Optimal Supply Networks I: Branching

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Optimal transportation

Optimal branching

Murray's law Murray meets Tokunaga







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Outline

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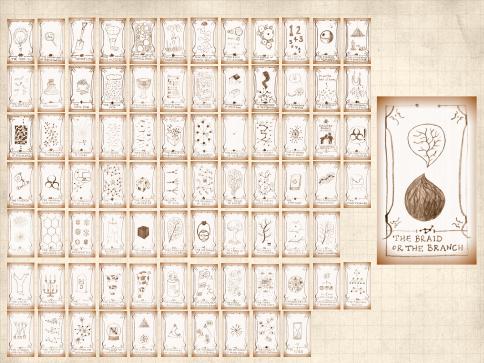
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Optimal

What's the best way to distribute stuff?

- 🙈 Stuff = medical services, energy, people, ...
- Some fundamental network problems:
 - 1. Distribute stuff from a single source to many sinks
 - 2. Distribute stuff from many sources to many sinks
 - 3. Redistribute stuff between nodes that are both sources and sinks
- Supply and Collection are equivalent problems





Basic question for distribution/supply networks:



How does flow behave given cost:

$$C = \sum_j I_j^{\,\gamma} Z_j$$

where

 I_i = current on link jand Z_i = link j's impedance.



 \Longrightarrow Example: $\gamma = 2$ for electrical networks.

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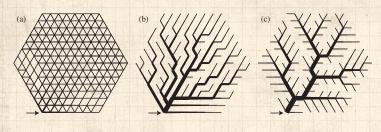
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(a) $\gamma > 1$: Braided (bulk) flow

(b) γ < 1: Local minimum: Branching flow

(c) $\gamma < 1$: Global minimum: Branching flow

Note: This is a single source supplying a region.

From Bohn and Magnasco [3] See also Banavar *et al.* [1]: "Topology of the Fittest Transportation Network"; focus is on presence or absence of loops—same story PoCS @pocsvox

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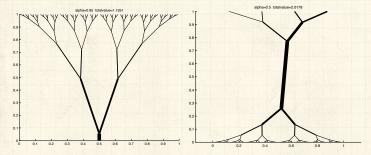
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Optimal paths related to transport (Monge) problems ::





"Optimal paths related to transport problems" 🗹

Qinglan Xia, Communications in Contemporary Mathematics, **5**, 251–279, 2003. [19] PoCS @pocsvox

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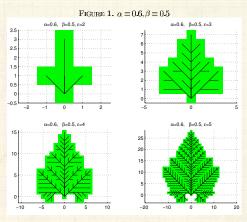
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Growing networks—two parameter model: [20]



- \ref{approx} Parameters control impedance ($0 \le \alpha < 1$) and angles of junctions ($0 < \beta$)
- \clubsuit For this example: $\alpha=0.6$ and $\beta=0.5$

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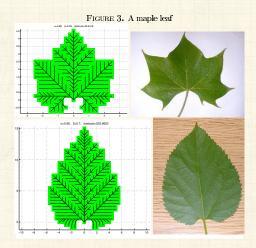
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Growing networks: [20]



 α Top: $\alpha = 0.66$, $\beta = 0.38$; Bottom: $\alpha = 0.66$, $\beta = 0.70$

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An immensely controversial issue ...

The form of natural branching networks: Random, optimal, or some combination? [6, 18, 2, 5, 4]

River networks, blood networks, trees, ...

Two observations:

Self-similar networks appear everywhere in nature for single source supply/single sink collection.

Real networks differ in details of scaling but reasonably agree in scaling relations.

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River network models

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Optimality:

Optimal channel networks [13]

Thermodynamic analogy [14]

versus ...

Randomness:

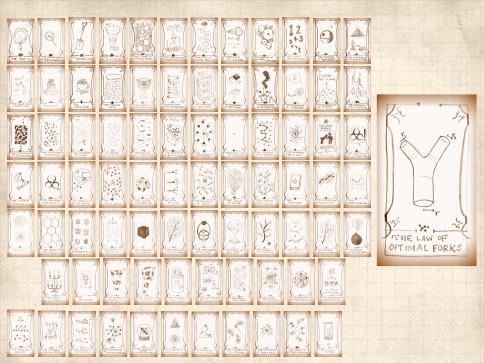
Scheidegger's directed random networks

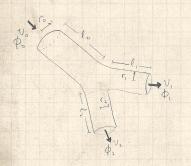
Undirected random networks





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Murray's law (1926) connects branch radii at forks: [11, 10, 12, 7, 16]

$$r_0^3 = r_1^3 + r_2^3$$

where r_0 = radius of main branch, and r_1 and r_2 are radii of sub-branches.

- Holds up well for outer branchings of blood networks.
- Also found to hold for trees [12, 8] when xylem is not a supporting structure [9].
- See D'Arcy Thompson's "On Growth and Form" for background and general inspiration [15, 16].

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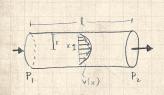




Use hydraulic equivalent of Ohm's law:

$$\Delta p = \Phi Z \Leftrightarrow V = IR$$

where Δp = pressure difference, Φ = flux.



Fluid mechanics: Poiseuille impedance for smooth Poiseuille flow in a tube of radius r and length ℓ :

$$Z = \frac{8\eta\ell}{\pi r^4}$$



 \Re η = dynamic viscosity \square (units: $ML^{-1}T^{-1}$).



Power required to overcome impedance:

$$P_{\mathsf{drag}} = \Phi \Delta p = \Phi^2 Z.$$



Also have rate of energy expenditure in maintaining blood given metabolic constant c:

$$P_{\rm metabolic} = c r^2 \ell$$

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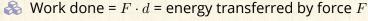
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Aside on P_{drag}



 $lap{R}$ Power = P = rate work is done = $F \cdot v$

 Δp = Pressure differential = Force per unit area

 Φ = Volume flow per unit time (current) = cross-sectional area · velocity

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Murray's law:

Total power (cost):

$$P = P_{\text{drag}} + P_{\text{metabolic}} = \Phi^2 \frac{8\eta \ell}{\pi r^4} + cr^2 \ell$$

- \red Observe power increases linearly with ℓ
- $\ensuremath{\&}$ But r's effect is nonlinear:
 - increasing r makes flow easier but increases metabolic cost (as r^2)
 - decreasing r decrease metabolic cost but impedance goes up (as r^{-4})

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Murray's law:

 \mathbb{A} Minimize P with respect to r:

$$\frac{\partial P}{\partial r} = \frac{\partial}{\partial r} \left(\Phi^2 \frac{8\eta \ell}{\pi r^4} + cr^2 \ell \right)$$

Flow rates at each branching have to add up (else) our organism is in serious trouble ...):

$$\Phi_0 = \Phi_1 + \Phi_2$$

where again 0 refers to the main branch and 1 and 2 refers to the offspring branches





Murray's law:



$$\Phi = kr^3$$

- A Insert question from assignment 16 🗹
- All of this means we have a groovy cube-law:

$$r_0^3 = r_1^3 + r_2^3$$

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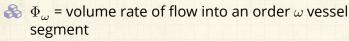
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Optimization

Murray meets Tokunaga:



Tokunaga picture:

$$\Phi_{\omega} = 2\Phi_{\omega-1} + \sum_{k=1}^{\omega-1} T_k \Phi_{\omega-k}$$

$$\left(r_{\omega}\right)^{3}=2\left(r_{\omega-1}\right)^{3}+\sum_{k=1}^{\omega-1}T_{k}\left(r_{\omega-k}\right)^{3}$$

Same form as:

$$n_{\omega} = \frac{2n_{\omega+1}}{\text{generation}} + \sum_{\omega'=\omega+1}^{M} \underbrace{T_{\omega'-\omega}n_{\omega'}}_{\text{absorption}}$$

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Murray meets Tokunaga:

- $\ref{eq:special}$ Find R_r^3 satisfies same equation as R_n and R_v (v is for volume):

$$R_r^3 = R_n = R_v$$

Is there more we could do here to constrain the Horton ratios and Tokunaga constants?





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Murray meets Tokunaga:

- $\red {\$}$ Isometry: $V_\omega \propto \ell_\omega^3$
- Gives

$$R_{\ell}^3 = R_r^3 = R_n = R_v$$

- & We need one more constraint ...
- West *et al.* (1997) [18] achieve similar results following Horton's laws (but this work is a disaster).
- So does Turcotte *et al.* (1998) [17] using Tokunaga (sort of).

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