

LECTURE III QUANTIFYING AND APPLYING NON-MARKOVIANITY





Turku Centre for Quantum Physics

Non-Markovian Processes and Complex Systems Group



Lecture I

- I. General framework: Open quantum systems
- 2. Local in time master equations

Lecture II

3. Solving local in time master equations: Markovian and non-Markovian quantum jumps

Lecture III

- 4. Measures of non-Markovianity
- 5. Applications of non-Markovianity
- Q1: How much memory?
- Q2: Anything useful can be done?









IV Measures of non-Markovianity

- Background
- Information flow/trace distance based measure
- Other measures of non-Markovianity





Any realistic quantum system coupled to its environment



The open system exchanges energy and information with its environment

Thermalization

Simple example: 2-level atom on excited state coupled to vacuum electromagnetic field (T=0) - spontaneous emission of a photon



Consequence: decoherence

- Quantum systems lose their quantum properties (e.g. quantum superpositions lost)
 Transition from quantum to classical world
- From Schrödinger equation and wave functions to master equations and density matrices
- Eindblad master equation (1975, Markovian, semigroup)

$$\frac{d}{dt}\rho_{S}(t) = -i\left[H,\rho_{S}(t)\right] + \mathcal{D}(\rho_{S}(t))$$

$$\mathcal{D}(\rho_{S}) \equiv \sum_{\substack{k \ \uparrow \ decay \, rate}} \gamma_{k} \left(A_{k}\rho_{S}A_{k}^{\dagger} - \frac{1}{2}A_{k}^{\dagger}A_{k}\rho_{S} - \frac{1}{2}\rho_{S}A_{k}^{\dagger}A_{k}\right)$$



Background

Non-Markovian dynamics: quantum memory effects

- Memoryless Markovian dynamics (e.g. exponential decay) is an approximation of more general theory of non-Markovian open system dynamics
- Classical vs quantum stochastic processes
- Eg. atom spontaneously emitting a photon: free space vs photonic band gap materials
- Is the loss of quantum properties a one-way process, role of memory effects?
- Can we control and exploit memory effects?



Background

Early experimental work on decoherence (Markovian):

Haroche and co-workers, PRL 1996 (observing):
Light field in a cavity, preparation of a

- Schrödinger's cat-like state
- Measuring decoherence by atoms



 Wineland and co-workers, Nature 2000 (engineered):
 Trapped ions, superpositions of coherent states and Fock-states

 Engineering the decoherence by applying noise to trap electrodes



Often decoherence considered as an obstacle, for example to quantum information... however, can be also exploited...



Background

No universally agreed definition of non-Markovianity for quantum processes

Earlier non-Markovianity associated, e.g., to:

- Anything not following Lindblad-equation
- Non-exponential decay
- Memory-kernel equations
- Negative decay rates...

$$\frac{d\rho(t)}{dt} = \int_0^t k(t') \mathcal{L}\rho(t-t') dt'$$

Is it possible to define and quantify non-Markovianity
Independently of the used mathematical formalism
Intuitively clear interpretation
Physically motivated approach (instead of formal mathematical approach)
What does memory mean in quantum dynamics?

Our starting point: information flow and its direction between the system and environment...



Information flow: Markovian case

Cartoon of the information flow: Markovian case (no memory).

"Small" system. "Large" environment.



System

System-environment interaction → System reaches steady state

Any initial state of the system leads to same steady state → The system loses information

on its initial state.

Markovian system: Information flow from the system to the environment.



Distance measures for quantum states

 ρ_2

Q: What is a suitable distance measure *P*₁ to quantify non-Markovianity?

Criteria for arbitrary distance measure: $L(\rho_1, \rho_2)$

- Symmetric? $L(\rho_1, \rho_2) = L(\rho_2, \rho_1)$
- Contractivity? $L(\Phi_t(\rho_1), \Phi_t(\rho_2)) \leq L(\rho_1, \rho_2)$

• Triangle equality? $L(\rho_1, \rho_2) \leq L(\rho_1, \rho_3) + L(\rho_3, \rho_2) \rho_3$ $\rho_1 \bullet \rho_2$



Distance measures for quantum states

Choices (for example):

 $L(\rho_1, \rho_2) = L(\rho_2, \rho_1)$ $L \bullet \Phi \bullet \rho_1 \bullet, \Phi \bullet \rho_2 \bullet \leq L \bullet \rho_1, \rho_2 \bullet$

Relative entropy: $S(\rho_1 || \rho_2) = \text{Tr}[\rho_1(\log \rho_1 - \log \rho_2)]$ $L(\rho_1, \rho_2) \leq L(\rho_1, \rho_3) + L(\rho_3, \rho_2)$

Not symmetric

Ooes not satisfy triangular inequality

Singularities

Fidelity:

$$F(\rho_1, \rho_2) = \text{Tr}\sqrt{\rho_1^{1/2}\rho_2\rho_1^{1/2}}$$

• Does not satisfy triangular inequality Hilbert-Schmidt distance: $HS(\rho_1, \rho_2) = \sqrt{Tr[(\rho_1 - \rho_2)^2]}$

Generally not contractive

Trace distance:

$$D(\rho_1, \rho_2) = \frac{1}{2} \text{Tr} |\rho_1 - \rho_2|$$

• OK



Trace distance

Distance measure for two states ρ_1 and ρ_2 : Trace distance D:

$$D(\rho_1, \rho_2) = \frac{1}{2} \text{Tr} |\rho_1 - \rho_2| \qquad 0 \le D \le 1$$

- ${\small \bigcirc}$ For identical states D=0 , for orthogonal states D=1
- Physical interpretation: measure of distinguishability The max probability to distinguish the two states is equal to $\frac{1}{2}$
- In terms of information:
 - The larger D, the higher the probability to distinguish, more information which state we have
- Invariant under unitary transformations
- ullet Contractive for all CPT-maps Φ

 $D(\Phi\rho_1, \Phi\rho_2) \le D(\rho_1, \rho_2)$





⁽Nielsen, Chuang)



Information flow and distinguishability of states (2)

Notation: $\sigma(t) = dD(t)/dt$ the change of the trace distance

In terms of the information flow:

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σ(t) < 0
Decreasing trace distance:
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More and more difficult to distinguish the different states. →

The information flows from the system to the environment.

 $\sigma(t) > 0$

 \rightarrow

Increasing trace distance:

More and more easy to distinguish the different states.



The information flows from the environment to the system.

The aim: quantify memory by calculating the reverse flow of information from the environment to the system.



Non-Markovianity: whenever period of increase of trace distance: backflow of information from the environment to the open system

Allows to define a measure for non-Markovianity:

$$\mathcal{N}(\Phi) = \max_{\rho_{1,2}(0)} \int_{\sigma>0} dt \ \sigma(t, \rho_{1,2}(0)).$$

• Gives the total increase of the trace distance during the time evolution

• The total amount of information that has flown from the environment to the system during the time evolution.

General definition, independent of the used formalism to solve the open system dynamics.



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Example: 2-level atom

Two-level atom with Lorentzian spectral density $\dot{\rho}(t) = \frac{1}{i}\lambda(t)[\sigma_+\sigma_-,\rho(t)] + \Delta(t)\sigma_-\rho(t)\sigma_+ - \frac{1}{2}\Delta(t)\{\rho(t),\sigma_+\sigma_-\}.$ DECAY RATE [r] Decay rate oscillates (a) having periods of negativity 0 (b) • Excited state population ground 0.8 POPULATIONS NMQJ: ρ_{aa} oscillates $\text{NMQJ:}\,\rho_{\text{bb}}$ 0.6 $\rho_{aa}(t) = e^{-D_1(t)}\rho_{aa}(0)$ analytical 0.4 excited 0.2 0 NMQJ: Ipah (C) **Decoherence** -0.4 analytical COHERENCES re-coherence cycles 0.3 $\rho_{ab}(t) = e^{-D_1(t)/2} \rho_{ab}(0)$ 0.2 0.1 $D_i(t) = \int ds \Delta_i(s)$ 0 10 2 8 9 3 5 7 TIME $[1/\Gamma]$



Example: 2-level atom





Other measures



 $\Phi(t,0) = \Phi(t,s)\Phi(s,0)$

Our view: PRL 2009:

Recycling of info between S and E, quantify backflow

Xiao-Ming Lu, Xiaoguang Wang, C. P. Sun: PRA 2010

Quantum Fisher Information based measure

Bylicka, Chruscinski, Maniscalco: arXiv1301:2585

Definition based on channel capacity

time time time time

...also others exist



Comparing to Wolf, Eisert, Cubitt, Cirac: PRL 2008

- Markovianity vs non-Markovianity from a snapshot of the evolution.
- Non-zero value when the state can have reached by non-Markovian evolution, zero when both Markovian and non-Markovian possible

Pure dephasing and rephasing evolution:

$\rho_{++}(t) = \rho_{++}(0)$	$ \rho_{}(t) = \rho_{}(0) $
$\rho_{-+}(t) = g(t)\rho_{-+}(0)$	$\rho_{+-}(t) = g(t)\rho_{+-}(0)$
$g(t) = \frac{1}{2} \left(1 + \cos^2(\omega t) \right)$ Snapshot measure = 0:	

at all points of time, the map can be written as an element of Markovian semigroup \longrightarrow snapshot measure = 0

$$\Phi(t_0, 0) = \exp(\mathcal{L}) \qquad \mathcal{L}\rho = \frac{1}{2}(\sigma_3\rho\sigma_3 - \rho) \ \Gamma = -\ln g(t_0)$$

Our trace distance measure:

Laine, Piilo, Breuer: Physical Review A 81, 062115 2010

 $\mathcal{N}=\infty~$ (decoherence-recoherence-decoherence cycles till time ∞)



\int	Semigroup	, , , , , , , , , , , , , , , , , , ,	Information backflow (trace distance meas.)
Φ	$\Phi(t_1 + t_2, 0) = \Phi(t_1, 0)\Phi(t_2, 0)$	$\Phi(t,0) = \Phi(t,s)\Phi(s,0)$	$\mathcal{N}(\Phi)$

- Semigroup: constant decay rates
- Semigroup broken when decay rates become time dependent
- Divisibility broken when at least one of the decay rates becomes negative
- Trace distance measure: backflow of information
- Output Can break divisibility but no backflow of information



General classification based on the information flow





V Applications of non-Markovianity

- Controlling the memory effects
- Nonlocal memory effects
- Non-Markovian probes



$$\frac{d\rho(t)}{dt} = -i\frac{\epsilon(t)}{2}\left[\sigma_z,\rho\right] + \frac{\gamma(t)}{2}\left(\sigma_z\rho\sigma_z-\rho\right)$$

- Time evolution and the map: two state system $|H\rangle$, $|V\rangle$ $\rho_{\rm H,H}(t) = \rho_{\rm H,H}(0), \quad \rho_{\rm V,V}(t) = \rho_{\rm V,V}(0),$ $\rho_{\rm H,V}(t) = \kappa^*(t)\rho_{\rm H,V}(0), \quad \rho_{\rm V,H}(t) = \kappa(t)\rho_{\rm V,H}(0)$
- ${\ensuremath{\, \bullet }}$ Connection between the decoherence function $\kappa(t)$ and the rates in the master equation

$$\kappa(t) = \exp\left(-\int_{0}^{t'} \gamma(t') + i\epsilon(t')dt'\right)$$
$$\epsilon(t) = -\Im\left[\frac{\dot{\kappa}(t)}{\kappa(t)}\right], \qquad \gamma(t) = -\Re\left[\frac{\dot{\kappa}(t)}{\kappa(t)}\right]$$



$$\frac{d\rho(t)}{dt} = -i\frac{\epsilon(t)}{2}\left[\sigma_z,\rho\right] + \frac{\gamma(t)}{2}\left(\sigma_z\rho\sigma_z-\rho\right)$$

Single photons in dephasing reservoir

ullet The system: polarization states of the photon: $|H
angle\;|V
angle$

• Environment: frequency degrees of freedom $\{|\omega\rangle\}_{\omega\in\mathbb{R}}$ $|\omega_i\rangle$

• Dephasing by quartz plates (birefringent material)



• Initial total system states $|\psi_{1,2}(0)\rangle = |\varphi_{1,2}\rangle \otimes |\chi\rangle$

Initial open system states (polarization)

 $|\varphi_{1,2}\rangle = \frac{1}{\sqrt{2}} \left(|H\rangle \pm |V\rangle\right)$

• Initial environmental states (frequency distribution) $|\chi\rangle = \int d\omega f(\omega) |\omega\rangle$

Modified by the FP cavity

• Total system evolution in the quartz plate $U(t)|\lambda\rangle \otimes |\omega\rangle = e^{in_{\lambda}\omega t}|\lambda\rangle \otimes |\omega\rangle$

- H and V acquire different phase due to the different refraction indices (birefringent quartz plate)







The tilting of the FP cavity modifies the frequency spectrum.



Markovian - non-Markovian transition experiment



Tilting of the cavity modifies the initial environmental state



Non-Markovianity

• The optimal trace distance $D(\rho_1(t), \rho_2(t)) = |\kappa(t)|$.

$$D(\rho_1(t), \rho_2(t)) = \sqrt{a^2 + |\kappa(t)b|^2}$$

$$b = \rho_1^{12}(0) - \rho_2^{12}(0)$$

$$a = \rho_1^{11}(0) - \rho_2^{11}(0)$$

• Two Gaussian peaks, relative weights

$$A_1 = \frac{1}{1+A}, \ A_2 = \frac{A}{1+A}$$

Decoherence function

$$|\kappa(t)| = \frac{e^{-\frac{1}{2}\sigma^2(\Delta nt)^2}}{1+A}\sqrt{1+A^2+2A\cos(\Delta\omega\cdot\Delta nt)}$$

 $\label{eq:one-peak} \bullet A = 0 $$ |\kappa(t)|$ monotonically decreasing $$$



Trace distance dynamics -

transition from monotonic to non-monotonic behavior





Markovian - non-Markovian transition



Experimental control on the amount and direction of the information flow between the system and the environment.



Nonlocal memory effects and a non-Markovian quantum probe

Laine, Breuer, Piilo, C.-F. Li, Guo, Phys. Rev. Lett. 108, 210402 (2012)

Liu, Cao, Huang, C.-F. Li, Guo, Laine, Breuer, Piilo "Photonic realization of nonlocal memory effects and non-Markovian quantum probes", arXiv:1208.1358



2-qubits interacting with their local environments





2-qubits interacting with their local environments



What happens when the environments are initially correlated?

$$\rho_S^{12}(t) = \Phi_{12}(t)(\rho_S^{12}(0)) = \operatorname{tr}_E \left[(U_1(t) \otimes U_2(t)) \rho_S^{12}(0) \otimes \rho_E^{12}(0) (U_1^{\dagger}(t) \otimes U_2^{\dagger}(t)) \right]$$



• Considered open system: two photons

- polarizations H,V
- initial pure state

 $\ket{\psi_{12}} = a \ket{HH} + b \ket{HV} + c \ket{VH} + d \ket{VV}$

Environment: frequency degrees of freedom

$$\chi
angle = \int d\omega_1 d\omega_2 \, g(\omega_1, \omega_2) \ket{\omega_1, \omega_2}$$

- joint probability distribution $P(\omega_1, \omega_2) = |g(\omega_1, \omega_2)|^2$
- Initial system-environment product state $|\Psi(0)\rangle = |\psi_{12}\rangle \otimes \int d\omega_1 d\omega_2 g(\omega_1, \omega_2) |\omega_1, \omega_2\rangle$



General dephasing for 2 qubits (photons, polarization)



• Local states for qubit I and 2 [trace out qubit 2 (I)]

$$\rho_{1}(t) = \begin{pmatrix} |a|^{2} + |b|^{2} & (ac^{*} + bd^{*})\kappa_{1}(t) \\ (ca^{*} + db^{*})\kappa_{1}^{*}(t) & |c|^{2} + |d|^{2} \end{pmatrix}$$
$$\rho_{2}(t) = \begin{pmatrix} |a|^{2} + |c|^{2} & (ab^{*} + cd^{*})\kappa_{2}(t) \\ (ba^{*} + dc^{*})\kappa_{2}^{*}(t) & |b|^{2} + |d|^{2} \end{pmatrix}$$

- Global 2-qubit decoherence functions: $\kappa_{12}(t)$, $\Lambda_{12}(t)$
- **Local** decoherence functions: $(\kappa_1(t), \kappa_2(t))$



• ...however, the open system map is a product of local maps $\Phi_{12}(t) = \Phi_1(t) \otimes \Phi_2(t)$ if and only if $\kappa_{12}(t) = \kappa_1(t)\kappa_2(t)$ $\Lambda_{12}(t) = \kappa_1(t)\kappa_2^*(t)$

• Local interaction Hamiltonian for photon i $H_{i} = -\int d\omega_{i} \,\omega_{i} \left[n_{V} |V\rangle \langle V| + n_{H} |H\rangle \langle H| \right] \otimes |\omega_{i}\rangle \langle \omega_{i}|$

 $H_{\rm int}(t) = \chi_1(t)H_1 + \chi_2(t)H_2$

Oecoherence functions $\kappa_{1}(t) = \int d\omega_{1}d\omega_{2}P(\omega_{1},\omega_{2})e^{-i\Delta n\omega_{1}t_{1}}$ $\kappa_{2}(t) = \int d\omega_{1}d\omega_{2}P(\omega_{1},\omega_{2})e^{-i\Delta n\omega_{2}t_{2}}$ $\kappa_{12}(t) = \int d\omega_{1}d\omega_{2}P(\omega_{1},\omega_{2})e^{-i\Delta n(\omega_{1}t_{1}+\omega_{2}t_{2})}$ $\Lambda_{12}(t) = \int d\omega_{1}d\omega_{2}P(\omega_{1},\omega_{2})e^{-i\Delta n(\omega_{1}t_{1}-\omega_{2}t_{2})}$

Frequency correlations give nonlocal map and non-Markovian dynamics



- Consider
 - -Gaussian two-photon frequency distribution
 - -Variance C
 - Correlation coefficient K
 - initial open system states

 $|\psi_{12}^{\pm}
angle~=~\left(|HH
angle\pm|VV
angle
ight)/\sqrt{2}$



$$D(t) = \exp\left[-\frac{1}{2}\Delta n^2 C\left(t_1^2 + t_2^2 - 2|K|t_1t_2\right)\right]$$



• ...and the non-Markovianity measure is $\mathcal{N} = e^{-\frac{1}{2}C_{11}(\Delta nT)^2} \left[e^{\frac{1}{2}C_{11}(\Delta nT)^2 K^2} - 1 \right]$

Direct connection between the amount of non-Markovianity of the open system and the correlations between the local environments

Laine, Breuer, Piilo, C.-F. Li, Guo: Phys. Rev. Lett. 108, 210402 (2012)





Nonlocal memory effects

Correlations, anticorrelations, and non-Markovianity



Max non-Markovianity for

Correlated frequency, anticorrelated polarization
 Anticorrelated frequency, correlated polarization

Anticorrelated frequency, correlated polariz

Laine, Breuer, Piilo, C.-F. Li, Guo, Phys. Rev. Lett. 108, 210402 (2012)





Local Markovian dynamics - global non-Markovian dynamics



Laine, Breuer, Piilo, C.-F. Li, Guo, Phys. Rev. Lett. 108, 210402 (2012)





Experimental setup



Liu, Cao, Huang, C.-F. Li, Guo, Laine, Breuer, Piilo

"Photonic realization of nonlocal memory effects and non-Markovian quantum probes", arXiv:1208.1358



Nonlocal memory effects experiment



...this controls the frequency anticorrelations of the down converted photon pair (energy conserved, $w_0 = w_1 + w_2$)

Liu, Cao, Huang, C.-F. Li, Guo, Laine, Breuer, Piilo "Photonic realization of nonlocal memory effects and non-Markovian quantum probes", arXiv:1208.1358



Nonlocal memory effects experiment

Amount of frequency correlations K and the degree of non Markovianity N



- Correlations in frequency induce non-Markovian polarization dynamics
- By measuring polarization we probe the frequency correlations

Liu, Cao, Huang, C.-F. Li, Guo, Laine, Breuer, Piilo

"Photonic realization of nonlocal memory effects and non-Markovian quantum probes", arXiv:1208.1358



Applications of the information flow based measure

Energy transfer in light-harvesting complexes

P. Rebentrost and A. Aspuru-Guzik

J. Chem. Phys. 134, 101103 (2011).

Spin environments

T. J. G. Apollaro, C. Di Franco, F. Plastina, and M. Paternostro Phys. Rev. A 83, 032103 (2011)

Cold atomic gases

P. Haikka, S. McEndoo, G. de Chiara, M. Palma, and S. Maniscalco PRA 2011

Chaotic quantum systems

M. Znidaric, C. Pineda, I. Garcia-Mata PRL 2011

Ising model in transverse field

Non-Markovianty pinpoints the quantum phase transition from paramagnetic to ferromagnetic ground state Haikka, Maniscalco PRA 2012



MARKOVIAN: Constant info flow rate from the system to the environment.

TIME-DEPENDENT MARKOVIAN:

Time-dependent info flow rate from the system to the environment.



NON-MARKOVIAN: Recycling of info from the environment to the system.





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