

Fluctuations in Extractable Work Bound the Charging Power of Quantum Batteries

Luis Pedro García-Pintos^{1,2,*}, Alioscia Hama^{2,†}, and Adolfo del Campo^{3,4,2,‡}

¹*Joint Center for Quantum Information and Computer Science and Joint Quantum Institute, NIST/University of Maryland, College Park, Maryland 20742, USA*

²*Department of Physics, University of Massachusetts, Boston, Massachusetts 02125, USA*

³*Donostia International Physics Center, E-20018 San Sebastián, Spain*

⁴*IKERBASQUE, Basque Foundation for Science, E-48013 Bilbao, Spain*



(Received 19 September 2019; accepted 8 July 2020; published 22 July 2020)

We study the connection between the charging power of quantum batteries and the fluctuations of the extractable work. We prove that in order to have a nonzero rate of change of the extractable work, the state ρ_W of the battery cannot be an eigenstate of a “free energy operator,” defined by $\mathcal{F} \equiv H_W + \beta^{-1} \log(\rho_W)$, where H_W is the Hamiltonian of the battery and β is the inverse temperature of a reference thermal bath with respect to which the extractable work is calculated. We do so by proving that fluctuations in the free energy operator upper bound the charging power of a quantum battery. Our findings also suggest that quantum coherence in the battery enhances the charging process, which we illustrate on a toy model of a heat engine.

DOI: [10.1103/PhysRevLett.125.040601](https://doi.org/10.1103/PhysRevLett.125.040601)

The study of the thermodynamics of quantum systems can be traced back to the beginnings of quantum theory, but a recent surge in interest has been spurred by major advances [1] in which the use of quantum information theoretic tools plays a key role [2–8]. Despite this progress, fundamental issues continue to be a matter of debate. Prominent examples include the notion of thermodynamical work in the quantum realm [9–13], and the role of quantum coherences and their potential use as a thermodynamical resource [14–21].

In this context, the study of heat engines and quantum machines has proven useful, helping to illustrate the capabilities and limits of quantum devices [22–31]. The analysis of these devices has established a trade-off between work fluctuations and dissipation in heat engines [32,33], an increase in the charging power [34–37] and decrease in fluctuations [38] with collective operations acting on storage devices in parallel, the advantage on charging power over many cycles of a heat engine [39] and of indistinguishable heat engines [40], as well as an increased efficiency when considering correlated thermal machines [41]. On the other hand, it has also been shown that entanglement is not indispensable for work extraction [42], and that within the set of Gaussian operations to charge a battery there are inevitably fluctuations in the stored work [43].

Along parallel lines, Mandalstam and Tamm considered the limits that quantum mechanics imposes on the rate of quantum evolution [44]. By considering the minimum time necessary for a state to evolve to an orthogonal one, their work inspired a wide range of results [45–54], that are usually encompassed under the term of quantum speed limits to evolution [55].

In quantum thermodynamics, speed limits were first discussed to bound the output power of heat engines [27]. They are known to govern work and energy fluctuations in superadiabatic processes [33,56,57] and can be used to analyze advantages from many-particle effects in quantum batteries [34,36]. Articles studying the limits to increasing the energy of batteries in unitary protocols appeared recently [58–61]. In Ref. [6], general bounds on the efficiency of an engine and the extractable work are derived as a function of the work fluctuations, for a battery modeled by a weight that starts and ends in classical (diagonal) states.

In this Letter, we investigate the fundamental limits that quantum mechanics imposes on the process of charging a battery, where we take a battery to be any system that can store energy for subsequent extraction. We show that in order to have a nonzero charging power there must exist fluctuations of a free energy operator that characterizes the extractable work in the battery. In particular, we provide bounds on the charging power that (a) explicitly take into consideration the entropic aspects of the amount of extractable work as opposed to limiting the study to energy storage, and (b) are valid for arbitrary states and a wide range of protocols, including those based on unitary evolution as well as protocols involving open-system dynamics, and hold regardless of the nature of the battery, its state, and the properties of the extractable work of the battery.

In a work extraction protocol any system out of thermal equilibrium is a resource, from which work can be extracted if one has access to a thermal bath. If the resource system is in state ρ , and τ_β denotes the thermal state at

inverse temperature β , the maximum amount of work that can be extracted from the resource state ρ is given by

$$W_{\max} = F(\rho) - F(\tau_\beta), \quad (1)$$

where the free energy is $F(\rho) = U - S/\beta$, and U and $S = -\text{Tr}(\rho \ln \rho)$ are the average energy and von Neumann entropy of the system, respectively [3,4,62,63]. We refer to W_{\max} as the ‘‘extractable work.’’ In Ref. [3] this is shown to be the case, on average, in the limit of asymptotically many copies of ρ , while Ref. [62] constructs a protocol that works for a single copy of ρ , albeit with nonzero fluctuations in the extracted work and with a toy model as a work reservoir. Interestingly, Ref. [63] proves that, if the resource system is allowed to become correlated to catalytic systems, then W_{\max} exactly determines the extractable work in single-shot scenarios as well. It is worth clarifying that the notion of work used in Refs. [3,4,62,63] is an operational one, in which the extracted work is stored in reservoirs, for posterior extraction. Specifically, Refs. [3,63] show that an amount of energy equal to the extractable work can be used to excite two-level systems from the ground state, while in Refs. [4,62] it is used to raise the energy of an (unbounded) harmonic oscillator. Importantly, these protocols also provide a mechanism to extract the same amount of energy from the reservoirs, thus providing an operational way in which work is defined in this optimal-protocol context (albeit with the possible need of auxiliary resources, such as a system that serves as entropy sink in Ref. [63]).

In this Letter we focus on the limitations that quantum mechanics imposes on any process of storing or extracting the extractable work, as illustrated in Fig. 1.

Isolated system analysis.—A protocol to extract work from a system and store it in a battery necessarily couples, at some point, the battery with the bath, the system, and/or auxiliary ancillas \mathcal{A} used in order to implement the necessary operations. For instance, ancillas can be used to describe an arbitrary map acting on the system. We denote the system used as a work reservoir, i.e., the *battery*, by \mathcal{W} , and its Hamiltonian by $H_{\mathcal{W}}$. We assume that the total system, consisting of all of the above, is isolated and evolves unitarily with a total Hamiltonian

$$H = H_{SBA} \otimes \mathbb{1}_{\mathcal{W}} + \mathbb{1}_{SBA} \otimes H_{\mathcal{W}} + V, \quad (2)$$

where H_{SBA} accounts for all terms that correspond to bath, system, and ancilla evolution, V incorporates all terms corresponding to interactions with the battery, and $\mathbb{1}_{\{\cdot\}}$ denotes identities on the subspaces indicated by the subindex. Note that, while we allow H_{SBA} and V to be explicitly time dependent, we assume $H_{\mathcal{W}}$ to be time independent, motivated by the idea of having a battery whose physical characteristics, e.g., Hamiltonian, do not change. Then, the full state ρ of $SBA\mathcal{W}$ evolves according to

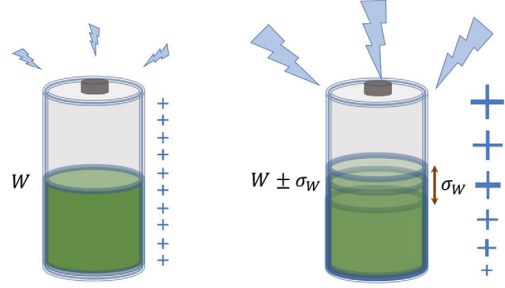


FIG. 1. Fluctuations in extractable work limit the charging power of batteries. A quantum battery is a system that follows the laws of quantum physics, and that can store energy to be extracted later in order to perform work. The most important quantity in a battery is the maximum amount of work W that can be extracted from it, i.e., the extractable work, which in certain cases is directly characterized by the free energy. However, due to the quantum nature of the battery, this free energy is a random quantity with uncertainties. We prove that the rate at which any battery can be charged is linked to such fluctuations, via fundamental bounds that hold for any charging protocol. For a battery with a well-defined value of the extractable work W , depicted on the left, the maximum rate at which the extractable work changes is zero. Fluctuations allow for faster charging rate, as illustrated on the right.

$$\frac{d\rho}{dt} = -i[H, \rho]. \quad (3)$$

For ease of notation we omit the explicit time dependence of the evolving state ρ .

A central quantity behind the results of this Letter is the free energy operator \mathcal{F} characterizes the amount of work that can be extracted from the battery, as from Eq. (1) it follows that

$$\mathcal{F} \equiv H_{\mathcal{W}} + \beta^{-1} \log \rho_{\mathcal{W}}, \quad (4)$$

where $\rho_{\mathcal{W}} = \text{Tr}_{SBA}(\rho)$ is the state of the battery. The free energy operator \mathcal{F} characterizes the amount of work that can be extracted from the battery, as from Eq. (1) it follows that

$$W_{\max} = \langle \mathcal{F} \rangle_{\mathcal{W}} - \langle \mathcal{F} \rangle_{\beta}, \quad (5)$$

where $\langle \mathcal{F} \rangle_{\mathcal{W}} \equiv \text{Tr}(\mathcal{F} \rho_{\mathcal{W}})$ and $\langle \mathcal{F} \rangle_{\beta} \equiv \text{Tr}(\mathcal{F} \tau_\beta)$. Note that, while \mathcal{F} is a Hermitian operator, it depends on the state of the battery and it need not correspond to a physical observable. It is also worth stressing, though, that \mathcal{F} is distinctively different to the process-dependent ‘‘work operator’’ considered in Refs. [64–66]. Unlike the work operator, \mathcal{F} corresponds to a state function, and its average characterizes the *extractable work* [3,4,62,63].

The rate at which the battery’s extractable work changes, that we refer to as the charging power, is given by

$$P(t) = \frac{dW_{\max}}{dt} = \frac{d\langle \mathcal{F} \rangle_{\mathcal{W}}}{dt} \quad (6)$$

$$= \frac{d\text{Tr}(\rho_{\mathcal{W}} H_{\mathcal{W}})}{dt} - \frac{1}{\beta} \frac{dS(\rho_{\mathcal{W}})}{dt}, \quad (7)$$

where we used that $H_{\mathcal{W}}$ is time independent. As proven in Refs. [67,68], it holds that $dS(\rho_{\mathcal{W}})/dt = -i\text{Tr}(V[\log \rho_{\mathcal{W}} \otimes \mathbb{1}_{SBA}, \rho])$. Note that $P(t)$ limits the power needed in realistic charging protocols. Specifically, the energy E needed to increase a battery's extractable work by $\Delta W \equiv W_{\max}(t) - W_{\max}(0)$ in realistic charging protocols is bounded by the charging power, as $E \geq \Delta W = \int_0^t P(t') dt'$.

Using Eqs. (2) and (3), it further follows that $d\text{Tr}(\rho_{\mathcal{W}} H_{\mathcal{W}})/dt = -i\text{Tr}([\rho, H_{\mathcal{W}} \otimes \mathbb{1}_{SBA}]V)$. As a result,

$$\begin{aligned} P(t) &= -i\text{Tr}([\rho, H_{\mathcal{W}} \otimes \mathbb{1}_{SBA}]V) \\ &\quad - i \frac{1}{\beta} \text{Tr}([\rho, \log \rho_{\mathcal{W}} \otimes \mathbb{1}_{SBA}]V) \\ &= -i\text{Tr}([\rho, \mathcal{F} \otimes \mathbb{1}_{SBA}]V). \end{aligned} \quad (8)$$

At a qualitative level, this simple derivation suggests that coherence in the eigenbasis of the free energy operator—which in turn necessitates coherence in the energy eigenbasis—serves to enhance the charging process. Indeed, if one considers a battery initially uncorrelated from other systems, $\rho = \rho_{\mathcal{W}} \otimes \rho_{SBA}$, the charging power is zero unless the state is coherent in the eigenbasis of \mathcal{F} .

Our main result sets bounds on the charging power. Defining $\delta\mathcal{F} \equiv \mathcal{F} - \langle \mathcal{F} \rangle_{\mathcal{W}}$ and $\delta V = V - \langle V \rangle$, it holds that

$$\begin{aligned} P^2(t) &\leq |\text{Tr}(\rho[\delta\mathcal{F} \otimes \mathbb{1}_{SBA}, \delta V])|^2 \leq 4|\text{Tr}(\rho\delta\mathcal{F}\delta V)|^2 \\ &\leq 4\text{Tr}(\rho(\delta\mathcal{F})^2 \otimes \mathbb{1}_{SBA})\text{Tr}(\rho(\delta V)^2), \end{aligned} \quad (9)$$

where we use that $H_{\mathcal{W}}$ is time independent and denote the averages of the extractable work and battery interaction energy by $\langle \mathcal{F} \rangle_{\mathcal{W}} = \text{Tr}(\rho_{\mathcal{W}} \mathcal{F})$ and $\langle V \rangle \equiv \text{Tr}(\rho V)$, respectively. For the last step of the calculation we used the fact that for Hermitian operators $A > 0$, B and C , the Cauchy-Schwarz inequality implies that $\text{Tr}^2(ABC) = \text{Tr}^2(\sqrt{A}BC\sqrt{A}) \leq \text{Tr}(AB^2)\text{Tr}(AC^2)$. Denoting the standard deviations of \mathcal{F} and of the battery interaction Hamiltonian by $\sigma_{\mathcal{F}}$ and σ_V , respectively,

$$\sigma_{\mathcal{F}}^2(t) \equiv \langle \mathcal{F}^2 \rangle_{\mathcal{W}} - \langle \mathcal{F} \rangle_{\mathcal{W}}^2, \quad (10)$$

$$\sigma_V^2(t) \equiv \langle V^2 \rangle - \langle V \rangle^2, \quad (11)$$

Equation (9) implies

$$|P(t)| \leq 2\sigma_{\mathcal{F}}(t)\sigma_V(t). \quad (12)$$

This proves the existence of a trade-off between charging power and the fluctuations of the free energy operator \mathcal{F} : for

a fixed interaction with the battery, a desired power input necessarily comes with fluctuations of the operator \mathcal{F} whose mean characterizes the extractable work of the battery. In contrast, attempting to charge a battery with a deterministic amount of extractable work, such that $\sigma_{\mathcal{F}} = 0$, leads to a null instantaneous charging power. It is straightforward to see that this implies, for instance, zero charging power for batteries in eigenstates of energy. We stress that this bound applies, in particular, to protocols that charge the battery via unitary evolution with a time-dependent perturbation Hamiltonian $V(t)$ [31,34–36,58–60].

Open system analysis.—Evaluating bound (12) for charging protocols involving unitary, time-dependent, control of the battery is straightforward, as it solely involves knowledge of the state of the battery, its Hamiltonian $H_{\mathcal{W}}$, and the control Hamiltonian V . However, evaluating the factor $\sigma_V(t)$ may be hard for protocols that involve contact of the battery with secondary systems, as evaluating it requires specifying the full state ρ of the battery and all the systems it interacts with. In the light of this, it is of practical importance to extend the analysis to encompass open-system descriptions of the dynamics of the battery, which we do next.

The formalism introduced so far allows us to extend the analysis to include an open-system description of the dynamics of the battery. From Eq. (3), the state of the battery evolves according to

$$\begin{aligned} \frac{d}{dt}\rho_{\mathcal{W}} &= -i\text{Tr}_{SBA}([H, \rho]) \\ &= -i[H_{\mathcal{W}}, \rho_{\mathcal{W}}] - i\text{Tr}_{SBA}([V, \rho]). \end{aligned} \quad (13)$$

In the Markovian and weak-coupling limits the evolution of a system interacting with an environment is well approximated by

$$\begin{aligned} \frac{d}{dt}\rho_{\mathcal{W}} &\approx -i[H_{\mathcal{W}}, \rho_{\mathcal{W}}] - i[\tilde{H}_{\mathcal{W}}, \rho_{\mathcal{W}}] \\ &\quad + \sum_j \gamma_j \left(L_j \rho_{\mathcal{W}} L_j^\dagger - \frac{1}{2} \{L_j^\dagger L_j, \rho_{\mathcal{W}}\} \right), \end{aligned} \quad (14)$$

where $\tilde{H}_{\mathcal{W}}$ accounts for the unitary part of the evolution due to the interactions, the Lindblad operators L_j characterize the nonunitary effect of the interaction of the battery with the remaining systems, and the rates γ_j are non-negative [69]. Then,

$$\begin{aligned} \text{Tr}_{SBA}([V, \rho]) &\approx [\tilde{H}_{\mathcal{W}}, \rho_{\mathcal{W}}] \\ &\quad + i \sum_j \gamma_j \left(L_j^\dagger \rho_{\mathcal{W}} L_j - \frac{1}{2} \{L_j^\dagger L_j, \rho_{\mathcal{W}}\} \right). \end{aligned} \quad (15)$$

With this, we prove in the Supplemental Material [70] that the rate at which the extractable work of the battery changes is upper bounded by

$$|P(t)| \leq 2\sigma_{\mathcal{F}}(t)\sigma_{\tilde{H}_W}(t) + \sum_j \gamma_j \sqrt{\langle |[\delta\mathcal{F}, L_j]|^2 \rangle} \|L_j\|, \quad (16)$$

where the operator norm $\|A\|$ is given by the largest modulus of the eigenvalues of an operator A , and $|A|^2 = AA^\dagger$. This sets a bound valid for charging protocols based on open-system approaches. Importantly, the bound depends solely on the state ρ_W of the battery, and the decay rates γ_j and operators \tilde{H} and L_j , fixed by the master equation that governs the dynamics of the battery.

Remarkably, for the open-system case it also holds that the charging power is null unless there exist fluctuations in the free energy operator \mathcal{F} . In order to see this, let $|j\rangle$ denote the eigenbasis of $\delta\mathcal{F}$, with $\delta\mathcal{F} = \sum_j w_j |j\rangle\langle j|$, $\rho_{jk} = \langle j|\rho_W|k\rangle$ and $L_{jk} = \langle j|L|k\rangle$. One then finds that

$$\langle |[\delta\mathcal{F}, L_j]|^2 \rangle = \sum_{jkl} \rho_{jk} L_{kl} L_{lj}^\dagger (w_l^2 - w_l w_j - w_l w_k + w_j w_k). \quad (17)$$

As a result, for states $\rho_W = |j\rangle\langle j|$ with a deterministic amount of free energy characterized by w_j , one has that $\langle |[\delta\mathcal{F}, L_j]|^2 \rangle = 0$ and that $\sigma_{\mathcal{F}} = 0$, leading to a null charging rate $|P(t)| = 0$. By contrast, states with support on more than one eigenstate of \mathcal{F} can sustain a higher charging power, as both $\langle |[\delta\mathcal{F}, L_j]|^2 \rangle \neq 0$ and $\sigma_{\mathcal{F}} \neq 0$.

Finally, in the case of Hermitian Lindblad operators $L_j = L_j^\dagger$ we prove in the Supplemental Material [70] that the bound (16) simplifies to

$$|P(t)| \leq \sigma_{\mathcal{F}}(t) \left(2\sigma_{\tilde{H}_W}(t) + \sum_j 2\gamma_j \|L_j\|^2 \right). \quad (18)$$

Illustration with a heat engine.—We consider a minimal, self-contained, model of a heat engine that stores work in a quantum battery, as studied in detail in Refs. [71,72]. The engine consists of thermal baths at different temperatures, T_h and T_c , as a resource. Note that these baths are internal to the workings of the protocol to extract energy; in particular, the temperatures T_h and T_c are unrelated to the inverse temperature β of the reference bath with respect to which work is defined in Eq. (1).

In the engine, heat flow from the hot to the cold bath is exploited to extract work and store it in a toy-model battery that consists of a harmonic oscillator unbounded from below, with an energy gap ϵ for a Hamiltonian

$$H_W = \sum_{n=-\infty}^{\infty} n\epsilon |n\rangle_w \langle n|. \quad (19)$$

The storage device is indirectly coupled to the heat baths via a “switch,” consisting of two qubits. Qubit 1, with energy gap E_1 , is coupled to the cold bath, while qubit 2,

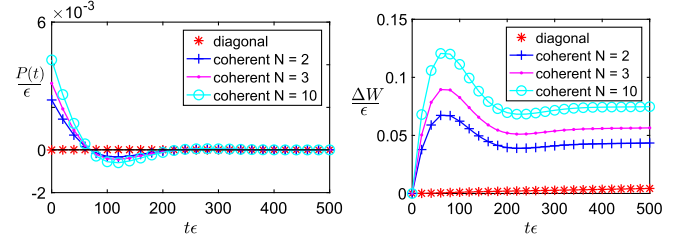


FIG. 2. Charging power and extractable work. Charging power $P(t)$ (left) and change in extractable work $\Delta W \equiv W_{\max}(t) - W_{\max}(0)$ (right) as a function of time for various initial states (for the values of the parameters used see Ref. [70]). The incoherent state ρ_{diag} (red asterisk) has null charging power initially, as expected from bound (12). This charging power is slowly increased as the state supports higher fluctuations in extractable work. A significant increase in the charging power is seen for a coherent superposition of eigenstates of \mathcal{F} . For intermediate times this initial increase is penalized with a regime in which the battery is being discharged. Nevertheless, there is a net increase in the extractable work with respect to an incoherent initial state. Taking coherent superpositions between more than two states gives a further increase in charging power and extractable work.

with energy gap E_2 , is coupled to the hot bath. The qubits have a free Hamiltonian given by $H_S = E_1|1\rangle_1\langle 1| + E_2|1\rangle_2\langle 1|$, with energies taken such that $E_2 - E_1 = \epsilon$, and they interact with the battery via

$$V = g \sum_n (|01, n\rangle\langle 10, n+1| + |10, n+1\rangle\langle 01, n|), \quad (20)$$

where g is a coupling constant. Qubits 1 and 2 are assumed to thermalize due to the interaction with the thermal baths at rates p_1 and p_2 , respectively. With these considerations, the evolution of the mean energy in the battery is solved analytically in Refs. [71,72], where it is found that choosing the parameters of the model correctly make the device work as a heat engine, storing work in W .

We consider for simplicity a reference bath of zero temperature to calculate the extractable work of the battery. For this case, we derive the equations governing the dynamics of the charging power, extractable work, and its fluctuations in the Supplemental Material [70]. We are interested in the charging power for initial states with uncertain amounts of free energy.

Consider first a state without uncertainty in the \mathcal{F} . For instance, consider qubits 1 and 2 initially in states τ_1 and τ_2 in thermal equilibrium with their respective thermal baths, and the battery in an eigenstate $|0\rangle_W$ of its Hamiltonian,

$$\rho_{\text{diag}} = \tau_1 \otimes \tau_2 \otimes |0\rangle_W \langle 0|. \quad (21)$$

Naively, this would appear to be an ideal initial state for the battery, with a well-defined deterministic initial amount of energy and extractable work, without fluctuations. However, for such a state, diagonal in both interaction

and battery Hamiltonians, Eq. (8) implies that the engine initially functions with null power. This is illustrated in Fig. 2.

In order to have nonzero charging power for product states, a coherent superposition in the free energy operator and interaction Hamiltonian is needed. Consequently, we consider both the qubits and battery in a pure state, the latter in a superposition between N energy levels $|n\rangle_{\mathcal{W}}$, with equal weights for simplicity:

$$\begin{aligned}
 |\Psi_N\rangle &= (\sqrt{r_1}|0\rangle_1 + e^{i\theta}\sqrt{1-r_1}|1\rangle_1) \\
 &(\sqrt{r_2}|0\rangle_2 + \sqrt{1-r_2}|1\rangle_2) \frac{1}{\sqrt{N}} \sum_{n=0}^{N-1} |n\rangle_{\mathcal{W}}. \quad (22)
 \end{aligned}$$

The phase $e^{i\theta}$ fixes whether the device works as an engine or a refrigerator.

With such coherent superposition as the initial state, one has $\sigma_{\mathcal{F}} \geq 0$ and $\sigma_v \geq 0$, such that inequality (8) allows us to have a nonzero charging power. This is to be compared with the null charging power of state ρ_{diag} . Figure 2 compares the charging power $P(t)$ and the change in the extractable work $\Delta W \equiv W_{\text{max}}(t) - W_{\text{max}}(0)$ as a function of time, for initial states given by ρ_{diag} and $|\Psi_N\rangle\langle\Psi_N|$, for different values of N . A superposition between eigenstates of \mathcal{F} causes an increase in the charging power, with respect to the null charging power of the incoherent state. Superpositions between more levels results in an even higher power. This is reflected in the total extractable work as well, with a considerable increase for coherent superpositions, with the most noticeable advantage achieved when going from incoherent state to $|\Psi_2\rangle\langle\Psi_2|$.

Discussion.—We have established a direct relationship between the statistics of the extractable work and the charging power of batteries: for the latter to be nonzero the extractable work of the battery has to fluctuate. To this end, we have introduced bounds to the charging power in terms of a “free energy operator” $\mathcal{F} \equiv H_{\mathcal{W}} + \beta^{-1} \log(\rho_{\mathcal{W}})$ whose expectation value equals the amount of maximum amount of work that can be extracted from the battery. Batteries in an eigenstate of \mathcal{F} suffer from a null charging power, while fluctuations of \mathcal{F} allow for higher charging rates. The bounds hold for a variety of battery charging protocols, including unitary dynamics of the battery via time-dependent perturbation Hamiltonians, isolated dynamics for a battery-system bath, as well as open-system dynamics. Our results also identify coherence as a resource in the process of work storage and should be of relevance to the engineering of quantum thermodynamic devices in a variety of platforms.

This work was funded by the John Templeton Foundation, UMass Boston Project No. P20150000029279, and DOE Grant No. DE-SC0019515. L. P. G. P. also acknowledges partial support by AFOSR MURI project “Scalable

Certification of Quantum Computing Devices and Networks,” DOE ASCR Accelerated Research in Quantum Computing program (Award No. DE-SC0020312), DOE BES Materials and Chemical Sciences Research for Quantum Information Science program (Award No. DE-SC0019449), DOE ASCR Quantum Testbed Pathfinder program (Award No. DE-SC0019040), NSF PFCQC program, AFOSR, ARO MURI, ARL CDQI, and NSF PFC at JQI.

*Corresponding author.

lpgp@umd.edu

†alioscia.hamma@umb.edu

‡adolfo.delcampo@dipc.org

- [1] J. Anders and M. Esposito, *New J. Phys.* **19**, 010201 (2017).
- [2] A. E. Allahverdyan, R. Balian, and T. M. Nieuwenhuizen, *Europhys. Lett.* **67**, 565 (2004).
- [3] F. G. S. L. Brandão, M. Horodecki, J. Oppenheim, J. M. Renes, and R. W. Spekkens, *Phys. Rev. Lett.* **111**, 250404 (2013).
- [4] J. Åberg, *Phys. Rev. Lett.* **113**, 150402 (2014).
- [5] J. Goold, M. Huber, A. Riera, L. del Rio, and P. Skrzypczyk, *J. Phys. A* **49**, 143001 (2016).
- [6] J. G. Richens and L. Masanes, *Nat. Commun.* **7**, 13511 (2016).
- [7] L. Masanes and J. Oppenheim, *Nat. Commun.* **8**, 14538 (2017).
- [8] A. M. Alhambra, G. Styliaris, N. A. Rodríguez-Briones, J. Sikora, and E. Martín-Martínez, *Phys. Rev. Lett.* **123**, 190601 (2019).
- [9] J. Åberg, *Nat. Commun.* **4**, 1925 (2013).
- [10] M. F. Frenzel, D. Jennings, and T. Rudolph, *Phys. Rev. E* **90**, 052136 (2014).
- [11] C. Jarzynski, H. T. Quan, and S. Rahav, *Phys. Rev. X* **5**, 031038 (2015).
- [12] A. M. Alhambra, L. Masanes, J. Oppenheim, and C. Perry, *Phys. Rev. X* **6**, 041017 (2016).
- [13] R. Gallego, J. Eisert, and H. Wilming, *New J. Phys.* **18**, 103017 (2016).
- [14] K. Brandner, M. Bauer, M. T. Schmid, and U. Seifert, *New J. Phys.* **17**, 065006 (2015).
- [15] M. Lostaglio, K. Korzekwa, D. Jennings, and T. Rudolph, *Phys. Rev. X* **5**, 021001 (2015).
- [16] R. Uzdin, A. Levy, and R. Kosloff, *Phys. Rev. X* **5**, 031044 (2015).
- [17] K. Korzekwa, M. Lostaglio, J. Oppenheim, and D. Jennings, *New J. Phys.* **18**, 023045 (2016).
- [18] A. Streltsov, G. Adesso, and M. B. Plenio, *Rev. Mod. Phys.* **89**, 041003 (2017).
- [19] G. Francica, J. Goold, and F. Plastina, *Phys. Rev. E* **99**, 042105 (2019).
- [20] C. L. Latune, I. Sinayskiy, and F. Petruccione, *Sci. Rep.* **9**, 3191 (2019).
- [21] G. M. Andolina, M. Keck, A. Mari, M. Campisi, V. Giovannetti, and M. Polini, *Phys. Rev. Lett.* **122**, 047702 (2019).
- [22] N. Brunner, N. Linden, S. Popescu, and P. Skrzypczyk, *Phys. Rev. E* **85**, 051117 (2012).

- [23] N. Linden, S. Popescu, and P. Skrzypczyk, *Phys. Rev. Lett.* **105**, 130401 (2010).
- [24] L. A. Correa, J. P. Palao, G. Adesso, and D. Alonso, *Phys. Rev. E* **87**, 042131 (2013).
- [25] P. Skrzypczyk, A. J. Short, and S. Popescu, *Nat. Commun.* **5**, 4185 (2014).
- [26] R. Kosloff and A. Levy, *Annu. Rev. Phys. Chem.* **65**, 365 (2014).
- [27] A. del Campo, J. Goold, and M. Paternostro, *Sci. Rep.* **4**, 6208 (2015).
- [28] R. Alicki and D. Gelbwaser-Klimovsky, *New J. Phys.* **17**, 115012 (2015).
- [29] P. P. Hofer, M. Perarnau-Llobet, J. B. Brask, R. Silva, M. Huber, and N. Brunner, *Phys. Rev. B* **94**, 235420 (2016).
- [30] W. Niedenzu, D. Gelbwaser-Klimovsky, A. G. Kofman, and G. Kurizki, *New J. Phys.* **18**, 083012 (2016).
- [31] F. Caravelli, G. Coulter-De Wit, L. P. Garcia-Pintos, and A. Hama, *Phys. Rev. Research* **2**, 023095 (2020).
- [32] K. Funo and M. Ueda, *Phys. Rev. Lett.* **115**, 260601 (2015).
- [33] K. Funo, J.-N. Zhang, C. Chatou, K. Kim, M. Ueda, and A. del Campo, *Phys. Rev. Lett.* **118**, 100602 (2017).
- [34] F. C. Binder, S. Vinjanampathy, K. Modi, and J. Goold, *New J. Phys.* **17**, 075015 (2015).
- [35] J. Jaramillo, M. Beau, and A. del Campo, *New J. Phys.* **18**, 075019 (2016).
- [36] F. Campaioli, F. A. Pollock, F. C. Binder, L. Céleri, J. Goold, S. Vinjanampathy, and K. Modi, *Phys. Rev. Lett.* **118**, 150601 (2017).
- [37] D. Ferraro, M. Campisi, G. M. Andolina, V. Pellegrini, and M. Polini, *Phys. Rev. Lett.* **120**, 117702 (2018).
- [38] M. Perarnau-Llobet and R. Uzdin, *New J. Phys.* **21**, 083023 (2019).
- [39] G. Watanabe, B. P. Venkatesh, P. Talkner, and A. del Campo, *Phys. Rev. Lett.* **118**, 050601 (2017).
- [40] G. Watanabe, B. P. Venkatesh, P. Talkner, M.-J. Hwang, and A. del Campo, *Phys. Rev. Lett.* **124**, 210603 (2020).
- [41] M. P. Müller, *Phys. Rev. X* **8**, 041051 (2018).
- [42] K. V. Hovhannisyan, M. Perarnau-Llobet, M. Huber, and A. Acín, *Phys. Rev. Lett.* **111**, 240401 (2013).
- [43] N. Friis and M. Huber, *Quantum* **2**, 61 (2018).
- [44] L. Mandelstam and I. Tamm, *J. Phys. (USSR)* **9**, 1 (1945).
- [45] J. Anandan and Y. Aharonov, *Phys. Rev. Lett.* **65**, 1697 (1990).
- [46] N. Margolus and L. B. Levitin, *Physica (Amsterdam)* **120D**, 188 (1998).
- [47] S. Lloyd, *Nature (London)* **406**, 1047 (2000).
- [48] S. Lloyd, *Phys. Rev. Lett.* **88**, 237901 (2002).
- [49] V. Giovannetti, S. Lloyd, and L. Maccone, *Phys. Rev. A* **67**, 052109 (2003).
- [50] M. M. Taddei, B. M. Escher, L. Davidovich, and R. L. de Matos Filho, *Phys. Rev. Lett.* **110**, 050402 (2013).
- [51] A. del Campo, I. L. Egusquiza, M. B. Plenio, and S. F. Huelga, *Phys. Rev. Lett.* **110**, 050403 (2013).
- [52] S. Deffner and E. Lutz, *Phys. Rev. Lett.* **111**, 010402 (2013).
- [53] I. Marvian, R. W. Spekkens, and P. Zanardi, *Phys. Rev. A* **93**, 052331 (2016).
- [54] B. Shanahan, A. Chenu, N. Margolus, and A. del Campo, *Phys. Rev. Lett.* **120**, 070401 (2018).
- [55] S. Deffner and S. Campbell, *J. Phys. A* **50**, 453001 (2017).
- [56] S. An, D. Lv, A. del Campo, and K. Kim, *Nat. Commun.* **7**, 12999 (2016).
- [57] S. Campbell and S. Deffner, *Phys. Rev. Lett.* **118**, 100601 (2017).
- [58] K. Ito and T. Miyadera, *arXiv:1711.02322*.
- [59] G. M. Andolina, D. Farina, A. Mari, V. Pellegrini, V. Giovannetti, and M. Polini, *Phys. Rev. B* **98**, 205423 (2018).
- [60] S. Julia-Farre, T. Salamon, A. Riera, M. N. Bera, and M. Lewenstein, *Phys. Rev. Research* **2**, 023113 (2020).
- [61] G. M. Andolina, M. Keck, A. Mari, V. Giovannetti, and M. Polini, *Phys. Rev. B* **99**, 205437 (2019).
- [62] P. Skrzypczyk, A. J. Short, and S. Popescu, *Nat. Commun.* **5**, 4185 (2014).
- [63] M. P. Müller, *Phys. Rev. X* **8**, 041051 (2018).
- [64] A. E. Allahverdyan and T. M. Nieuwenhuizen, *Phys. Rev. E* **71**, 066102 (2005).
- [65] P. Talkner, E. Lutz, and P. Hänggi, *Phys. Rev. E* **75**, 050102 (2007).
- [66] M. Perarnau-Llobet, E. Bäumer, K. V. Hovhannisyan, M. Huber, and A. Acín, *Phys. Rev. Lett.* **118**, 070601 (2017).
- [67] C. A. Rodríguez-Rosario, G. Kimura, H. Imai, and A. Aspuru-Guzik, *Phys. Rev. Lett.* **106**, 050403 (2011).
- [68] S. Das, S. Khatri, G. Siopsis, and M. M. Wilde, *J. Math. Phys. (N.Y.)* **59**, 012205 (2018).
- [69] H. P. Breuer and F. Petruccione, *The Theory of Open Quantum Systems* (Oxford University Press, New York, 2007).
- [70] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.125.040601> for proof of bounds for open systems and details of the illustration on a model for a heat engine.
- [71] N. Linden, S. Popescu, and P. Skrzypczyk, *arXiv:1010.6029*.
- [72] N. Brunner, N. Linden, S. Popescu, and P. Skrzypczyk, *Phys. Rev. E* **85**, 051117 (2012).