

Optimal Supply Networks I: Branching

Last updated: 2023/08/22, 11:48:21 EDT

Principles of Complex Systems, Vols. 1, 2, & 3D
 CSYS/MATH 6701, 6713, & a pretend number,
 2023-2024 | @pocsvox

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Single source optimal supply

Basic question for distribution/supply networks:

How does flow behave given cost:

$$C = \sum_j I_j^\gamma Z_j$$

where

I_j = current on link j

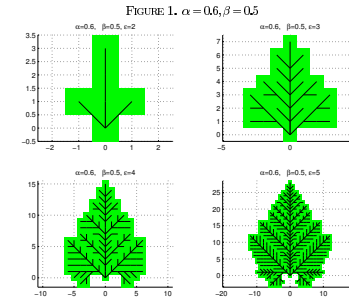
and

Z_j = link j 's impedance.

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

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Growing networks—two parameter model: [21]



Parameters control impedance ($0 \leq \alpha < 1$) and angles of junctions ($0 < \beta$)

For this example: $\alpha = 0.6$ and $\beta = 0.5$

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Outline

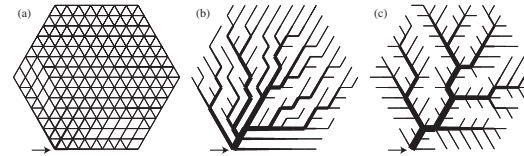
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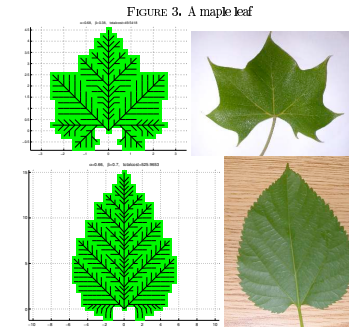


- (a) $\gamma > 1$: Braided (bulk) flow
 - (b) $\gamma < 1$: Local minimum: Branching flow
 - (c) $\gamma < 1$: Global minimum: Branching flow
- Note: This is a single source supplying a region.

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

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Growing networks: [21]



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

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Optimal supply networks

What's the best way to distribute stuff?

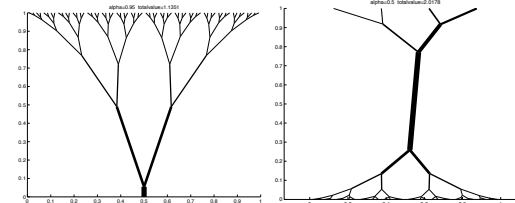
- Stuff = medical services, energy, people, ...
- Some fundamental network problems:
 1. Distribute stuff from a single source to many sinks
 2. Distribute stuff from many sources to many sinks
 3. Redistribute stuff between nodes that are both sources and sinks

Supply and Collection are equivalent problems

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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. [20]

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Single source optimal supply

An immensely controversial issue ...

- The form of natural branching networks: Random, optimal, or some combination? [6, 19, 2, 5, 4]
- River networks, blood networks, ...

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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Optimality:

- Optimal channel networks^[13]
- Thermodynamic analogy^[14]

versus ...

Randomness:

- Scheidegger's directed random networks
- Undirected random networks

Optimization—Murray's law

Aside on P_{drag}

- Work done = $F \cdot d$ = energy transferred by force F
- 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
- So $\Phi \Delta p$ = Force · velocity

Optimization—Murray's law

Murray's law:

Find:

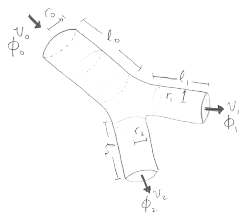
$$\Phi = kr^3$$

Insert assignment question

All of this means we have a groovy cube-law:

$$r_{parent}^3 = r_{offspring1}^3 + r_{offspring2}^3$$

Optimization—Murray's law



Murray's law (1926) connects branch radii at forks: ^[11, 10, 12, 7, 17]

$$r_{parent}^3 = r_{offspring1}^3 + r_{offspring2}^3$$

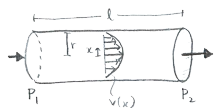
where r_{parent} = radius of 'parent' branch, and $r_{offspring1}$ and $r_{offspring2}$ are radii of the two 'offspring' sub-branches.

- Holds up well for outer branchings of blood networks^[15].
- 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^[16, 17].

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 :

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

- η = dynamic viscosity (units: $ML^{-1}T^{-1}$).
- Power required to overcome impedance:

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

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

$$P_{metabolic} = cr^2\ell$$

Optimization—Murray's law

Murray's law:

Total power (cost):

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

- Observe power increases linearly with ℓ
- 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})

Murray meets Tokunaga:

Φ_ω = volume rate of flow into an order ω vessel segment

Tokunaga picture:

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

Using $\phi_\omega = kr_\omega^3$

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

Same form as:

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

Optimization—Murray's law

Murray's law:

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

Optimization

Murray meets Tokunaga:

- Find Horton ratio for vessel radius $R_r = r_\omega / r_{\omega-1}$.
- 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?

Optimization

Murray meets Tokunaga:

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)^[19] achieve similar results following Horton's laws (but this work is a disaster).

So does Turcotte *et al.* (1998)^[18] using Tokunaga (sort of).

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