

Optimal Supply Networks I: Branching

Complex Networks | @networksvox
CSYS/MATH 303, Spring, 2016

Optimal
transportation

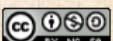
Optimal
branching

Murray's law
Murray meets Tokunaga

References

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Optimal
transportation

Optimal
branching

Murray's law

Murray meets Tokunaga

References



Outline

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References

Optimal transportation

Optimal branching

Murray's law

Murray meets Tokunaga

References



Optimal supply networks

Optimal
transportation

Optimal
branching

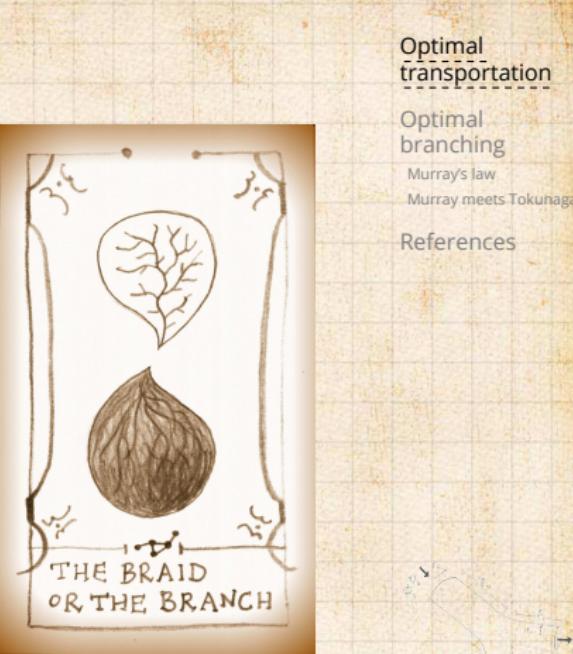
Murray's law
Murray meets Tokunaga

References

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





Single source optimal supply

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References

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

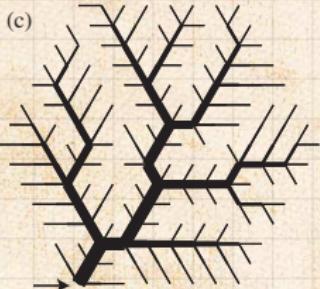
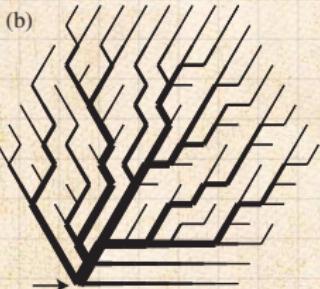
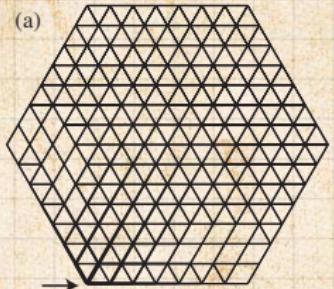
and

Z_j = link j 's impedance?

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



Single source optimal supply



- (a) $\gamma > 1$: **Braided** (bulk) flow
- (b) $\gamma < 1$: Local minimum: **Branching** flow
- (c) $\gamma < 1$: Global minimum: **Branching** flow

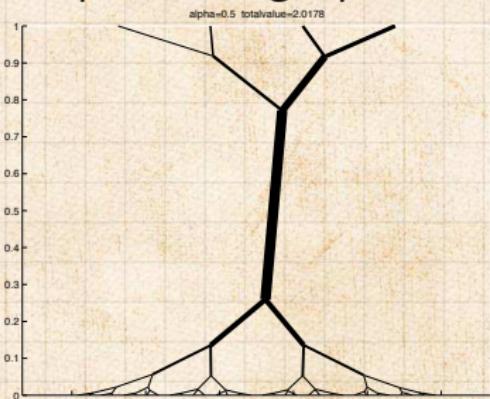
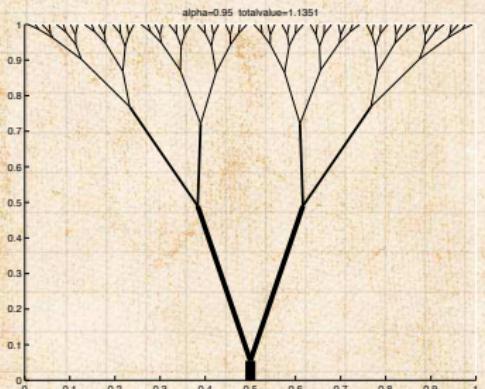
From Bohn and Magnasco [3]
See also Banavar et al. [1]

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Optimal
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Murray's law
Murray meets Tokunaga
References



Single source optimal supply

Optimal paths related to transport (Monge) problems:



Xia (2003) [19]

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Optimal
branching

Murray's law

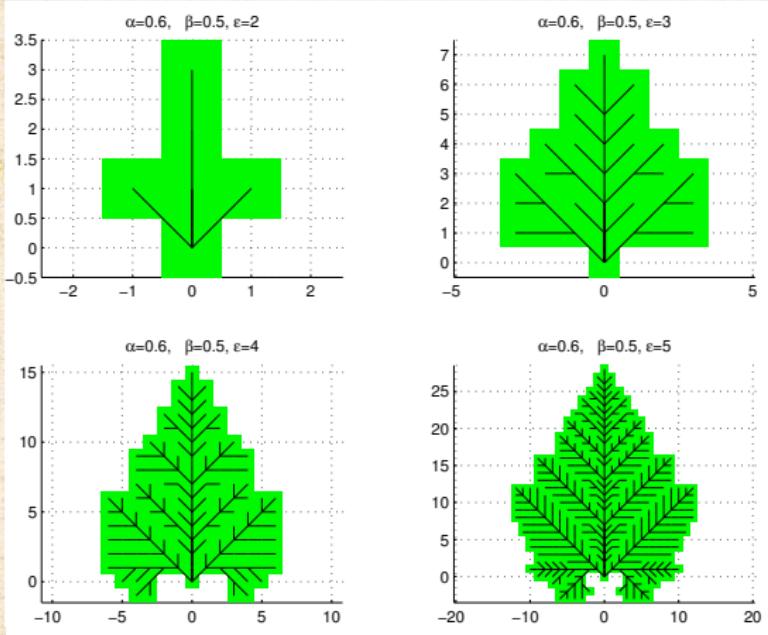
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References



Growing networks:

FIGURE 1. $\alpha = 0.6, \beta = 0.5$



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branching

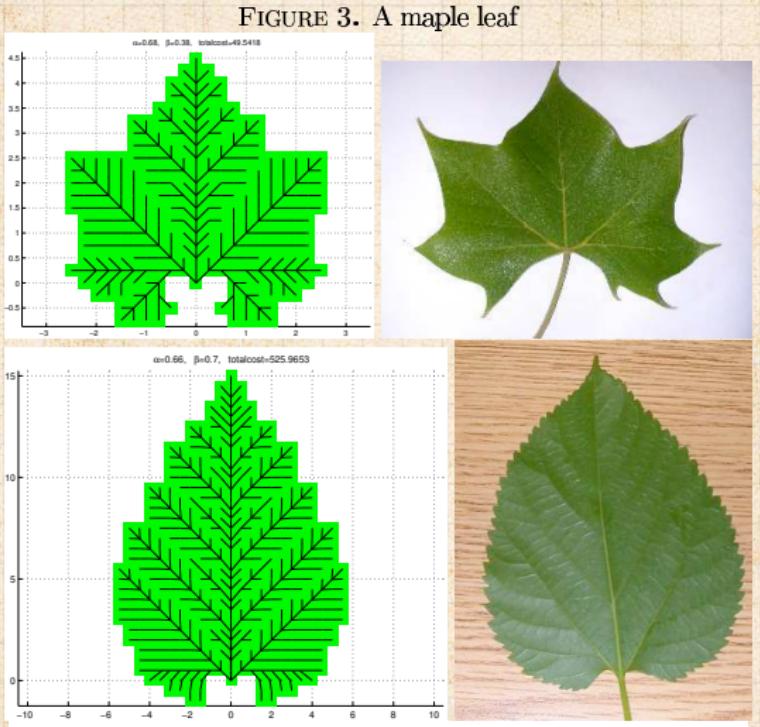
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Murray meets Tokunaga

References



Xia (2007)^[18]

Growing networks:



Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References



Single source optimal supply

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References

An immensely controversial issue...

- ▶ The form of river networks and blood networks:
optimal or not? [17, 2, 5, 4]

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**.



River network models

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Murray's law
Murray meets Tokunaga

References

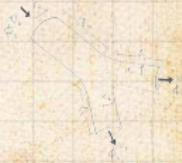
Optimality:

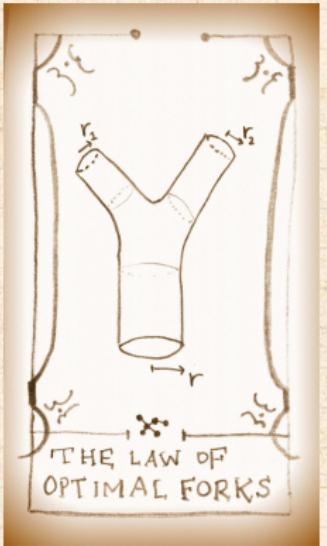
- ▶ Optimal channel networks [12]
- ▶ Thermodynamic analogy [13]

versus...

Randomness:

- ▶ Scheidegger's directed random networks
- ▶ Undirected random networks





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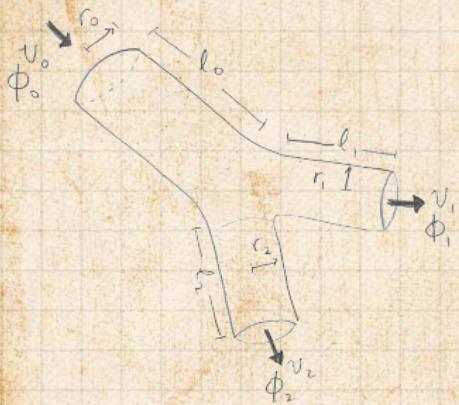
Murray's law

Murray meets Tokunaga

References



Optimization—Murray's law



- ▶ Murray's law (1926) connects branch radii at forks: [10, 9, 11, 6, 15]

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

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 [11, 7, 8].
- ▶ See D'Arcy Thompson's "On Growth and Form" for background inspiration [14, 15].

Optimal transportation

Optimal branching

Murray's law

Murray meets Tokunaga

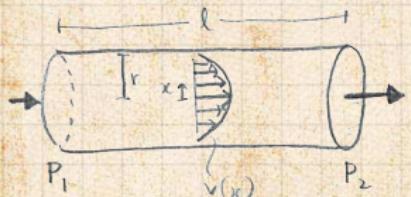
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- ▶ Use hydraulic equivalent of Ohm's law:

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

where Δp = pressure difference, Φ = flux.



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

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

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

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

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

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

Optimal transportation

Optimal branching

Murray's law
Murray meets Tokunaga

References



Optimization—Murray's law

Optimal
transportation

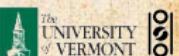
Optimal
branching

Murray's law
Murray meets Tokunaga

References

Aside on P_{drag}

- ▶ Work done = $F \cdot d$ = energy transferred by force F
- ▶ Power = P = rate work is done = $F \cdot v$
- ▶ Δp = Force per unit area
- ▶ Φ = Volume per unit time
= cross-sectional area · velocity
- ▶ So $\Phi \Delta p$ = Force · velocity



Optimization—Murray's law

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References

Murray's law:

- ▶ Total power (cost):

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

- ▶ 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})



Optimization—Murray's law

Murray's law:

- Minimize P with respect to r :

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

$$= -4\Phi^2 \frac{8\eta\ell}{\pi r^5} + c2r\ell = 0$$

- Rearrange/cancel/slap:

$$\Phi^2 = \frac{c\pi r^6}{16\eta} = k^2 r^6$$

where $k = \text{constant}$.

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transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References



Optimization—Murray's law

Murray's law:

- ▶ So we now have:

$$\Phi = kr^3$$

- ▶ 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

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

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

Optimization

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

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

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

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References



Optimization

Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References

Murray meets Tokunaga:

- ▶ 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

Optimal
transportation

Optimal
branching

Murray's law

Murray meets Tokunaga

References

Murray meets Tokunaga:

- ▶ Isometry: $V_\omega \propto \ell_\omega^3$
- ▶ Gives

$$R_\ell^3 = R_v = R_n$$

- ▶ We need one more constraint...
- ▶ West et al (1997)^[17] achieve similar results following Horton's laws.
- ▶ So does Turcotte et al. (1998)^[16] using Tokunaga (sort of).



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Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

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Optimal transportation

Optimal branching

Murray's law

Murray meets Tokunaga

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Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

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Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

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Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

References



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Optimal
transportation

Optimal
branching

Murray's law
Murray meets Tokunaga

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