Berry phase quantum thermometer

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Abstract. We show how the Berry phase can be used to construct a high-precision quantum thermometer. An important advantage of our scheme is that there is no need for the thermometer to acquire thermal equilibrium with the sample. This reduces measurement times and avoids precision limitations.

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1. Introduction

The state of a point-like discrete energy level quantum system (e.g. an atom) interacting with a quantum field acquires a geometric phase [1] that is dependent on the state of the field. For pure field states, the phase encodes information about the number of particles in the field [2]. In particular, for initial squeezed states, the phase also depends on the squeezing strength [3]. If the field is in a thermal state, this geometric phase encodes information about its temperature [4], and so is used in a proposal to measure the Unruh effect at low accelerations [4].

In this paper, we use the Berry phase to construct a high-precision thermometer. The thermometer consists of an atomic interferometer and measures the temperature of a cold medium by comparison with a hotter thermal source of approximately known temperature. Since our scheme does not require the thermometer to reach thermal equilibrium with the sample, measurement times are shorter and precision limitations due to thermalization are avoided [5].

In principle, the thermometer that we introduce can be physically implemented in different ways. Here, as a simple example, we consider the point-like system to be an atom whose internal level structure is described by a quantum harmonic oscillator. The sample will be a thermal source modeled as a quantum scalar field contained in a cavity. The sampled field could correspond to many possible different physical situations: a phonon field, an infinite number of harmonic oscillators coupled to the atom (a way to model thermal reservoirs), the electromagnetic (EM) field in one dimension or a fixed polarization component of a full three-dimensional EM field.

In the scenario we consider, the most favorable precision is obtained for temperatures of cold sources three orders of magnitude below the reference temperature of the hot source. Moreover, our thermometer is very sensitive to temperature variations of the cold sample but almost insensitive to variations of the hot source, making this setting a very precise and reliable method of measuring low temperatures with high precision. The temperature range in which the thermometer operates optimally is determined by the natural frequency of the harmonic oscillator (in this case the atom). Therefore, the thermometer will be tuned to a particular temperature range by choosing an atom with a given energy gap. It is possible to make measurements within different temperature ranges by using atoms with various energy gaps. If one employs atoms with energy gaps going from 1 MHz to 1 GHz, it is possible to measure with high precision temperatures over a range of $10^{-1}$–$10^{-6}$ K. Our proposal is applicable in non-relativistic settings and is readily scalable using atoms with energy gaps of radio-frequency width to measure nanokelvin temperatures.
2. Geometrical phase

To construct the Berry-phase-based thermometer, consider an atomic interferometer in which a single atom follows two different paths (see figure 1). In one branch the atom goes through a cavity containing the cold sample, while in the second branch the atom moves through a cavity where the field is in a thermal state of a known temperature. In both arms the atom will acquire a Berry phase due to the interaction with the field, which will depend on the field’s temperature. The geometric phase difference measured at the output of the interferometer will determine the temperature of the cold sample.

We now proceed to compute the Berry phase acquired by an atom coupled to a quantum field inside a cavity. Let the atom follow a trajectory given by $x(t)$ and let its internal harmonic degrees of freedom be described by creation and annihilation operators $d^\dagger$ and $d$. We consider the interaction Hamiltonian between the atom and the field to be given by

$$H_I = \sum_m \lambda_m (d^\dagger e^{i\Omega t} + d e^{-i\Omega t})(f^\dagger_m u_m(x(t)) + h.c.).$$

The $\lambda_m$ are coupling constants; $\Omega$ and $t$ are the atom’s frequency and proper time, respectively. The field operators $f^\dagger_m$ are associated with the field mode solutions $u_m(x(t))$, which are labeled by the frequency index $m$. These functions are evaluated at the atom’s position, implying that the interaction between the field and the atom takes place locally. For simplicity, we are considering a $(1+1)$-dimensional field.

This model is a type of Unruh–DeWitt detector [6–10] that has been previously studied in other contexts [4, 11, 12] and whose effectiveness and suitability has been deeply analyzed in [13–15]. It generally describes a multi-level quantum emitter (typically an atom) interacting with an infinite number of harmonic oscillators that may correspond to a scalar field. This model describes any harmonic detector coupled to the field (e.g. the electronic or vibrational spectrum of a many-level atom is very well approximated by harmonic oscillators) and is strongly related to the standard two-level approaches typically used to model atoms coupled to a scalar or EM field. In fact, there is a well-known mapping between this model and the standard Jaynes–Cummings model via the Holstein–Primakoff transformations [16]; the equivalence of the two models for the relevant regimes can be seen in [17, 18].

Since we consider the field to be inside a cavity, the mode frequencies $m$ are discrete. For small cavities the mode separation is large, and in this case it is possible to assume that the atom
interacts effectively only with a single near-resonant cavity mode. In this case the analysis is considerably simplified and the Hamiltonian becomes

$$H_I = \lambda (d^\dagger e^{i\Omega t} + d e^{-i\Omega t})(f^\dagger u(x(t)) + h.c.)$$

(2)

and has been empirically verified [19, 20]. For a cavity of length $L$ with walls placed at $x = \pm(L/2)$, the normalized mode function is given by

$$u(x(t)) = \frac{1}{\sqrt{\omega L}} \sin(\omega [x + (L/2)]) e^{-i\omega t},$$

(3)

where $\omega$ is the field frequency. We assume the atom’s trajectory is orthogonal to $x$ such that $x(t) = \text{const}$. The effective coupling $\lambda$ seen by the atom would depend on the configuration of the cavity. We will work in the standard coupling regime $\lambda \approx 1–1.5 \text{kHz}$ [19].

The most convenient procedure for computing the Berry phase is to operate in a mixed picture [4], where the atom’s free evolution is absorbed in the atom’s operators. This situation is mathematically more convenient for Berry phase calculations and the Hamiltonian (2) can be diagonalized analytically; its eigenstates are the dressed states $U^\dagger|m, n\rangle$, where $|m, n\rangle$ are eigenstates of the non-interacting Hamiltonian $H_0(\nu_f, \nu_d) = \frac{1}{2} \nu_d d^\dagger d + \nu_f f^\dagger f$ and $U = S_f S_d D_f d \hat{S}_d R_f$. The operators

$$D_{f/d} = \exp \left[ s(f^\dagger d - fd^\dagger) \right], \quad S_f = \exp \left[ \frac{1}{2} u(f^\dagger f - f^2) \right],$$

$$S_d = \exp \left[ \frac{1}{2} v(d^2 - d^\dagger)^2 \right], \quad \hat{S}_d = \exp \left[ p (d^2 - d^\dagger)^2 \right],$$

$$R_f = \exp \left( -i\varphi f^\dagger f \right)$$

(4)

are well known in quantum optics and are the two-mode displacement, single-mode squeezing and phase rotation operators [19], respectively.

Obtaining the values of the parameters characterizing the squeezing, displacement and rotation as functions of $(\lambda, \Omega, \omega)$ entails diagonalization of the Hamiltonian. Since the eigenstates are $U^\dagger|m, n\rangle$, this is a straightforward but tedious way of carrying out basic algebraic manipulations, yielding four independent parameters — $v$, $\varphi$, $\nu_d$ and $\nu_f$ — as relatively complicated functions of $(\lambda, \Omega, \omega)$. With these results it is now possible to calculate the Berry phase acquired by the atom due to its interaction with the field.

Berry showed that an eigenstate of a quantum system acquires a phase, in addition to the usual dynamical phase, when the parameters of the Hamiltonian are adiabatically and cyclically varied with time [1, 21, 22]. In the case of a point-like atom interacting with a quantum field, the motion of the atom in space-time gives rise to Berry’s phase, provided that the changes in the Hamiltonian are adiabatic and cyclic. This phenomenon was recently exploited to propose an experiment to detect the Unruh effect with much smaller accelerations than previously considered [4]. Here we use it to build a quantum thermometer: when an atom interacts with a thermal state of a bosonic field, the geometric phase acquired is a function of the temperature of the probed state.

The Berry phase acquired by the eigenstate $|\psi(t)\rangle$ of a system, whose Hamiltonian depends on $k$ cyclically, is obtained by adiabatically varying the vector of parameters: $R = (R_1(t), \ldots, R_k(t))$:

$$i\gamma = \oint_R A \cdot dR,$$

(5)

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where $R(t)$ is a closed trajectory in the parameter space and

$$A = \left( (\psi(t)| \partial_R \psi(t)) , \ldots , (\psi(t)| \partial_R |\psi(t)) \right).$$

(6)

As a first step, we calculate the Berry phase acquired by an eigenstate of the Hamiltonian under cyclic and adiabatic evolution of the parameters $\{v, \varphi, v_f, v_d\}$. The only non-zero component of $A$ is $A_v = \langle 0, n | S_a S_b D_{ab} R_a \partial_\varphi (R_b D_{ab}^{\dagger} S_a^{\dagger} S_b^{\dagger}) |0, n \rangle$, and so the Berry phase acquired by an eigenstate $U^\dagger |0, n\rangle$ is

$$\gamma_{ln} = 2\pi \left[ \frac{v_d n \sinh(2v) \cosh[2(C - v)]}{v_f \sinh[2(C - v)] + v_d \sinh(2v)} + \gamma_{l0} \right],$$

(7)

where $C = \frac{1}{2} \ln \left( \frac{v_f}{v_d} \right)$ with $v_f/v_d > e^{2v}$ and

$$\gamma_{l0} = \frac{v_f \sinh^2 v \sinh[2(C - v)] + v_d \sinh(2v) \sinh^2(C - v)}{v_f \sinh[2(C - v)] + v_d \sinh(2v)}.$$ (8)

Note that the time dependence of the Hamiltonian that gives rise to the Berry phase is given by an eigenstate $U^\dagger |\psi_i\rangle \otimes |\psi_f\rangle$, where $|\psi_i\rangle$ is the Berry phase accrued by an eigenstate $|\psi\rangle$ of the Hamiltonian when it interacts with a field in a thermal state. The density matrix of a thermal state with temperature $T$ is given by [4]

$$\rho_T = \bigotimes_\omega \frac{1}{\cosh^2 r_T} \sum_{n_\omega} \tanh^{2n} r_T |n_\omega\rangle \langle n_\omega| ,$$

where

$$\tanh r_T = \exp \left( -\frac{\hbar \omega}{2k_B T} \right).$$

We assume that, before the atom–field interaction is turned on, the system is in the mixed state $|0\rangle \langle 0| \otimes \rho_T$. If the interaction is then turned on adiabatically, the state evolves into $\rho = U^\dagger |0\rangle \langle 0| \otimes \rho_T) U$.

3. Quantum thermometer

We are now interested in calculating the Berry phase acquired by the atom when it interacts with a field in a thermal state. The density matrix of a thermal state with temperature $T$ is given by [4]

$$\rho_T = \bigotimes_\omega \frac{1}{\cosh^2 r_T} \sum_{n_\omega} \tanh^{2n} r_T |n_\omega\rangle \langle n_\omega| ,$$

where

$$\tanh r_T = \exp \left( -\frac{\hbar \omega}{2k_B T} \right).$$

We assume that, before the atom–field interaction is turned on, the system is in the mixed state $|0\rangle \langle 0| \otimes \rho_T$. If the interaction is then turned on adiabatically, the state evolves into $\rho = U^\dagger |0\rangle \langle 0| \otimes \rho_T) U$.

The geometric phase $\gamma = \text{Re} \left( \eta \right)$ acquired by a mixed state $\rho = \sum_i \omega_i |i\rangle \langle i|$ after a cyclic and adiabatic evolution is given by $e^{i\eta} = \sum_i \omega_i e^{i\gamma_i}$ [23], where $\gamma_i$ is the Berry phase acquired by an eigenstate $|i\rangle$ of the Hamiltonian. We find that after a single cycle the state $\rho_T$ acquires the phase

$$e^{i\eta} = \frac{e^{i\gamma_{l0}}}{\cosh^2 r_T - e^{2\pi i G} \sinh^2 r_T} ,$$

(9)

where

$$G = \frac{v_d \sinh(2v) \cosh[2(C - v)]}{v_f \sinh[2(C - v)] + v_d \sinh(2v)}.$$ (10)

Therefore the Berry phase accrued by the state of the atom interacting with a thermal state is

$$\gamma_T = \text{Re} \eta = \gamma_{l0} - \text{Arg} \left( \cosh^2 r_T - e^{2\pi i G} \sinh^2 r_T \right).$$ (11)
Let us compare the Berry phase acquired by the same multi-level system interacting with two thermal sources at different temperatures. After a complete cycle in the parameter space (near resonance this means waiting a time \(2\pi \Omega^{-1}\)), the phase difference between these two atoms is equal to

\[
\delta = \text{Arg} \left( 1 - e^{-\frac{\hbar \nu}{k_B T_1}} - 2\pi iG \right) - \text{Arg} \left( 1 - e^{-\frac{\hbar \nu}{k_B T_2}} - 2\pi iG \right). 
\]

For realistic coupling values for atoms in cavities, the phase difference \(\delta\) is of considerable magnitude as shown in figure 2; it is also very sensitive to a particular range of temperatures, depending on the atomic gap. Adjusting this gap, we tune \(\delta\) to a particular temperature range, where we assume the hot source to have the temperature three orders of magnitude greater than the temperature of the target source. While \(\delta\) is quite sensitive to variations of the cold source, it is rather insensitive to changes in the hot source. Consequently, we can obtain high-precision measurements of the temperature of the cold source with almost no need to control the temperature of the hot one. In figure 3, we illustrate the effects of underestimating the temperature of the hot source on the precision of the measurement of \(\delta\). Large variations of the hot source temperature translate into very small variations of the measured phase, providing us with a high-precision thermometer, able to measure temperatures three orders of magnitude smaller than the hot source, over which no high-precision temperature control is required.

The Berry phase is always a global phase. In order to detect it, it is necessary to prepare an interferometric experiment. For example, a multi-level system in a superposition of trajectories that go through different thermal sources would allow for detection of the phase. Any
**Figure 3.** Relative error in the Berry phase (and therefore the determination of the temperature for the cold source) as a function of the relative error in determining the temperature of the hot source. As we see, the setting is very robust: huge changes of temperature of the hot source translate into small changes in the phase $\delta$.

Experimental setup in which such a superposition can be implemented would serve our purpose. A possible scenario can be found in the context of atomic interferometry. This technology has already been successfully employed to measure with great precision general relativistic effects such as time dilation due to the Earth’s gravitational field [24]. Paths (of slightly different length) can be chosen such that the dynamical relative phase cancels. It is sufficient that the dynamical phase difference throughout both trajectories be equal to or smaller than the geometric phase to allow for its detection.

Although such cancellation depends on the specific experimental setup, we find that even for a simple setting with current length metrology technology$^9$, we can control the relative dynamical phase with a precision several orders of magnitude smaller than the Berry phase acquired in one cycle.

Consider an atom moving with speed $v$ along a trajectory of length $L$ in the interferometer. Changing the path length with a precision of $\Delta L$ translates into changing the time of flight $T = L/v$ with a precision $\Delta T = \frac{v}{L} \Delta L = \frac{\Delta L}{v}$. Since the dynamical phase $\phi \propto \Omega T$, the resulting precision is approximately $\Delta \phi = \Omega \Delta T \approx \Omega \Delta T = \frac{\Omega \Delta L}{v}$. For $v \approx 100 \text{ m s}^{-1}$, $\Omega \approx 2 \text{ GHz}$ and $\Delta L \approx 1 \times 10^{-11} \text{ m}$ (see [20]), we obtain $\Delta \phi = 2 \times 10^{-4}$.

### 4. Possible limitations of the results

One might expect the atoms to lose coherence upon interacting with thermal sources, thus rendering the interferometry experiment useless. However, for the energy gaps and temperatures considered here, to have decoherence times comparable with the time for one cycle the thermal sources must be at temperatures several orders of magnitude above the ones we wish to measure.

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$^9$ Laserscale®, [www.gebotech.de/pdf/LaserscaleGeneralCatalog_en_2010_04.pdf](http://www.gebotech.de/pdf/LaserscaleGeneralCatalog_en_2010_04.pdf).

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Furthermore, for this formalism to be valid we require that the probability of finding the atom in an excited state after one cycle of evolution should be much smaller than 1 (weak adiabaticity), in turn implying coherence is not lost. In general, we expect this condition to fail in three regimes: small atomic gap, high temperatures or high coupling. When it holds, the interaction time of the multi-level atom with the thermal state should be short enough so that the only change in the atom’s state is that it acquires a global phase (dynamical + geometrical).

To investigate under what regimes the time evolution of the coupled field–atom system fulfills this weak adiabaticity condition, we check that the ground state of the atom for the Hamiltonian $H(t_0)$ evolves after a time $t-t_0$ (equal to one cycle of evolution) to the ground state of the Hamiltonian $H(t)$ when the field is in a thermal state\textsuperscript{10} [4], a fact easily demonstrated for realistic values of the coupling $\lambda$.

Solving the exact Schwinger equation (in the interaction picture)

$$\frac{d}{dt} \rho = -i[H_I, \rho]$$

numerically\textsuperscript{11}, we find that indeed, after a short period of time (approximately $10^4 \times 2\pi \Omega^{-1}$ for realistic couplings), the probability of finding the atom in the excited state cannot be distinguished from thermal noise, as expected since for short times a ground-state atom in a thermal bath transiting to an excited state will start a Rabi-like oscillation. For longer times the atom will thermalize.

However, for our purposes, we will only have the atoms interacting with the thermal bath for very short times (one cycle of evolution $t \approx 2\pi \Omega^{-1}$), and for these times we can see that for realistic values, the hypothesis holds perfectly. In figure 4, we can see that for

\textsuperscript{10} Note that for realistic values of the coupling, the energy gap between the ground state and the first excitation of the interaction Hamiltonian is almost equal (different by approximately one part per million) to the free Hamiltonian.

\textsuperscript{11} Numerical computations were carried out with Matlab differential equation solvers.

Figure 4. Probability of atomic excitation in a time of a small number of cycles.
(One cycle means $t = 2\pi/\omega$. For the 1 GHz case, $P < 10^{-9}$; for the MHz case, $P < 10^{-3}$.)
the cases considered, even in the worst-case scenario (1 MHz gap and 1 mK temperature) the probability of excitation is \( P \approx 10^{-3} \ll 1 \). In the best-case scenario considered (1 GHz gap and 1 K temperature) the probability is \( P \approx 10^{-9} \ll 1 \) and the approximation holds very well. In other words, the atoms interact with the thermal bath for a time short enough to avoid being excited; the only effect of the interaction is that they acquire a phase dependent on the bath’s temperature.

Furthermore, the probability remains negligible for many evolution cycles. Even in the worst-case scenario considered, with realistic couplings and for an atomic gap of 1 GHz, the approximation remains valid for more than 100 cycles until thermal fluctuations render the hypothesis invalid. Considering one cycle of evolution, we need to demand that the transition probability \( P \ll 1 \). For the scenarios considered, we find for the best-case scenario \( P \approx 10^{-9} \) (GHz gap), while for the worst-case scenario \( P \approx 10^{-3} \) (MHz gap).

As a final remark, the thermometer has a target temperature range for which it is highly sensitive. As seen in figure 2, this range is typically of two to three orders of magnitude. To see an interference pattern in an atomic interference experiment that allows measurement of the target temperature, we also need a reference source at a fixed temperature for which we need to have some control but whose temperature may experience variations. To satisfy the necessary condition for reducing the influence of fluctuations in the reference source, in a manner shown in figure 3, we need the temperature of the source to be three orders of magnitude different from the target temperature. In principle, the source could be hotter or colder than the target, but of course, it is much easier to control warmer thermal sources than colder ones. As a final note, geometric phases can be generalized to non-adiabatic and non-cyclic cases. We are currently considering generalizations of the thermometer in these directions.

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Appendix. Diagonalization of the Hamiltonian

In this appendix, we will show how to diagonalize a single-mode Unruh–Dewitt-like Hamiltonian describing the interaction of a harmonic oscillator with a scalar field. We will do it in a way that will be very convenient in order to deal with the Berry phase calculations.

We consider a point-like detector endowed with an internal structure which couples linearly to a scalar field \( \phi(x(t)) \) at a point \( x(t) \) corresponding to the world line of the detector. The interaction Hamiltonian is of the form \( H_I \propto \hat{X}(x(t)) \) where we have chosen the detector to be modeled by a harmonic oscillator with frequency \( \Omega_1 \). In this case the operator \( \hat{X} \propto (d^\dagger + d) \) corresponds to the detector’s position where \( b^\dagger \) and \( b \) are creation and annihilation operators. Considering that the detector couples only to a single mode of the field with frequency \( |k| = \omega \), the field operator takes the form \( \phi(x(t)) \approx \hat{\phi}_k(x(t)) \propto [f e^{i(kx - \omega t)} + f^\dagger e^{-i(kx - \omega t)}] \), where \( a^\dagger \) and \( a \) are creation and annihilation operators associated with the field mode \( k \). The Hamiltonian
is therefore

\[ H_T = \omega f^\dagger f + \Omega d^\dagger d + \lambda (d + d^\dagger)[f^\dagger e^{i(kx - \omega t)} + f e^{-i(kx - \omega t)}], \]

where \( \lambda \) is the coupling frequency. This Hamiltonian resembles the Unruh–DeWitt detector in the case where the atom interacts with a single mode of the field. Standard calculations involving the Unruh–DeWitt detector employ the interaction picture since this is the most convenient picture to calculate transition probabilities. However, we employ a mixed picture where the detector’s operators are time independent since this situation is mathematically more convenient for Berry phase calculations.

The Hamiltonian can be diagonalized analytically. The eigenstates are given by \( U^\dagger |N_a N_b\rangle \) where \( U = S_a(u, \theta_a) S_b(v, \theta_b) D_{ab}(s, \phi) \hat{S}_b(p) R_a(\varphi) \) and \( |N_a N_b\rangle \) are eigenstates of the diagonal Hamiltonian \( H_0(\omega_a, \omega_b) = \omega_a f^\dagger f + \omega_b d^\dagger d \) which determines the energy spectrum of the system. The operators are slightly more general versions of those given in equation (4):

\[
D_{ab} = D_{ab}(\chi) = \exp(\chi f^\dagger d - \chi^* f d^\dagger), \\
S_a = S_a(\alpha) = \exp(\alpha^* f^\dagger^2 - \alpha f^2), \\
S_b = S_b(\beta) = \exp(\beta^* d^\dagger^2 - \beta d^2), \\
\hat{S}_b = \hat{S}_b(p) = \exp[p (d^\dagger^2 - d^2)], \\
R_a = R_a(\varphi) = \exp(-i \varphi f^\dagger f),
\]

where \( \chi = s e^{i \varphi}, \alpha = \frac{1}{2} u e^{i \theta_a} \) and \( \beta = \frac{1}{2} v e^{i \theta_b} \). In order to satisfy the equation

\[ H_T(\omega_a, \omega_b, \alpha, \beta, \chi, \varphi) = U^\dagger H_0 U, \]

the following constraints must be satisfied:

\[
s = \arctan \left( \frac{\omega_a \sinh 2u}{\omega_b \sinh 2v} \right), \quad \phi = \theta_a = 0, \quad \theta_b = \pi, \quad u = C - v,
\]

where \( C = \frac{1}{2} \ln(\omega_a/\omega_b) \) with \( \omega_a/\omega_b > e^{2v} \). The dependence of the Hamiltonian parameters on the parameters in the unitaries which diagonalize it are given by

\[
\omega = \frac{\sinh 2v \left[ \cosh[2(C - v)] + \frac{\sinh[2(C - v)]}{\tanh 2v} \right]}{\omega_a^{-1} \sinh 2v + \omega_b^{-1} \sinh[2(C - v)]}, \\
\Omega = \sqrt{\Omega^2 - 4Z^2}, \\
\lambda = e^{i\varphi} \sqrt{\frac{\omega_a \omega_b \sinh[2(C - v)] \sinh 2v}{\omega_b \sinh 2v + \omega_a \sinh[2(C - v)]}} \left[ \omega_a \cosh[2(C - v)] - \omega_b \cosh 2v \right], \\
\varphi = kx - \omega t, \\
\text{where } 2p = \arctanh[-2Z/\Omega] \text{ and } \\
\hat{\Omega} = \frac{\sinh 2v \left[ \omega_a^2 \sinh[4(C - v)]/\sinh 2v + \omega_b^2 \cosh 2v \right]}{\omega_b \sinh 2v + \omega_a \sinh[2(C - v)]} \left[ \omega_a \cosh[2(C - u)] - \omega_b \cosh 2u \right], \\
Z = \frac{1}{2} \frac{\sinh 2v \left( \omega_a^2 \sinh[4(C - v)]/\sinh 2v - \omega_b^2 \sinh 2v \right)}{\omega_b \sinh 2v + \omega_a \sinh[2(C - v)]}.
\]
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