

Fluctuation-dissipation relations for thermodynamic distillation processes

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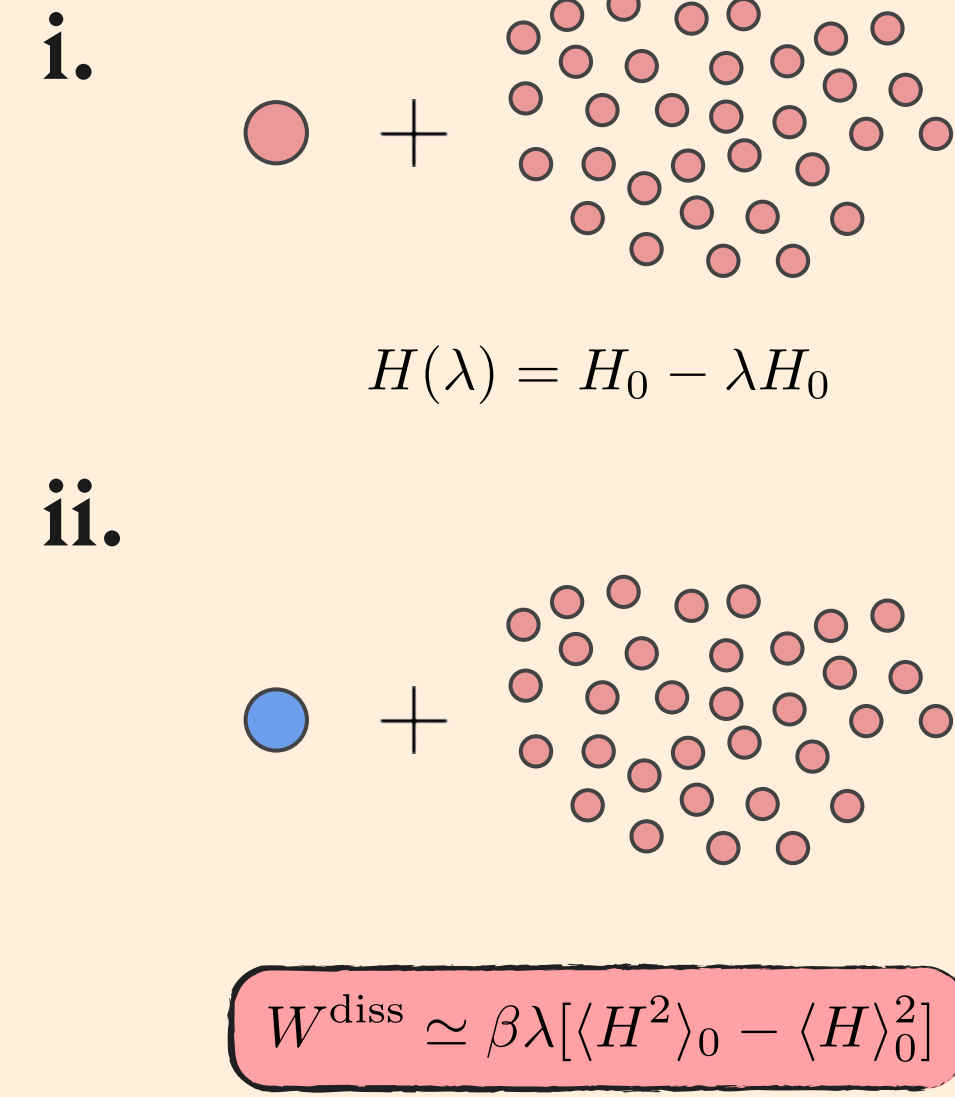
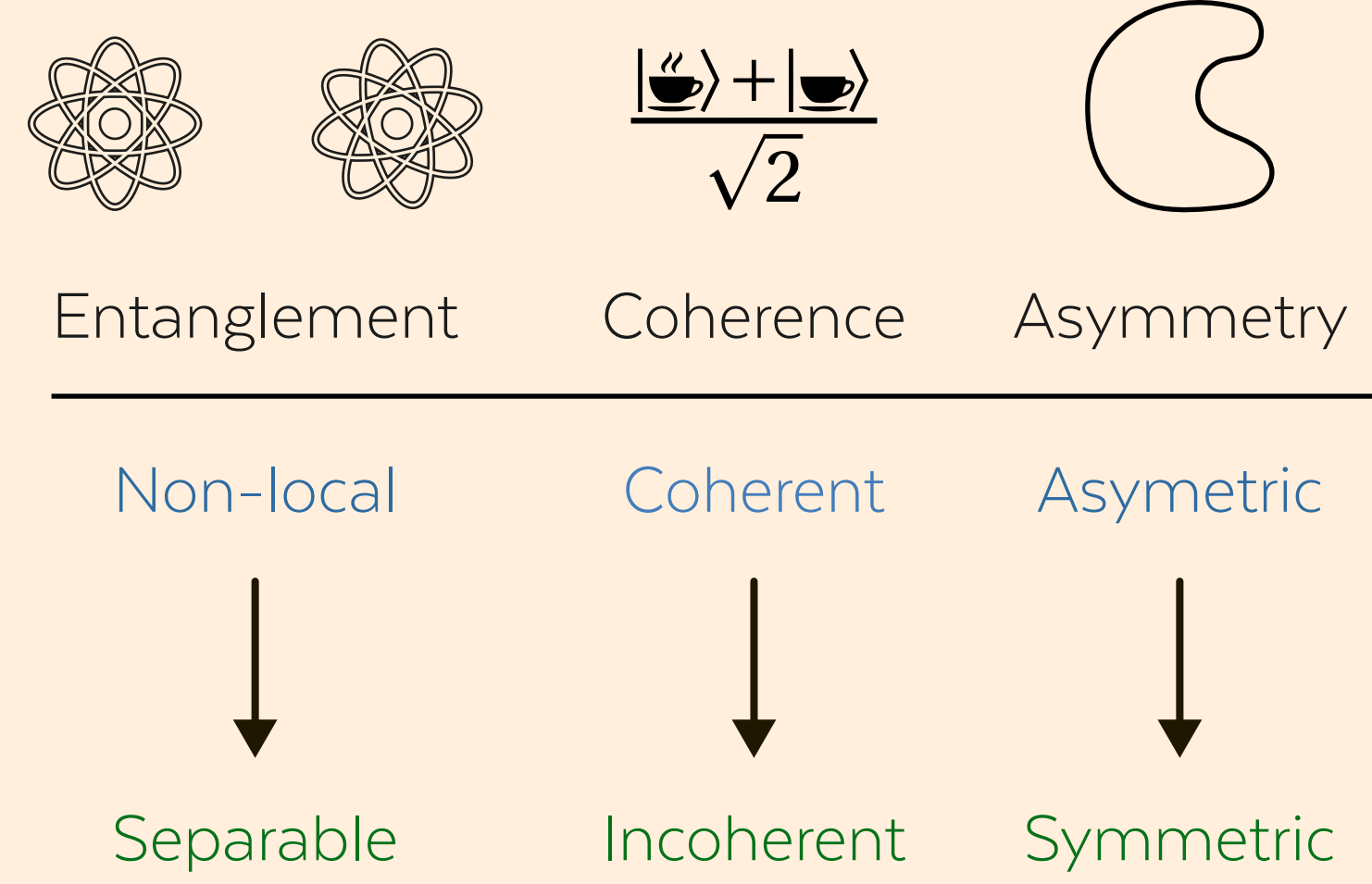
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Background

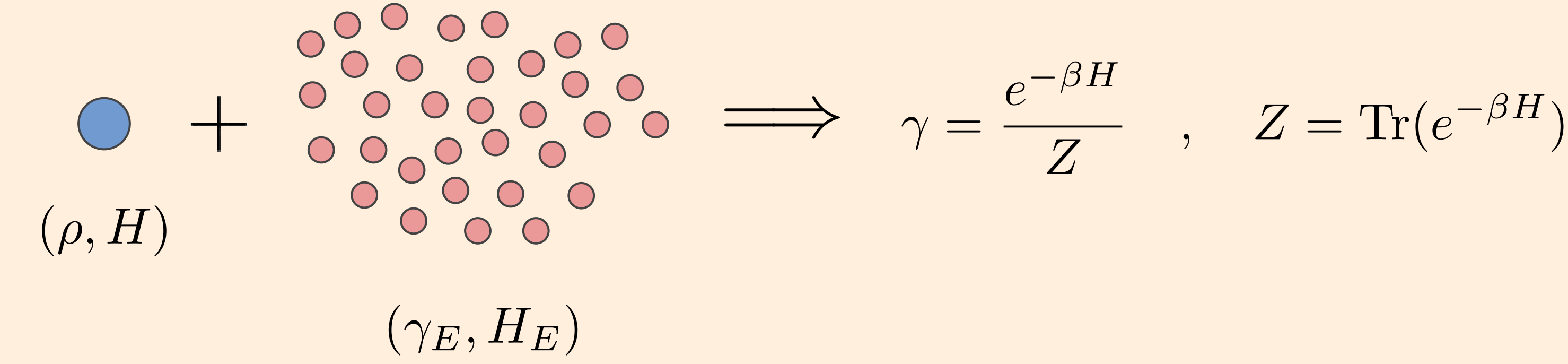
Goal: study resource dissipation and characterise optimal state transformation protocols that minimise dissipation

Framework: ordering among quantum states



Setting the scene

Identifying the set of **thermodynamically-free states**

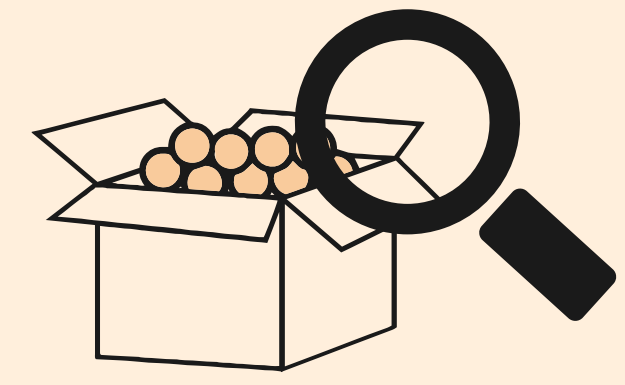


Thermodynamic transformations are modelled by **thermal operations**

$$\mathcal{E}(\rho) = \text{Tr}_E(U(\rho \otimes \gamma_E)U^\dagger) \quad \text{with} \quad [U, H \otimes \mathbb{1}_E + \mathbb{1}_E \otimes H_E] = 0$$

Energy-conserving interaction

Thermodynamic **monotone** $\phi : \mathcal{S}_d \rightarrow \mathbb{R}_+ \cup \{0\}$



$$\text{i. } \phi(\mathcal{E}(\rho)) \leq \phi(\rho)$$

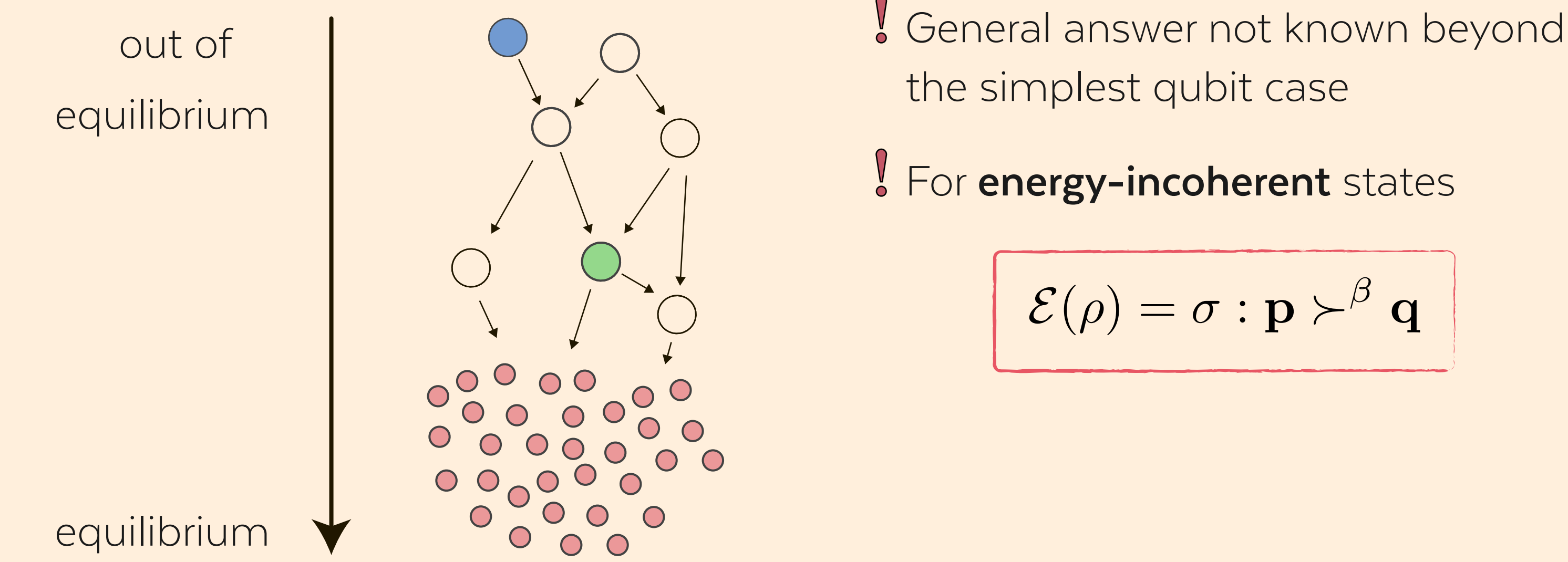
$$\text{ii. } \phi(\gamma) = 0$$

$$D(\rho \parallel \gamma) = \text{Tr}(\rho(\log \rho - \log \gamma))$$

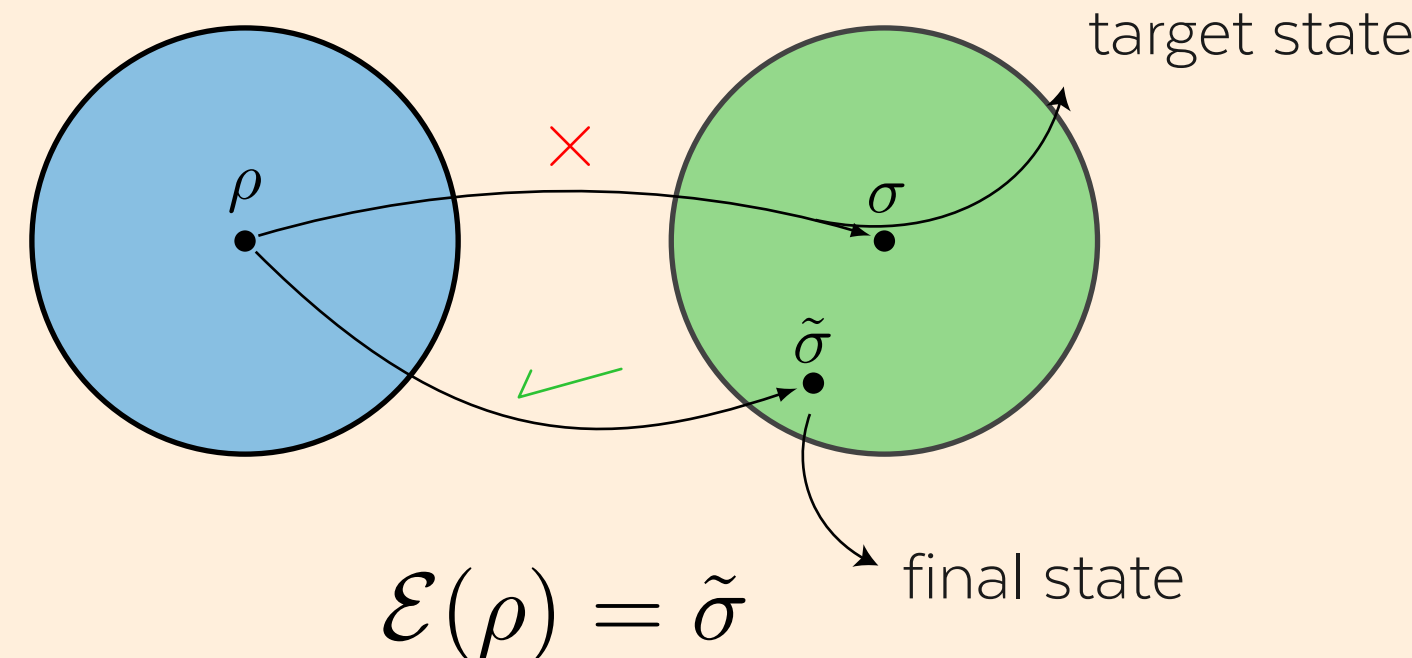
Generalised **free energy**

Thermodynamic distillation process

For initial state ρ , target state σ , thermal bath $\beta \implies \mathcal{E}(\rho) = \sigma$



ϵ -approximate **interconversion** problem:



$$\sigma \approx_\epsilon \tilde{\sigma} \text{ means } 1 - F(\sigma, \tilde{\sigma}) \leq \epsilon$$

ϵ -approximate **thermodynamic distillation process**: $(\rho, H) \xrightarrow{\mathcal{E}} (\tilde{\rho}, \tilde{H})$

$$\text{with } \tilde{\rho} = \bigotimes_{m=1}^{\tilde{N}} |\tilde{E}_{k_n}^{(n)}\rangle \langle \tilde{E}_{k_n}^{(n)}|$$

Eigenstate of $|\tilde{E}_{k_n}^{(n)}\rangle$ corresponding to energy $\tilde{E}_{k_n}^{(n)}$

$$\Delta F^N := \frac{1}{\beta} \left(\sum_{n=1}^N D(\rho_n^N \parallel \gamma_n^N) - D(\tilde{\rho}^N \parallel \tilde{\gamma}^N) \right)$$

Free energy **change**

$$\sigma^2(F^N) := \frac{1}{\beta^2} \sum_{n=1}^N V(\rho_n^N \parallel \gamma_n^N)$$

Free energy **fluctuations**

Results

Theorem 1. For a distillation setting with energy incoherent initial states, the transformation error of the approximate distillation process in the asymptotic limit is given by

$$\lim_{N \rightarrow \infty} \epsilon_N = \lim_{N \rightarrow \infty} \Phi \left(-\frac{\Delta F^N}{\sigma(F^N)} \right)$$

Moreover, for any N there exist an approximate distillation process with the transformation error bounded by

$$\epsilon_N \leq \Phi \left(-\frac{\Delta F^N}{\sigma(F^N)} \right) + \frac{C\kappa^3(F^N)}{\sigma^3(F^N)}$$

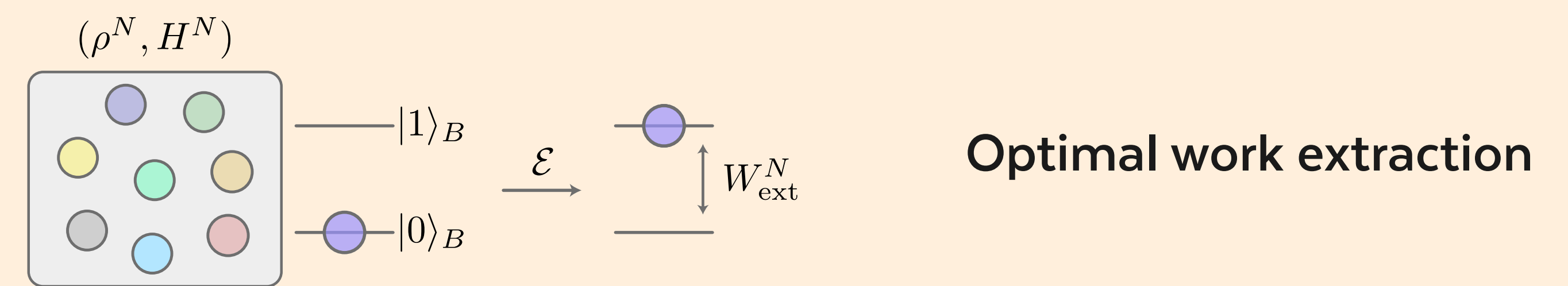
Theorem 2. For a distillation setting with N identical initial systems each in a pure state and described by the same Hamiltonian, the transformation error of the optimal approximate distillation process in the asymptotic limit is given by

$$\lim_{N \rightarrow \infty} \epsilon_N = \lim_{N \rightarrow \infty} \Phi \left(-\frac{\Delta F^N}{\sigma(F^N)} \right)$$

The amount of **free energy dissipated** in both settings satisfies:

$$F_{\text{diss}}^{\text{tot}} = a(\epsilon) \sigma^{\text{tot}}(F) \quad \text{with} \quad a(\epsilon) = -\Phi^{-1}(\epsilon)(1 - \epsilon) + \frac{\exp \left(\frac{-(\Phi^{-1}(\epsilon))^2}{2} \right)}{\sqrt{2\pi}}$$

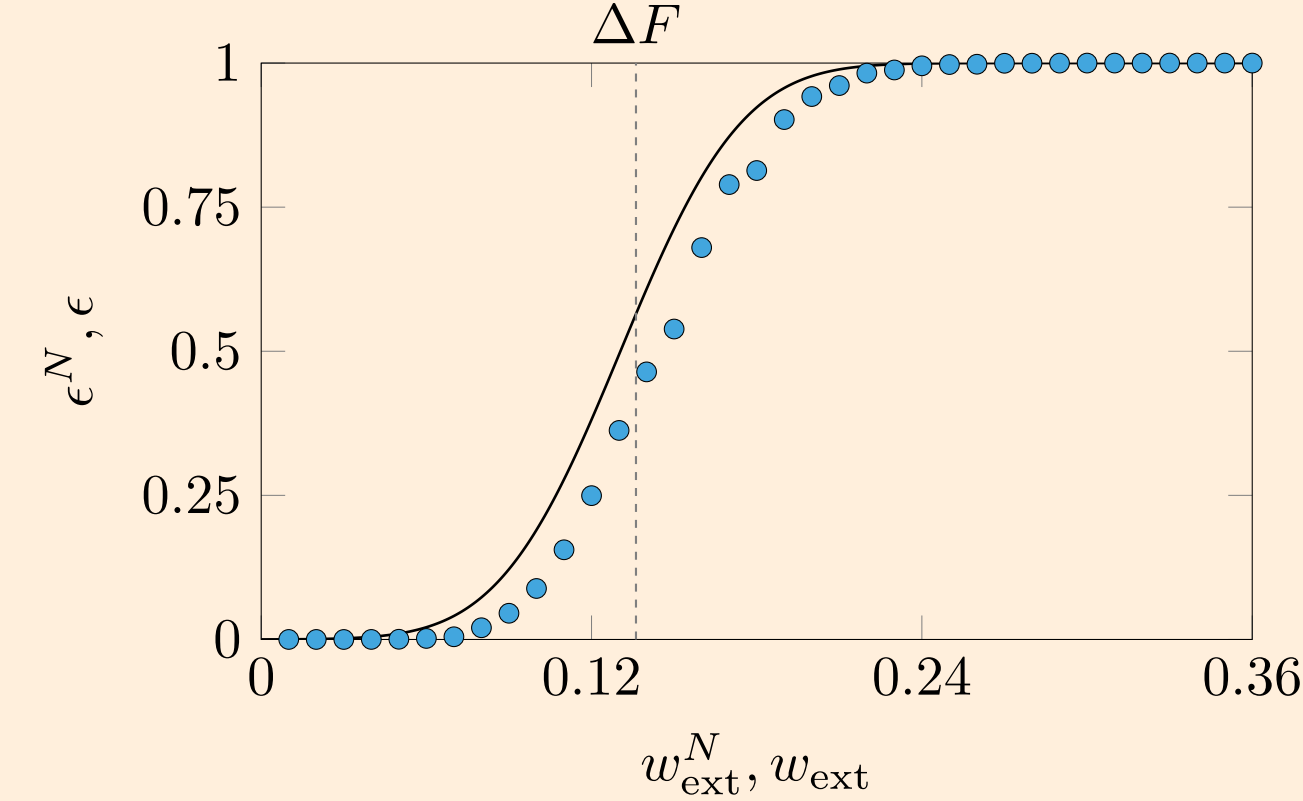
Applications



Optimal work extraction

■ **Theorem 1** tell us that the optimal transformation error for extracting the amount of work W_{ext}^N is:

$$W_{\text{ext}}^N \simeq F^N + \sigma(F^N) \Phi^{-1}(\epsilon)$$



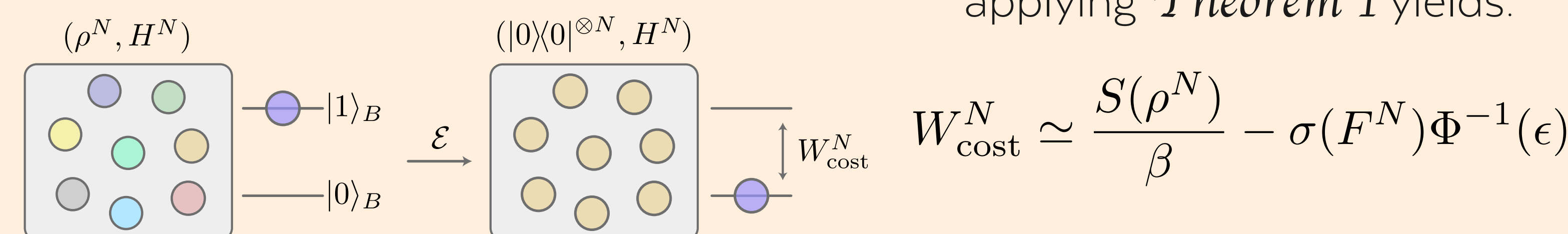
Initial system is composed of 100 two-level subsystems. The first 59 is prepared in the state $p = (0.9, 0.1)$ with thermal state $(0.6, 0.4)$ and remaining is prepared in a state $(0.7, 0.3)$ and thermal state $(0.75, 0.25)$.

■ From **Theorem 2**, the optimal amount of work extracted from N pure quantum systems up to second-order is given by

$$W_{\text{ext}} \simeq N \left(\langle H \rangle_\psi + \frac{\log Z}{\beta} + \frac{\langle H^2 \rangle_\psi - \langle H \rangle_\psi^2}{\sqrt{N}} \Phi^{-1}(\epsilon) \right)$$

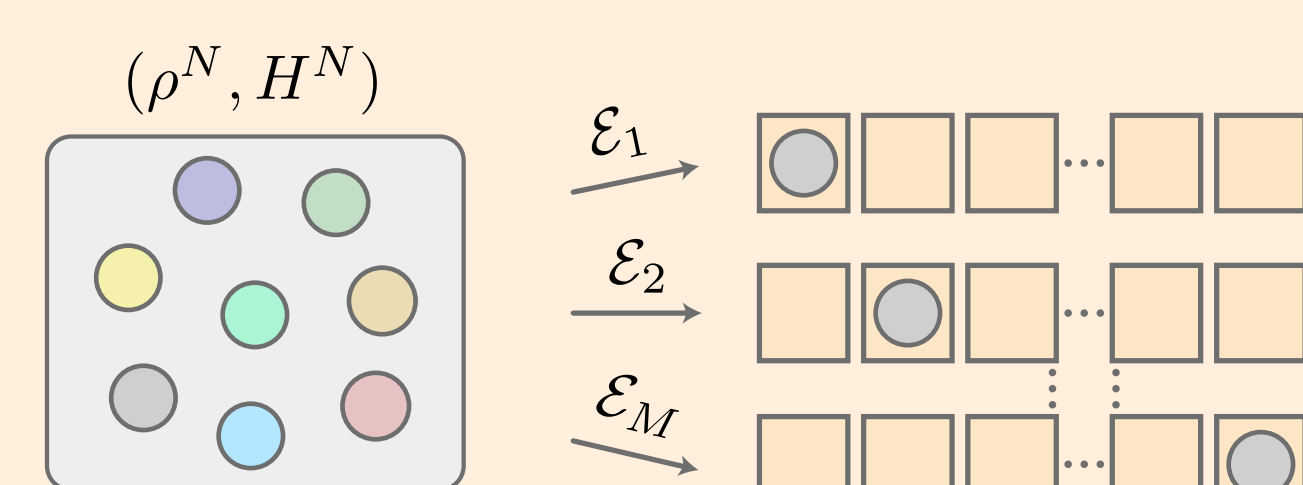
Optimal information erasure

applying **Theorem 1** yields:



$$W_{\text{cost}}^N \simeq \frac{S(\rho^N)}{\beta} - \sigma(F^N) \Phi^{-1}(\epsilon)$$

Optimal thermodynamically-free communication rate



The optimal number of messages that can be encoded into in a thermodynamically-free way:

$$R(\rho^{\otimes N}, \epsilon_d) \simeq D(\rho \parallel \gamma) + \frac{\sqrt{V(\rho \parallel \gamma)}}{\sqrt{N}} \Phi^{-1}(\epsilon_d)$$

! This result is valid for either a pure or incoherent state!