Thermal Bound Entanglement in Macroscopic Systems and Area Law

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Does bound entanglement naturally appear in quantum many-body systems? We address this question by showing the existence of bound-entangled thermal states for harmonic oscillator systems consisting of an arbitrary number of particles. By explicit calculations of the negativity for different partitions, we find a range of temperatures for which no entanglement can be distilled by means of local operations, despite the system being globally entangled. We offer an interpretation of this result in terms of entanglement-area laws, typical of these systems. Finally, we discuss generalizations of this result to other systems, including spin chains.

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Introduction.—Many quantum information tasks are based on several distant observers sharing entangled pure states. However, in practical situations, environmental noise is always present and then we unavoidably deal with partially entangled mixed states. The first attempt to overcome this degradation led to the idea of entanglement distillation [1,2], a sequence of local operations assisted by classical communication (LOCC) capable of extracting entangled pure states from a large set of entangled mixed states. However, in some cases, the noise degradation happens to be irreparable: given a system composed by \( n \) parties there exist entangled states that need pure-state entanglement in order to be generated, while no pure-state entanglement can be recovered back by the \( n \) parties via LOCC. These states are known as bound-entangled states [3] and they represent arguably the most striking manifestation of irreversibility in the context of quantum information science.

Although there exist several examples of bound-entangled states [4], to our knowledge they have mainly been inspired by mathematical intuitions and simple recipes to construct these states are still lacking. This poses doubts about whether bound-entangled states, despite their remarkable properties, are mainly a mathematical construction. Therefore, up until now a question has remained open: do bound-entangled states appear inherently in nature? In particular, do these states emerge in standard quantum many-body systems? By standard systems we specifically mean local interacting systems in a macroscopic thermal state characterized by only a few parameters. A first attempt to address this question has been given in Ref. [5], where bound entanglement was detected in the thermal state of spin systems of up to 9 qubits.

In this work we show that bound-entangled states naturally appear as thermal states of many-body systems composed by a macroscopic number of particles. The fundamental intuition behind our results comes from the entanglement-area law, a property satisfied by many condensed matter systems according to which the entanglement between two regions scales as the surface separating them [6]. By relating bound entanglement to area laws, we provide the first construction of bound-entangled states with a clear physical inspiration. In this sense, the route we take goes in the reverse direction with respect to the one usually pursued by many recent works [7]: we borrow concepts from the condensed matter field to get new insight on quantum information science.

Bound entanglement and area law.—Consider a quantum system of \( n \) particles described by a local Hamiltonian. For the sake of simplicity, we will restrict our analysis to translational invariant one-dimensional systems of \( n \) particles. If the system obeys an entanglement-area law, the ground-state entanglement for a bipartite splitting of the system into two groups, say \( A \) and \( B \), scales as the number of connections between them. This behavior has been observed in many noncritical quantum systems, while logarithmic corrections may appear in the critical case [6]. Consider now two different partitions of a system, one in which a contiguous half of the particles belongs to \( A \) and the other half to \( B \) (we will refer to such kind of partition as half-half), and another partition in which the particles with even label belong to \( A \) and the others to \( B \) (even-odd partition). Because of the area law, the entanglement will saturate for sufficiently large \( n \) for the half-half partition, while it will increase with \( n \) for the even-odd partition. In this configuration, it is reasonable to expect that, by increasing the temperature, the entanglement in the even-odd partition is more robust to thermal noise than in the half-half partition, and that this behavior is preserved for large systems.

A fundamental result in the understanding of bound entanglement has been to recognize that all distillable entangled states have a nonpositive partial transposition (NPPT) [8]. Thus if one finds a nonseparable state with positive partial transposition (PPT) it must be bound entangled [3]. Now, denote by \( T_{\text{dist}}^{\text{h-h}} \) (\( T_{\text{dist}}^{\text{e-o}} \)) the threshold temperatures at which the partial transposition with respect to all half-half (even-odd) partitions becomes positive [9]. Because of the area law, one can expect that \( T_{\text{dist}}^{\text{h-h}} \) is strictly smaller than \( T_{\text{dist}}^{\text{e-o}} \). Thus, it emerges a range of temperatures...
for which the system is still entangled (as detected by the entanglement in the even-odd partition), nevertheless single particles cannot distill pure entanglement (as the half-half partitions become PPT). Indeed, for any pair of particles, there is always a half-half partition for which they are in opposite sides and the partial transposition is positive according to this splitting. In other words, bound-entangled states should appear in general under these conditions. The rest of the Letter is devoted to put this intuition on solid grounds. We first consider systems of coupled harmonic oscillators and identify a temperature range for which bound entanglement is present. This behavior is then proven to persist in the macroscopic limit. We also performed the same calculations for spin-1/2 models and found similar results.

**Harmonic oscillators.**—Consider a system composed of \( n \) harmonic oscillators, each one associated with position and momentum operators \( x_i \) and \( p_i \), respectively \( (i = 1, \ldots, n) \), described by the Hamiltonian

\[
H = \frac{1}{2} \sum_i p_i^2 + \frac{1}{2} \sum_{i,j} x_i V_{i,j} x_j \tag{1}
\]

The diagonal elements of matrix \( V \) describe the potential energy in each oscillator, while the non-diagonal terms give the coupling between oscillators \( i \) and \( j \). In this scenario both the ground and the thermal states are Gaussian. In what follows the entanglement is measured by the log-negativity \( E_N \), which quantifies by how much the partial transpose with respect to a given partition fails to be positive [10]. Thus, when \( E_N = 0 \) the considered partition is PPT. In Ref. [11] the value of \( E_N \) between two complementary groups of oscillators, \( A \) and \( B \), of the thermal state \( \rho = \exp[-H/T]/\text{Tr}\{\exp[-H/T]\} \) at temperature \( T \) was obtained:

\[
E_N = \sum \log_2\{\text{max}[1, \lambda_k(Q)]\},
\tag{2}
\]

where \( Q = P\omega^- P\omega^+ \), \( \omega^\pm = W(T)^{-1/2} V^{\pm (1/2)} \), and \( W(T) = 1 + 2[\exp(V^{1/2}/T) - 1]^{-1} \). We denote by \( \{\lambda_k(Q)\}_{k=0}^{n-1} \) the spectrum of the matrix \( Q \), whereas \( P \) is an \( n \times n \) diagonal matrix with the \( i \)th entry given by \( 1 \) or \( -1 \) depending on which group, \( A \) or \( B \), oscillator \( i \) belongs to. An exact area law for the ground-state entanglement of this system was proven in Ref. [12]. As far as for thermal states, the entanglement for a given bipartition is upper bounded by the number of connecting points [13]. Here, we mainly consider harmonic chain systems with nearest-neighbor interactions and periodic boundary conditions. The corresponding Hamiltonian (1) is given by a circulant potential matrix \( V = \text{circ}(1, -c, 0, \ldots, 0, -c) \), with \( 0 \leq c < 1/2 \). The system is equivalent to a chain of harmonic oscillators coupled with a springlike interaction and is critical when \( c \to 1/2 \).

We used Eq. (2) to compute the log-negativity for the even-odd and the half-half partition [9] for different temperatures and number of particles. Our calculations show that the entanglement follows a strict area law for nonzero temperatures: it increases linearly with \( n \) for the even-odd case, while it saturates for the half-half partition. The temperature just defines the rate the entanglement increases with \( n \) for the even-odd partition and the entanglement saturation value for large \( n \) for the half-half partition. As shown in Fig. 1, \( T_{\text{dist}}^{\text{e-o}} \) is strictly larger than \( T_{\text{dist}}^{\text{h-h}} \), indicating, as discussed above, the presence of bound entanglement. We performed our calculations for systems composed by up to 800 oscillators and found that all the computed threshold temperatures, and so also the gap \( T_{\text{dist}}^{\text{e-o}} - T_{\text{dist}}^{\text{h-h}} \), are independent of the size of the system, as can be seen in the inset of Fig. 1.

**Macroscopic systems.**—The results displayed in Fig. 1 strongly suggest that bound entanglement persists in the macroscopic limit. Actually, we can prove this statement by establishing an analytical formula for \( E_N \) in the even-odd partition and an upper bound for it in the half-half partition, when \( n \to \infty \). We present here the main steps of this proof. A more detailed analysis is given in [14].

We start with the calculation of the even-odd log-negativity, following the one given in Ref. [11] for the ground state. Consider the Hamiltonian (1) with \( V \) describing nearest-neighbor interactions as shown before. Matrix \( V \) is circulant and the matrices \( \omega^\pm \) can be diagonalized by a

![FIG. 1.](image-url)
 discrete Fourier transformation, implemented by a matrix Ω. Namely, we have that \( \Omega \omega^2 \Omega^\dagger = D^2 \), with \( D_{ij} = \delta_{ij} d_i^2 \) and \( d_i^2 = \Lambda_i^{1/2} \tanh(\Lambda_i/2T) \), where \( \Lambda_i = 1 - 2c \cos(2\pi k/n) \) are the eigenvalues of \( V \). Concerning matrix \( P \), one has that
\[
\Omega P \Omega^\dagger = \tilde{P} = \begin{pmatrix} 0 & 1 \\ \mathbb{I}_{n/2} & 0 \end{pmatrix}.
\] (3)
The spectrum of \( Q \) coincides then with the one of \( PD^2 \tilde{P} \), which in turn is straightforward to calculate due to its block diagonal structure. Defining the function
\[
f(k, n, c, T) = \frac{\Lambda^k n/2}{\Lambda_k} \tanh(\Lambda_k/2T) \tanh(\Lambda_k^{1/2}/2T)
\]
one has that the eigenvalues of \( Q \) that can contribute to the log-negativity (2) are given by \( f(k, n, c, T) \), with double multiplicity and \( k = 0, \ldots, n/4 \) (for \( n \) multiple of 4, \( n \geq 4 \)). The log-negativity of the even-odd partition is different from zero when the temperature \( T \) is such that \( f(0, n, c, T) > 1 \). In particular, the curve \( f(0, n, c, T) = 1 \) gives the threshold temperature \( T_{\text{dist}}^{\text{th}} \) in formula
\[
\frac{\Lambda^k n/2}{\Lambda_k} \tanh(\Lambda_k/2T) \tanh(\Lambda_k^{1/2}/2T) = 1. \] (4)
The threshold above coincides with the one depicted in Fig. 1 (solid line) and it is independent of the total number of particles \( n \) (see also the solid line in the inset of Fig. 1); i.e., it holds in the macroscopic limit. For temperatures below \( T_{\text{dist}}^{\text{th}} \) there exists a \( k(n, c, T) \) such that \( f(k, n, c, T) > 1 \) for \( k < k(n, c, T) \), which in turn gives rise to the following expression for the log-negativity
\[
E_N = \sum_{k=0}^{k(n,c,T)} \log_2 f(k, n, c, T)
\]
\[
\approx \frac{n}{2\pi} \int_{\mathbb{R}} d\xi \log_2 f(x, c, T),
\]
where, for large \( n \), we have replaced the sum over \( k \) by an integral over \( x = 2\pi k/n \). As a consequence, we see that the log-negativity grows linearly with the system size also for nonzero temperatures.

Regarding the half-half partition it is possible to find an exact upper bound for the threshold temperature \( T_{\text{dist}}^{\text{th}} \). Remarkably, this allows us to identify a range of temperatures for which the presence of bound entanglement can be guaranteed also in the macroscopic limit. We proceed as follows. Let us define the matrix
\[
X_{ij} = \omega_{ij} n/2 - 1 \sum_{k=0}^{n-1} (\delta_{ik} \delta_{jk} + \delta_{jk} \delta_{ih}),
\] (5)
where \( \delta_{ij} \) denotes the Kronecker delta. Following Ref. [13], the log-negativity is zero when
\[
\lambda_{\min}(W(T))^2 + 2\max_i |\lambda_i(X\omega^+)| < 1,
\] (6)
where
\[
\lambda_{\min}(W(T)) = e^{\sqrt{1+2c^2/T} + 1} - e^{\sqrt{1+2c^2/T} - 1}.
\] (7)
Recognizing that the second term in the left-hand side of (6) is twice the spectral radius \( r(X\omega^+) \) of the matrix \( X\omega^+ \), we can use any matrix norm to bound it from above [15]. An upper bound for \( r(X\omega^+) \) is then given by \( r(X\omega^+) \leq ||X\omega^+|| \leq ||X|| ||\omega^+|| \). We consider the maximum row sum matrix as a representative norm: \( ||A|| = \max_j \sum_i |A_{ij}| \). The goal, now, is to bound \( ||\omega^+|| \) and \( ||X|| \). Before proceeding, recall that \( \omega^+ \) are circulant matrices, hence completely specified by their first row \( \omega^+_n = \text{circ}(v_0^+, \ldots, v_{n-1}^+) \). One can show that
\[
v_i^+ = \frac{1}{2\pi} \int_0^{2\pi} dx d\xi (x) e^{i\xi l},
\] (8)
when \( n \to \infty \). As a consequence, for any integer \( s \), by integrating, by parts \( s \) times we have
\[
|v_i^+| \leq \frac{1}{2\pi l^2} \int_0^{2\pi} dx \left| \frac{d}{dx} d\xi(x) \right| = C_s \frac{\pi}{2\pi l^2}.
\] (9)
Let us bound first \( ||\omega^+|| \). Being \( \omega^+ \) a circulant matrix, it follows that there is no need to look for the maximum over the rows. Then one can write, for any integer \( m \), \( ||\omega^+|| = S_m^+ + E_m^+ \), where we defined the partial sum and the residual term, respectively, as follows:
\[
S_m^+ = \sum_{l=-m}^m |v_l^+|, \quad E_m^+ = 2 \sum_{l=-m+1}^\infty |v_l^+|.
\] (10)
In order to obtain a bound on \( ||\omega^+|| \) one can now fix \( m \), calculate explicitly \( S_m^+ \) and bound \( E_m^+ \) from above. This latter step can be achieved by using the bound in Eq. (9), leading to \( E_m^+ \leq C_s \zeta(s, m + 1)/\pi \) where \( \zeta(s, m + 1) \) is the generalized Riemann zeta function. Summarizing, for any integer \( s \) and \( m \), we have proven that
\[
||\omega^+|| \leq S_m^+ + C_s \frac{\pi}{2\pi l^2} \zeta(s, m + 1) \equiv K_m^+, \]
the bound above being tighter for large \( m \) and \( s \).

Concerning the upper bound for \( ||X|| \) we have that \( ||X|| \) coincides with \( ||B|| \), where \( B \) is the symmetric Toeplitz matrix with the first row given by \( (v_0^+, v_0^+/2, \ldots, v_0^+/2, v_1^+, \ldots) \). We then need to determine which row of \( B \) has the maximum sum. Fortunately, when \( n \to \infty \) the periodic boundary conditions can be disregarded and the first row of \( B \) determines its norm. This is because going from the first to the second row we simply remove the term \( |v_1^+| \), and so on for the other rows. Similarly as we did for \( ||\omega^+|| \) we obtain that for any integer \( s \) and \( m \)
\[
||X|| \leq S_m^+ + C_s \frac{\pi}{2\pi l^2} \zeta(s, m + 1) \equiv K_m^+, \]
where \( S_m^+ = \sum_{l=-m}^m |v_l^+| \). As done before, \( S_m^+ \) can be calculated explicitly and the bound is tighter for large \( m \) and \( s \).
depicted in Fig. 1 the region in the
Based on the formula above and on the threshold (4), we
ing inequality is satisfied
Summarizing, considering Eqs. (6), (7), (11), and (12), we
have shown that, in the macroscopic limit, the log-
\begin{equation}
2K_{m,+}^{m,+}K_{m,-}^{-} + \left( e^{\frac{1+2c}{T}} - \frac{1}{2} \right)^2 < 1. \tag{13}
\end{equation}
Based on the formula above and on the threshold (4), we
depicted in Fig. 1 the region in the c-T plane for which bound entanglement is present in the macroscopic limit
( shaded region). We see that for any coupling we can
guarantee that there is a range of temperatures for which the log-negativity in the half-half partition is zero, never-
thless the state is entangled. Thus the temperature, a
single macroscopic and measurable quantity, clearly deter-
mines the distillability properties of the system.

Concluding remarks.—First of all, note that our results,
when combined with those of Ref. [16], prove that en-
tangled states such that all bipartitions are PPT, cannot be
obtained for the harmonic systems studied here [17]. This
is because the thermal states become PPT and fully sepa-
rate at the same temperature.

Finally, we have considered other models of harmonic
chains as well as spin systems. All the obtained results are
consistent with the previous reasoning [14]: there is a
temperature range for which the negativity in the half-
half partitions is zero, nevertheless the system is still en-
tangled as proven by the even-odd negativity. For spin
systems we could not go beyond 12 particles due to com-
putational hardness, but the observed numerical results
again support the existence of thermal-state bound entan-
glement in the macroscopic limit (see Fig. 2). Because of
the generality of the area laws, the present results are
expected to hold for a variety of other systems. These
findings, besides showing novel aspects of entanglement in
many-body systems, are of relevance from an
application-oriented perspective. As shown here, quantum
correlations emerge “gradually” in a many-body system:
starting at high temperatures in a classically correlated
scenario, the system passes through a regime where quan-
tum correlations start to be present, but in a bound form.
Then, for lower temperatures, the system enters a truly
quantum correlated regime where free distillable entangle-
ment is present and potentially available for quantum
information applications.

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Note added.—By completing this work, bound entan-
glement was independently found in three-qubit reduced
states of the XY model at nonzero temperature [18]. Here
we have shown the existence of bound entanglement in the
whole thermal state of the macroscopic system.

[4] For a review, see R. Horodecki, P. Horodecki, M.
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