

# Optimal Speed of Quantum Control in Open Quantum Systems

**Sarfraj Fency** (PhD student)

Collaborator: Dr. Riddhi Chatterjee (IISc Bangalore)

Supervisor: Prof. Rangeet Bhattacharyya



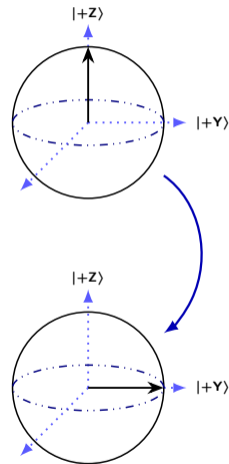
Indian Institute of Science Education and  
Research Kolkata, India

7th February, 2025



# The motivation

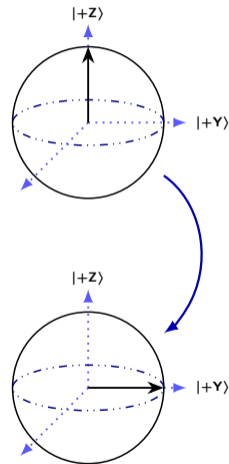
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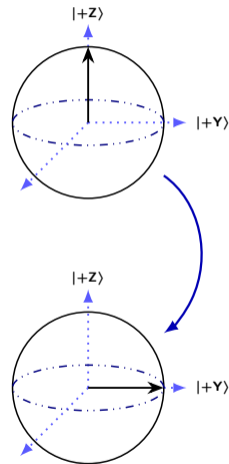


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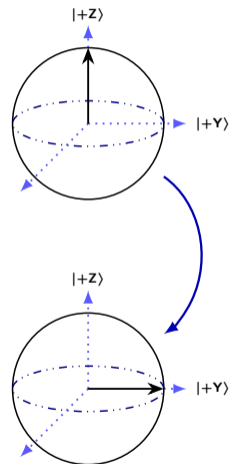
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- **Practical challenge**: Environmental dissipation and noise complicate evolution in open quantum systems.
- Strong external drives, necessary for fast operations, introduce **drive-induced dissipation (DID)**.

*Chakrabarti et al., EPL 121, 57002 (2018)*

*Chakrabarti et al., Phys. Rev. A 97 6 (2018)*

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- Can we achieve **fast** and **high-fidelity** quantum operations in open quantum systems while accounting for these dissipation?
- Can we design **optimal control strategies** to mitigate these challenges?



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- The thermal bath is modeled as a two-level system:

$$\mathcal{H}_{SL} = \omega_{SL} (\sigma_+ L_- + \sigma_- L_+)$$

# Quantum master Equation

- The standard quantum master equations  $\implies$  environmental dissipation
  - Bloch Equation, Phys. Rev. 89, 728 (1953)
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  - Several theoretical papers have shown the existence and implication of DID
    - Chanda *et al.*, PRA 104, 022436 (2021)
    - Saha *et al.*, PRA 107, 022206 (2023)
    - Chatterjee *et al.*, The European Physical Journal D 78, 44 (2024)
    - Das *et al.*, PRA 110, 062211 (2024)
- and more....

# Fluctuation Regulated Quantum Master Equation

$$\dot{\rho}_s = -i \text{Tr}_L[H_{\text{eff}}, \rho]^{\text{sec}} - \int_0^\infty d\tau e^{-\frac{\tau}{\tau_c}} \text{Tr}_L[H_{\text{eff}}(t), [H_{\text{eff}}(t - \tau), \rho]]^{\text{sec}}$$

Chakrabarti *et al.*, Phys. Rev. A 97 6 (2018)

$$H_{\text{eff}} = H_{\text{dr}} + H_{\text{SL}},$$

$$\tau_c = \frac{2}{\kappa^2},$$

$\rho$  = total density matrix,

$\rho_s$  = system's density matrix

$\text{Tr}_L$  = partial trace taken over the bath degrees of freedom

“sec” = secular approximation where only the slow oscillating terms are retained

Cohen-Tannoudji, Atom-photon interactions: basic processes and applications

# The dynamical equation

The entire dynamical equation is scaled with  $\omega_{SL}$

$$\dot{\rho}'_s = \boxed{-i\left(\alpha'(t') [\sigma_+, \rho'_s] e^{-i\Delta'_- t'} + h.c.\right)} + \boxed{\chi\left(P_1 \mathcal{D}[\sigma_+] + P_2 \mathcal{D}[\sigma_-]\right)}$$

$$+ \boxed{2|\alpha'(t')|^2 \beta\left(\mathcal{D}[\sigma_+] + \mathcal{D}[\sigma_-]\right) - \alpha'_1(t') \beta_2\left(\sigma_+ \rho'_s \sigma_+ e^{-2i\Delta'_- t'} + h.c.\right)}$$

$$\alpha(t) = \frac{u_1(t) - iu_2(t)}{4},$$

$$t' = \omega_{SL} t,$$

$$\beta = \beta_1 + \beta_2, \quad \beta_1 = J[\Delta'_+], \quad \beta_2 = J[\Delta'_-],$$

$$\alpha'(t') = \frac{\alpha(t')}{\omega_{SL}},$$

$$\Delta_- = \omega - \Omega = \text{frequency of the co-rotating frame},$$

$$\Delta'_- = \frac{\Delta_-}{\omega_{SL}},$$

$$\alpha'_1(t') = \alpha'(t')^2 + \alpha'^*(t')^2,$$

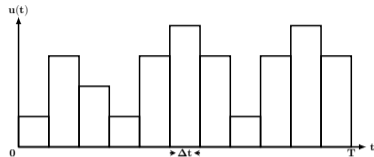
$$\mathcal{D}[\mathcal{O}] = \mathcal{O} \rho'_s \mathcal{O}^\dagger - \frac{1}{2} \{\mathcal{O}^\dagger \mathcal{O}, \rho'_s\},$$

$$\chi = \omega_{SL} \tau_c = \text{environmental correlation time}$$



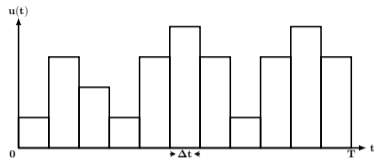
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- GRAPE = GRadient Ascent Pulse Engineering



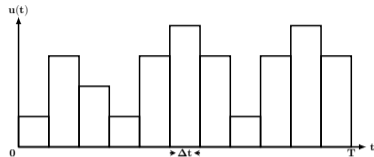
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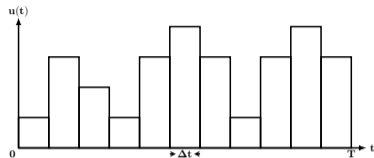
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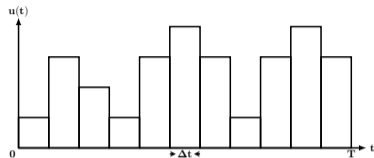
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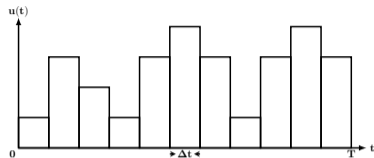
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- The Hamiltonian at  $j^{th}$  time step is:

$$H(j) = H_o + \sum_{k=1}^m u_k(j) H_k$$



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$$F(\rho_0, \rho_\tau) = \left[ \text{Tr} \left\{ \sqrt{\sqrt{\rho_0} \rho_\tau \sqrt{\rho_0}} \right\} \right]^2$$



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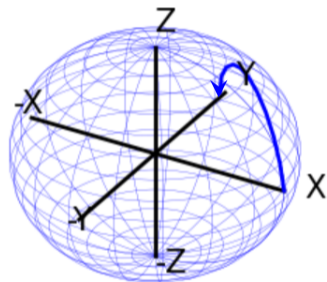
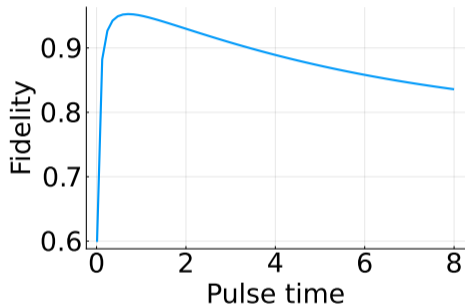
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  - It gives us an **optimal pulse profile** for the given number of steps

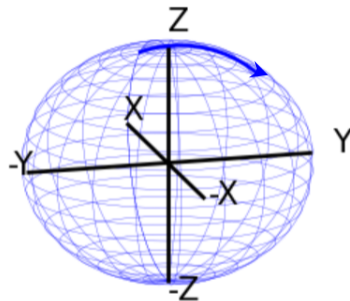
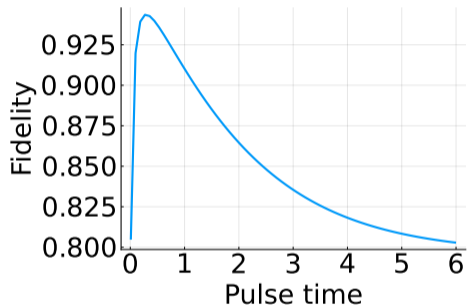
# The results: Pulse Time vs Fidelity



$$|\psi_i\rangle = +X = \frac{1}{\sqrt{2}}(|0\rangle + |1\rangle),$$

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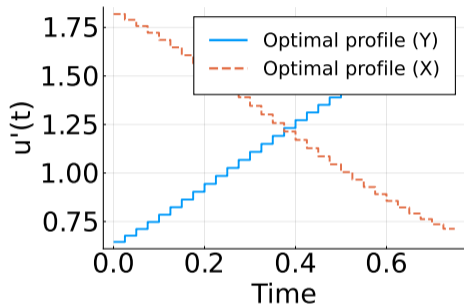
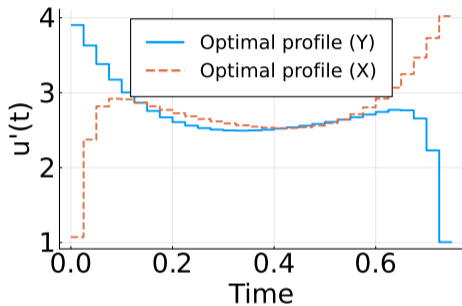
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# The results: Optimal Pulse profile

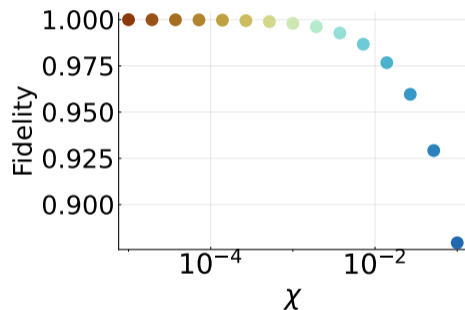
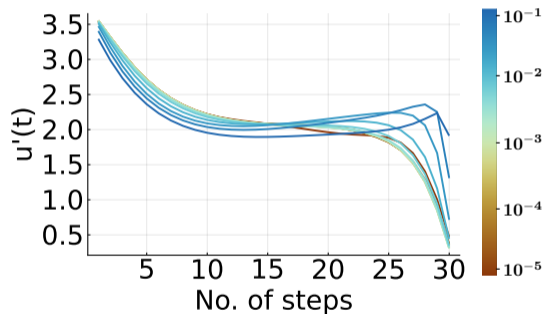


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Varying environmental correlation time ( $\chi$ ) from  $10^{-5}$  to  $10^{-1}$



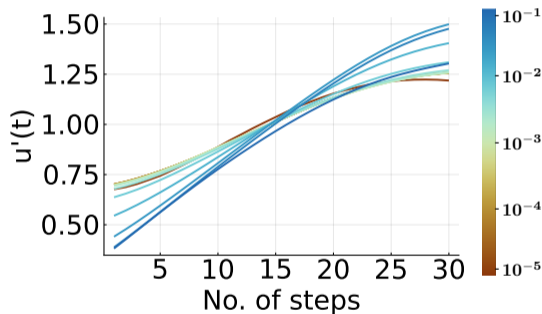
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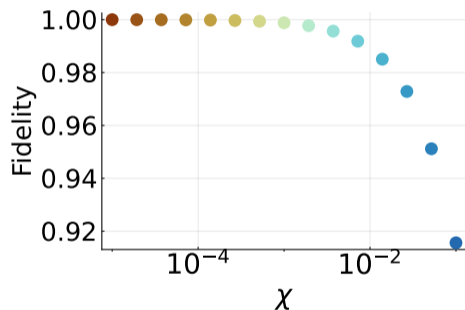


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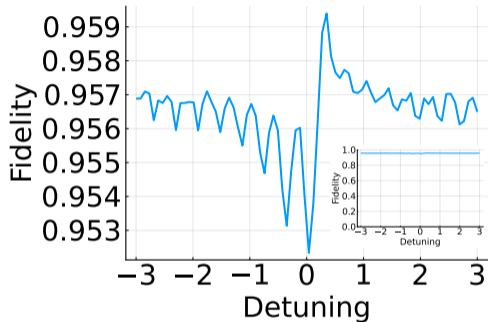


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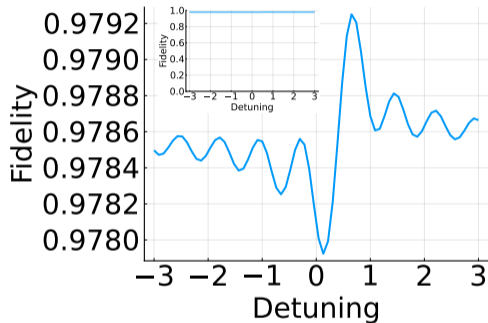


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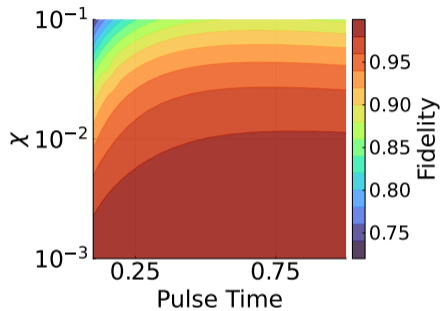


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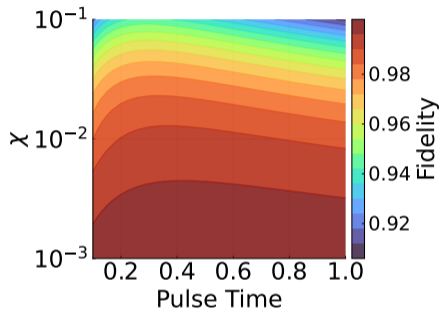


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# Thank You

Optimal Speed of Quantum Operations in Open Quantum Systems

**Sarfraj Fency**, Riddhi Chatterjee and Rangeet Bhattacharyya

*Manuscript under preparation*

- The total Hamiltonian in the lab frame is given as follows:

$$\mathcal{H}(t) = \mathcal{H}_S^\circ + \mathcal{H}_L^\circ + \mathcal{H}_{SL} + \mathcal{H}_S(t) + \mathcal{H}_L(t)$$

where  $\mathcal{H}_S^\circ$  = static Hamiltonian of the quantum system

$\mathcal{H}_L^\circ$  = static Hamiltonian of local-environment

$\mathcal{H}_{SL}$  = Coupling between system and local environment

$\mathcal{H}_S(t)$  = external drive applied on the quantum system.

- The total Hamiltonian of the system in interaction picture of  $\mathcal{H}_S^\circ + \mathcal{H}_L^\circ$  takes the following form:

$$H = H_S + H_L + H_{SL}$$

- The thermal noise from the environment were chosen to be diagonal in the eigen basis of the static Hamiltonian of the environment

$$\mathcal{H}_L = \sum_j f_j |\phi_j\rangle\langle\phi_j|$$

here  $f_j$  is assumed to be Gaussian,  $\delta$  correlated stochastic variable with zero mean and standard deviation  $\kappa$ .

- Finite propagator  $\implies$  regulator from the thermal fluctuations (Schrödinger equation)
- The propagator = system Hamiltonian (infinitesimal) + thermal fluctuations (finite)
- Time scale separation:  $\tau_c \ll \Delta t \ll \tau_s$
- Born approximation:  $\rho(t) = \rho_s(t) \otimes \rho_L^{\text{eq}}$
- Secular approximation:  $|\omega_r| \Delta t \ll 1$

# Final form of FRQME

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$$\dot{\rho}_s = -i \text{Tr}_L[H_{\text{eff}}, \rho]^{\text{sec}} - \int_0^\infty d\tau e^{-\frac{\tau}{\tau_c}} \text{Tr}_L[H_{\text{eff}}(t), [H_{\text{eff}}(t - \tau), \rho]]^{\text{sec}}$$

here,  $\tau_c = \frac{2}{\kappa^2}$ ,  $\rho =$  total density matrix,  $\rho_s =$  system's density matrix  
“sec” = secular approximation

U. Haeberlen, *High Resolution NMR in solids selective averaging* (Elsevier, 2012)

$\text{Tr}_L \implies$  partial trace taken over the bath degrees of freedom

- The effective Hamiltonian:  $H_{\text{eff}} = H_S + H_{\text{SL}}$ .
- The bath is assumed to be isotropic in nature and hence

$$\text{Tr}_L[H_{\text{SL}}, \rho] = 0$$

- Second term of FRQME, predicts the presence of Drive induced Dissipation (DID)