

Thermalization of Quantum Systems II.: Results from Quantum Information Theory

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ELTE Statistical Physics Seminar
17 May 2017

- A central question about thermalization: Why is the Gibbs state special?

$$\rho_\beta = \frac{1}{Z} e^{-\beta H}, \quad Z = \text{Tr}(e^{-\beta H}).$$

We discuss three points:

- **1st lecture** (*Pure-state Statistical Physics*, Márton Kormos):
It naturally emerges from subsystem expectmany-body dynamics.
- **2nd lecture** (*Results from Quantum Information Theory*, ZZ):
It naturally emerges when we introduce actors who can apply different unitary transformations from a given set.
- **3rd lecture** (*Integrable Systems*, Balázs Pozsgay):
Counter-examples, Generalized Gibbs Ensembles.

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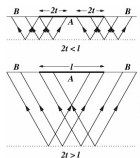
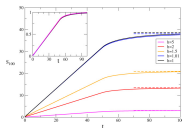
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Last seminar talk: emergence of Gibbs state from unitary dynamics

- Given a Hamiltonian H and a generic pure state with finite energy density. During the time-evolution the entanglement entropy increases linearly in time until $t \sim \ell$:

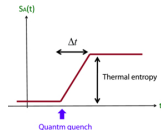


- The state on the local subsystems will be Gibbs states. The entanglement entropy will play the role of thermal entropy:

$$\hat{\rho}_t = e^{\hat{H}_t/\hbar} \hat{\rho}_0 e^{-\hat{H}_t/\hbar}$$

Diagram illustrating the subsystem Ω and its complement Ω^\perp . The entanglement entropy $S_E(t)$ is shown to play the role of thermal entropy.

$\lim_{t \rightarrow \infty} \text{Tr}_{\Omega^\perp} \hat{\rho}_t = ?$



- The *Gibbs state* at *inverse temperature* β corresponding to a Hamiltonian H is given as

$$\rho_\beta = \frac{e^{-\beta H}}{\text{Tr}(e^{-\beta H})}.$$

- We should define a *free energy functional* on the set of density matrices (corresponding inverse temperature β and Hamiltonian H)

$$F(\rho) = \text{Tr}(\rho H) - \beta^{-1} S(\rho),$$

such that the state that maximizes this functional is exactly the Gibbs state ρ_β .

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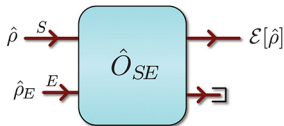
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Quantum information approach: introducing actors who can use ancillary systems and unitaries

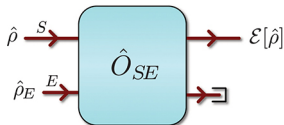
- The starting state $\rho_\beta = \rho_S \otimes \rho_E$ (or at least $\rho \approx \rho_S \otimes \rho_E$) that evolves unitarily.



- Why could $\rho = \rho_S \otimes \rho_E$ be justified? Doesn't that contradict our previous set-up (the entropy is built up from the entanglement with environment)?

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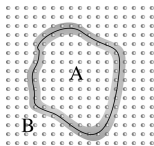
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Subsystem are not very much entangled or correlated

- Monogamy of entanglement
- For **any short-ranged Gibbs** there is an area law for the mutual information of $I(A : B) = S(\rho_A) + S(\rho_B) - S(\rho_{AB})$ of neighboring regions and this mutual information will even decay exponentially with the distance (M.M. Wolf, F. Verstraete, M.B. Hastings, and J.I. Cirac, *Phys. Rev. Lett.* **100**, 070502 (2008)).



- The above results also holds for other correlation measures:
 - Negativity V. Eisler, *ZZ, New J. Phys.* **16**, 123020 (2014); *Phys. Rev. B* **93**, 115148 (2016).
 - Rényi mutual information: $D_i(\rho_{AB} \| \rho_A \otimes \rho_B)$ where

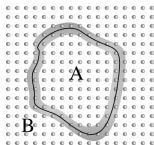
$$D_1^{(\alpha)}(\rho \| \sigma) = \frac{1}{\alpha - 1} \log \text{Tr} (\rho^\alpha \sigma^{1-\alpha}) ,$$

$$D_2^{(\alpha)}(\rho \| \sigma) = \frac{1}{\alpha - 1} \log \text{Tr} \left(\sigma^{\frac{1-\alpha}{2\alpha}} \rho \sigma^{\frac{1-\alpha}{2\alpha}} \right)^\alpha .$$

(Martin Müller-Lennert et al, *J. Math. Phys.* **54**, 122203 (2013); R. L. Frank, E. H. Lieb, *J. Math. Phys.* **54**, 122201 (2013).)

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One counter-example: nonequilibrium steady state generated by a temperature gradient

- The initial state:

$$\rho_0 = \frac{1}{Z_L} e^{-\beta_\ell \mathcal{H}_\ell} \otimes \frac{1}{Z_R} e^{-\beta_r \mathcal{H}_r},$$

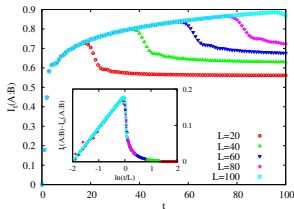
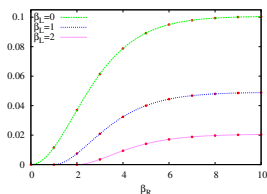
- Time evolution: $\rho(t) = e^{-itH} \rho_0 e^{itH}$, where

$$\mathcal{H} = -\frac{1}{2} \sum_{m=-\infty}^{\infty} \left(c_{m+1}^\dagger c_m + c_m^\dagger c_{m+1} \right)$$



- Result: There is a logarithmic violation of the area law,
 $I(A : B) = \sigma \log L + k$,

$$\sigma = \frac{1}{\pi^2} \left[a \operatorname{Li}_2 \left(\frac{a-b}{a} \right) + (1-a) \operatorname{Li}_2 \left(\frac{b-a}{1-a} \right) + b \operatorname{Li}_2 \left(\frac{b-a}{b} \right) + (1-b) \operatorname{Li}_2 \left(\frac{a-b}{1-b} \right) \right],$$

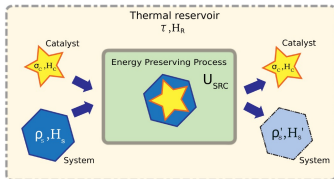


V. Eisler, Z. Zimborás, *Phys. Rev. A* **89**, 032321 (2014),
M. Kormos, Z. Zimborás, [arXiv:1612.04837](https://arxiv.org/abs/1612.04837) (2016).

- Similar counter examples appeared after ours:

M. Hoogeveen, B. Doyon, *Nucl. Phys. B* **898**, 78-112 (2015), S. Ajişaka, F. Barra, B. Zunkovic, *New J. Phys.* **16** 033028 (2014); F. Ares, J. G. Esteve, F. Falceto, E. Burillo, *J. Phys. A* **47**, 245301 (2014).

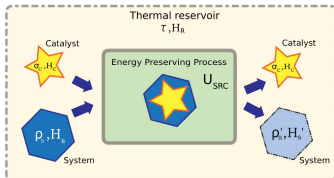
Catalytic thermal operations



$$\rho_S \otimes \tau^{\otimes n} \otimes \sigma \rightarrow U \rho_S \otimes \tau^{\otimes n} \otimes \sigma U^\dagger,$$
$$[U, H_S + H_R + H_C] = 0.$$

- Borrow a catalyst that is returned to itself (or ϵ close to itself).
- Thermal reservoir: add arbitrary number of copies of a thermal state τ belonging to a Hamiltonian H_R .
- Perform any unitary operation U such that $[U, H_S + H_R + H_C]$.
- Perform the partial trace over the reservoir and the catalyst.

A Zeroth Law and Gibbs states



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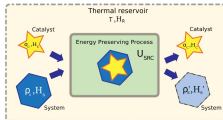
$$[U, H_S + H_R + H_C] = 0.$$

- A type of **Zeroth Law** of Thermodynamics:

If we choose for τ any other form than $\frac{1}{Z} e^{-\beta H_R}$,
then any $\rho_S \rightarrow \rho'_S$ transformation is possible.

(F. Brandao, M. Horodecki, J. Oppenheim, J. Renes and R.W. Spekkens, *Phys. Rev. Lett.* **111**, 250404 (2013).)

A Second Law and the von Neumann entropy



- A type of **Second Law** of Thermodynamics:
If in the above set-up a $\rho_S \rightarrow \rho'_S$ transformation is possible, then

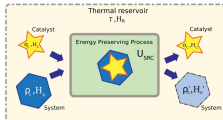
$$F(\rho'_S) \leq F(\rho_S)$$

$$\text{Tr}(H_S \rho'_S) - \beta^{-1} S(\rho'_S) \leq \text{Tr}(H_S \rho_S) - \beta^{-1} S(\rho_S)$$

$$\beta^{-1} D(\rho'_S \| \rho_\beta) - \beta^{-1} \log Z \leq \beta^{-1} D(\rho_S \| \rho_\beta) - \beta^{-1} \log Z$$

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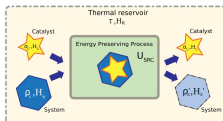
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The Second Laws of Thermodynamics



- However, the decrease of free energy is not the only restriction!
- **Second Laws of Thermodynamics:**
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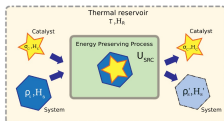
$$\beta^{-1} D_1^{(\alpha)}(\rho'_S \| \rho_B) - \beta^{-1} \log Z \leq \beta^{-1} D_1^{(\alpha)}(\rho_S \| \rho_B) - \beta^{-1} \log Z \text{ for } \frac{1}{2} \leq \alpha \leq 1,$$

$$\beta^{-1} D_2^{(\alpha)}(\rho'_S \| \rho_B) - \beta^{-1} \log Z \leq \beta^{-1} D_2^{(\alpha)}(\rho_S \| \rho_B) - \beta^{-1} \log Z \text{ for } \alpha \geq 1.$$

It is then customary to introduce

$$F_i(\rho \| \rho_B) = D_i^{(\alpha)}(\rho'_S \| \rho_B) - \beta^{-1} \log Z.$$

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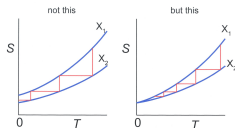
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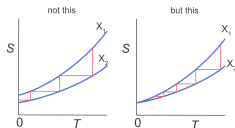
Extension of the Third Law of Thermodynamics



- Nernst's argument states that: at zero temperature, a finite size system has an entropy S , which is independent of any external parameters X , that is $S(T, X_1) - S(T, X_2) \rightarrow 0$ as the temperature $T \rightarrow 0$.
- As the entropy at zero temperature should be the logarithm of the ground state degeneracy, the validity of the third law is contingent on whether the degeneracy changes for different parameters.
- By modeling a physical process using two main ingredients:
 - finite information (entropy) propagation speed,
 - bound on the entropy $S(E) \leq \alpha V^{1-\nu} E^\nu$ with $\nu \in (1/2, \frac{d}{d+1})$,one can derive a temperature-time bound

$$T_S \geq \frac{\text{const.}}{t^\gamma}.$$

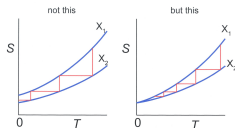
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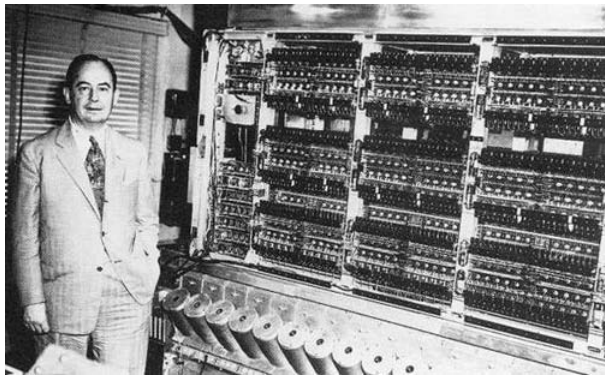
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Quantum Computers with a Neumann Architecture

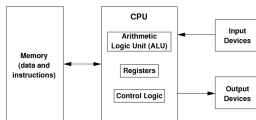
Thermodynamics
and
Quantum
Information

Zoltán
Zimborás

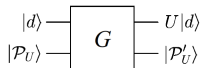


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- The classical Neumann architecture:



- The quantum mechanical version: [Universally Programmable Quantum Computer](#)

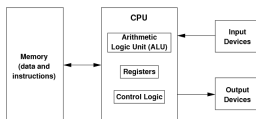


- One needs a program input size that scales between $\exp(d)$ and $\exp(\exp(d))$.

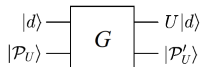
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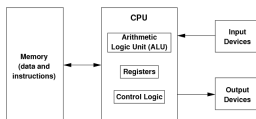


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**14TH JOINT EUROPEAN
THERMODYNAMICS
CONFERENCE - MAY 21-25
2017 - BUDAPEST, HUNGARY**

Josiah Willard Gibbs
(1839-1903)
American



Aims & scope
Committees
Scientific Program
Mini-Symposia

AIMS & SCOPE

Methods and concepts of equilibrium and non-equilibrium thermodynamics appear in various areas of physics, engineering, sciences