COHERENT & DISSIPATIVE
TRANSPORT
in
APERIODIC MEDIA

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Main References


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I - Why Revisiting Transport?

- It is a very old problem
  - **Boltzmann** (1872-1880) for classical systems;
  - **Drude** (1900) for electrons.

- It is treated in textbooks: phenomenology, perturbation theory, numerical calculations.
I.1) Motivations

Conceptual Difficulties

1. No mathematically rigorous proof of the Kubo formulæ for transport coefficients.
   (However substantial progress for classical systems (Lebowitz’s school)
   and for quantum ones (Pillet-Jaksic, Fröhlich et al.).)

2. Low temperature effects are difficult to describe
   \textbf{ex.} Mott’s variable range hopping
   (see e.g. Efros & Schklovsky)

3. Aperiodic materials escape Bloch theory: need for
   a more systematic treatment
   \textbf{(ex.} quasicrystals).

4. Aperiodic media exhibit anomalous quantum diffusion
Transport is complex

- Thermodynamic quantities are much easier to measure: experiments are cleaner, easier to control. **Ex.** : heat capacity, magnetic susceptibility, structure factors... . But they do not separate various mechanisms.

- Transport measurements are mostly indirect: harder to interpret (especially at low temperature). Too many mechanisms occur at once.
Few mechanisms

1. For metals, $\sigma(T)$ increases as temperature decreases

$$\sigma(T) \xrightarrow{T \downarrow 0} T^{-2}, \quad (\text{Fermi liquid theory}).$$

2. For a thermally activated process

$$\sigma(T) \xrightarrow{T \downarrow 0} e^{-\Delta/T} \quad (\text{If a gap holds at Fermi level}).$$

3. For weakly disordered systems

$$\sigma(T) \xrightarrow{T \downarrow 0} \sigma(0) > 0 \quad (\text{residual conductivity}).$$

4. For strongly disordered systems in 3D

$$\sigma(T) \xrightarrow{T \downarrow 0} e^{-(T_0/T)^{1/4}} \quad (\text{variable range hopping}).$$
I.2)- Mott’s variable range hopping

N. Mott, (1968).


- Strongly localized regime, dimension $d$
- Low electronic DOS, Low temperature
• Absorption-emission of a phonon of energy $\varepsilon$

$$\text{Prob} \propto e^{-\varepsilon/k_B T}$$

• Tunnelling probability at distance $r$

$$\text{Prob} \propto e^{-r/\xi}$$

• Density of state at Fermi level $n_F$,

$$\varepsilon n_F r^d \approx 1$$

• Optimizing, the conductivity satisfies

$$\sigma \propto e^{-(T_0/T)^{1/d+1}} \quad \text{Mott’s law}$$

• Optimal energy $\varepsilon_{opt} \sim T^{d/(d+1)} \gg T$

• Optimal distance $r_{opt} \sim 1/T^{1/(d+1)} \gg \xi$
I.3)- Transport in Quasicrystals

Lectures on Quasicrystals,  
S. Roche, D. Mayou and G. Trambly de Laissardièrè,  

Quasicrystalline alloys:

Metastable QC’s:  
AlMn (Shechtman D., Blech I., Gratias D. & Cahn J., PRL 53, 1951 (1984))  
AlMnSi  
AlMgT ($T = Ag, Cu, Zn$)

Defective stable QC’s:  
AlLiCu (Sainfort-Dubost, (1986))  
GaMgZn (Holzen et al., (1989))

High quality QC’s:  
AlCuT ($T = Fe, Ru, Os$)  
( Hiraga, Zhang, Hirakoyashi, Inoue, (1988); Gurnan et al., Inoue et al., (1989); Y. Calvayrac et al., (1990) )

“Perfect” QC’s:  
AlPdMn  
AlPdRe
Resistivity ($\mu\Omega$cm)

- **Semiconductors**
- **Doped semiconductors**
- **Stable perfect quasicrystals** \( (AlPdRe) \)
- **High quality quasicrystals** \( (AlPdMn, AlCuFe, AlCuRu) \)
- **Defective stable quasicrystals** \( (AlCuLi, GaMgZn) \)
- **Metastable quasicrystals** \( (AlMn, AlMgZn,...) \)
- **Amorphous metals** \( (CuZr,...) \)
- **Metallic crystals** \( (Al,...) \)

**Typical values of the resistivity**

\[ \rho_{Mott} \]

\[ \text{Taken from C. Berger in ref. Lectures on Quasicrystals} \]
Conductivity of Quasicrystals vs Temperature

\[ \sigma \approx \sigma_0 + a T^\gamma \text{ with } 1 < \gamma < 1.5 \]

for \(.01 \text{ } K \leq T \leq 1000 \text{ } K\)
For Quasicrystals

1. $Al, Fe, Cu, Pd$ are very good metals: why is the conductivity of quasicrystalline alloys so low? Why is it decreasing?

2. At high enough temperature

\[ \sigma \propto T^\gamma \quad 1 < \gamma < 1.5 \]

There is a new mechanism here!

3. At low temperature for $Al_{70.5}Pd_{22}Mn_{7.5}$,

\[ \sigma \approx \sigma(0) > 0 \]

4. At low temperature for $Al_{70.5}Pd_{21}Re_{8.5}$,

\[ \sigma \propto e^{-(T_0/T)^{1/4}} \quad C. \text{Berger et al. (1998)} \]

Is disorder playing any role at very low temperature?
II - Coherent Transport

- Transport before collisions destroys quantum coherence.
- In aperiodic solids this transport can be anomalous.
II.1)- Mathematical Framework

1. Closing suitably the set of translated of the set of atomic positions leads to the Hull: it is a compact metrizable space $\Omega$ endowed with an $\mathbb{R}^d$-action.

2. An invariant ergodic probability measure $\mathbb{P}$ is provided by the Gibbs state at zero temperature.

3. Observables are random operators $A = (A_\omega)_{\omega \in \Omega}$ acting on the Hilbert space $\mathcal{H}$ of quantum states (such as $L^2(\mathbb{R}^d)$ for spinless electrons) with:

   (a) Covariance: $T(a) A_\omega T(a)^{-1} = A_{\tau - a\omega}$.

   (b) $\omega \mapsto A_\omega$ is strongly continuous.

4. The trace per unit volume, defined by $\mathbb{P}$, exists:

   $$ \mathcal{T}_\mathbb{P}(A) = \lim_{\Lambda \uparrow \mathbb{R}^d} \frac{1}{|\Lambda|} \text{Tr}(A_\omega \upharpoonright \Lambda) = \int_{\Omega} d\mathbb{P}(\omega) \langle x | A_\omega | x \rangle $$

5. Differential: $(\vec{\nabla} A)_\omega = -\nu [\vec{X}, A_\omega]$
II.2)- Local Exponents

Given a positive measure $\mu$ on $\mathbb{R}$:

$$\alpha_{\mu}^{\pm}(E) = \lim \left\{ \sup_{\varepsilon \downarrow 0} \inf \left\{ \frac{\ln \int_{E-\varepsilon}^{E+\varepsilon} d\mu}{\ln \varepsilon} \right\} \right\}$$

For $\Delta$ a Borel subset of $\mathbb{R}$:

$$\alpha_{\mu}^{\pm}(\Delta) = \mu-\text{ess sup}_{E \in \Delta} \left\{ \sup_{E \in \Delta} \alpha_{\mu}^{\pm}(E) \right\}$$

1. For all $E$, $\alpha_{\mu}^{\pm}(E) \geq 0$.
   $\alpha_{\mu}^{\pm}(E) \leq 1$ for $\mu$-almost all $E$.

2. If $\mu$ is $ac$ on $\Delta$ then $\alpha_{\mu}^{\pm}(\Delta) = 1$,
   if $\mu$ is $pp$ on $\Delta$ then $\alpha_{\mu}^{\pm}(\Delta) = 0$.

3. If $\mu$ and $\nu$ are equivalent measures on $\Delta$, then
   $\alpha_{\mu}^{\pm}(E) = \alpha_{\nu}^{\pm}(E)$ $\mu$-almost surely.

4. $\alpha_{\mu}^{+}$ coincides with the packing dimension.
   $\alpha_{\mu}^{-}$ coincides with the Hausdorff dimension.
II.3)- Fractal Exponents

For $p \in \mathbb{R}$:

$$D_{\mu, \Delta}^{\pm}(q) = \lim_{q' \to q} \frac{1}{q' - 1} \lim_{\varepsilon \downarrow 0} \sup_{\inf} \left\{ \ln \left( \frac{\int_{\Delta} d\mu(E) \left\{ \int_{E + \varepsilon}^{E - \varepsilon} d\mu \right\}^{q'-1}}{\ln \varepsilon} \right) \right\}$$

1. $D_{\mu, \Delta}^{\pm}(q)$ is a non decreasing function of $q$.

2. $D_{\mu, \Delta}^{\pm}(q)$ is not an invariant of the measure class, in general.

3.(a) If $\mu$ is ac on $\Delta$ then $D_{\mu, \Delta}^{\pm}(q) = 1$.

(b) If $\mu$ is pp on $\Delta$ then $D_{\mu, \Delta}^{\pm}(q) = 0$. 
II.4)- Spectral Exponents

Given a Hamiltonian $H = (H_\omega)_{\omega \in \Omega}$, namely a selfadjoint observable, we define:

1. The *local density of state* (LDOS) is the spectral measure of $H_\omega$ relative to a vector $\varphi \in \mathcal{H}$.
2. The corresponding local exponent is obtained after maximizing (+) or minimizing (-) over $\varphi$. It is denoted $\alpha_{\text{LDOS}}^\pm$. It is $\mathbb{P} - a. s.$ independent of $\omega$.
3. The *density of states* (DOS) as the measure defined by

$$\int d\mathcal{N}_\mathbb{P}(E)f(E) = \mathcal{T}_\mathbb{P}(f(H))$$

4. The local exponent associated with the DOS is denoted by $\alpha_{\text{DOS}}^\pm$.
5. Inequality : $\alpha_{\text{LDOS}}^\pm(\Delta) \leq \alpha_{\text{DOS}}^\pm(\Delta)$.
6. The fractal exponents for the LDOS are defined in the same way, provided we consider the average over $\omega$ before taking the logarithm and the limit $\varepsilon \downarrow 0$. 


II.5)- Transport Exponents

1. For $\Delta \subset \mathbb{R}$ Borel, let $P_{\Delta, \omega}$ be the corresponding spectral projection of $H_\omega$. Set:

$$\vec{X}_\omega(t) = e^{itH_\omega} \vec{X} e^{-itH_\omega}$$

2. The averaged spread of a typical wave packet with energy in $\Delta$ is measured by:

$$L^{(p)}_{\Delta}(t) = \left( \int_0^t ds \int_\Omega d\mathbb{P} \langle x | P_{\Delta, \omega} | \vec{X}_\omega(t) - \vec{X}^p P_{\Delta, \omega} | x \rangle \right)^{1/p}$$

3. Define $\beta = \beta_p^+(\Delta)$ similarly so that $L^{(p)}_{\Delta}(t) \sim t^\beta$.

4. $\beta_p^-(\Delta) \leq \beta_p^+(\Delta)$.

$\beta_p^+(\Delta)$ are non decreasing in $p$.

5. **Heuristic**

- $\beta = 0 \rightarrow$ absence of diffusion \textit{(ex: localization)}.
- $\beta = 1 \rightarrow$ ballistic motion \textit{(ex: in crystals)}.
- $\beta = 1/2 \rightarrow$ quantum diffusion \textit{(ex: weak localization)}.
- $\beta < 1 \rightarrow$ subballistic regime,
- $\beta < 1/2 \rightarrow$ subdiffusive regime \textit{(ex: in quasicrystals)}. 
II.6)- Inequalities

1. *Guarneri’s inequality:*  \( (Guarneri \, '89, \, Combes \, , \, Last \, '96)\)

\[
\beta_p^\pm(\Delta) \geq \frac{\alpha_{\text{LDOS}}^\pm(\Delta)}{d}
\]

2. *BGT inequalities:*  \( (Barbaroux, \, Germinet, \, Tcheremchantsev \, '00)\)

\[
\beta_p^\pm(\Delta) \geq \frac{1}{d} \, D_{\text{LDOS}, \Delta}^\pm \left( \frac{d}{d + p} \right)
\]

3. *Heuristics:*

(a) *ac* spectrum implies \( \beta \geq 1/d. \)

(b) *ac* spectrum implies ballistic motion in \( d = 1 \)

(c) *ac* spectrum is compatible with quantum diffusion in \( d = 2. \) This is expected in weak localization regime.

(d) *ac* spectrum is compatible with subdiffusion for \( d \geq 3. \)
II.7)- Results in Models

1. For Jacobi matrices (1D chains), the position operator is defined by the spectral measure (orthogonal polynomials) \( \Rightarrow \) transport exponents should be defined through the spectral ones.

2. For Jacobi matrices of a Julia set, with \( \mu \) the \( \sigma \)-balanced measure (Barbaroux, Schulz-Baldes '99)

\[
\beta_p^+ \leq D_\mu(1 - p) \quad \text{for all} \quad 0 \leq p \leq 2
\]

3. If \( H_1, \cdots, H_d \) are Jacobi matrices, \( \eta_1, \cdots, \eta_d \) are positive numbers and if

\[
H(\eta) = \sum_{j=1}^{d} \eta_j \ 1 \otimes \cdots \otimes H_j \otimes \cdots \otimes 1
\]

then (Schulz-Baldes, Bellissard '00)
\[
\beta_p^+(H^{(\eta)}) = \max_j \beta_p^+(H_j)
\]
\[
\alpha_{\text{LDOS}}(H^{(\eta)}) = \min\{1, \sum_j \alpha_{\text{LDOS}}(H_j)\}
\]

for a.e. \( \eta \). In addition if \( \sum_j \alpha_{\text{LDOS}}(H_j) > 1 \), \( H^{(\eta)} \) has \textit{a.c. spectrum}.

4. For any \( \epsilon > 0 \), there is a Jacobi matrix \( H_0 \) such that if \( H_j = H_0, \ \forall j \), \( H^{(\eta)} \) has \textit{a.c. spectrum} for \( d \geq 3 \) and spectral exponent \( \leq 1/d - \epsilon \) for a.e. \( \eta \).
\[(\text{Schulz-Baldes, Bellissard '00})\]

5. There is a class of models of Jacobi matrices on an infinite dimensional hypercube with \textit{a.c. spectrum} and vanishing transport exponents.
\[(Vidal, Mosseri, Bellissard '99)\]
III - Dissipative Transport

- Transport beyond the collision time.
- Several mechanisms of dissipation may be considered: *electron-electron* collisions, interactions with *acoustic* phonons or *optical* ones. *etc..*
III.1)- The Drude Model

Assumptions:

1. Electrons in a metal are free classical particles of mass $m_*$ and charge $q$.

2. They experience collisions at random Poissonnnian times $\cdots < t_n < t_{n+1} < \cdots$, with average relaxation time $\tau_{rel}$.

3. If $p_n$ is the electron momentum between times $t_n$ and $t_{n+1}$, then the $p_{n+1}$’s is updated according to the Maxwell distribution at temperature $T$.

Then the conductivity follows the Drude formula

$$\sigma = \frac{q^2n}{m_* \tau_{rel}}$$
The Drude Kinetic Model
III.2)- The Kubo Formula

(Bellissard, Schulz-Baldes ’95)

1. Replace the classical dynamics by the quantum one electron dynamic in the aperiodic solid.

2. At each collision, force the density matrix to come back to equilibrium. (Relaxation time Approximation or RTA).

3. There is then one relaxation time $\tau_{rel}$. The electric conductivity is then given by Kubo’s formula:

$$
\sigma_{i,j} = \frac{q^2}{\hbar} \mathcal{T}_p \left( \partial_j \left( \frac{1}{1 + e^{\beta(H-\mu)}} \right) \frac{1}{1/\tau_{rel} - \mathcal{L}_H \partial_i H} \right)
$$

Here $q$ is the charge of the carriers, $\beta = 1/k_B T$, $\mu$ is the chemical potential and $\mathcal{L}_H = \imath/\hbar [H, .]$.

4. For the Hilbert-Schmidt inner product defined by $\mathcal{T}_p$, $\mathcal{L}_H$ is anti-selfadjoint. Thus as $\tau_{rel} \uparrow \infty$, the resolvent of $\mathcal{L}_H$ is evaluated closer to the spectrum near 0.
III.3) The Anomalous Drude Formula

(Mayou '92, Sire '93, Bellissard, Schulz-Baldes '95)

\[
\sigma \sim \tau_{rel}^{2\beta_F - 1} \quad \text{where } \beta_F \text{ is the transport exponent } \beta_2(E_F) \text{ evaluated at Fermi level.}
\]

1. In practice, \( \tau_{rel} \uparrow \infty \) as \( T \downarrow 0 \).

2. If \( \beta_F = 1 \) (ballistic motion), \( \sigma \sim \tau_{rel} \) (Drude). The system behaves as a conductor.

3. For \( 1/2 < \beta_F \leq 1 \), \( \sigma \uparrow \infty \) as \( T \downarrow 0 \): the system behaves as a conductor.

4. If \( \beta_F = 1/2 \) (quantum diffusion), \( \sigma \sim \text{const.} \): residual conductivity at low temperature.

5. For \( 0 \leq \beta_F < 1/2 \), \( \sigma \downarrow 0 \) as \( T \downarrow 0 \): the system behaves as an insulator.
III.4)- Time Scales

1. The phonon mediated electron-electron interaction gives \( \tau_{rel} \sim T^{-2} \) (Fermi liquid theory).

2. Electron interaction with acoustic phonons, gives \( \tau_{rel} \sim T^{-5} \) (Bloch theory).

3. Quantum chaos in the one electron spectrum makes the Hamiltonian looks like a random matrix like in weak localization regime (Bellissard, Magnen, Rivasseau, ‘02). This leads to a conductivity independent of the temperature.

4. Optical phonons are important in aperiodic system. They produce a band similar to the spectrum of a random matrix (quantum chaos). The influence of optical phonons on the relaxation time is unknown.
5. In the conducting regime, the *shortest* dissipative time scale dominates, favoring Fermi liquid theory:

\[ \sigma \sim T^{1/0} T^{-2(2\beta_F-1)} \quad \text{if } \beta_F > 1/2, \]

6. In the insulator regime, the *longest* dissipative time scale dominates. Thus Bloch theory is likely to dominate

\[ \sigma \sim T^{1/0} T^{5(1-2\beta_F)} \quad \text{if } \beta_F < 1/2, \]

7. If there is an infinite number of time scales the low temperature behaviour is not a scaling law: for example in the Mott variable range hopping regime.
Conductivity of Quasicrystals vs Temperature

\[ \sigma \approx \sigma_0 + a T^\gamma \text{ with } 1 < \gamma < 1.5 \]

for \(.01 \text{ } K \leq T \leq 1000 \text{ } K\)
III.5)- Conductivity in Quasicrystals


1. LMTO ab initio computations for $i-AlCuCo$ give $\beta_F \approx 0.375 < 1/2$. Thus Bloch’s law dominates giving

$$\sigma \approx T^{0} T^{5/4} \text{ if } \beta_F < 1/2,$$

compatible with experimental results!

2. If disorder dominates at low temperature
   (a) for $AlPdMn$, the occurrence of a residual conductivity implies weak localisation. Thus there should be a high density of defects or impurities
   (b) $AlPdRe$ behaves like an insulator with Mott’s variable range hopping conductivity. This is a sign for strong localization, implying a low density of defects or impurities.

This seems with these materials being structurally similar and samples being high quality.
III.6) - Quantum Chaos in Quasicrystals

J.B. & D.Mayou, speculations ’99


2. For a sample of size $L$ in dimension $d$:
   Mean level spacing $\Delta \sim L^{-d}$.
   Thus Heisenberg time $\tau_H \sim L^d$.

3. Time necessary to reach the boundary (Thouless) $L \sim \tau_Th^{\beta_F}$. Thus $\tau_Th^{\beta_F} \sim L^{1/\beta}$.

4. Hence:
   (a) if $\beta_F > 1/d$ level repulsion dominates implying
       - quantum diffusion $\langle x^2 \rangle \sim t$
       - residual conductivity
       - absolutely continuous spectrum at Fermi level;
   (b) if $\beta_F < 1/d$ level repulsion can be ignored and
       - anomalous diffusion dominates $\langle x^2 \rangle \sim t^{2/\beta_F}$
       - insulating behaviour with scaling law
       - singular continuous spectrum near Fermi level.
III.7)- Beyond the RTA

1. At low temperature, the RTA is invalid. There is a spectrum of relaxation times.

2. A kinetic model of quantum jumps has been proposed leading to the validity of linear response. (Spehner, Bellissard ’00, Bellissard, Rebolledo, Spehner, von Waldenfels ’00).

3. The current admits two parts: the coherent one, induced by $\vec{J} = i[\vec{X}, H]$, and a dissipative one including other effects like phonon drag, etc.

4. The Kubo formula becomes more involved and can be decomposed into five contributions in general.

5. Applied to strongly localized electrons, this formalism gives rise to a justification of the Abrahams and Miller random resistor network model,(Spehner, Thesis ’00, Spehner, Bellissard, ’00) describing the Mott variable range hopping.
IV)- Conclusions :

1. The electron dynamics in an aperiodic solid can be described by using random operators and rules of Non Commutative Calculus.

2. The quantum evolution of a typical wave packet leads to anomalous diffusion, described through various spectral and transport exponents.

3. These exponents are related by inequalities that allow subdiffusion together with absolutely continuous spectrum for $d \geq 3$.

4. Dissipative mechanisms, such as electron-phonon interaction, may be described through kinetic models, generalizing the Drude model.

5. The interplay between coherent and dissipative transport is revealed at low temperature. Anomalous diffusion then leads to an anomalous Drude formula within the RTA.

6. The anomalous Drude formula may explain the behaviour of quasicrystals.

7. Beyond the RTA, the kinetic models are still valid but involve more conditions. One consequence is the justification of the Abraham-Miller random resistor network which usually leads to a better understanding of the Mott variable range hopping conductivity, in strongly disordered systems.