Angular momentum plays a pivotal role in physics. In settings with a continuous rotational symmetry, Noether’s theorem \[1, 2\] dictates that this quantity is conserved, an aspect that governs both the macroscopic and microscopic behaviors of a multitude of physical systems. These could range from the dynamics of spiral galaxies \[3\], pulsars \[4\] and neutron stars to the intriguing properties of quantum vortices in superfluids \[5\] and superconductors. That fact that the electromagnetic field carries spin and/or orbital angular momentum (OAM) was recognized early on with the advent of Maxwell’s electrodynamics \[6, 7\]. Yet, it is only recently that the angular momentum of light was recognized as a new degree of freedom through which a wealth of opportunities could open up within the discipline of optics and photonics \[8–11\]. In this respect, the orbital angular momentum of a light beam propagating in a nonlinear cylindrical multimode optical waveguide \[12\] can be thermalized in a nonlinear multimode nonlinear bosonic systems that display additional conservation laws.

We show that the orbital angular momentum (OAM) of a light field can be thermalized in a nonlinear cylindrical multimode optical waveguide. We find, that upon thermal equilibrium, the maximization of the optical entropy leads to a generalized Rayleigh-Jeans distribution that governs the power modal occupancies with respect to the discrete OAM charge numbers. This distribution is characterized by a temperature that is by nature different from that associated with the longitudinal electromagnetic momentum flow of the optical field. Counterintuitively and in contrast to previous results, we demonstrate that even under positive temperatures, the ground state of the fiber is not always the most populated in terms of power. Instead, because of OAM, the thermalization processes may favor higher order modes. A new equation of state is derived along with an extended Euler equation — resulting from the extensivity of the entropy itself. By monitoring the nonlinear interaction between two multimoded optical wavefronts with opposite spins, we show that the exchange of angular momentum is dictated by the difference in OAM temperatures, in full accord with the second law of thermodynamics. The theoretical analysis presented here is corroborated by numerical simulations that take into account the complex nonlinear dynamics of hundreds of modes. Our results may pave the way towards high power optical sources with controllable orbital angular momenta, and at a more fundamental level, could shed light on the physics of other complex multimoded nonlinear bosonic systems that display additional conservation laws.
no longer separable with respect to the conservation laws of longitudinal momentum and OAM. Even more importantly, upon attaining thermal equilibrium, the OAM is now governed by its own optical temperature $T_L$ — a temperature that has nothing to do with that assigned to its energy counterpart $T_U$ (Fig. 1). In this respect, a generalized Rayleigh-Jeans distribution can be derived, that even under positive temperature ($T_L$) conditions, does not always favor the ground state. New equations of state are obtained that relate the extensive variables with their corresponding generalized forces. Finally, based on entropic principles, we show that the OAM temperature is a true thermodynamic quantity in the sense that it dictates the OAM exchange between two optical beams of opposite spin when traversing a nonlinear medium. The predicted theoretical results are in excellent agreement with numerical simulations.

To this end, let us consider a weakly guiding, nonlinear cylindrical waveguide having a normalized refractive index distribution $V(r)$, where $r$ represents the cylindrical radial coordinate. In Kerr nonlinear media, the optical field $u(x, y, z)$ evolution along the $z$ direction is governed by the normalized nonlinear Schrödinger equation:

$$i \frac{\partial u}{\partial z} + \left( \frac{\partial^2 u}{\partial x^2} + \frac{\partial^2 u}{\partial y^2} \right) + V(x, y)u + |u|^2 u = 0. \quad (1)$$

In this system, the field orbital angular momentum is given by $L = \int_{-\infty}^{\infty} \hat{r} \times \hat{p} \, dxdy$, where $\hat{r} = \frac{1}{2}(u \nabla_u u^* - u^* \nabla_u u)$ is the transverse momentum density, and $\nabla_u = \hat{x} \frac{\partial}{\partial x} + \hat{y} \frac{\partial}{\partial y}$. Given that $L \parallel \hat{z}$, from this point on, $L$ will be treated as a scalar quantity. The conservation laws $C$ associated with Eq. (1) are known to satisfy $\{C, H\}_{u, \pi} = \int_{-\infty}^{\infty} \left( \frac{\delta C}{\delta \pi} - \frac{\delta H}{\delta u} \right) \, dxdy = 0$, where $C = \rho(\pi_1 + \pi_2 + \frac{i}{2} u^2)$ is the canonical field momentum, $\{\cdot\}$ denotes a Poisson bracket and $\delta$ represent functional derivatives. In addition to the total power $P = \int_{-\infty}^{\infty} |u|^2 dxdy$ that remains constant throughout propagation, one can show that in a fully cylindrical structure $[V(x, y) = V(r)]$, the field orbital angular momentum $L$ is also an invariant in this nonlinear system (see Supplemental Material [35]).

In general, an optical field propagating in the aforementioned cylindrical waveguide (supporting $M$ modes) can be represented as a superposition of its underlying bound states $\psi_n(r, \phi, z) = R_n(r)e^{i(\pi r/R + \alpha_n \phi)}$, i.e.,

$$u = \sum_{n=1}^{M} c_n \psi_n.$$  

Here, $c_n$ stands for the propagation constant or eigenvalue of the eigenmode $\psi_n$, $l_n \in \{0, \pm 1, \pm 2, \ldots\}$ denotes its discrete OAM charge, $R_n(r)$ is the radial part of the eigenfunction while the complex number $c_n(z)$ describes the power occupancy ($|c_n|^2$) of this state during propagation. In this representation the power and OAM invariants can be written as $P = \sum_{n=1}^{M} |c_n|^2$ and $L = \sum_{n=1}^{M} l_n |c_n|^2$ [34]. On the other hand, the Hamiltonian constant of the motion $H = \int \mathcal{H} \, dxdy$ involves both a linear ($-U$) and a nonlinear component ($\mathcal{H}_{NL}$), that is $H = -U + \mathcal{H}_{NL}$. Given that the system is operating in the weakly nonlinear regime, one finds that the Hamiltonian is dominated by its linear contribution, and as a result the quantity $U = -\sum_{n=1}^{M} c_n |c_n|^2$ now assumes the role of the third invariant. Interestingly, this “internal optical energy” $U$ is associated with the Minkowski longitudinal electromagnetic momentum flowing in this guiding arrangement [35].

The existence of these three invariants ($P, U, L$) in this weakly nonlinear heavily multimode optical fiber now allows one to deploy principles from statistical mechanics. What facilitates this approach is the absence of required conservation laws ($M - 3$), an aspect that leads to chaotic behavior in the evolution of the modal occupancies $|c_n|^2$ and therefore allows the system to ergodically explore its phase space. In other words, all possible microstates ($c_1, c_2, \ldots, c_M$) that lie on the manifolds of constant angular momentum ($L$), power ($P$) and energy ($U$) are accessed with equal probability because of nonlinearity. To derive the state functions that relate the thermodynamically extensive variables ($P, U, L, M$) to the entropy $S$, we adopt a grand canonical description in a phase space constructed by $J_n \equiv |c_n|^2$ ($n = 1, 2, \ldots, M$) [39]. In this grand canonical frame, the local state of the system is described by a normalized probability density distribution $\rho(J_1, \ldots, J_M)$ where \[ \int_0^\infty \rho(J_1, \ldots, J_M) \prod_{n=1}^{M} dJ_n = 1. \] Once the system attains thermal equilibrium, the average values of $P, L$ and $U$ serve as the invariants of this grand canonical system. In turn, the probability density $\rho$ can be used to construct the Gibbs entropy

$$S = -\int_0^\infty \rho(J_1, \ldots, J_M) \ln \rho(J_1, \ldots, J_M) \prod_{n=1}^{M} dJ_n, \quad (2)$$

which can be maximized by means of Lagrange multipli-
\[
\rho(J_1, \ldots, J_M) = \frac{e^{-\alpha P(J_1, \ldots, J_M) - \beta U(J_1, \ldots, J_M) - \gamma L(J_1, \ldots, J_M)}}{Z},
\]

where \(\alpha, \beta\) and \(\gamma\) represent constants associated with the three invariants. In this case, the generalized grand partition function is given by [36]

\[
Z = \int_0^\infty e^{-\alpha \sum_{n=1}^M J_n + \beta \sum_{n=1}^M \varepsilon_n J_n - \gamma \sum_{n=1}^M l_n J_n} \prod_{n=1}^M dJ_n
\]  

(4a) \[
= \prod_{n=1}^M \frac{1}{-\alpha + \beta \varepsilon_n - \gamma l_n}.
\]  

(4b)

At equilibrium, the mean value \(\langle J_m \rangle = \langle |c_m|^2 \rangle = \int_0^\infty \rho(J_1, \ldots, J_M)J_m \prod_{n=1}^M dJ_n\) can be directly obtained from the generalized partition function \(Z\) through the relations: \(\langle J_m\rangle = \frac{1}{\beta} \frac{\partial \ln Z}{\partial \varepsilon_m} = -\frac{1}{\gamma} \frac{\partial \ln Z}{\partial l_m}\) [36]. From here, one can find that \(\langle J_m\rangle = \frac{-\beta}{\alpha \beta + \gamma l_m}\). We then introduce two intensive quantities by defining \(\theta = -1/T_U\) and \(\gamma = -1/T_L\), i.e., an energy temperature \(T_U\) and an OAM temperature \(T_L\), that are conjugate to the extensive variables \(U\) and \(L\), respectively. Here, \(\alpha\) is another intensive quantity that is conjugate to \(P\), which serves as the chemical potential of the system. As a result, upon thermalization, the average power occupancy of each mode is found to obey a generalized Rayleigh-Jeans distribution

\[
\langle |c_n|^2 \rangle = \frac{-1}{\alpha + \beta \varepsilon_n - \gamma l_n}.
\]

(5)

The generalized Rayleigh-Jeans distribution in Eq. (5) resulting from a generalized Gibbs ensemble [40] [41] is central in this work. It implies that not only the internal energy \(U\) can be thermalized (via eigenvalue \(\varepsilon_n\)), but also the OAM since it explicitly involves the topological charge of the mode \(l_n\) — an aspect that has a profound effect on the modal power distributions (Fig. 1). Because of OAM, the possibility exists that the ground state (the lowest order mode) is no longer the most populated in terms of power. Even more importantly, \(T_L\) is a true thermodynamic quantity that governs the flow of OAM between two optical beams. From this generalized distribution, the following global equation of state can be derived [30]

\[
-\alpha P + \frac{U}{T_U} + \frac{L}{T_L} = M,
\]

(6)

which relates the three intensive variables \(\alpha, T_U, \text{ and } T_L\) to the four extensive quantities \(P, U, L\text{ and } M\).

Based on these premises, one can formally show that the entropy \(S\) of this optical multimode system can be directly expressed in terms of the modal occupancies via \(S = \sum_{n=1}^M \ln(\langle |c_n|^2 \rangle)\) [30] [39]. In addition, the fundamental equation of thermodynamics demands that \(S = \)

\[
\mathcal{S}(P, U, L, M).
\]

Hence, two temperatures can be entropically defined through \(T_{\alpha} \equiv \partial S/\partial P,\) and \(T_L \equiv \partial S/\partial L\). Similarly, a generalized chemical potential \(\alpha\) and an optical thermodynamic pressure \(\tilde{\alpha}\) can also be introduced using \(\alpha \equiv -\partial S/\partial P\) and \(\tilde{\alpha} \equiv -\partial S/\partial M\) [30]. From here, a corresponding Euler equation can be obtained which is a direct manifestation of the extensivity of the entropy with respect to \((P, U, L, M)\).

\[
S = -\alpha P + \frac{U}{T_U} + \frac{L}{T_L} + \tilde{\alpha} M.
\]

(7)

In what follows, we corroborate our theoretical formalism by performing a series of numerical simulations in nonlinear parabolic and step-index optical fibers. In this respect, the results presented in Eqs. (5) and (6) will be employed to predict the OAM thermalization once equilibrium conditions are attained. As a first example, let us consider a parabolic silica fiber having \(M = 120\) modes, conveying in total 100 kW of power at a wavelength of 1064 nm [36]. For this case, 62 modes of this fiber are
evenly excited [Fig. 2(a)]. In normalized units, these launching conditions correspond to \( P = 100, U = -1468 \) and \( L = -140 \). For this scenario, our theory predicts that at thermal equilibrium, the propagating optical wavefront is expected to settle into an OAM temperature \( T_L = 14.5 \), an energy temperature \( T_U = 15.2 \) and a generalized chemical potential \( \alpha = -2.26 \). To verify these predictions, we numerically solved Eq. (1) under the same initial conditions. For each run, we allowed the phases of the input \( e_u \) coefficients to statistically vary in order to construct the statistical ensembles. Figures 2(b) and (c) show that an excellent agreement exists between the theoretically anticipated results and the numerical computations after OAM thermal equilibrium is reached at \( \sim 1 \) m of propagation. In this case, the modal occupancies \( \langle |c_n|^2 \rangle \) are displayed on a two-dimensional triangular map that is specified by the accessible normalized propagation constants \( \varepsilon_n \) and OAM numbers \( l_n \) associated with the optical modes. For these initial conditions, the energy temperature \( T_U \) is positive and therefore the fundamental mode \( \text{LP}_{01} \) is the one mostly populated. Meanwhile, the OAM temperature \( T_L \) so happens to be negative, in which case the optical power resides mostly in the right-handed rotating modes with positive OAM charges \( (l_n > 0) \). It is important to note that, in all cases, the initial conditions uniquely determine the equilibrium intensive variables \( (T_{L}, T_{U}, \alpha) \) of this system. As Fig. 2(d) shows, both positive and negative \( T_L \) temperatures are possible, favoring left-handed and right-handed OAM groups of modes, respectively. Given that the OAM numbers in a cylindrical structure always come in pairs \( (\pm l) \), if the input OAM is zero, the system will relax into an infinite \( T_L \) temperature [Fig. 2(d)], which leads to OAM equipartition and hence the Rayleigh-Jeans distribution of Eq. (5) assumes a standard form [22].

Our theory is universal in the sense that it can be deployed to predict OAM thermalization in any cylindrical multimode nonlinear structure. Figure 3(a) shows the results of OAM thermalization in a step-index fiber having \( M = 234 \) modes. For the step-index case, the \( \varepsilon - l \) map now assumes a parabolic-like shape. For the excitation conditions used in this simulation, \( P = 3200, U = -75420 \) and the \( L = 8776 \), for which our theory predicts \( T_U = 104, T_L = -204 \) and \( \alpha = -0.04 \). As Figs. 3(a) and (b) show, there is an excellent agreement between the numerical simulations and the theoretically anticipated results. Counterevintuitively, in this case, while the energy temperature \( T_U \) is positive, the fundamental mode \( \text{LP}_{01} \) is no longer the most populated. Instead, the \( 8^\text{th} \) mode \( \text{LP}_{+3,1} \) \( (l = +3) \) carries most of the power. This anomaly in the distribution results from the interplay between the more convolved \( \varepsilon - l \) map that corresponds to a step-index fiber and the non-separability of \( U \) and \( L \) in the generalized Rayleigh-Jeans distribution. To intuitively understand this behavior, one can assign effective eigen-energies in this map according to \( \varepsilon_{n,\text{eff}} = \varepsilon_n - l_n T_U / T_L \) that angularly \( \theta = \arctan(T_U / T_L) \) project this data into a standard Rayleigh-Jeans curve. This is illustrated in Fig. 3(b) where the \( 8^\text{th} \) mode is the first in line to assume the role of an “effective ground state”. In this respect, the ratio of the two temperatures \( T_U / T_L \) can be used to select the highest populated mode in this system.

Next, we show that the OAM temperature \( T_L \) is an actual thermodynamic force that governs the direction of OAM exchange between two subsystems, just like its energy counterpart. Figure 4(a) illustrates a scenario where two circularly polarized beams are launched in the same nonlinear parabolic silica fiber. In this case, the total electric field with components \( |R \rangle \) and \( |L \rangle \) can be written in terms of slowly varying envelopes \( (u,v) \) as \( \tilde{E} = u |R \rangle + v |L \rangle \), whose evolution equations can be found in [36]. In this arrangement, one can formally identify the following conservation laws [36]: \( P_u = \text{const.} \), \( P_v = \text{const.} \), \( L_{\text{total}} = L_u + L_v = \text{const.} \), \( U_{\text{total}} = U_u + U_v = \text{const.} \). In the example that follows the right-hand circularly (RHC) polarized light (red) is hotter in energy temperature \( T_U \), while the left-handed (LHC, blue) is hotter in terms of its OAM. Moreover, at the input \( L_u(0) = -74, L_v(0) = 4 \) \( (L_{\text{total}} = -70) \), \( P_u = 50, P_v = 40, U_u(0) = -800, U_v(0) = -774 \) \( (U_{\text{total}} = -1574) \). While the two circular polarizations do not exchange power \( P \), they can exchange energy \( U \) and OAM \( L \) according to the second law of thermodynamics

\[
\frac{dS_T}{T} = \left( \frac{1}{T_{U,u}} - \frac{1}{T_{U,v}} \right) dU_u + \left( \frac{1}{T_{L,u}} - \frac{1}{T_{L,v}} \right) dL_u \geq 0 \tag{8}
\]

given that \( dU_u = -dU_v \) and \( dL_u = -dL_v \). As a result, during co-propagation, both energy and OAM are ex-
FIG. 4. (a) When a RHC beam conveying significant negative OAM (red) and a LHC beam having almost 0 OAM (blue) are launched into the same fiber, they eventually settle into the same energy temperature $T_U \approx 4.9$ and OAM temperature $T_L \approx 8.0$. During propagation, the longitudinal momentum $U$ is transferred from the RHC to LHC polarization (b), whereas the OAM flows in the opposite direction given that initially $T_{U,u} > T_{U,v} > 0$ and $T_{L,v} < 0 < T_{L,u}$ (negative temperatures are hotter than positive). (d) Meanwhile, the total optical entropy monotonically increases until it stabilizes as demanded by the second law of thermodynamics. (e) The final state can be uniquely predicted from the initial conditions $P_u, P_v, U_{total}$ and $L_{total}$, through the intersection of the $T_U$, $T_L$ manifolds in the $(U, L)$ plane [36].

In conclusion, we have shown that the OAM of a light beam propagating in a nonlinear cylindrical waveguide structure can be thermalized. A generalized Rayleigh-Jeans distribution has been derived that explicitly depends on the OAM charge. The interplay between energy and OAM can lead to an anomaly in the power distributions among modes, thus allowing higher-order modes to be favored. The exchange of OAM between two subsystems formally indicates that the OAM temperature acts like a true thermodynamic force.

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