to adding a coherent term³⁹

$$H_{\text{lock}} = i\hbar K_0 (ba^{\dagger} e^{-i(\omega - \omega_0)t} - ab^{\dagger} e^{i(\omega - \omega_0)t})$$
 (6)

to the master equation (1), so that the first term is $-\frac{i}{\hbar}[H_{\rm int} + H_{\rm lock}, \rho]$. Here ω_0 is the frequency of the (slave) laser, and we work in the rotating frame $a \equiv a_S e^{-i\omega_0 t}$, where a_S is the operator in the stationary frame. b is the annihilation operator for the injected laser, and ω is its frequency which has been factored out. We assume that bright, on-resonant $(\omega_0 = \omega)$ coherent light is used for the injected master laser, so that we may replace $b \to B e^{-i\psi}$, where B > 0 is a c number. Assuming without loss of generality that the injection locking laser has phase $\psi = 0$, and converting (1) into a Fokker-Planck equation for the Wigner function, we obtain

$$\begin{split} \frac{dW}{dt} &= -K \left(\cos \theta - \frac{\sin \theta}{r} \right) \frac{\partial W}{\partial r} \\ &+ \frac{U}{\hbar} \left[(1 - r^2) \frac{\partial W}{\partial \theta} + \frac{1}{16} \left(\frac{1}{r} \frac{\partial^2 W}{\partial r \partial \theta} + \frac{\partial^3 W}{\partial r^2 \partial \theta} \right. \\ &+ \left. \frac{1}{r^2} \frac{\partial^3 W}{\partial \theta^3} \right) \right] - \frac{1}{2r} \frac{\partial}{\partial r} [r^2 (A - \gamma - Br^2) W] \\ &+ \frac{(A + \gamma)}{8} \left[\frac{1}{r} \frac{\partial}{\partial r} \left(r \frac{\partial W}{\partial r} \right) + \frac{1}{r^2} \frac{\partial^2 W}{\partial \theta^2} \right], \end{split}$$
(7)

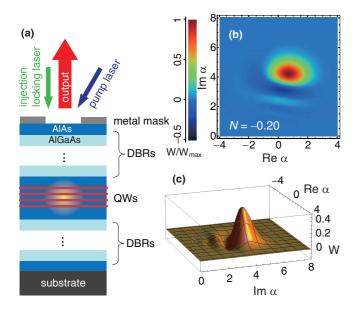


FIG. 2. (Color online) (a) The schematic device configuration. A standard exciton-polariton microcavity with distributed Bragg reflectors (DBRs) is pumped to form a condensate in the quantum wells (QWs). Here we show the example of an AlAs/AlGaAs structure that could be used for the device, although other materials could be used. An injection locking master laser is injected perpendicularly in order to add a phase fixing term to the condensate. The results of this Rapid Communication are concerned with light labeled as "output" emerging from the microcavity. (b), (c) Typical steady-state output of the injection locked polariton system. Parameters used are $D_2/D_1 = 10$, $G_1/D_1 = 1$, $G_2 = 0$, $U/D_1 = 10$, $K/D_1 = 10^3$.

where $K = K_0 B$. The steady-state solution of such an equation is shown in Fig. 1(b). We see that the phase fixing combined with an interaction U leads to a steady-state Wigner function with non-Gaussian features, signaled by the presence of a negative region. This negativity is very stable with respect to initial conditions and remains a feature of the distribution once the steady-state regime is reached. Of course, such a laser with a built-in nonlinearity will in practice have exceedingly small values of U, creating extremely small negativities in the Wigner function. We shall, however, see that for exciton-polaritons such nonlinearities are within experimental reach.

We first describe our proposed device configuration [Fig. 2(a)]. A standard microcavity quantum well system is used, where exciton-polaritons are pumped by an excitation laser or electrical injection. A secondary injection locking laser is introduced at normal incidence of variable intensity. Metal deposition on the surface of the microcavity allows for a trapping potential which confines the condensate area A. Metal deposition is known to create an effective potential which traps the exciton-polariton condensate. A lternatively, the microcavity can be etched into either micropillars or mesas in order to produce the spatial confinement. The emerging photoluminescence is then used as the output light with non-Gaussian properties.

To describe this system, our starting point is the theory of Refs. 31 and 32, where a master equation for the polariton condensate is derived. Including the injection locking terms, the density matrix of the condensate evolves in time

according to

$$\frac{d\rho}{dt} = -\frac{i}{\hbar} [H_{\text{int}} + H_{\text{lock}}, \rho] - \Gamma_2 \mathcal{L}[aa, \rho] - \Delta_2 \mathcal{L}[a^{\dagger}a^{\dagger}, \rho]
- (\Gamma_1 + \gamma_0) \mathcal{L}[a, \rho] - \Delta_1 \mathcal{L}[a^{\dagger}, \rho].$$
(8)

Here a is the annihilation operator for a k=0 polariton, and ω_0 is its frequency (which may include a mean-field energy shift due to interactions). $\Gamma_{1,2}$ are one and two polariton loss terms due to polariton scattering, $\Delta_{1,2}$ are one and two polariton gain terms due to scattering, and γ_0 is a loss rate due to leakage through the microcavity mirrors. From (8) it is possible to derive a Fokker-Planck equation for the Wigner function.³³ Using standard techniques³ we obtain

$$\begin{split} \frac{\partial W}{\partial t} &= -2K \left(\cos \theta - \sin \theta \frac{1}{r} \right) \frac{\partial W}{\partial r} - \frac{U}{\hbar} (r^2 - 1) \frac{\partial W}{\partial \theta} \\ &+ \frac{U}{16\hbar} \left(\frac{1}{r} \frac{\partial^2 W}{\partial r \partial \theta} + \frac{\partial^3 W}{\partial r^2 \partial \theta} + \frac{1}{r^2} \frac{\partial^3 W}{\partial \theta^3} \right) \\ &- (2G_2 + 8G_1 r^2) W + (-G_2 + 2D_1 - 2G_1 r^2) r \frac{\partial W}{\partial r} \\ &+ \left(\frac{D_2}{4} + D_1 r^2 - \frac{G_1}{2} \right) \left[\frac{1}{r} \frac{\partial W}{\partial r} + \frac{\partial^2 W}{\partial r^2} + \frac{1}{r^2} \frac{\partial^2 W}{\partial \theta^2} \right], \end{split}$$

where we have retained only derivatives up to second order in the Lindblad terms in (8), since this gives a very small correction to the overall dynamics. Here $G_1 = \Delta_1 - \Gamma_1$, $G_2 = \Delta_2 - \Gamma_2 - \gamma_0$, $D_1 = \Delta_1 + \Gamma_1$, $D_2 = \Delta_2 + \Gamma_2 + \gamma_0$ are the net gain and diffusion coefficients. The precise expressions for G_i and D_i may be calculated using the expressions in Refs. 31–33. While different materials give rise to different parameters, the integral expressions given in the above references possess certain properties which are common to all systems. As can be deduced from dimensional analysis, the order of the parameters are approximately

$$\Delta_1, \Gamma_1 \sim \frac{V_0^2 A}{4\pi\hbar E_0 a^2}, \quad \Delta_2, \Gamma_2 \sim \frac{V_0^2 A^2}{8\pi^3 \hbar E_0 a^4},$$
(9)

where $V_0 = 6e^2a_B/4\pi \epsilon A$ is the polariton-polariton interaction energy, A is the quantization area, a is a temperature length scale set according to $E_0 = \hbar^2/2m_{\rm exc}a^2 = k_BT$, where T is the temperature, and a_B is the exciton Bohr radius. We can observe that the order $O(\Delta_1/\Delta_2) \sim O(\Gamma_1/\Gamma_2) \sim 2\pi^2a^2/A$ is typically a small parameter such that $\Delta_1, \Gamma_1 \ll \Delta_2, \Gamma_2$. However, Δ_1, Γ_1 cannot be neglected as without these condensation is unstable. Meanwhile the interaction can be estimated to be⁴⁶

$$U \sim \frac{30e^2a_B|X|^4}{\pi^3\epsilon A},$$
 (10)

where X is the excitonic Hopfield coefficient. From the expressions in Ref. 31 it can be deduced that $G_1 < 0$ and $D_{1,2} > 0$, but G_2 can change sign, which can be taken to be the criterion for condensation assuming $\Delta_1 \ll G_2$.³¹

In Figs. 2(b) and 2(c) we plot the typical Wigner distribution for the steady-state operation of the injection locked polariton BEC. The distribution typically consists of a bright peak that is phase fixed due to the injection locking. Adjacent to the peak, there is a region of negativity, approximately in the direction of origin. The negativity appears as soon as the injection locking is switched on, and at first grows

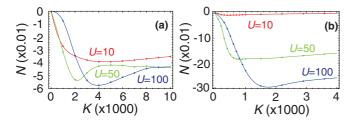


FIG. 3. (Color online) Dependence of the total negativity N with injection locking strength K for various interactions U (in units of D_1) for the polariton BEC. Steady-state values are taken for (a) estimated experimental parameters $D_2/D_1=100$, (b) very high Q cavity/small trap case $D_2/D_1=10$. Other parameters are $G_1/D_1=1$, $G_2=0$.

with both injection locking strength K and interaction strength U. For all parameters K>0, U>0 we see stable negativities of the Wigner function, quantified by $N=\int d^2\alpha [W(\alpha)-|W(\alpha)|]/2$. Figure 3(a) shows the behavior of the amount of steady-state negativity with K. At large enough K, there is a saturation of the negativity, beyond which the total negativity no longer increases, and in fact decreases slowly beyond this point. Although the overall negativity decreases for very large K, the Wigner distribution tends to become more sharply defined for large K. For smaller values of K, the negative region tends to be shallower, but distributed over a larger area. In Fig. 3(b) we plot the negativities for $D_2/D_1=10$, which corresponds to very small $(r=0.16~\mu\text{m})$ traps or very high-Q cavities giving very long polariton lifetimes $(1/\gamma_0=1~\text{ns})$. We see that the reduced phase diffusion improves the negativity of

the Wigner function. We generally also observe that close to threshold gives the largest negativities.

In conclusion, we have proposed a device for deterministically producing bright coherent non-Gaussian light from an exciton-polariton condensate. It was shown that steady-state generation of such light is possible using currently accessible experimental parameters when a suitable method of phase fixing is imposed on the polariton condensate. Although we have considered injection locking here, other methods such as feedback control of the phase should accomplish the same task. The phase fixing overcomes the phase diffusion which tends to dephase the condensate and leads to a completely mixed state; with phase fixing, non-Gaussian states are created exploiting the natural nonlinearity of the polaritons. The completely deterministic nature of the production of these non-Gaussian states, as well as their brightness, is potentially useful in the context of continuous-variable quantum information processing, for which at least some resource states must possess a negative Wigner function. On-demand sources will save quantum memories and help scalability, while mean photon numbers above unity allow for making use of larger Hilbert spaces encoded into the infinite-dimensional space of an optical mode.

T.B. thanks Howard Wiseman, Terry Rudolph, and Hui Deng for discussions. This work is supported by the Special Coordination Funds for Promoting Science and Technology, Navy/SPAWAR Grant No. N66001-09-1-2024, MEXT, and the JSPS through its FIRST program.

¹R. J. Glauber, Phys. Rev. **130**, 2529 (1963).

²M. O. Scully and M. S. Zubairy, *Quantum Optics* (Cambridge University Press, Cambridge, UK, 1997).

³D. F. Walls and G. J. Milburn, *Quantum Optics* (Springer, Berlin, 2008).

⁴H. A. Bachor and T. C. Ralph, *A Guide to Experiments in Quantum Optics* (Wiley-VCH, Weinheim, 2004).

⁵A. Ourjoumtsev, R. Tualle-Brouri, J. Laurat, and P. Grangier, Science **312**, 83 (2006).

⁶J. S. Neergaard-Nielsen, B. M. Nielsen, C. Hettich, K. Molmer, and E. S. Polzik, Phys. Rev. Lett. **97**, 083604 (2006).

⁷M. D. Eisaman, J. Fan, A. Migdall, and S. V. Polyakov, Rev. Sci. Instrum. **82**, 071101 (2011).

⁸S. L. Braunstein and P. van Loock, Rev. Mod. Phys. 77, 513 (2005).
⁹C. Weedbrook, S. Pirandola, R. García-Patrón, N. J. Cerf, T. C. Ralph, J. H. Shapiro, and S. Lloyd, Rev. Mod. Phys. 84, 621 (2012).

¹⁰J. Eisert and M. Plenio, Int. J. Quantum Inf. 1, 479 (2003).
¹¹G. Adesso and F. Illuminati, J. Phys. A: Math. Theor. 40, 7821

¹¹G. Adesso and F. Illuminati, J. Phys. A: Math. Theor. **40**, 782 (2007).

¹²A. Ferraro, S. Olivares, and M. G. A. Paris, *Gaussian States in Quantum Information* (Bibliopolis, Napoli, 2005).

¹³A. Furusawa, J. L. Sorensen, S. L. Braunstein, C. A. Fuchs, H. J. Kimble, and E. S. Polzik, Science 282, 706 (1998).

¹⁴S. Lloyd and S. L. Braunstein, Phys. Rev. Lett. **82**, 1784 (1999).

¹⁵S. D. Bartlett and B. C. Sanders, Phys. Rev. A **65**, 042304 (2002).

¹⁶S. Sefi and P. van Loock, Phys. Rev. Lett. **107**, 170501 (2011).

¹⁷T. Broecker and R. F. Werner, J. Math. Phys. **36**, 62 (1995).

¹⁸A. Mandilara, E. Karpov, and N. J. Cerf, Phys. Rev. A **79**, 062302 (2009).

¹⁹D. Gross, J. Math. Phys. **47**, 122107 (2006).

²⁰A. Mari and J. Eisert, Phys. Rev. Lett. **109**, 230503 (2012).

²¹V. Veitch, C. Ferrie, D. Gross, and J. Emerson, New J. Phys. 15, 013037 (2013).

²²K. Nemoto and W. J. Munro, Phys. Rev. Lett. **93**, 250502 (2004).

²³M. Bartkowiak, L.-A. Wu, and A. Miranowicz, arXiv:1210.2384.

²⁴H. Deng, H. Haug, and Y. Yamamoto, Rev. Mod. Phys. **82**, 1489 (2010).

²⁵J. Kasprzak *et al.*, Nature (London) **443**, 409 (2006).

²⁶H. Deng, G. Weihs, C. Santori, J. Bloch, and Y. Yamamoto, Science 298, 199 (2002).

²⁷T. Byrnes, T. Horikiri, N. Ishida, M. Fraser, and Y. Yamamoto, Phys. Rev. B **85**, 075130 (2012).

²⁸A. Amo et al., Nature (London) **457**, 291 (2009).

²⁹B. Nelsen, G. Liu, M. Steger, D. W. Snoke, R. Balili, K. West, and L. Pfeiffer, arXiv:1209.4573.

³⁰N. Y. Kim *et al.*, Phys. Status Solidi **245**, 1076 (2008).

³¹P. Schwendimann and A. Quattropani, Phys. Rev. B **74**, 045324 (2006).

³²P. Schwendimann and A. Quattropani, Phys. Rev. B **77**, 085317 (2008)

³³P. Schwendimann, A. Quattropani, and D. Sarchi, Phys. Rev. B 82, 205329 (2010).

- ³⁴M. Sargent, M. O. Scully, and W. E. Lamb, *Laser Physics* (Addison-Wesley, Boston, 1978).
- ³⁵A. G. White, T. C. Ralph, and H.-A. Bachor, J. Opt. Soc. Am. B 13, 1337 (1996).
- ³⁶R. Schack, A. Sizmann, and A. Schenzle, Phys. Rev. A **43**, 6303 (1991).
- ³⁷A. Sizmann, R. Schack, and A. Schenzle, Europhys. Lett. 13, 109 (1990).
- $^{38}\text{T.}$ Rudolph and B. C. Sanders, Phys. Rev. Lett. $\pmb{87},\,077903$ (2001).
- ³⁹H. A. Haus and Y. Yamamoto, Phys. Rev. A **29**, 1261 (1984).
- ⁴⁰H. M. Wiseman and G. J. Milburn, Phys. Rev. Lett. **70**, 548 (1993).
- ⁴¹H. M. Wiseman and G. J. Milburn, Phys. Rev. A **49**, 1350 (1994).

- ⁴²M. Stobinska, G. J. Milburn, and K. Wódkiewicz, Phys. Rev. A 78, 013810 (2008).
- ⁴³S. I. Tsintzos, P. G. Savvidis, G. Deligeorgis, Z. Hatzopoulos, and N. T. Pelekanos, Appl. Phys. Lett. **94**, 071109 (2009).
- ⁴⁴A. Löffler, J. P. Reithmaier, G. Sek, C. Hofmann, S. Reitzenstein, M. Kamp, and A. Forchel, Appl. Phys. Lett. **86**, 111105 (2005).
- ⁴⁵R. I. Kaitouni, O. El Daif, A. Baas, M. Richard, T. Paraiso, P. Lugan, T. Guillet, F. Morier-Genoud, J. D. Ganiere, J. L. Staehli, V. Savona, and B. Deveaud, Phys. Rev. B 74, 155311 (2006).
- ⁴⁶T. Byrnes, P. Recher, and Y. Yamamoto, Phys. Rev. B **81**, 205312 (2010).