# Operator Algebras, Non-Equilibrium Thermodynamics and Conformal Field Theory

#### Roberto Longo



#### Noncommutative Geometry: State of the Art and Future Prospects

#### Alain Connes' 70th birthday

Shanghai, March-April 2017

Based on a joint work with S. Hollands

and previous works with Bischoff, Kawahigashi, Rehren and Camassa, Tanimoto, Weiner

# Thermal equilibrium states

Thermodynamics concerns heat and temperature and their relation to energy and work. A primary role is played by the equilibrium distribution.

#### **Gibbs states**

Finite quantum system:  $\mathfrak{A}$  matrix algebra with Hamiltonian H and evolution  $\tau_t = \mathrm{Ad} e^{itH}$ . Equilibrium state  $\varphi$  at inverse temperature  $\beta$  is given by the Gibbs property

$$\varphi(X) = rac{\operatorname{Tr}(e^{-eta H}X)}{\operatorname{Tr}(e^{-eta H})}$$

What are the equilibrium states at infinite volume where there is no trace, no inner Hamiltonian?

# KMS states (HHW, Baton Rouge conference 1967)

Infinite volume.  $\mathfrak{A}$  a  $C^*$ -algebra,  $\tau$  a one-par. automorphism group of  $\mathfrak{A}$ . A state  $\varphi$  of  $\mathfrak{A}$  is KMS at inverse temperature  $\beta > 0$  if for  $X, Y \in \mathfrak{A} \exists F_{XY} \in A(S_{\beta})$  s.t.

(a)  $F_{XY}(t) = \varphi(X\tau_t(Y))$ (b)  $F_{XY}(t+i\beta) = \varphi(\tau_t(Y)X)$ 

where  $A(S_{\beta})$  is the algebra of functions analytic in the strip  $S_{\beta} = \{0 < \Im z < \beta\}$ , bounded and continuous on the closure  $\overline{S}_{\beta}$ .

(Note: it is sufficient to check (a) and (b) for X, Y in a dense \*-subalgebra  $\mathfrak{B}$ .)

KMS states have been so far the central objects in <u>Equilibrium</u> Quantum Statistical Mechanics, for example in the analysis of phase transition.

# Modular theory and Connes cocycles

Let  $\mathcal{M}$  be a von Neumann algebra and  $\varphi$  a normal faithful state on  $\mathcal{M}$ . The Tomita-Takesaki theorem gives a *canonical evolution*:

 $t \in \mathbb{R} \mapsto \sigma_t^{\varphi} \in \operatorname{Aut}(\mathcal{M})$ 

By a remarkable historical coincidence, Tomita announced the theorem at the 1967 Baton Rouge conference. Soon later Takesaki charcterised the modular group by the KMS condition.

The Connes Radon-Nikodym cocycle relates the modular groups of different states

$$u_t = (D\psi: D\varphi)_t \in \mathcal{M} \ , \quad \sigma_t^\psi = u_t \sigma_t^\varphi(\cdot) u_t^*$$

a first step towards the celebrated Connes classification of factors.

# Non-equilibrium thermodynamics

*Non-equilibrium thermodynamics*: study physical systems not in thermodynamic equilibrium but basically described by thermal equilibrium variables. Systems, in a sense, near equilibrium; but, in general, the system is <u>non-uniform</u> in space and time.

Non-equilibrium thermodynamics has been effectively studied for decades with important achievements, yet the general theory is still missing. The framework is even more incomplete in the quantum case, *non-equilibrium quantum statistical mechanics*.

We aim provide a general, model independent scheme for the above situation in the context of quantum, two dimensional *Conformal Quantum Field Theory*. As we shall see, we provide the general picture for the evolution towards a *non-equilibrium steady state*.



Figure: Non-equilibrium steady state

#### A typical frame described by Non-Equilibrium Thermodynamics:



Two infinite reservoirs  $R_1$ ,  $R_2$  in equilibrium at their own temperatures  $T_1 = \beta_1^{-1}$ ,  $T_2 = \beta_2^{-1}$ , and possibly chemical potentials  $\mu_1$ ,  $\mu_2$ , are set in contact, possibly inserting a probe.

As time evolves, the system should reach a non-equilibrium steady state.

This is the situation we want to analyse. As we shall see the *Operator Algebraic approach to CFT* provides a model independent description, in particular of the asymptotic steady state, and exact computation of the expectation values of the main physical quantities.

## Non-equilibrium steady states (Ruelle)

A non-equilibrium steady state NESS  $\varphi$  of  $\mathfrak{A}$  satisfies property (a) in the KMS condition, for all X, Y in a dense \*-subalgebra of  $\mathfrak{B}$ , but not necessarily property (b).

For any X, Y in  $\mathfrak{B}$  the function

 $F_{XY}(t) = \varphi \big( X \tau_t(Y) \big)$ 

is the boundary value of a function holomorphic in  $S_{\beta}$ .

Example: the tensor product of two KMS states at temperatures  $\beta_1$ ,  $\beta_2$  is a NESS with  $\beta = \min(\beta_1, \beta_2)$ .

Problem: describe the NESS state  $\omega$  and show that the initial state  $\psi$  evolves towards  $\omega$ 

$$\lim_{t \to +\infty} \psi \cdot \tau_t = \omega$$

Möbius covariant nets (Haag-Kastler nets on  $S^1$ ) A local Möbius covariant net A on  $S^1$  is a map

$$I \in \mathcal{I} 
ightarrow \mathcal{A}(I) \subset \mathcal{B}(\mathcal{H})$$

 $\mathcal{I} \equiv$  family of proper intervals of  $S^1$ , that satisfies:

- ▶ **A.** *Isotony.*  $I_1 \subset I_2 \implies \mathcal{A}(I_1) \subset \mathcal{A}(I_2)$
- ▶ **B.** Locality.  $I_1 \cap I_2 = \emptyset \implies [\mathcal{A}(I_1), \mathcal{A}(I_2)] = \{0\}$
- ► C. Möbius covariance. ∃ unitary rep. U of the Möbius group Möb on H such that

$$U(g)\mathcal{A}(I)U(g)^* = \mathcal{A}(gI), g \in \mathrm{M\"ob}, I \in \mathcal{I}.$$

- D. Positivity of the energy. Generator L<sub>0</sub> of rotation subgroup of U (conformal Hamiltonian) is positive.
- E. Existence of the vacuum. ∃! U-invariant vector Ω ∈ H (vacuum vector), and Ω is cyclic for V<sub>I∈T</sub> A(I).

## Consequences

- Irreducibility:  $\bigvee_{I \in \mathcal{I}} \mathcal{A}(I) = B(\mathcal{H}).$
- Reeh-Schlieder theorem: Ω is cyclic and separating for each A(I).
- Bisognano-Wichmann property (KMS property of ω|<sub>A(I)</sub>): The modular operator/conjugation Δ<sub>I</sub> and J<sub>I</sub> of (A(I), Ω) are

$$U(\delta_I(2\pi t)) = \Delta_I^{-it}, \ t \in \mathbb{R},$$
 dilations  
 $U(r_I) = J_I$  reflection

(Fröhlich-Gabbiani, Guido-L.)

- Haag duality:  $\mathcal{A}(I)' = \mathcal{A}(I')$
- ► Factoriality: A(I) is III<sub>1</sub>-factor (in Connes classification)

### Local conformal nets $Diff(S^1) \equiv group \text{ of orientation-preserving smooth diffeomorphisms of } S^1$

$$\operatorname{Diff}_{I}(S^{1}) \equiv \{g \in \operatorname{Diff}(S^{1}) : g(t) = t \,\,\forall t \in I'\}.$$

A local conformal net  $\mathcal{A}$  is a Möbius covariant net s.t.

**F.** Conformal covariance.  $\exists$  a projective unitary representation U of  $\text{Diff}(S^1)$  on  $\mathcal{H}$  extending the unitary representation of Möb s.t.

$$egin{aligned} U(g)\mathcal{A}(I)U(g)^* &= \mathcal{A}(gI), \quad g\in \mathrm{Diff}(S^1), \ U(g)xU(g)^* &= x, \quad x\in \mathcal{A}(I), \ g\in \mathrm{Diff}_{I'}(S^1), \end{aligned}$$

 $\longrightarrow$  unitary representation of the Virasoro algebra

$$[L_m, L_n] = (m - n)L_{m+n} + \frac{c}{12}(m^3 - m)\delta_{m,n}$$

 $\longrightarrow$  stress-energy tensor:

$$T(z) = \sum_{n \in \mathbb{Z}} L_n z^{-n-2}$$

### Representations

A (DHR) representation  $\rho$  of local conformal net  $\mathcal{A}$  on a Hilbert space  $\mathcal{H}$  is a map  $I \in \mathcal{I} \mapsto \rho_I$ , with  $\rho_I$  a normal rep. of  $\mathcal{A}(I)$  on  $\mathcal{H}$  s.t.

$$ho_{\widetilde{I}} \upharpoonright \mathcal{A}(I) = 
ho_{I}, \quad I \subset \widetilde{I}, \quad I, \widetilde{I} \subset \mathcal{I} \;.$$

Index-statistics relation (L.):

$$d(\rho) = \left[\rho_{I'}(\mathcal{A}(I'))':\rho_{I}(\mathcal{A}(I))\right]^{\frac{1}{2}}$$

DHR dimension =  $\sqrt{\text{Jones index}}$ 

# (Complete) rationality (Kawahigashi, Müger, L.)



The representations of A form a *modular tensor category*. (Feng Xu in the loop group case)

# Circle and real line picture



We shall frequently switch between the two pictures.

### KMS and Jones index

#### Kac-Wakimoto formula (conjecture)

Let  ${\mathcal A}$  be a conformal net,  $\rho$  representations of  ${\mathcal A},$  then

$$\lim_{t\to 0^+} \frac{\operatorname{Tr}(e^{-tL_{0,\rho}})}{\operatorname{Tr}(e^{-tL_0})} = d(\rho)$$

#### Analog of the Kac-Wakimoto formula (theorem)

 $\rho$  a representation of  $\mathcal{A}$ :

$$(\xi, e^{-2\pi K_{\rho}}\xi) = d(\rho)$$

where  $K_{\rho}$  is the generator of the dilations  $\delta_{I}$  and  $\xi$  is any vector cyclic for  $\rho(\mathcal{A}(I'))$  such that  $(\xi, \rho(\cdot)\xi)$  is the vacuum state on  $\mathcal{A}(I')$ .

## Basic conformal nets

#### U(1)-current net

In the real line picture  $\mathcal{A}$  is given by

 $\mathcal{A}(I) \equiv \{W(f) : f \in C^{\infty}_{\mathbb{R}}(\mathbb{R}), \text{ supp } f \subset I\}''$ 

where W is the representation of the Weyl commutation relations

$$W(f)W(g) = e^{-i\int fg'}W(f+g)$$

associated with the vacuum state  $\boldsymbol{\omega}$ 

$$\omega(W(f)) \equiv e^{-||f||^2}, \quad ||f||^2 \equiv \int_0^\infty p|\widetilde{f}(p)|^2 \mathrm{d}p$$

where  $\tilde{f}$  is the Fourier transform of f.

$$W(f) = \exp\left(-i\int f(x)j(x)dx\right), \quad [j(f),j(g)] = i\int fg'dx$$

with j(x) the U(1)-current.

There is a one parameter family  $\{\gamma_q, q \in \mathbb{R}\}$  of irreducible sectors and all have index 1 (Buchholz, Mack, Todorov)

$$\gamma_q(W(f)) \equiv e^{i\int Ff}W(f), \quad F\in C^\infty, \quad rac{1}{2\pi}\int F=q \; .$$

q is the called the charge of the sector.

The positive energy, irr. unitary representations of the Virasoro algebra are indexed by the central charge c and the lowest weight  $h \ge 0$ . They correspond to irr. reps of  $Diff(S^1)$ .

For every possible value of c, let  $U_c$  the irreducible rep. of Diff $(S^1)$  with h = 0.

 $Vir_c(I) \equiv U_c(Diff_I(S^1))''$ 

Universal property of the Virasoro net  $Vir_c$ : By diffeomorphism covariance,  $Vir_c$  is contained in every conformal net.

A classification of KMS states (Camassa, Tanimoto, Weiner, L.)

#### **Rational case**

Only one KMS state (w.r.t. translations)  $\beta = 2\pi$ exp: net on  $\mathbb{R} \ \mathcal{A} \rightarrow$  restriction of  $\mathcal{A}$  to  $\mathbb{R}^+$ 

 $\exp \restriction \mathcal{A}(I) = \operatorname{Ad} U(\eta)$ 

 $\eta$  diffeomorphism,  $\eta{\restriction} l={\rm exponential}$ 

 $arphi_{ ext{geo}}$  geometric KMS state on  $\mathcal{A}(\mathbb{R}), \omega$  vacuum state on  $\mathcal{A}(\mathbb{R}^+)$ 

 $\varphi_{\text{geo}} = \omega \circ \exp$ 

Scaling with dilation, we get the unique KMS state at any given  $\beta > 0$ .

## Non-rational case: U(1)-current model

The primary (locally normal) KMS states of the U(1)-current net are in one-to-one correspondence with real numbers  $q \in \mathbb{R}$ ;

Geometric KMS state:  $\varphi_{\text{geo}} = \varphi^{0}$ 

Any primary KMS state:

 $\varphi^{\boldsymbol{q}} = \varphi_{\text{geo}} \circ \gamma_{\boldsymbol{q}}.$ 

where

$$\gamma_q(W(f)) = e^{iq \int f(x)dx} W(f)$$

 $\gamma_q$  is equivalent to the BMT q-sector.

(With c < 1 there is only one KMS state: the net is rational)

Extremal KMS states of the Vir<sub>c</sub> net determined by the value on the stress-energy tensor T:

$$\varphi^{|q|}\left(T\left(f\right)\right) = \left(\frac{\pi}{12\beta^2} + \frac{q^2}{2}\right)\int f\,dx.$$

and the geometric KMS state corresponds to  $q = \frac{1}{\beta} \sqrt{\frac{\pi(c-1)}{6}}$  and energy density  $\frac{\pi c}{12\beta^2}$ .

# Chemical potential (cf. AHKT)

 $\mathcal{A}$  a local conformal net on  $\mathbb{R}$  and  $\varphi$  an extremal  $\beta$ -KMS state on  $\mathfrak{A} \equiv \overline{\bigcup_{I \subset \mathbb{R}} \mathcal{A}(I)}$  w.r.t. the time translation group  $\tau$  and  $\rho$  an irreducible DHR localised endomorphism of  $\mathfrak{A}$ . Assume that  $\rho$  is normal and  $d(\rho) < \infty$  (automatic in rational case).

U time translation unitary covariance cocycle in  $\mathfrak{A}$ :

$$\operatorname{Ad} U(t) \cdot \tau_t \cdot \rho = \rho \cdot \tau_t , \quad t \in \mathbb{R} ,$$

(unique by Möb covariance).

U is equal up to a phase to a Connes Radon-Nikodym cocycle:

$$U(t) = e^{-i2\pi\mu_{\rho}(\varphi)t} d(\rho)^{-i\beta^{-1}t} (D\varphi \cdot \Phi_{\rho} : D\varphi)_{-\beta^{-1}t}$$

 $\mu_{\rho}(\varphi) \in \mathbb{R}$  is the *chemical potential* of  $\varphi$  w.r.t. the charge  $\rho$ .

Here  $\Phi_{\rho}$  is the left inverse of  $\rho$ .

The geometric  $\beta$ -KMS state  $\varphi_0$  has zero chemical potential. By the holomorphic property of the Connes Radon-Nikodym cocycle:

 $e^{2\pi\beta\mu_
ho(arphi)} = {
m anal.\,cont.\,} arphiig(U(t)ig)ig/{
m anal.\,cont.\,} arphi_0ig(U(t)ig) \;.$ 

Example, BMT sectors:

With  $\varphi_{\beta,q}$  the  $\beta$ -state associated with the charge q, the chemical potential w.r.t. the charge q is given by

 $\mu_p(\varphi_{\beta,q}) = qp/\pi$ 

By linearity  $\mu_p$  is determined at p = 1, so we may put  $\mu(\varphi_{\beta,q}) = q/\pi$ .

# 2-dimensional CFT

 $M = \mathbb{R}^2$  Minkowski plane.

 $\begin{pmatrix} T_{00} & T_{10} \\ T_{01} & T_{11} \end{pmatrix}$  conserved and traceless stress-energy tensor.

As is well known,  $T_+ = \frac{1}{2}(T_{00} + T_{01})$  and  $T_- = \frac{1}{2}(T_{00} - T_{01})$  are chiral fields,

$$T_+ = T_+(t+x), \quad T_- = T_-(t-x).$$

Left and right movers.

 $\Psi_k$  family of conformal fields on *M*:  $T_{ij}$  + relatively local fields  $\mathcal{O} = I \times J$  double cone, *I*, *J* intervals of the chiral lines  $t \pm x = 0$ 

$$\mathcal{A}(\mathcal{O}) = \{e^{i\Psi_k(f)}, \mathrm{supp} f \subset \mathcal{O}\}''$$

then by relative locality

$$\mathcal{A}(\mathcal{O}) \supset \mathcal{A}_L(I) \otimes \mathcal{A}_R(J)$$

 $\mathcal{A}_L, \mathcal{A}_R$  chiral fields on  $t \pm x = 0$  generated by  $\mathcal{T}_L, \mathcal{T}_R$  and other chiral fields

Rational case:  $\mathcal{A}_L(I) \otimes \mathcal{A}_R(J) \subset \mathcal{A}(\mathcal{O})$  has finite Jones index.

Phase boundaries (Bischoff, Kawahigashi, Rehren, L.)

 $M_L \equiv \{(t,x): x < 0\}, M_R \equiv \{(t,x): x > 0\}$  left and right half Minkowski plane

A transpartent phase boundary is given by specifying two local conformal nets  $\mathcal{B}^L$  and  $\mathcal{B}^R$  on  $M_{L/R}$  on the same Hilbert space  $\mathcal{H}$ ;

 $M_L \supset O \mapsto \mathcal{B}^L(O) ; \qquad M_R \supset O \mapsto \mathcal{B}^R(O) ,$ 

 $\mathcal{B}^L$  and  $\mathcal{B}^R$  both contain a common Virasoro or larger chiral net  $\mathcal{A}$  and by causality:

 $\left[\mathcal{B}^{L}(O_{1}),\mathcal{B}^{R}(O_{2})
ight]=0, \quad O_{1}\subset M_{L}, \ O_{2}\subset M_{R}, \ O_{1}\subset O_{2}'$ 

i.e. a Connes bimodule with symmetries.

We consider the von Neumann algebras

$$\mathcal{D}(O) \equiv \mathcal{B}^L(O) \lor \mathcal{B}^R(O) \;, \quad O \in \mathcal{K} \;.$$

In the rational case,  $\mathcal{A}(O) \subset \mathcal{D}(O)$  has finite Jones index, so the center of  $\mathcal{D}(O)$  is finite dimensional; we may cut down by a minimal projection of the center (a defect) and assume  $\mathcal{D}(O)$  to be a factor.

Universal construction and classification (rational case) is done by considering the *braided product* of the *Q*-systems associated with  $\mathcal{A}_+ \otimes \mathcal{A}_- \subset \mathcal{B}_L$  and  $\mathcal{A}_+ \otimes \mathcal{A}_- \subset \mathcal{B}_R$ .

Cf. Fröhlich, Fuchs, Runkel, Schweigert (Euclidean setting)

## Non-equilubrium states in CFT (S. Hollands, R.L.)

Two local conformal nets  $\mathcal{B}^L$  and  $\mathcal{B}^R$  on the Minkowski plane M, both containing the same chiral net  $\mathcal{A} = \mathcal{A}_+ \otimes \mathcal{A}_-$ . For the moment  $\mathcal{B}^{L/R}$  is rational, so the KMS state is unique, later we deal wih chemical potentials.

**Before contact.** The two systems  $\mathcal{B}^L$  and  $\mathcal{B}^R$  are, separately, each in a thermal equilibrium state. KMS states  $\varphi_{\beta_{L/R}}^{L/R}$  on  $\mathfrak{B}^{L/R}$  at inverse temperature  $\beta_{L/R}$  w.r.t.  $\tau$ , possibly with  $\beta_L \neq \beta_R$ .  $\mathcal{B}^L$  and  $\mathcal{B}^R$  live independently in their own half plane  $M_L$  and  $M_R$ 

and their own Hilbert space. The composite system on  $M_L \cup M_R$  is given by

 $M_L \supset O \mapsto \mathcal{B}^L(O), \qquad M_R \supset O \mapsto \mathcal{B}^R(O)$ 

with  $C^*$ -algebra  $\mathfrak{B}^L(M_L)\otimes\mathfrak{B}^R(M_R)$  and state

$$\varphi = \varphi_{\beta_L}^L|_{\mathfrak{B}^L(M_L)} \otimes \varphi_{\beta_R}^R|_{\mathfrak{B}^R(M_R)};$$

 $\varphi$  is a stationary but not KMS.

#### Soon after contact.

At time t = 0 we put the two systems  $\mathcal{B}^L$  on  $M_L$  and  $\mathcal{B}^R$  on  $M_R$  in contact through a totally transmissible phase boundary. We are in the phase boundary case,  $\mathcal{B}^L$  and  $\mathcal{B}^R$  are now nets on M acting on a common Hilbert space  $\mathcal{H}$ ; the algebras  $\mathfrak{B}^L(W_L)$  and  $\mathfrak{B}^R(W_R)$  commute.

We want to describe the *initial state*  $\psi$  of the global system at time t = 0. As above, we set

$$\mathcal{D}(O) \equiv \mathcal{B}^{L}(O) \vee \mathcal{B}^{R}(O)$$

 $\psi$  should be a natural state on the global algebra  ${\mathfrak D}$  that satisfies

$$\psi|_{\mathfrak{B}^{L}(W_{L})} = \varphi_{\beta_{L}}^{L}|_{\mathfrak{B}^{L}(W_{L})}, \quad \psi|_{\mathfrak{B}^{R}(W_{R})} = \varphi_{\beta_{R}}^{R}|_{\mathfrak{B}^{R}(W_{R})}.$$

Since  $\mathfrak{B}^L(M_L)$  and  $\mathfrak{B}^R(M_R)$  are not independent, the existence of such state  $\psi$  is not obvious.

 $\exists$  a state  $\psi \equiv \psi_{\beta_L,\beta_R}$  on  $\mathfrak{D}$  s.t.  $\psi|_{\mathfrak{B}(W_{L/R})}$  is  $\varphi_{\beta_L/\beta_R}^{L/R}$ 

the initial state  $\psi$  is *normal, natural* on  $W_L$  and  $W_R$ , essentially arbitrary on the probe.

The state  $\psi$  is given by  $\psi \equiv \varphi \cdot \alpha_{\lambda_L,\lambda_R}$ , where  $\varphi$  is the geometric state on  $\mathfrak{D}$  (at inverse temperature 1) and  $\alpha = \alpha_{\lambda_L,\lambda_R}$  is the above doubly scaling automorphism with  $\lambda_L = \beta_L^{-1}$ ,  $\lambda_R = \beta_R^{-1}$  (local diffeomorphism construction)



Figure 1: Spacetime diagram of our setup. The initial state  $\psi$  is set up in the shaded region before the system is in causal contact with the phase boundaries. In the shaded regions to the left/right of the probe, we have a thermal equilibrium state at inverse temperatures  $\beta_L/\beta_R$ . In the diamond shaped shaded region O, we have an essentially arbitrary probe state.

**The large time limit.** After a large time we expect the global system to reach *a non equilibrium steady state*  $\omega$ .

The final state  $\omega$ : Let  $\varphi_{\beta_L}$ ,  $\varphi_{\beta_R}$  be the geometric KMS states respectively on  $\mathfrak{A}_+$  and  $\mathfrak{A}_-$  with inverse temperature  $\beta_L$  and  $\beta_R$ ; we define

 $\omega \equiv \varphi_{\beta_L} \otimes \varphi_{\beta_R} \cdot \varepsilon \; ,$ 

so  $\omega$  is the state on  $\mathfrak{D}$  obtained by extending  $\varphi_{\beta_L} \otimes \varphi_{\beta_R}$  from  $\mathfrak{A}$  to  $\mathfrak{D}$  by the conditional natural expectation  $\varepsilon : \mathfrak{D} \to \mathfrak{A}$ . Clearly  $\omega$  is a stationary state, indeed:

 $\omega$  is a NESS on  $\mathfrak{D}$  with  $\beta = \min\{\beta_L, \beta_R\}$ .

We now want to show that the evolution  $\psi \cdot \tau_t$  of the initial state  $\psi$  of the composite system approaches the non-equilibrium steady state  $\omega$  as  $t \to +\infty$ .

Note that:

$$\psi|_{\mathcal{D}(\mathcal{O})} = \omega|_{\mathcal{D}(\mathcal{O})} \text{ if } \mathcal{O} \in \mathcal{K}(V_+)$$

We have:

For every  $Z \in \mathfrak{D}$  we have:

$$\lim_{t\to+\infty}\psi(\tau_t(Z))=\omega(Z)\;.$$

Indeed, if  $Z \in \mathcal{D}(O)$  with O bounded and  $t > t_O$ , we have  $\tau_t(Z) \in \mathfrak{D}(V_+)$  as said, so

$$\psi( au_t(Z)) = \omega( au_t(Z)) = \omega(Z) , \quad t > t_O ,$$

because of the stationarity property of  $\omega$ .



Figure 2: Spacetime diagram of simplified setup. There is just one phase boundary and no probe. Every time-translated diamond will eventually enter the future lightcone  $V^+$ .

#### Case with chemical potential

In this case have chiral U(1)-currents  $J^{\pm}$  (non rational, c = 1 case).

Initial state  $\psi$ :

$$\psi|_{\mathfrak{B}^{L}(W_{L})} = \varphi_{\beta_{L},q_{L}}|_{\mathfrak{B}^{L}(W_{L})}, \qquad \psi|_{\mathfrak{B}^{R}(W_{R})} = \varphi_{\beta_{R},q_{R}}|_{\mathfrak{B}^{R}(W_{R})}.$$

Final NESS state  $\omega = \varphi_{\beta_L, q_L} \otimes \varphi_{\beta_R, q_R} \cdot \varepsilon$ 

$$\varphi_{\beta_L,q_L}(J^+(0)) = q_L , \qquad \varphi_{\beta_R,q_R}(J^-(0)) = q_R .$$

and for every  $Z \in \mathfrak{D}$  we have:

$$\lim_{t\to+\infty}\psi\bigl(\tau_t(Z)\bigr)=\omega(Z)\;.$$

We can explicitly compute the expected value of the asymptotic NESS state  $\omega$  on the stress energy tensor and on the current

In presence of chemical potentials  $\mu_{L/R} = \frac{1}{\pi}q_{L/R}$ , the large time limit of the two dimensional current density expectation value (x-component of the current operator  $J^{\mu}$ ) in the state  $\psi$  is, with  $J^{x}(t,x) = J^{-}(t+x) - J^{+}(t-x)$ 

 $\lim_{t\to+\infty}\psi(J^{\mathsf{x}}(t,x))=\varphi_{\beta_{L},q_{L}}^{-}(J^{-}(0))-\varphi_{\beta_{R},q_{R}}^{+}(J^{+}(0))=-\pi(\mu_{L}-\mu_{R}),$ 

whereas on the stress energy tensor

$$\lim_{t \to +\infty} \psi (T_{tx}(t,x)) = \varphi_{\beta_L,q_L}^+ (T^+(0)) - \varphi_{\beta_R,q_R}^- (T^-(0))$$
$$= \frac{\pi}{12} (\beta_L^{-2} - \beta_R^{-2}) + \frac{\pi^2}{2} (\mu_L^2 - \mu_R^2) ,$$

(cf. Bernard-Doyon)

Happy birthday again