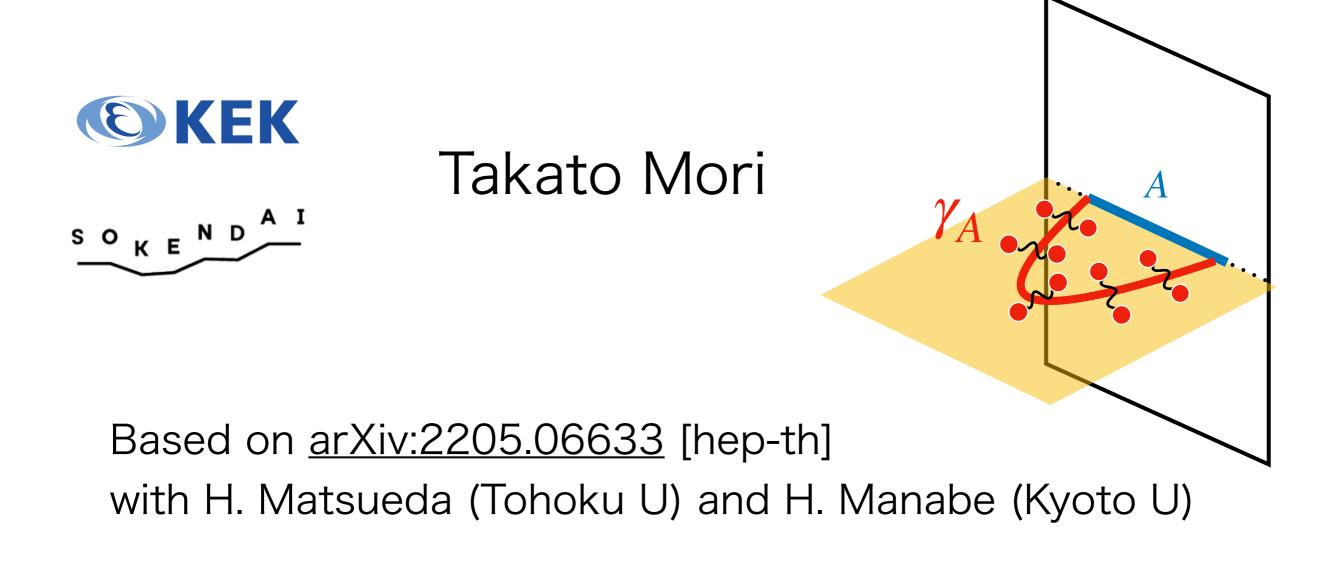
Entanglement distillation in tensor networks



熱場の量子論とその応用,2022年9月21日(オンライン参加)

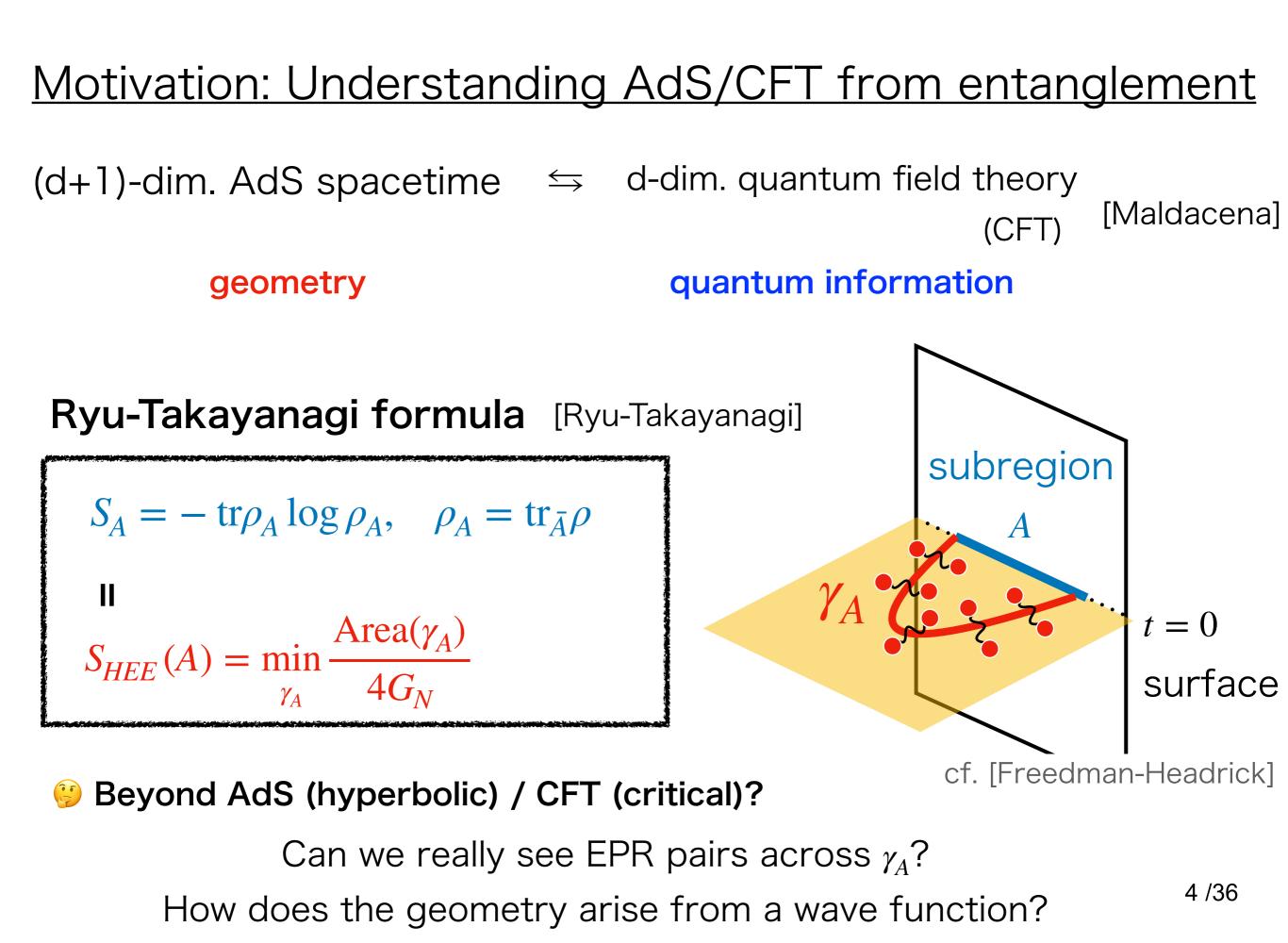
Take-home message

Tensor network defines a set of reduced *transition* matrices They describe entanglement distillation via geometry

The method works for arbitrary tensor networks; a systematic, quantitative study of states vs. geometry in tensor networks is now possible

Outline

- Introduction AdS/CFT, tensor networks as toy models
- 2. Entanglement distillation in MERA
- 3. Numerical results for random MERA
- 4. Entanglement distillation in MPS
- 5. Summary



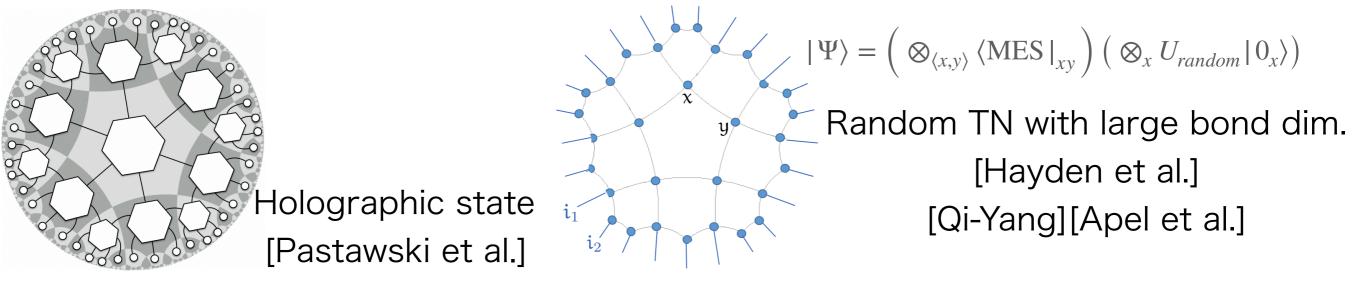
Tensor networks as toy models of holography

 Tensor networks (=variational wave function) provide a qualitative picture [Swingle]

 $S_{TN}(A) \lesssim \min_{\gamma_A} (\# \text{ bond cut by } \gamma_A) \times \log \chi \sim \text{RT formula}?$

Multi-scale entanglement renormalization ansatz (MERA) [Vidal]

• Some proposals try to mimic holography (esp. RT formula)



CONS: Lack of expressivity; TN state ≠ conformally invariant

 γ_A

Entanglement distillation in holographic tensor networks

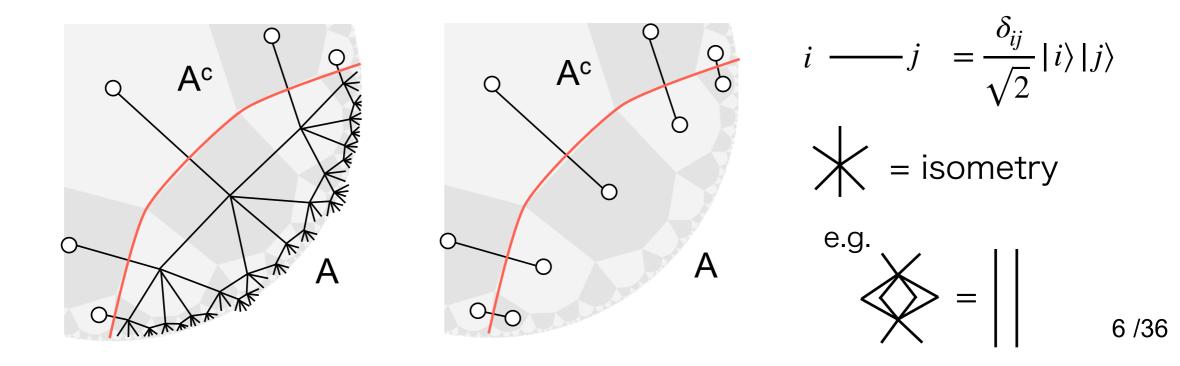
[Pastawski et al.]

(Similar work: [Bao-Penington-Sorce-Wall], [Lin-Sun-Sun])

 A holographic state (or isometric tensor networks in general) is known to geometrize <u>entanglement distillation</u>

= extracting S_A bits of EPR pairs from the state

• Removing tensors, we obtain EPR pairs across the minimal surface $:: S_A(V|\Psi\rangle) = S_A(|\Psi\rangle)$



Entanglement distillation in holographic tensor networks

[Pastawski et al.]

(Similar work: [Bao-Penington-Sorce-Wall], [Lin-Sun-Sun])

 A holographic state (or isometric tensor networks in general) is known to geometrize entanglement distillation

= extracting S_A bits of EPR pairs from the state

• Removing tensors, we obtain EPR pairs across the minimal surface $:: S_A(V|\Psi\rangle) = S_A(|\Psi\rangle)$

So far this manipulation is limited to isometric TNs.

Geometrization of entanglement distillation in other types of TNs?

→ Clarify the operational role of internal d.o.f. after optimization!

Entanglement distillation in tensor networks

The important aspect for entanglement distillation is

1. Conservation of entanglement (entropy)

2. Extracting strongly entangled pairs

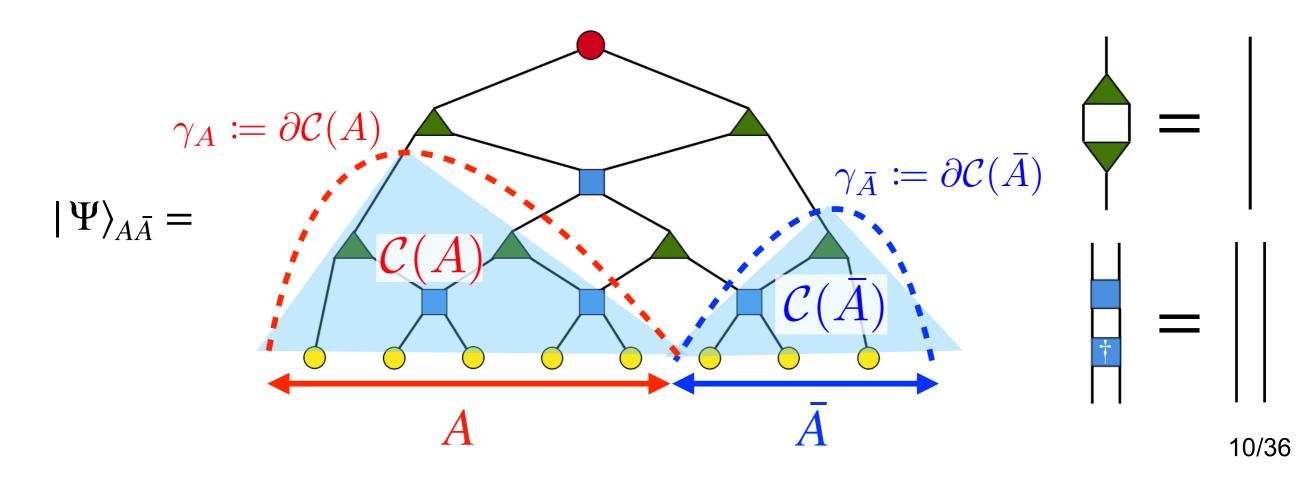
Entanglement distillation in tensor networks

The important aspect for entanglement distillation is

- 1. Conservation of entanglement (entropy)
 - Reduced transition matrix instead of reduced density matrix
- 2. Extracting strongly entangled pairs
 - Nontrivial for non-isometic TNs but can be systematically studied

Multi-scale entanglement renormalization ansatz (MERA) [Vidal]

- MERA has minimal bond cut surface(s) $\gamma_* = \min(\gamma_A, \gamma_{\bar{A}})$
- What we want to see: A state on $\gamma_* \stackrel{?}{=} EPR$ pairs
- Need to provide a method to properly define a state on a bond cut surface γ



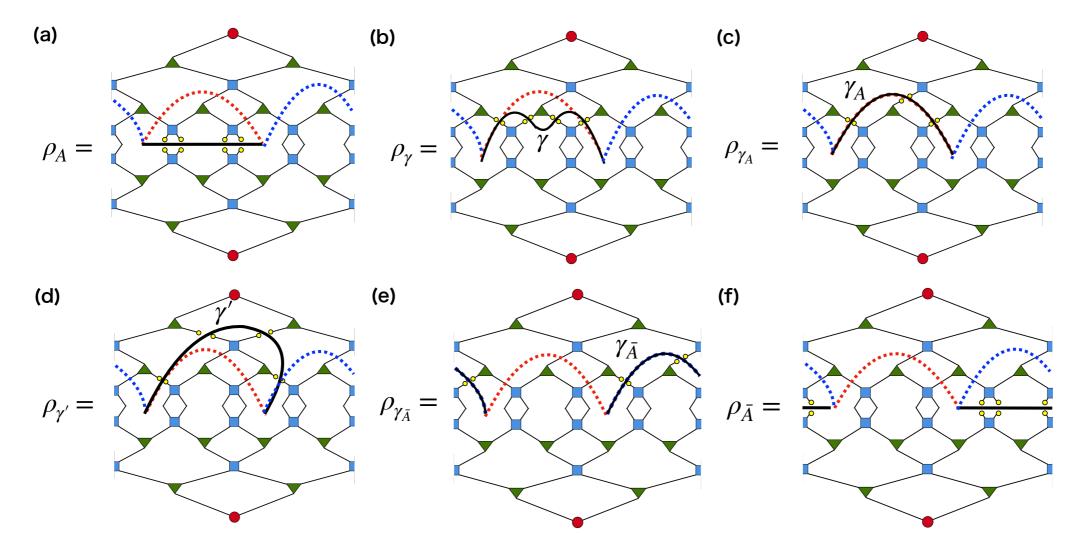
[TM-Manabe-Matsueda]

- Cut internal bonds across a bond cut surface γ
 instead of removing tensors from
 This defines a reduced transition matrix ρ_γ = tr_Ā (|Ψ(γ)⟩⟨Φ(γ) |) on ℋ_γ
- To relate it with entanglement distillation, we consider foliations $\{\gamma\}$ (a) s.t. $\partial \gamma = \partial A$. $\in \mathcal{H}_{\gamma} \otimes \mathcal{H}_{\bar{A}}$ (b` $\rho_{\gamma} =$ $=|\langle \psi(\phi)\rangle| =$ $\in \mathscr{H}^*_{\gamma} \otimes \mathscr{H}^*_{\bar{A}}$ $\equiv \langle \Psi | M_{\gamma}$

[TM-Manabe-Matsueda]

- Changing the location of foliations γ , we obtain a family of states on various bond cut surfaces (e.g. ρ_A for $\gamma = A$, ρ_{γ_*} for $\gamma = \gamma_*$)
- Entanglement conservation w.r.t. $\forall \gamma : S(\rho_{\gamma}) = S(\rho_{A})$, where $S(\rho_{\gamma})$ is

the pseudo entropy [Nakata-Takayanagi et al.] $S(\rho_{\gamma}) = -\operatorname{tr} \rho_{\gamma} \log \rho_{\gamma}$



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[TM-Manabe-Matsueda]

- Changing the location of foliations γ , we obtain a family of states on various bond cut surfaces (e.g. ρ_A for $\gamma = A$, ρ_{γ_*} for $\gamma = \gamma_*$)
- Entanglement conservation w.r.t. $\forall \gamma : S(\rho_{\gamma}) = S(\rho_A)$, where $S(\rho_{\gamma})$ is the pseudo entropy $S(\rho_{\gamma}) = -\operatorname{tr} \rho_{\gamma} \log \rho_{\gamma}$
 - This is owing to the common eigenvalue distribution between ρ_{γ} and ρ_{A} ; the same entanglement spectrum!
- Furthermore, when M_{γ} is isometric, we can show $\langle \Phi(\gamma) | = \langle \Psi(\gamma) |$ \Rightarrow For isometric TNs, we obtain EPR pairs across the minimal bond cut surface (i.e. $\rho_{\gamma_*} \propto 1$) when $\gamma = \gamma_*$

[TM-Manabe-Matsueda]

- To go beyond the isometric case, we need to define a state from a reduced transition matrix.
 - We use the purification technique (a.k.a. channel-state duality)

$$|\rho_{\gamma}^{1/2}\rangle \equiv \mathscr{N}_{\gamma}\sqrt{\dim \mathscr{H}_{\gamma}}(\rho_{\gamma}^{1/2}\otimes\mathbf{1}) |\operatorname{EPR}_{\gamma}\rangle,$$

where $\mathscr{N}_{\gamma} = \left[\operatorname{tr}(\rho_{\gamma}^{\dagger 1/2}\rho_{\gamma}^{1/2})\right]^{-1/2}$ and $|\operatorname{EPR}_{\gamma}\rangle = (\dim \mathscr{H}_{\gamma})^{-1/2}\sum_{i=1}^{\dim \mathscr{H}_{\gamma}} |i\rangle \otimes |i\rangle.$

This will be regarded as a geometrically distilled state up to γ by TN.

Now we can make a quantitative comparison w.r.t. EPR pairs!

→ Trace distance from the EPR pair: $D_{\gamma} \equiv \sqrt{1 - \left| \langle \text{EPR}_{\gamma} | \rho_{\gamma}^{1/2} \rangle \right|^2}$

(related to Rényi-1/2 entropy
$$\left| \langle EPR_{\gamma} | \rho_{\gamma}^{1/2} \rangle \right|^2 = \frac{\mathcal{N}_{\gamma}^2}{\dim \mathcal{H}_{\gamma}} e^{S_{1/2}}$$
) 14/36

Random MERA

- Random TN is expected to reproduce RT formula in the large bond dimension limit [Hayden et al.]
- Why is the minimal bond cut surface is special? (RT formula only tells us about entanglement entropy)
- The large bond dimension limit is essential?

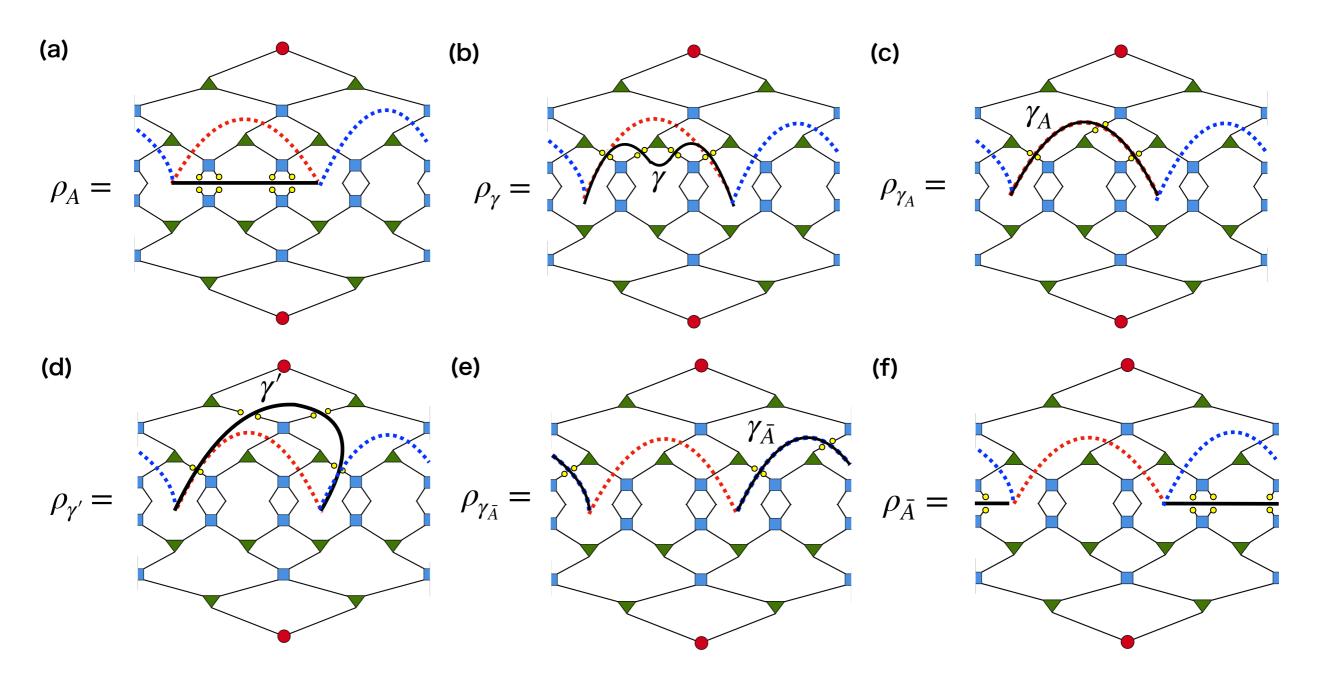
(All tensors are Haar random)

 $|0\rangle$

Haai

Numerical results for random MERA [TM-Manabe-Matsueda]

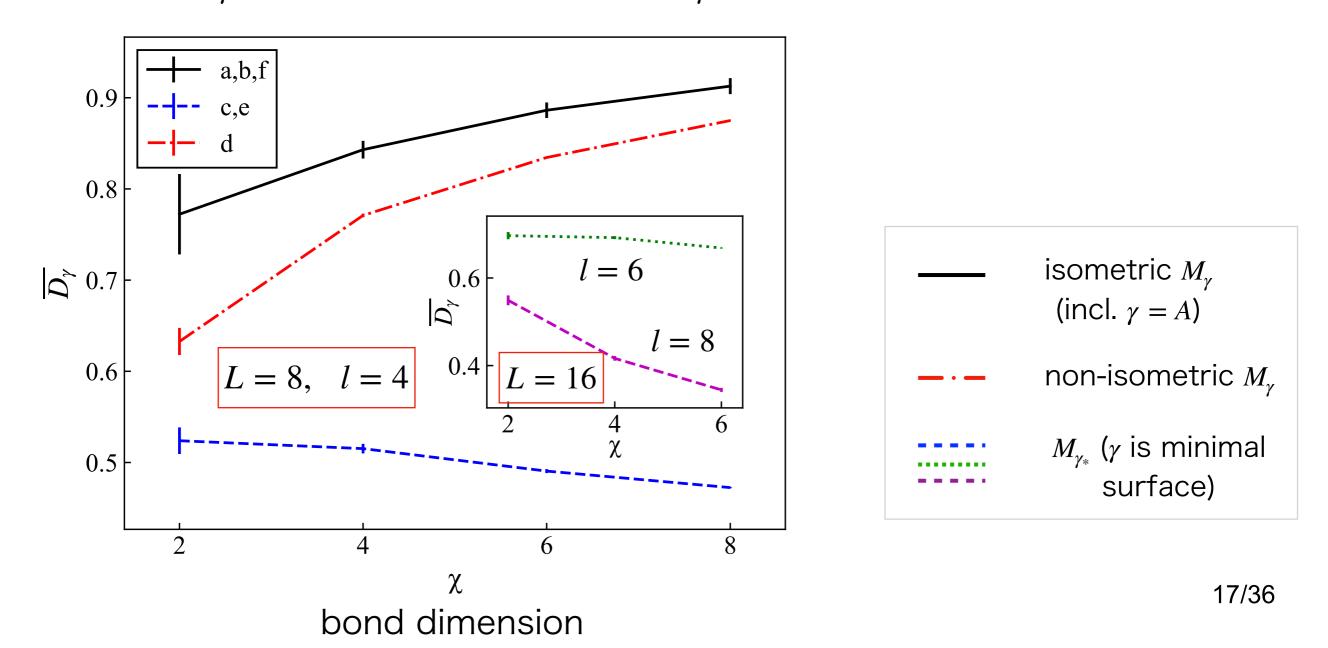
Choice of foliations:



Numerical results for random MERA [TM-Manabe-Matsueda]

We compared the (averaged) trace distance $D_{\gamma} \equiv \sqrt{1 - \left| \langle \text{EPR}_{\gamma} | \rho_{\gamma}^{1/2} \rangle \right|^2}$

between $|\rho_{\gamma}^{1/2}\rangle$ and the EPR pair $|\text{EPR}_{\gamma}\rangle$.

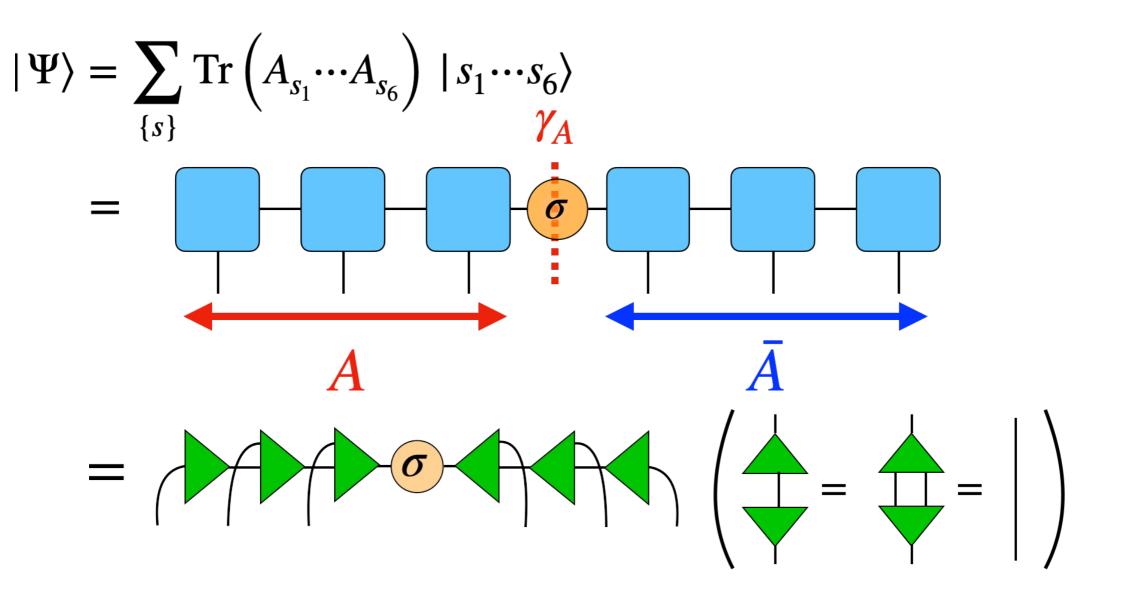


[TM-Manabe-Matsueda]

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Similarly consider pushing the foliation towards the minimal bond cut surface in matrix product states (MPS)

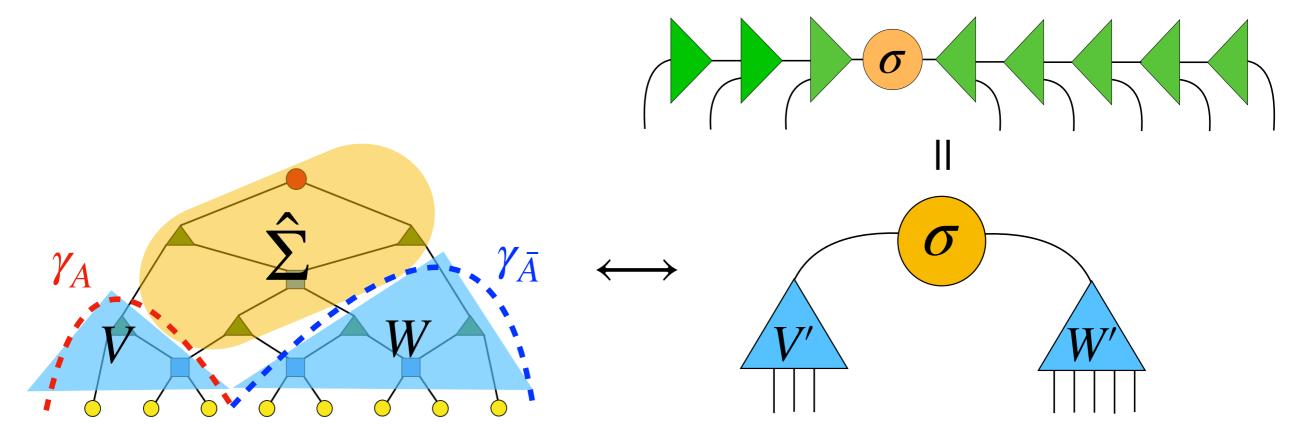
Let us focus on the MPS in a mixed canonical form



Entanglement distillation in MPS [TM-Manabe-Matsueda]

<u>Note:</u>

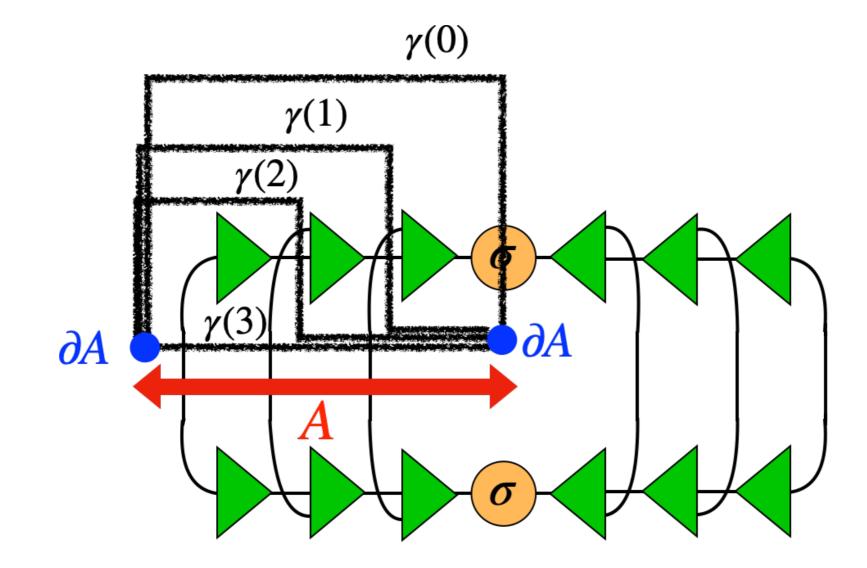
An MPS in a mixed canonical form is an analog of MERA (regarding its structure)



[**TM**-Manabe-Matsueda]

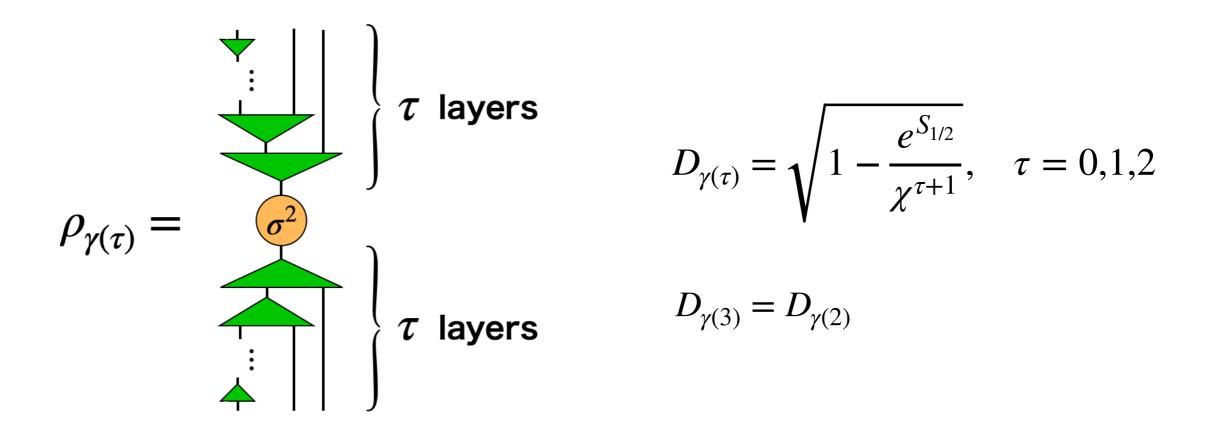
We can consider the following foliations $\gamma = \gamma(\tau)$, $\tau = 0,1,2,3$

($\tau \sim$ distance from the minimal bond cut surface)



Entanglement distillation in MPS [TM-Manabe-Matsueda]

The reduced transition matrix and the trace distance are



The distillation by pushing the foliation equals removing redundant tensors.

The entanglement spectrum σ remains unchanged.

Summary

- . By pushing towards the minimal surface γ_* , strongly entangled pairs are geometrically distilled in tensor networks while retaining the entanglement spectrum
- It is essential to consider reduced transition matrices rather than a reduced density matrix
- Our method works for non-holographic TNs: This suggests geometry of TN is intimately related to distillation for generic TNs
 ➡ Holography beyond AdS/CFT Emergent geometry from distillation

Future directions

- Operational interpretation of geometric distillation: local conf. trf., corner transfer matrix, modular flow? [Milsted-Vidal; Nishino-Okunishi; Okunishi-Seki]
- Analytic "proof" beyond MPS using analytic MERA rep. [Evenbly-White]
- CFT realization? Quant. adiabatic comp. (~annealing) with $T\bar{T}$

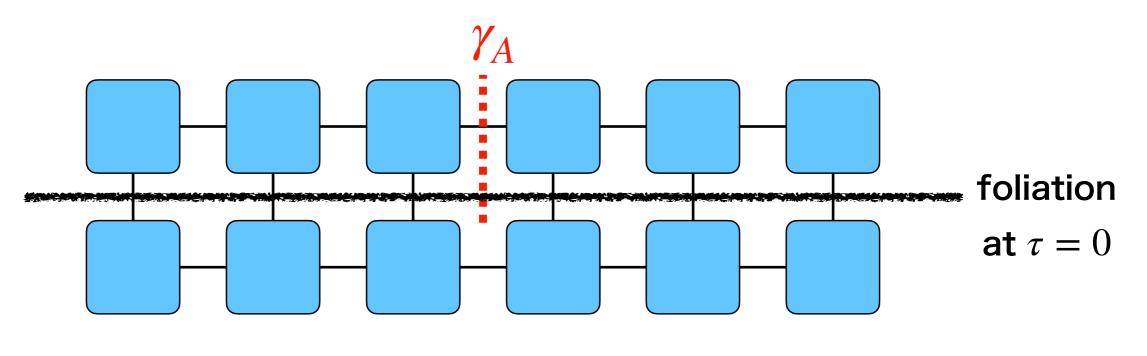
[McGough-Mezei-Verlinde]

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Appendix

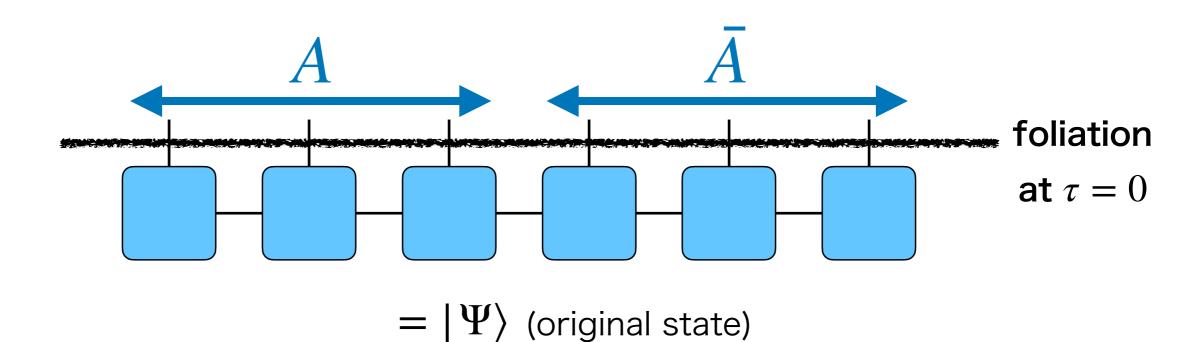
Example: Matrix Product States $|\Psi\rangle = \sum_{\{s\}} \operatorname{Tr} \left(A_{s_1} \cdots A_{s_6} \right) |s_1 \cdots s_6 \rangle$ = 4 A \overline{A}

- 1. Push the boundaries for each A and \overline{A} towards the RT (min bond cut) surface (cf. surface/state correspondence [Miyaji-Takayanagi])
- 2. Define a new state on a pushed boundary ("foliation") by removing the unreached part of TN and taking an inner product with the original TN

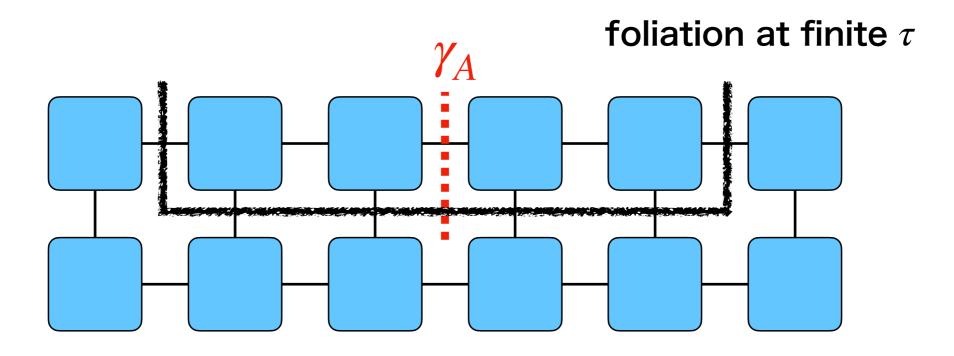


(τ : flow time)

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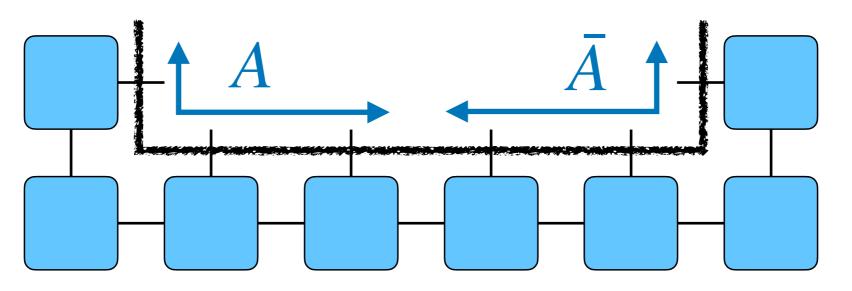


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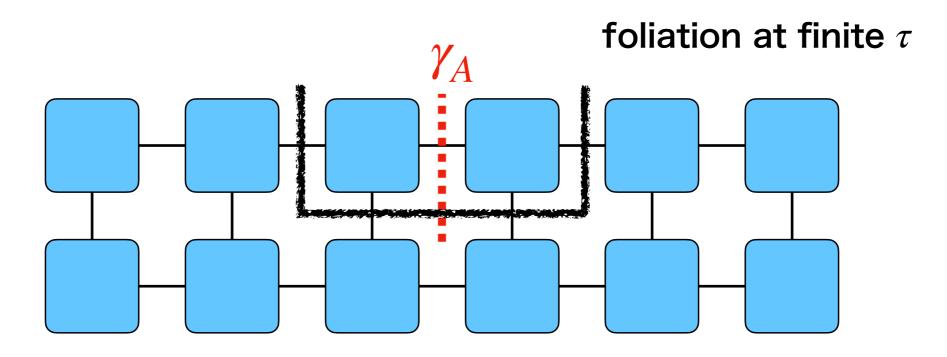


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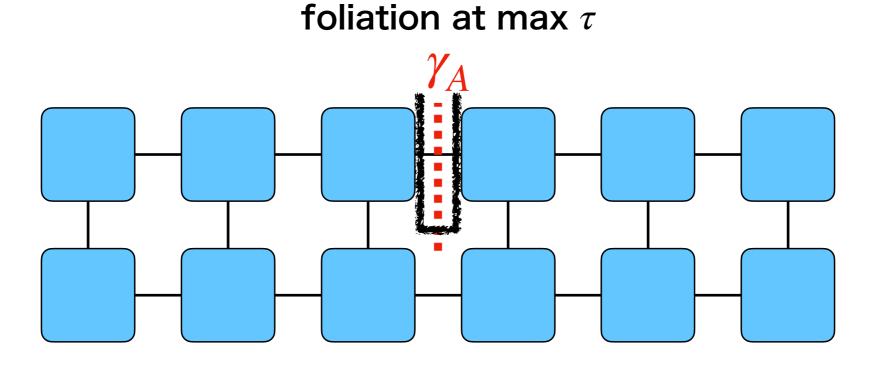
foliation at finite τ



- 1. Push the boundaries for each A and \overline{A} towards the RT (min bond cut) surface (cf. surface/state correspondence [Miyaji-Takayanagi])
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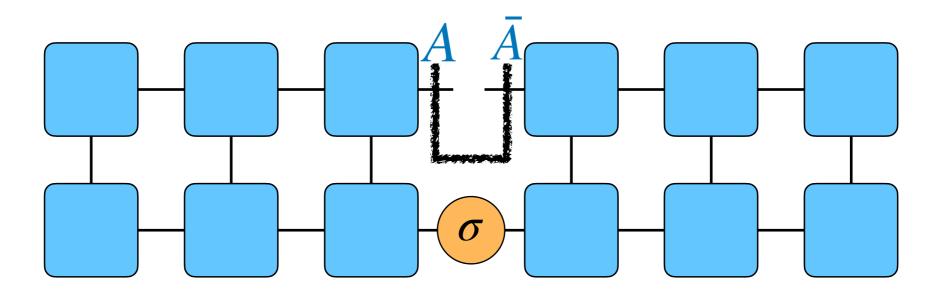


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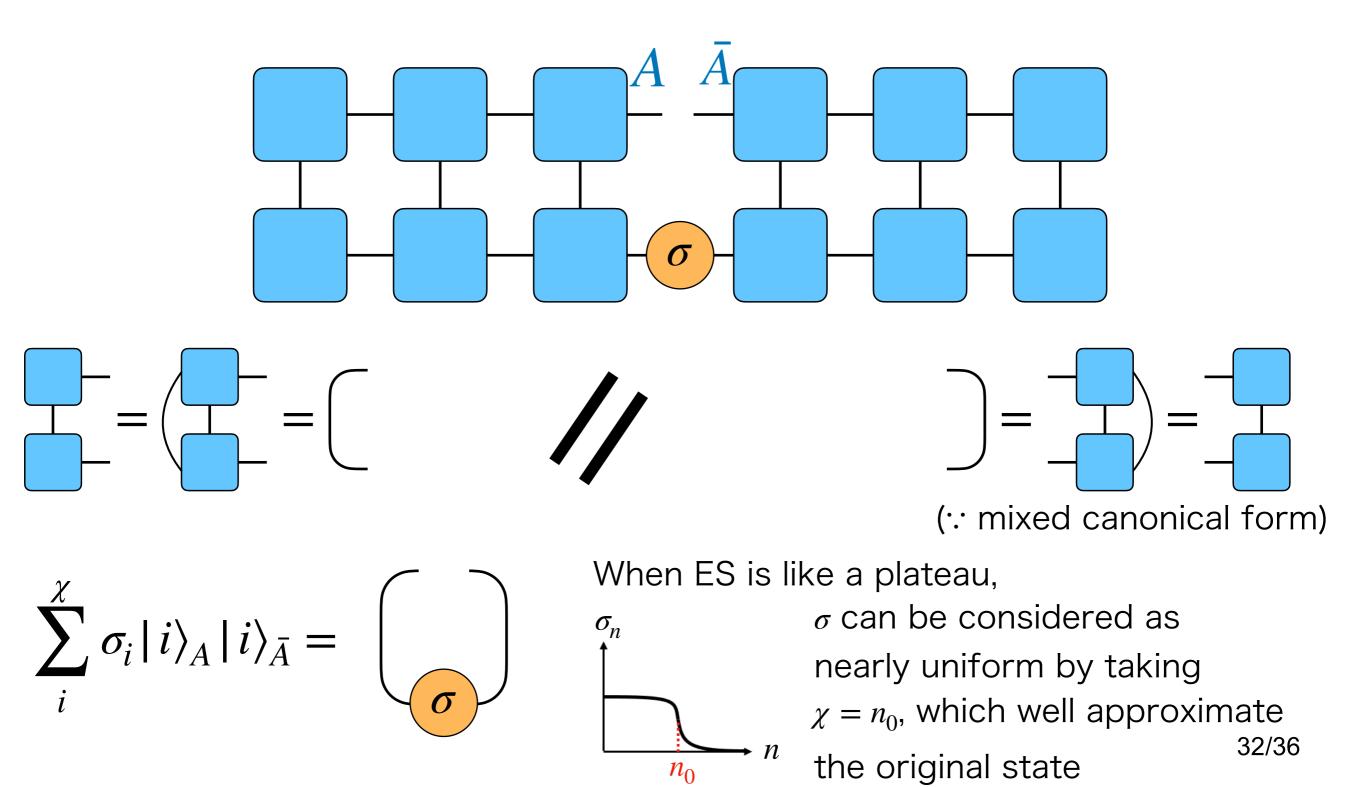
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foliation at max τ

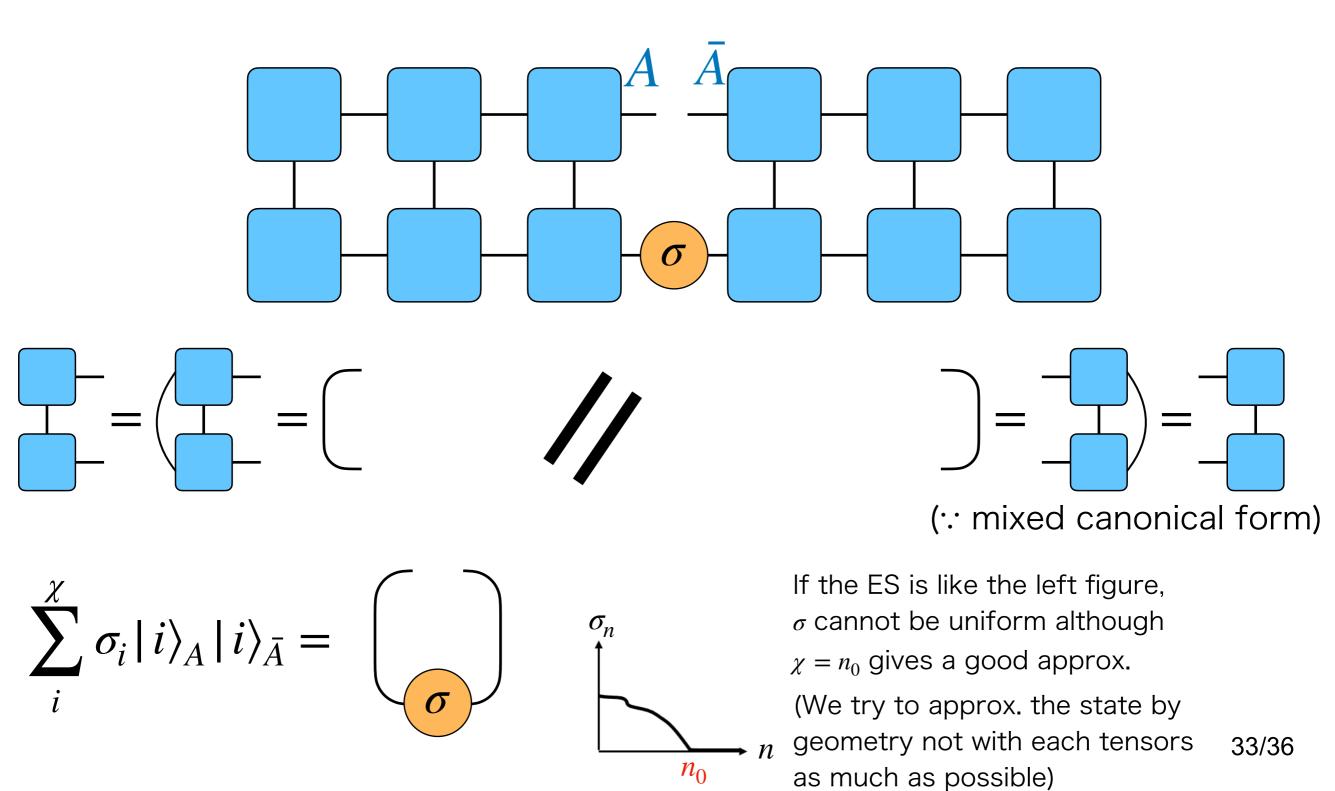


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- 2. Define a new state on a pushed boundary ("foliation") by removing the unreached part of TN and taking an inner product with the original TN
- 3. To retain the amount of entanglement, remove singular value matrix σ_A at γ_A

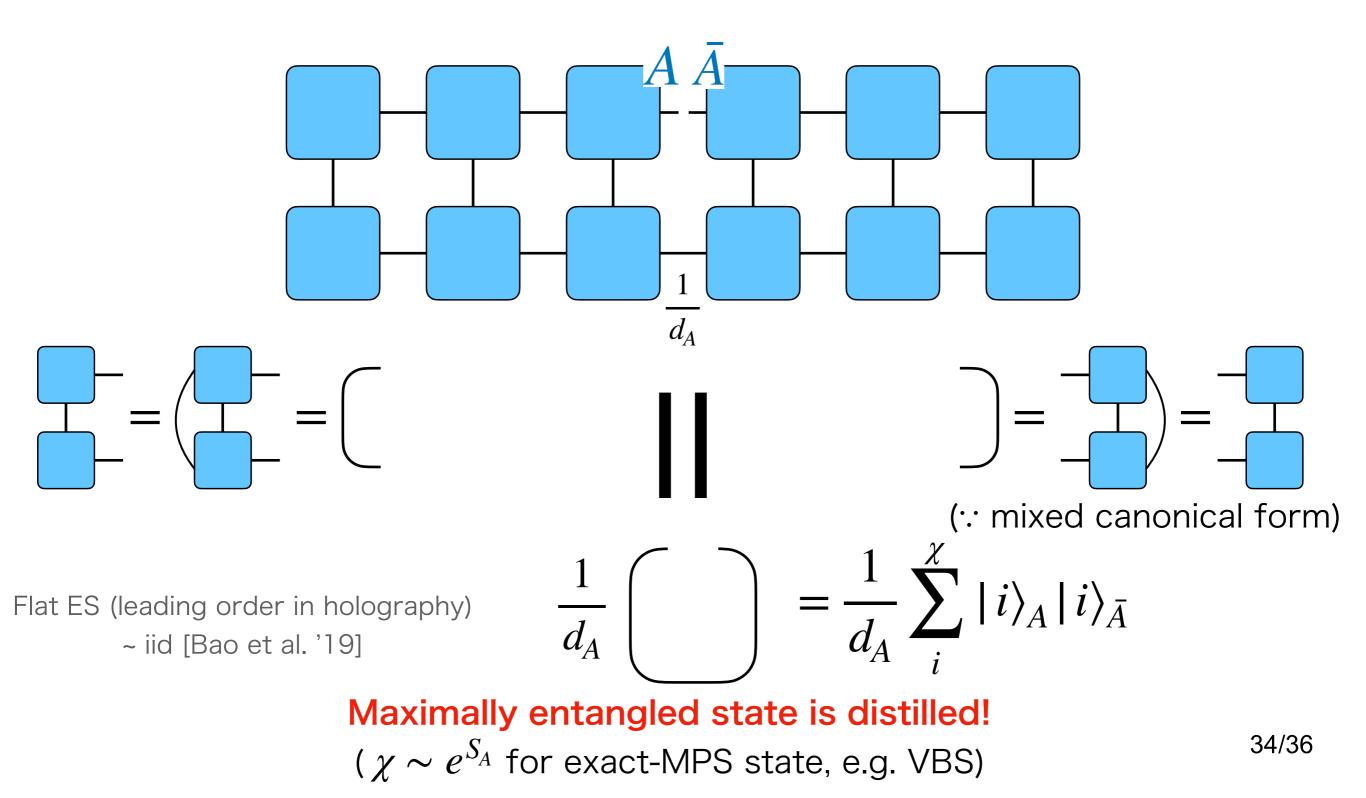
foliation at max τ

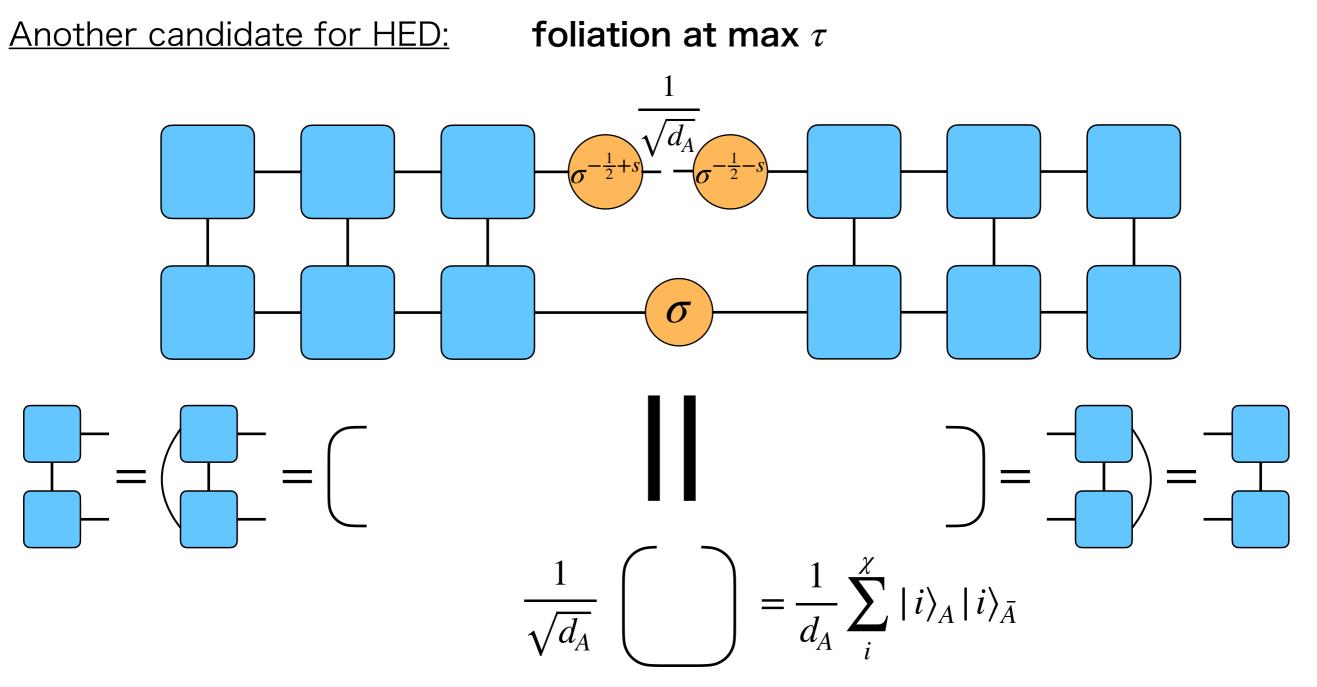


foliation at max τ



foliation at max τ





 \checkmark This is nothing but a distillation by bulk modular flow

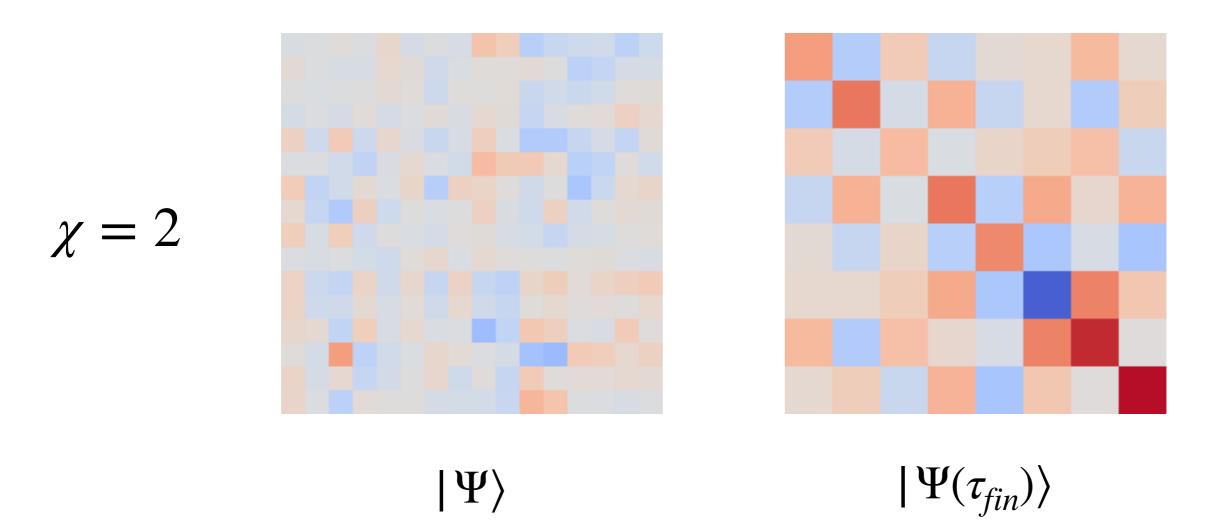
 \checkmark The amount of entanglement is not preserved; it maximizes EE

✓ This distillation is consistent with iid limit (inverse needs ∞-ly many copies) 35/36 [Quintino, et al. '19][Yoshida, et al. '21]

Example 2: Random MERA

• The distribution of the real part of coefficients (wave function) in $|\Psi\rangle$ (original state) and $|\Psi(\tau_{fin})\rangle$ (distilled one) for \mathscr{H}_A and $\mathscr{H}_{\bar{A}}$:

(larger: red, lower: blue, almost zero: white)

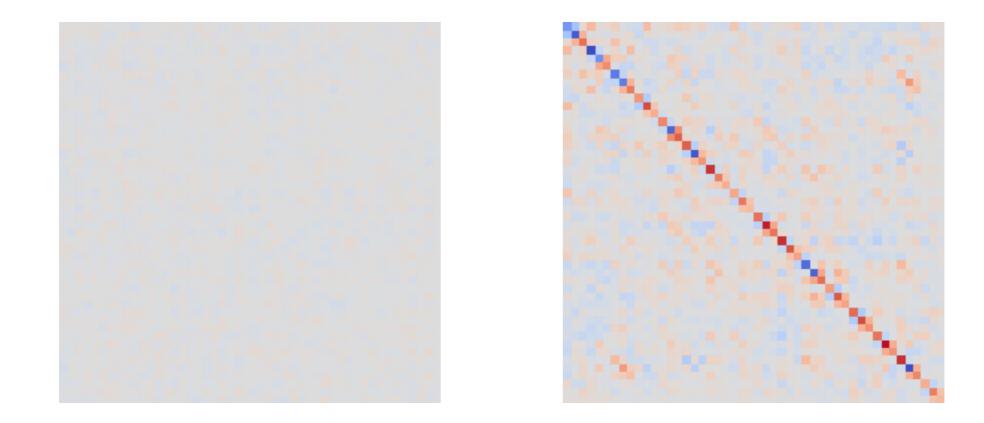


(Images are coarse-grained by 48×48 for visualization)

• The distribution of the real part of coefficients (wave function) in $|\Psi\rangle$ (original state) and $|\Psi(\tau_{fin})\rangle$ (distilled one) for \mathscr{H}_A and $\mathscr{H}_{\bar{A}}$:

(larger: red, lower: blue, almost zero: white)

 $\chi = 4$

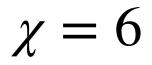


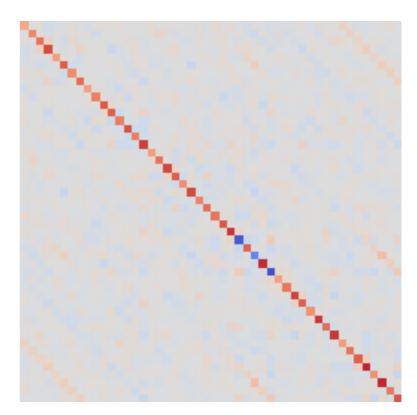
 $|\Psi
angle$ (too scattered)

 $|\Psi(\tau_{fin})\rangle$

• The distribution of the real part of coefficients (wave function) in $|\Psi\rangle$ (original state) and $|\Psi(\tau_{fin})\rangle$ (distilled one) for \mathscr{H}_A and $\mathscr{H}_{\bar{A}}$:

(larger: red, lower: blue, almost zero: white)



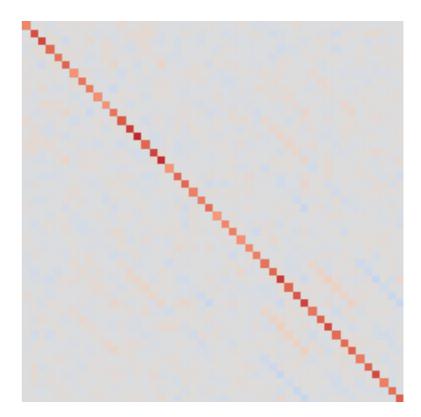


 $|\Psi(\tau_{fin})\rangle$

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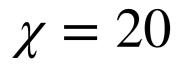
$$\chi = 8$$

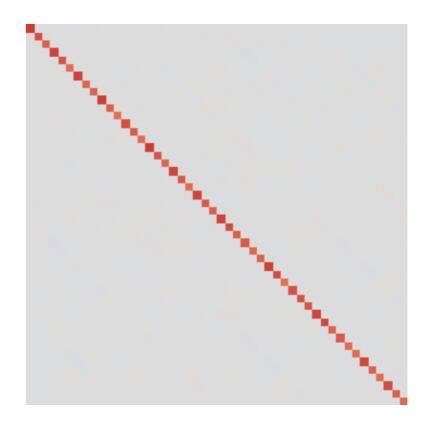


 $|\Psi(\tau_{fin})\rangle$

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(larger: red, lower: blue, almost zero: white)





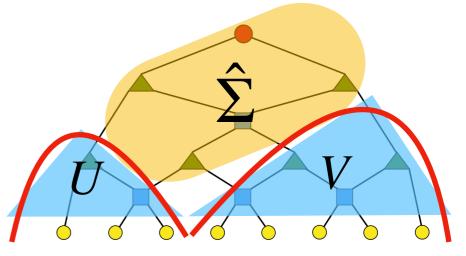
 $|\Psi(\tau_{fin})\rangle$

• What's important here: RT surface (min bond cut surface) in TN is NOT unique

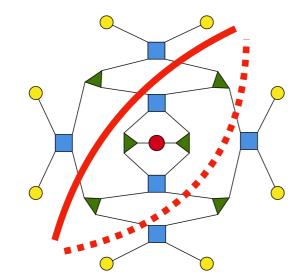
Why does this happen and can we see this in the AdS/CFT?

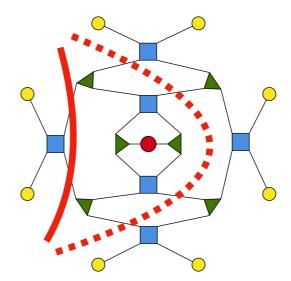
Identifying this phenomenon must be extremely important

as $\hat{\Sigma}$ includes all the information about singular values

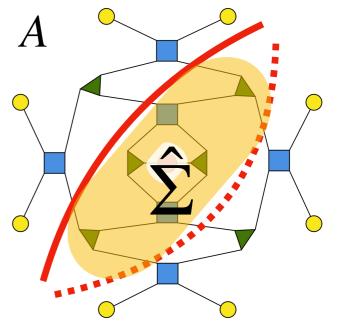


- ✓ The enclosed region by each boundary and each RT surface is isometric; beyond that (even in EW) is NOT isometric (in the direction from one boundary to the other)
- $\checkmark~\hat{\Sigma}$ region necessarily contains the top tensor
- ✓ By taking the infinite volume and continuum limit, two RT surfaces should get closer and each slope at boundary should becomes orthogonal; But the each RT surface is bounded by the location of the top tensor
- \checkmark By smoothly changing the subregion, other possible candidates fail to give minimum

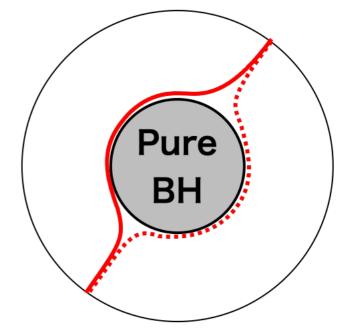


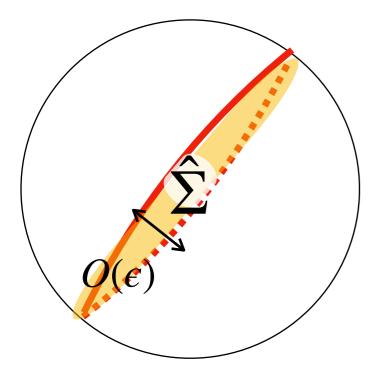


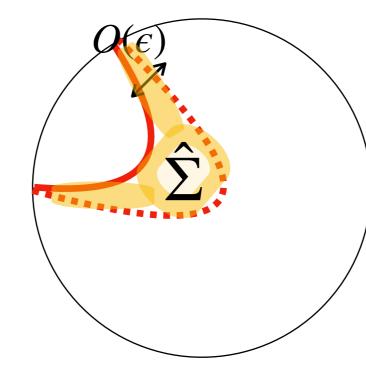
- This picture quite resembles to the pure BH. An RT surface can continuously move from one side to the other without any obstructions (because the state is pure)
- TN suggests Σ̂ region can exists at the origin of the bulk even for pure AdS as a point-like region.

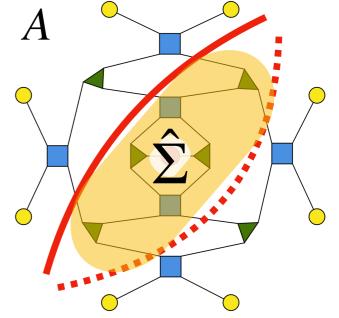


Dashed: causal cone for A









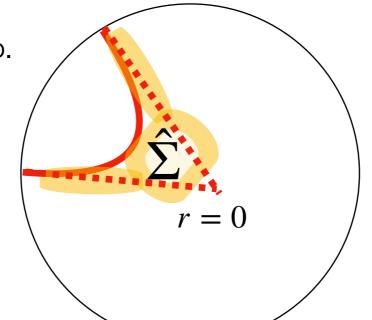
Dashed: causal cone for A

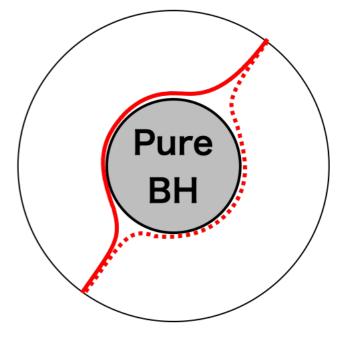
Python's lunch (region sandwiched by extremal surfaces) or entanglement shadow appear due to discretization

This is very close to EWCS=EoP setup. The important differences are:

 $\begin{array}{l} \cdot \mbox{ This exists even in the pure state} \\ \cdot \mbox{ If you take sufficiently small} \\ \mbox{ subregion, the RT surface is get} \\ \mbox{ disconnected for minimization} \\ \mbox{ condition whereas } \hat{\Sigma} \mbox{ remains finite} \end{array}$







Entanglement distillation (concentration)

Suppose we want to extract maximally entangled pairs from the following state: $|\Psi\rangle_{AB}^{\otimes n} = [\cos\theta|00\rangle + \sin\theta|11\rangle]_{AB}^{\otimes n}$ Expanding terms: $|00\cdots00\rangle, \cdots, |11\cdots11\rangle$ n+1 different coefficients. Regard them as n+1 different orthogonal states. Within each state, there are $\binom{n}{k}$ basis.

The probability to extract *k*-th density matrix (via projective measurement $|\phi_k\rangle = \sum_{i=1}^{\binom{n}{k}} |i\rangle |i'\rangle$): $p_k = \binom{n}{k} \cos^{2(n-k)} \theta \sin^{2k} \theta$

Then the averaged # of EPRs (ebits) are

$$\sum_{k=0}^{n} p_k \log \binom{n}{k} = \sum_{k=0}^{n} \exp \left[\log \binom{n}{k} + (n-k) \log \cos^2 \theta + k \log \sin^2 \theta \right] \log \binom{n}{k}$$

(Stirling's formula $\log n! \sim n \log n - n \Rightarrow \log \binom{n}{k} \sim n \log n - k \log k - (n-k) \log(n-k) = k \log \frac{n}{k} + (n-k) \log \frac{n}{n-k}$)

$$= \sum_{k=0}^{n} \exp\left[k \log\left(\frac{n}{k} \sin^2 \theta\right) + (n-k) \log\left(\frac{n}{n-k} \cos^2 \theta\right)\right] \log\binom{n}{k}$$

The saddle point approx. for the blue part: $k = n \sin^2 \theta$ ~ $-n \cos^2 \theta \log \cos^2 \theta - n \sin^2 \theta \log \sin^2 \theta = nS_A$

Entanglement distillation in holography

 In contrast, holography (or tensor network), we only have a single state. Then, why can one argue distillation?

First, the distillation or modular flow is not state-independent. (TTbar might offer a state-independent (but Hamiltonian-dependent) construction.)

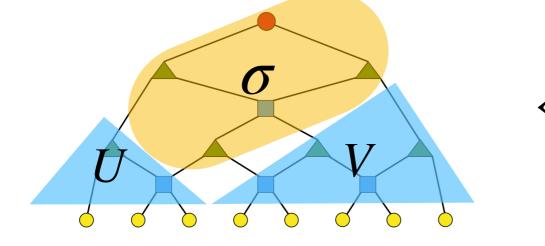
Second, it has been argued that a holographic theory has a flat entanglement spectrum, i.e. $\partial_n S_n = 0$, (to leading order). This is equivalent to preparing iid. [Bao, et al.]

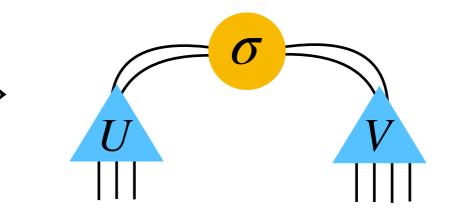
(In contrast, when we consider back reaction flat ES is false because gravitational Renyi entropy is affected by backreacting cosmic brane.)

Finally, holographic toy models like HaPPY code, random TN have flat ES [Dong-Harlow-Marolf]. Thus it is convincing that they offer a complete distillation. In general, we expect quasi randomness or quasi perfectness (~approx. QEC); such cases might result decrease in the success prob. for distillation.

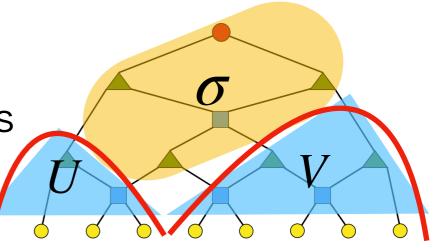
- So far this works for MPS, or more generally the state of the type $|\psi\rangle = (U_A \otimes V_{\bar{A}}) \sum \sigma_{\alpha} |\alpha\alpha\rangle_{A\bar{A}}$
- Q. Can we extend our analysis for MERA?
 A. At least we can make a guess!

• The domain of dependence (\neq entanglement wedge) for each region corresponds to U and V for MERA

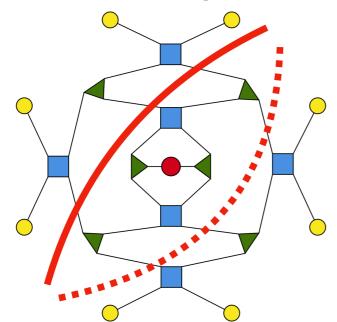


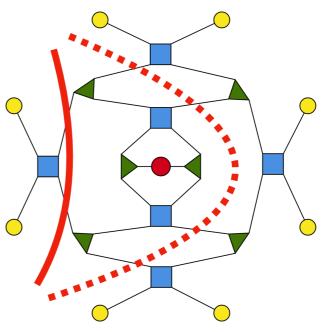


- The region σ can appear because we take $D(A), D(\overline{A})$, not EW
- RT surface is not unique in TN;
 Even within the pure state 2 RT surfaces seemingly does not match

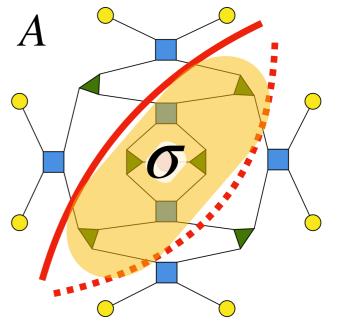


 But by smoothly changing the subregion, other possible candidates fail to give minimum

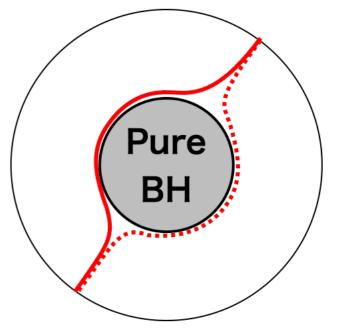




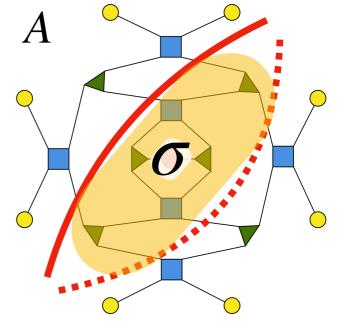
- This picture quite resembles to the pure BH. An RT surface can continuously move from one side to the other without any obstructions (because the state is pure)
- But now, the pure BH region is very imp; it accounts for *σ*. Beyond that region, the channel from the boundary is no more isometry (unless tensors have special properties like perfect, dual-unitary, etc.)
- TN suggests σ region can exists at the origin of the bulk even for pure AdS (but maybe just because of sub-AdS breakdown)



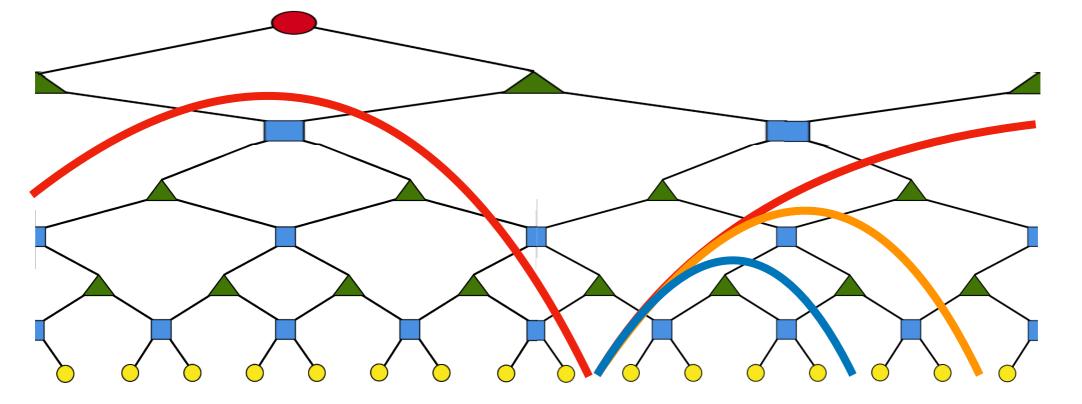
Dashed: causal cone for A



 This is special to TN. Since the angle between the minimal bond cut surface and the boundary near the endpoint of the subregion is always bounded by one unit of the isometry. It means the every RT surface sharing a common endpoint looks locally the same near the point.



Dashed: causal cone for A



Global coordinates?

Coordinate trf (for a parameter $\forall \alpha \in \mathbb{R}$: $r = 0 \Leftrightarrow z^2 - t^2 = \alpha^2, x = 0$):

$$\sqrt{1+r^{2}}\cos\tau = \frac{z^{2}+x^{2}-t^{2}+\alpha^{2}}{2\alpha z}; \quad \sqrt{1+r^{2}}\sin\tau = \frac{t}{z};$$
$$-r\cos\theta = \frac{z^{2}+x^{2}-t^{2}-\alpha^{2}}{2\alpha z}; \quad r\sin\theta = \frac{x}{z};$$

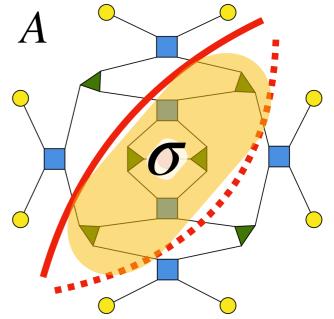
Geodesic in Poincare coordinates: $z^2 + (x - l_0)^2 = (l + l_0)^2$ where $A : [-l, l + 2l_0]$

 $\epsilon = 0$ makes the all the slopes of those geodesics same but at the same time the slope becomes orthogonal to the boundary. Furthermore, for a finite ϵ , the slopes are different from each other. This is different from TN.

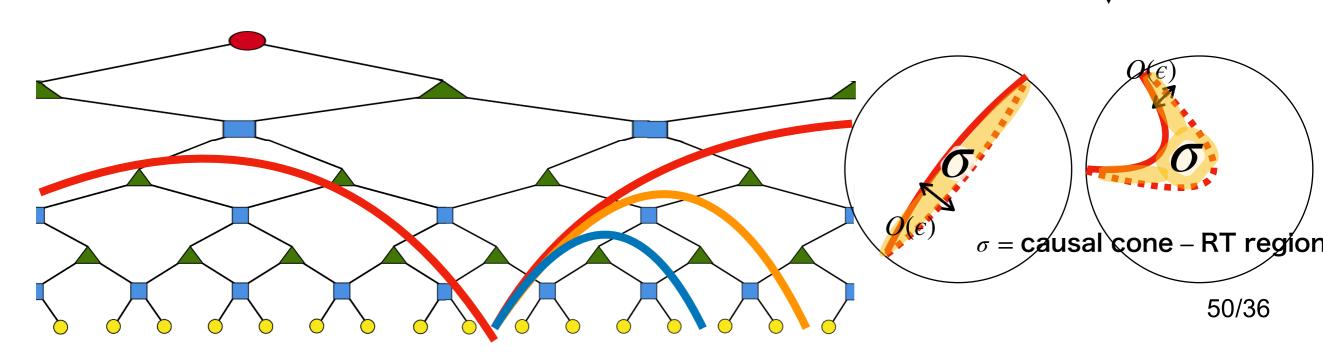
But isn't this natural from RG? Different lattice regularization leads different theory

up to correction vanishing as $\epsilon \to 0$.

 \rightarrow Then, the sigma region becomes thinner and thinner.



Dashed: causal cone for A



Continuum limit

Path integral approach to distillation in MERA

- As we have discussed, modular flow works for a certain circumstance. In such a case, the (open boundary) MERA can be thought as a state prepared by Euclidean path integral on a semi-disc.
- Then, the modular flow=HED
- Path integral optimization-> Liouville action

 $|0\rangle$

 $|0\rangle$

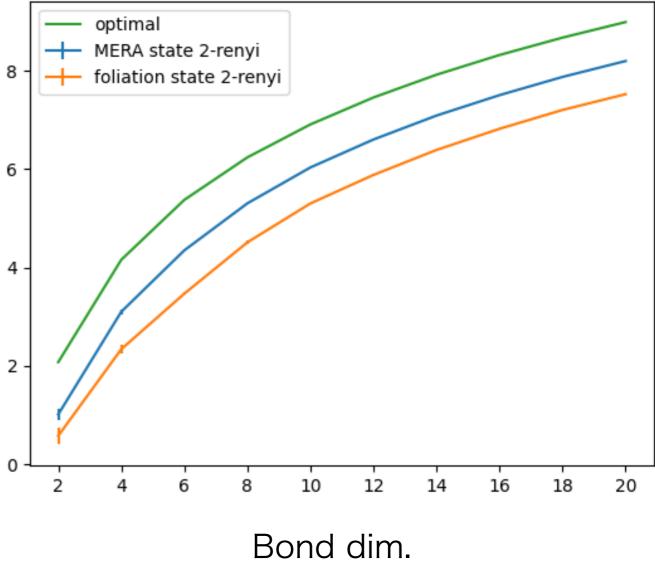
 $|0\rangle$

Random MERA: Renyi-2 entropy

- In general, Renyi-2 entropy of a TN state (blue) is less than the expected result from RT formula (green)
- In the large bond dim limit, they are expected to coincide
- The Renyi-2 entropy of holographically distilled state on γ_A (orange) is close to the original one (blue) probably due to doubled singular value

Renyi-2 entropy

$$S_2 = -\log \mathrm{Tr}\rho_A^2$$



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