Experimental quantum Hamiltonian identification from measurement time traces

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ABSTRACT

Identifying Hamiltonian of a quantum system is of vital importance for quantum information processing. In this article, we realized and benchmarked a quantum Hamiltonian identification algorithm recently proposed (Zhang and Sarovar, 2014). we realized the algorithm on a liquid nuclear magnetic resonance quantum information processor using two types of working media with different forms of Hamiltonian. Our experiment realized the quantum identification algorithm based on free induction decay signals. We also showed how to process data obtained in a practical experiment. We studied the influence of decoherence by numerical simulations. Our experiments and simulations demonstrate that the algorithm is effective and robust.

1. Introduction

One critical task in quantum information processing is to characterize a quantum system so that it can be used for tasks, such as quantum teleportation [1,2], quantum cryptography [3,4], quantum computation [5,6], quantum simulation [7–11] and quantum metrology [12,13]. One way of fully characterizing a quantum system is doing quantum state tomography (QST) and quantum process tomography (QPT) [14–19]. The QPT approach requires an exponential number of experiments, which makes it difficult to be realized for even a small sized quantum system [20–23].

Meanwhile, various methods based on measurement time traces for Hamiltonian identification have been proposed for general quantum systems. Fourier transformation (FT) of only one measurement observable is used for a single qubit Hamiltonian identification [24]. Temporal evolution of concurrence measure of entanglement is employed to identify arbitrary two-qubit Hamiltonian [25]. Hamiltonian identification using dynamical decouplings was proposed [26]. Schemes of estimating the coupling parameters for a complex quantum network based on measurements of a small part of the network were proposed [27,28]. A basic and general quantum system identification framework has been established on how much knowledge that is attainable about a quantum system for a given experimental setup [29].

Recently, Zhang and Sarovar [30] proposed an efficient approach (the ZS approach) for identifying arbitrary Hamiltonian quantum dynamics, taking advantage of available prior knowledge of the system.

One typical dynamical system is the nuclear magnetic resonance (NMR) system, which is well described by quantum mechanics. Moreover, its control technology has been well developed during the 50 years since the birth of NMR. These factors make the NMR system an appealing quantum system for sophisticated manipulation. Therefore, NMR systems are widely used for quantum information processing [31,32]. To obtain the information of an NMR system, modern NMR spectrometers acquire the free induction decay (FID) signals, which are the measurement time traces of certain observables. Schemes based on FT (e.g. FT-NMR) of the FID signals, which is one of the most robust ways of processing FID, have been developed [33,34]. Because the ZS approach is based on measurement time traces for an arbitrary quantum system, the NMR spectrometer provides a practical and controllable system for demonstrating and benchmarking the ZS approach.

In this article, we implemented the ZS approach on an NMR quantum information processor and compared the result with that of FT approach. The experiments were performed with two types of working media with different Hamiltonian forms. Because of different Hamiltonian forms, we have to choose different measurement observables which require distinct experimental setups. Unlike works of NMR quantum computing in last two decades,
we started from the thermal equilibrium state rather than the pseudo-pure state, and directly processed the FID signals. Our experiments demonstrated that the ZS is an very efficient approach for Hamiltonian identification. We also analysed the influences of imperfect experiment conditions and decoherence on the results using numerical simulations.

2. Algorithm

Here we briefly describe the ZS approach. Suppose now we have an n-qubit quantum system with Hamiltonian $\hat{H}$. With the initial state $\rho(0)$, the system evolves, governed by the Hamiltonian $\hat{H}$. During the evolution, the expectation value of an observable $O$ at time $t$ is measured and recorded as $y(t)$. $y(t)$ is also called the measurement time trace of observable $O$. The Hamiltonian can be written in a parametrized form,

$$\hat{H} = \sum_m a_m \hat{x}_m,$$  
\[ (1) \]

where $a_m$ is the unknown parameters to be acquired, $\hat{x}_m \in S = \{ |x_1 \rangle \otimes |x_2 \rangle \otimes \cdots \otimes |x_N \rangle | x_i | = 1 \}$, and $|x_1 \rangle, |x_2 \rangle, \ldots, |x_N \rangle$ are the Pauli matrices $\sigma_x, \sigma_y, \sigma_z$, and $2 \times 2$ identity operator $I_2$. The number of the elements in $S$ is $4^n - 1$. However, if taken into consideration of a practical physical system, the number of the non-zero $a_m$'s can be significantly decreased.

All the elements in set $S$ are Hermitian operators. However, because of the physical constraints, only some of them can be easily measured, e.g., only the transverse magnetization in NMR, denoted by operator $\sigma_y$, can be observed. The temporal record of the expectations of such an observable $O$ can be collected, which is called the measurement time trace, and denoted by $y(t)$, then $y(t) = \text{Tr}(O(t))$, with $O(t)$ being the density matrix at time $t$. Let $y(t)$ be treated as an output of a linear system, and if we can find a set of $\{ C_0, A_0, x_0(0) \}$, which satisfies $y(t) = C_0 e^{A_0 t} x_0(0)$, we call this set a realization. For a certain output, various realizations can be obtained. Among these realizations, an invariant function, called transfer function $Y(s)$ exists, which is actually the Laplace transformation of the output, i.e.,

$$Y(s) = L(y(t)) = C_0 (sI - A_0)^{-1} x_0(0),$$ \[ (2) \]

where $s$ is Laplace variable, $L$ denotes Laplace transformation, and $I$ is the density matrix with the same dimension of $A_0$. The basic idea of the ZS approach is to find two realizations, one with all the unknown parameters $a_m$ (called realization 1, denoted by $\{ C_0, A_0, x_0(0) \}$) and the other (called realization 2, denoted by $\{ \mathbf{C}, \mathbf{A}, \mathbf{x}_0(0) \}$) with completely known numbers. With these two realizations, the coefficients of the Laplace variable $s$ can be compared, hence the unknown parameters can be obtained, i.e.,

$$\mathbf{C}(sI - \mathbf{A})^{-1} \mathbf{x}_0(0) = \mathbf{C}(sI - \mathbf{A})^{-1} \mathbf{x}_0(0).$$  
\[ (3) \]

A schematic of the ZS approach is shown in Fig. 1.

Realization 1 is obtained from the parameterized Hamiltonian $\hat{H}$, the observable $O$ and the initial state $\rho(0)$. The observable and the initial state are appropriately chosen arbitrarily according to the structure of a physical system. In such case, the vector $x_0(0)$ describes the initial system state $\rho(0)$, the matrix $\mathbf{A}$ describes the dynamical evolution driven by $\hat{H}$, and the matrix $\mathbf{C}$ predicts the measurement outcome $y(t)$ for the system state $x_0$.

Realization 2 is obtained solely by performing numerical methods, without relying on specific knowledge of the underlying system. One way to do so is the eigenstate realization algorithm (ERA) [35]. Technical details on how to obtain the two realizations is out of the scope of this article and shown in the Electronic Supplementary Material.

3. Experiments setup and results

The ZS approach was tested in a two-qubit and a three-qubit NMR system, which were implemented with $^{13}$C-labelled trichloroethylene (TCE) and $^{13}$C-labelled L-alanine (ALA) as the working media, respectively. The molecular structures and the thermal spectra of ALA and TCE are shown in Fig. 2.

The Hamiltonian of a liquid NMR system is $(h = 1)$

$$\hat{H}_{\text{NMR}} = \sum_{j=1}^{N} \pi_j \sigma_j^0 + \sum_{j=1}^{N} \frac{\pi_j}{2} \sigma_j^0 \cdot \sigma_j^1,$$ \[ (4) \]

where $2\pi\nu_i$ is the Larmor frequency for the $i$th spin, $J_{ij}$ is the indirect coupling constant between the $i$th and $j$th spin. In weak coupling, $| \nu_i - \nu_j | \gg | J_{ij} |$, which is valid for ALA, only the secular components of the scalar coupling survive. Hence the Hamiltonian of ALA is parameterized as

$$\hat{H}_{\text{ALA}} = a_1^0 \sigma_1^0 + a_3^0 \sigma_3^0 + a_2^0 \sigma_2^0 + a_1^2 \sigma_1^2 + a_2^2 \sigma_2^2 + a_3^2 \sigma_3^2,$$ \[ (5) \]

TCE is strongly coupled, and its parameterized Hamiltonian has a more complicated form,

$$\hat{H}_{\text{TCE}} = a_1^0 \sigma_1^0 + a_2^0 \sigma_2^0 + a_3^0 \sigma_3^0 + a_1^x \sigma_1^x + a_2^x \sigma_2^x + a_3^x \sigma_3^x,$$ \[ (6) \]

Once the Hamiltonian is parameterized, the observable can be decided. For TCE, $\sigma_3^y$ is chosen, and one qubit time trace is sufficient to identify the whole Hamiltonian because of strong coupling. Whereas for ALA, $\sigma_3^x + \sigma_3^y + \sigma_3^z$ has to be chosen as the observable because of the nature of the weak coupling Hamiltonian.

Then we prepare the initial states. Different forms of Hamiltonian require different experimental strategies. Different from NMR quantum computing with pseudo-pure initial state, Hamiltonian characterization should start directly from states that are easily prepared without knowing the Hamiltonian details, e.g., state $\rho(0) = \sum_i \sigma_i^0$. For ALA, we repeated the experiments for three times with three different initial states $\sigma_i^0, \sigma_i^2$, and $\sigma_i^z$, each corresponds to a different $x_0(0)$ for ALA. Choosing three initial states instead of one simplifies the data processing procedure. For TCE, a single input state $\sigma_3^z$ is enough.

After preparation, $\rho(0)$ starts to evolve under the system Hamiltonian, hence the macroscopic magnetization in NMR rotates. The rotation of the magnetization induces an electromagnetic wave which is received by a coil, and the signal received is called the free induction decay (FID) signal. The FID signal acquired by modern NMR spectrometers contains real and imaginary parts. The real part is

$$V_R(t) = x \text{Tr}(F_x \rho(t)),$$ \[ (7) \]

where $x$ is a coefficient related to the spectrometer, $\rho(t)$ is the density matrix of the system at time $t$ and $F_x = \sum_i \sigma_i^x$, where the summation includes the spins in the chosen observable. For TCE, only the second spin is observed and for ALA, all three qubits are observed. Thus the observable for TCE is $F_1^x = \sigma_1^x$ and for ALA is $F_3^x = \sigma_1^z + \sigma_2^z + \sigma_3^z$, which are exactly the observables we chose for ZS approach. The state $\rho(t)$ evolves as $\rho(t) = U \rho(0) U^\dagger$ with $U = \exp(-i \hat{H}_{\text{NMR}} t)$. From this discussion, we can see that the real
part of FID is the measurement time traces required by the ZS approach. After the FID signals are acquired, one can obtain a realization through ERA. However, data processing is not that simple due to the imperfections of a spectrometer. We have to deal with two problems, the dead time of the spectrometer and the unknown coefficient $x$. The dead time problem can be solved by doing phase correction on a spectrometer, and the unknown coefficient problem can be solved by scaling the FID properly. Detailed discussion about processing the experimental data before ERA is shown in the Electronic Supplementary Material.

Then ERA is performed to obtain a realization of the system on the processed FID signal. In our experiments and numerical simulations, 1600 points were used in ERA. Compared with the number of points (more than 16,000 in our experiments) FT-NMR utilized, the number of points ERA used is quite small. Once the realization is obtained, the transfer function can be calculated. By comparing the transfer functions, the parameters of the Hamiltonian are obtained. The results of the Hamiltonians of TCE and ALA are shown in Tables 1 and 2, respectively. From Tables 1 and 2, we can see that the parameters obtained from FT and that from the ZS approach agree well with each other. The absolute difference between the experimental values and the theoretical values of the chemical shifts for TCE is about 1 Hz (note there is a multiplication of $\pi$ between the chemical shift and the parameters $a_x$), which is only slightly larger than 0.5 Hz, the resolution of a modern FT-NMR spectrometer. The relative errors for the parameters related to the chemical shifts of ALA ($a_1^x - a_2^x$) are smaller than that of TCE ($a_1^x$), while at the same time, the relative errors for the parameters related to the spin-spin couplings of ALA ($a_1^y$, $a_2^y$) are larger than that of TCE ($a_2^z$). From these comparisons, we can say that the larger the value of a parameter, the more robust of the result. Despite the small relative error, the absolute differences of the chemical shifts of ALA are about 2 – 4 Hz, which is tiny but can be identified on modern NMR spectrometers. The extremely weak coupling can not be identified through looking at the whole spectrum (which means the observable is $\sigma_1^2 + \sigma_2^2 + \sigma_3^2$), both by using FT and the ZS approach. This is caused by the low resolution of a spectrum with a large spectral width (SW) and the decoherence time $T_2$. SW is a parameter decided before acquisition. Since the difference between the chemical shifts of C1 and C3 of alanine is large, to obtain the whole spectrum, the spectral width in Hz (SWH) has to be very large. The FID resolution in Hertz is $\text{FIDRES}=\text{SWH}/TD$. As $\text{FIDRES}$ is proportional to SWH, a large SWH means a large $\text{FIDRES}$, hence low resolution. Combined with the line broadening caused by decoherence, the weak coupling can not be identified on the spectrum of FT-NMR, which means that the information of $F_{13}$ is lost during the acquisition. To identify weak couplings, a small SWH (for ALA, it can be set to less than 100 Hz) should be chosen and the transmitter frequency should be set to the frequency of C1 or C3.

It is worth noting that only the absolute value of the parameters is provided in the Tables 1 and 2. For a signal $V(t) = \cos(\omega t)$, the result of FT will give frequencies $\omega$ and $-\omega$. Therefore, to identify the sign of chemical shifts, modern NMR spectrometers use quadrature detection [34], where the imaginary FID is generated and employed to assist identifying the frequencies, while during our experiments with the ZS approach, only $F_x$ is observed. The signs of the spin-spin couplings can not be identified by just obtaining the 1D spectrum. To identify these signs, additional experiments such as COSY-45 or spin polarization transfer are required. From the above discussion, we can see that the ZS approach gives almost the same amount of information as that provided by FT.

Besides FT, other methods, such as maximum entropy method [36,37], linear prediction [38,39] are also in common use. Though not as robust as FT, these methods have advantages in certain cases 

\(\text{See Bruker manuals of Topspin for details.}\)
such as when there are only a few data points. Since the data points used by the ZS approach is much smaller than that used by FT, our experiments show that the ZS approach is also applicable in such cases.

4. Decoherence

In NMR systems, decoherence is a common feature, which causes error in the ZS approach and leads information loss of parameter $T_2$. For practical experiments, the decoherence parameters are decided according to the spectrometer and the preparation of the sample. Therefore, to benchmark the ZS approach with the presence of decoherence, numerical simulations are employed.

![Molecule structure and spectrum for TCE (a) and ALA (b).](image)

Table 1
Experimental results for TCE. Relative error is defined as $|\text{FT result - ZS result}|/\text{FT result}$. The values of the parameters in FT result are obtained by numerically fitting the Fourier transformed spectra.

| $a_m$ | $X_m$ | FT result | ZS result | Relative error |
|-------|-------|-----------|-----------|----------------|
| $a_1$ | $\sigma_1^1$ | 1180.6     | 1179.4    | $1.05 \times 10^{-3}$ |
| $a_2$ | $\sigma_1^2$ | 1081.2     | 1082.5    | $1.25 \times 10^{-3}$ |
| $a_3$ | $\sigma_1^1\sigma_1^2 + \sigma_1^3\sigma_1^2$ | 161.9      | 162.6     | $4.24 \times 10^{-3}$ |

Table 2
Experimental results for ALA. Relative error is defined as $|\text{FT result - ZS result}|/\text{FT result}$. The values of the parameters in FT result are obtained by numerically fitting the Fourier transformed spectra.

| $a_m$ | $X_m$ | FT result | ZS result | Relative error |
|-------|-------|-----------|-----------|----------------|
| $a_1$ | $\sigma_2^1$ | 25723.3    | 25721.2   | $7.98 \times 10^{-6}$ |
| $a_2$ | $\sigma_2^2$ | 13876.7    | 13881.5   | $3.44 \times 10^{-4}$ |
| $a_3$ | $\sigma_3^1$ | 24745.6    | 24749.9   | $1.71 \times 10^{-4}$ |
| $a_4$ | $\sigma_3^1\sigma_3^2$ | 84.8       | 84.3      | $5.5 \times 10^{-3}$ |
| $a_6$ | $\sigma_3^2\sigma_3^3$ | 54.8       | 55.7      | $1.59 \times 10^{-2}$ |

In this article, all $T_2$ is actually $T_{2a} = T_{2b}$ and the inhomogeneity of the magnetic field. The inhomogeneity can be refocused by pulses. But when we acquire the FID, the inhomogeneity always exists and can not be eliminated. Thus all the $T_2$’s described in this article is $T_{2a}$, which can be straightly obtained by fitting the spectra.
Numerical simulations are performed to benchmark the influence of $T_2$. Using the $T_2$ model presented in Ref. [33], the output FID reads

$$V_{rd}(t) = \sum_{rs} F_{rs} \rho_{rs}(0) e^{i\omega_{rs} - \omega_{rs}t}, \quad (8)$$

where $F_{rs}$ ($\rho_{rs}(0)$) denotes the $r$-th and $s$-th ($s$-th and $r$-th) entry of $F$, $\omega_{rs}$ denotes the frequency between energy level $r$ ans $s$ and $\lambda_{rs}$ is the relaxation rate, i.e., $\omega_{rs} = \langle r | \hat{H}_{\text{NMR}} | r \rangle - \langle s | \hat{H}_{\text{NMR}} | s \rangle$ and $\lambda_{rs} = 1/T_2^s$. For simplicity and without loss of generality, all the spins relax at the same rate, so $T_2 = T_2$. In our simulations, the $T_2$'s are chosen to be from 0.01 to 0.1 s, which are shorter than $T_2$ of the systems. The couplings of ALA's are much smaller than that of TCE's, hence the errors brought by $T_2$ to the couplings of ALA's are much larger than that of TCE's, while on the contrary, the value of the chemical shifts of ALA's are much smaller than that of TCE's, hence the errors brought by the chemical shifts hardly exists between two spins. In summary, we have realized the ZS approach using an NMR quantum information processor with different work media. We showed that by choosing suitable observables for a given form of coupling, weak or strong coupling, the Hamiltonian can be efficiently identified using the ZS approach. Our experiments show that the ZS approach simplifies for the strong coupling systems, thus it can be used to identify these systems, such as solid-state and liquid crystal NMR systems, whose Hamiltonians are difficult to identify using FT approach. We also studied the influence of $T_2$ on the result of the ZS approach. The numerical simulation indicated that a very short $T_2$, implying a strong decoherence, can downgrade the ZS approach. However for a reasonable $T_2$, the ZS approach is efficient and robust.

Conflict of interest

The authors declare that they have no conflict of interest.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.scib.2017.05.013.

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The number of non-zero $a_n$’s for strong coupling system is $4^n - 1$, which is exponential, making QPT extremely hard to be performed. If without any prior knowledge of the system, all these $a_n$’s have to be identified using the ZS approach, which means the scale of the problem grows exponentially and the ZS approach is hardly scalable. However, because of the physical constraints on system energy, locality and structure, the number of the non-zero $a_n$’s would decrease rapidly, which significantly simplifies the problem. Taking the liquid NMR system Hamiltonian of Eq. (4) with weak coupling as an example, the number of the non-zero $a_n$’s is $O(n^3)$, which is polynomial, and can be further reduced because the indirect spin-spin coupling hardly exists between two spins that are four chemical bonds away. From this point of view, the ZS approach for the liquid NMR Hamiltonians is scalable as the number of qubits grows.

\*\*\* The number of non-zero $a_n$’s for strong coupling system is $(3 \times n^2 - n)/2$, which is also polynomial.
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