Two-qubit quantum probes for the temperature of an Ohmic environment

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We address a particular instance where open quantum systems may be used as quantum probes for an emergent property of a complex system, as the temperature of a thermal bath. The inherent fragility of the quantum probes against decoherence is the key feature making the overall scheme very sensitive. The specific setting examined here is that of quantum thermometry, which aims to exploit decoherence as resource to estimate the temperature of a sample. We focus on temperature estimation for a bosonic bath at equilibrium in the Ohmic regime (ranging from sub-Ohmic to super-Ohmic), by using pairs of qubits in different initial states and interacting with different environments, consisting either of a single thermal bath, or of two independent ones at the same temperature. Our scheme involves pure dephasing of the probes, thus avoiding energy exchange with the sample and the consequent perturbation of temperature itself. We discuss the interplay between correlations among the probes and correlations within the bath, and show that entanglement improves thermometry at short times whereas, if the interaction time is not constrained, coherence rather than entanglement, is the key resource in quantum thermometry.

I. INTRODUCTION

Quantum sensing techniques are among the most advanced quantum technologies and have led to major changes in the field of metrology in the last two decades. Upon exploiting the peculiar features of quantum systems, several novel enhanced sensors and measuring devices have been indeed suggested and demonstrated. In particular, quantum sensing based on quantum coherence and entanglement have been exploited to overcome precision bounds of classical sensors [1–4]. Among quantum sensing techniques, the use of quantum probes has recently gained attention as a non-invasive technique to estimate parameters of interest without perturbing too much the system under investigation. The basic scheme is the following: a simple quantum system, say a qubit or a pair of qubits, is prepared in a given initial state, and then interacts with an external system under investigation. After the interaction, which imprints information of some parameter onto the state of the quantum probe, the latter is measured in order to extract such information [5–9].

In this paper, we address the use of quantum probes to estimate an emergent property of a complex system, i.e. its temperature [10–15]. In particular, in view of its importance for several fields of quantum information science, we consider here quantum thermometry of a bosonic bath in the Ohmic regime, ranging from sub-Ohmic to super-Ohmic. At first sight, quantum features may not be expected to play a role in building an effective thermometer. After all, temperature is an inherently classical parameter and any change in the temperature of a sample is just changing its (classical) equilibrium distribution. On the other hand, quantum probing involves out-of-equilibrium states and since temperature is governing the amount of thermal fluctuations, the inherent fragility of quantum systems against decoherence is the key feature making the overall thermometric scheme very sensitive.

The use of single qubit as a thermometer has been analyzed recently, illustrating the interplay between the dephasing dynamics and the Ohmic structure of the environment in determining the overall precision [17, 18]. Here, we devote attention to the role of correlations in the estimation procedure, and consider both the use of correlated probes as well as thermometry of correlated environments. In particular, we investigate the use of two-qubit quantum probes to estimate the temperature of different class of bosonic systems at equilibrium, either made of a single thermal bath interacting with the two qubits, or of two independent environments having the same temperature, each one interacting locally with one of the qubits. We compare performances with those obtained with a single qubit probe, and analyze in some details the interplay among the structure of the baths, the interaction time and the quantum correlations between the qubits in determining the overall thermometric precision.

The paper is structured as follows. In section II we briefly review the main concepts of quantum estimation theory that will be used throughout the work. In Section III we present the physical model of a two-qubit system interacting with a bosonic thermal bath and we analyze how its dynamics is affected by a common environment or two independent and identical baths. In Section IV we assess the role of entanglement in the precision of the probing strategy and then in Section V we investigate the form of the POVM that guarantees optimal performances in possible experimental implementations. Moreover we test the robustness of the probes against initial perturbations and we compare the performances of Bell/product states with more general probes. Section VI closes the paper with some final remarks.
II. TOOLS OF QUANTUM ESTIMATION THEORY

In several sensing schemes, there is no direct access to the physical quantities of interest, which need to be evaluated by means of indirect measurements. In turn, inferring the value of the quantity of interest by inspecting a set of data coming from the measurement of a different observable is precisely the goal of estimation theory.

Here we discuss how to optimally estimate the temperature $T$ of a (bosonic) thermal bath by performing a measurement on a quantum probe which is let interact with the bath, considered as its environment. The probe state is described by a density operator $\rho$ which, thanks to the interaction with the environment, becomes a function of the temperature of the bath $\rho \rightarrow \rho_T$. In other words, $T$, the thermodynamical temperature of the bath, becomes just a parameter of the probe state having nothing to do with a possible probe temperature. This situation is at variance with classical thermometry, where the probe is let interact with the sample until it reaches equilibrium. Then the temperature read from the probe represents the thermodynamical temperature of both the sample and the probe. The larger set of available states and the inherent fragility of quantum states against decoherence make quantum thermometry potentially more precise than any classical protocol.

In order to determine the parameter $T$, $M$ repeated measurements of a probe observable $X$ are performed on identical preparations of the probe and the outcomes $x = \{x_1, x_2, \ldots, x_M\}$ are then used to construct an estimator $\hat{T}(x)$ of the temperature $T$. Hence, data will be distributed around the mean value of the estimator according to the probability distribution $p(x|T) = \prod_{k=0}^{M} p(x_k|T)$ (which is the conditional probability of obtaining the outcome $x_k$ when the parameter has value $T$):

$$\langle T \rangle = \int d\mathcal{P} p(\mathcal{P}|T) \hat{T}(\mathcal{P}),$$

with a certain variance $\sigma_T^2$, which characterizes the precision of the parameter estimation. At a classical level, for unbiased estimators, the variance of such distribution is bounded from below according by the Cramèr-Rao inequality [20]:

$$\sigma_T^2 \geq \frac{1}{MF(T)},$$

where $F(T)$ is the Fisher information for a single outcome,

$$F(T) = \int dx \frac{1}{p(x|T)} \left( \frac{\partial p(x|T)}{\partial T} \right)^2.$$  

In turn, $F(T)$ quantifies the amount of information carried by the random variable $x$ about the unknown parameter $T$. Upon optimising over all the possible quantum measurements one obtains the quantum Cramèr-Rao bound

$$\sigma_T^2 \geq \frac{1}{MF(T)} \geq \frac{1}{MH(T)},$$

where $H(T)$ is the quantum Fisher information (QFI from now on). One of its explicit forms is:

$$H(T) = 2 \sum_{m,n} \frac{|\langle \psi_m | \partial_T \rho_T | \psi_n \rangle|^2}{\lambda_m + \lambda_n},$$

with $\psi_n$ and $\lambda_n$ being respectively the $T$-dependent eigenvectors and eigenvalues of $\rho_T$. Clearly, the optimal measurement is the one maximizing the QFI, while the the optimal estimator is that saturating the inequality.

The QFI can be written in the following form as well:

$$H(T) = \text{tr}(\rho_T L_T^2),$$

where the symmetric logarithmic derivative (SLD) is given by

$$L_T = 2 \sum_{m,n} \frac{\langle \psi_m | \partial_T \rho_T | \psi_n \rangle}{\lambda_m + \lambda_n} |\psi_m\rangle\langle \psi_n|.$$  

This object is particularly useful since it can be proven that the optimal POVM is the spectral measure of the SLD. Related to the QFI, there is another figure of merit that gives a quantification of the precision of the estimation. This is the quantum signal-to-noise ratio (QSNR), expressed as:

$$R(T) = T^2 H(T).$$

The expression of the QSNR in Eq. [3] is derived from the ratio between the parameter $T^2$ and the (single-measure) variance of the estimator. In this way, a large value for $R$ means that the estimator has a small relative error, i.e. the error is small compared to the value of the parameter to be estimated. The QFI, QSNR and the SLD are indeed the main tools that will be used in the following.

III. PHYSICAL MODEL

The single-qubit case has already been described in previous works, dealing with the purely dephasing bath, related both to the estimation of temperature or other bath parameters [18][19]. Here we focus, under the same conditions, on the two-qubit probe scenario, which allows us to explore the role of quantum correlations and the number of qubits in inferring the temperature. We assume that initially the global state is separable $\rho(0) = \rho_s(0) \otimes \rho_B(0)$, where $\rho_B(0)$ is a thermal state of the environment at inverse temperature $\beta$ and characterized by the spectral density:

$$J_s(\omega, \Omega) = \frac{\omega^\beta}{\Omega^\beta} e^{-\omega/\Omega},$$

where $\Omega^2 = \frac{4\hbar}{k_B T}$ and $\hbar$ is the reduced Planck constant. We assume that the bath can be described by the following master equation:

$$\dot{\rho} = -i[H_{\text{eff}}, \rho] - \frac{1}{\tau} (\rho - \rho_0),$$

where $\tau$ is the bath relaxation time and $H_{\text{eff}}(t)$ is the effective Hamiltonian taking into account the interaction of the probe with the environment. The effective Hamiltonian is a function of the temperature $T$, which is, in turn, a function of the bath parameters $\{\beta, \Omega\}$.

The goal is to find an optimal measurement that allows us to estimate the temperature $T$ with the highest possible precision. The optimal measurement is the one that maximizes the QFI, which, in the case of a single-qubit probe, is given by

$$H(T) = 2 \sum_{m,n} \frac{\langle \psi_m | \partial_T \rho_T | \psi_n \rangle^2}{\lambda_m + \lambda_n}.$$
where $\Omega$ is the cutoff frequency and $s$ is a Ohmic parameter which distinguishes between sub-Ohmic ($s < 1$), Ohmic ($s = 1$) and super-Ohmic ($s > 1$) regime. The probe is composed of two qubits, which can be in a product or entangled initial state. Moreover, the two qubits can be embedded in a common environment or in independent local baths.

### A. Common bath

The Hamiltonian of two qubits interacting with a same bosonic environment can be written as:

$$H = \sum_{j=1}^{2} \frac{\omega_j}{2} \sigma_3^{(j)} + \sum_{k=0}^{\infty} \omega_k b_k^\dagger b_k + \sum_{j=1}^{2} \sum_{k=0}^{\infty} \sigma_3^{(j)} (g_k b_k^\dagger + g_k^* b_k),$$  \hspace{1cm} \text{(10)}$$

where the index $j$ labels the qubits and the index $k$ labels the modes of the bath. $\sigma_3^{(j)}$ is the third Pauli matrix and $b(b^\dagger)$ are the annihilation (creation) bosonic operators. Moving to the continuum $\sum_k g_k \to \int d\omega J_\omega(\omega, \Omega)(2 g(\omega))^{-1}$ and letting the compound system probe plus bath evolve up to rescaled dimensionless time $\tau = \Omega t$ according to the above Hamiltonian, after tracing over the bath’s degrees of freedom, one finds that the dynamics of the two-qubit reduced density matrix can be expressed as [21]:

$$\rho_{\text{red}}(\tau, T) = \mathcal{V}(\tau, T) \circ \mathcal{R}(\tau) \circ \rho,$$  \hspace{1cm} \text{(11)}$$

where $\mathcal{V}(\tau, T)$ is given by:

$$\mathcal{V}(\tau, T) = \begin{pmatrix} 1 & e^{-\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & e^{-4\Gamma_s(\tau, T)} \\ e^{-\Gamma_s(\tau, T)} & 1 & 1 & e^{-\Gamma_s(\tau, T)} \\ e^{-\Gamma_s(\tau, T)} & 1 & 1 & e^{-\Gamma_s(\tau, T)} \\ e^{-4\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & 1 \end{pmatrix},$$  \hspace{1cm} \text{(12)}$$

and

$$\mathcal{R}(\tau) = \begin{pmatrix} 1 & e^{2i f(\tau)} & e^{2i f(\tau)} & 1 \\ e^{-2i f(\tau)} & 1 & 1 & e^{-2i f(\tau)} \\ e^{-2i f(\tau)} & 1 & 1 & e^{-2i f(\tau)} \\ 1 & e^{2i f(\tau)} & e^{2i f(\tau)} & 1 \end{pmatrix},$$  \hspace{1cm} \text{(13)}$$

and $\circ$ is the Hadamard (entrywise product), while $\rho$ is the initial state of the qubits. The decoherence function depends on time and the dimensionless temperature $T = (\Omega \beta)^{-1}$. By expressing also $\omega$ in unit of $\Omega$, i.e. $\omega \rightarrow \omega/\Omega$, we may write

$$\Gamma_s(\tau, T) = \int_0^{\infty} d\omega \ e^{-\omega} \frac{1 - \cos \omega \tau}{\omega^2} \coth \left( \frac{\omega}{2T} \right),$$  \hspace{1cm} \text{(14)}$$

whose analytic expression can be found in [13]. The function $f(\tau)$ is instead a temperature-independent quantity

$$f(\tau) = \frac{1}{2} \int_0^{\infty} d\omega J_\omega(\omega) \frac{\omega \tau - \sin \omega \tau}{\omega^2}.$$  \hspace{1cm} \text{(15)}$$

The matrix $\mathcal{V}(\tau, T)$ is dominant at short times and the matrix $\mathcal{R}(\tau)$ is able to create entanglement between the qubits.

### B. Independent baths

In the case the two qubits interact with identical but independent local environments, the Hamiltonian takes the form:

$$\mathcal{H} = \sum_{j=1}^{2} \frac{\omega_j}{2} \sigma_3^{(j)} + \sum_{k=0}^{\infty} \omega_k b_k^\dagger b_k + \sum_{j=1}^{2} \sum_{k=0}^{\infty} \sigma_3^{(j)} (g_k b_k^\dagger + g_k^* b_k),$$

Let the initial density matrix of the bath be factorized $\rho_B(0) = \rho_B^{(1)} \otimes \rho_B^{(2)}$ where $\rho_B^{(1,2)}$ are thermal Gibbs states at dimensionless temperature $T$. In this case, the evolution of the probe is given by:

$$\rho_{\text{red}}(\tau, T) = \mathcal{W}(\tau, T) \circ \rho,$$  \hspace{1cm} \text{(16)}$$

where $\mathcal{W}(\tau, T)$ is the tensor product of two single qubit dephasing maps

$$\mathcal{W}(\tau, T) = \begin{pmatrix} 1 & e^{-\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & e^{-2\Gamma_s(\tau, T)} \\ e^{-\Gamma_s(\tau, T)} & 1 & e^{-2\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} \\ e^{-2\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & 1 & e^{-\Gamma_s(\tau, T)} \\ e^{-\Gamma_s(\tau, T)} & e^{-2\Gamma_s(\tau, T)} & e^{-\Gamma_s(\tau, T)} & 1 \end{pmatrix}. $$  \hspace{1cm} \text{(17)}$$

### IV. TWO-QUBIT THERMOMETRY

We now compare the performances of two-qubit quantum probes for the temperature by analyzing the behaviour of the QFI. We consider two different initial states for the qubits, i.e. initially entangled and initially separable probes and we study the role of local against common baths.

#### A. Entangled qubits

We start by analyzing the performances of quantum probes initially prepared in one of the Bell states: $|\varphi^{\pm}\rangle = \frac{1}{\sqrt{2}}(|00\rangle \pm |11\rangle)$ and $|\psi^{\pm}\rangle = \frac{1}{\sqrt{2}}(|01\rangle \pm |10\rangle)$. We firstly consider the scenario with a common environment: the Bell states $|\psi^{\pm}\rangle$ are not changed by the map [11], while $|\varphi^{\pm}\rangle$ are affected by the environment. Indeed, for both states $|\varphi^{\pm}\rangle$ the QFI is:

$$H_{\text{CB}}^{\text{ent}}(\tau, T, s) = \frac{\partial^2}{\partial \Gamma_s^2} \left( \Gamma_s(\tau, T) \right)^2 \frac{2}{e^{4\Gamma_s(\tau, T) - 1}.} $$  \hspace{1cm} \text{(18)}$$

Notice that this quantity does not depend on $f(\tau)$. On the other hand, when the interaction of the probe with two identical and independent local baths is considered, all four Bell states lead to the same QFI:

$$H_{\text{LB}}^{\text{ent}}(\tau, T, s) = \frac{2}{e^{4\Gamma_s(\tau, T) - 1}.} $$  \hspace{1cm} \text{(19)}$$
A numerical evaluation of the QFI shows that the Bell states are optimal within the subset of states

$$|\Phi_\alpha\rangle = \cos \alpha |00\rangle + \sin \alpha |11\rangle \quad (20)$$

$$|\Psi_\alpha\rangle = \cos \alpha |01\rangle + \sin \alpha |10\rangle \quad (21)$$

with $\alpha \in [0, 2\pi]$ a real parameter.

B. Separable qubits

Let us consider two qubits initially prepared in the product state $|++\rangle$, where $|+\rangle = \frac{1}{\sqrt{2}}(|0\rangle + |1\rangle)$, which is a particular instance of the one-parameter family of states

$$|\psi_\alpha\rangle := |\psi_\alpha\rangle \otimes |\psi_\beta\rangle \quad , \quad |\psi_\alpha\rangle = \cos \alpha |0\rangle + \sin \alpha |1\rangle .$$

In the common-bath case, the QFI has the expression:

$$H_{\text{CB}}(\tau, T, s) = \frac{2(2 - e^{4\Gamma_s(\tau, T)} + 2e^{4\Gamma_s(\tau, T)} + e^{6\Gamma_s(\tau, T)})[\partial_T \Gamma_s(\tau, T)]^2}{3 e^{8\Gamma_s(\tau, T)} - 2 e^{4\Gamma_s(\tau, T)} - 1} \quad (23).$$

Again, the quantity $f(\tau)$ does not play a role in the estimation of the temperature. In the case of local independent baths, instead, the QFI reads:

$$H_{\text{LB}}^{\text{sep}}(\tau, T, s) = \frac{2[\partial_T \Gamma_s(\tau, T)]^2}{e^{2\Gamma_s(\tau, T)} - 1} \quad (24).$$

As expected it is twice the QFI of a single-qubit probe \[18\]. Both results \[23\] and \[24\] can be obtained by initializing the probe also in one of the states $|--\rangle$, $|+\rangle$ and $|--\rangle$.

Let us now compare the behaviour of the quantum signal-to-noise ratio (QSNR) given in \[8\], that for sake of simplicity we shall denote by $R$ in the following, for different initial states and bath parameters. In figure Fig. 1 we show the behaviour of $R$ for different values of the Ohmicity parameter as a function of dimensionless time $\tau$ and temperature $T$. The four different colors correspond to the possible scenarios analyzed in this work, i.e. common/local bath and entangled/separable initial state. Right column: Ratio $R_{\text{CB}}^{\text{ent}}/R_{\text{LB}}$ for the same three values of $s$. FIG. 1. Left column: behaviour of the QSNR for three different values of $s = 0.5$, $1$, $3$, as a function of dimensionless time $\tau$ and temperature $T$. The four different colors correspond to the possible scenarios analyzed in this work, i.e. common/local bath and entangled/separable initial state. Right column: Ratio $R_{\text{CB}}^{\text{ent}}/R_{\text{LB}}$ for the same three values of $s$. In Fig. 1 we also analyze the ratio $R_{\text{CB}}^{\text{ent}}/R_{\text{LB}}$ to emphasize the optimal procedures in the $(\tau, T)$-space. As we see from the graphs, the Ohmicity parameter $s$ only affects the qualitative behaviors of the ratio, and the two different estimation strategies are clearly displayed. In order to better understand why the change from one optimal strategy to the other occurs, we examine the behaviour of the quantum Fisher information $H$ for small values of $\tau$:

$$H(\tau, T, s) \simeq \tau^2 \left\{ \begin{array}{ll}
\frac{1}{\gamma(\tau, s)} [\partial_T \gamma(\tau, s)]^2 & \text{for } H_{\text{CB}}^{\text{ent}} \\
\frac{1}{2\gamma(\tau, s)} [\partial_T \gamma(\tau, s)]^2 & \text{other cases,}
\end{array} \right. \quad (25)$$

where

$$\Gamma_s(T, \tau) \simeq \frac{\tau^2}{2} \int_0^\infty dx \ x^s e^{-x} \coth \frac{x}{2T} \equiv \frac{\tau^2}{2} \gamma(T, \tau) \quad (26)$$

at short times. So, $H_{\text{CB}}^{\text{ent}}$ outperforms by a factor 2 the other strategies in the $\tau \ll 1$ regime.

In Fig. 2 we show the maximum value of the QSNR, called $R_{\text{opt}}$, as a function of the temperature and for different values of the Ohmicity parameter $s$. Notice that
this maximum is obtained in the case where two separable qubits are embedded in local baths. We notice that, as the Ohmicity parameter $s$ grows, the maximal values of the QSNR decrease for a fixed value of the temperature, except for $T$ becoming large and the dependency on $s$ disappears. The time corresponding to the maximum of $R$, $R_{\text{opt}}$, is called optimal time $\tau_{\text{opt}}$, notice that, especially for low temperatures, it strongly depends on the Ohmicity parameter. Furthermore, Fig. 2 shows agreement with the behaviour found in the case of single qubit thermometry [18]. This is expected since the maximum of the QSNR is obtained employing two qubits in a product states that evolve in local baths. This procedure is the same as sending twice a single qubit into the quantum environment.

V. IMPLEMENTATIONS

In this section we analyze possible ways to experimentally implement the optimal strategies. We focus on entangled qubits in a common bath at short times, and separable qubits in independent baths for longer interaction times. By calculating the symmetric logarithmic derivative, we derive the optimal POVM to be performed on the qubits in order to infer the value of temperature in a bosonic environment.

A. SLD for Bell states in a common bath

We begin by considering the scenario of the optimal Bell states $|\varphi^{\pm}\rangle$ in a common bath. The SLD is derived using equation (7):

$$L_{\text{CB}}^\text{opt} = a_- |\varphi^-\rangle\langle\varphi^-| - a_+ |\varphi^+\rangle\langle\varphi^+|,$$

where

$$a_{\pm} = \frac{4 \partial_1 \Gamma_\xi (T, \tau)}{e^{4 \Gamma_\xi (T, \tau)} + 1} \pm 1$$

B. SLD for $|++\rangle$ states in local baths

Regarding the separable state $|++\rangle$ coupled with two independent local baths, the SLD has the expression:

$$L_{\text{LB}}^\text{sep} = b_+ \Pi_+ - b_- \Pi_- + c \left(|\varphi^-\rangle\langle\varphi^-| + |\psi^-\rangle\langle\psi^-|\right)$$

where

$$b_{\pm} = \frac{2 \partial_1 \Gamma_\xi (T, \tau)}{e^{4 \Gamma_\xi (T, \tau)} + 1} \pm 1 \quad c = \frac{2 \partial_1 \Gamma_\xi (T, \tau)}{e^{4 \Gamma_\xi (T, \tau)} - 1}$$

and $\Pi_{\pm} = |\pm\rangle\langle\pm|$. Although the presence of Bell states in Eq. (29) may led to think that global measurements on both qubits are needed in order to discriminate the various projections, actually this is not the case: the eigenstates of $L_{\text{LB}}^\text{sep}$ are eigenvectors of $\sigma_1 \otimes \sigma_1$, an observable that can be accessed locally, namely by separate measurements on the two qubits. This scenario is sketched in Fig. 4.
C. Robustness

We want to check the robustness of the optimal states, namely we want to quantify how the maximal QFI is affected by small deviations $\delta$ from the optimal states. We start by focusing on the two optimal scenarios. As already mentioned, sending two qubits in a product state in independent local environments corresponds to repeating the same experiment twice; therefore we can write the QFI for a generic initial state $|\psi_{\alpha\beta}\rangle$ defined in (22) as the sum of the QFI of the two single qubits:

$$H_{\text{LB}}^{\alpha\beta}(\tau, T) = \left[ \sin^2(2\alpha) + \sin^2(2\beta) \right] \frac{(\partial_T \Gamma_s(\tau, T))^2}{e^{2\Gamma_s(\tau, T) - 1}}. \tag{31}$$

We now consider the perturbed initial state $|\psi_{\alpha+\delta\beta+\delta\beta}\rangle$ around the optimal choice ($\alpha = \beta = \pi/4$), we find the perturbed QFI:

$$H_{\text{LB}}^{\delta\beta}(\tau, T) = H_{\text{LB}}^{\alpha\beta}(\tau, T) \left[ 1 - 2 \left( \delta^2_{\alpha} + \delta^2_{\beta} \right) \right]. \tag{32}$$

On the other side, the general entangled state $|\Phi_\alpha\rangle$ defined in (20) in a common bath yields

$$H_{\text{CB}}^{\delta}(\tau, T) = \frac{16 \sin^2(2\alpha) \left[ \partial_T \Gamma_s(\tau, T) \right]^2}{e^{8\Gamma_s(\tau, T) - 1}}. \tag{33}$$

If we perturb the state $|\Phi_\alpha\rangle$ around $\alpha = \pi/4$, we find the following QFI:

$$H_{\text{CB}}^{\delta}(\tau, T) \approx H_{\text{CB}}^{\alpha}(\tau, T) \left( 1 - 4\delta^2 \right). \tag{34}$$

In both cases the deviation from the optimal QFI are at second order on the magnitude of the perturbation, i.e. the QFI obtained from both $|\varphi^+\rangle$ and $|++\rangle$ is robust with respect to the probe preparation.

D. Performances of other probes

So far we maximized the QFI within specific subsets of state, $|\Phi_\alpha\rangle$ and $|\psi_{\alpha\beta}\rangle$ for entangled and separable probes and we found that, within these families, the states $|\varphi^+\rangle$ and $|++\rangle$ are optimal regardless of $\tau$ and $T$. We now consider the normalized superposition:

$$|S_{\alpha}\rangle = C \left( \cos \alpha |\varphi^+\rangle + \sin \alpha |++\rangle \right), \tag{35}$$

where $C$ is a normalization constant. In Fig. (5) we compare the behaviour of the QSNR for $s = 1$ and $T = 1$ for different values of the parameter $\alpha$ in Eq. (35). Both the local and global scenarios are analyzed. The largest value of the QSNR is obtained for $\alpha = \pi/2$, corresponding to initially separable qubits. At shorter times, the $\alpha = 0$ case, i.e. entangled qubits, is the best probe, but only in the common-bath case. In the common-environment scenario, moreover, there is also a small time-interval where other probes perform better, while this behaviour is never seen in the local-bath case.

VI. CONCLUSIONS

The aim of thermometry is to estimate the temperature of a certain object without altering its properties, temperature included. When the object is a quantum system, the idea of finding a non-invasive probe becomes a necessity, in order to avoid decoherence. In this paper, we have addressed optimal quantum thermometry for bosonic thermal baths by means of two-qubits quan-
tum probes. We have analyzed the case of two independent baths, each interacting locally with a qubit, and the case of a common reservoir acting upon the two qubits. Moreover, we have considered two families of initial states for the probes: Bell states and product states of the form $|\sigma, \eta\rangle$ where $\sigma = \pm$ and $\eta = \pm$. In particular, we have analyzed the behaviour of the QFI and the QSNR as a function of time, temperature and Ohmicity parameter. Our results show that at short times the best way to probe the temperature of the bath is to employ maximally entangled qubit interacting with the same global bath, independently from the Ohmicity of the environment. However, if time is not considered a resource, we found that the best estimate is obtained upon using two qubits in a product states, each one subject to a local environment, i.e. the best strategy is to repeat twice a single-qubit probe measurement [17][18]. Notice that the notions of same bath or two independent replicas of the same bath do not require multiple physical systems to be implemented in practice. In a realistic environment, the two situations correspond respectively to the two qubits propagating one close to each other, in order to sense the same portion of the environment, or far away, such that possible spatial correlations of the bath may be neglected. We have analysed possible implementations of the optimal estimation schemes discussed above and we found that the entangled case boils down to a Bell measurement over the entangled quits, while the case of factorized probe may be implemented by a local measurement performed separately on the two quits. We have analyzed the behaviour of the estimation precision against perturbations in the preparation of the initial state and have shown the they contribute to decrease the QFI with second order corrections in the perturbations, i.e. our scheme is robust.

We have also compared the performances of the Bell and product states with other probes. Our intent was to analyze whether a combination of the Bell and separable states could improve the precision of the estimation. We showed that the absolute maximum of the QFI is indeed obtained for independent qubits in a product state. However, there exists temporal regions where a superposition of the two probes perform better, though only in the common-bath scenario. In order to gain more insight into the role of other probes, we have randomly generated pure states and analyzed the maximum value of their QFI versus the entanglement of the generated state. We found numerical evidence that the state $|++\rangle$ is the optimal one in the case of local environments, while for the common bath there exist initial entangled states of the probe that outperform the product state in a certain temporal region that depends upon the temperature and the Ohmicity parameter. The corresponding optimal measurement, however, do not correspond to an easy implementable one and thus it may be challenging to achieve the corresponding enhancement in a practical scenario.

Summarizing our results, we have that entanglement improve thermometry at short times whereas, if the interaction time is not constrained, coherence rather than entanglement, is the key resource to improve precision. We also emphasise that our scheme for quantum thermometry is based on pure dephasing and it does not involve energy exchanges between the probes and the bath. The corresponding measuring protocols are thus inherently non invasive, and they may be of value whenever a direct inspection threatens to destroy the sample, or to perturb the temperature itself, as it would happen in the case of very cold samples.

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