Perturbative expansions for the fidelities and spatially-correlated dissipation of quantum bits

Lu-Ming Duan* and Guang-Can Guo*
Department of Physics and Nonlinear Science Center, University of Science and Technology of China, Hefei, Anhui 230026, People’s Republic of China

Abstract

We construct generally applicable short-time perturbative expansions for some fidelities, such as the input-output fidelity, the entanglement fidelity, and the average fidelity. Successive terms of these expansions yield characteristic times for the damping of the fidelities involving successive powers of the Hamiltonian. The second-order results, which represent the damping rates of the fidelities, are extensively discussed. As an interesting application of these expansions, we use them to study the spatially-correlated dissipation of quantum bits. Spatial correlations in the dissipation are described by a correlation function. Explicit conditions are derived for independent decoherence and for collective decoherence.

PACS numbers: 03.65.Bz, 89.70.+c, 42.50.Dv

*Electronic address: gcguo@sunlx06.nsc.ustc.edu.cn
In various fields of physics, the study of open quantum systems plays an important role. For example, decoherence was recognized as a major problem in realizing quantum computation \[1,2\]. Decoherence in quantum computers mainly results from the coupling of the quantum bit (qubits) to the environment. In general, it is not practical to look for exact solutions of these complicated systems, which consists of many spatially-correlated qubits interacting with a reservoir.

In this paper, we propose a perturbative approach to the study of open quantum systems. We construct generally applicable short-time perturbative expansions for some fidelities, such as the input-output fidelity, the entanglement fidelity, and the average fidelity. A similar perturbative expansion for coherence loss has been proposed in a recent paper \[3\], where the coherence loss is measured by the quantity \( \delta(t) = tr [\rho(t) - \rho^2(t)] \), in which \( \rho(t) \) indicates the reduced density of the system. Since \( \delta(t) \) has no direct physical meanings, in this paper, we choose the fidelities rather than \( \delta(t) \) as measures of decoherence. The fidelities are important quantities and have been widely used in quantum coding theory \[4-7\]. Through the perturbative expansions, we can get some general relations between the fidelities, such as the demonstration that the entanglement fidelity decays more rapidly than the average fidelity.

As an interesting application of this perturbative approach, we use it to study spatially-correlated dissipation of the qubits. The latter question is of great practical importance in designing quantum computers. Spatial correlations in pure dephasing of the qubits have been analyzed in Refs. \[8\] and \[9\]. However, the calculations of spatial correlations in general dissipation of the qubits, including the dephasing and the relaxation, are far more involved. This question is hard to solve either by any existing exact approaches or by solving the master equation.
[10]. Fortunately, here we show, through the perturbative expansions, spatial correlations in general dissipation of the qubits can be easily analyzed. Spatial correlations are described by a correlation function. From this result, we derive explicit conditions for independent decoherence and for collective decoherence, which are two important ideal circumstances in dissipation of the qubits.

1 Perturbative expansions for the fidelities

We consider an open quantum system, the total Hamiltonian of which is expressed as

$$H_T = H_0 + H_I + H_{env},$$

where $H_0$ and $H_{env}$ indicate the free Hamiltonians of the system and of the environment, respectively. $H_I$ is the interaction Hamiltonian between the system and the environment. First, we suppose that the system is initially in a pure state $|\Psi_0\rangle$. If there is no coupling between the system and the environment, at time $t$ the system is in the state

$$|\Psi (t)\rangle = e^{-iH_0t/\hbar} |\Psi_0\rangle = U_0 (t) |\Psi_0\rangle.$$  

But in reality, coupling of the system to the environment is inevitable. So at time $t$ the system is in fact described by the reduced density

$$\rho (t) = tr_{env} [\exp (-iH_T t/\hbar) \rho_{env} (0) \otimes |\Psi_0\rangle \langle \Psi_0| \exp (iH_T t/\hbar)],$$

where $\rho_{env} (0)$ is the initial density of the environment. Decoherence of the system due to this inevitable coupling can therefore be measured by the fidelity between the state (2) and (3), which has the form

$$F (t) = \langle \Psi_0| U_0^+ (t) \rho (t) U_0 (t) |\Psi_0\rangle.$$
The input-output fidelity is first defined in Ref. [4]. Here the definition is slightly modified to Eq. (4) to make it more suitable for measuring decoherence of the system.

In general, the reduced density $\rho (t)$ in Eq. (3) is hard to calculate. So there is difficulty to obtain the fidelity (4). Fortunately, in practice, the short-time behaviors of the system under dissipation are of most interests. Hence we do not need to exactly calculate the fidelity (4), but expand it into a short-time power series

$$F (t) = 1 - \frac{t}{\tau_1} - \frac{t^2}{\tau_2^2} - \cdots .$$

(5)

Following Eqs. (2), (3) and (4), it is not difficult to obtain the expansion coefficients. Up to order $t^2$ they read explicitly

$$\frac{1}{\tau_1} = 0$$

(6)

$$\frac{\hbar^2}{2\tau_2^2} = \langle \Psi_0 | tr_{env} \{ [H_I, [H_I, \rho_{env} (0) \otimes |\Psi_0\rangle \langle \Psi_0|] + \} [H_I, H_0] , \rho_{env} (0) \otimes |\Psi_0\rangle \langle \Psi_0|] |\Psi_0\rangle$$

(7)

$$= \langle H_I^2 \rangle_{s,env} - \langle \langle H_I \rangle_{s,env}^2 \rangle ,$$

where the symbol $\langle \cdots \rangle_{s,env}$ stands for the average value over the system and the environment, i.e., $\langle H_I \rangle_s = \langle \Psi_0 | H_I | \Psi_0\rangle$, and $\langle H_I \rangle_{s,env} = tr_{env} [\rho_{env} (0) \langle H_I \rangle_s]$. It is obvious that the expansion coefficients in Eq. (5) furnish characteristic times $\tau_n$ associated with decoherence processes involving $H_I$ to order $n$. The second-order coefficient $\frac{1}{\tau_2}$ is of special interest. It is the coefficient of the first non-trivial term in the expansion. Under the short-time approximation, $\frac{1}{\tau_2}$ measures the damping rate of the fidelity. In most cases, it is sufficient to consider the expansion up to order $t^2$.

In the above, we have assumed that the system is initially in a pure state. The input-output fidelity (4) does not apply for the mixed states. For a mixed
input state of the system, there are some fidelities defined, such as the entangle-
ment fidelity and the average fidelity [5]. These two fidelities are widely used in
quantum coding theory. Suppose $|\Psi_{rs}\rangle$ is a purification of the density $\rho_s$, i.e., the
state $|\Psi_{rs}\rangle$ satisfies $tr_r(\langle \Psi_{rs} | \langle \Psi_{rs} |) = \rho_s$, where the symbol $r$ denotes an ancillary
system. Similar to Eq. (4), the entanglement fidelity $F_e$ is defined as
\begin{equation}
F_e(t) = \langle \Psi_{rs}| U_0^+ (t) tr_{env} [\exp (-iH_T t/\hbar) \rho_{env}(0)] \otimes |\Psi_{rs}\rangle \langle \Psi_{rs}| \exp (iH_T t/\hbar)] U_0(t) |\Psi_{rs}\rangle
= 1 - \frac{t}{\tau_{1e}} - \frac{t^2}{\tau_{2e}} - \cdots,
\end{equation}
where $F_e(t)$ is furthermore subjected to a short-time power series expansion.

In Ref. [5], it has been proven that the entanglement fidelity defined above is
an intrinsic quantity of the system $s$, i.e., it does not depend on the specific
purification $|\Psi_{rs}\rangle$. From Eq. (5), we easily obtain
\begin{equation}
\frac{1}{\tau_{1e}} = 0
\end{equation}
\begin{equation}
\frac{\hbar^2}{2\tau_{2e}} = \langle H_I^2 \rangle_{s,env} - \langle \langle H_I \rangle^2 \rangle_{s,env}.
\end{equation}
Comparing Eqs. (6) (7) with Eqs. (9) (10), we see, the characteristic times
for the entanglement fidelity have the same forms as those for the input-output
fidelity. Only change is that the symbol $\langle H_I \rangle_s$ now means $\langle H_I \rangle_s = tr_s(\rho_s H_I)$.

So the entanglement fidelity is a natural extension of the input-output fidelity to
include the mixed input states. Eqs. (9) and (10) also show, the characteristic
times for the entanglement fidelity are all intrinsic properties of the system $s$.

The initial density of the system can be expressed as a mixture of pure states,
i.e.,
\begin{equation}
\rho_s = \sum_i p_i |\Psi_i\rangle \langle \Psi_i|,
\end{equation}
where $p_i$ satisfy $\sum_i p_i = 1$. The average fidelity $F_a$ is defined as

$$F_a(t) = \sum_i p_i F(|\Psi_i\rangle) = 1 - \frac{t}{\tau_{1a}} - \frac{t^2}{\tau_{2a}^2} - \cdots,$$

(12)

where $F(|\Psi_i\rangle)$ denotes the input-output fidelity for the pure state $|\Psi_i\rangle$. Since the expression (11) for the density $\rho_s$ is not unique, unlike the entanglement fidelity, the average fidelity is not solely defined for a definite $\rho_s$. From Eqs. (12) and (4), we get the characteristic times for the average fidelity

$$\frac{1}{\tau_{1a}} = 0,$$

(13)

$$\frac{\hbar^2}{2\tau_{2a}^2} = \langle H_I^2 \rangle_{\text{s,env}} - \left\langle \sum_i p_i \langle H_I \rangle_i^2 \right\rangle_{\text{env}},$$

(14)

where the symbol $\langle H_I \rangle_i$ indicates $\langle \Psi_i | H_I | \Psi_i \rangle$. Comparing Eq. (10) with Eq. (14), we get an interesting inequality. For any interaction Hamiltonians $H_I$, there is the operator inequality

$$\sum_i p_i \langle H_I \rangle_i^2 \geq \left( \sum_i p_i \langle H_I \rangle_i \right)^2 = \langle H_I \rangle_s^2.$$  

(15)

Hence Eqs. (10) and (14) yield

$$\frac{1}{\tau_{2e}} \geq \frac{1}{\tau_{2a}},$$

(16)

which suggests, the entanglement fidelity decays more rapidly than any average fidelities. In Ref. [5] it has been proven that the entanglement fidelity is always less than the average fidelity. Here we show this fact from another aspect.

2 Spatially-correlated dissipation of the qubits

As an interesting application of the perturbative approach to open quantum systems, we consider a practical question: decoherence in quantum computers. This
decoherence is due to the inevitable coupling of the qubits to the external environment. A few papers [1,2,8,9,11-14] have been published on this subject with some simplifications, such as omitting spatial correlations in the decoherence [11-14], or omitting relaxation of the qubits [8,9], or omitting both of them [1,2]. However, in real circumstances, such as in the ion-trapped quantum computers [15], relaxation of the qubits has notable contributions to decoherence [11-13]. On the other hand, spatial correlation properties of decoherence play an important role in the choice of the decoherence-reducing strategies. For independent decoherence and for collective decoherence, the decoherence-reducing strategies are quite different. So here we consider a more practical model of decoherence. This decoherence is described by a spatially-correlated amplitude damping, which includes both the dephasing and the relaxation of the qubits. The environment is modelled by a bath of oscillators with infinite degrees of freedom and the mode functions of the bath field are chosen as plane waves. The Hamiltonian describing this decoherence process has the form

\[ H = H_0 + \sum_{l=1}^L \sum_k \left[ \hbar g_k \left( e^{-ikr_l} a_k + e^{ikr_l} a_k^+ \right) \left( \lambda_1 \sigma_l^x + \lambda_2 \sigma_l^y \right) \right] + \sum_k \left( \hbar \omega_k a_k^+ a_k \right), \]  

(17)

where the Pauli operator \( \sigma_l \) represents the \( l \) qubit and \( a_k \) is the annihilation operator of the both mode \( k \). The symbol \( r_l \) denotes the site of the \( l \) qubit, and \( \lambda_1, \lambda_2 \) and \( g_k \) are coupling constants. \( H_0 \) in Eq. (17) describes the free evolution and the internal interaction of the qubits. The qubits, whether in memory or in quantum gate operations, can all be described by the Hamiltonian (17).

The Hamiltonian (17) is very complicated and it is hard to find its exact solutions. Fortunately, with the perturbative approach developed in the previous section, this complex system can be easily treated. To analyze the decoherence, we use the perturbative expansion for the entanglement fidelity, which returns to
the input-output fidelity if the initial state of the qubits is pure. The environment is supposed in thermal equilibrium, i.e., the initial density $\rho_{\text{env}}(0)$ of the bath has the following form in the coherent representation

$$\rho_{\text{env}}(0) = \prod_k \int d^2\alpha_k \frac{1}{\pi \langle N_{\omega_k} \rangle} \exp \left( - \frac{||\alpha_k||^2}{\pi \langle N_{\omega_k} \rangle} \right) |\alpha_k\rangle \langle \alpha_k|,$$

where the mean photon (or phonon) number

$$\langle N_{\omega_k} \rangle = \frac{1}{\exp \left( \frac{\hbar \omega_k}{k_B T} \right) - 1}.$$ 

With this density, and substituting the Hamiltonian (17) into Eq. (10), we obtain the decoherence rate $\frac{1}{\tau_{2e}}$ for this system

$$\frac{1}{\tau_{2e}} = \sum_{l_1, l_2 = 1}^L \Omega^2 (r_{l_1} - r_{l_2}) \langle \Delta A_{l_1} \Delta A_{l_2} \rangle_s,$$

where the operator

$$A_{l_i} = \lambda_1 \sigma_{l_i}^x + \lambda_2 \sigma_{l_i}^y \ (i = 1, 2)$$

and the spatial correlation function

$$\Omega^2 (r_{l_1} - r_{l_2}) = 2 \sum_k \left| g_k \right|^2 \cos \left[ k (r_{l_1} - r_{l_2}) \right] \coth \left( \frac{\hbar \omega_k}{2k_B T} \right).$$

Spatial correlation properties of the decoherence is completely determined by the correlation function $\Omega^2 (r_{l_1} - r_{l_2})$.

In the following, we discuss two important ideal circumstances for the correlation function. Eq. (22) can be rewritten as

$$\Omega^2 (r_{l_1} - r_{l_2}) = x \sum_k f(k) \cos \left[ k (r_{l_1} - r_{l_2}) \right],$$

where $f(k)$ is a normalized distribution satisfying $\sum_k f(k) = 1$ and $x$ is the normalization constant

$$x = 2 \sum_k \left| g_k \right|^2 \coth \left( \frac{\hbar \omega_k}{2k_B T} \right).$$
The expression of $f(k)$ is given by comparing (23) with (22). Its explicit form depends on the coupling coefficient $|g_k|^2$, whereas the latter is determined by the specific physical model for quantum computers. As a simplification, here we assume that the distribution $f(k)$ can be approximated by a Gaussian function with an expectation value $\bar{k}$ and a variance $\Delta k$, respectively. With this simplification, Eq. (23) reduces to

$$\Omega^2 (r_{l_1} - r_{l_2}) \approx x \cos [\bar{k}(r_{l_1} - r_{l_2})] \exp \left[ -\frac{1}{2} (\Delta k)^2 (r_{l_1} - r_{l_2})^2 \right].$$  \hspace{1cm} (25)

Suppose $d$ is distance between the adjacent qubits. If $d$ satisfies

$$(\Delta k) d >> 1,$$  \hspace{1cm} (26)

Eq. (25) yields

$$\Omega^2 (r_{l_1} - r_{l_2}) \approx x \delta_{l_1l_2},$$  \hspace{1cm} (27)

and from Eq. (20) the decoherence rate is simplified to

$$\frac{1}{\tau_{2e}} = \sum_{l=1}^{L} x \langle (\Delta A_l)^2 \rangle_s,$$  \hspace{1cm} (28)

which suggests, the total decoherence rate of $L$ qubits equals the sum of the decoherence rates of individual qubits. So in this circumstance, the qubits are decohered independently. Eq. (26) is the condition for independent decoherence. Most of the existing quantum error correction schemes are designed to correct for the errors induced by independent decoherence [16-24].

Apart from the independent decoherence, there is another ideal circumstance for the spatial correlation function. Suppose there are $2L$ qubits. Two adjacent qubits make up a qubit-pair. So we have $L$ qubit-pairs. The two qubits in the $l$ qubit-pair are indicated by $l$ and $l'$, respectively. If distance $d$ between the adjacent qubits satisfies the condition

$$\bar{k}d << 1 \text{ and } (\Delta k) d << 1,$$  \hspace{1cm} (29)
form Eq. (25) the spatial correlation function remains a constant for the two qubits in each qubit-pair. The decoherence rate $\frac{1}{\tau_{2e}}$ for $L$ qubit-pairs thus becomes

$$\frac{1}{\tau_{2e}^2} = \sum_{l_1, l_2=1}^{L} \Omega^2 (r_{l_1} - r_{l_2}) \langle \Delta (A_{l_1} + A'_{l_1}) \Delta (A_{l_2} + A'_{l_2}) \rangle_s.$$  \tag{30}

So under the condition (29) the two qubits in each qubit-pair are decohered collectively. In the collective decoherence, the decoherence rate is sensitive to the type of the initial states. If the initial state of the qubit-pairs is a co-eigenstate of all the operators $A_l + A'_l$, form Eq. (30), the second-order decoherence rate $\frac{1}{\tau_{2e}}$ reduces to zero. With these states decoherence of the qubits can therefore be much reduced. The co-eigenstates of the operators $A_l + A'_l$ are called the subdecoherent states. In fact, an arbitrary input state of $L$ qubits can be transformed into the corresponding subdecoherent state of $L$ qubit-pairs by the following encoding

$$| -1 \rangle \rightarrow | -1, +1 \rangle,$$
$$| +1 \rangle \rightarrow | +1, -1 \rangle,$$  \tag{31}

where $| -1 \rangle$ and $| +1 \rangle$ are two eigenstates of the operator $A_l$. Obviously, the encoded state is a co-eigenstate of the operators $A_l + A'_l$ with the eigenvalue 0. This encoding has been mentioned in [8] and [25] and extended in [26] and [27] to reduce decoherence in general circumstances. Here we derive the working condition (29) for this encoding.

In the above, $\bar{k}$ and $\Delta k$ are introduced phenomenologically. It is assumed that we have no knowledge about the coupling coefficient $|g_k|^2$. There is an interesting case in which the spatial correlation function can be calculated exactly. Consider one dimensional quantum computers. In the continuum limit, it can be assumed that

$$\sum_k |g_k|^2 \cdots \propto \int_0^\infty d\omega_k \omega_k e^{-\omega_k/\omega_c} \cdots$$  \tag{32}

where $\omega_c$ is the cut-off frequency whose specific value depends on the particular nature of the physical qubit under investigation. The form (32) of $|g_k|^2$ was also
used in Refs. [1] and [8]. We consider two circumstances. In the high temperature limit, i.e., \( T \gg \hbar \omega_c/k_B \), the correlation function (22) is simplified to

\[
\Omega^2 (r_{l_1} - r_{l_2}) \propto \frac{4k_B T}{\hbar \omega_c} \int_0^\infty d\omega_k e^{-\omega_k/\omega_c} \cos \left[ \frac{\omega_k}{v} (r_{l_1} - r_{l_2}) \right] \approx 4k_B T \frac{\omega_c^2}{1 + \omega_c^2 (r_{l_1} - r_{l_2})^2 / v^2},
\]

where \( v \) indicates velocity of the noise field. If distance \( d \) between the adjacent qubits satisfies \( d \gg v/\omega_c \), Eq. (33) tends to a delta-function and the qubits are therefore decohered independently. On the other hand, if \( d \ll v/\omega_c \), we have \( \Omega^2 (d) = \Omega^2 (0) \) and the adjacent qubits are then decohered collectively. In the low temperature limit, things are much similar. If \( T \ll \hbar \omega_c/k_B \), \( \coth \left( \frac{\hbar \omega_c}{2k_BT} \right) \approx 1 \) and Eq. (22) yields

\[
\Omega^2 (r_{l_1} - r_{l_2}) \propto 2 \int_0^\infty d\omega_k \omega_k e^{-\omega_k/\omega_c} \cos \left[ \frac{\omega_k}{v} (r_{l_1} - r_{l_2}) \right] = 2\omega_c^2 \frac{1 - \omega_c^2 (r_{l_1} - r_{l_2})^2 / v^2}{1 + \omega_c^2 (r_{l_1} - r_{l_2})^2 / v^2}. \]

Eq. (34) suggests, the above conditions for independent decoherence and for collective decoherence still hold at low temperature. The type of decoherence is mainly determined by distance between the adjacent qubits and by the cut-off frequency. Temperature of the environment hardly influences the decoherence type, though it determines the decoherence rate.

3 Summary

In this paper, we develop short-time perturbative expansions for some widely-used fidelities. From the expansions, we demonstrate some interesting relations between the fidelities. Perturbative expansions for the fidelities can be used for studying open quantum systems. As an example, we consider spatially-correlated dissipation of the qubits in the quantum computer. Spatial correlations in the
dissipation are described by a correlation function and we successfully derive the explicit conditions for independent decoherence and for collective decoherence. This example suggests, the perturbative expansions for the fidelities may be proven as a useful tool for studying open quantum systems.

Acknowledgment

This project was supported by the National Natural Science Foundation of China.
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