











Reduced dynamics

$$\hat{\rho}_{S}(t) := \hat{\mathcal{T}}(t, t_{0})\hat{\rho}_{S}(t_{0}) = \operatorname{Tr}_{E}\left[\hat{\rho}_{SE}(t)\right]$$

 $\hat{\mathcal{T}}(t,t_0)$ is a linear map which transforms the input state $\hat{\rho}_s(t_0)$ onto the output state $\hat{\rho}_s(t)$.

The question: how we can determine (reconstruct) the master equation which governs the time evolution of the reduced density operator $\hat{\rho}_s(t)$

$$\frac{d}{dt}\hat{
ho}(t) = \hat{\mathcal{L}}(t,t_0)\hat{
ho}(t)$$

Liouvillian superoperator

This master equation can be written in the *convolutionless* form

$$\frac{d}{dt}\hat{\rho}(t) = \hat{\mathcal{L}}(t, t_0)\hat{\rho}(t).$$

due to the fact that in the *finite-dimensional* Hilbert spaces matrix elements of density operators are analytic functions. Consequently, $\hat{\mathcal{T}}(t, t_0)$ are non-singular operators (except may be for a set of *isolated* values of t) in which case the inverse operators $\hat{\mathcal{T}}(t, t_0)^{-1}$ exist and the Liouvillian superoperator can be expressed as

$$\hat{\mathcal{L}}(t,t_0) := \left[\frac{d}{dt}\hat{\mathcal{T}}(t,t_0)\right]\hat{\mathcal{T}}^{-1}(t,t_0)$$

 $\hat{\mathcal{T}}(t,t_0)$ is uniquelly specified by $\hat{H}_{\scriptscriptstyle SE}$ and by the initial state $\hat{\rho}_{\scriptscriptstyle E}(t_0)$ of the environment.

Reconstruction of the map $\hat{\mathcal{T}}(t, t_0)$

Let us assume that the system S has been initially prepared in a pure state

$$|\Psi(t_0)\rangle = \sum_{i_1=0}^N c_{i_1}|i_1\rangle$$

where $|i_1\rangle$ are basis vectors in the (N+1)-dimensional Hilbert space $\mathcal{H}_{\scriptscriptstyle S}$ of the system under consideration.

The environment is initially prepared in a state $\hat{\rho}_E(t_0) = \sum_{\alpha_1 \alpha_2} d_{\alpha_1 \alpha_2} |\alpha_1\rangle_E \langle \alpha_2 |$, where $|\alpha_i\rangle_E$ are basis vectors in the Hilbert space \mathcal{H}_E of the environment.

Reconstruction of the map $\hat{\mathcal{T}}(t, t_0)$

The physical process $\hat{\mathcal{T}}(t_k, t_0)$ is determined by a transformation acting on basis vectors of the system and the environment

$$|i_1\rangle_S |\alpha_1\rangle_E \xrightarrow{\hat{\mathcal{T}}(t_k)} \sum_{j_1=0}^N \sum_{\beta_1} E_{(i_1j_1)(\alpha_1\beta_1)}(t_k) |j_1\rangle_S |\beta_1\rangle_E.$$

The output density operator $\hat{\rho}(t_k)$

 $\hat{\rho}(t_k) = \sum_{i_1, i_2=0}^{N} c_{i_1}(c_{i_2})^* \hat{R}_{(i_1, i_2)}(t_k),$

where $(N+1)^2$ operators $\hat{R}_{(i_1,i_2)}(t_k)$

$$\hat{R}_{(i_1,i_2)}(t_k) = \sum_{j_1,j_2=0}^N D_{(i_1,i_2)(j_1,j_2)}(t_k) |j_1\rangle \langle j_2|,$$

with

$$D_{(i_1,i_2)(j_1,j_2)}(t_k) = \sum_{\alpha_1,\alpha_2,\gamma} d_{\alpha_1\alpha_2} \times E_{(i_1j_1)(\alpha_1\gamma)}(t_k) E^*_{(i_2j_2)(\alpha_2\gamma)}(t_k).$$

Properties of $\hat{R}_{(i_1,i_2)}(t_k)$

The process $\hat{\mathcal{T}}(t_k)$ for a given time t_k is *completely* determined by $(N+1)^2$ operators $\hat{R}_{(i_1,i_2)}(t_k)$, which in turn are specified by the $(N+1)^2 \times (N+1)^2$ matrix elements $D_{(i_1,i_2)(j_1,j_2)}(t_k)$.

 $\hat{R}_{(i_1,i_2)}(t_k)$ have the properties

$$\operatorname{Tr}\hat{R}_{(i_1,i_2)}(t_k) = \delta_{i_1,i_2}; \qquad (\hat{R}_{(i_1,i_2)}(t_k))^{\dagger} = \hat{R}_{(i_2,i_1)}(t_k),$$

or, equivalently,

$$\sum_{j=0}^{N} D_{(i_1,i_2)(j,j)}(t_k) = \delta_{i_1,i_2}; \qquad D^*_{(i_1,i_2)(j_1,j_2)}(t_k) = D_{(i_2,i_1)(j_2,j_1)}(t_k)$$

Determination of $\hat{R}_{(i_1,i_2)}(t_k)$

In order to specify the $(N + 1)^2$ operators $\hat{R}_{(i_1,i_2)}(t_k)$ one has to consider $(N+1)^2$ specific initial conditions $|\Psi^{(k_1,k_2)}\rangle_{in} = \sum_{i_1=0}^N c_{i_1}^{(k_1,k_2)} |i_1\rangle$ where $k_1, k_2 = 0, 1, ..., N$ and to measure the corresponding $(N + 1)^2$ output density operators $\hat{\rho}^{(k_1,k_2)}(t_k)$ which can be expressed as

$$\hat{\rho}^{(k_1,k_2)}(t_k) = \sum_{i_1,i_2=0}^N M_{(k_1,k_2)(i_1,i_2)} \hat{R}_{(i_1,i_2)}(t_k),$$

where

$$M_{(k_1,k_2)(i_1,i_2)} = c_{i_1}^{(k_1,k_2)} (c_{i_2}^{(k_1,k_2)})^*$$

If the $(N + 1)^2$ initial conditions $|\Psi^{(k_1,k_2)}\rangle_{in}$ are chosen so, that the matrix $M_{(k_1,k_2)(i_1,i_2)}$ is invertible, then the set of equations can be solved with respect of the operators $\hat{R}_{(i_1,i_2)}(t_k)$.

Determination of $\hat{R}_{(i_1,i_2)}(t_k)$

To make the reconstruction possible the matrix M has to be invertible. Obviously, there are many choices of such matrix. For instance M is given by complex amplitudes $c_i^{(k_1,k_2)}$ specified as

$$c_i^{(k_1,k_2)} = \begin{cases} (\delta_{i,k_1} + \delta_{i,k_2})/\sqrt{2} & \text{if } k_1 > k_2 \\ \delta_{i,k_1} & \text{if } k_1 = k_2 \\ (\delta_{i,k_1} + i\delta_{i,k_2})/\sqrt{2} & \text{if } k_1 < k_2 \end{cases}$$
(1)

The reconstruction process described above gives us a set of operators $\hat{R}_{(i_1,i_2)}(t_k)$ which describe the transition of the system from the state $\hat{\rho}(t_0)$ to the state $\hat{\rho}(t_k)$ at a given time t_k . In principle, one can perform a whole sequence of such reconstructions at different times t_1, t_2, \dots, t_K so that the *reduced dynamics* of the studied system can be reconstructed from the measured data.

Reconstruction of $\hat{\mathcal{L}}(t)$

Now our task is to determine (reconstruct) from a set of measurements of the output states $\hat{\rho}^{(k_1,k_2)}(t)$ for given input states $\hat{\rho}^{(k_1,k_2)}(t_0)$, the form of the Liouvillian superoperator $\hat{\mathcal{L}}(t)$.

The operators $\hat{R}_{(i_1,i_2)}(t)$ are also governed by the same master equation

$$\frac{d}{dt}\hat{R}_{(i_1,i_2)}(t) = \hat{\mathcal{L}}(t)\hat{R}_{(i_1,i_2)}(t),$$

Alternatively, for matrix elements $D_{(i_1,i_2)(k_1,k_2)}(t)$

$$\frac{d}{dt}D_{(i_1,i_2)(k_1,k_2)}(t)\sum_{j_1,j_2=0}^N D_{(i_1,i_2)(j_1,j_2)}(t)G_{(j_1,j_2)(k_1,k_2)}(t),$$

Here the matrix $G_{(j_1,j_2)(k_1,k_2)}(t)$ is defined as

$$G_{(j_1,j_2)(k_1,k_2)}(t) = \langle k_1 | \left(\hat{\mathcal{L}}(t) | j_1 \rangle \langle j_2 | \right) | k_2 \rangle,$$

and it uniquely determines the Liouvillian superoperator $\hat{\mathcal{L}}(t)$.

Reconstruction of $\hat{\mathcal{L}}(t)$

We already know how to reconstruct matrices D from the measured data for arbitrary time t (from these data we can also evaluate the corresponding time derivatives). Providing the matrix $D_{(i_1,i_2)(j_1,j_2)}(t)$ is not singular its inverse $\tilde{D}_{(j_1,j_2)(i_1,i_2)}(t)$ can be found and then the reconstructed matrix $G_{(j_1,j_2)(k_1,k_2)}(t)$ is given by a simple expression

$$G_{(j_1,j_2)(k_1,k_2)}(t) = \sum_{i_1,i_2=0}^{N} \tilde{D}_{(j_1,j_2)(i_1,i_2)}(t) \frac{d}{dt} D_{(i_1,i_2)(k_1,k_2)}(t)$$

from which the superoperator $\hat{\mathcal{L}}(t)$ at time t can be determined.

The end

Let us consider a two-level system (a two-level atom, a spin-1/2, or a qubit) with a two-dimensional Hilbert space \mathcal{H}_s spanned by two vectors $|1\rangle$ and $|0\rangle$. In order to specify the Liouvillian superoperator $\hat{\mathcal{L}}(t)$ for the two-level atom we have to know the time evolution of four initial states. Let us assume that from the measured data it is found that these states evolve as

$$\begin{split} \hat{\rho}^{(0,0)}(t) &= \begin{pmatrix} 0 & 0 \\ 0 & 1 \end{pmatrix}; \quad \hat{\rho}^{(1,1)}(t) = \begin{pmatrix} e^{-\Gamma t} & 0 \\ 0 & 1 - e^{-\Gamma t} \end{pmatrix}, \\ \hat{\rho}^{(0,1)}(t) &= \frac{1}{2} \begin{pmatrix} e^{-\Gamma t} & ie^{-\Gamma t/2} \\ -ie^{-\Gamma t/2} & 2 - e^{-\Gamma t} \end{pmatrix}; \\ \hat{\rho}^{(1,0)}(t) &= \frac{1}{2} \begin{pmatrix} e^{-\Gamma t} & e^{-\Gamma t/2} \\ e^{-\Gamma t/2} & 2 - e^{-\Gamma t} \end{pmatrix}. \end{split}$$

Example 1: Decay of 2-leve atom

Now we can apply our reconstruction scheme and we find for the matrix $G_{(j_1,j_2)(k_1,k_2)}(t)$ the expression

$$G_{(j_1,j_2)(k_1,k_2)}(t) = \begin{pmatrix} -\Gamma & 0 & 0 & \Gamma \\ 0 & -\Gamma/2 & 0 & 0 \\ 0 & 0 & -\Gamma/2 & 0 \\ 0 & 0 & 0 & 0 \end{pmatrix}.$$

This matrix corresponds to the Liouvillian which defines the master equation

$$\frac{d}{dt}\hat{\rho} = \hat{\mathcal{L}}\hat{\rho} = \frac{\Gamma}{2} \left[2\hat{\sigma}_{-}\hat{\rho}\hat{\sigma}_{+} - \hat{\sigma}_{+}\hat{\sigma}_{-}\hat{\rho} - \hat{\rho}\hat{\sigma}_{+}\hat{\sigma}_{-} \right],$$

describing the decay of a two-level atom into a zero-temperature reservoir. The Liouvillian is time independent which reflects the fact that the state of the reservoir does not change in time under the influence of the system.

Example 2: Decay of 2-leve atom

Let us consider a single two-level atom coupled to K modes of the electromagnetic field in a one-dimensional cavity of the length L. The spectrum of modes is discrete with frequencies $\omega_k = k\pi c/L$. The corresponding total Hamiltonian in the dipole and rotating-wave approximations reads

$$\hat{H} = \omega_A \hat{\sigma}_z + \sum_{k=1}^K \omega_k \hat{a}_k^{\dagger} \hat{a}_k + \sum_{k=1}^K \lambda_k (\hat{\sigma}_+ \hat{a}_k + \hat{\sigma}_- \hat{a}_k^{\dagger}).$$

The field is assumed to be initially in the *vacuum* state. By applying our algorithm we find the master equation for the atom

$$\frac{d}{dt}\hat{\rho} = \hat{\mathcal{L}}\hat{\rho} = \frac{\gamma(t)}{2} \left[2\hat{\sigma}_{-}\hat{\rho}\hat{\sigma}_{+} - \hat{\sigma}_{+}\hat{\sigma}_{-}\hat{\rho} - \hat{\rho}\hat{\sigma}_{+}\hat{\sigma}_{-}\right],$$

except the "decay" rate $\Gamma \to \gamma(t)$ is now explicitly time dependent. It can be expressed in terms of the "measured" probability $P(t) = \langle 1 | \hat{\rho}_A(t) | 1 \rangle$ that the upper atomic level is excited:

$$\gamma(t) = -\left(\frac{dP(t)}{dt}\right)P(t)^{-1}.$$



The time evolution of the decay rate $\gamma(t)$ (thin line) and the population of the excited atomic level P(t) (thick line). We assume the atom to be in the center of the 1-D cavity, so it is coupled only to the odd modes (i.e. $\lambda_{2k} = 0$). We assume $L = 2\pi$ and c = 1 so that $\omega_{2k+1} = k+1/2$, and $\lambda_{2k+1} = \lambda = 0.3$. The effective density of modes which interact with the atom is $d_{eff}(\omega) = L/2c\pi = 1$. Therefore the decay rate $\Gamma = 2\pi\lambda^2 d_{eff}(\omega) \simeq 0.564$. We consider K = 400 modes of the field initially in the vacuum state and the atom (with $\omega_A = 101$) in its upper state $|1\rangle$.

















