

Broadband pseudothermal states with tunable spectral coherence generated via nonlinear optics

Nicolás Quesada and Agata M. Brańczyk*

Perimeter Institute for Theoretical Physics, Waterloo, Ontario, Canada N2L 2Y5

(Received 15 October 2018; published 17 January 2019)

It is well known that the reduced state of a two-mode squeezed vacuum state is a thermal state, i.e., a state whose photon-number statistics obey a geometric distribution. More exotic *broadband* states can be realized as the reduced state of two spectrally entangled beams generated using nonlinear optics. We show that these broadband “pseudothermal” states are tensor products of states in spectral Schmidt modes, whose photon-number statistics obey a geometric distribution. We study the spectral and temporal coherence properties of these states and show that their spectral coherence can be tuned—from perfect coherence to complete incoherence—by adjusting the pump spectral width. In the limit of a cw pump, these states are tensor products of true thermal states but with different temperatures at each frequency. This could be an interesting state of light for investigating the interplay between spectral, temporal, and photon-number coherences.

DOI: [10.1103/PhysRevA.99.013830](https://doi.org/10.1103/PhysRevA.99.013830)**I. INTRODUCTION**

Thermal states are of fundamental and practical interest. Although they are diagonal in the photon-number and coherent-state bases, they can behave nonclassically. They can be used for generating nonclassical states [1,2] or for mediating entanglement between quantum systems [3]. They can also be used for quantum information protocols such as continuous-variable quantum key distribution [4] and improving the efficiency of quantum state tomography [5].

In quantum information, one often deals with single-mode thermal states. There, the relevant property is the photon-number statistics—the thermal state density matrix is diagonal with a geometric probability distribution.

In quantum optics, when considering many radiation modes, the temperature T takes a more central role, as it determines the light’s spectral radiance according to Planck’s law. For multimode thermal light, one can speak about its spatial, spectral, temporal, and momentum coherence.

Incoherence in these degrees of freedom can be useful. Spatially incoherent light has been used for ghost imaging [6–12], subwavelength lithography [13], and improving diffraction pattern visibility [14] and spatial resolution [15]. Broadband spectrally incoherent light has been used for resolution-enhanced optical coherence tomography [16], optical guiding of microscopic particles [17], and noisy-light spectroscopy [18,19].

Various methods exist for generating incoherent light. Spatially incoherent, pseudothermal light can be generated by sending a cw laser through a rotating ground-glass disk [20,21]. Broadband, spectrally incoherent light can be generated by a thermal source, e.g., a hollow cathode lamp [9]. Other approaches use amplified spontaneous emission from quantum dots, [22,23], warm atomic vapor [24], and modified dye lasers [25]. Here, we focus on light generated in one

arm of a broadband twin-beam state, such as that generated via spontaneous parametric downconversion (SPDC) [26,27] or spontaneous four-wave mixing (SFWM) [28]. Spectral entanglement between the beams makes each individual beam spectrally incoherent.

Photon statistics and coherence can be studied by measuring correlation functions. Various groups measured such functions on light generated via nonlinear optics. These include measurements of time-resolved temporal correlation functions in cw-pumped SPDC [29], time-averaged temporal correlation functions in two-mode SPDC [26,30], frequency cross- and autocorrelation functions in two-mode SPDC [31], multiphoton statistics of single-mode SPDC [32], and frequency-resolved spectral correlation functions and multiphoton statistics in harmonic generation [27]. Theory has also been done on photon-number statistics [33] and spatial correlation functions [6,7] of downconverted light. But to the best of our knowledge, no one has written down the reduced density operator for one arm of an arbitrary spectrally entangled twin-beam state, nor has anyone computed its time-resolved temporal and frequency-resolved spectral correlation functions.

In this paper, we show that the density operator describing one arm of a twin-beam state with arbitrary spectral entanglement can be decomposed into a tensor product of states prepared in spectral Schmidt modes, each with geometric photon-number statistics. We therefore refer to the state as a broadband “pseudothermal” (BPT) state.

We then write down Heisenberg-picture operators for the Schmidt modes and compute the BPT state’s frequency-resolved spectral and time-resolved first- and second-order correlation functions. These functions reveal the light’s spectral and temporal coherence, as well as its intensity-intensity correlations. We find that the spectral coherence of BPT states can be tuned—from perfect coherence to full incoherence—by adjusting the pump spectral width. This is consistent with recent experiments [30]. (The spectral coherence can also be tuned by changing the spectral phase of the pump, e.g., by

*abrancyk@pitp.ca

applying a chirp [34], although we do not consider this here.) From these correlation functions, one can identify interesting regimes not yet explored experimentally, such as partial spectral coherence.

In summary, the main contributions of this paper are three-fold. First, we derive the reduced density operator of one beam of a twin-beam state with arbitrary spectral entanglement in terms of its Schmidt modes. This expression has conceptual value and provides intuition about the nature of the light in each beam. Second, we derive time- and frequency-resolved correlation functions for one beam of a twin-beam state in terms of its Schmidt modes and explore the relationship between interbeam spectral entanglement and single-beam temporal coherence. Third, we make clear the connection between these correlation functions and results from classical coherence theory. Our results should therefore be useful for both experimental and theoretical studies of partially spectrally coherent light.

II. BROADBAND TWO-MODE SQUEEZED VACUUM STATE

We consider broadband light, generated by, e.g., SPDC or SFWM, emitted into two orthogonal modes. In a one-dimensional propagation geometry where the longitudinal wave vector is specified by the frequency and assuming that the pump beam remains undepleted, the twin-beam state is

$$|\psi\rangle = \hat{U}_{\text{SQ}} |\text{vac}\rangle, \quad (1)$$

$$\hat{U}_{\text{SQ}} = e^{[\iint d\omega_a d\omega_b J(\omega_a, \omega_b) \hat{a}^\dagger(\omega_a) \hat{b}^\dagger(\omega_b) - \text{H.c.}]}, \quad (2)$$

where $J(\omega_a, \omega_b)$ is known as the *joint spectral amplitude* (JSA) of the generated beams [35–37] and \hat{U}_{SQ} is the broadband two-mode squeezing operator. The operators $\hat{a}(\omega)$ and $\hat{b}(\omega)$ are single-frequency annihilation operators that satisfy the commutation relations $[\hat{a}(\omega), \hat{a}^\dagger(\omega')] = [\hat{b}(\omega), \hat{b}^\dagger(\omega')] = \delta(\omega - \omega')$ (all other commutators are zero). The JSA depends on the properties of the pump field(s) and the nonlinear material. For purposes of this paper, we leave it quite general.

To simplify calculations, the JSA can be decomposed as $J(\omega_a, \omega_b) = \sum_k r_k \phi_k(\omega_a) \varphi_k(\omega_b)$ in what is known as the Schmidt decomposition. The twin-beam state can then be written as [26]

$$|\psi\rangle = e^{\sum_k r_k \hat{A}_k^\dagger \hat{B}_k^\dagger - \text{H.c.}} |\text{vac}\rangle, \quad (3)$$

where the operators

$$\hat{A}_k = \int d\omega_a \phi_k^*(\omega_a) \hat{a}(\omega_a), \quad (4a)$$

$$\hat{B}_k = \int d\omega_b \varphi_k^*(\omega_b) \hat{b}(\omega_b), \quad (4b)$$

are broadband annihilation operators that satisfy the commutation relations $[\hat{A}_k, \hat{A}_{k'}^\dagger] = [\hat{B}_k, \hat{B}_{k'}^\dagger] = \delta_{k, k'}$ (all other commutators are zero), and ϕ_k and φ_k are known as Schmidt modes; these functions satisfy completeness and orthogonality relations [38].

Analytical forms for the Schmidt decomposition are known only for two-dimensional Gaussian functions [39]. For more general functions, one can use approximate numerical methods, e.g., computing the singular value decomposition of a truncated, discretized JSA. The impact of such approximations in SPDC source characterization was recently discussed in [40].

One can also invert the relations in Eqs. (4) and find

$$\hat{a}(\omega) = \sum_k \phi_k(\omega) \hat{A}_k, \quad (5a)$$

$$\hat{b}(\omega) = \sum_k \varphi_k(\omega) \hat{B}_k. \quad (5b)$$

The twin-beam state in Eq. (3) can be rewritten as

$$|\psi\rangle = \bigotimes_k |r_k\rangle, \quad (6)$$

where

$$|r_k\rangle = e^{r_k \hat{A}_k^\dagger \hat{B}_k^\dagger - \text{H.c.}} |\text{vac}\rangle \quad (7)$$

is a two-mode squeezed vacuum (TMSV) state prepared in two Schmidt modes ϕ_k and φ_k , with *squeezing parameter* $r_k \geq 0$. Note that here $J(\omega_a, \omega_b)$ is not necessarily normalized and thus $\sum_k r_k^2$ does not necessarily equal 1. Each TMSV state can be represented in the number basis

$$|r_k\rangle = \sum_{n_k=0}^{\infty} \frac{(\tanh r_k)^{n_k}}{\cosh(r_k)} |n_k\rangle_{\phi_k} |n_k\rangle_{\varphi_k}. \quad (8)$$

In the Heisenberg picture, the operators \hat{A}_k and \hat{B}_k transform as

$$\begin{aligned} \hat{A}_k &\rightarrow \hat{U}_{\text{SQ}}^\dagger \hat{A}_k \hat{U}_{\text{SQ}} \\ &= \cosh(r_k) \hat{A}_k + \sinh(r_k) \hat{B}_k^\dagger, \end{aligned} \quad (9a)$$

$$\begin{aligned} \hat{B}_k &\rightarrow \hat{U}_{\text{SQ}}^\dagger \hat{B}_k \hat{U}_{\text{SQ}} \\ &= \cosh(r_k) \hat{B}_k + \sinh(r_k) \hat{A}_k^\dagger. \end{aligned} \quad (9b)$$

Later, we will use the relations in Eqs. (5) and the transformation in Eqs. (9) to compute correlation functions.

III. BROADBAND PSEUDOTHERMAL STATES

We are interested in the quantum state of the individual beams; we thus compute the reduced density matrices for modes a and b by tracing out the other mode. The density operator for mode a is

$$\rho_a = \text{Tr}_b[|\psi\rangle\langle\psi|] = \bigotimes_k \rho_{\phi_k}, \quad (10)$$

where

$$\rho_{\phi_k} = \text{Tr}_{\varphi_k}[|r_k\rangle\langle r_k|] \quad (11)$$

$$= \sum_{n_k=0}^{\infty} P_{n_k} |n_k\rangle_{\phi_k} \langle n_k|_{\phi_k} \quad (12)$$

is a state prepared in a single Schmidt mode ϕ_k . The states $|n_k\rangle_{\phi_k} = (n_k!)^{-1/2}(\hat{A}_k^\dagger)^{n_k}|0\rangle$ are broadband Fock states and are distributed according to

$$P_{n_k} = \frac{1}{1 + \bar{n}_k} \left(\frac{\bar{n}_k}{1 + \bar{n}_k} \right)^{n_k}, \quad (13)$$

where

$$\bar{n}_k = \sinh^2(r_k). \quad (14)$$

The state ρ_{ϕ_k} is like a single-mode thermal state in the sense that it is diagonal in the photon-number basis and the probability distribution is geometric. But since the mode is not at a well-defined frequency, it doesn't make sense to talk about an associated temperature T .

Similarly, the beam in mode b has the state $\rho_b = \bigotimes_k \rho_{\phi_k}$, where $\rho_{\phi_k} = \sum_{n_k=0}^{\infty} P_{n_k} |n_k\rangle_{\phi_k} \langle n_k|_{\phi_k}$. While both ρ_a and ρ_b obey the same statistics given by P_{n_k} , the states will have different spectral and temporal properties because the spectral properties of $|n_k\rangle_{\phi_k}$ differ from those of $|n_k\rangle_{\varphi_k}$.

The states in Eq. (12) can be written more succinctly as (see Appendix A)

$$\rho_{\phi_k} = \frac{1}{Z_k} e^{-\alpha_k \hat{A}_k^\dagger \hat{A}_k}, \quad (15)$$

$$Z_k = \text{Tr}(e^{-\alpha_k \hat{A}_k^\dagger \hat{A}_k}) = \frac{1}{1 - e^{-\alpha_k}}, \quad (16)$$

where Z_k is the partition function of mode k and

$$e^{-\alpha_k} = \tanh^2(r_k) = \frac{\bar{n}_k}{1 + \bar{n}_k}. \quad (17)$$

Using this notation we can also write the full state in mode a as

$$\rho_a = \frac{1}{Z} e^{-\sum_k \alpha_k \hat{A}_k^\dagger \hat{A}_k}, \quad (18)$$

$$Z = \text{Tr}(e^{-\sum_k \alpha_k \hat{A}_k^\dagger \hat{A}_k}). \quad (19)$$

IV. PROPERTIES OF BROADBAND PSEUDOTHERMAL STATES

To study the coherence properties of the broadband states, we compute various temporal and spectral correlation functions. The expressions for mode a can be mapped to those for mode b by making the substitution $\phi_k \rightarrow \varphi_k$.

A. Spectral correlation function

To study the spectral coherence properties of the light, we introduce a spectral correlation function $S(\omega, \omega')$. Using the procedure outlined in Appendix B, one finds that

$$S_a(\omega, \omega') = \langle \hat{a}^\dagger(\omega) \hat{a}(\omega') \rangle_\psi \quad (20)$$

$$= \sum_k \sinh^2(r_k) \phi_k^*(\omega) \phi_k(\omega'), \quad (21)$$

and also that $\langle \hat{a}(\omega) \hat{a}(\omega') \rangle_\psi = \langle \hat{a}^\dagger(\omega) \hat{a}^\dagger(\omega') \rangle_\psi = 0$.

The same-frequency correlation function is

$$S_a(\omega, \omega) = \sum_k |\phi_k(\omega)|^2 \bar{n}_k, \quad (22)$$

which can be interpreted as the spectral density.

B. First-order temporal correlation function

The first-order temporal correlation function for mode a is [41]

$$G_a^{(1)}(t_1, t_2) = \langle \hat{a}^\dagger(t_1) \hat{a}(t_2) \rangle_\psi. \quad (23)$$

In the limit of the state $|\psi\rangle$ having sufficiently narrow frequency support, this quantity is proportional to $\langle \hat{E}^{(-)}(t_1) \hat{E}^{(+)}(t_2) \rangle_\psi$, where $\hat{E}_a^{(\pm)}(t)$ are the usual positive or negative frequency components of the electric field operator in mode a . The operators $\hat{a}(t)$ are nothing but the Fourier transform of the operators $\hat{a}(\omega)$

$$\hat{a}(t) = \frac{1}{\sqrt{2\pi}} \int d\omega \hat{a}(\omega) e^{i\omega t}. \quad (24)$$

We thus have

$$G_a^{(1)}(t_1, t_2) = \langle \hat{a}^\dagger(t_1) \hat{a}(t_2) \rangle_\psi \quad (25)$$

$$= \int \frac{d\omega d\omega'}{2\pi} S(\omega, \omega') e^{-i\omega t_1 + i\omega' t_2} \quad (26)$$

$$= \sum_k \tilde{\phi}_k^*(t_1) \tilde{\phi}_k(t_2) \bar{n}_k, \quad (27)$$

where \bar{n}_k is defined in Eq. (14), and where

$$\tilde{\phi}_k(t) = \frac{1}{\sqrt{2\pi}} \int d\omega \phi_k(\omega) e^{i\omega t}. \quad (28)$$

This expression is derived in Appendix C. To see the temporal distribution, we compute

$$G_a^{(1)}(t, t) = \sum_k |\tilde{\phi}_k(t)|^2 \bar{n}_k, \quad (29)$$

which can be interpreted as the probability per unit time that a photon is absorbed by an ideal detector at time t [41].

We can also compute a normalized first-order correlation function

$$g^{(1)}(t_1, t_2) = \frac{G^{(1)}(t_1, t_2)}{\sqrt{G^{(1)}(t_1, t_1) G^{(1)}(t_2, t_2)}}. \quad (30)$$

In the special case where the decomposition has only one Schmidt mode, the normalized first-order correlation function is $|g^{(1)}(t_1, t_2)| = 1$. In other words, this is a state with finite bandwidth and thermal photon-number statistics, and yet it has perfect first-order coherence [41].

It is known from classical coherence theory that correlation functions for partially coherent light can be decomposed as sums of coherent mode functions [42]. The connection between such spatial coherent modes and spatial Schmidt modes in SPDC was suggested and explored experimentally in [43]. Here, we show this explicitly in terms of spectral coherent modes. Equation (27) shows that for a single beam of a twin-beam state with arbitrary spectral entanglement, these mode functions are indeed the (spectral) Schmidt modes.

C. Second-order temporal correlation function

To see intensity-intensity correlations, we look at the second-order correlation function [41]:

$$G^{(2)}(t_1, t_2) = \langle \hat{a}^\dagger(t_1)\hat{a}^\dagger(t_2)\hat{a}(t_1)\hat{a}(t_2) \rangle_\psi \quad (31)$$

$$= G_a^{(1)}(t_1, t_1)G_a^{(1)}(t_2, t_2) + G_a^{(1)}(t_1, t_2)G_a^{(1)}(t_2, t_1), \quad (32)$$

where we again replaced the electric field operators with photon-number creation and destruction operators, and where $G_a^{(1)}(t_1, t_2)$ is defined in Eq. (26). This expression is derived in Appendix C. Equation (32) can be interpreted as the probability per unit (time)² that one photon is recorded at time t_1 and another at time t_2 [41].

V. CONTINUOUS-WAVE LIMIT

In the case of a cw laser driving an SPDC process at frequency $\bar{\omega}_p$ (or a SFWM process at $\bar{\omega}_p/2$), energy is conserved according to $\omega_a + \omega_b = \bar{\omega}_p$. The two-mode squeezed state has the form

$$|\psi\rangle = \hat{U}_{\text{SQ}} |\text{vac}\rangle, \quad (33)$$

$$\hat{U}_{\text{SQ}} = e^{[\int d\omega r(\omega)\hat{a}^\dagger(\omega)\hat{b}^\dagger(\bar{\omega}_p-\omega) - \text{H.c.}]}. \quad (34)$$

We can also construct Heisenberg-picture transformation (similar to the pulsed-pump case):

$$\begin{aligned} \hat{a}(\omega) &\rightarrow \hat{U}_{\text{SQ}}^\dagger \hat{a}(\omega) \hat{U}_{\text{SQ}} \\ &= \cosh(r(\omega))\hat{a}(\omega) + \sinh(r(\omega))\hat{b}^\dagger(\omega_p - \omega) \end{aligned} \quad (35)$$

$$\begin{aligned} \hat{b}(\omega) &\rightarrow \hat{U}_{\text{SQ}}^\dagger \hat{b}(\omega) \hat{U}_{\text{SQ}} \\ &= \cosh(r(\omega))\hat{b}(\omega) + \sinh(r(\omega))\hat{a}^\dagger(\omega_p - \omega), \end{aligned} \quad (36)$$

and also write states for mode a (or b)

$$\rho_a = \frac{1}{Z} e^{-\int d\omega \alpha(\omega)\hat{a}^\dagger(\omega)\hat{a}(\omega)}, \quad (37)$$

$$Z = \text{Tr}(e^{-\int d\omega \alpha(\omega)\hat{a}^\dagger(\omega)\hat{a}(\omega)}), \quad (38)$$

where $\alpha(\omega) = \ln[1/\tanh^2(r(\omega))]$. By generalizing the result in Appendix B, from a discrete to a continuum index, the spectral correlation function becomes

$$S_{\text{CW}}(\omega, \omega') = \bar{n}_{\text{CW}}(\omega)\delta(\omega - \omega'), \quad (39)$$

$$\bar{n}_{\text{CW}}(\omega) = \sinh^2(r(\omega)). \quad (40)$$

Equation (39) tells us that there are no spectral correlations between two different frequency modes.

The first-order correlation function for cw broadband thermal light is the two-dimensional Fourier transform of $S_{\text{CW}}(\omega, \omega')$. After integrating out the Dirac delta function, this becomes

$$G_{\text{CW}}^{(1)}(t_1, t_2) = \frac{1}{2\pi} \int d\omega e^{i\omega(t_1-t_2)} \bar{n}_{\text{CW}}(\omega), \quad (41)$$

as expected from the Wiener-Khinchin theorem. From this we can compute

$$G_{\text{CW}}^{(1)}(t, t) = \frac{1}{2\pi} \int d\omega \bar{n}_{\text{CW}}(\omega). \quad (42)$$

That Eq. (42) is constant also shows that the light is fully spectrally incoherent, since time-varying intensities arise from well-defined phases between frequencies. Similarly, the second-order correlation function for cw-pumped BPT states is

$$G_{\text{CW}}^{(2)}(t_1, t_2) = G_{\text{CW}}^{(1)}(t_1, t_1)G_{\text{CW}}^{(1)}(t_2, t_2) + G_{\text{CW}}^{(1)}(t_1, t_2)G_{\text{CW}}^{(1)}(t_2, t_1), \quad (43)$$

where $G_{\text{CW}}^{(1)}(t_1, t_2)$ is defined in Eq. (41).

If one sets $\alpha(\omega) = \hbar\omega/k_B T$, then ρ_a can be thought of as a multimode thermal state in the traditional sense. Equation (43) then becomes identical to the expression for $G^{(2)}(t_1, t_2)$ for chaotic light, derived by, e.g., Loudon [44]. The shape of $\alpha(\omega)$ depends on $r(\omega)$, which in turn depends on the nonlinearity profile of the material. It would therefore be difficult to make light that exactly matches the Planck spectrum. But one could match the Planck spectrum over a finite bandwidth or make more general states where each frequency corresponds to a thermal state at a different temperature T .

VI. EXAMPLES

To illustrate the pump laser's impact on the coherence of BPT states, we examine the concrete case of SPDC. We look at three examples: short pulse, long pulse, and cw.

For simplicity, we consider pump fields with Gaussian spectral distributions (e.g., shaped using optical pulse shaping [45]) and crystals with Gaussian nonlinearity profiles (e.g., engineered using nonlinearity shaping methods [46–50]). We consider the system to be in the low-gain regime and satisfy symmetric group-velocity matching [51,52].

We take

$$J(\Omega_a, \Omega_b) = A e^{-\frac{(\Omega_a + \Omega_b)^2}{2\sigma_p^2}} e^{-\frac{(\Omega_a - \Omega_b)^2}{2\sigma_c^2}} \quad (44)$$

for the pulsed case, where $\Omega_j = \omega_j - \bar{\omega}_j$ (with $\bar{\omega}_j$ the mean frequency of the downconverted beam), σ_p is the spectral width of the pump amplitude function, and σ_c is the width of the phase-matching function (given as the Fourier transform of the longitudinal shape of the material nonlinearity). We also take

$$r(\Omega) = A e^{-\frac{(2\Omega)^2}{2\sigma_c^2}} \quad (45)$$

for the cw case, where $\Omega = \omega - \bar{\omega}_a$. Figure 1 compares three examples. For easy comparison, we chose σ_p and σ_c to make BPT states with similar spectral distributions (the choice of A does not affect the normalized functions). These are shown in Table I.

Figure 1(a) shows the JSA of the twin-beam state. The JSA for the shorter-pulse case is separable (note that this only happens for a pair of *Gaussians* of the same width [53]). The longer-pulsed laser leads to a slightly correlated JSA, and the cw laser leads to a strongly correlated JSA.

Figure 1(b) shows the spectral distributions in mode a , chosen to be the same for all cases. Figure 1(c) shows the

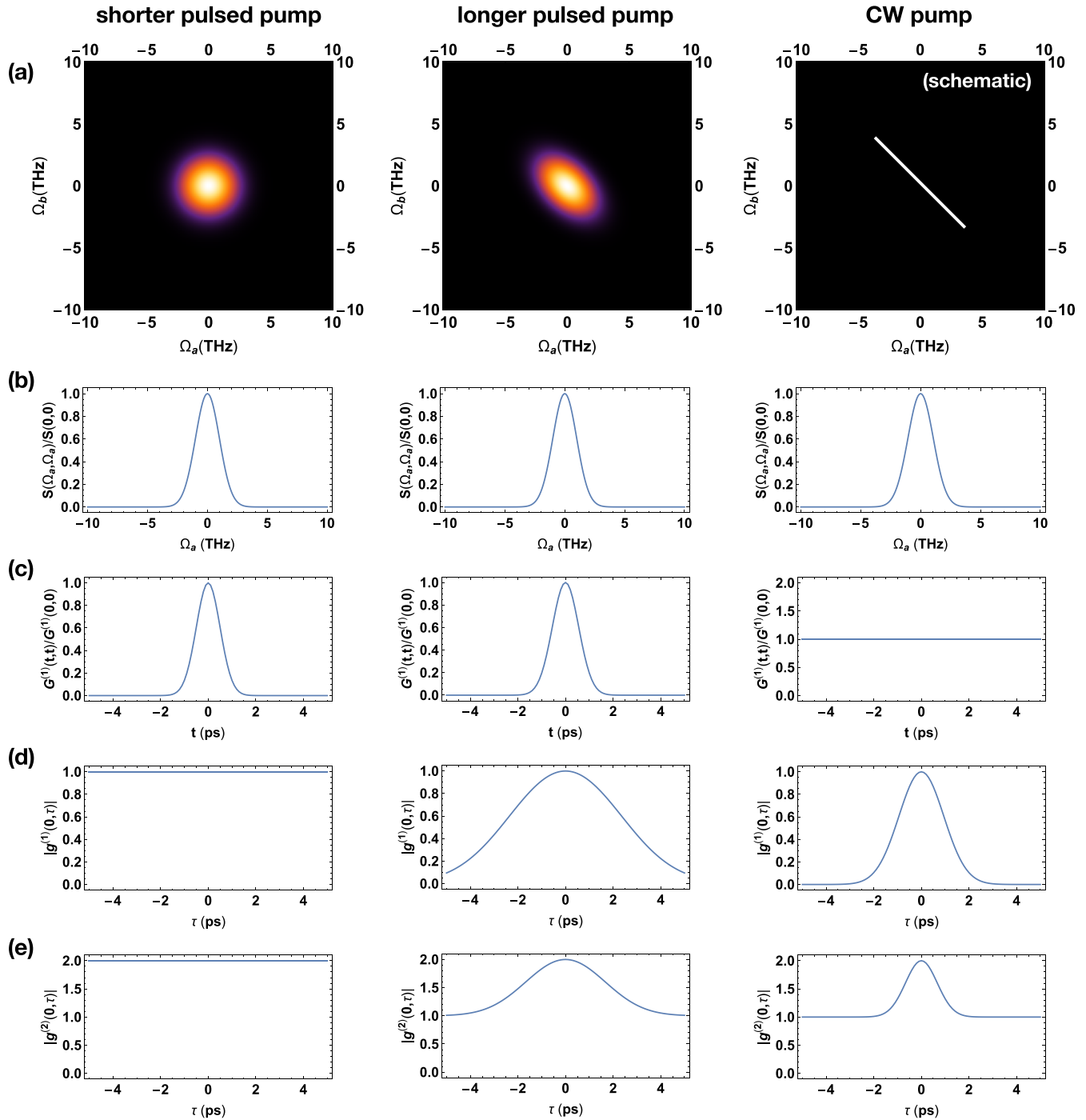


FIG. 1. Comparison between sources pumped by a shorter-pulsed, longer-pulsed, and cw laser: (a) the joint spectral amplitude $J(\omega_1, \omega_2)$; (b) equal-frequency spectral correlation function (normalized), (c) equal-time first-order temporal correlation function (normalized); (d) time-separated normalized first-order temporal correlation function; and (e) time-separated normalized second-order temporal correlation function $g^{(2)}(t_1, t_2) = 1 + g^{(1)}(t_1, t_2)g^{(1)}(t_2, t_1)$. Notice that only the cw-pumped case satisfies the Wiener-Khinchin theorem. Also, notice that the time-integrated $g^{(2)}$ decreases with increasing number of Schmidt modes, as was shown in [26].

TABLE I. Parameters used to generate Fig. 1.

Pump	σ_p	σ_c
Shorter	$2 \times 10^{12} \text{ s}^{-1}$	$2 \times 10^{12} \text{ s}^{-1}$
Longer	$1.5 \times 10^{12} \text{ s}^{-1}$	$2.4 \times 10^{12} \text{ s}^{-1}$
cw	n/a	$3 \times 10^{12} \text{ s}^{-1}$

temporal distributions in mode a . These vary between the three examples. Because the short pump yields a separable JSA, the BPT state is prepared in a single, broad, yet coherent spectral mode. This means that the frequencies have fixed relative phases, which leads to a spectrally coherent pulse of pseudothermal light. As the pump pulse length increases slightly, the pseudothermal pulse gets slightly longer. In turn,

the relative phases of the frequencies become slightly less fixed. In the limit of a cw pump, the BPT state is also continuous, but broadband, and the frequencies have *no* fixed relative phase relationship.

Figure 1(d) shows the normalized time-resolved first-order temporal correlation function in mode a . The light in all three examples has drastically different coherence times, despite having very similar spectra. For a separable JSA, $|g^{(1)}(t_1, t_2)| = 1$, but as the spectral correlations increase, the coherence time decreases. We note that since a pulse is not stationary, it does not have to satisfy the Wiener-Khinchin theorem.

Figure 1(e) shows the normalized time-resolved second-order temporal correlation function in mode a . For a separable JSA, $|g^{(2)}(t_1, t_2)| = 2$, but as the spectral correlations increase, the second-order coherence time decreases.

In summary, these examples show that by varying the spectrum of the pump field as well as the shape of the nonlinearity function of the material, it is possible to generate various states of light with the same spectrum but drastically different spectral and first- and second-order temporal coherence properties.

VII. DISCUSSION

Broadband states generated via nonlinear optical processes, such as SPDC or SFWM, have interesting coherence properties. We calculated their spectral and temporal correlation functions in terms of the Schmidt modes of the joint spectral amplitude and showed that these states can be thought of as pseudo-thermal states. We also showed that these states have tuneable spectral coherence—which can be tuned from perfectly spectrally coherent to completely incoherent. Regardless of the level of spectral coherence, these states can be decomposed into tensor products of states with geometric photon-number statistics.

High-gain SPDC can be extremely bright for a parametric process—up to hundreds of mW mean power [27,54,55]. This is close to the ~ 1 W achievable via nonparametric processes [18] and is bright enough to study interesting nonclassical phenomena such as the interplay between spectral and photon-number coherence. Furthermore, a source based on SPDC is extremely customizable. Beyond tuning the spectral coherence, the spectral shape can be customized using optical pulse shaping [45] or nonlinearity shaping methods [46–50].

Broadband spectrally incoherent light generated by nonlinear optics may also have application in studying the dynamics of photoinduced processes, such as the time scales and mechanisms underlying the initial step of photosynthesis in light-harvesting complexes. Some researchers have questioned whether dynamics initiated by sunlight excitation might be different from those detected in femtosecond laser experiments performed on light-harvesting complexes [56–60]. A broadband source with tuneable spectral coherence might help answer this question.

On the theory side, this work is relevant to the study of decompositions of thermal light into broadband coherent modes. Thermal light cannot be represented as a statistical mixture of single pulses [61], but one can construct mixtures of single pulses that yield the same first-order temporal correlation

function as thermal light [62]. In a one-dimensional waveguide, thermal light was shown to decompose into a statistical mixture of sets of coherent pulses [63]. Here, we show that partially spectrally coherent light can be decomposed into a tensor product of states prepared in spectral Schmidt modes, each with geometric photon-number statistics. Our work also connects with decompositions, into Schmidt-like modes, of correlation functions for partially spatially coherent thermal light [43].

Our formalism also provides a convenient tool for computing any observables of broadband pseudo-thermal states with partial spectral coherence—even for states created with completely different experimental methods, such as [22–25,64]. This is due to a well-known result in quantum information theory, the Stinespring dilation theorem [65], that states that any mixed state can be represented as a pure state in a higher-dimensional Hilbert space. Any broadband pseudo-thermal state can therefore be written as the reduced state of a hypothetical broadband twin-beam state.

The relationship between spectral, temporal, and photon-number coherence is of fundamental and practical interest. We hope that our analysis of broadband pseudo-thermal states generated via nonlinear optics provides a useful way of exploring it theoretically and experimentally.

ACKNOWLEDGMENTS

The authors thank J. Sipe for helpful discussions and J. Donohue for helpful comments on the manuscript. Research at Perimeter Institute is supported by the Government of Canada through Industry Canada and by the Province of Ontario through the Ministry of Research and Innovation. We acknowledge the support of the Natural Sciences and Engineering Research Council of Canada.

APPENDIX A: WRITING THE SINGLE-SCHMIDT-MODE THERMAL STATE IN GIBBS FORM

Consider the number operator of the k th Schmidt mode,

$$\hat{n}_k = \hat{A}_k^\dagger \hat{A}_k. \quad (\text{A1})$$

We can write its eigendecomposition and the resolution of the identity as follows:

$$\hat{n}_k |n_k\rangle = n_k |n_k\rangle, \quad \mathbb{I}_k = \sum_{n_k=0}^{\infty} |n_k\rangle \langle n_k|. \quad (\text{A2})$$

Using these expressions we calculate

$$\exp(-\alpha_k \hat{n}_k) = \exp(-\alpha_k \hat{A}_k^\dagger \hat{A}_k) = \sum_{n_k=0}^{\infty} e^{-\alpha_k n_k} |n_k\rangle \langle n_k|, \quad (\text{A3})$$

$$\text{Tr}[\exp(-\alpha_k \hat{n}_k)] = \sum_{n_k=0}^{\infty} e^{-\alpha_k n_k} = \frac{1}{1 - e^{-\alpha_k}}, \quad (\text{A4})$$

where in the last expression we use the geometric series sum. Consider now their ratio,

$$\frac{\exp(-\alpha_k \hat{A}_k^\dagger \hat{A}_k)}{\text{Tr}[\exp(-\alpha_k \hat{A}_k^\dagger \hat{A}_k)]} = \sum_{n_k=0}^{\infty} (1 - e^{-\alpha_k}) (e^{-\alpha_k})^{n_k} |n_k\rangle \langle n_k|. \quad (\text{A5})$$

We can compare this with Eqs. (11) and (13) and identify

$$e^{-\alpha_k} = \frac{\bar{n}_k}{\bar{n}_k + 1}, \quad (\text{A6})$$

easily verifying that $1 - e^{-\alpha_k} = 1/(\bar{n}_k + 1)$, completing the derivation of Eq. (15).

APPENDIX B: DERIVATION OF $G^{(1)}(\omega, \omega')$ AND $G^{(2)}(\omega, \omega')$

In this section, we derive the frequency second- and fourth-order moments of the a fields. For the second-order moment, we use the relations in Eqs. (5) to write

$$\langle \hat{a}^\dagger(\omega) \hat{a}(\omega') \rangle_\psi = \sum_{kl} \phi_k^*(\omega) \phi_l(\omega) \langle A_k^\dagger A_l \rangle_\psi. \quad (\text{B1})$$

We then use the linear Bogoliubov transformations in Eqs. (9) to write

$$\langle A_k^\dagger A_l \rangle_\psi = \langle \text{vac} | \hat{U}_{\text{SQ}}^\dagger \hat{A}_k \hat{U}_{\text{SQ}} \hat{U}_{\text{SQ}}^\dagger \hat{A}_l \hat{U}_{\text{SQ}} | \text{vac} \rangle \quad (\text{B2})$$

$$= \langle \text{vac} | (\cosh(r_k) \hat{A}_k + \sinh(r_k) \hat{B}_k^\dagger) (\cosh(r_l) \hat{A}_l + \sinh(r_l) \hat{B}_l^\dagger) | \text{vac} \rangle \quad (\text{B3})$$

$$= \sinh(r_k) \sinh(r_l) \langle \text{vac} | \hat{B}_k \hat{B}_l^\dagger | \text{vac} \rangle \quad (\text{B4})$$

$$= \sinh^2(r_k) \delta_{k,l}. \quad (\text{B5})$$

Plugging this result into Eq. (B1), we obtain

$$\langle \hat{a}^\dagger(\omega) \hat{a}(\omega') \rangle_\psi = \sum_k \sinh^2(r_k) \phi_k^*(\omega) \phi_k(\omega). \quad (\text{B6})$$

Now let us consider the fourth-order moment

$$\langle \hat{a}^\dagger(\omega) \hat{a}^\dagger(\omega') \hat{a}(\omega) \hat{a}(\omega') \rangle_\psi = \sum_{k,l,m,n} \phi_k(\omega) \phi_l(\omega') \phi_m^*(\omega) \phi_n^*(\omega') \langle \hat{A}_k^\dagger \hat{A}_l^\dagger \hat{A}_m \hat{A}_n \rangle_\psi. \quad (\text{B7})$$

As before, we look at the expectation value

$$\langle \hat{A}_k^\dagger \hat{A}_l^\dagger \hat{A}_m \hat{A}_n \rangle_\psi = \langle \text{vac} | \hat{U}_{\text{SQ}}^\dagger \hat{A}_k^\dagger \hat{U}_{\text{SQ}} \hat{U}_{\text{SQ}}^\dagger \hat{A}_l^\dagger \hat{U}_{\text{SQ}} \hat{U}_{\text{SQ}}^\dagger \hat{A}_m \hat{U}_{\text{SQ}} \hat{U}_{\text{SQ}}^\dagger \hat{A}_n \hat{U}_{\text{SQ}} | \text{vac} \rangle \quad (\text{B8})$$

and use the linear Bogoliubov transformation, and then expand to obtain

$$\langle \hat{A}_k^\dagger \hat{A}_l^\dagger \hat{A}_m \hat{A}_n \rangle_\psi = \sinh(r_k) \sinh(r_l) \sinh(r_m) \sinh(r_n) \langle \text{vac} | \hat{B}_k \hat{B}_l \hat{B}_m^\dagger \hat{B}_n^\dagger | \text{vac} \rangle \quad (\text{B9})$$

$$= \sinh^2(r_k) \sinh^2(r_l) \delta_{k,m} \delta_{l,n} + \sinh^2(r_k) \sinh^2(r_l) \delta_{k,n} \delta_{l,m}. \quad (\text{B10})$$

Plugging these results into Eq. (B7), we find

$$\langle \hat{a}^\dagger(\omega) \hat{a}^\dagger(\omega') \hat{a}(\omega) \hat{a}(\omega') \rangle_\psi = \langle \hat{a}^\dagger(\omega) \hat{a}(\omega) \rangle_\psi \langle \hat{a}^\dagger(\omega') \hat{a}(\omega') \rangle_\psi + \langle \hat{a}^\dagger(\omega) \hat{a}(\omega') \rangle_\psi \langle \hat{a}^\dagger(\omega') \hat{a}(\omega) \rangle_\psi, \quad (\text{B11})$$

where the terms in the right-hand side are given by Eq. (B6).

APPENDIX C: DERIVATION OF $G^{(1)}(t, t')$ AND $G^{(2)}(t, t')$

The first-order temporal correlation function for mode a is given by [41]

$$G_a^{(1)}(t_1, t_2) = \langle \hat{E}_a^{(-)}(t_1) \hat{E}_a^{(+)}(t_2) \rangle_\psi, \quad (\text{C1})$$

where $\hat{E}_a^{(\pm)}(t)$ are the usual positive and/or negative frequency components of the electric field operator in mode a . To simplify calculations, we follow Christ *et al.* [26] and replace the electric field operators by photon-number creation and destruction operators [$\hat{E}_a^{(+)} \propto \hat{a}(t)$]. This is valid when the spectra of the beams are not too broad. We thus have

$$G_a^{(1)}(t_1, t_2) = \langle \hat{a}^\dagger(t_1) \hat{a}(t_2) \rangle_\psi. \quad (\text{C2})$$

We now use the relation

$$\hat{a}(t) = \frac{1}{\sqrt{2\pi}} \int d\omega \hat{a}(\omega) e^{i\omega t} = \frac{1}{\sqrt{2\pi}} \int d\omega \left(\sum_k \phi_k^*(\omega) \hat{A}_k \right) e^{i\omega t} \equiv \sum_k \hat{A}_k \tilde{\phi}_k^*(t), \quad (\text{C3})$$

where $\tilde{\phi}_k(t) = \frac{1}{\sqrt{2\pi}} \int d\omega \phi_k(\omega) e^{-i\omega t}$. This gives

$$G_a^{(1)}(t_1, t_2) = \left\langle \left(\sum_k \hat{A}_k^\dagger \tilde{\phi}_k(t_1) \right) \left(\sum_l \hat{A}_l \tilde{\phi}_l^*(t_2) \right) \right\rangle_\psi \quad (\text{C4})$$

$$= \sum_{k,l} \tilde{\phi}_k(t_1) \tilde{\phi}_l^*(t_2) \langle \hat{A}_k^\dagger \hat{A}_l \rangle_\psi. \quad (\text{C5})$$

Using the result in Eq. (B5), we have

$$G_a^{(1)}(t_1, t_2) = \sum_{k,l} \tilde{\phi}_k(t_1) \tilde{\phi}_l^*(t_2) \sinh^2(r_k) \delta_{k,l} \quad (\text{C6})$$

$$= \sum_k \tilde{\phi}_k(t_1) \tilde{\phi}_k^*(t_2) \sinh^2(r_k) \quad (\text{C7})$$

$$= \sum_k \tilde{\phi}_k(t_1) \tilde{\phi}_k^*(t_2) \bar{n}_k, \quad (\text{C8})$$

where \bar{n}_k is defined in Eq. (14).

The second-order temporal correlation function for mode a is given by [41]

$$G_a^{(2)}(t_1, t_2) = \langle \hat{a}^\dagger(t_1) \hat{a}^\dagger(t_2) \hat{a}(t_1) \hat{a}(t_2) \rangle_\psi. \quad (\text{C9})$$

Using a similar procedure as above, we can write this as

$$G_a^{(2)}(t_1, t_2) = \sum_{k,l,m,n} \tilde{\phi}_k(t_1) \tilde{\phi}_l(t_2) \tilde{\phi}_m^*(t_1) \tilde{\phi}_n^*(t_2) \langle \hat{A}_k^\dagger \hat{A}_l^\dagger \hat{A}_m \hat{A}_n \rangle_\psi. \quad (\text{C10})$$

Using Eq. (B10), we have

$$G_a^{(2)}(t_1, t_2) = \sum_{k,l,m,n} \tilde{\phi}_k(t_1) \tilde{\phi}_l(t_2) \tilde{\phi}_m^*(t_1) \tilde{\phi}_n^*(t_2) [\sinh^2(r_k) \sinh^2(r_l) \delta_{k,m} \delta_{l,n} + \sinh^2(r_k) \sinh^2(r_l) \delta_{k,n} \delta_{l,m}] \quad (\text{C11})$$

$$= \sum_k \tilde{\phi}_k(t_1) \tilde{\phi}_k^*(t_1) \bar{n}_k \sum_l \tilde{\phi}_l(t_2) \tilde{\phi}_l^*(t_2) \bar{n}_l + \sum_k \tilde{\phi}_k(t_1) \tilde{\phi}_k^*(t_2) \bar{n}_k \sum_l \tilde{\phi}_l(t_2) \tilde{\phi}_l^*(t_1) \bar{n}_l, \quad (\text{C12})$$

where \bar{n}_k is defined in Eq. (14). Comparing with the result in Eq. (C8), we find that

$$G_a^{(2)}(t_1, t_2) = G_a^{(1)}(t_1, t_1) G_a^{(1)}(t_2, t_2) + G_a^{(1)}(t_1, t_2) G_a^{(1)}(t_2, t_1). \quad (\text{C13})$$

-
- [1] A. Zavatta, V. Parigi, and M. Bellini, *Phys. Rev. A* **75**, 052106 (2007).
- [2] R. Tahira, M. Ikram, H. Nha, and M. S. Zubairy, *Phys. Rev. A* **79**, 023816 (2009).
- [3] M. S. Kim, *Fortschr. Phys.* **50**, 652 (2002).
- [4] C. Weedbrook, S. Pirandola, and T. C. Ralph, *Phys. Rev. A* **86**, 022318 (2012).
- [5] G. Harder, D. Mogilevtsev, N. Korolkova, and C. Silberhorn, *Phys. Rev. Lett.* **113**, 070403 (2014).
- [6] A. Gatti, E. Brambilla, M. Bache, and L. A. Lugiato, *Phys. Rev. A* **70**, 013802 (2004).
- [7] A. Gatti, E. Brambilla, M. Bache, and L. A. Lugiato, *Phys. Rev. Lett.* **93**, 093602 (2004).
- [8] A. Valencia, G. Scarcelli, M. D'Angelo, and Y. Shih, *Phys. Rev. Lett.* **94**, 063601 (2005).
- [9] D. Zhang, Y.-H. Zhai, L.-A. Wu, and X.-H. Chen, *Opt. Lett.* **30**, 2354 (2005).
- [10] I. N. Agafonov, M. V. Chekhova, T. S. Iskhakov, and L.-A. Wu, *J. Mod. Opt.* **56**, 422 (2009).
- [11] X.-H. Chen, I. N. Agafonov, K.-H. Luo, Q. Liu, R. Xian, M. V. Chekhova, and L.-A. Wu, *Opt. Lett.* **35**, 1166 (2010).
- [12] P.-A. Moreau, E. Toninelli, T. Gregory, and M. J. Padgett, *Laser Photon. Rev.* **12**, 1700143 (2017).
- [13] D.-Z. Cao, G.-J. Ge, and K. Wang, *Appl. Phys. Lett.* **97**, 051105 (2010).
- [14] G. Lu, X. Jun, Z. Shu-Heng, W. Wei, and W. Kai-Ge, *Chin. Phys. Lett.* **25**, 1277 (2008).
- [15] J. Sprigg, T. Peng, and Y. Shih, *Sci. Rep.* **6**, 38077 (2016).
- [16] H. Lajunen, V. Torres-Company, J. Lancis, and A. T. Friberg, *J. Opt. Soc. Am. A* **26**, 1049 (2009).
- [17] C. López-Mariscal and J. C. Gutiérrez-Vega, *J. Opt.* **12**, 075702 (2010).
- [18] D. J. Ulness, *J. Chem. Phys.* **107**, 8111 (2003).
- [19] D. B. Turner, P. C. Arpin, S. D. McClure, D. J. Ulness, and G. D. Scholes, *Nat. Commun.* **4**, 2298 (2013).
- [20] G. Li, T. Zhang, Y. Li, and J. Wang, in *Quantum Optics and Applications in Computing and Communications II* (International Society for Optics and Photonics, SPIE, Bellingham, WA, 2005), Vol. 5631, pp. 134–143.
- [21] T. Iskhakov, A. Allevi, D. Kalashnikov, V. Sala, M. Takeuchi, M. Bondani, and M. Chekhova, *Eur. Phys. J.: Spec. Top.* **199**, 127 (2011).

- [22] A. Jechow, M. Seefeldt, H. Kurzke, A. Heuer, and R. Menzel, *Nat. Photon.* **7**, 973 (2013).
- [23] S. Hartmann, F. Friedrich, A. Molitor, M. Reichert, W. Elsäß, and W. R., *New J. Phys.* **17**, 043039 (2015).
- [24] J. Mika, L. Podhora, L. Lachman, P. Obšil, J. Hloušek, M. Ježek, R. Filip, and L. Slodička, *New J. Phys.* **20**, 093002 (2018).
- [25] T. F. Schulz, P. P. Aung, L. Weisel, K. Cosert, M. W. Gealy, and D. J. Ulness, *J. Opt. Soc. Am. B* **22**, 1052 (2005).
- [26] A. Christ, K. Laiho, A. Eckstein, K. N. Cassemiro, and C. Silberhorn, *New J. Phys.* **13**, 033027 (2011).
- [27] K. Y. Spasibko, D. A. Kopylov, V. L. Krutyanskiy, T. V. Murzina, G. Leuchs, and M. V. Chekhova, *Phys. Rev. Lett.* **119**, 223603 (2017).
- [28] Z. Vernon, M. Menotti, C. Tison, J. Steidle, M. Fanto, P. Thomas, S. Preble, A. Smith, P. Alsing, M. Liscidini *et al.*, *Opt. Lett.* **42**, 3638 (2017).
- [29] B. Blauensteiner, I. Herbauts, S. Bettelli, A. Poppe, and H. Hübel, *Phys. Rev. A* **79**, 063846 (2009).
- [30] A. Eckstein, A. Christ, P. J. Mosley, and C. Silberhorn, *Phys. Rev. Lett.* **106**, 013603 (2011).
- [31] K. Y. Spasibko, T. S. Iskhakov, and M. V. Chekhova, *Opt. Express* **20**, 7507 (2012).
- [32] K. Wakui, Y. Eto, H. Benichi, S. Izumi, T. Yanagida, K. Ema, T. Numata, D. Fukuda, M. Takeoka, and M. Sasaki, *Sci. Rep.* **4**, 4535 (2014).
- [33] J. Huang and P. Kumar, *Phys. Rev. A* **40**, 1670 (1989).
- [34] V. Ansari, J. M. Donohue, M. Allgaier, L. Sansoni, B. Brecht, J. Roslund, N. Treps, G. Harder, and C. Silberhorn, *Phys. Rev. Lett.* **120**, 213601 (2018).
- [35] W. P. Grice and I. A. Walmsley, *Phys. Rev. A* **56**, 1627 (1997).
- [36] N. Quesada and J. E. Sipe, *Phys. Rev. A* **90**, 063840 (2014).
- [37] N. Quesada and J. E. Sipe, *Phys. Rev. Lett.* **114**, 093903 (2015).
- [38] N. Quesada, G. Triginer, M. D. Vidrighin, and J. E. Sipe (unpublished).
- [39] F. G. Mehler, *Journal für die reine und angewandte Mathematik* **66**, 161 (1866).
- [40] F. Graffitti, J. Kelly-Massicotte, A. Fedrizzi, and A. M. Brańczyk, *Phys. Rev. A* **98**, 053811 (2018).
- [41] R. J. Glauber, *Phys. Rev.* **130**, 2529 (1963).
- [42] L. Mandel and E. Wolf, *Optical Coherence and Quantum Optics* (Cambridge University Press, Cambridge, UK, 1995).
- [43] I. B. Bobrov, S. S. Straupe, E. V. Kovlakov, and S. P. Kulik, *New J. Phys.* **15**, 073016 (2013).
- [44] R. Loudon, *The Quantum Theory of Light* (Clarendon Press, Oxford, 1983).
- [45] A. M. Weiner, *Opt. Commun.* **284**, 3669 (2011).
- [46] A. M. Brańczyk, A. Fedrizzi, T. M. Stace, T. C. Ralph, and A. G. White, *Opt. Express* **19**, 55 (2011).
- [47] P. B. Dixon, J. H. Shapiro, and F. N. C. Wong, *Opt. Express* **21**, 5879 (2013).
- [48] A. Dosseva, L. Cincio, and A. M. Brańczyk, *Phys. Rev. A* **93**, 013801 (2016).
- [49] J.-L. Tambasco, A. Boes, L. G. Helt, M. J. Steel, and A. Mitchell, *Opt. Express* **24**, 19616 (2016).
- [50] F. Graffitti, D. Kundys, D. T. Reid, A. M. Brańczyk, and A. Fedrizzi, *Quantum Sci. Technol.* **2**, 035001 (2017).
- [51] V. Ansari, J. M. Donohue, B. Brecht, and C. Silberhorn, *Optica* **5**, 534 (2018).
- [52] O. Kuzucu, F. N. C. Wong, S. Kurimura, and S. Tovstonog, *Phys. Rev. Lett.* **101**, 153602 (2008).
- [53] N. Quesada and A. M. Brańczyk, *Phys. Rev. A* **98**, 043813 (2018).
- [54] A. M. Pérez, K. Y. Spasibko, P. R. Sharapova, O. V. Tikhonova, G. Leuchs, and M. V. Chekhova, *Nat. Commun.* **6**, 7707 (2015).
- [55] K. Y. Spasibko, D. A. Kopylov, T. V. Murzina, G. Leuchs, and M. V. Chekhova, *Opt. Lett.* **41**, 2827 (2016).
- [56] X.-P. Jiang and P. Brumer, *J. Chem. Phys.* **94**, 5833 (1991).
- [57] T. Maňal and L. Valkunas, *New J. Phys.* **12**, 065044 (2010).
- [58] K. Hoki and P. Brumer, *Procedia Chem.* **3**, 122 (2011).
- [59] P. Brumer and M. Shapiro, *Proc. Natl. Acad. Sci. USA* **109**, 19575 (2012).
- [60] I. Kassal, J. Yuen-Zhou, and S. Rahimi-Keshari, *J. Phys. Chem. Lett.* **4**, 362 (2013).
- [61] A. Chenu, A. M. Brańczyk, G. D. Scholes, and J. E. Sipe, *Phys. Rev. Lett.* **114**, 213601 (2015).
- [62] A. Chenu, A. M. Brańczyk, and J. E. Sipe, *Phys. Rev. A* **91**, 063813 (2015).
- [63] A. M. Brańczyk, A. Chenu, and J. E. Sipe, *J. Opt. Soc. Am. B* **34**, 1536 (2017).
- [64] Y. Zhou, X. Zhang, Z. Wang, F. Zhang, H. Chen, H. Zheng, J. Liu, F.-I. Li, and Z. Xu, *Optics Communications* **437**, 330 (2019).
- [65] W. F. Stinespring, *Proc. Am. Math. Soc.* **6**, 211 (1955).