Title: Phenomenological thermodynamics with multiple quantities of interest Speakers: Lidia del Rio Series: Quantum Foundations Date: October 21, 2022 - 2:00 PM URL: https://pirsa.org/22100138 Abstract: Joint work (in progress) with Ladina Hausmann, Nuriya Nurgalieva and Renato Renner

We can classify contemporary approaches to thermodynamics in roughly four camps:

(1) Top-down microscopic approaches. These are for example resource-theoretical approaches to quantum thermodynamics: they have a microscopic model of states and systems, and which microscopic restrictions implement macroscopic properties. For instance, in the resource theory of quantum thermodynamics, states are represented by density operators, thermal states in particular have a specific micro-canonical form, and constraints like energy preservation are enforced by forcing quantum transformations to commute with a global Hamiltonian. These approaches success at deriving thermodynamic laws in general settings that satisfy the microscopic model (like non-relativistic quantum systems.

(2) Bottom-up microscopic approaches. These also start from a microscopic model, but rather than looking for universal restrictions, they search for explicit thermodynamics protocols: this is the case of recent proposals for quantum work extraction or nano quantum heat engines.

(3) Top-down phenomenological approaches. These try to derive thermodynamic laws from first principles independently of a microscopic model. In principle the results derived in this framework can be applied to a wider variety of explicit systems, and the challenge is then to find the right implementations. The first derivations of thermodynamics were naturally phenomenological, and some modern information-inspired derivations follow this approach.

(4) Bottom-up phenomenological approaches. These approaches try to find explicit thermodynamic protocols independently of the microscopic model, based only on operational properties of the systems at hand. It was the case for Carnot's original engines and more recently for some approaches to deriving black hole thermodynamics, or thermodynamics of new materials; some experimental results also fit in this camp.

In this work we generalize top-down phenomenological approaches to the case of multiple conserved quantities. Note that multiple conserved quantities have been studied in top-down and bottom-up microscopic approaches to quantum thermodynamics. We argue that our framework is more general, in that it can be applied to systems for which we don't have an explicit microscopic model; in particular we will apply the results of this framework to black hole thermodynamics. Moreover, having a phenomenological axiomatic approach to thermodynamics allows us to identify which properties are specific to a microscopic model like quantum physics, and which hold in any physical theory: our results can be applied to study the thermodynamics of generalized process theories, and other generalizations and foils of quantum mechanics. This generalization makes us reconsider the second law of thermodynamics, adapting for an exchange of different conserved quantities, for example, energy and angular

momentum, or energy and spin. Our guiding principle here is to use information as a universal token of exchange to convert between different quantities via Landauer's principle.

Zoom Link: https://pitp.zoom.us/j/96001094153?pwd=YTArTGpPdEJ1NFBMcnFqV1dIRTVyZz09

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n Formation + Energy + Angular mom. + Particle numer +



















Step 2: Erase BB - > spond W=-kTeln2H = Qc



 $W = (T_H - T_c) k_b \ln a \delta H$ $=(T_{H}-T_{c})$ Qh La Carnot's efficiency

× Energy
Angular momentum
Information

$$E_{1----}$$

 $e_{1+1+1+1}$
 $deg.$ Spin lattice
 $(E)=0$
 L_{2}
 L











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Elabal view of local processors frames VS, VR: SAR=p-19 and GR + [PIS= | Pxi] YPE 3, HOREZR, 3

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∀S ∀B, PEP → W^B_s(P) → additive: W^b_s(P)= ≥ W_f(P) First Law YS, YP, OEZS, JPERS: P=0, Hor JPERS: OSP Instant $\mathcal{O}_{\mathcal{D}} = (\mathcal{O}_{\mathcal{O}_{\mathcal{O}}})$

∀S ∀B, PEP → W^B_s(P) → additione: W_s(P)= ? First Law HS, HP, OEES, JPEPS: p=0, HPEPS, W^B_S(P)=Function (TPT, LPJ,) And in case of the local division of the loc $D = (\mathcal{D}_1, \mathcal{O}_2)$

US UB, PEP -> WB(P) -> addition: Ws (P)= ? First Law HS, HP, OEES, JPEPS: p=0, HPEPS, W^B_s(P)=Function (TPTs, LPJs) I NAME OF TAXABLE $=(\mathcal{D}_{1},\mathcal{O}_{2})$

Conserved quantities D33 $\forall S \forall B, P \in P \rightarrow W_{S}^{B}(P) \rightarrow addition W_{S}^{B}(P) = \ge$ W (P) First JPER: p=0 $\forall s, \forall p, \sigma \in \overline{Z}_s$ HPEPs, WS(P)=Function (TPT, LPJs)

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OUB Fully Legenerate For BEB (UR) BEB' injective on set or states, each UR= IR Passive B, US, UPEPRS, - IF TPTS=LPJS =>W(P)>0 .9

CB nvariani ¥ SR EZ Legenerate For BEEB 1Ly injective on set or states, each UR= IR Ĩr'Js FB, FS, FPEPRS, -iF FPTS = LPJS=> $W_{R}^{S}(P) \ge 0$ 3 CAUTION



First Law YS, YP, OEZS, JPEPS: p=5 Jor JPEPS: o≤p YPEPS, WS(P)=Function (IPTS, LPJS) $G_{R} = (\sigma_{1}, \sigma_{2})$