

A low-control and robust quantum refrigerator, and applications with electronic spins in diamond

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We propose a general protocol for low-control refrigeration and thermometry of thermal qubits, which can be implemented using electronic spins in diamond. The refrigeration is implemented by a probe, consisting of a network of spins with two-body XXZ interactions. The protocol involves two operations: (i) free evolution of the probe; and (ii) a swap gate between one spin in the probe and the thermal qubit we wish to cool. We show that if the initial state of the probe falls within a suitable range, then the cooling protocol will always succeed, with an efficiency that depends on the rate of spin dephasing and the swap gate fidelity. Furthermore, measuring the probe after it has cooled many qubits provides an estimate of their temperature. We provide a specific example where the probe is a Heisenberg spin chain, and suggest a physical implementation using electronic spins in diamond. Here the probe is constituted of nitrogen vacancy (NV) centers, while the thermal qubits are dark spins. By using a novel pulse sequence, a chain of NV centers can be made to evolve according to a Heisenberg Hamiltonian. This proposal allows for a range of applications, such as NV-based nuclear magnetic resonance of photosensitive molecules kept in a dark spot on a sample, and it opens up new possibilities for the study of quantum thermodynamics, environment-assisted sensing, and many-body physics.

I. INTRODUCTION

Quantum mechanics and thermodynamics are arguably two of the most successful physical theories to date. Quantum thermodynamics [1–3] is the interdisciplinary field that studies how the two theories influence one-another. For example, the thermodynamic laws of macroscopic physics are thought to *emerge* from the laws of quantum mechanics, when the number of quantum particles in a system grows to be infinitely large [4]. On the other hand, thermodynamic protocols have been shown to operate differently at the scale of few-particle quantum systems [5–8]. A central goal of quantum thermodynamics concerns the design of efficient and robust quantum mechanisms to cool such quantum systems: i.e., the development of “quantum refrigerators.” Cooling is an essential component for many emerging quantum technologies, including fault tolerant quantum computation [9] and quantum metrology at the Heisenberg limit of sensitivity [10]. This is because many of the salient features

of quantum mechanics only emerge when the system is in a low-entropy state, and cooling is the most natural method of entropy reduction. The cooling mechanisms that have been developed so far can be classified into three major groups: (i) dissipative cooling, where the system is cooled by bringing it into thermal equilibrium with a reservoir of lower temperature, which can be prepared with an absorption refrigerator [11–13]; (ii) dynamical cooling, where the dynamics of the system-plus-reservoir composite is controlled [14–16]; and (iii) measurement-assisted cooling, where entropy is reduced through projective measurements, followed by conditional unitary gates that transform the post-measurement state of the system to, say, the ground state of its Hamiltonian [17, 18]. All of these strategies suffer from different drawbacks. For example, while dissipative dynamics with a reservoir requires the least degree of control, it is normally slow. Moreover, the colder the initial temperature of a reservoir is, the more time and power is required to cool it further. This is due to the third law of thermodynamics [19]. Dynamical cooling, on the other hand, can cool at a faster rate, but generally requires a very high degree of control and reservoir-engineering. Finally, although measurement-assisted cooling can be

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fast, measuring one system can disturb others that are nearby. Furthermore, measurement-assisted cooling requires single-shot measurements, but these are often difficult or even impossible to implement experimentally. For example, although single-shot measurement of spins associated with the diamond nitrogen vacancy (NV) center were recently achieved at room [20] and low temperature [21], they are still limited in fidelity. A cooling strategy that combines the benefits of being fast, requiring low control, and acting locally on small systems would therefore be of great use.

Controlling the dynamics of non-equilibrium many-body systems has been proven to be efficient for information transfer [22–24], entanglement generation [25], and quantum gate operations [26, 27]. This relies on the unitary evolution of the system, generated by its Hamiltonian, to perform the desired state transformation. Consequently, the system must be initialized in a non-equilibrium state, such as a superposition of energy eigenstates. The speed of the unitary dynamics is determined by the couplings between the particles and can be engineered to be fast. One may wonder if it is possible to exploit the coherent dynamics of a non-equilibrium quantum system, which we call a refrigeration probe, to cool another system that is in thermal equilibrium. There are three major questions that need to be addressed: (i) will the refrigeration protocol be robust, i.e., will it always cool the thermal system, or could it possibly heat the system instead?; (ii) how much control is required for the probe to function as a refrigerator?; (iii) what is the maximum amount of entropy that the probe can extract from the thermal systems – if the initialization time of the probe is long, and we may only extract a small quantity of entropy with it, then this will limit any potential benefits that fast, coherent dynamics may offer.

This paper addresses these questions. We consider how to use the coherent dynamics of a probe to cool quantum bits (qubits) with temperature T . The setup is shown in Fig. 1, where the probe is a system of interacting spin-half systems (red spheres), and the black spheres are a system of thermal qubits. We prove that if any given pair of spins in the probe have an XXZ interaction (a generalization of a Heisenberg interaction, but with an anisotropy in the z -direction), and if the probe is initialized in an appropriate “cold” state, then it will always cool the thermal qubits it interacts with. We show that minimal control is required – one only needs to engineer a time-controlled interaction Hamiltonian between one spin in the probe and the qubit to be cooled, which will generate a swap operation between them. Finally, a probe with multiple “cold” spins allows more entropy to be extracted from the thermal qubits. This will reduce the need for constant re-initialization of the probe. As an ad-

ditional benefit, we show that the probe can also act as a thermometer to estimate the temperature, T . While this mechanism is very general, we propose a specific model where the probe is a one-dimensional Heisenberg spin chain, and investigate the performance of this probe numerically. Furthermore, we offer an implementation of this model with electronic spins in diamond, where the probe is composed of nitrogen vacancy centers (NVs) and the thermal qubits are dark spins. The probe could allow cooling and sensing of a photosensitive target molecule if one end of the spin chain is in proximity to the target molecule, in the dark, and the other end is cooled by optical pumping. A novel pulse sequence, consisting of a modified version of the WAHUA [28], is proposed to achieve a Heisenberg spin chain with NVs.

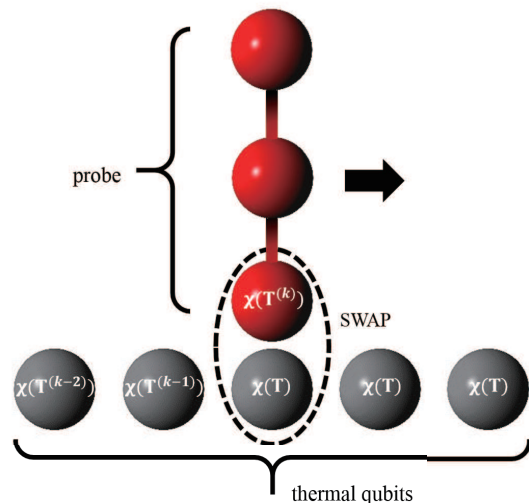


FIG. 1: Schematic of the cooling process. The probe (red spheres), is a network of spin-half systems, coupled through an XXZ interaction. The black spheres are a collection of thermal qubits that are initially in the state $\chi(T)$, with temperature $T > 0$. The protocol cools the k^{th} thermal qubit by: (i) first allowing the probe to evolve freely, for a time τ_k , so that the target spin is prepared in the state $\chi(T^{(k)})$, where $T^{(k)} \leq T$; and (ii) subsequently, swapping the target spin of the probe with the k^{th} thermal qubit, thus cooling it.

II. THEORETICAL MODEL

A. The set-up

We consider a probe \mathcal{P} , consisting of a network of N spin-half systems. The free evolution of the probe is generated by the Liouville super-operator

$$\mathcal{L} : \rho_{\mathcal{P}} \mapsto i[\rho_{\mathcal{P}}, H_{\mathcal{P}}]_{-} + \Gamma \sum_{n=1}^N (\sigma_n^z \rho_{\mathcal{P}} \sigma_n^z - \rho_{\mathcal{P}}). \quad (\text{II.1})$$

Here $[\cdot, \cdot]_-$ is the commutator and the first term on the right hand side accounts for the unitary evolution of the probe due to its XXZ Hamiltonian, given by

$$H_{\mathcal{P}} := \sum_{n=1, m>n}^N J_{n,m} (\Delta_{n,m} \sigma_n^z \otimes \sigma_m^z + \sigma_n^x \otimes \sigma_m^x + \sigma_n^y \otimes \sigma_m^y). \quad (\text{II.2})$$

Here $\{\sigma_n^i | i \in \{x, y, z\}\}$ are the Pauli operators on the n^{th} spin, $J_{n,m}$ is the interaction strength between the n^{th} and m^{th} spins, and $\Delta_{n,m}$ is the anisotropy parameter in the z direction. Throughout, we work in units of $\hbar = 1$. The second term of Eq. (II.1) accounts for local dephasing of the probe, with strength $\Gamma \geq 0$.

We use this probe to cool a collection of thermal qubits. Each thermal qubit \mathcal{Q} has the Hamiltonian $H_{\mathcal{Q}} := \sigma^z$, and is initially in the state

$$\chi(T) := \frac{e^{-H_{\mathcal{Q}}/K_B T}}{Z}. \quad (\text{II.3})$$

Here, K_B is Boltzmann's constant, T is the temperature, and $Z := \text{tr}[e^{-H_{\mathcal{Q}}/K_B T}]$ is the partition function.

The composite system of the probe and the k^{th} thermal qubit is initially in the product state

$$\rho_{\mathcal{P}+\mathcal{Q}}^{(k)} := \rho_{\mathcal{P}}^{(k-1)} \otimes \chi(T). \quad (\text{II.4})$$

Allowing the probe to evolve freely for a duration of τ_k , and then swapping the target spin of the probe with the thermal qubit, produces the state

$$\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k) := (\text{SWAP} \circ e^{\tau_k \mathcal{L}}) [\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}]. \quad (\text{II.5})$$

Here, $e^{\tau_k \mathcal{L}}$ is the free evolution quantum channel (completely positive, trace preserving map) generated by the Liouville super-operator given in Eq. (II.1), and SWAP is a (possibly imperfect) swap operation between the k^{th} thermal qubit and the target spin of the probe. This can be generated by a time controlled Heisenberg interaction, making the swap operation perfect in the limit that this interaction is much stronger than the intra-probe couplings. After the joint evolution, the probe and thermal qubit have the new states

$$\begin{aligned} \rho_{\mathcal{P}}^{(k)} &:= \text{tr}_{\mathcal{Q}} [\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k)], \\ \rho_{\mathcal{Q}}^{(k)} &:= \text{tr}_{\mathcal{P}} [\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k)]. \end{aligned} \quad (\text{II.6})$$

We omit the τ_k dependence for simplicity. The probe will then be moved to the next thermal qubit and the process can begin anew.

Throughout Sec. II, unless stated otherwise, we assume \mathcal{P} is a network of spins coupled through the XXZ interaction. On the other hand, when we wish to analyze the system numerically, we restrict ourselves to the simplest class of an XXZ spin network, i.e., an isotropic Heisenberg spin chain. This has the simplified Hamiltonian,

$$H_{\mathcal{P}} := J \sum_{n=1}^{N-1} \boldsymbol{\sigma}_n \cdot \boldsymbol{\sigma}_{n+1}, \quad (\text{II.7})$$

where $\boldsymbol{\sigma}_n := (\sigma_n^x, \sigma_n^y, \sigma_n^z)$ is a vector of Pauli operators on the n^{th} spin, and J is the interaction strength between each nearest-neighbour spins.

B. Application 1: cooling

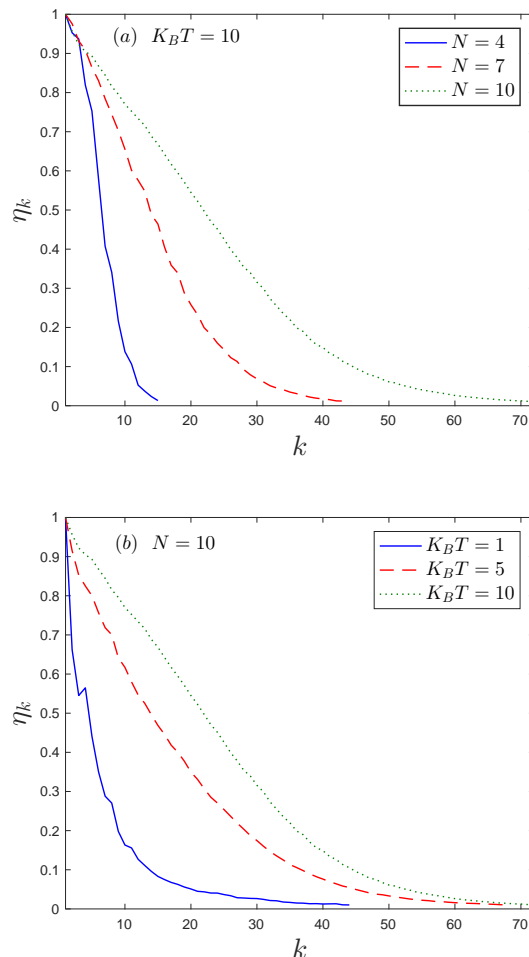


FIG. 2: The ideal cooling protocol, using a Heisenberg spin chain with $J = 1$ and $\epsilon = 0.01$. (a) and (b) show, respectively, the dependence of the cooling efficiency of the k^{th} thermal qubit, η_k , on the length of the chain, N , and temperature of the thermal qubits, T .

As shown in Appendix (A), if the initial state of the probe can be written as

$$\rho_{\mathcal{P}}^{(0)} = \bigotimes_{n=1}^N \chi(T_n), \quad (\text{II.8})$$

such that for all n , $T_n \leq T$, then irrespective of the thermal qubit number k , and the waiting times $\{\tau_k\}_k$, we have

$$\rho_{\mathcal{Q}}^{(k)} = \chi(T^{(k)}), \quad (\text{II.9})$$

with $T^{(k)} \leq T$. That is to say, so long as we can initialize the probe in an appropriate state, it will always cool the thermal qubits, or leave them at the same temperature. This will be true even in the presence of dephasing, or with imperfect swap operations. Note that Eq. (II.8) is not a thermal state of the probe. Each spin in the probe, however, can be thought of as being ‘‘colder’’ than the thermal qubits in a counterfactual sense – if the probe was also a system of non-interacting spins, each with Hamiltonian σ^z . To quantify the performance of each cooling process, we introduce the cooling efficiency, defined as

$$\eta_k := \frac{T - T^{(k)}}{T}. \quad (\text{II.10})$$

We wish to maximise the cooling efficiency at each stage by optimising the waiting times $\{\tau_k\}_k$. This can be done if we have prior knowledge of: the temperature, T ; the probe and qubit Hamiltonians, $H_{\mathcal{P}}$ and $H_{\mathcal{Q}}$, respectively; and the initial state of the probe, $\rho_{\mathcal{P}}^{(0)}$. By simulating the dynamics of the probe, we may find the shortest time τ_k that maximizes η_k . Even when the probe undergoes dephasing, we evaluate the optimal sequence $\{\tau_k\}_k$ for the case of zero dephasing only. This is because the dephasing strength cannot in general be known and it may change from shot-to-shot.

We stop the protocol when the efficiency drops below a threshold $\epsilon \in [0, 1]$. Let k_ϵ be the smallest k such that η_k is larger than ϵ , and where η_{k+1} is smaller than ϵ :

$$k_\epsilon := \min\{k | \eta_k \geq \epsilon \text{ and } \eta_{k+1} < \epsilon\}. \quad (\text{II.11})$$

At the k^{th} stage, the total entropy is reduced by

$$\Delta S_{\mathcal{Q}}^{\text{total}}(k) := \sum_{i=1}^k \Delta S_{\mathcal{Q}}^{(i)}, \quad (\text{II.12})$$

where

$$\Delta S_{\mathcal{Q}}^{(k)} := S_T - S_{T^{(k)}} \quad (\text{II.13})$$

is the entropy reduction of the k^{th} thermal qubit, and

$$S_T \equiv S(\chi(T)) := -\text{tr}[\chi(T) \ln(\chi(T))] \quad (\text{II.14})$$

is the von Neumann entropy of the thermal qubit at temperature T . There is a one-to-one correspondence between the efficiency η_k and the entropy reduction $\Delta S_{\mathcal{Q}}^{(k)}$, where a higher efficiency translates to a larger entropy reduction and vice versa. However, these quantities scale differently, as will become apparent when we discuss the effect of imperfections on the cooling protocol in Sec. IID. As shown in Appendix (C), the total entropy reduction of the thermal qubits is bounded by the entropy increase of the probe:

$$\Delta S_{\mathcal{Q}}^{\text{total}}(k) \leq S(\rho_{\mathcal{P}}^{(k)}) - S(\rho_{\mathcal{P}}^{(0)}) \leq N S_T, \quad (\text{II.15})$$

where a necessary condition for achieving the upper bound is for the probe to be initially prepared in the state

$$\rho_{\mathcal{P}}^{(0)} = |1\rangle\langle 1|^{\otimes N}. \quad (\text{II.16})$$

Here, we are using the notation $\sigma^z := |0\rangle\langle 0| - |1\rangle\langle 1|$, so comparing this with Eq. (II.8) shows that for all n , we have $T_n = 0$. Eq. (II.15) shows that the more spins are present in the probe, the more entropy one can extract from the collection of thermal qubits.

In Fig. 2 we numerically investigate the cooling efficiency of a Heisenberg spin chain of length N . For the moment, we will consider the optimal scenario where: the initial state of the probe is given by Eq. (II.16); the probe evolves in the absence of dephasing; and the swap operation is perfect and instantaneous. Furthermore, we limit the range of evolution times to $\tau_k \in [0, N/J]$. The upper limit N/J is chosen so that the excitations of the probe have enough time to travel from one end of the chain to the other. In Fig. 2(a) we plot η_k as a function of k for various N when the original temperature is fixed to $K_B T = 10$. As can be seen, the efficiencies decrease as the protocol progresses. However, larger chains will provide higher efficiencies over more iterations. Similarly, in Fig. 2(b) we plot η_k as a function of k for various temperatures when the probe length is fixed to $N = 10$. As before, the efficiencies decrease as the protocol progresses. The performance of the protocol improves for hotter qubits. Because of the one-to-one correspondence between entropy reduction and efficiency, the behavior of $\Delta S_{\mathcal{Q}}^{\text{total}}(k)$ will be qualitatively identical in this case.

C. Application 2: thermometry

In Appendix (B) we prove that, given the dynamics given in Eq. (II.5), the probe has a unique stationary state

$$\rho_{\mathcal{P}}^{(\infty)} = \chi(T)^{\otimes N}. \quad (\text{II.17})$$

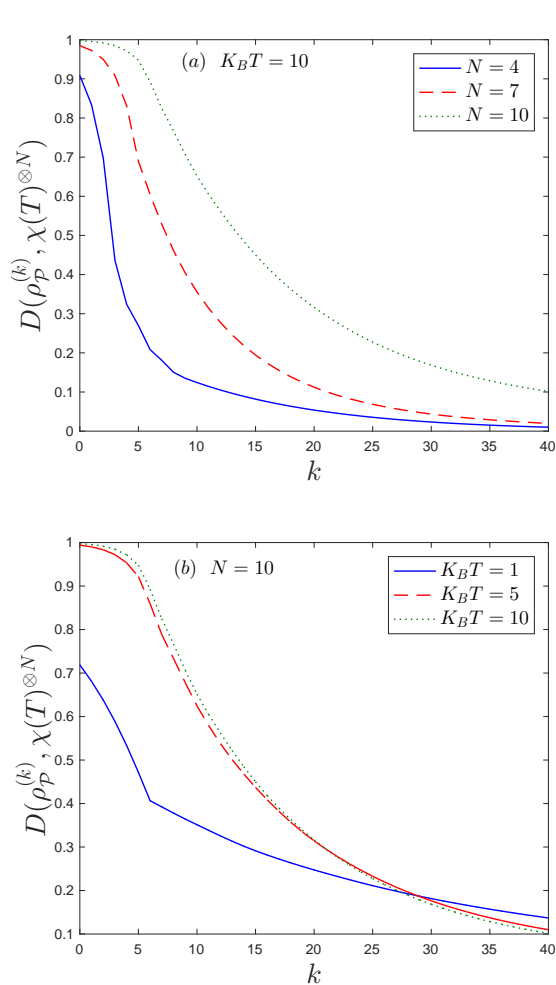


FIG. 3: Pseudo-thermalization of a Heisenberg spin chain as a result of the time dynamics, where $D(\rho_{\mathcal{P}}^{(k)}, \chi(T)^{\otimes N})$ is the trace distance between the state of the probe and the pseudo-thermal state, after the k^{th} thermal qubit has been cooled. In all cases, we set $J = \tau_k = 1$. (a) and (b) show, respectively, the dependence of pseudo-thermalization on the chain length N and temperature T , for the ideal case.

This follows from the fact that the above state is the only one that commutes with the generating Hamiltonian of the evolution. We say this is a pseudo-thermal state because it is not given as the Gibbs state of the probe Hamiltonian $H_{\mathcal{P}}$, but rather as N copies of the thermal qubits $\chi(T)$. This feature of the probe allows it to be used for thermometry; we may obtain an estimate for the temperature of the thermal qubits, T , from the measurement statistics of the σ^z observable on every spin of the probe. If the probe is prepared in the steady state $\rho_{\mathcal{P}}^{(\infty)}$, we will have N identical copies of $\chi(T)$ for our measurement statistics. In practical situations, however, we take the probe for thermometry just after a finite number of iterations, when the steady state has not yet been fully achieved.

The trace distance between the state of the probe and

the pseudo-thermal state, $D(\rho_{\mathcal{P}}^{(k)}, \chi(T)^{\otimes N})$, bounds the accuracy of our estimation of T . Due to the contractivity of the trace distance under quantum channels, this will never increase as we continue to interact with the thermal qubits [29]. How fast this quantity vanishes determines the performance of the probe for thermometry. Furthermore, unlike the case of cooling, we are by definition ignorant of the temperature T . Therefore, we cannot simulate the dynamics of the probe, and have no means of optimizing the waiting times $\{\tau_k\}_k$ between consecutive swaps. Accordingly, we must make an arbitrary choice.

In Fig. 3 we simulate the pseudo-thermalisation rate, given a Heisenberg spin chain of length N , and with $\tau_k = J = 1$. For the moment, we will consider the optimal scenario where: the initial state of the probe is given by Eq. (II.16); the probe evolves in the absence of dephasing; and the swap operation is perfect and instantaneous. In Fig. 3(a) we show the dependence of pseudo-thermalization on probe length N for a fixed temperature of $K_B T = 10$. It is evident that increasing N slows the rate of pseudo-thermalization. This implies a trade-off between the time required for thermometry, and the accuracy of thermometry; the more spins we have in the probe, the better our measurement statistics will be, but the longer we need to wait before making these measurements. In Fig. 3(b) we show the dependence of pseudo-thermalization on the temperature, for a fixed probe length of $N = 10$. For the first few iterations, the probe's distance to the pseudo-thermal state is greater for warmer temperatures T . However, this reverses after (approximately) the 30th qubit has been cooled, where increasing temperature decreases the trace-distance. Similarly with the probe length, this suggests a trade-off relation between time required for thermometry, and the temperature of the thermal qubits; colder temperatures require more time to be accurately determined.

D. Imperfections

There are two imperfections in the system that we study here: (i) presence of dephasing on the probe; and (ii) an imperfect swap operation implemented by a time-controlled Heisenberg interaction of finite strength. As Theorem A.1 and Theorem B.1 still apply in the case of imperfections, the cooling protocol will still function robustly, and the probe can still act as a thermometer. Moreover, because Theorem C.1 also applies, we know that the total entropy reduction will be bounded by the probe size. However, the efficiency of the protocols may change. To analyze the effect of such imperfections quantitatively, we will numerically investigate a probe consisting of a Heisenberg spin chain of $N = 10$ spins, initialized to the state

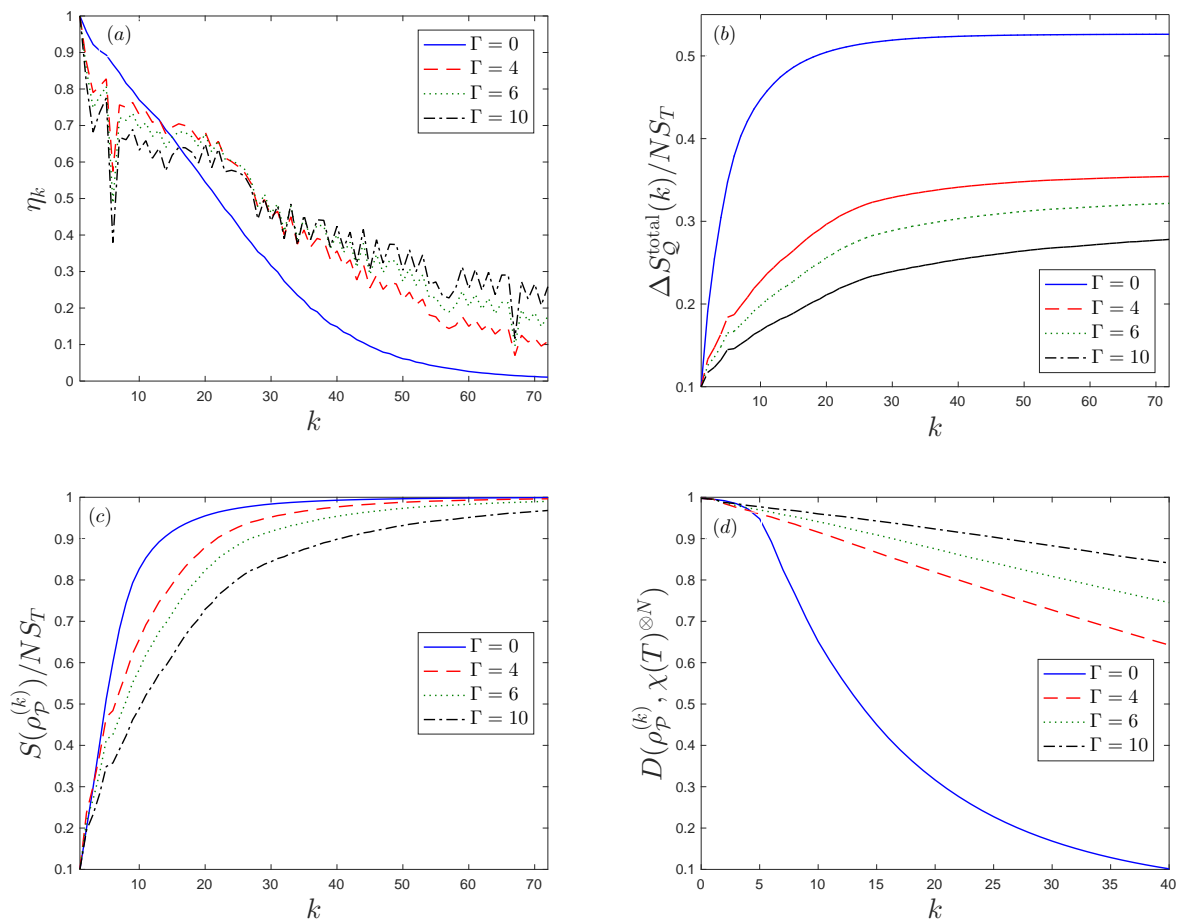


FIG. 4: (a) - (c) show the performance of the cooling protocol with a Heisenberg spin chain probe in the presence of local dephasing of strength Γ . All waiting times τ_k are calculated for the ideal case with $J = 1$, $N = 10$, $K_B T = 10$, and $\epsilon = 0.01$. (a) shows the dependence of the cooling efficiency of the k^{th} thermal qubit, η_k , on the dephasing strength. (b) and (c) show, respectively, how the dephasing strength affects the total entropy reduction of the thermal qubits, $\Delta S_Q^{\text{total}}(k)$, and the entropy of the chain, $S(\rho_P^{(k)})$, after the k^{th} thermal qubit has been cooled. S_T is the entropy of the thermal state $\chi(T)$. (d) shows the effect of dephasing on pseudo-thermalization, where $D(\rho_P^{(k)}, \chi(T)^{\otimes N})$ is the trace distance between the state of the probe and the pseudo-thermal state, after the k^{th} thermal qubit has been cooled. Here, we set $J\tau_k = 1$.

given by Eq. (II.16). The thermal qubits will be fixed to a temperature of $K_B T = 10$.

1. Dephasing

It is in general difficult to keep the probe fully isolated and, thus, the free evolution will not be unitary. To account for the interaction between the probe and its environment, we consider local dephasing with $\Gamma > 0$ in Eq. (II.1). For the cooling protocol, the waiting times $\{\tau_k\}_k$ are calculated for the ideal case, i.e. $\Gamma = 0$. In Fig. 4(a) we plot the efficiency η_k versus the step k for different values of Γ . As the figure shows, while an increase in dephasing strength results in a decrease in cooling efficiency for the first few iterations,

this is reversed at later stages. Fig. 4(b) shows that the total entropy reduction of the qubits, after stage k of the cooling protocol, is reduced by dephasing. In Fig. 4(c) we show how dephasing affects the entropy of the probe during the cooling protocol. The probe's entropy increases monotonically as the protocol progresses, but increasing dephasing strength decreases the probe's entropy at any stage k . Conforming with Eq. (II.15), the probe entropy is always larger than the total entropy reduction obtained on the thermal qubits. In Fig. 4(d) we plot the trace distance between the state of the probe and the pseudo-thermal state $\chi(T)^{\otimes N}$, as a function of k . Here, the time between consecutive swaps is fixed to $J\tau_k = 1$. As this figure shows, increasing dephasing strength slows the rate of pseudo-thermalization.

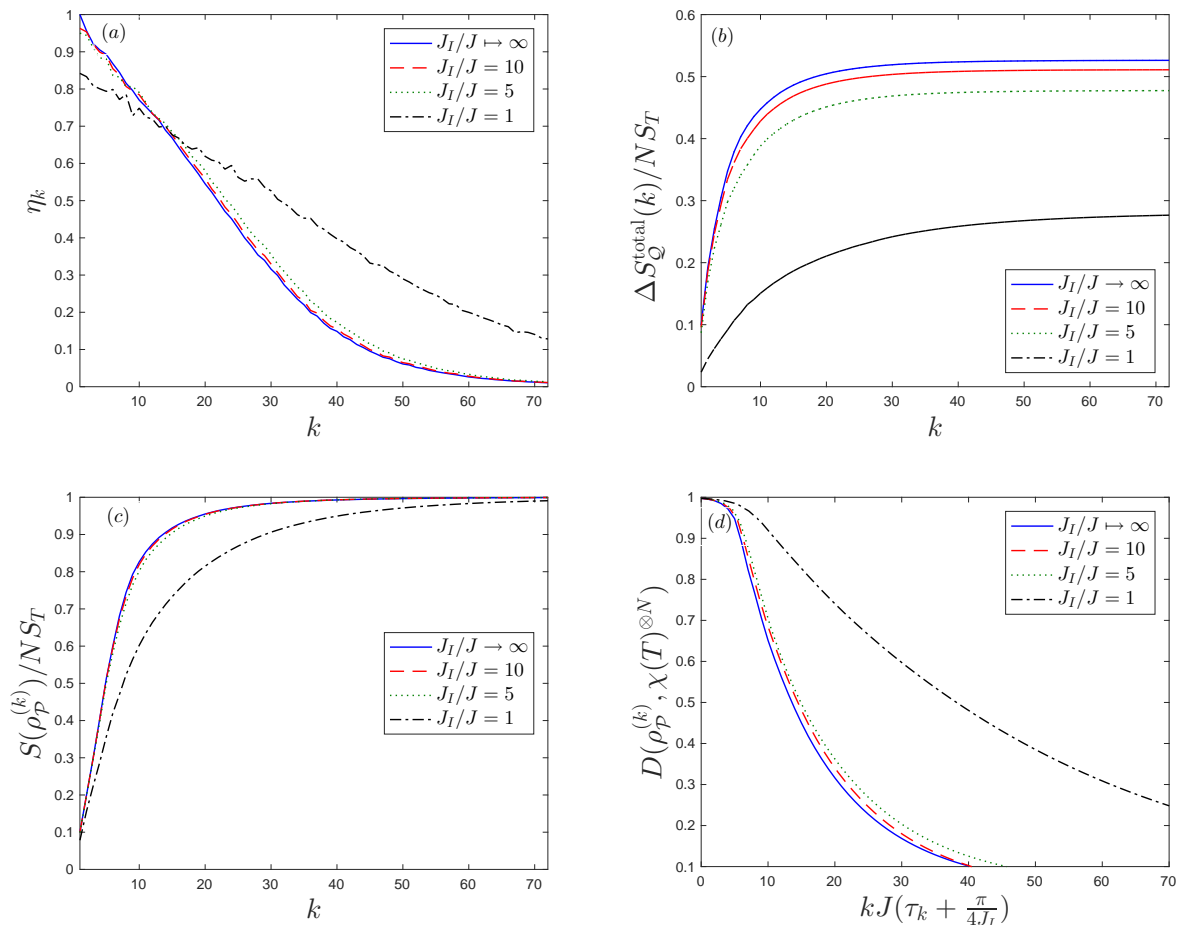


FIG. 5: (a) - (c) show the performance of the cooling protocol with a Heisenberg spin chain probe, with the swaps effected by a time-dependent Heisenberg interaction of strength J_I . All waiting times τ_k are calculated for the ideal case with $J = 1$, $N = 10$, $K_B T = 10$, and $\epsilon = 0.01$. (a) shows the dependence of the cooling efficiency of the k^{th} thermal qubit, η_k , on J_I . (b) and (c) show, respectively, how J_I affects the total entropy reduction of the thermal qubits, $\Delta S_Q^{\text{total}}(k)$, and the entropy of the chain, $S(\rho_P^{(k)})$, after the k^{th} thermal qubit has been cooled. S_T is the entropy of the thermal state $\chi(T)$. (d) shows the effect of J_I on the rate of pseudo-thermalization, where $D(\rho_P^{(k)}, \chi(T)^{\otimes N})$ is the trace distance between the state of the probe and the pseudo-thermal state, after the k^{th} thermal qubit has been cooled. Here, we set $J\tau_k = 1$.

2. Partial swaps

We now allow for the swap operation to be imperfect corresponding to a finite-duration interaction between the thermal qubit and probe spin. An imperfect swap may be realized by the time-dependent Hamiltonian of the compound system of probe and thermal qubit,

$$H_{\mathcal{P}+\mathcal{Q}}(t) := H_{\mathcal{P}} + H_{\mathcal{Q}} + H_I(t), \quad (\text{II.18})$$

with the interaction Hamiltonian

$$H_I(t) := f(t)J_I\sigma_{\mathcal{Q}} \cdot \sigma_{\mathcal{P}}. \quad (\text{II.19})$$

Here, $\sigma_{\mathcal{Q}}$ and $\sigma_{\mathcal{P}}$ are vectors of Pauli operators acting on the thermal qubit and the first spin of the probe, respectively, and J_I is the interaction strength

between these systems. In the absence of $H_{\mathcal{P}}$ and $H_{\mathcal{Q}}$, this Hamiltonian would induce a swap operation (with irrelevant phase factors) if $f(t) = 1$ for a period of $\pi/(4J_I)$, and zero otherwise.

To understand the effect of finite time-duration swap gates, we plot in Fig. 5(a) the cooling efficiency as a function of the normalized interaction strength J_I/J . We set dephasing to zero and, as before, the waiting times $\{\tau_k\}_k$ are calculated assuming for the ideal case, i.e., instantaneous and perfect swaps. Similarly to the case of dephasing, while a decrease in J_I/J results in a decrease in cooling efficiency for the first few iterations, this is reversed at later stages. However, as shown in Fig. 5(b), the total entropy reduction of the qubits, after stage k of the cooling protocol, is always less when J_I/J decreases. This means that decreasing J_I/J always decreases the overall perfor-

mance of the protocol. To see how the probe is affected by the strength of J_I/J , in Fig. 5(c) we depict the entropy of the probe after the k^{th} qubit has been cooled. The probe's entropy increases monotonically as the protocol progresses, but decreasing J_I/J lowers the probe's entropy at any stage k . Again, conforming with Eq. (II.15), the entropy of the probe always exceeds the total entropy reduction obtained on the thermal qubits. Finally, in Fig. 5(d) we plot the distance between the state of the probe and the pseudo-thermal state $\chi(T)^{\otimes N}$, as a function of k . The time between consecutive swaps is set to $J\tau_k = 1$. As the figure shows, while an increase in J_I/J from unity to five significantly improves the rate of pseudo-thermalization for the present case of evolution times, further increases in J_I/J have a much less noticeable effect.

III. DARK SPIN COOLING WITH A NITROGEN VACANCY SPIN CHAIN

Electronic spins in diamond are promising to realize our proposal discussed in Sec. II. Here the thermal qubits we wish to cool are environmental dark spins [30], and the probe is a Heisenberg spin chain composed of nitrogen vacancy (NV) color centers. In the negative charge state, the NV^- ground state constitutes a localized, spin-1 system with coherence times exceeding milliseconds even at room temperature [31, 32]. Its spin states can be initialized, manipulated, and measured with optical and microwave fields. The combined advantages of NVs — long coherence time, easy manipulation of spin states, and large gyromagnetic ratio (compared with nuclear spins) — make it a good candidate for quantum sensing [33–35]. Recent demonstrations on quantum sensing, such as paramagnetic centers in solids [36], single protein molecules [37], and a few nuclear spins [35], have shown the potential. The location of NVs within the diamond can be controlled in a variety of ways, including localized delta-doped growth [38], targeted implantation through a focused ion beam [39], and nano-masked implantation [40, 41]. These fabrication techniques have demonstrated the possibility of constructing NV spin chains with spatial precision on the 10 nm scale, as required for the realization of our method.

Besides NV centers, diamond is host to many different dark spins [30] — dark in the sense that they are not fluorescent. In particular, the low conversion efficiency from implanted (or native) nitrogen atoms in the diamond lattice to NV centers (~5% [42]) results in a large number of single-substitutional nitrogen defect centers (P1 centers) in the vicinity of NVs. These dark spins generally act as a spin bath, decohering the NV

centers [43]. However, some proximal spins can coherently interact with the NV centers [42, 44, 45]. If these proximal dark spins can be cooled down (initialized) efficiently, they can serve as a quantum resource in environment-assisted sensing [46]. There have been many attempts for dark spin cooling, but the polarization has been much lower than that of the NV so far. [47, 48].

Here, we propose an efficient method for dark spin cooling, which uses an NV center spin chain as a probe. The NV center that is closest to the dark spins takes the role of the first spin in the probe in Sec. II. The benefit of the spin chain is that it provides a cold reservoir and cooling conduit that can be cooled in one region. One exemplary application is magnetic resonance detection of photosensitive molecules, as illustrated in Fig. 6(a). In such circumstances, while nearby NV centers are able to interact with the dark spins and target molecules, they cannot be initialized constantly with a strong optical field. A possible solution is to use a chain of NVs to initialize the NVs far from the molecule, and let the chain transfer polarization to the dark spins. Through this method, one can gain more sensitivity from both coherence time ($\sim 1/\sqrt{T_2}$) and the number of spins ($\sim 1/\sqrt{N}$ for standard quantum limit or $1/N$ for Heisenberg limit depending on the sensing scheme [46]).

This spin chain cooling must meet several requirements: (i) initially, the NV spin chain should be cooled down, i.e., each NV must be cooled down with respect to its bare Hamiltonian; (ii) the dark spins should be decoupled from each other; (iii) a SWAP gate between the first NV and each individual dark spin should be applied when needed; and (iv) the spins in the chain should have a nearest-neighbor Heisenberg interaction. NVs can be optically initialized (polarized) with high fidelity [49]. A doughnut beam initializes NVs far from the photo-sensitive molecules, and the Heisenberg interaction distributes the polarization to the whole chain before transferring it to the dark spins. This automatically satisfies condition (i). In subsequent subsections, we will investigate a way to implement (ii) ~ (iv).

A. Probe NV - dark spin interaction

Several types of dark spins in diamond have been extensively studied [30]. Here, we focus on the P1 centers closely related with the NV center implantation process [45], but the argument is the same for other species.

The magnetic dipolar interaction between NV and P1 centers is captured by the interaction Hamiltonian

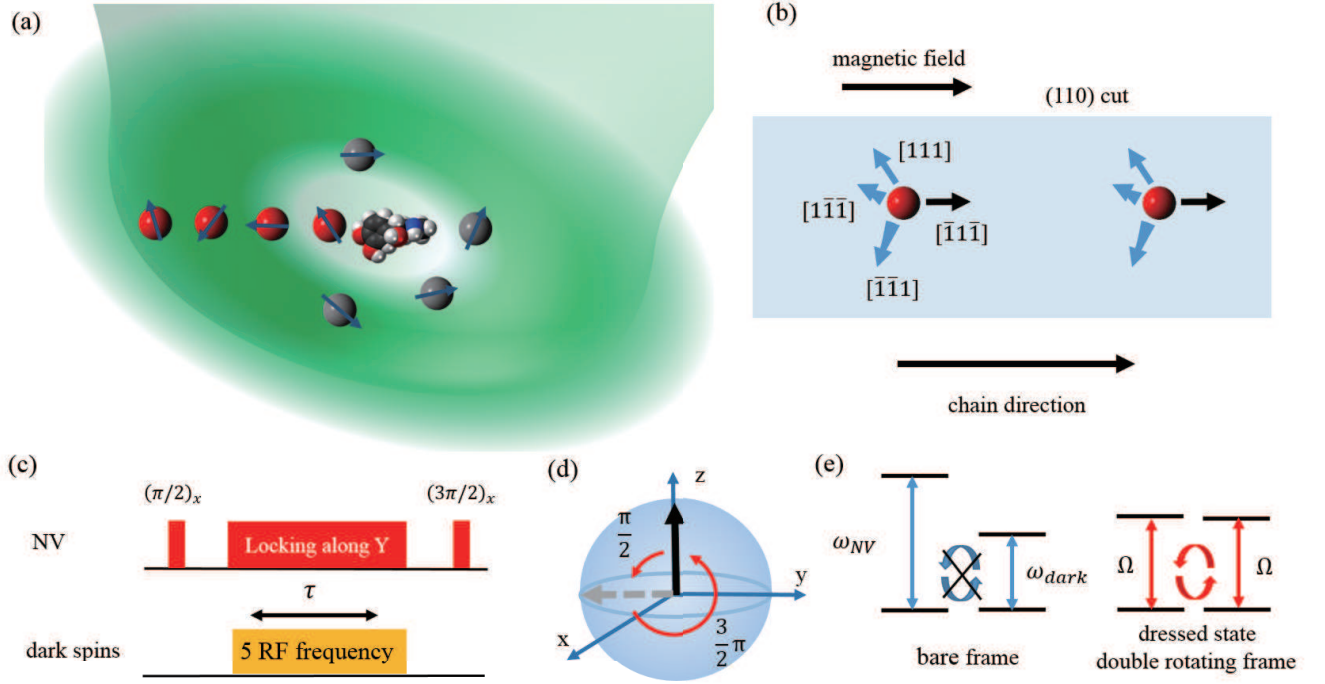


FIG. 6: (a) Dark spin cooling with an NV spin chain for single molecule NMR. Black spheres represent dark spins while red ones do NVs. (b) Orientation of NVs in the spin chain. To obtain a uniform interaction strength with optimal yield, (110)-cut diamond is presented. The magnetic field is aligned into the $[\bar{1}\bar{1}\bar{1}]$ direction (marked as black arrows), and the NV should be oriented into one of the other three directions – $[11\bar{1}]$, $[\bar{1}\bar{1}\bar{1}]$, or $[\bar{1}\bar{1}1]$ (marked as blue arrows). Nearest neighbor spins should have different directions. (c) Pulse sequences for the NV-dark spin interaction. For the dark spin, all 5 RF transitions are driven. For the spin-1 ^{14}N hyperfine axis parallel with magnetic field, the hyperfine splitting is $A_{\parallel} = 114$ MHz, while for the other three axes, $A_{\parallel} = 90$ MHz. (d) Bloch sphere representation of NV spin during pulse sequence. After $(\pi/2)_x$ pulse, the spin is locked into the y-direction (marked as a gray dotted arrow). (e) Dressed state resonant coupling. In the lab frame, energy difference between two spins prohibits energy exchange (spin flip-flop). In the double rotating frame with dressed states, energy can be exchanged between the two.

[50],

$$\begin{aligned}
 H_{dip} &= D_{NV,P1} S_{NV}^z \otimes S_{P1}^z \\
 &\xrightarrow{m_s=1,0} D_{NV,P1} \sigma_{NV}^z \otimes \sigma_{P1}^z \\
 &\xrightarrow{H.H.} \frac{D_{NV,P1}}{4} (\sigma_{NV}^+ \otimes \sigma_{P1}^- + \sigma_{NV}^- \otimes \sigma_{P1}^+)
 \end{aligned}
 \tag{III.1}$$

where: S_{NV}^z and S_{P1}^z are the electronic spin operators of NV centers and P1 centers respectively; σ^α with $\alpha \in \{x, y, z\}$ are Pauli operators on a pseudo spin- $\frac{1}{2}$ subspace spanned by $|m_s = 1\rangle$ and $|m_s = 0\rangle$, for the case of NVs; $\sigma^{+,-}$ are spin ladder operators; H.H. is the Hartmann–Hahn condition [51]; and $D_{NV,P1} = q \frac{\mu_0 \gamma_1 \gamma_2 \hbar^2}{4\pi}$ (Appendix D). Because of the energetic detuning between NV and P1 centers, terms related to spin flip-flops are suppressed in the secular approximation, resulting in an Ising interaction [52]. However, by locking the NV and P1 centers in the transverse direction with the same Rabi frequency (Hartmann-Hahn matching), a flip-flop interaction can be generated [50]. In the σ^z basis, this flip-flop operation is equivalent to the SWAP gate up

to an irrelevant phase factor.

Dressed-state resonant coupling has an advantage in that it is not sensitive to the intrinsic spin level energy. At first, all hyperfine levels of different species of dark spins can be driven, without regard to their associated nuclear spin state. In addition, the interaction can be easily switched on or off by locking or not locking spins. The relevant sequences for probe NV-dark spin coupling are shown in Fig. 6(c). The probe NV is locked while each dark spin (thermal qubit) is driven at the same time with the same Rabi frequency for all hyperfine levels. Demonstrated coupling strength (13MHz) [42] is more than a thousand times larger than the dephasing rate of NVs (~ 1 kHz) implying that the dephasing of the probe qubit is negligible during this SWAP operation.

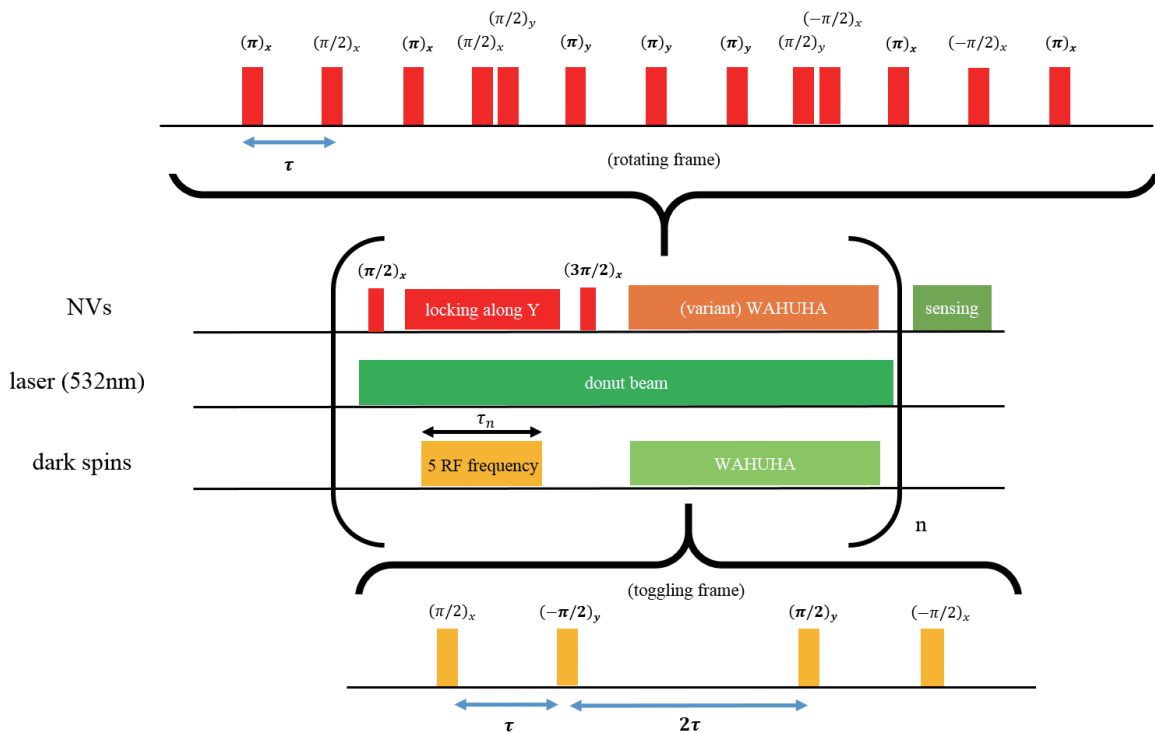


FIG. 7: Variant-WAHUHA sequence for Heisenberg interaction and the whole pulse sequence. Cooling dark spins and thermalizing spin chains are alternated multiple times. Environment-assisted sensing can be applied after this cooling step. Note that variant-WAHUHA is not in a toggling frame while WAHUHA is in a toggling frame.

B. NV Heisenberg spin chain

The interaction between NVs in the spin chain depends on the external magnetic field and the orientation of NVs within the chain, with many possibilities available [53]. In contrast, the required interaction for our method is the spin- $\frac{1}{2}$ Heisenberg interaction. We propose a novel way to form a Heisenberg interaction between NVs in the subspace spanned by $|m_s = 1\rangle$ and $|m_s = 0\rangle$. Here, we consider the situation where NVs are oriented in at most three different crystal orientations of (110) diamond, and that we can make the $|m_s = 1\rangle$ ground state energies of all NVs degenerate while splitting $|m_s = 1\rangle$ from the $|m_s = -1\rangle$ state. This can be achieved by the Zeeman effect by applying a uniform magnetic field in the fourth, $[\bar{1}\bar{1}\bar{1}]$ direction (Fig. 6(b)).

The nearest neighbor NV-NV interaction Hamiltonian after removing the non-energy conserving terms (Appendix E), is

$$H_{int} = -\frac{J_0}{r^3} [2g^+(\sigma_1^x \otimes \sigma_2^x + \sigma_1^y \otimes \sigma_2^y) + 2ih^-(\sigma_1^x \otimes \sigma_2^y - \sigma_1^y \otimes \sigma_2^x) + q\sigma_1^z \otimes \sigma_2^z] \quad (\text{III.2})$$

First, we ignore the $(\sigma_1^x \otimes \sigma_2^y - \sigma_1^y \otimes \sigma_2^x)$ term, and focus on the terms $\sigma_1^\alpha \otimes \sigma_2^\alpha$ for $\alpha = x, y, z$. Except in rare accidental configuration of NVs, which can be avoided in a implantation process, $4g^+ + q \neq 0$. As a result, globally rotating spins will feel an averaged isotropic interaction that is not canceled out. However, since $\sigma_1^x \otimes \sigma_2^x$, $\sigma_1^y \otimes \sigma_2^y$, and $\sigma_1^z \otimes \sigma_2^z$ are mutually non-commuting, simply rotating the spins will not result in the desired Heisenberg interaction. We can use Trotter-Suzuki decomposition [54] to approximate the Heisenberg Hamiltonian, and to minimize errors in the given order. 1st order Trotterization results in the WAHUHA pulse sequence [28] in effective Hamiltonian theory, widely used to nullify homonuclear interactions in solid state NMR. However, in the case of NVs, the application of the WAHUHA sequence results in an effective Heisenberg chain. This discrepancy comes from the difference of interaction Hamiltonians. NVs have a different Hamiltonian with homonuclear dipolar interaction ($\propto \sigma_1^x \otimes \sigma_2^x + \sigma_1^y \otimes \sigma_2^y - 2\sigma_1^z \otimes \sigma_2^z$) because we only use a two-dimensional subspace of the spin-1 Hilbert space (which is 3-dimensional) and express the interaction with pseudo-spin- $\frac{1}{2}$ Pauli operators. The resulting effective Hamiltonian (Appendix E) has the form of

$$H_{int} = -\frac{J_0}{3r^3}(4g^+ + q) \sum_{\alpha \in \{x,y,z\}} \sigma_1^\alpha \otimes \sigma_2^\alpha. \quad (\text{III.3})$$

The previously ignored term, $2ih^-(\sigma_1^x \otimes \sigma_2^y - \sigma_1^y \otimes \sigma_2^x)$ can be cancelled in the context of WAHUHA. Adding a π pulse in any direction does not change the Heisenberg interaction terms because two spins are flipped together. However, when a π pulse is applied in one of the transversal directions to the spin, the $h^-(\sigma_1^x \otimes \sigma_2^y - \sigma_1^y \otimes \sigma_2^x)$ terms change the sign. Therefore, when the evolution time of the two interactions are matched, they cancel each other. This π pulse also serves to dynamically decouple NVs from any slow-moving, detuned nuclear spins, reducing dephasing. The resulting pulse sequence with dark spin cooling is described in Fig. 7. Here, the original version of the WAHUHA is also applied to dark spins to prevent mutual interaction, resulting in a central-spin model.

Although we conjecture that a disorder of interaction strength in the chain does not degrade cooling efficiency dramatically, so that we can use the spin chain having randomly oriented NVs, we can fabricate a uniform interaction strength by post-selecting the chain with properly oriented NVs. Figure 6(b) shows a proper configuration of NVs in the chain. Here, we assume (110)-cut diamond with a static magnetic field in the $[\bar{1}\bar{1}\bar{1}]$ direction. The other three orientations of NVs – $[111]$, $[1\bar{1}\bar{1}]$, $[\bar{1}\bar{1}1]$ – are equivalent to each other in the sense that alternated NV orientations result in equal coupling strengths. The probability of creating a chain of N spins satisfying these properties is $P(N) = \frac{3}{4} \cdot (\frac{1}{2})^{N-1}$ assuming randomly oriented NVs resulting from implantation. Considering a separation of 25 nm between NVs, $N=6$ gives a spin chain of 150 nm length with $\sim 2.3\%$ yield, which could allow enough isolation to the doughnut beam [55, 56]. The T_1 time of NVs can be long, ~ 7.5 ms [57], while the coupling strength of this configuration reaches ~ 12.4 kHz. This allows for ~ 90 repetitive population transfers between nearest-neighbor NVs before relaxation.

IV. CONCLUSIONS

We proposed a method of refrigeration and thermometry of a collection of thermal qubits, each at temperature T , with a quantum many-body-probe composed of a network of spins with two-body XXZ interactions. We showed that minimal control is required; the protocol will succeed with just an imperfect swap gate between the target spin of the probe and the thermal qubit we wish to cool. Moreover, we analytically proved that the probe is a robust refrigerator if it is

initialized in an appropriate state; the thermal qubits will be cooled, or left at the same temperature, even in the presence of dephasing or with imperfect swap gates. Additionally, we showed that this many-body-probe can also be used as a quantum thermometer, providing an estimate of the absolute temperature of the thermal qubits. We numerically investigated a simple example of an XXZ spin network – a Heisenberg spin chain – and quantitatively analyzed how the cooling efficiency is affected by the size of the probe, presence of dephasing, and the fidelity of the swap gate between the probe and the thermal qubit. As the simulation with dephasing demonstrates, coherent dynamics improves the efficiency of the probe, serving a critical role in extracting entropy.

We considered an exemplary implementation using solid-state spin qubits, specifically nitrogen vacancy (NV) centers in diamond. Here, the probe can be constructed as a spin chain of NV centers, which can be used to efficiently cool down dark spins using a quantum refrigeration scheme based on a novel pulse sequence. This system is useful for environment-assisted quantum sensing, especially when the target is a photo-sensitive molecules such as a protein. Overall, our proposal for a low-control and robust quantum refrigerator opens new possibilities for low-entropy quantum-state preparation, useful for quantum metrology, quantum computation and for studying many-body quantum thermodynamics.

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M.H.M. and H.C. have contributed equally to this work.

Appendix A: Sufficient conditions ensuring that the thermal qubits are always cooled

Here we demonstrate the conditions that need to be satisfied by the initial state of the probe, so as to ensure that the cooling protocol will always cool the thermal qubits. To this end, we first give some useful definitions.

Definition A.1. *If a qubit is diagonal with respect to the eigenbasis of σ^z , we refer to it as σ^z -diagonal.*

Definition A.2. *If every eigenstate of a quantum state ρ has a Schmidt decomposition with respect to the eigenbasis of σ^z , i.e. $\{|0\rangle, |1\rangle\}$, we refer to it as σ^z -Schmidt decomposable.*

Definition A.3. *If a quantum channel (completely positive, trace preserving map) describing the time-evolution of a system composed of N spin-half systems for a period of $\tau > 0$, \mathcal{V}_τ , conserves $\sum_{n=1}^N \sigma_n^z$, we refer to it as σ^z -excitation conserving. As such, we can decompose the quantum channel as*

$$\mathcal{V}_\tau = \sum_{l=0}^N \mathcal{V}_\tau^l, \quad (\text{A.1})$$

where \mathcal{V}_τ^l is a quantum channel that acts only on the subspace \mathcal{H}_l , which is spanned by the product vectors containing l spins in the state $|0\rangle$, and $N-l$ spins in the state $|1\rangle$.

Now we prove the conditions under which the reduced state of every spin in an N -partite system will be σ^z -diagonal, which is a necessary condition for them to be thermal with respect to the σ^z Hamiltonian.

Lemma A.1. *Let a quantum system composed of N spin-half systems be prepared in a state*

$$\rho = \bigoplus_{l=0}^N \tilde{\rho}_l, \quad (\text{A.2})$$

where each $\tilde{\rho}_l$ is a subnormalised state on the subspace containing l excitations of σ_z , i.e., \mathcal{H}_l . Let every $\tilde{\rho}_l$ be σ^z -Schmidt decomposable, and let the system evolve according to a quantum channel \mathcal{V}_τ that is σ^z -excitation conserving. Then the reduced state of every spin, at all times $\tau > 0$, will be σ^z -diagonal. Furthermore, the only components of ρ that contribute to

the reduced state of any given spin are the diagonal elements with respect to the $\{|0\rangle, |1\rangle\}^{\otimes N}$ basis.

Proof. At initial time, we may write every eigenvector of $\tilde{\rho}_l$ as

$$|\psi\rangle = \sum_m \alpha_m |\psi_m^l\rangle. \quad (\text{A.3})$$

where $|\psi_m^l\rangle = \bigotimes_{n=1}^N |a_m^n\rangle$, with $a_m^n \in \{0, 1\}$. Each $|\psi_m^l\rangle$ has $a_m^n = 0$ for l spins and $a_m^n = 1$ for $N-l$ spins. By construction, $\langle \psi_m^l | \psi_k^l \rangle = 0$ if $m \neq k$. As such, the contribution of $|\psi\rangle$ to the reduced state of the first spin will be $\sum_m |\alpha_m|^2 |a_m^1\rangle \langle a_m^1|$, which is clearly σ^z -diagonal, and only involves the elements of $\tilde{\rho}_l$ that are diagonal with respect to the $\{|0\rangle, |1\rangle\}^{\otimes N}$ basis. As a convex combination of σ^z -diagonal states are also σ^z -diagonal, then the reduced state of the first spin will also be σ^z -diagonal. The same argument will hold, *mutatis mutandis*, for all other spins. To show that this will hold true for all times, given a σ^z -excitation conserving quantum channel \mathcal{V}_τ , it is sufficient to show that the state $\mathcal{V}_\tau^l(|\psi\rangle \langle \psi|)$ is itself σ^z -Schmidt decomposable. This is evidently true, as

$$\mathcal{V}_\tau^l(|\psi\rangle \langle \psi|) = \sum_i L_i |\psi\rangle \langle \psi| L_i^\dagger, \quad (\text{A.4})$$

with $L_i |\psi\rangle = \sum_m \alpha'_m |\psi_m^l\rangle$. \square

Now we prove a sufficient condition under which the reduced state of every spin in an N -partite system will be thermal with respect to the Hamiltonian σ^z , with a temperature less than or equal to T . For the proof it will be simpler to use the ratio of probabilities of thermal states instead of temperature. We therefore use the following equivalence:

$$\frac{\langle 1 | \chi(T') | 1 \rangle}{\langle 0 | \chi(T') | 0 \rangle} \geq q \iff T' \leq T, \quad (\text{A.5})$$

where $\chi(T)$ is defined as in Eq. (II.3), and q, T, T' are all non-negative numbers.

Lemma A.2. *Let a quantum system composed of N spin-half systems be prepared in a state*

$$\rho = \bigoplus_{l=0}^N \tilde{\rho}_l, \quad (\text{A.6})$$

where each $\tilde{\rho}_l$ is a subnormalised state on the subspace containing l excitations of σ_z , i.e., \mathcal{H}_l . Furthermore, let ρ be diagonal with respect to the basis $\{|0\rangle, |1\rangle\}^{\otimes N}$, with \mathbf{r}_l a vector composed of these diagonal elements. In this case, \mathbf{r}_l is the spectrum of $\tilde{\rho}_l$, i.e., $\mathbf{r}_l = \lambda(\tilde{\rho}_l)$. Finally, let the probe evolve according to a σ^z -excitation conserving quantum channel \mathcal{V}_τ

that is also unital, i.e., $\mathcal{V}_\tau(\mathbf{1}) = \mathbf{1}$. If for all i, j, l , the condition

$$\frac{\mathbf{r}_l(i)}{\mathbf{r}_{l+1}(j)} \geq q \quad (\text{A.7})$$

is satisfied, where $\mathbf{r}_l(i)$ signifies the i^{th} element of the vector \mathbf{r}_l , and $q > 0$, then the reduced state of every spin for all times $\tau > 0$, $\rho_n(\tau)$, will be thermal with respect to the Hamiltonian σ^z , and with a temperature less than or equal to T .

Proof. Due to Lemma A.1, the reduced state of every spin will be σ^z -diagonal at all times, which is a necessary condition for it to be assigned a temperature. Moreover the only elements of ρ contributing to the elements of the reduced state of any spin are given by the vectors \mathbf{r}_l . By ordering each of these vectors appropriately, we can show that

$$\frac{\langle 1|\rho_n|1\rangle}{\langle 0|\rho_n|0\rangle} = \frac{\sum_{l=0}^{N-1} \sum_{i=1}^{K_l} \mathbf{r}_l(i)}{\sum_{l=1}^N \sum_{i=1}^{K'_l} \mathbf{r}'_l(i)}. \quad (\text{A.8})$$

We note that \mathbf{r}_l and \mathbf{r}'_l have the same elements, but with a different ordering. Also, for each l , $K_l + K'_l$ equals the dimension of the l -excitation subspace, given as

$$\dim(\mathcal{H}_l) = \binom{N}{l} := \frac{N!}{l!(N-l)!}, \quad (\text{A.9})$$

with

$$K_l = \binom{N-1}{l}, \quad K'_l = \binom{N-1}{l-1}. \quad (\text{A.10})$$

We note that $K_N = K'_0 = 0$. From this observation, it will be simple to deduce that for all $l \in \{0, \dots, N-1\}$, we have $K_l = K'_{l+1}$. As a consequence of Eq. (A.7), and the above observations, it therefore follows that for each $l \in \{0, \dots, N-1\}$,

$$\sum_{i=1}^{K_l} \mathbf{r}_l(i) \geq q \sum_{i=1}^{K'_{l+1}} \mathbf{r}'_{l+1}(i). \quad (\text{A.11})$$

As such, Eq. (A.8) will obey the inequality

$$\frac{\langle 1|\rho_n|1\rangle}{\langle 0|\rho_n|0\rangle} \geq q \frac{\sum_{l=1}^N \sum_{i=1}^{K'_l} \mathbf{r}'_l(i)}{\sum_{l=1}^N \sum_{i=1}^{K'_l} \mathbf{r}'_l(i)} = q. \quad (\text{A.12})$$

Therefore, given the stated conditions on the initial state of the system, the reduced state of every spin will be thermal with respect to the Hamiltonian σ^z , with a temperature less than or equal to T . To show that this will be true at all later times, we note that the state of the probe, at time $\tau > 0$, will be given as

$$\begin{aligned} \mathcal{V}_\tau(\rho) &= \bigoplus_{l=0}^N \mathcal{V}_\tau^l(\tilde{\rho}_l), \\ &= \bigoplus_{l=0}^N \tilde{\rho}_l(\tau). \end{aligned} \quad (\text{A.13})$$

As \mathcal{V}_τ^l is unital, by Uhlmann's theorem [58] we know that the vector composed of the spectrum of $\tilde{\rho}_l$ majorizes that of $\tilde{\rho}_l(\tau)$, i.e.

$$\lambda(\tilde{\rho}_l) \succ \lambda(\tilde{\rho}_l(\tau)). \quad (\text{A.14})$$

Furthermore, it is trivial that the vector composed of the diagonal elements in any basis is majorized by that of the spectrum, i.e.

$$\lambda(\tilde{\rho}_l(\tau)) \succ \mathbf{r}_l(\tau). \quad (\text{A.15})$$

As $\mathbf{r}_l = \lambda(\tilde{\rho}_l)$, it follows therefore that for all l and $\tau > 0$,

$$\mathbf{r}_l \succ \mathbf{r}_l(\tau). \quad (\text{A.16})$$

Furthermore, since the above equation implies that

$$\mathbf{r}_l(\tau) = Q\mathbf{r}_l, \quad (\text{A.17})$$

where Q is a doubly stochastic matrix [59], then every element of $\mathbf{r}_l(\tau)$ is given as a convex combination of those in \mathbf{r}_l . Consequently, Eq. (A.7) is satisfied at all times and, hence, the reduced state of every spin in the system will be thermal with respect to σ^z , with a temperature less than or equal to T , at all times. \square

Now we show that if the initial state of the probe used in the cooling protocol is given as $\rho_{\mathcal{P}}^{(0)} = \bigotimes_{n=1}^N \chi(T_n)$, such that for all n , $T_n \leq T$, then the cooling protocol will always cool the thermal qubits, or leave them with the same temperature.

Theorem A.1. *Let the probe be initially prepared in the state*

$$\rho_{\mathcal{P}}^{(0)} = \bigotimes_{n=1}^N \chi(T_n), \quad (\text{A.18})$$

such that for all n , $T_n \leq T$. It follows that the cooling protocol will always cool a collection of K thermal qubits of temperature T , or leave them the same, irrespective of the waiting times $\{\tau_k\}_k$ and number of thermal qubits K .

Proof. We may write the composition of K thermal qubits and the probe as

$$\rho = \bigotimes_{n=1}^{K+N} \chi(T_n), \quad (\text{A.19})$$

such that for all $n \in \{1, \dots, K\}$, $T_n = T$, whereas for all $n \in \{K+1, \dots, K+N\}$, $T_n \leq T$. Clearly, the eigenvectors of ρ are product vectors from the basis $\{|0\rangle, |1\rangle\}^{\otimes N+K}$ and, as such, it can be decomposed into a direct sum of subnormalized states in different excitation subspaces, as in Eq. (A.6). Furthermore,

the vectors of the spectrum satisfy Eq. (A.7). Therefore as a consequence of Lemma A.2, if the total system of probe plus thermal qubits evolves according to a unital quantum channel that is σ^z -excitation conserving, then the reduced state of every thermal qubit will be thermal with respect to the Hamiltonian σ^z , with a temperature less than or equal to T . Every stage of the cooling protocol, of course, is determined by the quantum channel defined in Eq. (II.5) acting on the compound system of thermal qubit k and the probe. This quantum channel is both unital and σ^z -excitation conserving. \square

Appendix B: The unique stationary state of the probe

We wish to show that the only stationary state of the probe, given the dynamics it undergoes with the thermal qubits, is $\chi(T)^{\otimes N}$. We first introduce some notation. We take the probe \mathcal{P} to be a collection of spins labeled by the integers $\{1, \dots, N\}$. As such, the reduced state of any subset of spins X is defined as $\rho_X := \text{tr}_{\mathcal{P} \setminus X}[\rho_{\mathcal{P}}]$, where $\mathcal{P} \setminus X$ is the complement of X in the set \mathcal{P} .

Theorem B.1. *Consider the composition of a ther-*

mal qubit \mathcal{Q} and the probe \mathcal{P} , in the state $\rho_{\mathcal{P}+\mathcal{Q}} = \chi(T) \otimes \rho_{\mathcal{P}}$. Let the evolution of the system be determined by the quantum channel \mathcal{V}_τ defined in Eq. (II.5) as

$$\mathcal{V}_\tau = \text{SWAP} \circ e^{\tau \mathcal{L}}, \quad (\text{B.1})$$

where \mathcal{L} is defined in Eq. (II.1) and SWAP is a possibly imperfect swap operation between the thermal qubit and the first spin of the probe. Then $\mathcal{V}_\tau(\rho_{\mathcal{P}+\mathcal{Q}}) = \rho_{\mathcal{P}+\mathcal{Q}}$ if and only if $\rho_{\mathcal{P}} = \chi(T)^{\otimes N}$.

Proof. We note that the SWAP operation can be generated by a time-controlled Heisenberg interaction H_I between the thermal qubit and the first spin of the probe, becoming perfect in the limit as this interaction becomes infinitely stronger than the intra-probe couplings. As such, in the absence of dephasing, $\rho_{\mathcal{P}+\mathcal{Q}}$ is stationary with respect to \mathcal{V}_τ if and only if it commutes with the XXZ spin network Hamiltonian, given in Eq. (II.2), that governs the total compound system. Let us denote this total Hamiltonian as $H = H_I + H_{\mathcal{P}} + H_{\mathcal{Q}}$, and by defining $\mathcal{P} + \mathcal{Q} := \{0, \dots, N\}$, where the thermal qubit is labeled 0, expand $\rho_{\mathcal{P}+\mathcal{Q}}$ in the Pauli basis as

$$\rho_{\mathcal{P}+\mathcal{Q}} := \sum_{a=0}^3 r_n^a \sigma_n^a \otimes O_{\mathcal{P}+\mathcal{Q} \setminus n}^a, \quad (\text{B.2})$$

for any spin $n \in \mathcal{P} + \mathcal{Q}$. Here $\sigma^0 = \mathbf{1}$, $\sigma^1 = \sigma^x$, $\sigma^2 = \sigma^y$, and $\sigma^3 = \sigma^z$. It is simple to verify that $\sum_{a=0}^3 r_0^a \sigma_0^a = \chi(T)$. As such, we have

$$[\chi(T) \otimes \rho_{\mathcal{P}}, H]_- = \sum_{n=0, m>n}^N J_{n,m} \sum_{a,b=0}^3 \sum_{c=1}^3 (1 + \delta_{c,3}(\Delta_{n,m} - 1)) r_n^a r_m^b [\sigma_n^a \otimes \sigma_m^b, \sigma_n^c \otimes \sigma_m^c]_- \otimes O_{\mathcal{P}+\mathcal{Q} \setminus \{n,m\}}^b. \quad (\text{B.3})$$

The entire expression vanishes only if the summands vanish individually for each n and m . We now introduce the identity

$$[A \otimes B, C \otimes D]_- = \frac{1}{2} ([A, C]_- \otimes [B, D]_+) + \frac{1}{2} ([A, C]_+ \otimes [B, D]_-), \quad (\text{B.4})$$

where $[\cdot, \cdot]_+$ is the anti-commutator, and the relations

$$[\sigma^a, \sigma^b]_- = 2i\epsilon_{abc}\sigma^c, \quad (\text{B.5})$$

$$[\sigma^a, \sigma^b]_+ = 2\delta_{ab}\mathbf{1}. \quad (\text{B.6})$$

Here $\epsilon_{abc} = 1$ (respectively -1) with $\{a, b, c\}$ a cyclic (respectively anti-cyclic) permutation of $\{x, y, z\}$, and $\delta_{ab} = 1$ if $a = b$ and 0 otherwise. Using the above

identities, we see that the summand in the first line of Eq. (B.3), for $n = 0$ and $m = 1$ is

$$2i \sum_{a,b=1}^3 r_0^a r_1^b \epsilon_{abc} (\mathbf{1}_0 \otimes \sigma_1^c - \sigma_0^c \otimes \mathbf{1}_1) \otimes O_{\mathcal{P}+\mathcal{Q} \setminus \{0,1\}}^b. \quad (\text{B.7})$$

We only consider this term for the case of $n = 0$, as the only nonvanishing value of $J_{0,m}$ is when $m = 1$ by construction. The summands with $a = b$ clearly vanish, as in such cases we have $\epsilon_{aac} = 0$. The remaining summands cannot vanish if $O_{\mathcal{P} \setminus \{0,1\}}^b$ are not the same for all values of b . If $O_{\mathcal{P} \setminus \{0,1\}}^b$ are the same

for all values of b , however, then

$$\rho_{\mathcal{P}} = \sum_{b=0}^3 r_1^b \sigma_1^b \otimes \rho_{\mathcal{P} \setminus 1}, \quad (\text{B.8})$$

with $\sum_{b=0}^3 r_1^b \sigma_1^b = \rho_1$ describing a quantum state that could be different from $\chi(T)$. In such cases, however, $r_0^a r_1^b$ are all positive numbers, and the only way for Eq. (B.7) to vanish is if $r_0^a = r_1^a$ for all a . When this is satisfied, the summands with the values of a and b interchanged differ only by a sign change, and therefore cancel out. But this means that $\rho_1 = \chi(T)$. Hence, for the state to commute with the Hamiltonian, we must have

$$\rho_{\mathcal{P}} = \chi(T)_1 \otimes \rho_{\mathcal{P} \setminus 1}. \quad (\text{B.9})$$

Carrying out the same argument recursively for all $n \in \{1, \dots, N\}$, we prove that the only state $\rho_{\mathcal{P}}$, such that $\chi(T) \otimes \rho_{\mathcal{P}}$ commutes with the total XXZ Hamiltonian H , is

$$\rho_{\mathcal{P}} = \chi(T)^{\otimes N}. \quad (\text{B.10})$$

To including dephasing, the dissipator term of the Liouville super-operator in Eq. (II.1) must also vanish, i.e., we must show that

$$\sum_{n=1}^N \sigma_n^z \rho_{\mathcal{P}} \sigma_n^z - \rho_{\mathcal{P}} = 0. \quad (\text{B.11})$$

If $\rho_{\mathcal{P}} = \chi(T)^{\otimes N}$, with $\chi(T) = \frac{1}{2}\mathbb{1} + r^z \sigma^z$, then for each n we have

$$\begin{aligned} \sigma_n^z \rho_{\mathcal{P}} \sigma_n^z &= (\sigma_n^z \chi(T) \sigma_n^z) \otimes \left(\bigotimes_{m \in \mathcal{P} \setminus n} \chi(T)_m \right), \\ &= \rho_{\mathcal{P}}. \end{aligned} \quad (\text{B.12})$$

This shows that if $\rho_{\mathcal{P}} = \chi(T)^{\otimes N}$, then the probe is a stationary state with respect to the dynamics given in Eq. (B.1) also in the presence of dephasing. \square

Appendix C: Entropic inequalities

The von Neumann entropy of a system in state ρ is defined as

$$S(\rho) := \text{tr}[\rho \ln(\rho)], \quad (\text{C.1})$$

where $\ln(\cdot)$ is the natural logarithm. The increase in entropy of the probe at the k^{th} stage of the protocol is defined as

$$\Delta S_{\mathcal{P}}^{(k)} := S(\rho_{\mathcal{P}}^{(k)}) - S(\rho_{\mathcal{P}}^{(k-1)}), \quad (\text{C.2})$$

whereas the decrease in entropy of the k^{th} thermal qubit is

$$\Delta S_{\mathcal{Q}}^{(k)} := S_T - S(\rho_{\mathcal{Q}}^{(k)}), \quad (\text{C.3})$$

where we use $S_T := S(\chi(T))$ as the von Neumann entropy of the thermal qubit at temperature T . We now show that the increase in entropy of probe is at least as great as the decrease in entropy of the thermal qubit.

Lemma C.1. *Let the compound system of probe \mathcal{P} and k^{th} thermal qubit \mathcal{Q} be*

$$\rho_{\mathcal{P}+\mathcal{Q}}^{(k)} := \chi(T) \otimes \rho_{\mathcal{P}}^{(k-1)}. \quad (\text{C.4})$$

Let this system evolve according to the quantum channel defined in Eq. (II.5). The SWAP operation in this quantum channel is in general generated by a time-controlled Heisenberg interaction, and need not be perfect. Furthermore, let the initial state of the probe be

$$\rho_{\mathcal{P}}^{(0)} := \bigotimes_{n=1}^N \chi(T_n), \quad (\text{C.5})$$

such that for all n , $T_n \leq T$. It follows that

$$\Delta S_{\mathcal{P}}^{(k)} \geq \Delta S_{\mathcal{Q}}^{(k)} \geq 0. \quad (\text{C.6})$$

Proof. We denote the state of the compound system after the action of the quantum channel as $\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k)$. Due to the unitality of this quantum channel, and the fact that the compound system is initially in a product state, we know that

$$S(\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k)) \geq S(\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}) = S(\rho_{\mathcal{P}}^{(k-1)}) + S_T. \quad (\text{C.7})$$

This is due to Uhlmann's theorem [58, 59] and the subadditivity of the von Neumann entropy [9]. Furthermore, given the partial traces of the time-evolved compound system as given by Eq. (II.6), we can further use the subadditivity of the von Neumann entropy to show that

$$S(\rho_{\mathcal{P}}^{(k)}) + S(\rho_{\mathcal{Q}}^{(k)}) \geq S(\rho_{\mathcal{P}+\mathcal{Q}}^{(k)}(\tau_k)). \quad (\text{C.8})$$

By combining the above equations, we arrive at

$$\Delta S_{\mathcal{P}}^{(k)} \geq \Delta S_{\mathcal{Q}}^{(k)}. \quad (\text{C.9})$$

Finally, given the initial state of the probe and the dynamics in question, due to Theorem A.1 we know that $\Delta S_{\mathcal{Q}}^{(k)}$ is never negative. \square

Moreover, the total entropy increase of the probe poses an upper bound on the total entropy reduction

of the system of thermal qubits, for a cooling process of any length k . The total entropy reduction of the thermal qubits, up to stage k , is defined as

$$\Delta S_{\mathcal{Q}}^{\text{total}}(k) := \sum_{i=1}^k \Delta S_{\mathcal{Q}}^{(i)}. \quad (\text{C.10})$$

Theorem C.1. *Consider the setup of Lemma C.1. The total entropy reduction of the thermal qubits obeys the inequality*

$$\Delta S_{\mathcal{Q}}^{\text{total}}(k) \leq NS_T, \quad (\text{C.11})$$

the upper bound being realizable only if the probe is initially in the pure state $\rho_{\mathcal{P}}^{(0)} = |1\rangle\langle 1|^{\otimes N}$.

Proof. It follows from Lemma C.1 that for any k ,

$$\sum_{i=1}^k \Delta S_{\mathcal{P}}^{(i)} = S(\rho_{\mathcal{P}}^{(k)}) - S(\rho_{\mathcal{P}}^{(0)}) \geq \Delta S_{\mathcal{Q}}^{\text{total}}(k). \quad (\text{C.12})$$

As shown in Theorem B.1 the probe will pseudo-thermalize to a state with entropy NS_T . It therefore follows that the total entropy reduction of the thermal qubits obeys the inequality

$$\Delta S_{\mathcal{Q}}^{\text{total}}(k) \leq NS_T. \quad (\text{C.13})$$

The upper limit is achievable only if the probe is initially in a pure state. If this pure state is to satisfy the conditions required for always cooling, it has to be in the pure state $|1\rangle^{\otimes N}$. \square

Appendix D: NV-P1 interaction Hamiltonian

The quantization axis of P1 centers is the direction of magnetic field, while that of NVs is their orienta-

tion. The dipolar interaction between two spins after secular approximation has only an Ising interaction [50].

$$\begin{aligned} H_{int} &= -\frac{J_0}{r^3} [3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2] S_{NV}^z \otimes S_{P1}^z \\ &= -\frac{J_0}{r^3} q S_{NV}^z \otimes S_{P1}^z \end{aligned} \quad (\text{D.1})$$

where $J_0 = \frac{\mu_0 \gamma_1 \gamma_2 \hbar^2}{4\pi} \simeq (2\pi)52 \text{ MHz}\cdot\text{nm}^3$, \hat{z}_1 and \hat{z}_2 are quantization axis of NV and P1 centers respectively, \hat{r} is a unit vector directing from a NV to a P1 center, and $q = 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2$ (Appendix (E)).

Appendix E: NV-NV interaction Hamiltonian

We start with the universal dipolar interaction Hamiltonian with two spins labeled as 1 and 2 [52].

$$H_{int} = -\frac{J_0}{r^3} [3(\vec{S}_1 \cdot \hat{r})(\vec{S}_2 \cdot \hat{r}) - \vec{S}_1 \cdot \vec{S}_2] \quad (\text{E.1})$$

where $J_0 = \frac{\mu_0 \gamma_1 \gamma_2 \hbar^2}{4\pi} \simeq (2\pi)52 \text{ MHz}\cdot\text{nm}^3$ and \hat{r} is a unit vector directing from spin 1 to spin 2. Since the crystal field of diamond gives zero field splitting between $|m_s = 0\rangle$ and $|m_s = \pm 1\rangle$, a natural quantization axis of an NV is its own orientation. Therefore, we adopt the dual coordinate system $(\hat{x}_1, \hat{y}_1, \hat{z}_1), (\hat{x}_2, \hat{y}_2, \hat{z}_2)$ which corresponds to orientations of each NV. The interaction Hamiltonian can then be expanded with these vectors.

$$\begin{aligned} H_{int} &= -\frac{J_0}{r^3} \vec{S}_1^T \cdot \begin{bmatrix} 3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{x}_2) - \hat{x}_1 \cdot \hat{x}_2 & 3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{y}_2) - \hat{x}_1 \cdot \hat{y}_2 & 3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{z}_2) - \hat{x}_1 \cdot \hat{z}_2 \\ 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{x}_2) - \hat{y}_1 \cdot \hat{x}_2 & 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{y}_2) - \hat{y}_1 \cdot \hat{y}_2 & 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{z}_2) - \hat{y}_1 \cdot \hat{z}_2 \\ 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{x}_2) - \hat{z}_1 \cdot \hat{x}_2 & 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{y}_2) - \hat{z}_1 \cdot \hat{y}_2 & 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2 \end{bmatrix} \cdot \vec{S}_2 \\ &\simeq -\frac{J_0}{r^3} \vec{S}_1^T \cdot \begin{bmatrix} 3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{x}_2) - \hat{x}_1 \cdot \hat{x}_2 & 3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{y}_2) - \hat{x}_1 \cdot \hat{y}_2 & 0 \\ 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{x}_2) - \hat{y}_1 \cdot \hat{x}_2 & 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{y}_2) - \hat{y}_1 \cdot \hat{y}_2 & 0 \\ 0 & 0 & 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2 \end{bmatrix} \cdot \vec{S}_2 \end{aligned} \quad (\text{E.2})$$

The second line of E.2 is justified by the rotating wave

approximation (RWA) [60]. The NV coupling strength

is on the order of a few tens of kHz while zero field splitting (ZFS) of an NV is 2.88 GHz. Thus, the product of a fast rotating term (S_x, S_y) and a non-rotating term (S_z) is averaged out in the time evolution, which is consistent with the energy conservation.

To simplify further, terms in the Hamiltonian can be decomposed into symmetrical and anti-symmetrical combination of terms by introducing new variables - g^+, g^-, h^+, h^-, q [53]:

$$H_{int} = -\frac{J_0}{r^3} [(g^+ + g^-)S_1^x \otimes S_2^x + (g^+ - g^-)S_1^y \otimes S_2^y + (h^+ + h^-)S_1^x \otimes S_2^y + (h^+ - h^-)S_1^y \otimes S_2^x + qS_1^z \otimes S_2^z]. \quad (\text{E.3})$$

$$\begin{aligned} g^+ &= \frac{1}{2} [3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{x}_2) - \hat{x}_1 \cdot \hat{x}_2 + 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{y}_2) - \hat{y}_1 \cdot \hat{y}_2] \\ g^- &= \frac{1}{2} [3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{x}_2) - \hat{x}_1 \cdot \hat{x}_2 - 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{y}_2) + \hat{y}_1 \cdot \hat{y}_2] \\ h^+ &= \frac{1}{2} [3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{y}_2) - \hat{x}_1 \cdot \hat{y}_2 + 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{x}_2) - \hat{y}_1 \cdot \hat{x}_2] \\ h^- &= \frac{1}{2} [3(\hat{r} \cdot \hat{x}_1)(\hat{r} \cdot \hat{y}_2) - \hat{x}_1 \cdot \hat{y}_2 - 3(\hat{r} \cdot \hat{y}_1)(\hat{r} \cdot \hat{x}_2) + \hat{y}_1 \cdot \hat{x}_2] \\ q &= 3(\hat{r} \cdot \hat{z}_1)(\hat{r} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2 \end{aligned} \quad (\text{E.4})$$

Out of these, $g^-(S_1^x \otimes S_2^x - S_1^y \otimes S_2^y)$ and $h^+(S_1^x \otimes S_2^y + S_1^y \otimes S_2^x)$ are non-energy conserving terms that can be eliminated by RWA. The other two terms can also be simplified in our situation, where we restrict the dynamics to a two-dimensional subspace spanned by $|m_s = 0\rangle$ and $|m_s = 1\rangle$.

$$\begin{aligned} S_1^x \otimes S_2^x + S_1^y \otimes S_2^y &= (|01\rangle\langle 10| + |10\rangle\langle 01|) \\ S_1^x \otimes S_2^y - S_1^y \otimes S_2^x &= i(|01\rangle\langle 10| - |10\rangle\langle 01|). \end{aligned} \quad (\text{E.5})$$

The resulting Hamiltonian has the form of,

$$\begin{aligned} H_{int} &= -\frac{J_0}{r^3} [g^+(|01\rangle\langle 10| + |10\rangle\langle 01|) \\ &\quad + ih^-(|01\rangle\langle 10| - |10\rangle\langle 01|) \\ &\quad + q|11\rangle\langle 11|]. \end{aligned} \quad (\text{E.6})$$

Note that the strength of $S_1^x \otimes S_2^x$ and $S_1^y \otimes S_2^y$ are the same because the spin is rotating with the ZFS in the lab frame. This interaction Hamiltonian can be expressed in terms of pseudo spin- $\frac{1}{2}$ pauli operators for $|m_s = 0\rangle$ and $|m_s = 1\rangle$ states.

$$\begin{aligned} H_{int} &= -\frac{J_0}{r^3} [2g^+(\sigma_1^x \otimes \sigma_2^x + \sigma_1^y \otimes \sigma_2^y) \\ &\quad + 2ih^-(\sigma_1^x \otimes \sigma_2^y - \sigma_1^y \otimes \sigma_2^x) \\ &\quad + q\sigma_1^z \otimes \sigma_2^z] \\ &\quad + (\text{non-interacting terms}). \end{aligned} \quad (\text{E.7})$$

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