High Resolution non-Markovianity in NMR

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Memoryless time evolutions are ubiquitous in nature but often correspond to a resolution-induced approximation, i.e. there are correlations in time whose effects are undetectable. Recent advances in the dynamical control of small quantum systems provide the ideal scenario to probe some of these effects. Here we experimentally demonstrate the precise induction of memory effects on the evolution of a quantum coin (qubit) by correlations engineered in its environment. In particular, we design a collisional model in Nuclear Magnetic Resonance (NMR) and precisely control the strength of the effects by changing the degree of correlation in the environment and its time of interaction with the qubit. We also show how these effects can be hidden by the limited resolution of the measurements performed on the qubit. The experiment reinforces NMR as a test bed for the study of open quantum systems and the simulation of their classical counterparts.

Markov processes are defined as those in which the future of a system bears no correlation to its past. Such behavior is found in the description of phenomena as far apart as nuclear fission, the decay of an electronically excited atom, the growth of bacterial colonies or rabbit populations and the compound interests that dictate our debts, to name a few¹. In many situations, however, these time evolutions prove to be no more than a resolution-induced approximation, i.e. the interaction with external degrees of freedom (here called environment) produces correlations in the dynamics, but in a way that is too weak to be observed. Most of the times, such small deviations are either harmless and/or useless. However, there are meaningful counterexamples of both cases. For instance, many modern applications, such as the generation of large cryptographic keys or statistical sampling, rely on random number generators². Testing the independence of events in these devices is essential to guarantee the safety or fairness of these applications where even the tinniest correlations may be exploited. At the same time, controllable non-Markovian evolutions in small quantum systems have been thought of as promising tools for quantum information processing³⁷.

Markovian processes exist both in discrete time steps or as a continuous time evolution. A broadly used tool to discretize continuous Markovian dynamics are the so-called collisional models. In these models, the continuous stochastic process is replaced by a stroboscopic sequence formed by a series of discrete time steps defined, each one, by a collision between the system of interest and one of the particles forming its environment. The most famous application of collisional models in physics is the description of Brownian motion. In open quantum systems, collisional models have been extensively used as a test bed to investigate the details of system-environment interactions and the conditions under which non-Markovian dynamics arises⁸–¹⁷. Provided system and environment are initially uncorrelated, the evolution of the system \( \rho \) after \( j \) collisions with the environment \( \rho_\text{env} \) is given by

\[
\rho \rightarrow \rho' = \text{Tr}_\text{env} \left[ U_j U_{j-1} \ldots U_2 U_1 (\rho \otimes \rho_\text{env}) U_1^* U_2^* \ldots U_{j-1}^* U_j^* \right]
\]

where \( \text{Tr}_\text{env} \) denotes the partial trace over the environmental degrees of freedom and \( U_i \) is a global unitary operation due to the \( i \)-th collision. These models are simple to describe, easy to compute and have been recently simulated in optical setups where one degree of freedom of the photon is used as the quantum system and extra ones play the role of the environment¹⁸–²⁷.

In this paper, we implement a collisional model in Nuclear Magnetic Resonance (NMR) to investigate with high resolution the subtle non-Markovian evolution of a qubit subjected to the action of a structured and fully quantized external environment. This experimental technique has been very successful in the study of quantum

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between 0), the Ising coupling approximation was considered here. In the NMR experiment the spins being the natural Hamiltonian of the total system may be described as

\[
H = \sum_n \hbar (\omega_{0n} - \omega_z) I_n^z + \sum_{k=m} \hbar 2\pi J_{km} I_z^k \otimes I_z^m,
\]

where \(\omega_{0n}\) and \(\omega_z\) are the natural resonance frequency and the frequency of the rotating reference frame of the \(n\)-th spin, respectively. The first term represents the interaction of the spins with the static magnetic field applied along the \(z\) direction and the second term represents the interaction between the three spins, with \(J_{km}\) being the exchange integral. Here, \(I_n^z\) is the spin operator for the \(i\)-th spin, which for spin-1/2 corresponds to the Pauli matrix \(\sigma_z\) divided by 2. The physical parameters of our molecule are given in Methods. Since the individual chemical shifts of each spin place them far apart, i.e. their resonance frequencies are separated enough \(|\omega_{0n} - \omega_{0m}| \gg J_{km}\), the Ising coupling approximation was considered here. In the NMR experiment the spins also interact with oscillating magnetic fields that can be turned on and off at any frequency and different amplitudes, named radio frequency pulses (\(rf\)). These are responsible for the system control, and its interaction can be described by the Hamiltonian

\[
H_{rf} = \hbar \omega_z(t) [I_n^x \cos \phi + I_n^y \sin \phi],
\]

Information processes and in proofs of quantum principles as may be seen on refs 28–31. Here, the environment is composed of qubits that are initially correlated among themselves and collide, one at a time, with the target. We characterize non-Markovianity by calculating the Trace Distance \(D(\rho(t), \rho(t)) = \frac{1}{2} \text{Tr} \left| \rho(t) - \rho(t) \right|\) between two different states of the system as a function of time. It is well established that in a quantum stochastic evolution, \(D(\rho(t + \tau), \rho(t)) > D(\rho(t), \rho(t))\) implies non-Markovian behavior. The experiment features only two time steps because these are enough to establish all the properties we want to discuss. We measure \(D(\rho(t), \rho(t))\) after each collision as a function of the degree of correlation in the initial state of the environment and also as a function of the duration of each collision. Both quantities influence how much the dynamics deviates from a Markov process.

**Results**

We encoded the system–environment state in a sample of trifluoroiodoethylene (\(C_2F_3I\)). The NMR experiment was performed using a 500 MHz Varian spectrometer with the used sample diluted (~1%) in deuterated acetone (containing 97% of deuterium). A single molecule of \(C_2F_3I\) contains three atoms of fluorine with nuclear spin-1/2, each of them representing a qubit. One of the atoms will be regarded as the system and the other two as the environment. These spins interact with each other and with applied magnetic fields as well. We designed the experiment to implement the operations represented in the quantum circuit of Fig. 1: the first qubit, top line in the circuit, represents the system and the other lines are the qubits of the environment. The two qubits of the environment are initially prepared in a correlated state \(\rho_{env} = \frac{1}{2} [\{00\} \{00\} + \{11\} \{11\} + \frac{i}{\sqrt{2}} \{01\} \{01\} + \frac{1}{\sqrt{2}} \{10\} \{10\}]\) where “0” means spin up and “1” means spin down. In each collision, the state of the system undergoes a random-walk type of evolution, changing conditionally to the environment: environmental spin up induces a rotation in a certain direction (say, \(x\)), whereas environmental spin down induces a rotation in the orthogonal direction (\(y\)). This situation can be summed up by the unitary transformation \(U = e^{i\varphi x} \otimes [0] \{0\} + e^{i\varphi y} \otimes [1] \{1\}\) where \([j]\{k\}\) represents the internal state of an environmental particle and the exponentials dictate the system’s rotations.

In a NMR experiment the signal of each fluorine spin can be distinguished and singled manipulated. The natural Hamiltonian of the total system may be described as

\[
H = \sum_n \hbar (\omega_{0n} - \omega_z) I_n^z + \sum_{k=m} \hbar 2\pi J_{km} I_z^k \otimes I_z^m,
\]
Figure 2. The trace distance, $D(\rho_1(t), \rho_2(t)) = \frac{1}{2} \text{Tr} |\rho_1(t) - \rho_2(t)|$, was employed to witness non-Markovianity. The increase of $D(\rho_1(t), \rho_2(t))$ with time is a trace of non-Markovianity. (a) The plots show the dependence of $\Delta D$ with the strength of each collision $\eta$ for different values of the correlation parameter $q$ (0 for the solid blue line, 0.15 for the dashed orange line and 0.25 for the dash-dotted green line). $q = 0$ means that the two collisions were totally anti-correlated. (b) Detail of the $\Delta D$ curve for $\eta < 0.01$ and $q = 0$. In both plots the error bars were estimated taking into account every possible error source, as discussed in ref. 38. This led to the error determining the Bloch vector of the quantum state of $8\delta r = 5 \times 10^{-4}$, independent of the direction. By propagating this error, the error in $D(\rho_1(t), \rho_2(t))$ is given by $\Delta D = \delta r/\sqrt{2}$.

where $\phi(t)$ defines the pulse modulation and duration, and $\omega$ stands for its phase. The radio frequency pulses are used to perform rotations on the individual spins and by properly choosing these rotations, the quantum circuit from Fig. 1 can be implemented, as described in Methods.

The experiment was performed for two different initial states of the system, $\rho_1(0) = |0\rangle \langle 0|$ and $\rho_2(0) = |1\rangle \langle 1|$ (eigenstates of $\sigma_z$) and for the environment prepared in different states with different degrees of correlation $q$ ($q = 0, 0.15, and 0.25$). The theoretical analysis shows that for $q$ equals 0 and 0.15 these are the states that maximize $\Delta D$ for the chosen dynamics. For $q = 0.25$ there is a better pair of initial states but the eventual gain in $\Delta D$ (a factor of ~3) still lies within the current error bar. Using the rf pulses, the first and the second collisions were implemented, resulting in the system state $\rho_1(1) (\rho_2(1))$ and $\rho_1(2) (\rho_2(2))$, respectively. After each run, full state tomography was performed in the system.

Figure 2 shows the change in distance between the two initial states of the system after one and two collisions $\Delta D = D(\rho_1(2), \rho_2(2)) - D(\rho_1(1), \rho_2(1))$, as a function of the strength of each collision $\eta$ and for different degrees of correlation $q$ of the environmental state. For large enough interactions (larger $\eta$) and anti-correlation in the environment (smaller $q$), the collisions clearly generate a non-Markovian dynamics in the system ($\Delta D > 0$). The phenomenon, witnessed by the increase in $\Delta D$ also reflects a backflow of information from the environment to the system as time progresses.

It is also worth noticing that, as the non-Markovian effects induced by the collisions get smaller, either due to weaker collisions (small values of $\eta$) and/or environmental correlations ($q \rightarrow 1/2$), the uncertainties in $\Delta D$ due to the lack of resolution of the measurements become of the same order of $\Delta D$. This is true even for the strongest anti-correlation possible, $q = 0$. In this case, $\Delta D$, a sufficient but not necessary condition for non-Markovianity, does not provide a conclusive answer anymore, even though theory still predicts its vanishing only asymptotically with both parameters $\eta \rightarrow 0$ or $q \rightarrow 1/2$. Our resolution to conclusively guarantee non-Markovianity was limited to $\Delta D \sim 3.5 \times 10^{-4}$.

Discussions
Non-markovian dynamics of collisional models has been experimentally investigated in linear optical set-ups where the environment is simulated by semi-classical degrees of freedom. Usually, such experiments are performed in the photon counting regime and strong system-environment coupling. Our results put in evidence, in a controllable way, the role played by environmental correlations in the non-Markovian dynamics of quantum systems and also show how such effects may be hidden by the external noise brought in by unavoidable measurement uncertainties. We have shown how NMR can be exploited to study quantum dynamics of open systems using a bona fide quantum environment. NMR also presents a great advantage over previously used platforms when it comes to the versatility to design and control different environmental states and the precision to probe very small memory effects in the dynamics of quantum systems. For example, here we have observed variations of the Trace Distance of the order of 0.05%, a gain of at least two orders of magnitude over the typical optical experiments based on measurements of the polarization of photons. In photon polarization measurements, the signal-to-noise ratio is less than $\sqrt{N}$, where $N$ is the number of photons in each measurement. Such gain in precision can be essential, for example, if one would like to use a similar setup to test very small time correlations of black boxes such as quantum or classical random number generations, cryptographic machines and alike. Furthermore, one can also explore the capacity to dynamically change the environment during the interaction with the system and to directly interfere with the system during its time evolution in order to test and investigate very recent theoretical results in the field, such as the ones respectively published in refs 13 and 35.
Methods

Model. We consider a qubit $\rho$ that interacts with an environment from which it is initially decoupled. The interaction consists of consecutive collisions with qubits of the environment. During each collision, the system undergoes with equal probability an evolution $e^{i\eta xy}$ or $e^{i\eta yx}$ controlled by the internal state of the corresponding environmental particle. The map that describes the state of the system after collision one is $\Lambda_{10}(\cdot) = \frac{1}{2}[e^{i\eta xy}(\cdot) e^{-i\eta yx} + e^{i\eta yx}(\cdot) e^{-i\eta xy}]$. All the effects we are interested in can be observed with only two collisions described, in our case, by the general map $\Lambda_{20}(\cdot) = \frac{1}{2}\{e^{i\eta xy}(\cdot) e^{-i\eta yx} e^{-i\eta 00} + e^{i\eta yx}(\cdot) e^{-i\eta xy} e^{-i\eta 00} + e^{i\eta 00}(\cdot) e^{-i\eta 00} + e^{i\eta 00}(\cdot) e^{-i\eta 00}\}$, where $\eta$ defines their degree of correlation: for $q = 1$ (anti-correlated) and $q = 0$ (collisions are totally correlated (anti-correlated)). One possible way to generate this evolution is to consider two environments acting on the system in the following way $\rho_{\text{env}} = \frac{1}{2}\{\{00\} \{00\} + \{11\} \{11\}\} + \frac{1}{2}\{\{01\} \{01\} + \{10\} \{10\}\}$. (Note that a classically correlated state or the state $\frac{1}{2}\{\{00\} + \{11\}\} + \frac{1}{\sqrt{2}}\{\{01\} + \{10\}\}$ will render the same system evolution).

The interaction in each collision is described by a unitary transformation given by $U_{ii} = e^{i\eta xy} \otimes |0\rangle \langle 0| + e^{i\eta yx} \otimes |1\rangle \langle 1|$ which works as a conditional operation. The resulting state after one collision will be $\rho(1) = \Lambda_{10}(\rho(0)) = Tr_{\text{env}}[U_{ii} \otimes \rho(0) \otimes \rho_{\text{env}}(U_{ii}^\dagger \otimes I)] = \frac{1}{2}(e^{i\eta xy}(\rho(0))e^{-i\eta yx} + e^{i\eta yx}(\rho(0))e^{-i\eta xy})$, where $Tr_{\text{env}}$ denotes the partial trace over the environmental degrees of freedom. And after the second collision the evolution of the system reads $\rho(2) = \Lambda_{20}(\rho(0)) = Tr_{\text{env}}[U_{ii} U_{jj} \otimes \rho(0) \otimes \rho_{\text{env}}(U_{ii}^\dagger \otimes U_{jj}^\dagger)] = \frac{1}{2}[e^{i\eta xy}(\rho(0))e^{-i\eta yx} e^{-i\eta 00} + e^{i\eta yx}(\rho(0))e^{-i\eta xy} e^{-i\eta 00} + e^{i\eta 00}(\rho(0))e^{-i\eta 00} + e^{i\eta 00}(\rho(0))e^{-i\eta 00}]$. For simplicity, we chose these two specific operators, however, similar results would be observed had we used two different non-commuting operators. The parameters $\eta_{1}$ and $\eta_{2}$ are related to the strength of interaction and the time for one collision. We consider $\eta_{1} = \eta_{2} = \eta$, and considering a weak interaction and a short collision time, $\eta$ are assumed to be sufficiently small ($\eta < 0.1$) such that $e^{i\eta xy} \sim (1 + i\eta xy)$. In the first run of the experiment, the system interacts (collides with qubit one of the environment and, in the second run, the system repeats this collision and subsequently collides with qubit two of the environment. After each run, quantum state tomography is performed on the system in order to fully determine its state. Two orthogonal initial states of the system, $\rho(0) = |0\rangle \langle 0| + \rho(0) = |1\rangle \langle 1|$, are prepared so that the distance $D(\rho_{i}, \rho_{j})$ between them can be calculated after each run $j$ (orthogonality sets $D(\rho_{1}, \rho_{2}(0)) = 1$).

Being a qubit, the state of the system can also be represented by a vector $\vec{r} = \langle \sigma_{x} \rangle + i \langle \sigma_{y} \rangle$ in Bloch Sphere ($\langle \sigma_{z} \rangle = Tr(\sigma z)$) in which case the distance between $\rho_{1}$ and $\rho_{2}$ reduces to the geometric distance between the respective vectors $D(\rho_{1}, \rho_{2}) = \frac{1}{2}|\vec{r}_{2} - \vec{r}_{1}|$. This quantity can be easily calculated for the initial states $\rho_{1}(0) = |0\rangle \langle 0|$ and $\rho_{2}(0) = |1\rangle \langle 1|$ and for the rotations used in the experiment and gives $D(\rho_{1}1, \rho_{2}(0)) = \frac{1}{2}[3 + \cos(4\eta)]/2$ and $D(\rho_{1}(1), \rho_{2}(2)) = \frac{1}{2}[16 \cos \eta \sin \eta \cos \eta \sin \eta - \sin^{2} \eta]^{2} + \frac{1}{2}[4 \sin^{2} \eta \cos^{2} \eta (q + 1)]$ for the first and second collision, respectively. Expanding these results for $\eta \ll 1$ we get $\Delta D = -(1 - 4q)\eta^{2} + O(\eta^{4})$, making it clear why, for the range of $\eta$ used in the experiment, $\Delta D$ must be positive, i.e. the dynamics must be non-Markovian, for any $q < 1/4$ (highly anti-correlated environmental qubits). This result corresponds to the solid curves in Fig. 2.

Experimental details. The NMR experiment was performed encoding the system of interest and environment in a trifluoriodoethylene (CF$_3$I) sample diluted (1%) deuterated acetone (containing 97% of deuterium). The three nuclei spin-1/2 of the fluorine atoms were associated to system and environment qubits. The total system is described by the Hamiltonian

$$ \mathcal{H} = \sum_{n} \hbar(\omega_{on} - \omega_{n}) I_{z}^{n} + \sum_{k=m} \hbar 2\pi J_{km} I_{z}^{k} \otimes I_{z}^{m}, $$

where $\omega_{on}$ and $\omega_{n}$ are the natural resonance frequency and the frequency of the rotating reference frame of the $n$-th spin, $I_{z}^{n}$ is the $z$-component of the spin angular moment of nucleus $n$. The second term represents the interaction - being $J_{km}$ the exchange integral - between the three spins. Figure 3 shows the values of $(\omega_{on} - \omega_{n})$ and $J_{km}$.

Preparation of the initial state and unitary operations. The NMR implementation of the collisions between the system and particles of the environment are shown in Figs 4 and 5. The boxes with the symbols $R_{\alpha}(\theta)$ indicate that a rotation of the angle $\theta$ on that particular spin was performed around the $\alpha$ direction. The big boxes are free evolution of the system for the time $\tau$. The angles $\theta$ are related to the parameter $\eta$ of the collisional model.

Figure 3. Table of $(\omega_{on} - \omega_{n})/(2\pi)$ in the main diagonal elements and $J_{km}$ as the other elements, all values are given in Hz.
and they are of the order of 1°. For the steps used in the experiment the differences between them were approximately of 0.5°. The system we have used is homonuclear, which means that all of the spins are of the same species (fluorine in this case), and implementing single rotations were hard because they are close in frequency. Then, the length of the pulses for exciting only one spin (qubit) are long. This causes undesired evolutions of the quantum state due to the interactions between the spins, leading the whole system to evolve while the individual operations are applied. Furthermore, the rf pulses are also imperfect and they may affect others spins, besides the one which it is intended for. This means that at the end of the quantum algorithm lots of errors due to these undesirable evolutions and pulse imperfections will have been accumulated. In order to correct the state of the system the method described in ref. 33 was used.

For the experiment, the values of \( \eta \) were very small and this created a huge source of errors in the operations, since they had to be performed in tiny steps and, therefore, were not so different. In fact, the rotations angles were so small that the controlled operations, which simulate the collisions, were very hard to be performed properly. In order to achieve the necessary precision in the experiment the rotations of 0° had to be precisely adjusted for each value of \( \eta \) of the specific collision. For adjusting the angles values, two parameters may be varied, the amplitude and duration of the radio frequency pulses. However, varying these two parameters the sequence of operations needed for the corrections changes as well. Therefore, the correct pulse sequences were determined by combining the two analysis and testing in the spectrometer. A good precision could be achieved then and the small rotations, of the order of 0.30 degrees, could be implemented. For the state preparation the pulse sequence shown in Fig. 6 was performed, for more details see refs 36 and 37. The time \( \tau_2 \) of the free evolution here is related to the correlation parameter \( q \) as it is presented in Fig. 7. A the end of the circuit a magnetic field gradient in the \( z \)-direction is applied, which works as a transversal relaxation time killing the off-diagonal terms of the density matrix of system and environment. Finally, after each run, full state tomography was performed in the state of the system 34.

**Figure 4.** Quantum circuit representing the NMR pulse sequence for implementing the first collision. The boxes with the symbols \( R_{z}(\theta) \) indicate that a rotation of the angle \( \theta \) on that particular spin was performed around the \( z \) direction and \( R_{x}(\theta) = R_{z}(-\theta) \). The boxes with the symbol \( \tau_1 \) represent a free evolution of the system and environment for a period of time \( \tau_1 \). After the first collision, the resulting quantum state of system and environment is represented by \( \rho_1(1) \). \( \tau_1 \) is the time needed such that a rotation of \( \pi/2 \) occurs in the system and here it will be given by \( \tau_1 = 1/(4J_{r},_{\text{env}}) = 0.00358 \) s.

**Figure 5.** Quantum circuit representing the NMR pulse sequence for implementing the second collision. The input state is the resulting state from the first collision \( \rho_1(1) \). The boxes with the symbols \( R_{z}(\theta) \) indicate that a rotation of the angle \( \theta \) on that particular spin was performed around the \( z \) direction and \( R_{x}(\theta) = R_{z}(-\theta) \). The boxes with the symbol \( \tau_1 \) represent a free evolution of the system and environment for a period of time \( \tau_1 \). \( \tau_1 \) is the time needed such that a rotation of \( \pi/2 \) occurs in the system and here it will be given by \( \tau_1 = 1/(4J_{r},_{\text{env}}) = 0.00525 \) s.

**Figure 6.** Quantum circuit representing the NMR pulse sequence for the state preparation of the initial state of system and environment given by \( \rho(0) \otimes \rho_{\text{env}} \). The boxes with the symbols \( R_{z}(\theta) \) indicate that a rotation of the angle \( \theta \) on that particular spin was performed around the \( z \) direction and \( R_{x}(\theta) = R_{z}(-\theta) \). The boxes with the symbol \( \tau_2 \) represent a free evolution of the system and environment for a period of time \( \tau_2 \). The box with the symbol Grad stands for the magnetic field gradient that is applied in the \( z \)-direction.
Figure 7. Parameter $\tau_2$ has a relation to the correlation parameter $q$. The values used in the experiment are presented here.

Note that this experiment was done much faster (~0.03 s) than the typical relaxation times of the elements of the C$_2$F$_3$I molecule, which is of the order of 1 s.

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Author Contributions

N.K.B., C.H.M. and M.F.S. proposed the original idea. R.S.S., A.M.S., I.R. and I.S.O. designed the experimental setup. J.P.S.P. performed the experiment, N.K.B. and J.P.S.P. analyzed and interpreted the data. All authors contributed to the writing of the manuscript.

Additional Information

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