Experimental test of exchange fluctuation relations in an open quantum system

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Elucidating the energy transfer between a quantum system and a reservoir is a central issue in quantum nonequilibrium thermodynamics, which could provide novel tools to engineer quantum-enhanced heat engines. The lack of information on the reservoir inherently limits the practical insight that can be gained on the exchange process of open quantum systems. Here we investigate the energy transfer for an open quantum system in the framework of quantum fluctuation relations. As a novel toolbox, we employ a nitrogen-vacancy center spin qubit in diamond, subject to repeated quantum projective measurements and a tunable dissipation channel. In the presence of energy fluctuations originated by dissipation and quantum projective measurements, the experimental results, supplemented by numerical simulations, show the validity of the energy exchange fluctuation relation, where the energy scale factor encodes missing reservoir information in the system out-of-equilibrium steady-state properties. This result is complemented by a theoretical argument showing that, also for an open three-level quantum system, the existence of an out-of-equilibrium steady state dictates a unique time-independent value of the energy scale factor for which the fluctuation relation is verified. Our findings pave the way to the investigation of energy exchange mechanisms in arbitrary open quantum systems.

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I. INTRODUCTION

The connection among statistical properties of out-ofequilibrium dynamical systems, thermodynamics quantities, and information theory has been deeply investigated in classical and quantum systems and codified in terms of fluctuation relations [1–7]. However, in open quantum systems, despite several contributions [8–13], such connection is far from being completely understood, especially regarding the competition between thermal and quantum fluctuations. The latter assumes a paramount role at the nanoscale, for example, for developing quantum thermal engines [14,15] or studying information-energy conversion [16,17]. Accounting for the statistical fluctuations is the key to reformulate the second law of thermodynamics, usually expressed as inequalities, in terms of equalities. As a major example, the Jarzynski equality [18,19] relates the exponentiated negative work done on a system, averaged over a statistically relevant ensemble of realizations of the system dynamics, with the change in the free energy between two equilibrium thermal states. This framework has been also extended to describe the transport of energy and matter between different systems with different temperatures and chemical potentials [20,21]. Remarkably, these relations hold for any kind of process driving the system arbitrarily far from equilibrium, provided that the initial state is in thermodynamic equilibrium.

In quantum mechanical settings, the fluctuation relations can be recast in terms of the characteristic function— Fourier transform of the probability distribution function of the considered nonequilibrium quantity. This contains the full information about the fluctuations statistics and is obtained from two-time quantum correlations rather than by a



FIG. 1. Schematic illustration of the experiment. A two-level system, in our case a single NV center qubit in diamond, is subject to series of quantum projective measurements (QPMs), and controllable energy exchange with a reservoir. We investigate the energy exchange fluctuations occurred in the system from the beginning to the end, over a statistically relevant ensemble of realizations of the protocol.

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single-time expectation value [22]. Still, in the absence of a heat reservoir, where the internal energy variation ΔE is solely due to work, a formally equivalent quantum version of the Jarzynski equality (QJE) [23–25] has been verified in various experimental settings with no heat flux involved, ranging from single trapped ions [26] to liquid-state nuclear magnetic resonance platforms [27], atom chips [28], and superconducting Xmon qubits [29].

Opening an energy exchange channel from a quantum system to a reservoir poses challenges for describing the nonequilibrium thermodynamic processes that take place [4]. While a dephasing dynamics can be effectively traced back to the case of an isolated system [30], the energy transport between a system and its environment is expected to be described by an exchange fluctuation relation [3,4,8]. Such relation was originally introduced to describe the heat exchange between two bodies in thermal contact [20], each initially in thermal equilibrium. Evaluating the exchange fluctuation relation would then require information on the reservoir that is often not practically available. Since the practically accessible quantity is the system energy variation at two different times [27,29,30], the exchange fluctuation relation is conveniently expressed in terms of the characteristic function of the energy variation statistics $G(\varepsilon) \equiv \langle \exp(-\varepsilon \Delta E) \rangle$. Thus, we argue that the energy exchange fluctuation relation for an open quantum system with a time-independent Hamiltonian can be written as

$$G(\varepsilon) = 1, \tag{1}$$

where the scale factor ε has to be determined and is expected to be of the order of the system energy. Exemplarily, for a classical or quantum system under thermalizing dynamics, the energy scale ε is predicted to be related to the inverse temperatures β_{in} and β_{∞} of the initial and final states as $\varepsilon = \Delta\beta$, with $\Delta\beta = \beta_{in} - \beta_{\infty}$ [31]. For a *quantum* system coupled to a tunable and generically nonthermal environment, it is not yet specified whether ε such that $G(\varepsilon) = 1$ exists, and whether and how it is related to the system dynamics and its asymptotic behavior. Here we investigate these issues.

We experimentally test the energy exchange fluctuation relation in a quantum simulator, as sketched in Fig. 1, subject to repeated quantum projective measurements (QPMs) and energy dissipation, where the resulting dissipative dynamics drives the system toward an (out-of-equilibrium) energy steady state. We realize the simulator with the use of a single nitrogen-vacancy (NV) center qubit in diamond at room temperature, in the presence of trains of short laser pulses. Each absorbed laser pulse results in a QPM [32], and in an energy redistribution that can be modeled as a controlled energy exchange with a Markovian reservoir [14]. The time intervals between QPMs follow a stochastic distribution due to the finite absorption probability. Tuning the laser duration and power enables the control of the coupling strength between the quantum system and the reservoir.

The combined effect of QPMs and dissipation can create or destroy quantum coherence during the system dynamics, an effect that goes beyond the classical description. While QPMs affect the energy distribution of the system, they are expected to preserve the validity of quantum fluctuation relations [33,34], also for stochastic distributions of QPMs [35]. However, the energy fluctuations of a quantum system in the presence of QPMs and dissipative dynamics have not been studied yet. Measuring the statistics of the exponentiated energy fluctuations through a two-point measurement (TPM) protocol, we experimentally verify the exchange fluctuation relation for an open two-level quantum system. We find out a uniquely determined value of ε for which $G(\varepsilon) = 1$, incorporating missing information about the reservoir. For a two-level system, where any diagonal density matrix in the energy basis can be recast in terms of an effective temperature, ε encompasses the initial and final populations of the quantum system through their effective temperatures.

While these results have been obtained for an effective quantum two-level system, we provide a further analysis involving a three-level system that is affected by a spontaneous emission process. In the Appendix 2 we show with this example that the validity of the exchange fluctuation relation is conditioned by the existence of a unique nontrivial time-independent energy scale factor ε , without requiring thermalizing dynamics. Therefore, our results are representative of quantum systems with dimension larger than two subjected to dissipation dynamics.

II. PROTOCOL IMPLEMENTATION

Here we introduce our experimental platform and describe the protocol used for measuring the energy fluctuations. We show that this protocol can be broadly applied to any finitedimensional quantum system, and we detail it for our specific experimental setup.

The experimental platform is based on a negatively charged NV center—a localized impurity in the diamond lattice based on a nitrogen substitutional atom and a nearby vacancy—which forms an electronic spin S = 1 in its orbital ground state [see Fig. 2(a)]. The electronic spin can be initialized into the state $|0\rangle$, where $|m_s = 0, \pm 1\rangle$ stands for the eigenvalues of the spin operator S_z along the NV symmetry axis, via optical spin pumping under laser excitation [36]. A strong magnetic bias field removes the degeneracy of the spin states $|\pm 1\rangle$, allowing selective coherent manipulation of the transition $|0\rangle \leftrightarrow |+1\rangle$. The spin-dependent intensity of the emitted fluorescence enables the optical readout of the states $|0\rangle$ and $|+1\rangle$. A continuous nearly resonant microwave field sets the Hamiltonian \mathcal{H} of the two-level system in the frame rotating at the microwave frequency,

$$\mathcal{H} = \frac{\hbar\omega}{2} (\cos\alpha \ \sigma_z - \sin\alpha \ \sigma_x), \qquad (2)$$

with eigenstates $\{|\uparrow\rangle, |\downarrow\rangle\} = \{\cos\frac{\alpha}{2} |0\rangle - \sin\frac{\alpha}{2} |+1\rangle, \\ \sin\frac{\alpha}{2} |0\rangle + \cos\frac{\alpha}{2} |+1\rangle\}, \text{ and eigenvalues } E_{\uparrow} = \hbar\omega/2 \text{ and } E_{\downarrow} = -\hbar\omega/2. \text{ Here } \sigma_i \text{ are Pauli matrices, } \tan\alpha = -\Omega/\delta \text{ and } \omega = \sqrt{\delta^2 + \Omega^2}, \Omega = 1.3 \text{ MHz} \text{ being the bare Rabi frequency, } \\ \text{and } \delta \in [0, \Omega] \text{ the microwave detuning. The Hamiltonian (2) } \\ \text{remains unchanged during the protocol, and the nonunitary } \\ \text{dynamics is due to repeated QPMs and dissipation acting } \\ \text{along the } \sigma_z \text{ axis, as explained below.} \end{cases}$

The resulting quantum dynamics induces energy fluctuations of the spin system. To characterize the statistics of these energy fluctuations, we employ a TPM protocol, where the energy is measured at the initial and final times of the



FIG. 2. (a) Schematic representation of the photodynamics of a nitrogen-vacancy center. Photon absorption (green upward arrow) and spin preserving spontaneous emission (red downward arrow) between the electronic ground and excited levels realize quantum projective measurements of the spin state along the z axis (z-QPM). A nonradiative decay channel (left gray arrows) enables controlled optical spin pumping to the $|0\rangle$ state. Inset: Effective two-level system considered in the protocol, formed by two dressed spin states set by quasiresonant continuous driving [see Eq. (2)]. (b) Block diagram of the two-point measurement protocol implementation and experimental pulse sequence. After initialization in the $|0\rangle$ state, for each protocol repetition a Hamiltonian eigenstate $|\uparrow\rangle$ ($|\downarrow\rangle$) is prepared by applying a rotating microwave (mw) gate R_{ν}^{α} ($R_{\nu}^{\alpha+\pi}$). During the time t_{fin} , the mw-driven spin unitary evolution set by the Hamiltonian \mathcal{H} , defined in Eq. (2), is perturbed by equidistant short laser pulses acting as z-QPMs plus a dissipation channel (D). The interpulse time (270 $\leq \tau \leq$ 750 ns) is much longer than each pulse duration ($t_L = 41$ ns). At the end, a quantum projective measurement of the final-state energy (\mathcal{H} -QPM) is realized with a mw gate R^{α}_{μ} and a spin-selective fluorescence intensity measurement read out. (c) Exemplary spin-state evolution (red filled circles) on the Bloch sphere for a single realization of the protocol. (cI) Initially prepared Hamiltonian eigenstate, e.g., $|\uparrow\rangle$. (cII) $|\uparrow\rangle$ is projected in one of the σ_z eigenstates, e.g., $|+1\rangle$, due to photon absorption as denoted by the the green arrow, and then evolves under unitary dynamics (dashed circle) until a subsequent photon absorption. (cIII) Due to the spin amplitude damping, after several laser absorptions the system would most likely be in the |0> state. After the last short laser absorption the state follows a unitary evolution. On average over several realizations of the protocol, the state before the \mathcal{H} -QPM has the same energy than the state $|0\rangle$ but with an unknown phase in the Hamiltonian basis. (cIV) Applying a rotation R_{-v}^{α} allows us to obtain the energy of the system by measuring σ_z . Indeed, the final state is projected into one of the Hamiltonian eigenstates (e.g., $|\downarrow\rangle$).

process. We implement this protocol as shown in Fig. 2(b) and specified below. Figure 2(c) shows an example of a single trajectory followed by the system during the complete protocol.

(i) Preparation of initial thermal states, and energy quantum projective measurement (\mathcal{H} -QPM). Each realization starts by preparing the system in one of its energy eigenstates $|i\rangle\langle i|$. The experiment is then repeated P_iN times for each eigenstate over a statistical ensemble of N realizations, where P_i is the population fraction of each energy eigenstate following a thermal distribution [37]. Independently of the system dimension, this is equivalent to considering the statistical result of an initial energy measurement (\mathcal{H} -QPM) applied to the thermal state $\sum_i^n P_i |i\rangle\langle i|$, provided that N is large enough.

In the experiment, each of the two energy eigenstates (described by the density operators $\rho_{\uparrow} = |\uparrow\rangle\langle\uparrow|$ and $\rho_{\downarrow} = |\downarrow\rangle\langle\downarrow|)$ is prepared by optically pumping the NV into the $|0\rangle$ state and then applying a rotation R^{α} (respectively, $R^{\alpha+\pi}$) along σ_y , via a microwave (mw) gate as depicted in Fig. 2(b).

(ii) Evolution under repeated quantum projective measurements and dissipative dynamics. The system is repeatedly opened for short time intervals and stays closed otherwise, due to being subject to a series of QPMs of an operator noncommuting with the Hamiltonian, and a dissipative dynamics. These effects on top of the unitary evolution $U = e^{-i\mathcal{H}t/\hbar}$ lead the system into an asymptotic steady state.

In the experiment, we apply to the NV center trains of short laser pulses with duration t_L at intervals τ , as depicted in Fig. 2(b). The laser pulses trigger cycles of spin-preserving radiative transitions from the ground to the excited states [see Fig. 2(a) and Appendix 1]. On photon absorption, any superposition or mixed spin state is projected into either one of the two σ_z eigenstates $|0\rangle$ or $|+1\rangle$, while the state coherence imprinted by the microwave during the prior evolution is destroyed in the σ_z basis. This results in a quantum projective measurement of σ_z (z-QPM) [32], even when the measurement outcome is not recorded. Significantly, this mechanism produces coherence in other bases, such as the energy basis (for $\alpha \neq 0$), as shown in Fig. 2(c). Due to the finite photonabsorption probability, a train of equidistant laser pulses entails a stochastic time distribution of z-QPMs, for each single realization. In addition, the absorption of laser pulses induces a partial population transfer to $|0\rangle$, due to optical pumping [see Fig. 2(a)], with an effective rate that depends on the number of excitation-decay cycles performed by the system, which can be controlled by changing the laser pulse duration and power. This spin amplitude damping mechanism is equivalent to a controlled dissipative channel toward $|0\rangle$ in the two-level system, which together with z-QPMs, incorporates all the laser-induced NV photodynamics involving the ground and excited triplet states, and the metastable singlet state. The overall effect takes the system into an



FIG. 3. Conditional probabilities $P_{\uparrow\uparrow\uparrow}$ and $P_{\uparrow\downarrow\downarrow}$ to measure $E_f = E_{\uparrow}$ as a function of the evolution time $t_{\rm fin}$ starting from the Hamiltonian eigenstates $|\uparrow\rangle$ (blue circles) or $|\downarrow\rangle$ (gray squares), with $\alpha = \pi/2$ (a) and $\alpha = \pi/3$ (b). Vertical grids indicate the time position of each laser pulse, separated by $\tau = 5\pi/3\omega$, where the energy of the qubit jumps due to projective measurements and dissipative dynamics. The probability that a *z*-QPM occurs during each laser pulse is 0.18. Error bars denote the experimental uncertainty given by the photon shot noise. The solid lines are obtained by a Monte Carlo simulation of the dynamics (Sec. S.II in Ref. [38]). The horizontal dotted lines represent the conditional probabilities for the asymptotic (out-of-equilibrium) steady states (see text).

asymptotic out-of-equilibrium steady-state in the Hamiltonian basis.

(iii) Measurement of the final energy. The statistics of ΔE is provided by the conditional probabilities $P_{j|i} = P(E_{fin} =$ $E_j | E_{in} = E_i$) to measure E_j as the final energy, once known the initial energy E_i [see Eq. (A5) of the Appendix 3]. For an *n*-dimensional system, measuring n - 1 diagonal elements of the final density operator for each initial energy eigenstate $|i\rangle\langle i|$ bestows a full knowledge of all the conditional probabilities. For the NV qubit, we measure the conditional probabilities $P_{\uparrow|\uparrow}$ and $P_{\uparrow|\downarrow}$ for the spin to go in ϱ_{\uparrow} (with energy $E_{\text{fin}} = E_{\uparrow}$), when starting respectively from ρ_{\uparrow} or ρ_{\downarrow} . To implement the final energy measurement (\mathcal{H} -QPM), we apply a mw gate that maps \mathcal{H} into σ_z [as detailed in Figs. 2(b) and 2(cIV)], and then measure the σ_z operator by detecting the presence (or absence) of emitted photons. Low collection efficiency and photon shot noise impose the need of repeating the procedure several times ($\sim 1.6 \times 10^6$) and averaging over the detected intensity to reduce the readout uncertainty.

III. STATISTICS OF THE ENERGY VARIATION

Figure 3 shows the conditional probabilities $P_{\uparrow|\downarrow}$ and $P_{\uparrow|\downarrow}$ as a function of the evolution time t_{fin} , obtained in the experiment. The competing effects of *z*-QPMs and dissipation lead to a nontrivial dynamics, affecting the energy fluctuation distribution. To quantitatively support that the considered two-



FIG. 4. Statistics of the energy variation under different protocols: Probability to obtain $\Delta E = 0$, $+\hbar\omega$, $-\hbar\omega$, respectively, as a function of the number of laser pulses N_L experienced by the qubit, with initial probability $P_{\uparrow} = 1/(1 + e)$. For each considered N_L value, the experimental points represent the average over ~10 different values of final evolution times t_{fin} . The error bars are due to the uncertainty on the measured photoluminescence intensity. The solid lines are the joint probabilities obtained from the numerical simulation of the system dynamics.

level model provides an accurate description of the system dynamics, we performed a numerical Monte Carlo simulation of the dynamics (Sec. S.II in [38]) and we found excellent agreement with data. Note that the only fit parameter is the absorption probability, which depends on the laser power and characterizes the stochasticity of the protocol. In the absence of laser pulses, the spin qubit is a closed system and the energy eigenstates do not evolve in time (usually referred to as *spin lock*), while the absorption of laser pulses produce discrete energy jumps.

The energy variation occurred to the qubit after the process can assume one of the three values $\Delta E \in \{-\hbar\omega, 0, +\hbar\omega\}$. Figure 4 shows the distribution of energy variation $(P_{\Delta E=0},$ $P_{\Delta E=+\hbar\omega}$, and $P_{\Delta E=-\hbar\omega}$), for a fixed initial mixed state when varying the value of α and the power of the laser pulses. This result shows that in the presence of z-QPMs and dissipation, the energy distribution of the quantum system is modified. The system jumps between states with different coherences in the energy basis—as sketched in Fig. 2(c)—and finally reaches, for a large number of z-QPMs, an out-of-equilibrium steady state, which does not depend on the initial state. The photon absorption probability dictates how fast the system approaches the asymptotic steady state. Then, the final projective energy measurement returns, on average, a mixed state defined by the balance between the energy variation due to *z*-QPMs applied to the system and the dissipation channel.

For ideal equally spaced *z*-QPMs (perfect absorption, and only spin-preserving optical transitions) the asymptotic probability to find the spin in the $|\uparrow\rangle$ state $P^{\infty}_{\uparrow} = P^{\infty}_{\uparrow|\downarrow} = P^{\infty}_{\uparrow|\downarrow}$

can be analytically computed by modeling the spin temporal evolution with a master equation in the Lindblad formalism, yielding

$$P_{\uparrow}^{\infty} = \frac{1}{2} \left[1 - \frac{(1 - e^{-t_L \Gamma_{\mathcal{D}}}) \cos\alpha}{1 - e^{-t_L \Gamma_{\mathcal{D}}} \mu(\alpha, \tau)} \right], \tag{3}$$

with $\mu \equiv 1 - 2 (\sin \alpha \sin \frac{\omega \tau}{2})^2$ (Sec. S.II in Ref. [38]). Given the experimental dissipation rate Γ_D , the analytic prediction of P^{∞}_{\uparrow} matches the numerical simulations for ideal equally spaced *z*-QPMs. In the experiment, the stochasticity of the temporal distribution of *z*-QPMs—induced by the finite photon absorption—removes the strong dependence on τ (Sec. S.II in Ref. [38]). The analytical model is still a good approximation of the system dynamics, provided one replaces τ with an effective interpulse spacing (see Fig. IV.S in Ref. [38]).

In the absence of dissipation ($\Gamma_{\mathcal{D}} = 0$), z-QPMs bring the system into an equilibrium thermal state with infinite temperature [39,40] ($P^{\infty}_{\uparrow} = 1/2$). The cases $\alpha = \{0, \pi\}$ and $\tau = 2\pi/\omega$, i.e., $\mu = 1$, are exceptions [41]. The same asymptotic probability $P^{\infty}_{\star} = 1/2$ is observed also in presence of dissipation, when z-QPMs and amplitude damping act in a direction orthogonal to the Hamiltonian ($\alpha = \pi/2, \Gamma_D > 0$), as experimentally confirmed [see Fig. 3(a)]. Indeed for $\alpha = \pi/2$ the asymptotic state before the final energy measurement is a fully coherent state in the Hamiltonian basis such that $\langle \mathcal{H} \rangle =$ 0, thus the density matrix after the final energy measurement corresponds, in average, to a completely mixed state (that is, a thermal state with infinite effective temperature). In this configuration ($\alpha = \pi/2$), during a TPM protocol, dissipative dynamics is indistinguishable from unitary dynamics plus repeated measurements ($\Gamma_{\mathcal{D}} = 0$) [34].

IV. EXCHANGE FLUCTUATION RELATION

For two macroscopic systems S_1 and S_2 placed in thermal contact for a finite lapse of time, the statistics of exchanged heat Q is known to be described by the heat exchange fluctuation relation $\langle e^{-(\beta_{in,S_1}-\beta_{in,S_2})Q} \rangle = 1$, where β_{in,S_1} and β_{in,S_2} are the initial inverse temperatures of S_1 and S_2 and Q = $\Delta E_{S_1} = -\Delta E_{S_2}$ [20]. For thermalizing quantum dynamics, the non trivial value of ε for which Eq. (1) is verified is $\varepsilon = \Delta \beta \equiv \beta_{\rm in} - \beta_{\infty}$, where $\beta_{\rm in}$ and β_{∞} are the initial and asymptotic inverse temperatures of the system [31]. These two results are equivalent in the case of a quantum system interacting with a thermal reservoir at inverse temperature β_{∞} . However, the interaction with a nonthermal reservoir calls for a deeper understanding of the role played by the energy scale factor ε . With this aim, we implemented a numerical simulation to show (see Appendix 2) that, for a three-level quantum system asymptotically approaching a steady state in the energy basis (SSE), there exists one single timeindependent $\varepsilon \neq 0$ such that $\langle \exp(-\varepsilon \Delta E) \rangle = 1$. The system is brought into this steady state with constant energy in time, which does not depend on the initial state, by means of a dissipation channel that is modeled as a decay induced by spontaneous emission. After the second energy measurement of the TPM protocol, the quantum system is in a mixed state that is not necessarily thermal. This means that the effects of the dissipative channel on a quantum system may not be modeled by the effective interaction of the system with a thermal reservoir. However, also in this case, the validity of Eq. (1) is ensured and, specifically, related to the existence of a unique nontrivial constant value of ε when the system is at the SSE. It is worth noting that ε depends on the populations of the density matrix of both the initial state and the steady state. In the special case that the mixed state after the TPM protocol is thermal at inverse temperature β_{∞} , the fluctuation relation $\langle \exp(-\Delta\beta\Delta E)\rangle = 1$ is recovered [31]. The above considerations show that quantum fluctuation relations hold also in the steady-state regime [21], while in the transient regime a model-dependent behavior is expected. We observed that the energy scale ε does not depend on the nonunitary map defined by the applied protocol, but only depends on the asymptotic steady state.

In the experiment, the initial state is described by a density matrix diagonal in the energy basis and, since the system is effectively two-dimensional, can be written as a thermal state with an effective inverse temperature $\beta_{in}^{(eff)}$. Similarly, the stationary mixed state after the second energy measurement of the TPM protocol is, in average, equivalent to a thermal state with effective inverse temperature $\beta_{\infty}^{(eff)}$ [see Eq. (A7) of Appendix 3]. Notice that for each single realization and before the second energy measurement, the state evolves in time even in the asymptotic limit, marking a difference between dissipative and thermalizing dynamics for a two level system. Figure 5 shows that the experimental data and simulation always verify the relation

$$\langle \exp\left(-\varepsilon\Delta E\right)\rangle = 1,$$
 (4)

with

$$\varepsilon = \beta_{\rm in}^{\rm (eff)} - \beta_{\infty}^{\rm (eff)} \equiv \Delta \beta^{\rm (eff)},$$
 (5)

irrespective of the initial state and the applied protocol, i.e., relative orientation between the *z*-QPM operator and the system Hamiltonian, interpulse time intervals and photoabsorption probability. We emphasize that, albeit our measurements are done for a quantum two-level systems and therefore can be interpreted in terms of effective temperature, the formalism and the conclusion are expected to hold for a generic quantum system, including multilevel systems, as predicted by the numerical example in the Appendix 2.

V. CONCLUSIONS

We explored the quantum exchange fluctuation relation for an open quantum system coupled to a tunable dissipative channel. We investigated the interplay between quantum projective measurements and a dissipation channel, and we proposed a formulation of the energy exchange fluctuation relation, where the energy scaling factor depends on the populations of the stationary density matrix reached by the open quantum system. We showed that this formulation holds also for a three-level system that asymptotically reaches a steady state in the energy basis, suggesting that this result might be extended to a general finite-dimensional system. For the implemented two-level system, the energy scaling factor can be formulated in terms of the effective temperatures of the initial and final states of the system. At the steady state, the final effective temperature is indeed an invariant



FIG. 5. Experimental values of $\langle e^{-\Delta\beta^{(\text{eff})}\Delta E} \rangle$ [obtained as in Eq. (A6) in Appendix 2] as a function of the number of laser pulses N_L . Each dataset represents a different combination between the parameters, the angle α [see Eq. (2)], the time τ between laser pulses, and the photon-absorption probability p_{abs} . The top-to-bottom order in the legend symbols corresponds to the left-to-right one in the plots. These data constitute the experimental verification of the exchange fluctuation relation for an open quantum system Eq. (4). The initial probability is set to $P_{\uparrow} = 1/(1 + e)$, while the asymptotic probability P_{\uparrow}^{∞} is acquired from experimental data, as described in the Appendix 2. Both quantities define the value of $\Delta\beta^{(\text{eff})}$.

quantity, irrespective of the initial state. We have shown that this exchange fluctuation relation holds for any direction, with respect to the Hamiltonian, along which the intermediate quantum projective measurements are applied. In addition, we have observed that the exchange fluctuation relation is robust against the presence of randomness in the time intervals between measurements, as theoretically predicted in Ref. [35]. Our experimental study is enabled by the use of a single NV center in diamond at room temperature. We exploit the high control on the spin degrees of freedom, under the effect of trains of short laser pulses that perform quantum projective measurements and controllably open the two-level system, through a dissipation channel whose interaction coupling with the external surroundings can be tuned. This work, therefore, exploits NV centers in diamond as a quantum simulator to explore the physics of an out-of-equilibrium open quantum system, and to verify quantum fluctuation relations.

Our work paves the way for the investigation of Jarzynskilike equalities for general open quantum systems beyond two-dimensional Hilbert spaces. In addition, our results consolidate NV centers as a suitable platform to study phenomena related to open quantum systems, for example, to study the role of coherence in energy transport, or to experimentally verify different quantum fluctuation relations (QFR), such as the QFR for engines or refrigerators [42], or the so-called generalized QFR [43]. Finally, we hope that our work will stimulate further research to experimentally test our findings with other physical realizations, ranging from ions [26] to superconducting devices [29,44], and to ultracold gases [45]. Furthermore, our results can contribute to the implementation of heat engines working in out-of-equilibrium regimes [14].

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APPENDIX: METHODS

1. Experimental platform and modeling

We used a single NV center hosted in an electronic grade diamond sample, with 14-N concentration <5 ppb (Element Six). The NV center is optically addressed at ambient conditions with a home-built confocal microscope and its electronic spin is manipulated via resonant microwave driving. The NV center is chosen to be free from proximal 13-C nuclear spins. The 14-N spin is polarized due to a static bias magnetic field of 394 G, combined with electronic spin pumping [46]. The long coherence time of the nuclear spin ensures that it remains unaffected during the experiment. A microwave coherently manipulate the effective two-level system, composed by the $m_s = 0$ and $m_s = +1$ levels of the ground state. On the experiment timescales ($\sim \mu s$) spin-lattice relaxation is negligible ($T_1 \sim$ ms), while the Rabi driving prevents spin dephasing to occur [47].

The absorption of 532-nm laser light pulses excites the NV-center electronic spin from the ground to the excited triplet states. The decay involves (i) radiative transitions to the ground state, spin-preserving ($\sim 96.5\%$, see Sec. S.I in Ref. [38]), generating a red-shifted photoluminescence with zero-phonon line at 637 nm and (ii) nonradiative transitions through a singlet metastable state. Thus, the interaction with short laser pulses has a probability $(1 - p_{diss})$ to result in an ideal z-QPM, but also a finite probability ($p_{diss} < 1$) to destroy the state and force the resulting state to be $|0\rangle$, this process gives origin to the dissipative dynamics. Even when we cannot completely isolate each of these two effects, by changing the short laser pulses duration and intensity, we can control the value of $p_{\rm diss}$ to be between $\simeq 0.44$ and 1. The photodynamics of the NV center is thus well described with a seven-level model [32,48]. However, our experiments can be well reproduced by an effective two-level model. The neglected photodynamics occurring through the hidden physical states is re-absorbed through an effective photon absorption probability p_{abs} , an effective dissipation probability $p_{abs}\Gamma_{D}t_{L}$ and a correction on the *z*-QPM outcome that takes into account the nonspin conserving probability. In this regard, the simulations shown in the main text were realized by using a two-level system with absorption probability in the range 18–68 %, and 44% conditional probability to move populations to $|0\rangle$ once projected in $|+1\rangle$. The results of the analysis with a seven-level model and its comparison with the two-level one are reported in Sec. S.II in Ref. [38].

2. Energy steady-state regime enabling fluctuation relations: A three-level system case-study

From the analysis of the experimental data we have observed the connection between the stationary state SSE of a two-level quantum system and the existence of a unique, finite, time-independent value of ε , obeying the fluctuation relation $G(\varepsilon) = \langle \exp(-\varepsilon \Delta E) \rangle = 1$. In this section, we extend this result to to an exemplary case of a three-level system (3LS) subjected to dissipative dynamics.

Specifically, we have chosen a 3LS governed by the following Hamiltonian:

$$H = \omega_{12}(|1\rangle\langle 2| + |2\rangle\langle 1|) + \omega_{13}(|1\rangle\langle 3| + |3\rangle\langle 1|)$$
$$= \begin{pmatrix} 0 & \omega_{12} & \omega_{13} \\ \omega_{12} & 0 & 0 \\ \omega_{13} & 0 & 0 \end{pmatrix},$$
(A1)

where $|j\rangle$ denotes the *j*th level of the system and ω_{12} , ω_{13} are the coupling rates between the first level and the second and the third ones. The system is also characterized by a decay channel (mimicking a spontaneous emission process) with rate Γ between the first and second level of the 3LS. We describe the system dynamics by means of a Lindblad master equation, whereby the spontaneous decay is given by the following Lindbladian super-operator:

$$L = \sqrt{\Gamma} |1\rangle\langle 2| = \begin{pmatrix} 0 & \sqrt{\Gamma} & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix}.$$
 (A2)

Note that, albeit no intermediate QPMs are applied to the quantum system, the dynamics of the 3LS is dissipative due to the presence of a spontaneous decay term, here modelled by means of the Lindbladian formalism.

Under the hypothesis that the system reaches a steady state in its energy basis, the conditional probabilities $P_{j|i}$ are invariant with respect to the initially measured energy values obtained from the TPM scheme. This means that

$$P_{j|1} = P_{j|2} = P_{j|3} \equiv P_j \tag{A3}$$

for any j = 1, 2, 3, irrespective of the initial state ρ_0 and the transient properties of the dissipative dynamics acting on the 3LS. The validity of Eq. (A3) implies the following decomposition for the mean exponentiated energy variation $\langle \exp(-\eta \Delta E) \rangle$:

$$\langle \exp(-\eta \Delta E) \rangle = \left(\sum_{j=1}^{3} \widetilde{P}_{j} e^{-\eta E_{j}}\right) \left(\sum_{i=1}^{3} P_{i} e^{\eta E_{i}}\right),$$
 (A4)



FIG. 6. Exchange fluctuation relation for a three-level model subjected to spontaneous decay processes. (a) Plot of $g(\eta) \equiv G(\eta) - 1$ as a function of $\eta \in [0, 1]$ and $\omega_{13}/\omega_{12} \in [0, 3]$ at the steady state in the energy basis. (b) ε as a function of time $t \in [0, 80]$ for four different values of ω_{13}/ω_{12} , i.e., $\omega_{13}/\omega_{12} = \{0.21, 0.41, 0.91, 1.51, 2.28\}$, corresponding respectively to the green solid, red dashed, black dash-dotted, blue dotted, and magenta dashed lines. Inset: Time behavior of the probabilities P_{f_j} to measure the energy of the system at the final time instant of the TPM protocol, by taking $\omega_{13} = 0.21$ and $\omega_{12} = 1$. All the parameters used to numerically compute the system dynamics, namely ω_{13} , $\omega_{12} = 1$, $\Gamma = 1.5$, $\beta = 1$ with $\rho_0 = \exp(-\beta H)/\mathrm{Tr}[\exp(-\beta H)]$, are in units of $\hbar = 1$.

where $\{E_i\}$ and $\{E_j\}$ are, respectively, the set of measured energy at the initial and final time instants of the TPM scheme, while P_i denotes the probability of measuring E_i . We have numerically computed the system dynamics with a time duration long enough to ensure that the system has reached a steady state in the energy basis for each set of system parameters. In Fig. 6(a), we have plotted $g(\eta) \equiv G(\eta) - 1$ as a function of η and ω_{13}/ω_{12} at the SSE, by fixing the value of the other parameters (see caption). One can observe that, for each value of ω_{13} ($\omega_{12} = 1$), $g(\eta)$ always has two zeros, where one zero is always $\eta_0 = 0$. Instead, regarding the other zero of $g(\eta)$, in the numerical simulations it takes a unique time-independent value $\eta = \varepsilon$ that depends only on the initial state ρ_0 and the asymptotic steady state.

Finally, to conclude our analysis, we have fixed five different values of ω_{13} (with $\omega_{12} = 1$) and plotted ε [value of η such that $G(\eta) = 1$] as a function of time. In Fig. 6(b) one can observe that, as the time increases, ε becomes a constant (time-independent) value, depending on the specific steady-state reached by the 3LS. The same behavior occurs also by varying the other dynamics parameters, e.g., Γ and β .

3. Two-level systems: Energy variation distribution and asymptotic effective temperature

Measuring the conditional probabilities $P_{\uparrow|\uparrow}$ and $P_{\uparrow|\downarrow}$ gives access to the full statistics of ΔE

$$P_{\Delta E} \equiv \operatorname{Prob}(\Delta E) = \sum_{i,j} \delta(\Delta E - \Delta E_{i,j}) P_{j|i} P_i, \qquad (A5)$$

even without directly measuring the output energy for each experiment realization. The latter, indeed, cannot be achieved due to low photon collection efficiency from the NV center. In the previous equation P_i denotes the probability of measuring E_i at the beginning of the TPM protocol, $P_{j|i}$ is the conditional probability of measuring E_j at the end of the protocol, and the sum is performed over all the possible initial *i* and final *j* measured energies. Once obtained the distribution of the energy variation, we operatively evaluate the mean value of the exponentiated energy fluctuations of the two-level system as

$$\langle e^{-\Delta\beta^{(\text{eff})}\Delta E} \rangle = e^{-\Delta\beta^{(\text{eff})}\hbar\omega} P_{\hbar\omega} + P_0 + e^{\Delta\beta^{(\text{eff})}\hbar\omega} P_{-\hbar\omega}, \quad (A6)$$

where $\Delta \beta^{\text{(eff)}} = \beta_{\text{in}}^{\text{(eff)}} - \beta_{\infty}^{\text{(eff)}}$. For a two-level mixed state $\varrho^{\text{mix}} = p_{\uparrow} \varrho_{\uparrow} + (1 - p_{\uparrow}) \varrho_{\downarrow}$, an effective inverse temperature

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is defined as

$$\beta^{\text{(eff)}}(p_{\uparrow}) = \frac{2}{\hbar\omega} \arctan(1 - 2p_{\uparrow}). \tag{A7}$$

This picture of effective temperature does not capture the possible population inversion, and is therefore valid only for the half of the Bloch sphere containing the state at lower energy. In the experiment, $\beta_{in}^{(eff)}$ is defined by the choice of the initial mixed state. The asymptotic inverse temperature $\beta_{\infty}^{(eff)}$ is extracted from the experimental data after a large enough number of *z*-QPMs, whereby $P_{\uparrow\uparrow\uparrow}^{\infty} = P_{\uparrow\downarrow\downarrow}^{\infty} \equiv P_{\uparrow}^{\infty}$, thus ensuring the stationarity of the system final state. Otherwise, for low photon-absorption probability, the asymptotic temperature can be extracted by finding the initial state for which $\langle \Delta E \rangle = 0$ during the whole evolution.

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