

Déterminisme et indéterminisme multi-échelle, invariants et variance, des systèmes vivants.

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<https://www.afscet.asso.fr/pagesperso/Bricage.html>

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Bricage P. (2018) **Déterminisme et indéterminisme multi-échelle, invariants et variance des systèmes vivants. -présentation-**. Journées annuelles de l'AFSCET à Andé, France, 05 mai 2018, 84 p., http://afscet.asso.fr/Ande18/pbricage_Ande2018_interactions-PowerLaws_presentation.pdf

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UES-EUS, Bruxelles, Belgique, Europe <http://www.ues-eus.eu>

Directorate **World Organisation of Systems and Cybernetics**

WOSC, London, UK <http://www.wosc.co>

Secretary General **International Academy for Systems and Cybernetic Sciences**

IASCYS, Wien, Österreich, Europa <http://iascys.org>

Président Association **ALBA**, Trésorier Association **PELLEAS**

Lasclaveries, France, Europe <http://armsada.eu>



SURVIVRE

1. La MOBILISATION de la MATIÈRE et de l'ÉNERGIE
2. La CROISSANCE en masse et en nombre
3. La RÉPONSE à des STIMULATIONS
4. L'ORGANISATION interne, dans l'espace et dans le temps
5. L'INTÉGRATION exteme
6. Le MOUVEMENT

pour **SE SURVIVRE**

7. La REPRODUCTION



“ invariance de jauge des systèmes vivants “
Bricage P. (1991)

BRICAGE Pierre (1991) [Les caractéristiques des organismes vivants](#). 44 p.
Biologie, A.P.I.D.S., Fac. Sci. & Techniques, Univ. de Pau & Pays de l'Adour.

SURVIVRE

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7. La REPRODUCTION

La survie des organismes vivants.
04 février 2000, Paris, 33 p.
<http://www.afscet.asso.fr/SURVIVRE.pdf>

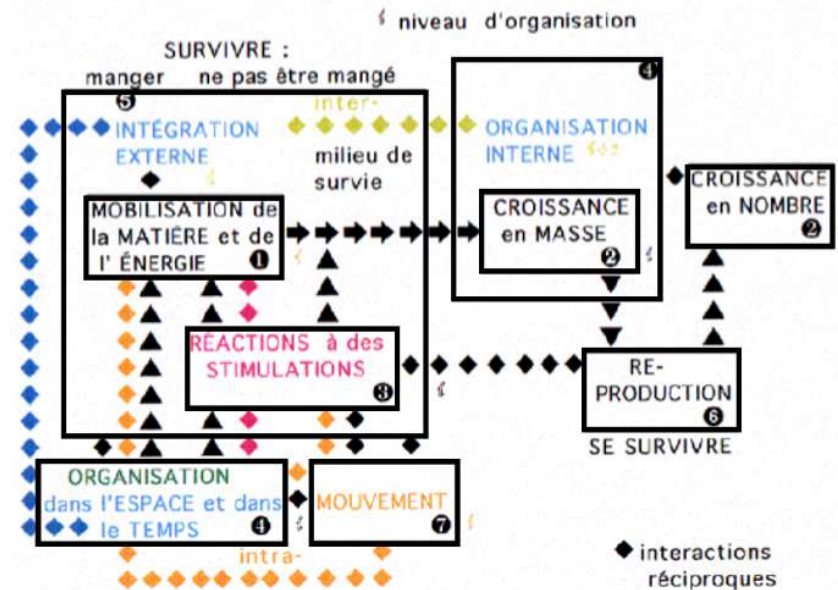
les 7 degrés de liberté de la vie

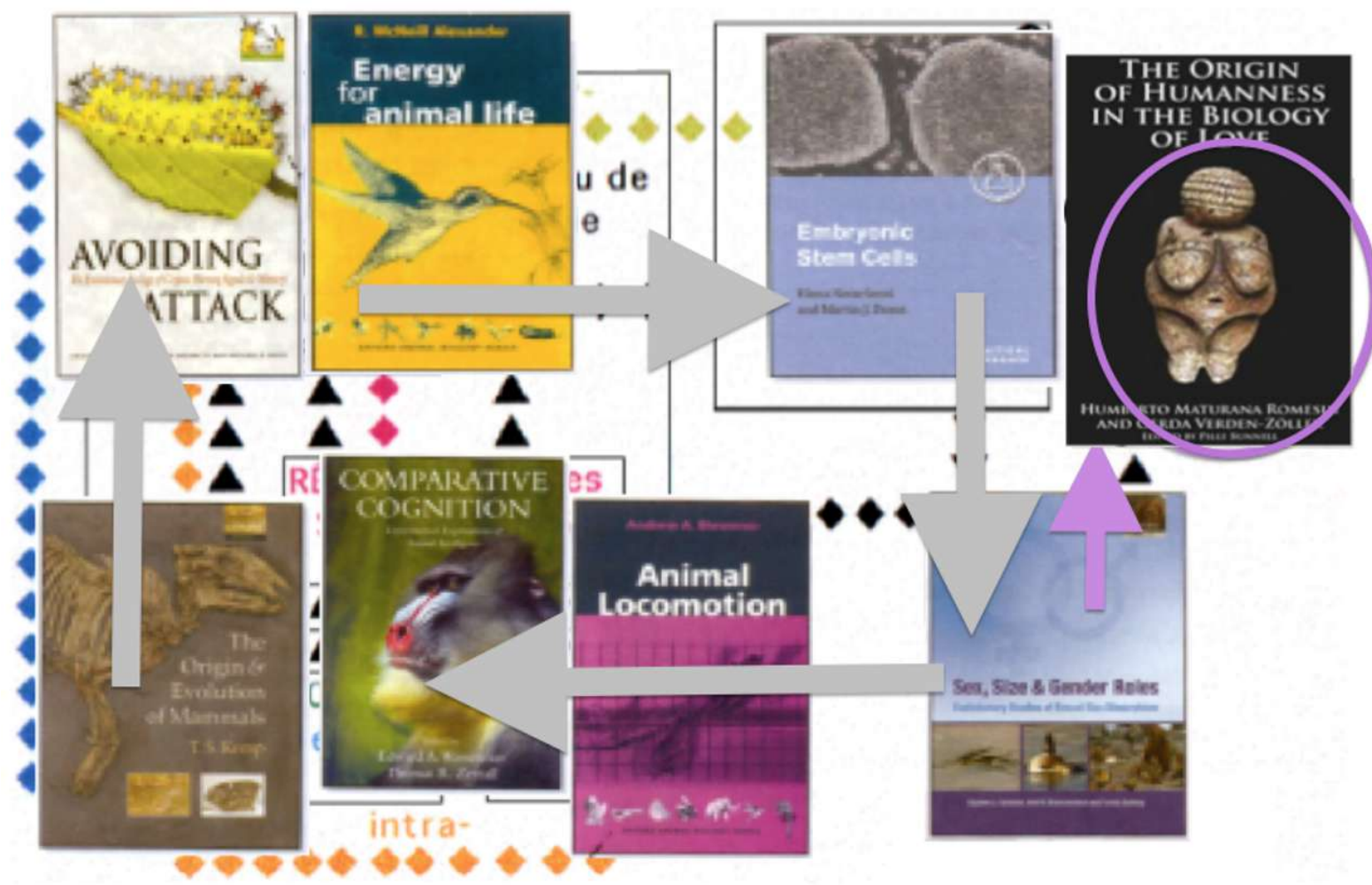
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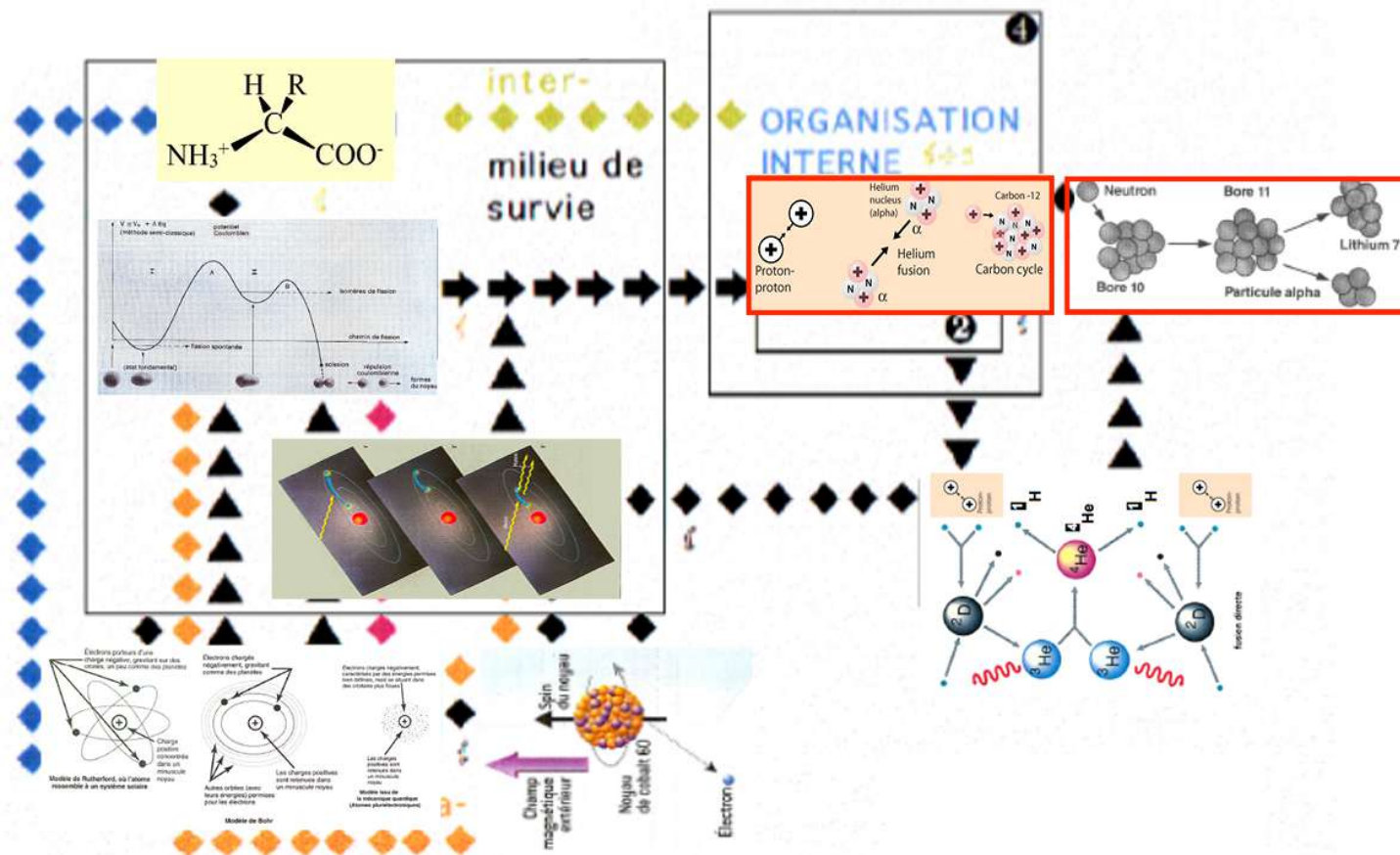
p. 2/11

L'évolution, du vivant au social. 1er décembre 2001, AFSCET, I.I.A.P., Paris

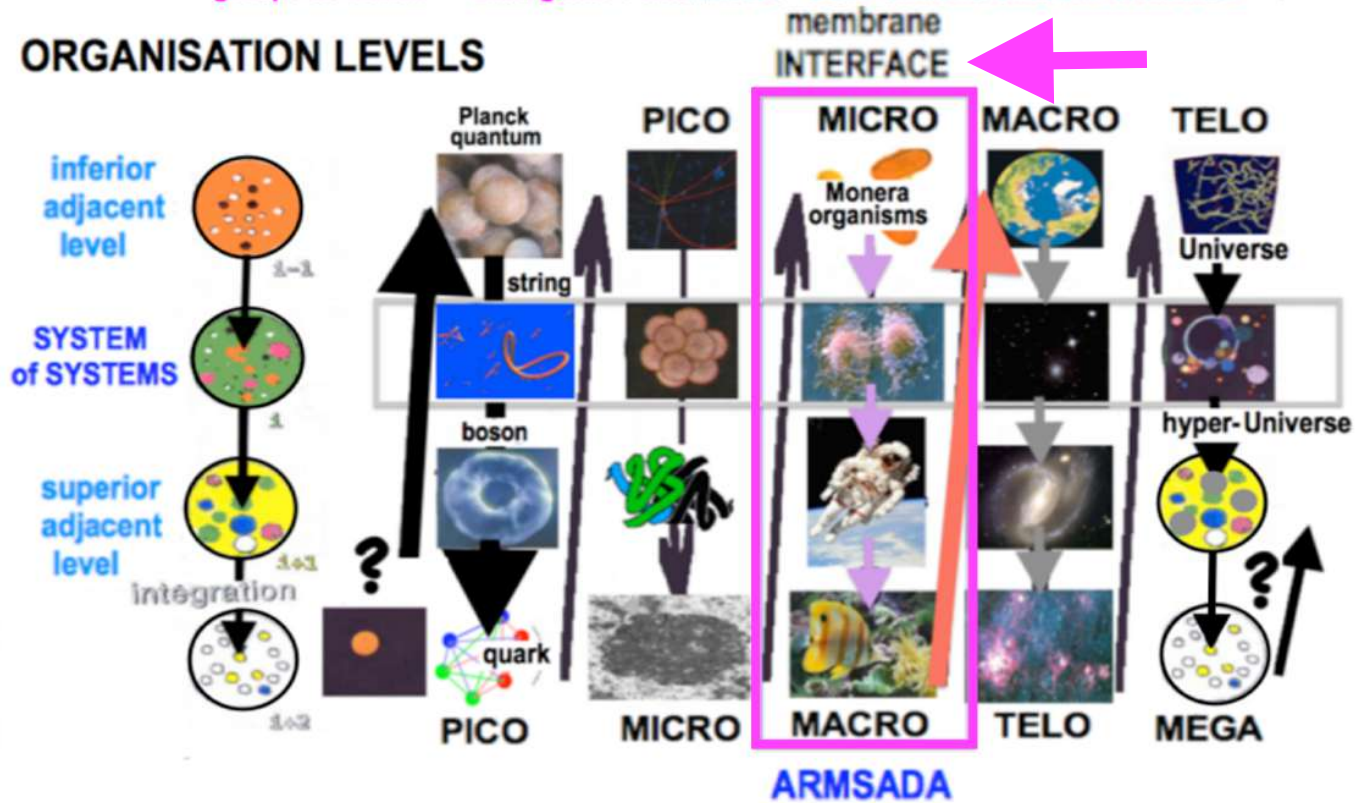
1- Les caractéristiques du vivant "vivant".







"groupe de travail" Emergence Paris; 11 fév 2013 maison des Arts et Métiers



intégration : écoexotopie - endophysiotopie

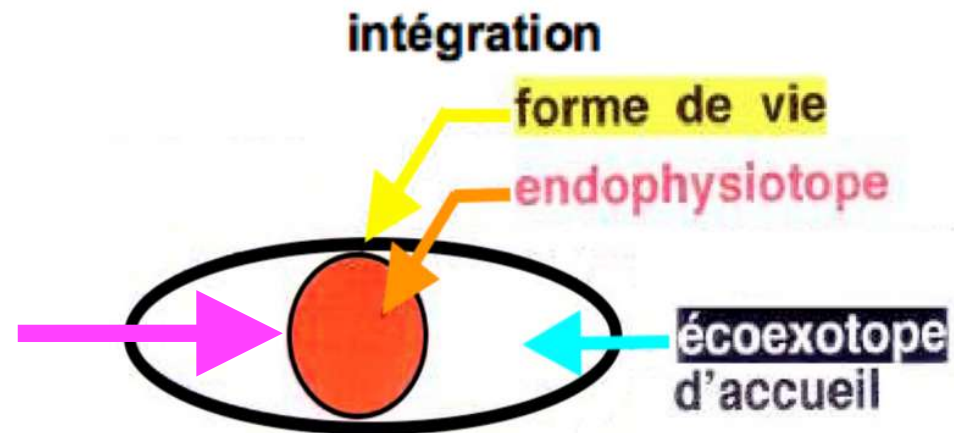
capacité d'accueil – capacité d'être accueilli

SURVIVRE

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pour **SE SURVIVRE**

7. La REPRODUCTION



La nature de la décision dans la nature ?

19 mai 2001, Andé, 16 p.

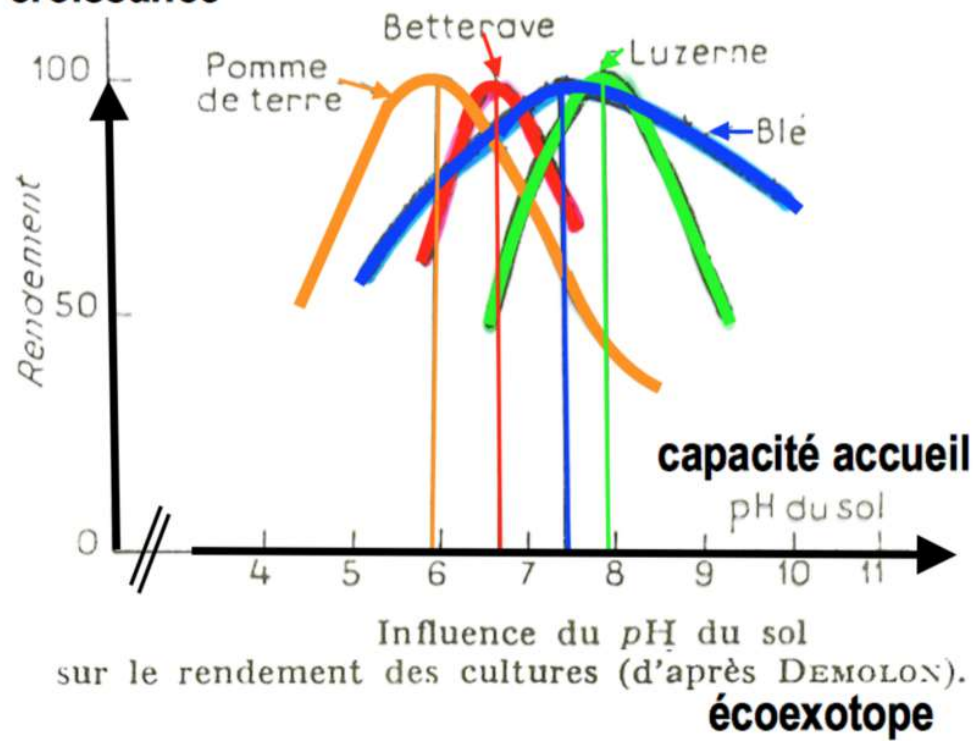
<http://www.afscet.asso.fr/Decision.pdf>

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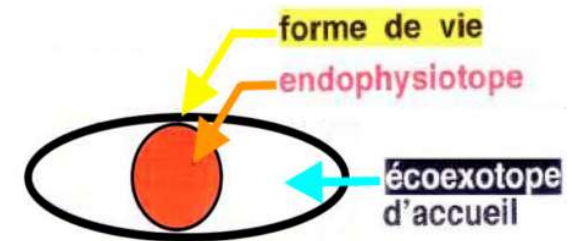
intégration : écoexotopie - endophysiologie

capacité d'accueil – capacité d'être accueilli

endophysiologie croissance

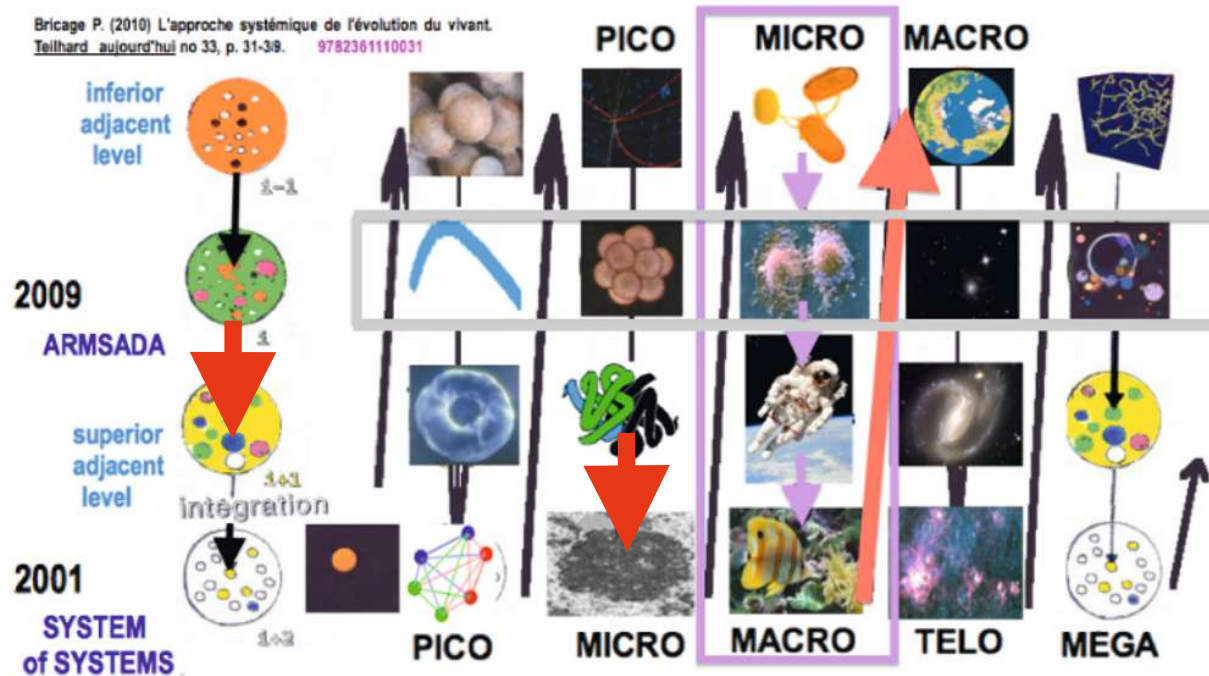


intégration



ORGANISATION LEVELS: PERIODIC CLASSIFICATION CHART

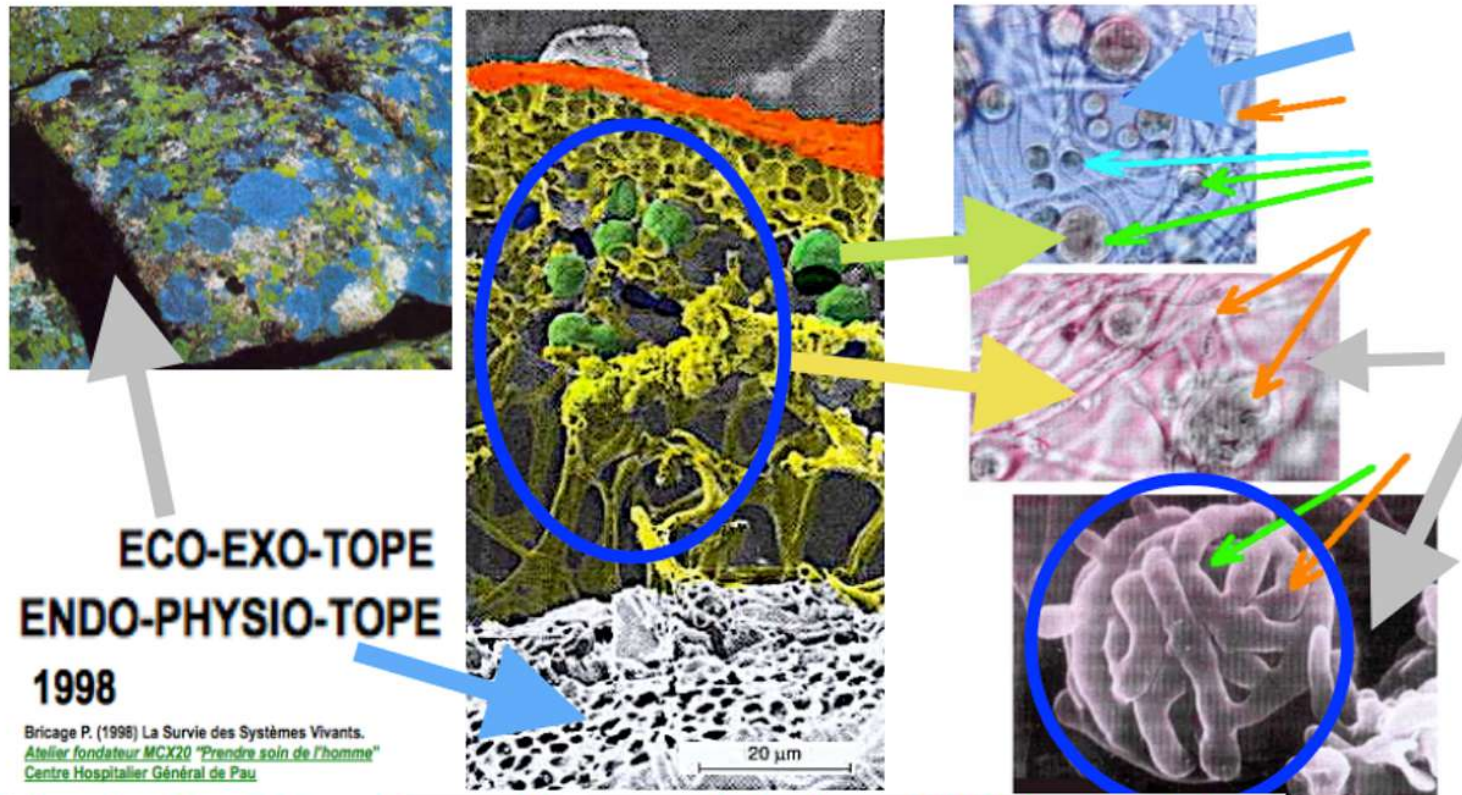
Bricage P. (2010) L'approche systémique de l'évolution du vivant.
 Teilhard aujourd'hui no 33, p. 31-39. 9782361110031



**Todos los nuevos planes de organización
 de los seres vivos son construidos por
 la implementación de una ARMSADA**

Associations for the Reciprocal and Mutual Sharing of Advantages and DisAdvantages

1.1. A LICHEN is NOT a WIN-WIN association BUT AN ARMSADA



El "endophysiotope" es a la vez el huésped (el invitado) y el rehén (el preso) de un ecoexotope para sobrevivir.

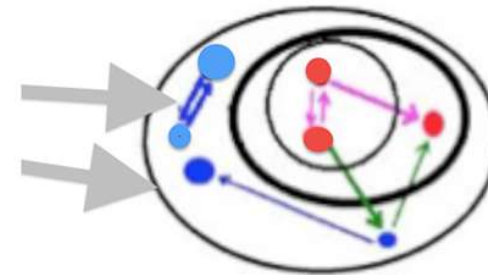
a lichen
AN ORGANISM MADE OF
A MULTICELL ORGANISM
A CELL ORGANISM POPULATION
A BACTERIAL POPULATION



AN ECOSYSTEM
WITH A FOOD CHAIN

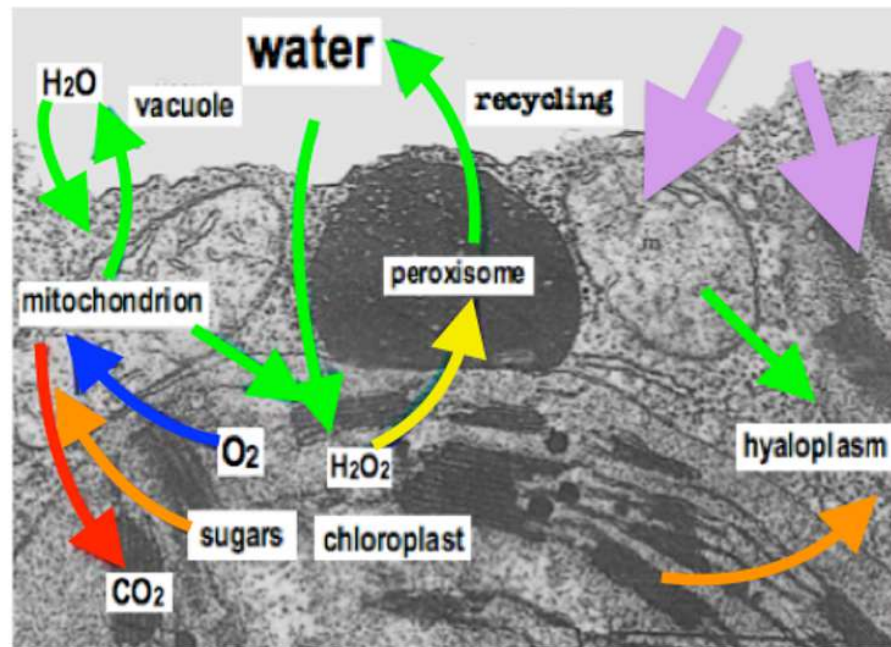
ARMSADA

Association for
the **R**eciprocal
and **M**utual
Sharing of
Advantages
and **D**is**A**dvantages



Associations for the Reciprocal and Mutual Sharing of Advantages and DisAdvantages

A CELL IS AN ENDOSYNCENOSIS, AN ECOSYSTEM of ORGANISMS

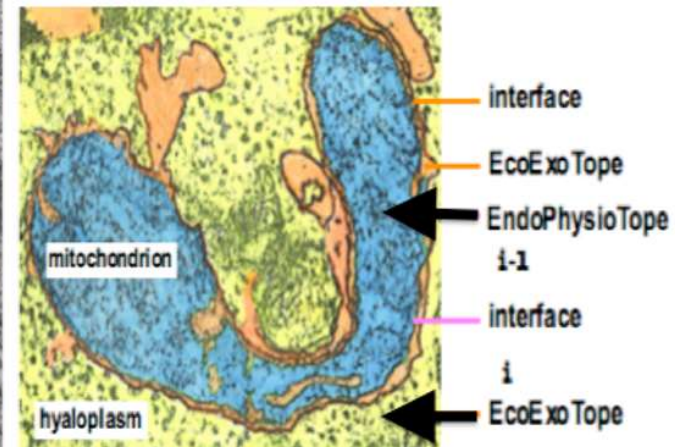


Bricage P. (1986) Isoperoxidases, markers of surrounding and physiological changes, in situ in leaves and in vitro in calli of *Pedilanthus tithymaloides* L. variegatus: cell compartmentation and polyfunctionality, control of activity by phenols, specific roles. p. 261-265. [Molecular & Physiological Aspects of Plant Peroxidases](#). Univ. Genève, (ISBN 2-88164-001-X)

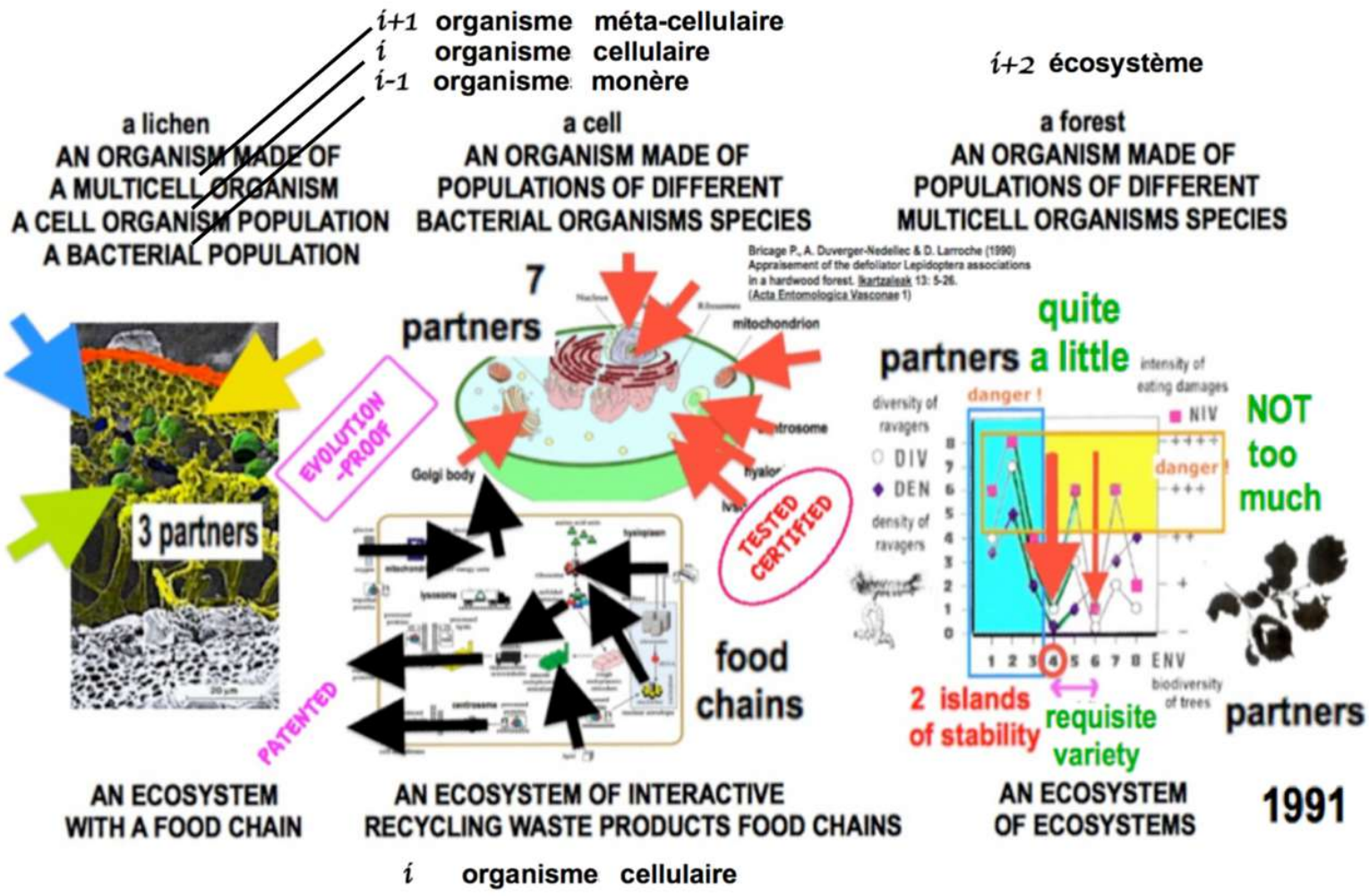
ISSS July 18-23, 2010, Wilfrid Laurier University, Waterloo, ON, Canada. Balancing Individualism and Collectivism: ARMSADA

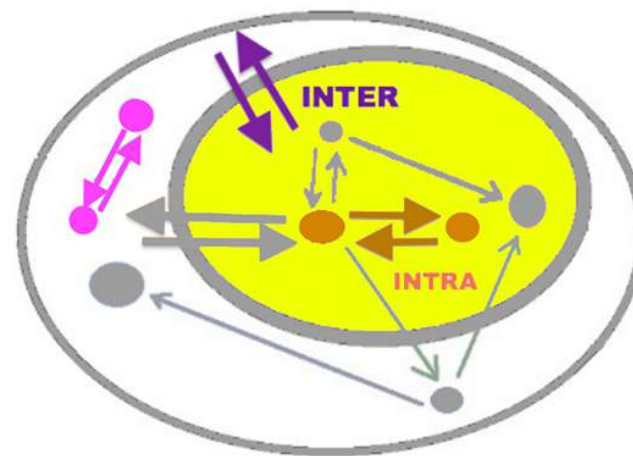
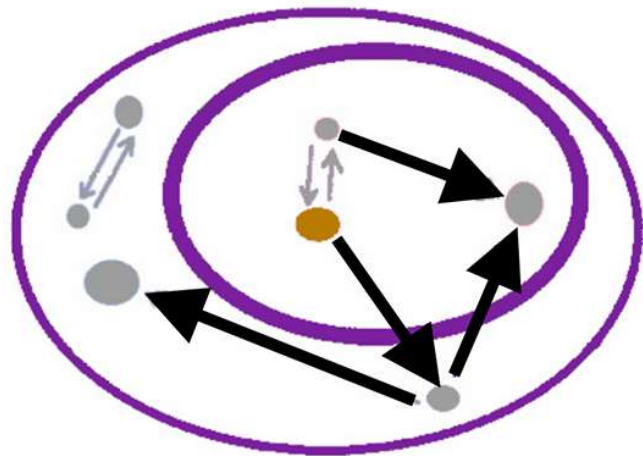
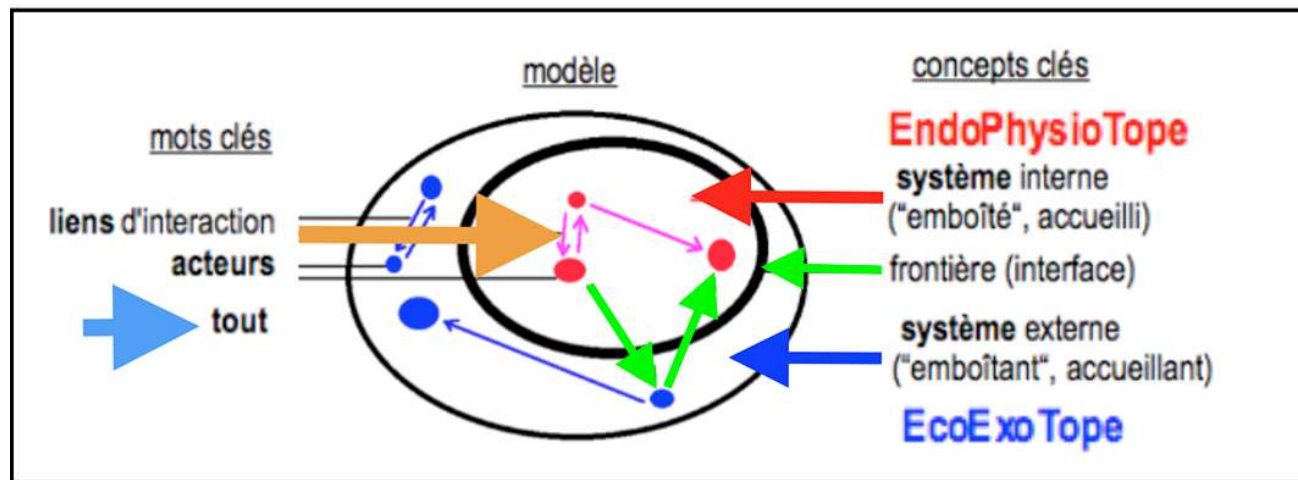
E PLURIBUS
UNUM
IN VARIETATE
CONCORDIA

UNUS PRO OMNIBUS
OMNES PRO UNO
UN POUR TOUS
TOUS POUR UN

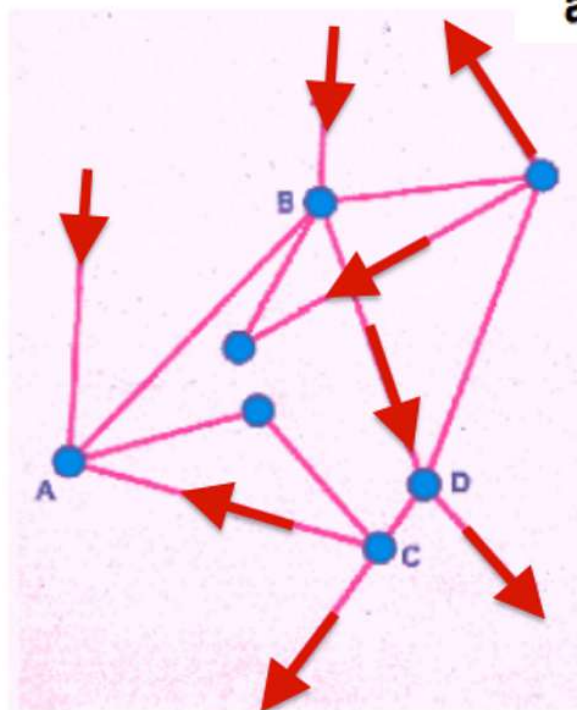


**FOR THE ONE TO SURVIVE
THE OTHER ONE MUST SURVIVE FIRST
AND RECIPROCALLY 2000**

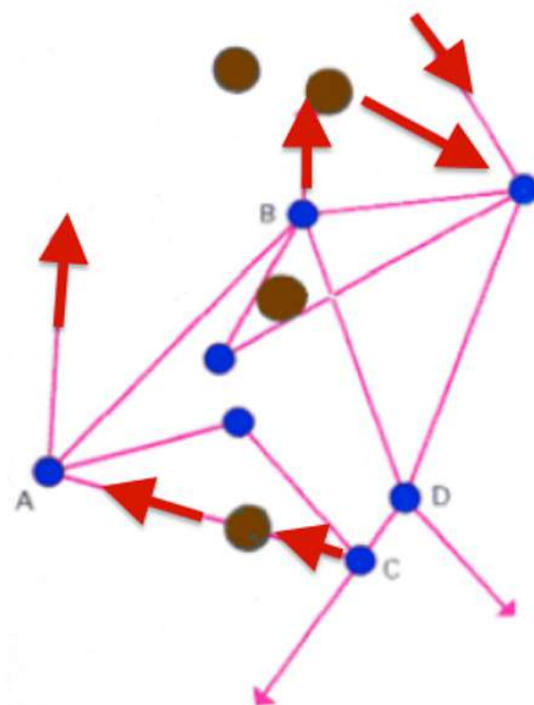




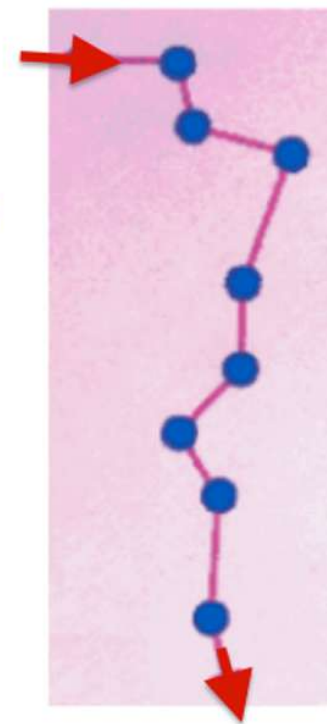
emergence of a new network
PERCOLATION
a metamorphosis process



before

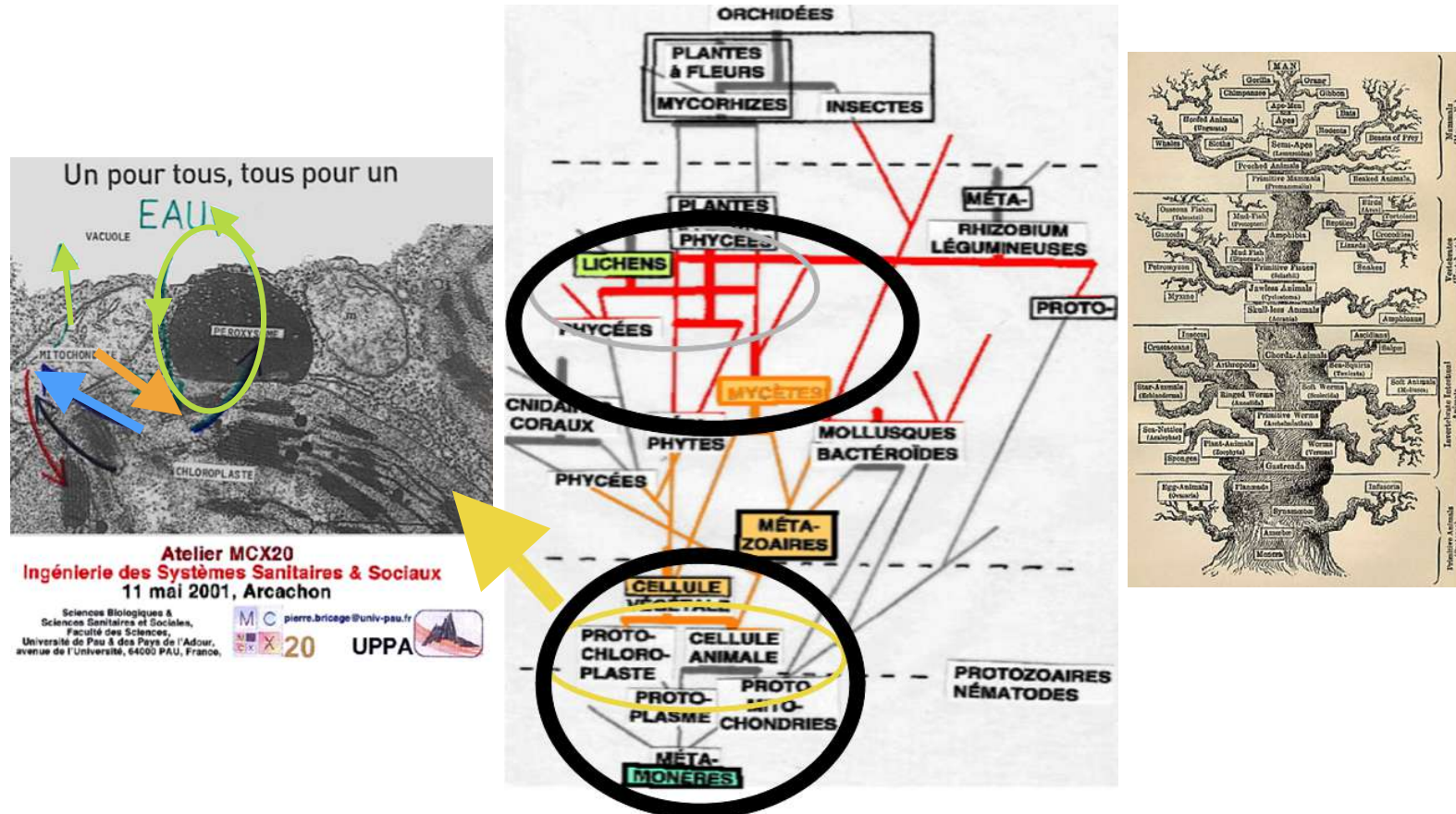


during



after

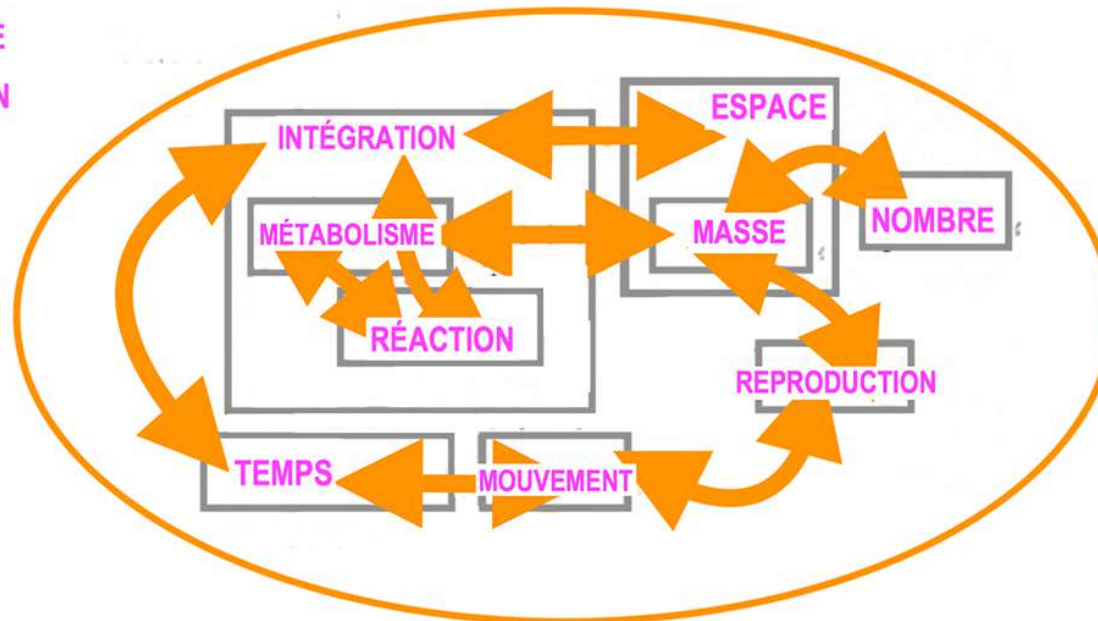
II.C. *“Life is a coral not a tree”*

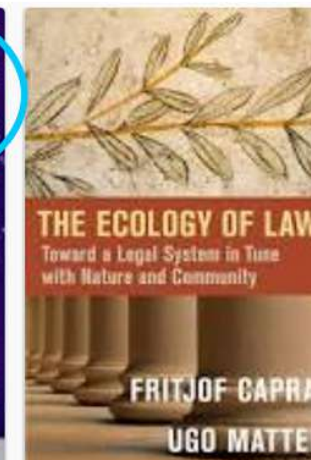
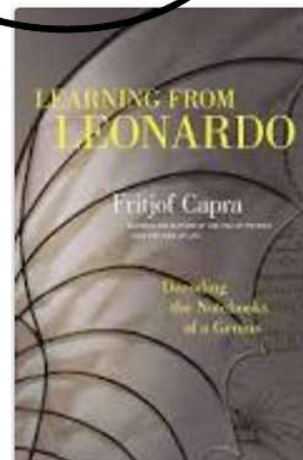
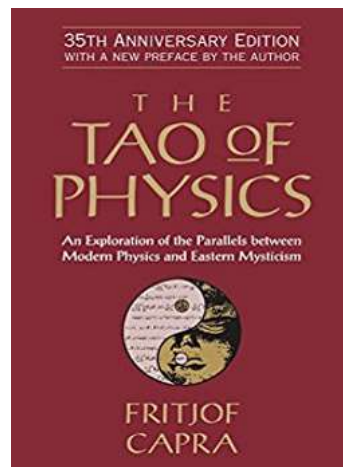
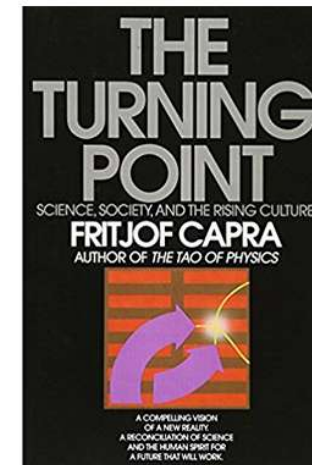
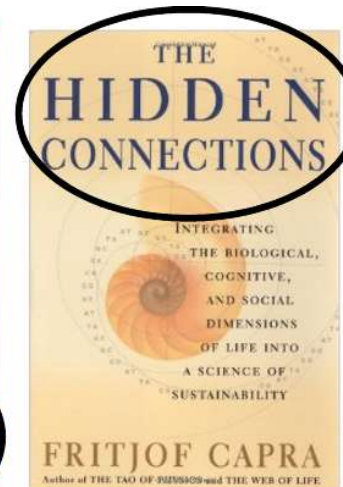
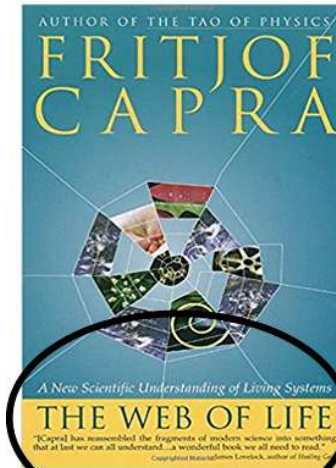
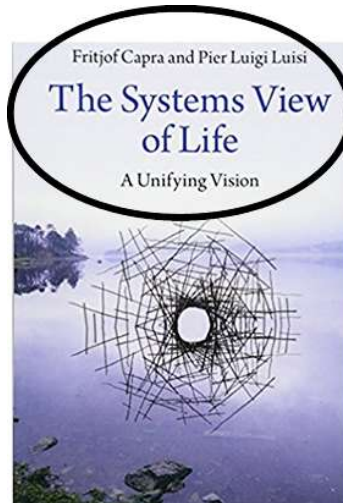


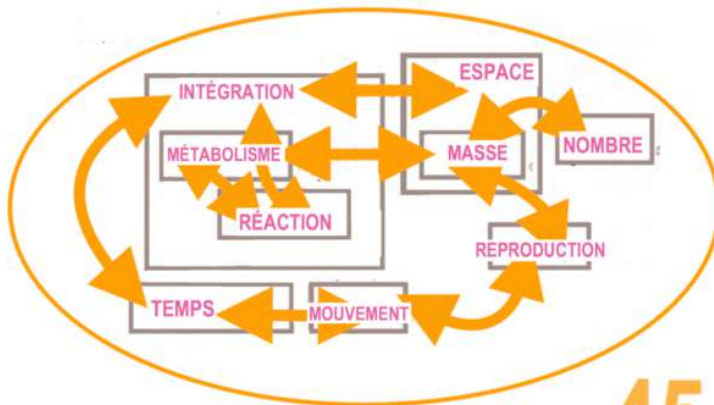
SURVIVRE

1. La MOBILISATION **MÉTABOLISME**
de la MATIÈRE et de l'ÉNERGIE
2. La CROISSANCE en masse **MASSE**
2. La CROISSANCE en nombre **NOMBRE**
3. La RÉPONSE à des STIMULATIONS **RÉACTION**
4. L'ORGANISATION interne dans l'espace et dans le temps **ESPACE**
TEMPS
5. L'**INTÉGRATION** externe
6. Le **MOUVEMENT**
pour **SE SURVIVRE**
7. La **REPRODUCTION**

systemique



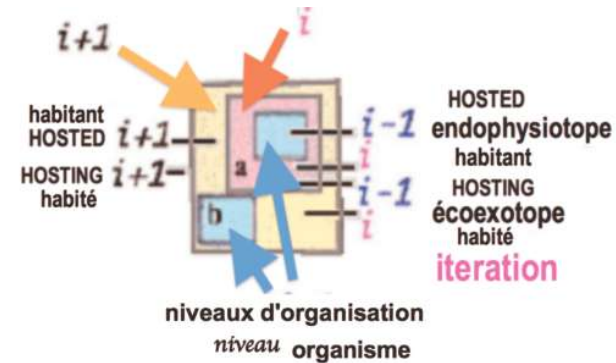




45 X

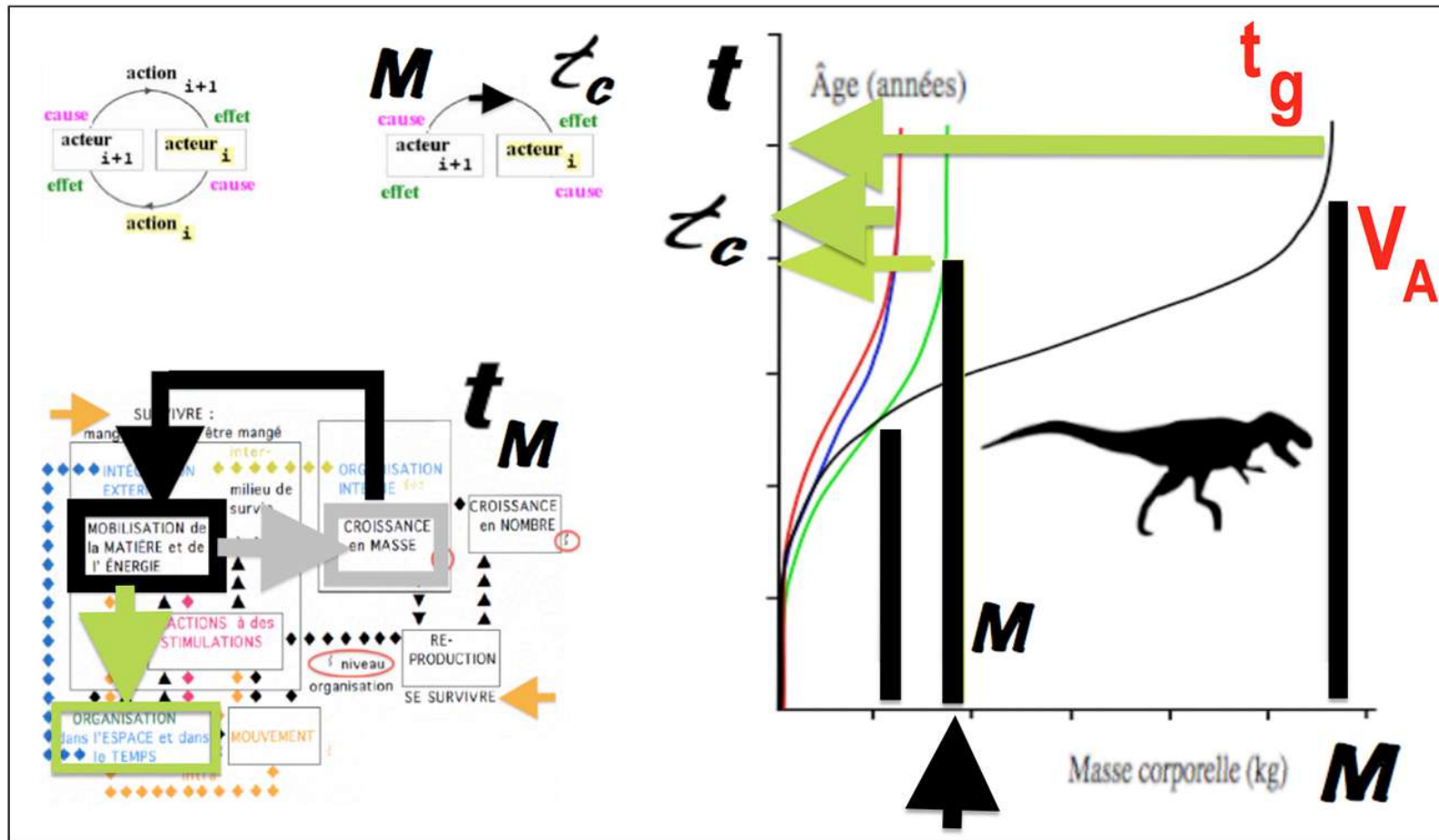
endophysiotope

métabolisme	↔	métabolisme
croissance en masse	↔	croissance en masse
réponses aux stimulations	↔	réponses aux stimulations
organisation spatiale	↔	organisation spatiale
organisation temporelle	↔	organisation temporelle
intégration	↔	intégration
mouvement	↔	mouvement
reproduction	↔	reproduction
croissance en nombre	↔	croissance en nombre



- i-10
- i-9 Planck quantum
- i-8 cordes
- i-7 super-cordes
- i-6 quarks
- i-5 particules
- i-4 atomes
- i-3 molécules
- i-2 amas de molécules
- i-1 organismes monères
- i organismes cellulaires
- i+1 organismes méta-cellulaires
- i+2 écosystèmes
- i+3 biosphères**
- i+4 systèmes stellaires
- i+5 galaxies
- i+6 amas de galaxies
- i+7 univers
- i+8 Hyper-univers
- i+9

18 X



Pierre BRICAGE

atemporalité, temporalité et intemporalité des systèmes vivants

Sensibilité différentielle et Temps de RÉACTION

Le Temps de Réaction Simple diminue selon une fonction de puissance de l'intensité du son.

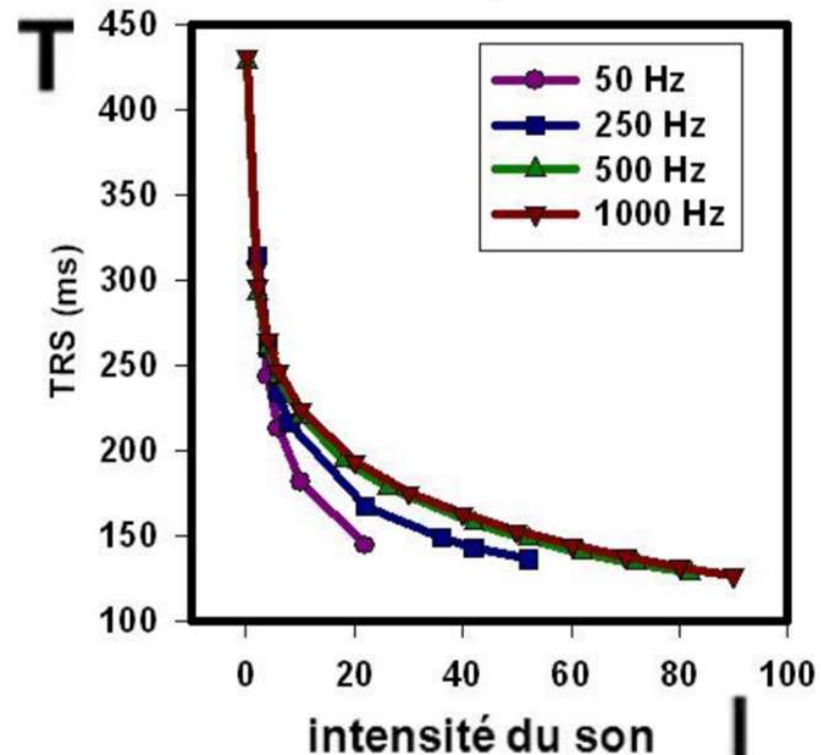
Fonction de Piéron :

$$T = (\beta I^{-\alpha}) + t_0$$

L'exposant de la fonction varie selon la fréquence du son.

La RÉPONSE à
des STIMULATIONS

Temps de Réaction Simple



$$T = (\beta I^{-\alpha}) + t_0$$

Loi de Piéron en audition

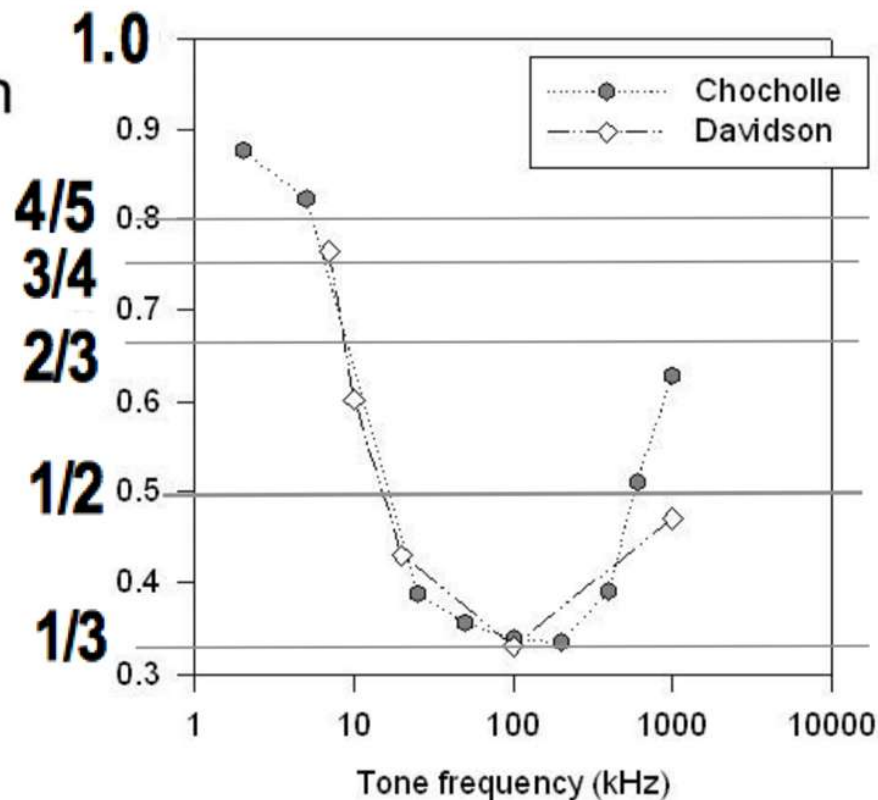
L'exposant varie curvilinéairement avec la fréquence du son avec un minimum dans la région 1000 – 2000 Hz

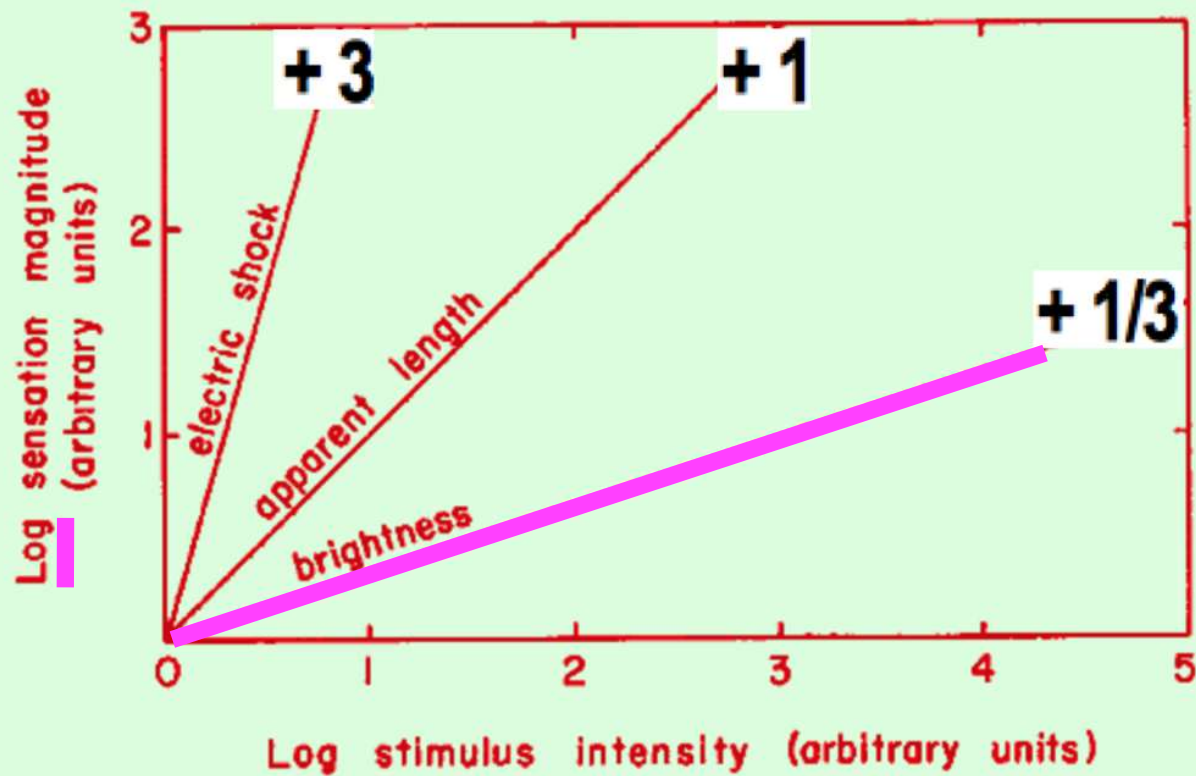
Chocholle 1940

Davidson 1990

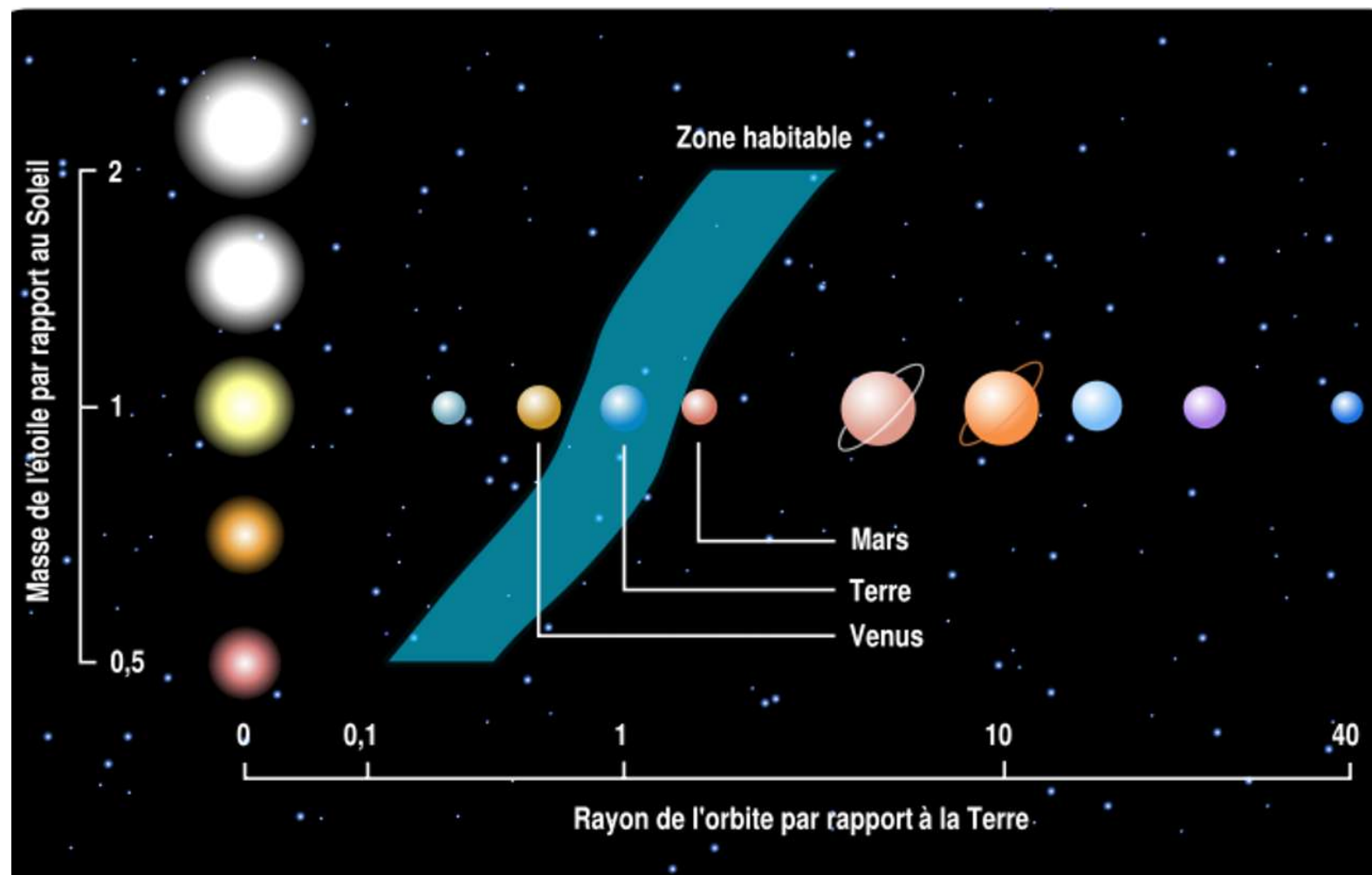
La courbe ressemble à celle des seuils de détection.

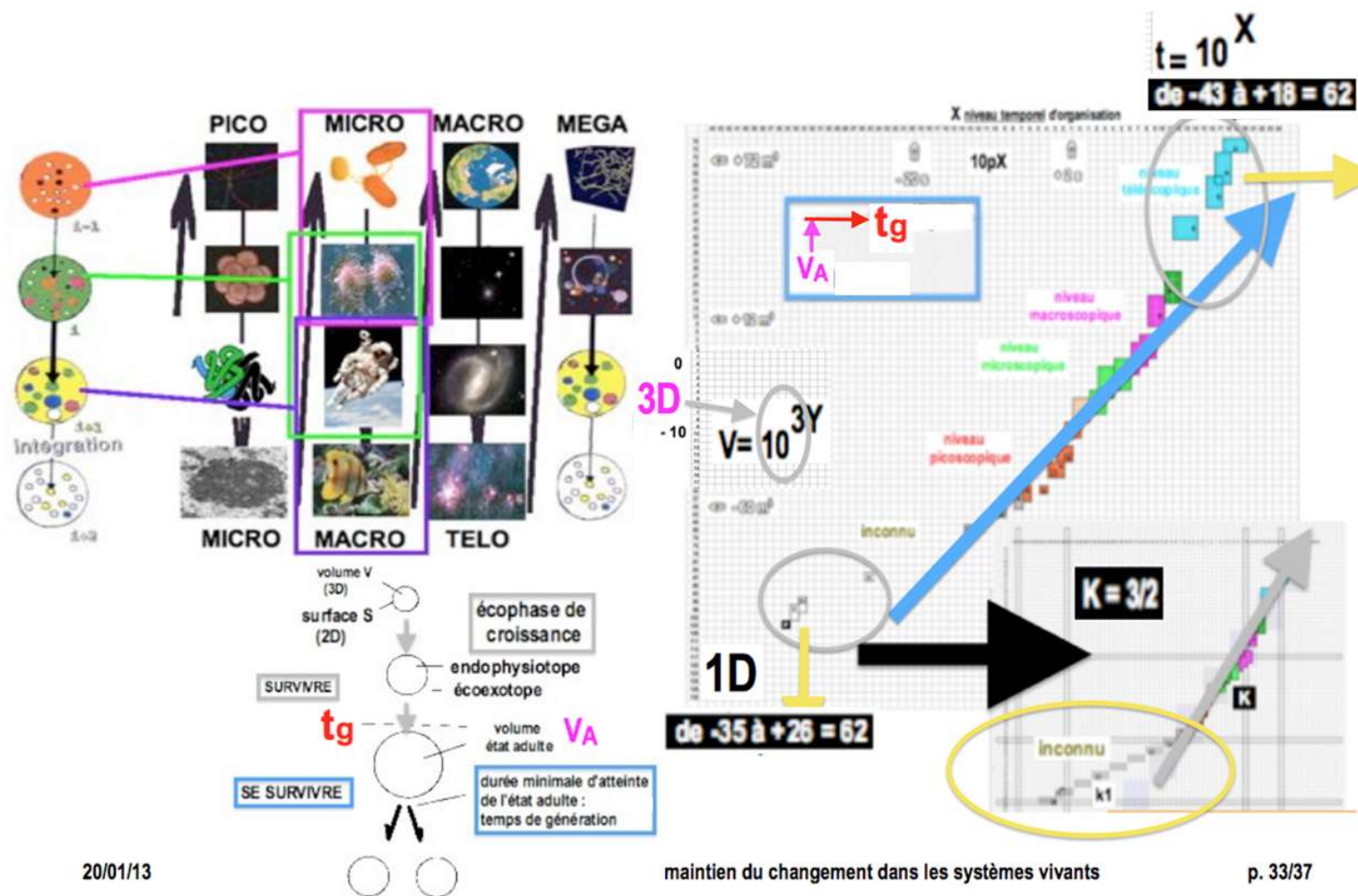
exposant α

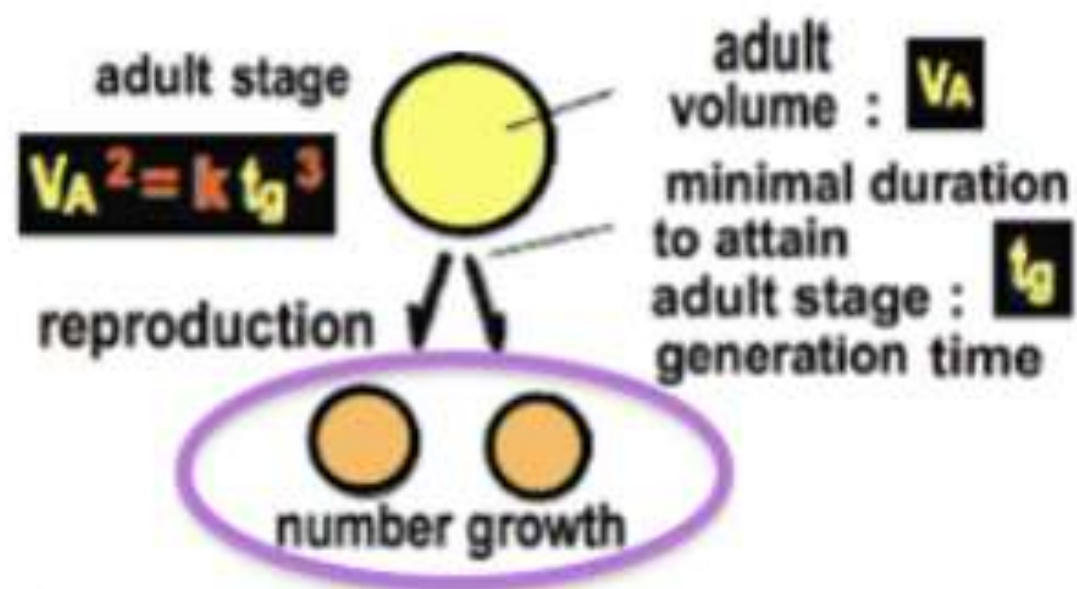




Psychophysical magnitude functions for three perceptual continua. The linearity of the functions on double logarithmic coordinates indicates that sensation magnitude is a power function of stimulus intensity. The slope of the line corresponds to the exponent of the power function.

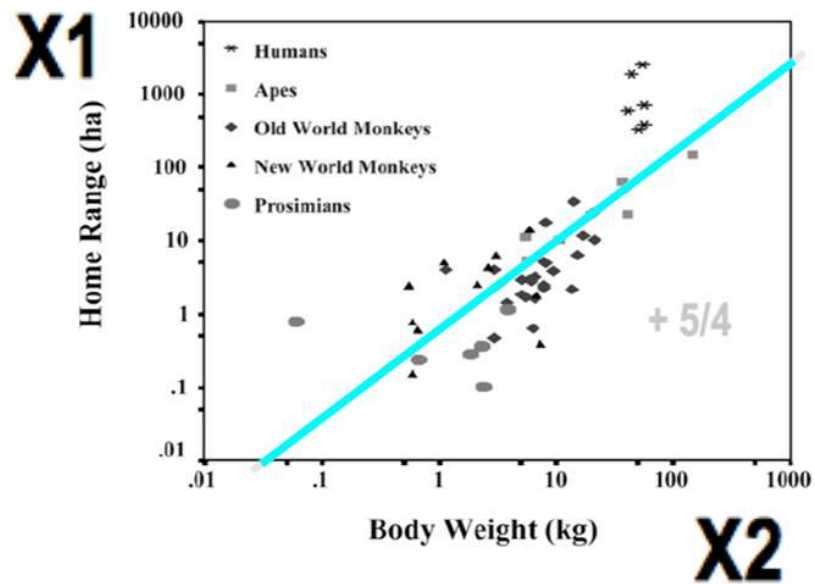






L'INTÉGRATION **exteme**

EcoExoTope

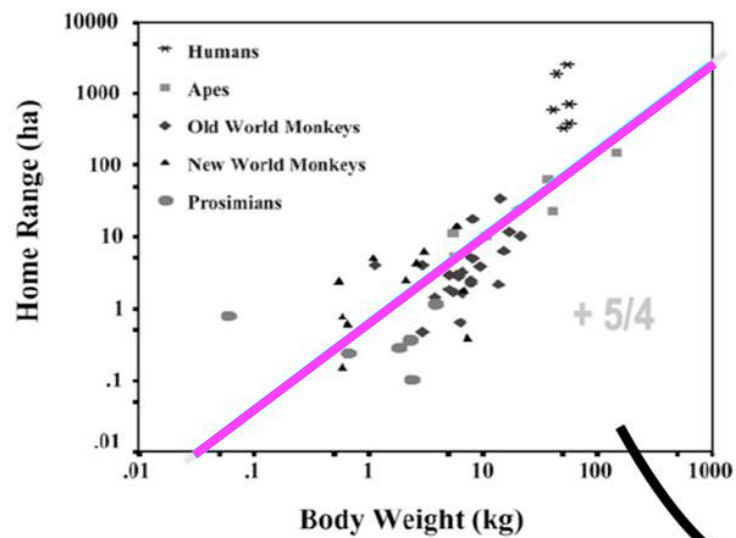


La CROISSANCE
en masse

MASSE

EndoPhysioTope

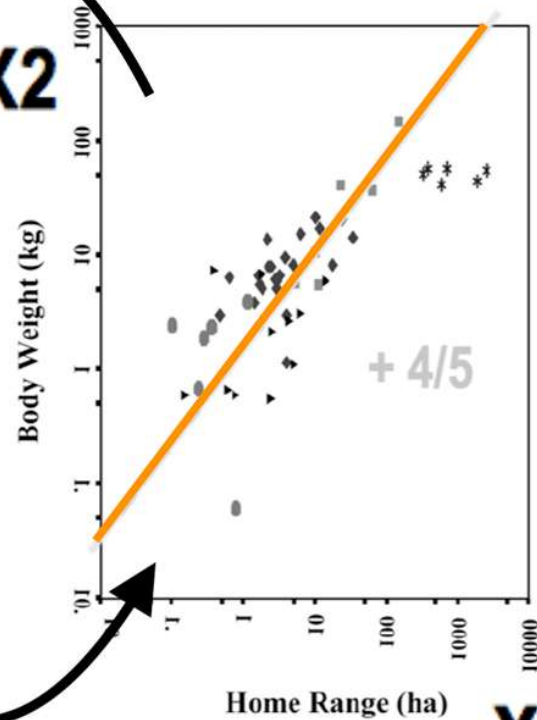
X1



EndoPhysioTope

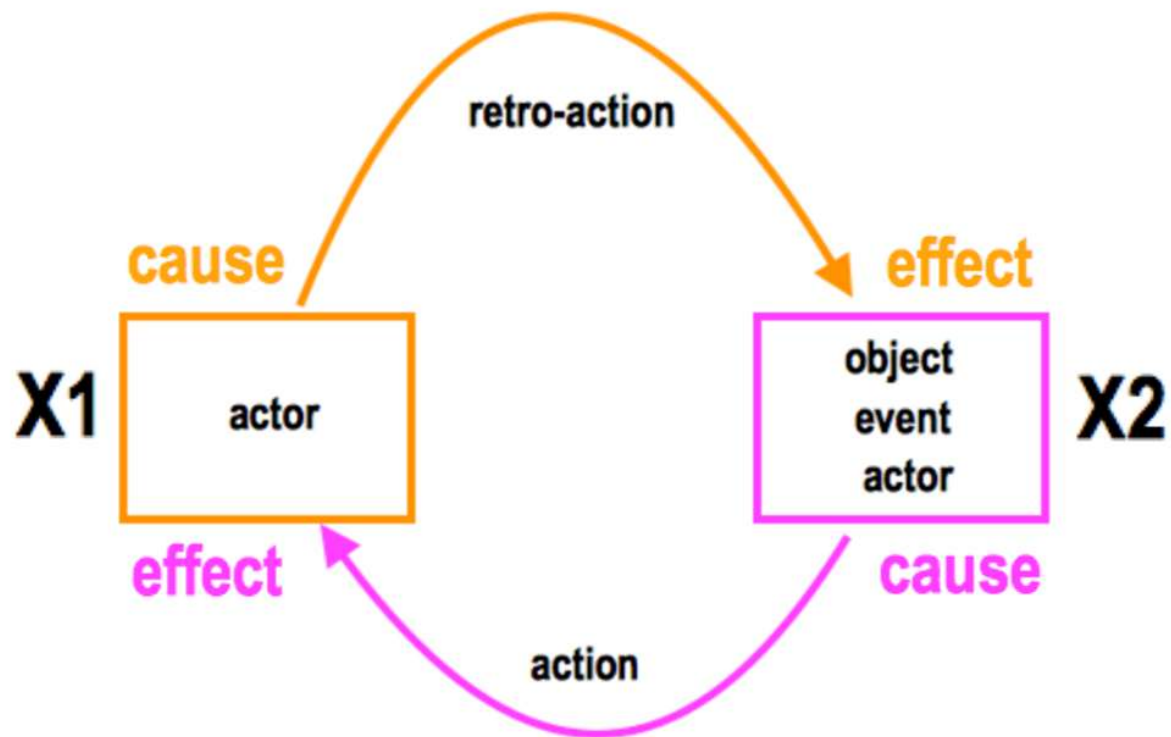
X2

X2



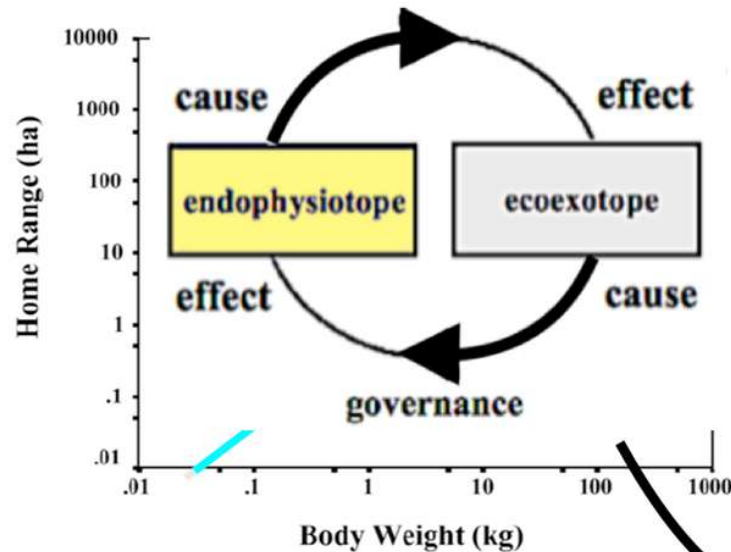
EcoExoTope

X1



loi systémique constructale

X1

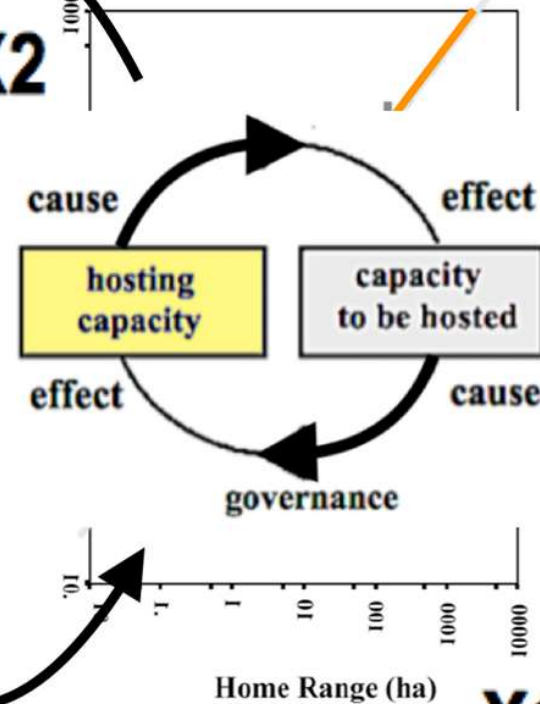


EndoPhysioTope X2

La cause conduit à l'effet (dépendance)
et l'effet retourne à la cause (**rétroaction**).

La fin est le commencement et
le commencement est la fin ("palindrome").

X2



EcoExoTope X1

$$V_A^2 = C.t_g^3$$

$$t_g = K.V_A^{2/3}$$

larval stage

duration in s

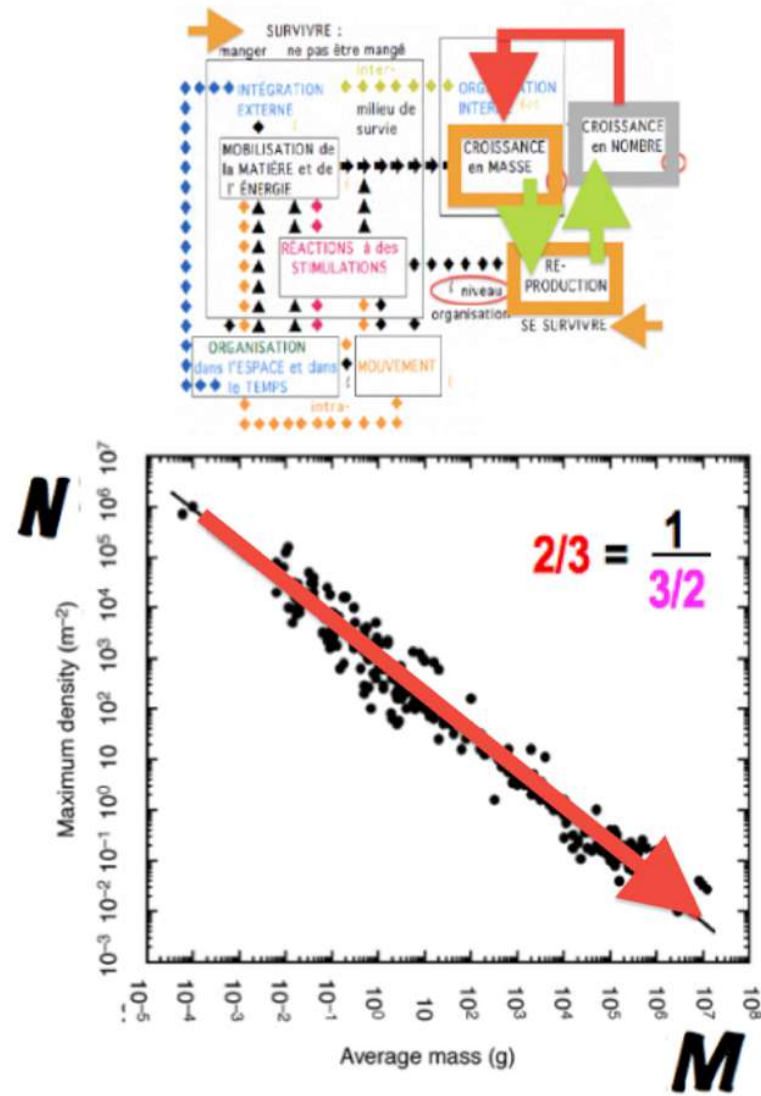
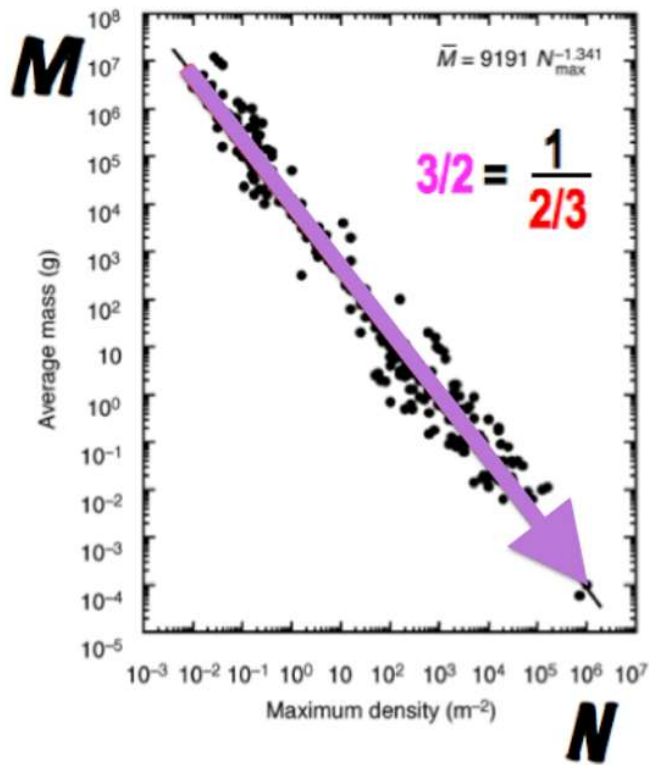
'generation time'

$$V_A = C.t_g^{3/2}$$

reproductive Adult

volume in m^3

Allometric scaling of plant energetics and population density



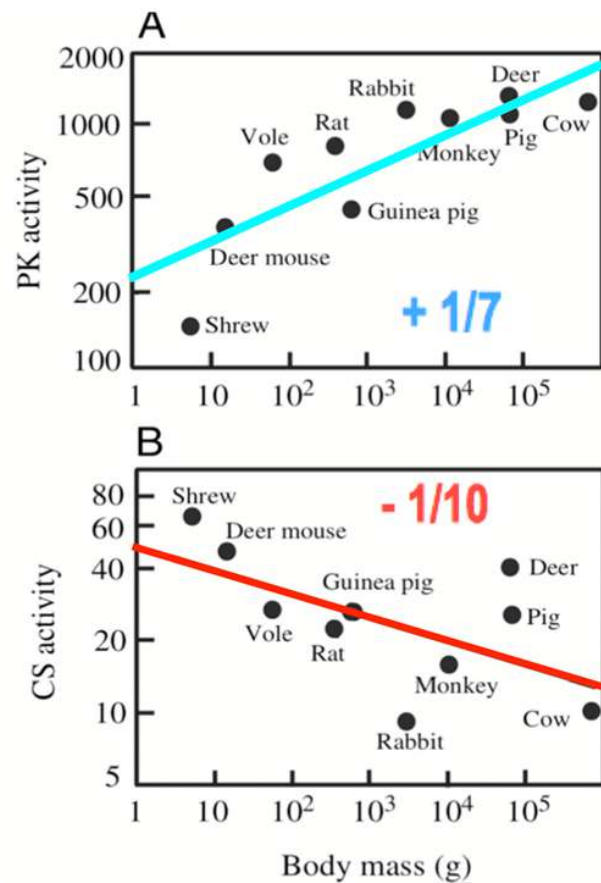


Fig. 1. Mitochondrial and glycolytic enzyme activities in muscles of mammals (adapted from Emmett and Hochachka, 1981). PK, pyruvate kinase (U g^{-1}); CS, citrate synthase (U g^{-1}); body mass (g).

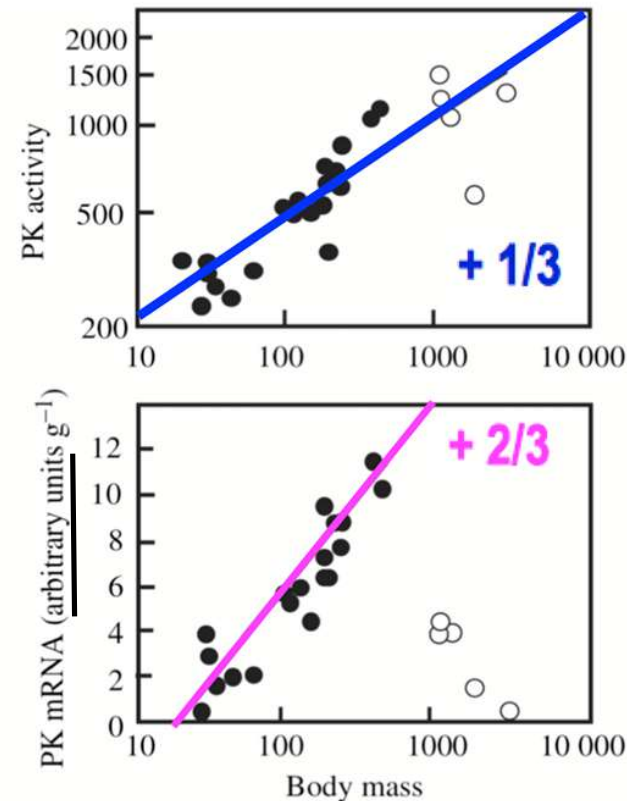
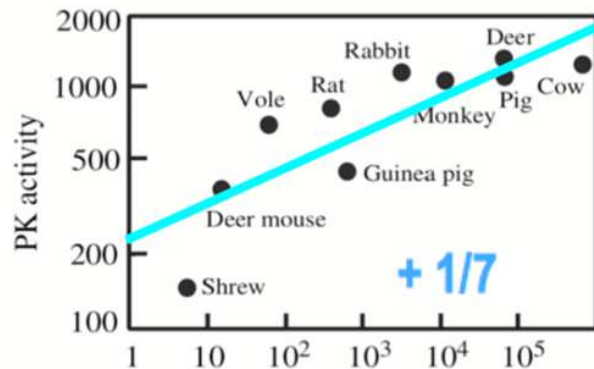


Fig. 2. Relationship between pyruvate kinase activity and mRNA. PK shows positive allometric scaling ($b=0.2$). Larger fish (open circles) are able to produce more protein from a lower amount of mRNA (adapted from Burness et al., 1999). PK, pyruvate kinase (U g^{-1}); body mass (g).

muscle bioenergetic gene expression: allometric scaling relationships of glycolytic and oxidative enzymes



https://en.wikipedia.org/wiki/Citrate_synthase

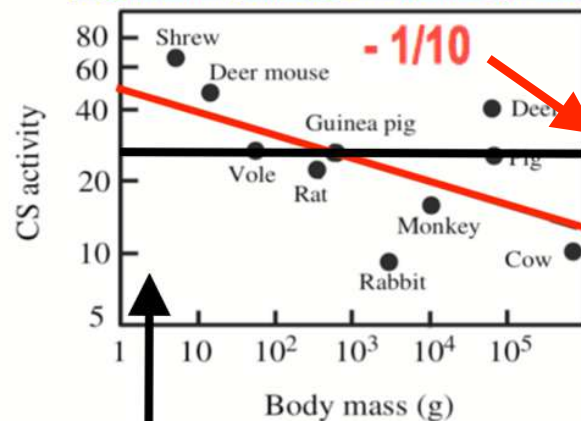


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https://en.wikipedia.org/wiki/Pyruvate_kinase

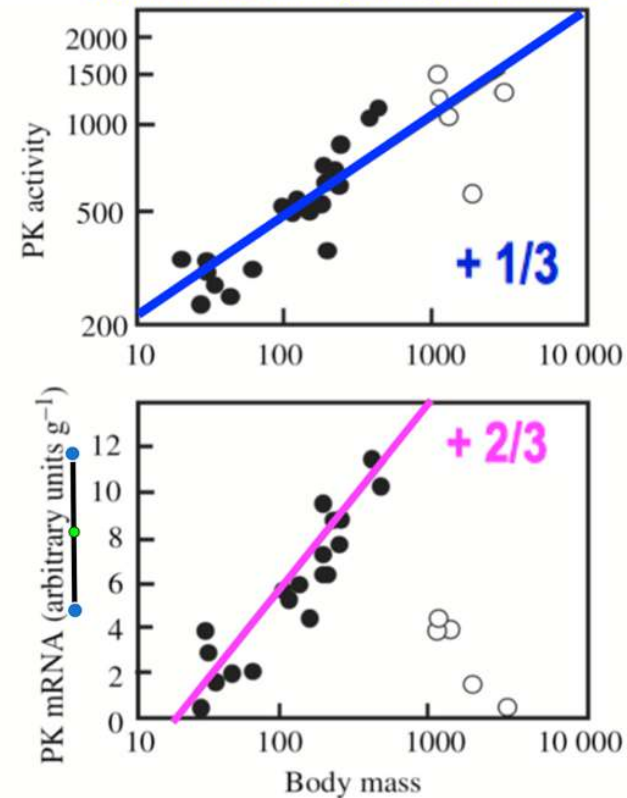


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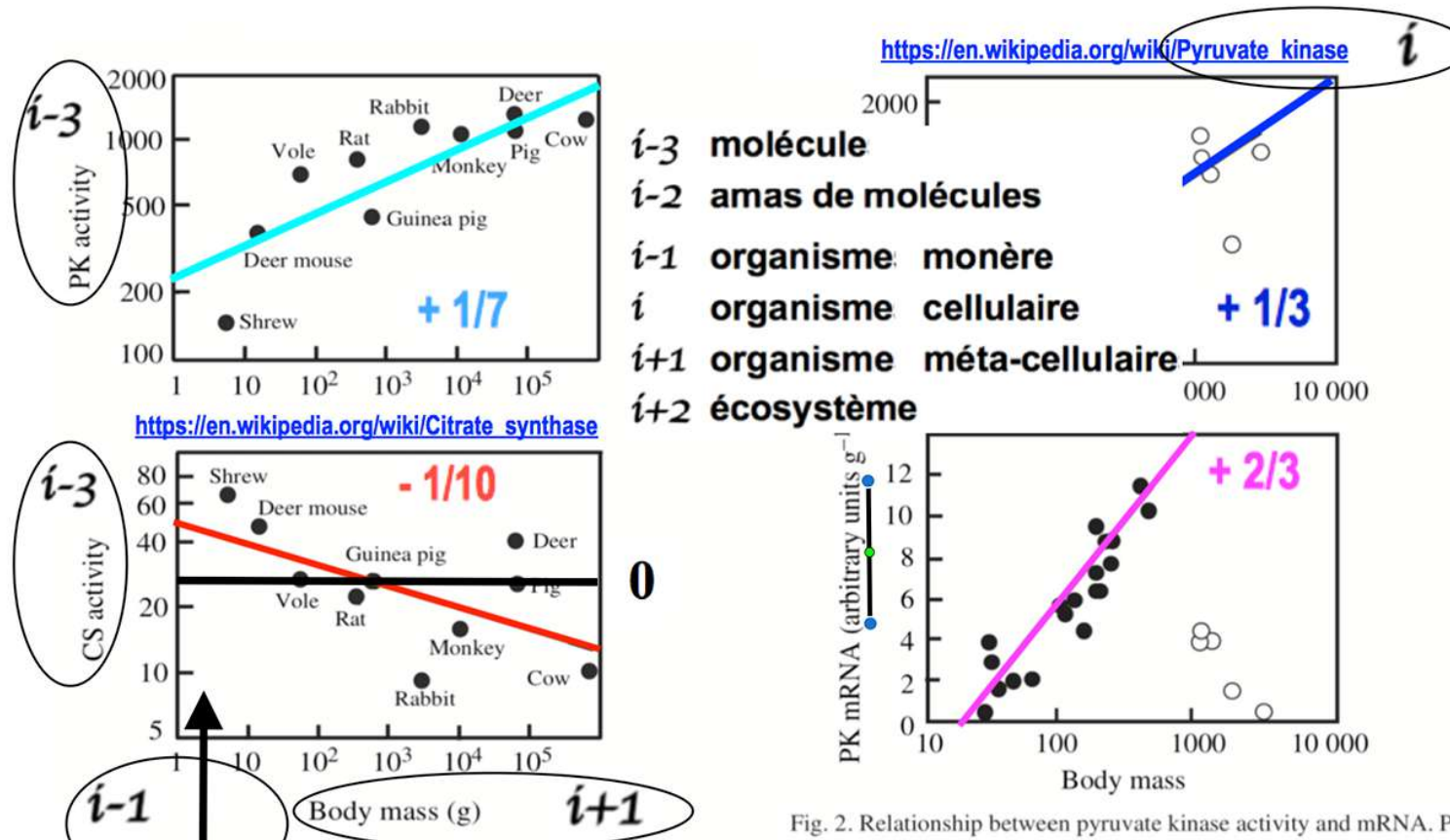


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muscle bioenergetic gene expression: allometric scaling relationships of glycolytic and oxidative enzymes

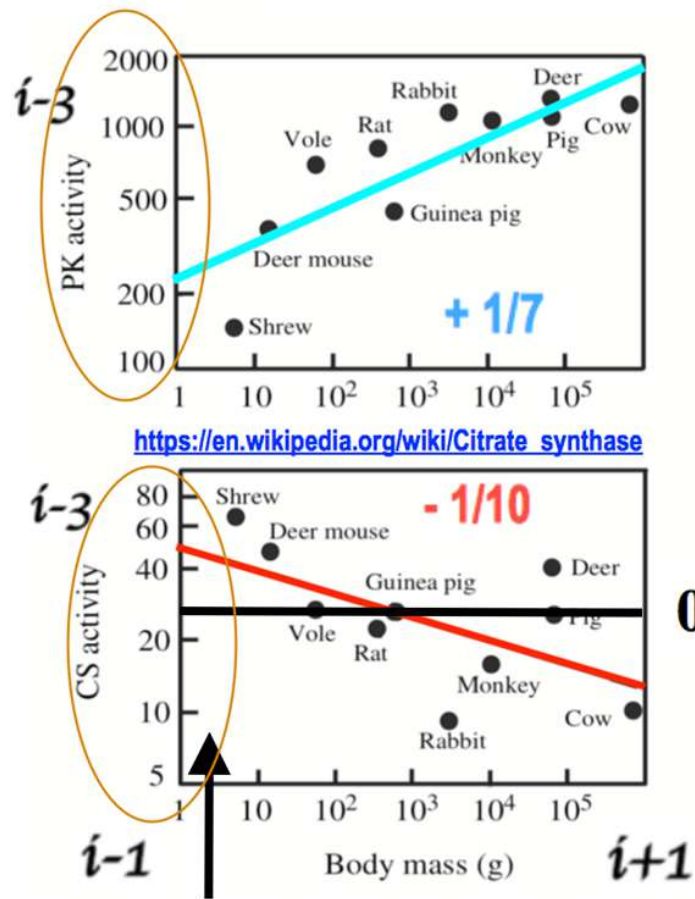


Fig. 1. Mitochondrial and glycolytic enzyme activities in muscles of mammals (adapted from Emmett and Hochachka, 1981). PK, pyruvate kinase (U g^{-1}); CS, citrate synthase (U g^{-1}); body mass (g).

muscle bioenergetic gene expression: allometric scaling relationships of glycolytic and oxidative enzymes

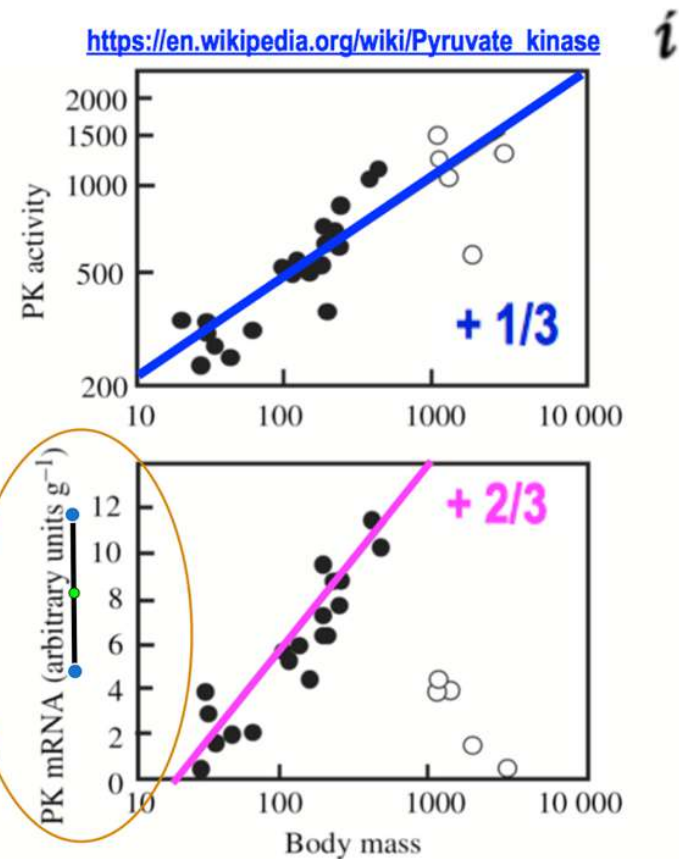
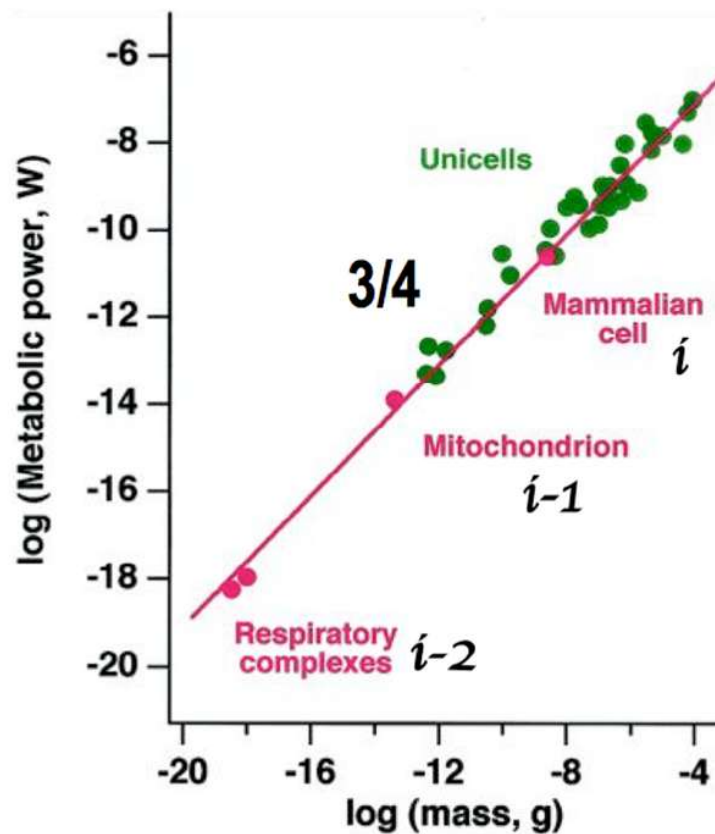


Fig. 2. Relationship between pyruvate kinase activity and mRNA. PK shows positive allometric scaling ($b=0.2$). Larger fish (open circles) are able to produce more protein from a lower amount of mRNA (adapted from Burness et al., 1999). PK, pyruvate kinase (U g^{-1}); body mass (g).



$i-9$	Planck quantum
$i-8$	cordes
$i-7$	super-cordes
$i-6$	quarks
$i-5$	particules
$i-4$	atomes
$i-3$	molécules
$i-2$	amas de molécules
$i-1$	organismes monères
i	organismes cellulaires
$i+1$	organismes méta-cellulaires
$i+2$	écosystèmes
$i+3$	biosphères
$i+4$	systèmes stellaires
$i+5$	galaxies
$i+6$	amas de galaxies
$i+7$	univers
$i+8$	Hyper-univers

Review

The origin of allometric scaling laws in biology from genomes to ecosystems: towards a quantitative unifying theory of biological structure and organization

Geoffrey B. West^{1,2,*} and James H. Brown^{1,3}

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Summary

Life is the most complex physical phenomenon in the Universe, manifesting an extraordinary diversity of form and function over an enormous scale from the largest animals and plants to the smallest microbes and subcellular units. Despite this many of its most fundamental and complex phenomena scale with size in a surprisingly simple fashion. For example, metabolic rate scales as the 3/4-power of mass over 27 orders of magnitude, from molecular and intracellular levels up to the largest organisms. Similarly, time-scales (such as lifespans and growth rates) and sizes (such as bacterial genome lengths, tree heights and mitochondrial densities) scale with exponents that are typically simple powers of 1/4. The universality and simplicity of these relationships suggest that fundamental universal principles underly much of the coarse-grained generic structure and organisation of living systems. We have proposed a set of

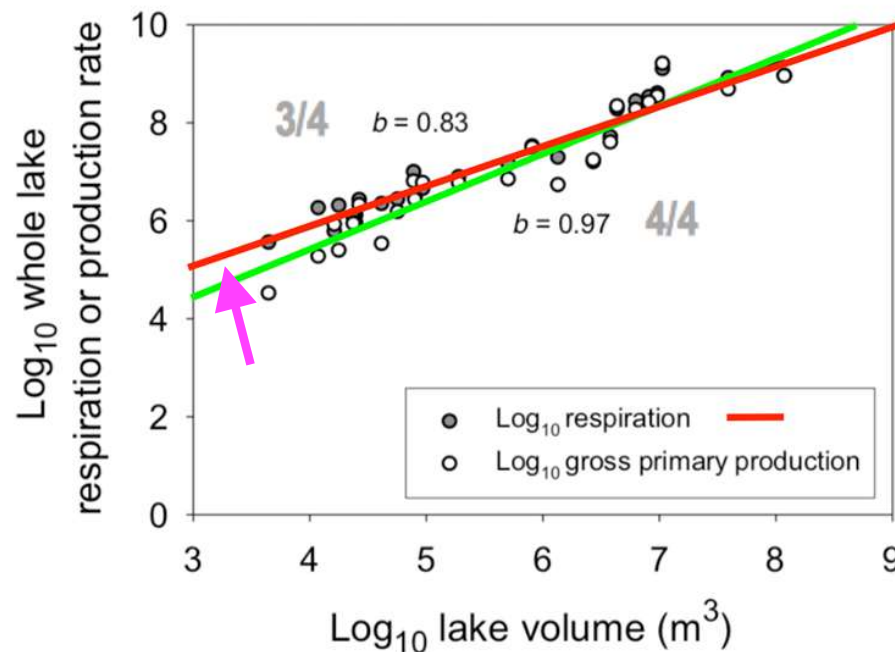
principles based on the observation that almost all life is sustained by hierarchical branching networks, which we assume have invariant terminal units, are space-filling and are optimised by the process of natural selection. We show how these general constraints explain quarter power scaling and lead to a quantitative, predictive theory that captures many of the essential features of diverse biological systems. Examples considered include animal circulatory systems, plant vascular systems, growth, mitochondrial densities, and the concept of a universal molecular clock. Temperature considerations, dimensionality and the role of invariants are discussed. Criticisms and controversies associated with this approach are also addressed.

Key words: allometry, quarter-power scaling, laws of life, circulatory system, ontogenetic growth.

Scaling of total community respiration (R , $\text{mmol}\cdot\text{O}_2\cdot\text{d}^{-1}$) and gross primary production (GPP , $\text{mmol}\cdot\text{O}_2\cdot\text{d}^{-1}$) with total water volume (V , m^3) above the thermocline in 25 lakes from Wisconsin and Michigan (data from [319]). R and GPP were estimated by multiplying measured rates per m^3 times V . The b values are scaling exponents (log-log slopes); and the lines refer to least squares regression equations (solid line: $\log_{10}R = 2.529 (\pm 0.568 \text{ 95\% CI}) + \log V(0.833 (\pm 0.99))$, $r = 0.964$, $p < 0.00001$; dashed line: $\log_{10}GPP = 1.553 (\pm 0.778) + \log V(0.966 (\pm 0.136))$, $r = 0.951$, $p < 0.00001$).

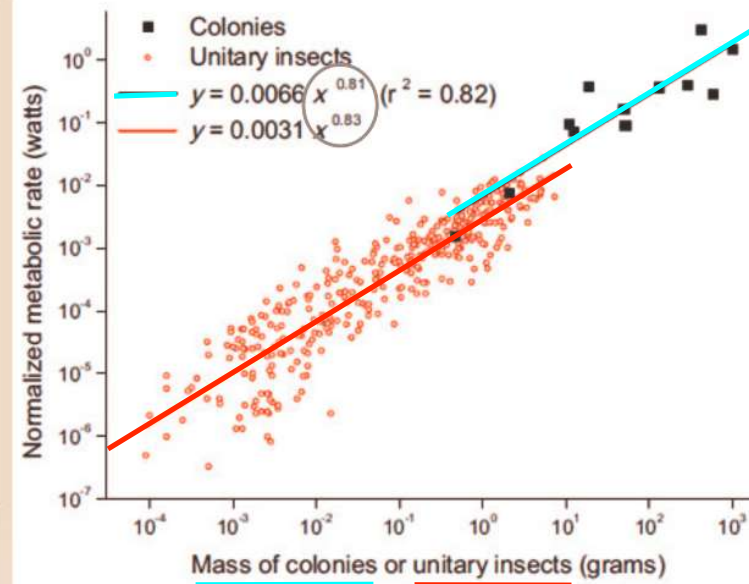
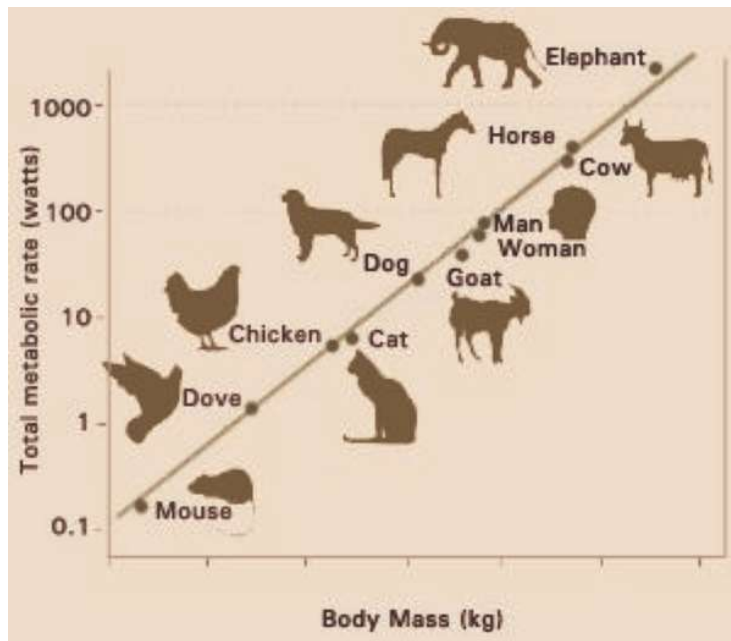
$$\log a^n = n \cdot \log a$$

$$\log a + \log b = \log a \cdot b$$



production - consommation
= énergie disponible
= masse $^{1/4}$

3/4

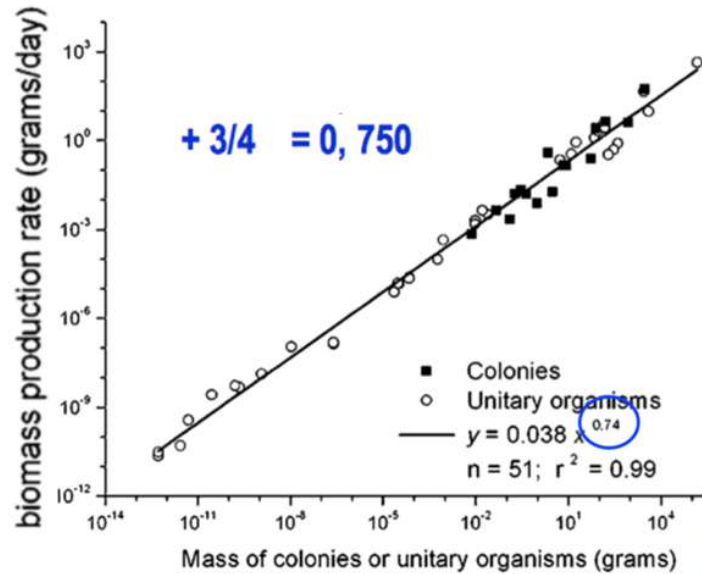


production < consommation

production = masse^{+0,75}

$$X1 = +0,75 X2$$

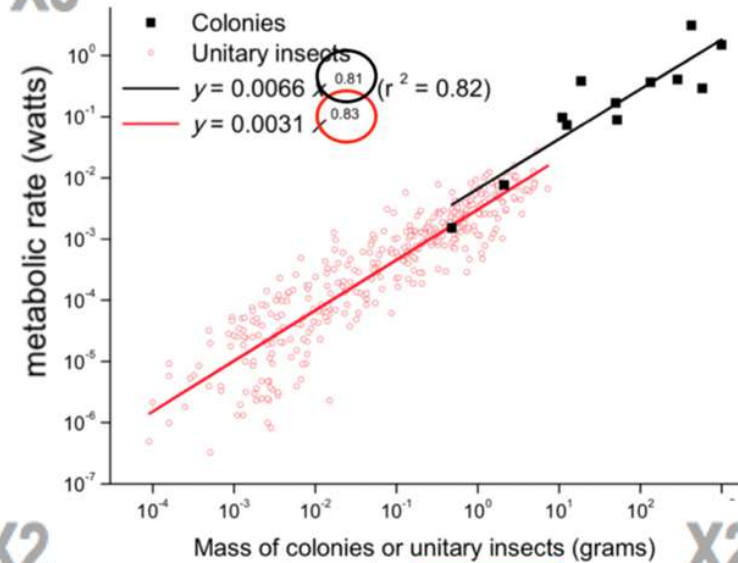
X1



consommation = masse^{+0,82}

$$X3 = +0,82 X2$$

X3



$$+ 3/4 = 0,750$$

$$+ 4/5 = 0,800$$

$$+ 5/6 = 0,833$$

$$+0,75 < +0,82$$

X2

X2

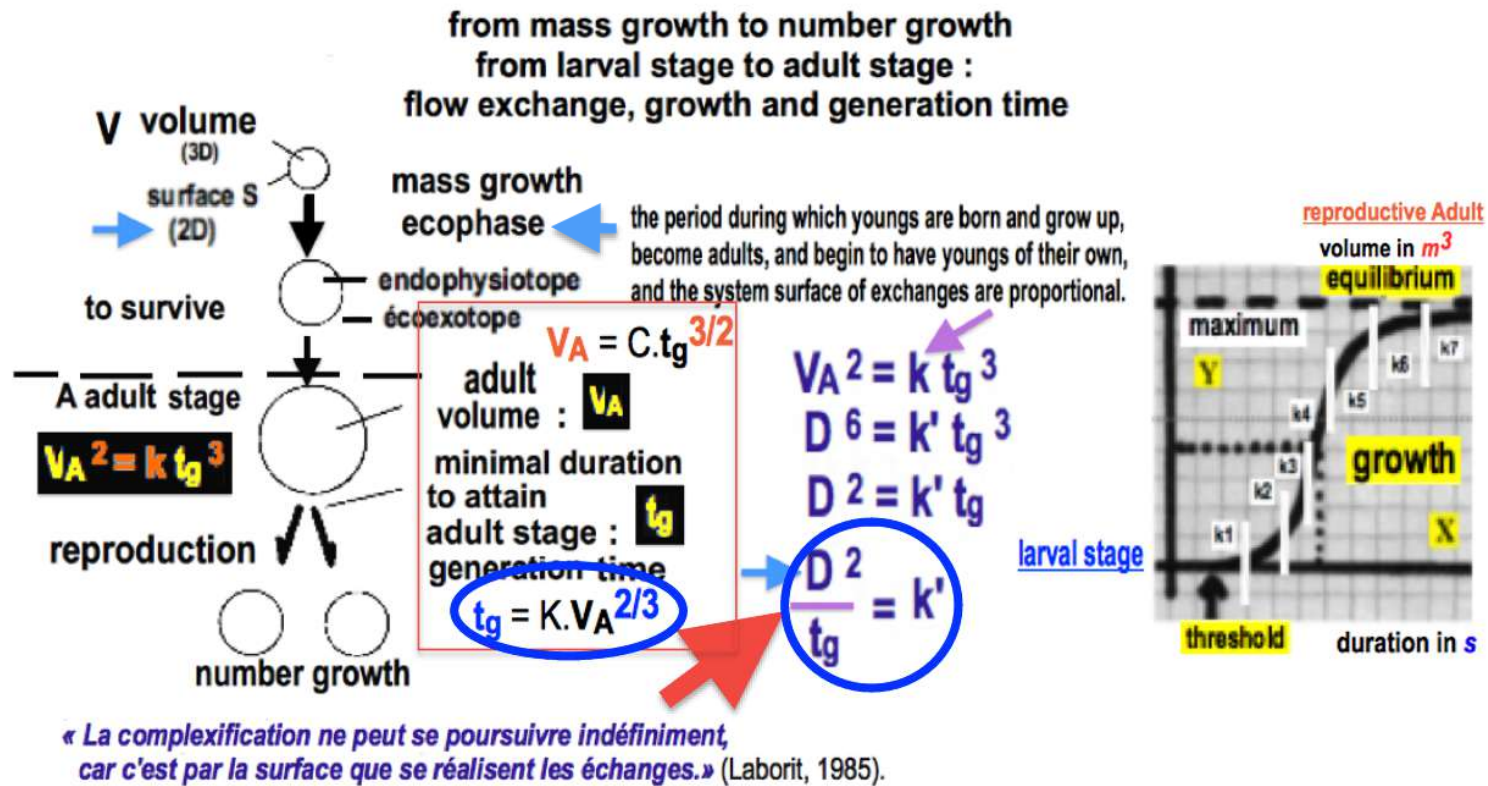
Apparently, the mysterious "3/4 law of metabolism" -- proposed by Max Kleiber in 1932, printed in biology textbooks for decades, explained theoretically in *Science* in 1997 and described in a 2000 essay in *Nature* as "extended to all life forms" from bacteria to whales -- is just plain wrong.

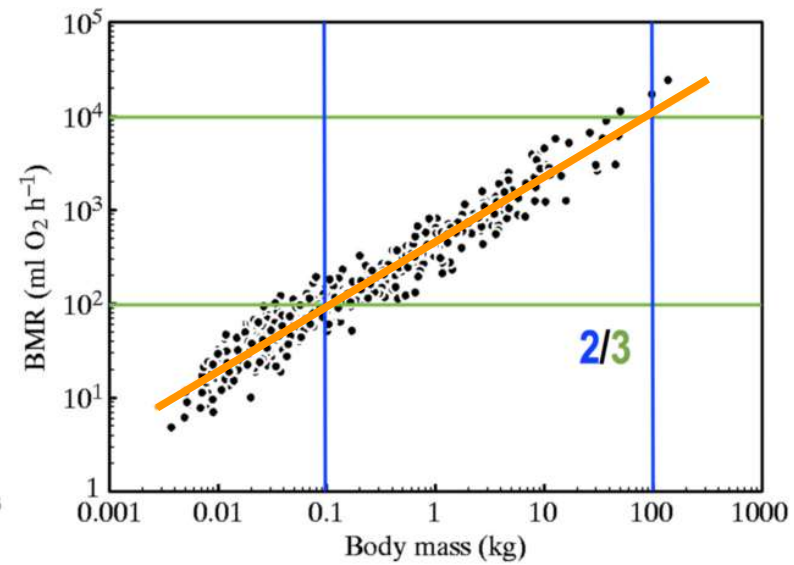
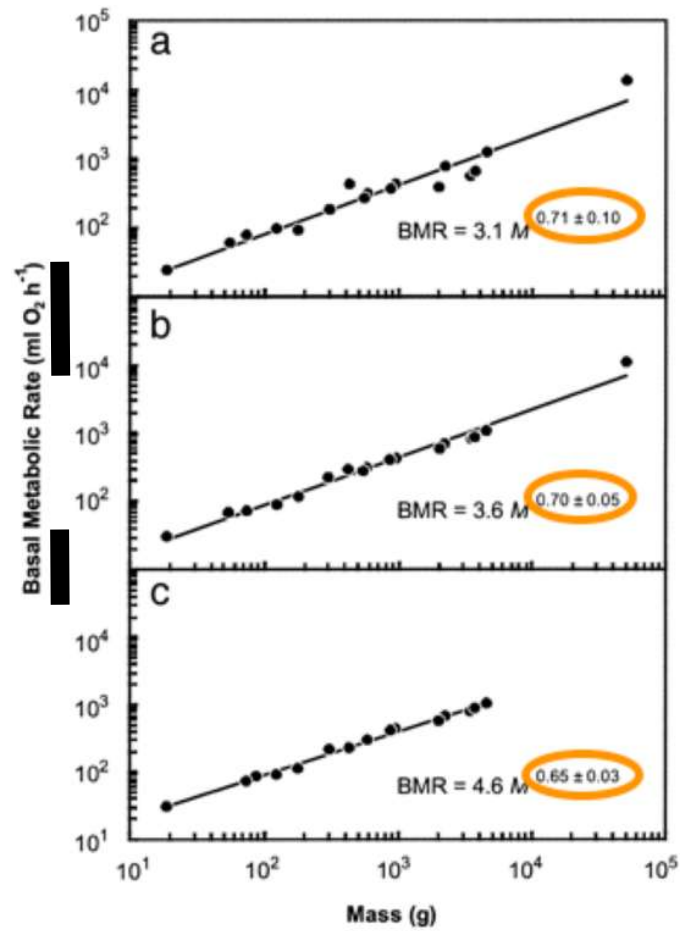
"Actually, it's two-thirds," says University of Vermont mathematician Peter Dodds. His paper in the January 29 edition of *Physical Review Letters* helps overturn almost eighty years of near-mystical belief in a 3/4 exponent used to describe the relationship between the size of animals and their resting metabolism.

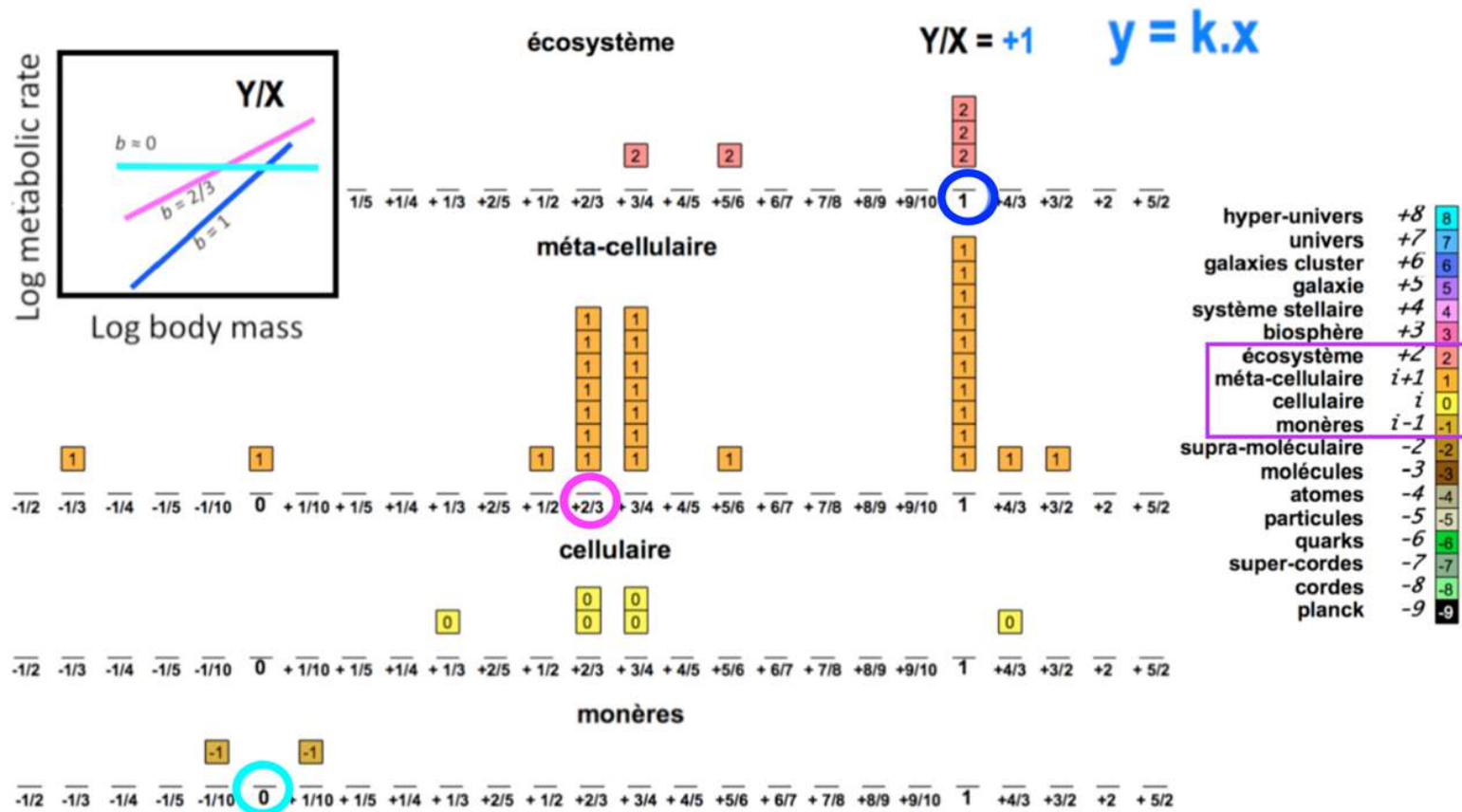
Two-thirds or three-quarters?

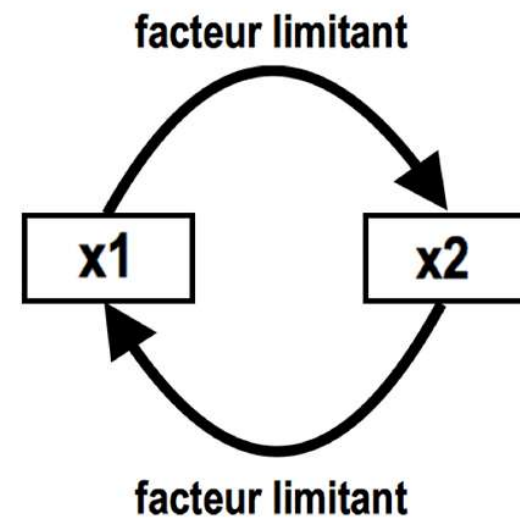
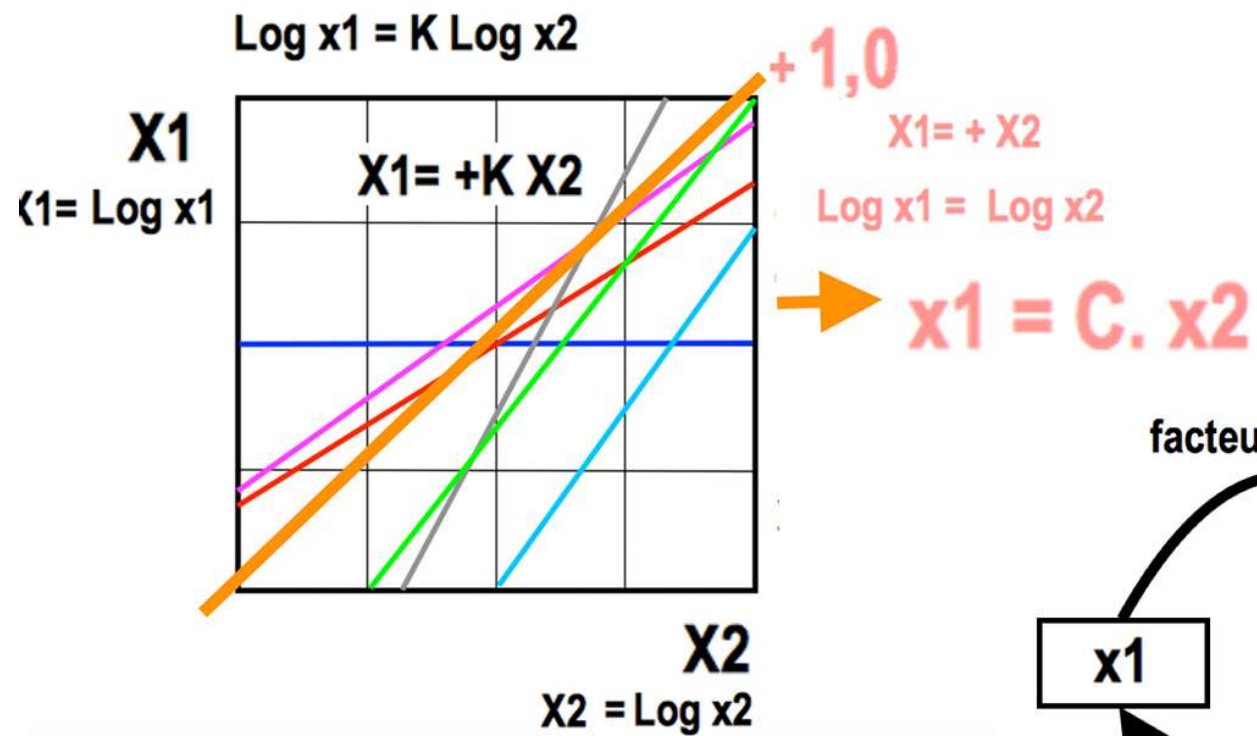
To understand the debate between 2/3 and 3/4, assume a spherical cow. "That's what a physicist would do," Dodds says, laughing. Basic geometry shows that the surface area of this difficult-to-milk creature would increase as the square of its radius while the volume would increase as the cube of the radius. In other words, the exponent that describes the ratio of surface area to volume is 2/3.

Associations for the Reciprocal and Mutual Sharing of Advantages and DisAdvantages



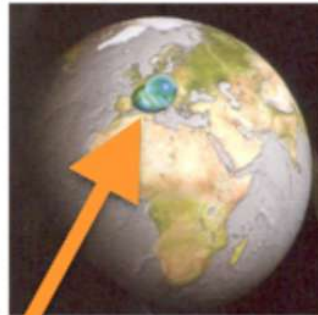






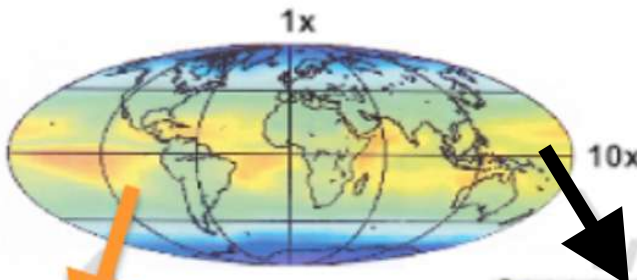
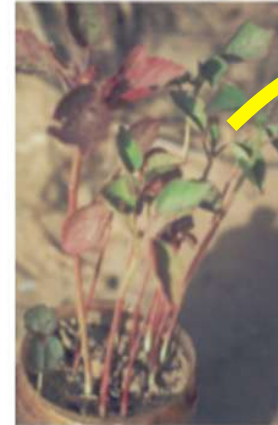


EcoExoTope & EndoPhysioTope

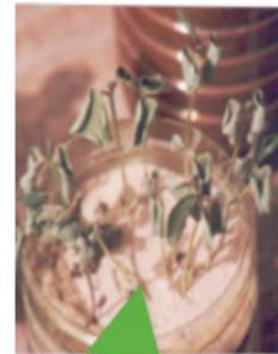
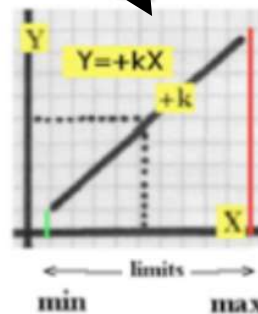


all the water of the earth

Limitations



solar energy
hosting capacity



capacity to be hosted

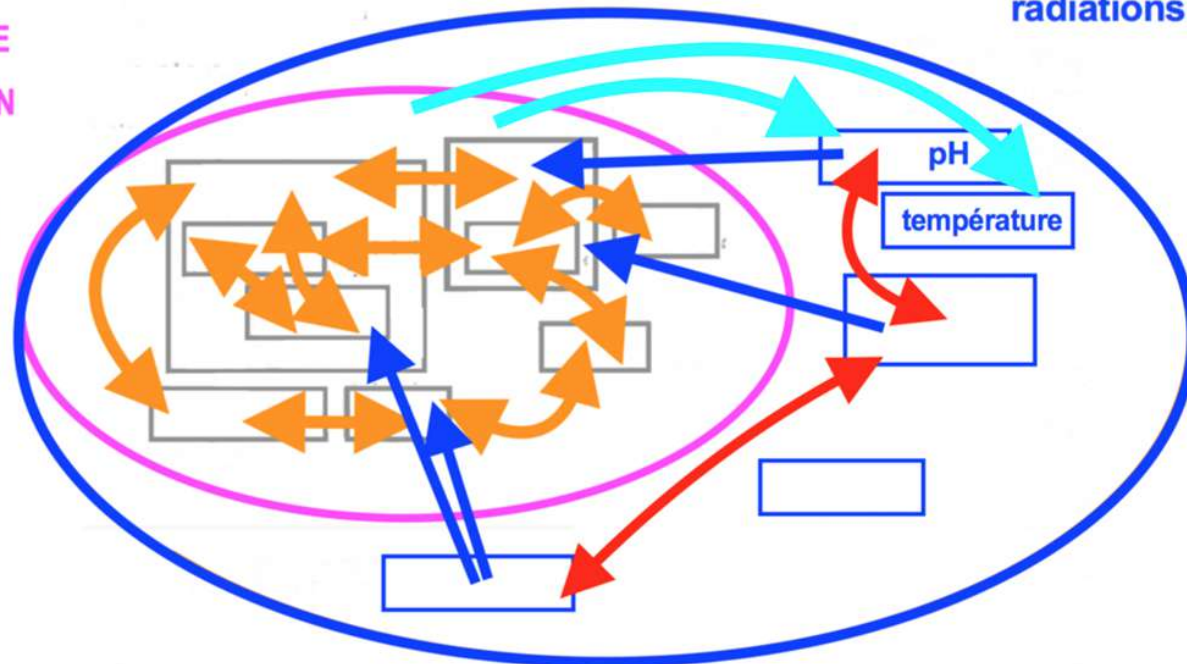
endophysiotope

SURVIVRE

1. La MOBILISATION de la MATIÈRE et de l'ÉNERGIE **MÉTABOLISME**
2. La CROISSANCE en masse **MASSE**
2. La CROISSANCE en nombre **NOMBRE**
3. La RÉPONSE à des STIMULATIONS **RÉACTION**
4. L'ORGANISATION interne dans l'espace et dans le temps **ESPACE TEMPS**
5. L'INTÉGRATION externe
6. Le MOUVEMENT
- pour **SE SURVIVRE**
7. La REPRODUCTION

écoexotope

pH
température
radiations



Scaling relationships for angiosperms of leaf nitrogen concentration N_L and leaf phosphorus concentration P_L in relation to SLA for all data pooled, for plants grouped by life form and for three biomes. (All relations were significant ($p < 0.0001$). All equations were fit using the log-log version of the equation: $Y = \beta X^\alpha$. Reduced major axis slopes (exponents) are shown, as well as the lower and upper 95% CI of the exponent, and r^2 . n , number of observations. Significant differences ($p < 0.05$) in exponents among groups (for appropriate contrasts separated by blank lines) are shown by the lack of shared letters.)

plant group	n	N_L			P_L		
		exponent	low CI, high CI	r^2	exponent	low CI, high CI	r^2
all	1819	3/4	0.73, 0.79	0.54	1,14	1.08, 1.12	0.45
<i>angiosperm</i>							
graminoid	34	0.59 a	0.41, 0.85	0.52	1.17 a	0.81, 1.69	0.51
forb	22	0.72 a	0.33, 1.56	0.34	1.01		ns
shrub	535	0.83 a	0.76, 0.89	0.55	1.19 a	1.08, 1.31	0.45
trees	1112	0.75 a	0.70, 0.80	0.45	1.17 a	1.09, 1.26	0.39
<i>biomes</i>							
temperate	449	0.67 a	0.59, 0.76	0.35	0.95 a	0.80, 1.12	0.24
Mediterranean	321	0.93 b	0.83, 1.03	0.54	1.41 b	1.26, 1.58	0.49
moist tropical	950	0.78 a	0.71, 0.85	0.34	1.21 ab	1.08, 1.35	0.24

endophysiotope

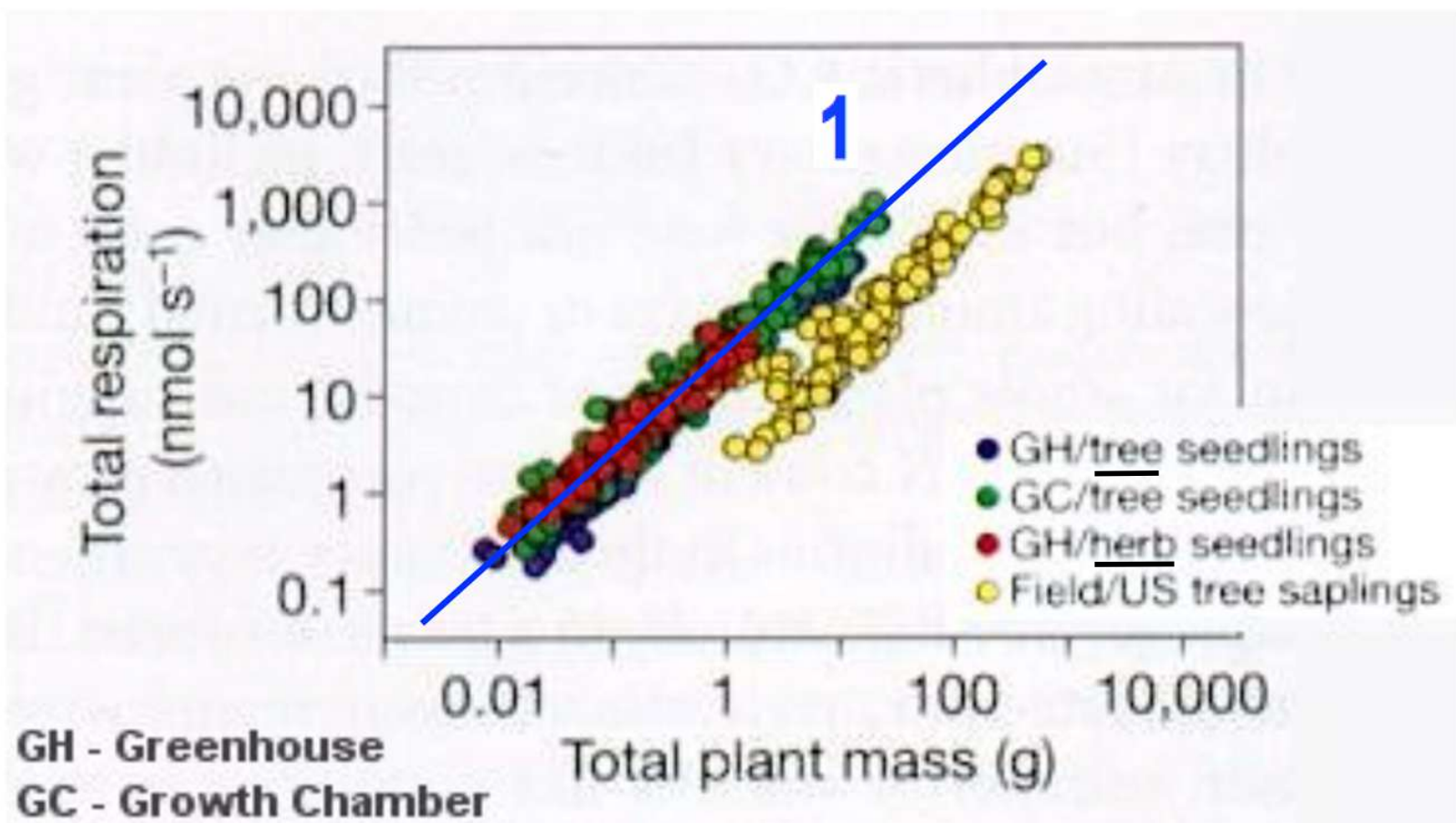
écoexotope

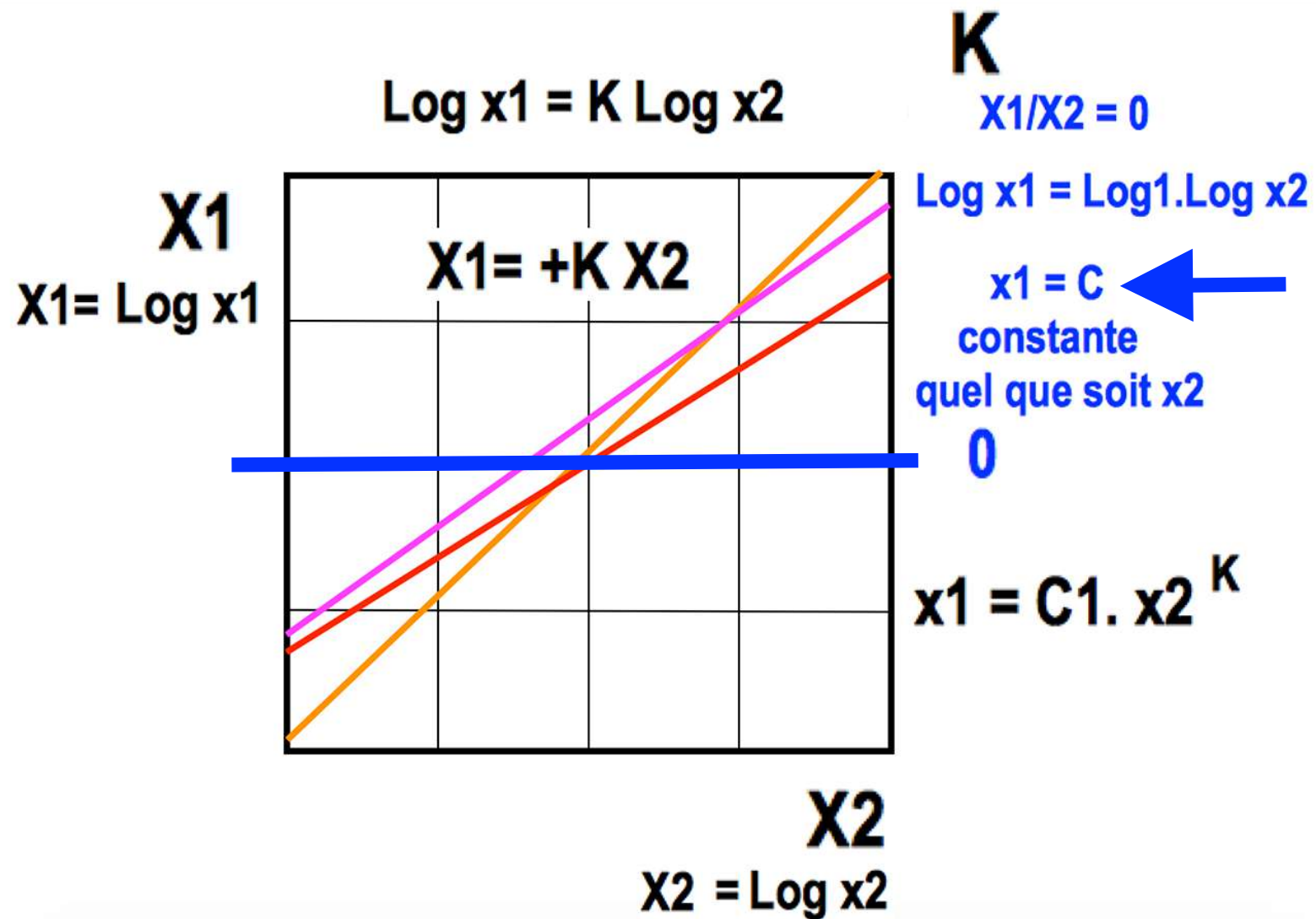
3/4

2/3

1

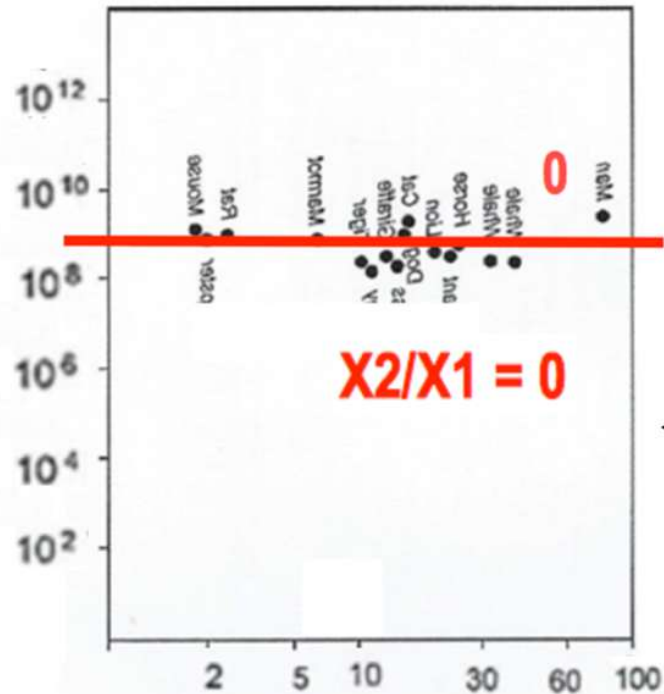
3/2





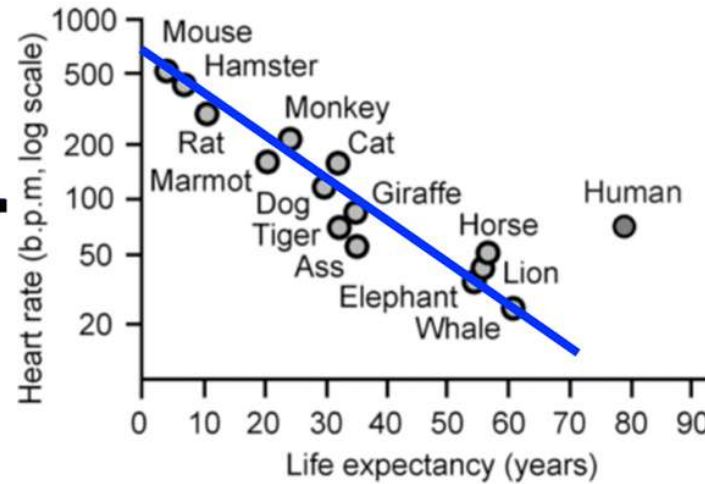
X2

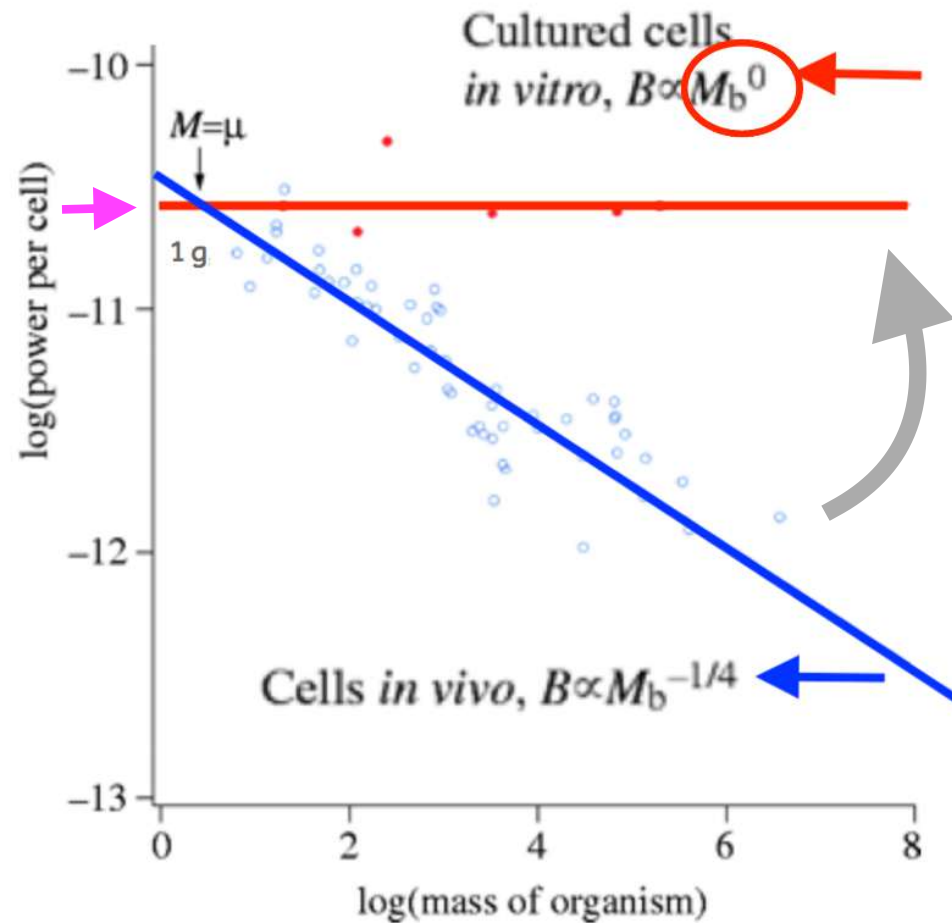
Nombre de battements cardiaque
cardiaque par vie



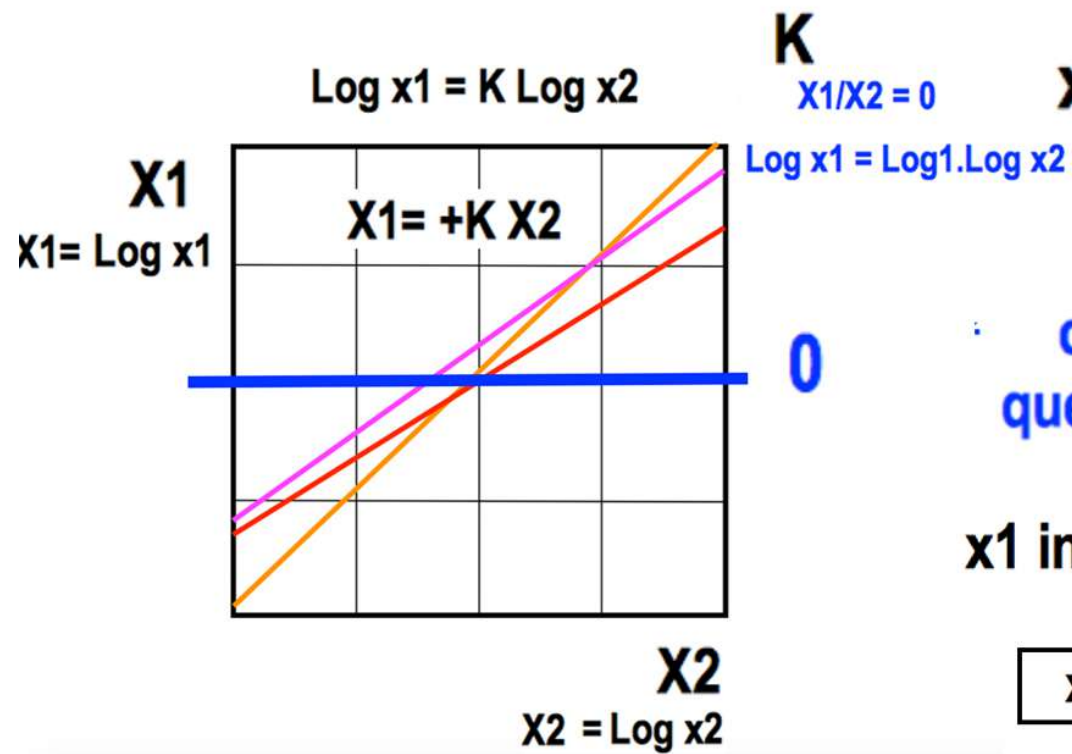
durée de vie X1
(en années)

X2 = cste





Metabolic rates (in W) of mammalian cells *in vivo* (blue line) and cultured *in vitro* (red line) plotted as a function of organism mass (M_b in g) on a log-log scale. While still in the body and constrained by vascular supply networks cellular metabolic rates scale as $M_b^{-1/4}$. When cells are removed from the body and cultured *in vitro*, their metabolic rates converge to a constant value predicted by theory (West et al., 2002b). The two lines meet at the mass of the smallest mammal



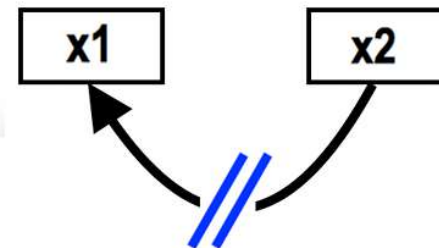
K

$$X_1/X_2 = 0$$

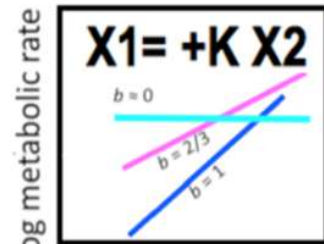
$$x_1 = C_1 \cdot x_2^0$$

$x_1 = C$ ←
constante
quel que soit x_2

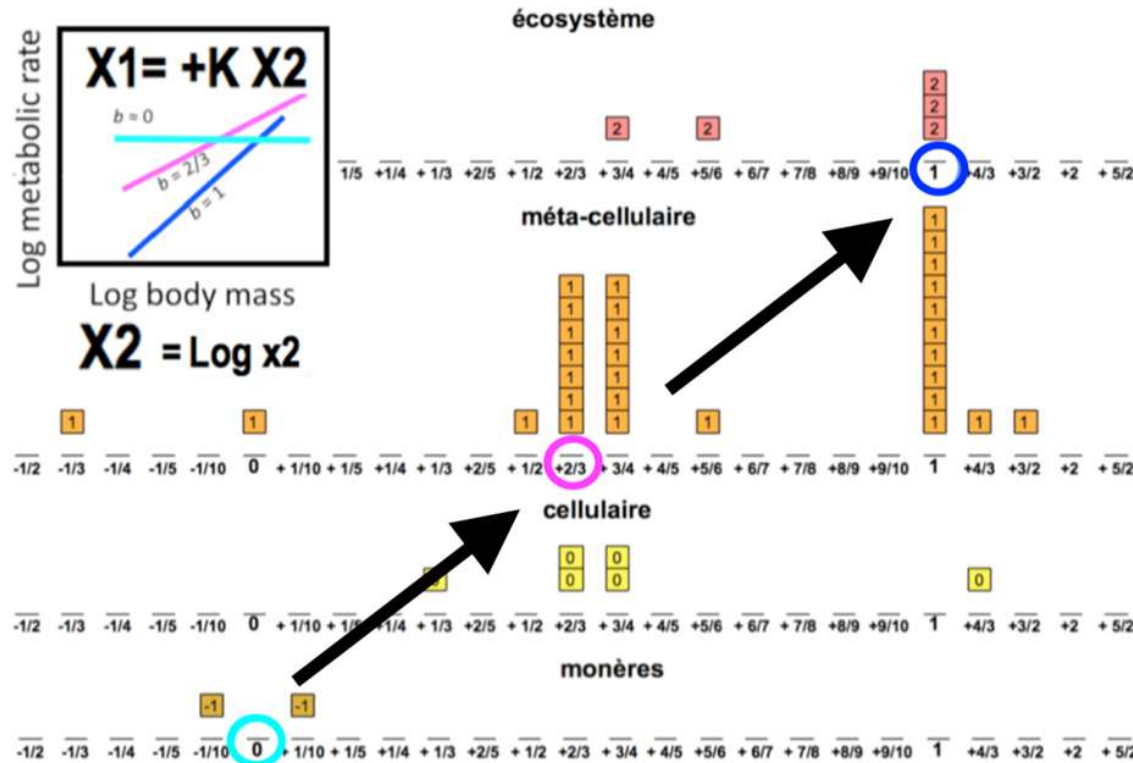
x_1 indépendant de x_2



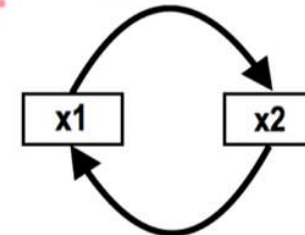
X1 $\text{Log } x1 = K \text{ Log } x2$



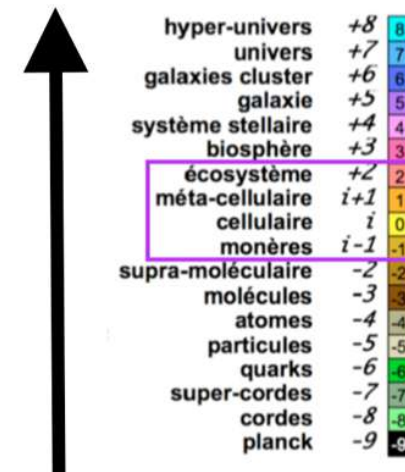
Log body mass
X2 = Log x2



$x1 = C \cdot x2$ facteur limitant

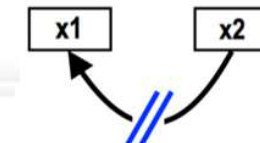


facteur limitant

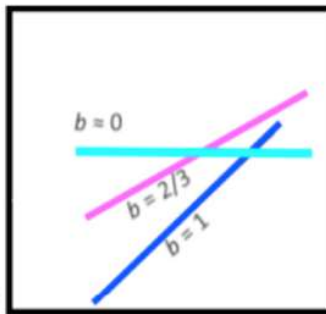


$x1$ indépendant de $x2$

$x1 = C$

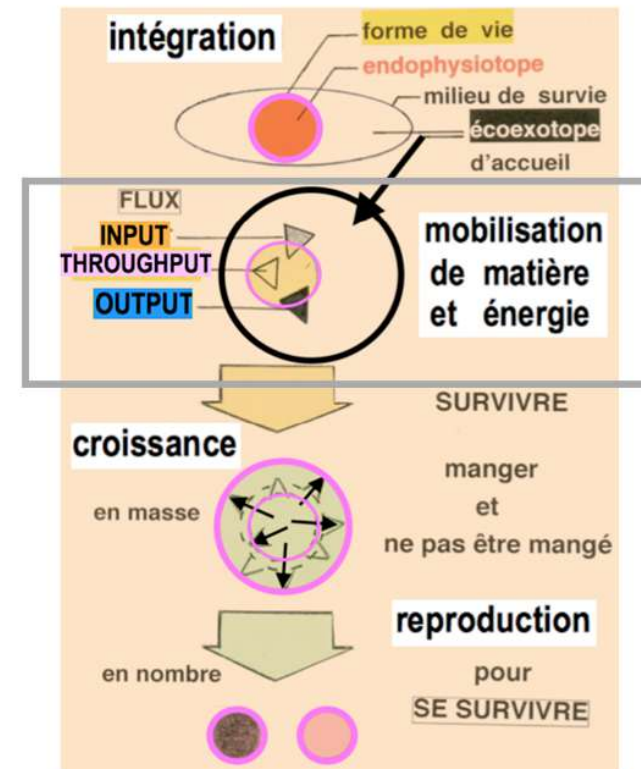


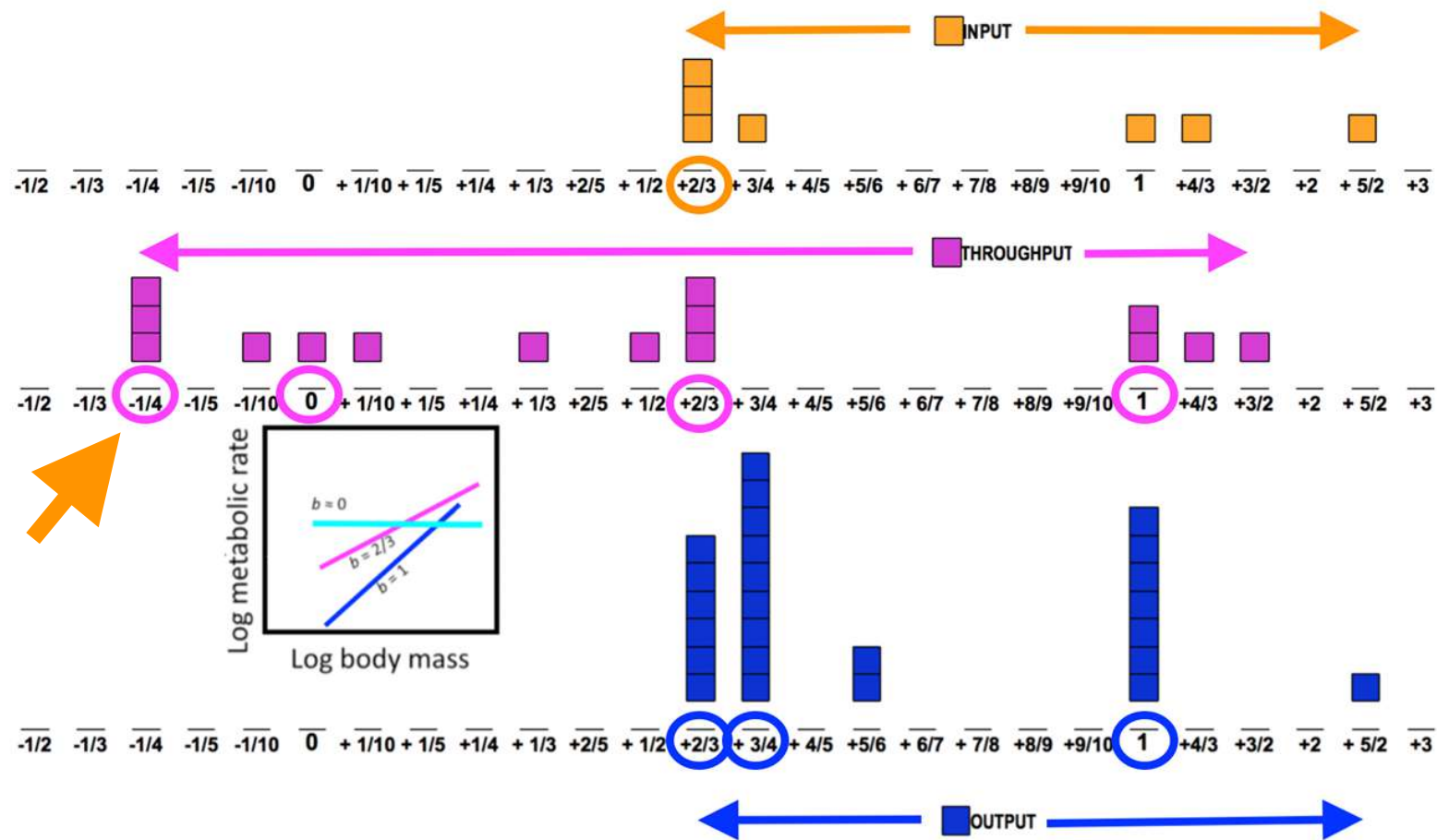
Log metabolic rate

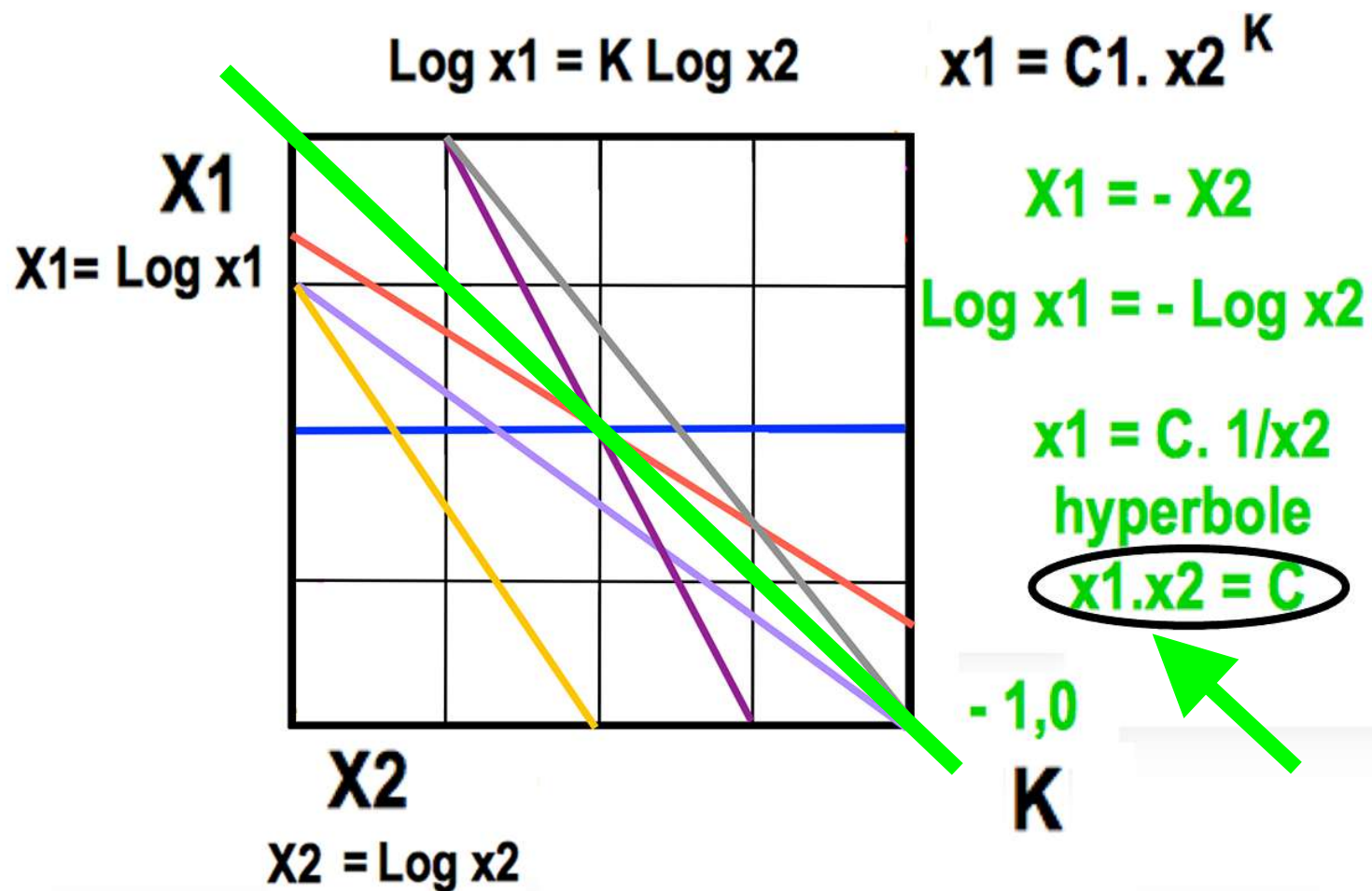


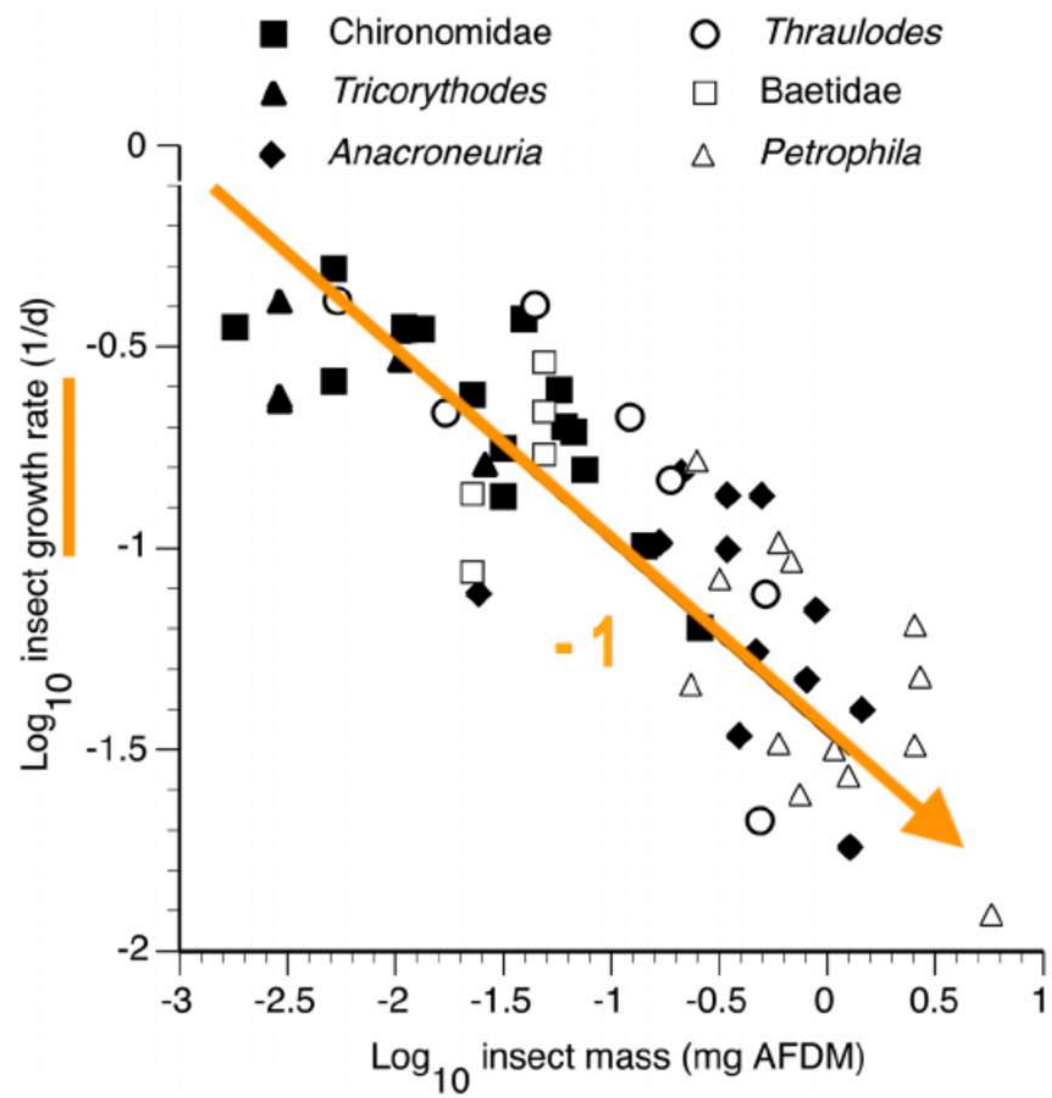
Log body mass

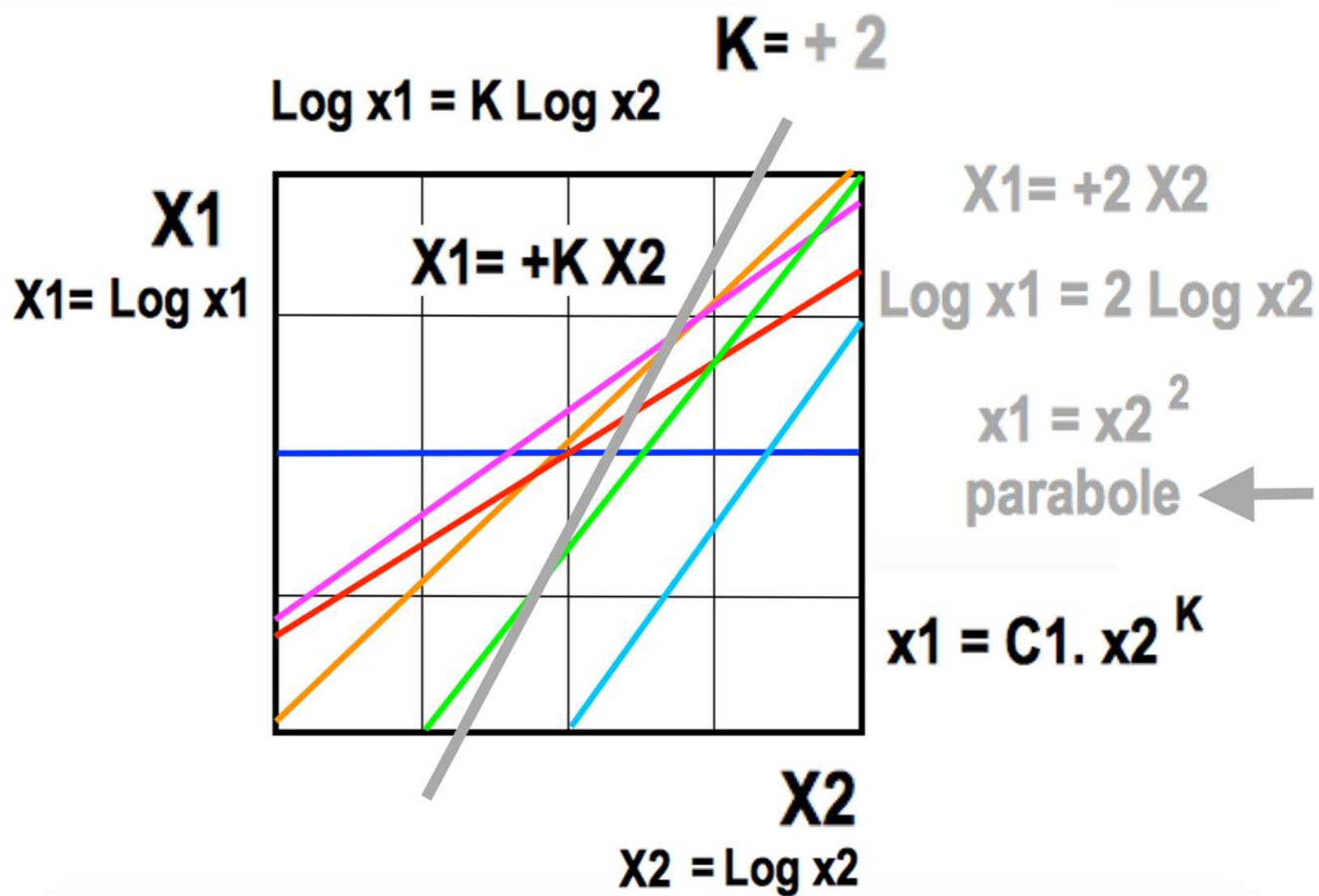
- $i-9$ Planck quantum
- $i-8$ cordes
- $i-7$ super-cordes
- $i-6$ quarks
- $i-5$ particules
- $i-4$ atomes
- $i-3$ molécules
- $i-2$ amas de molécules
- $i-1$ organismes monères
- i organismes cellulaires
- $i+1$ organismes méta-cellulaires
- $i+2$ écosystèmes
- $i+3$ biosphères
- $i+4$ systèmes stellaires
- $i+5$ galaxies
- $i+6$ amas de galaxies
- $i+7$ univers
- $i+8$ Hyper-univers







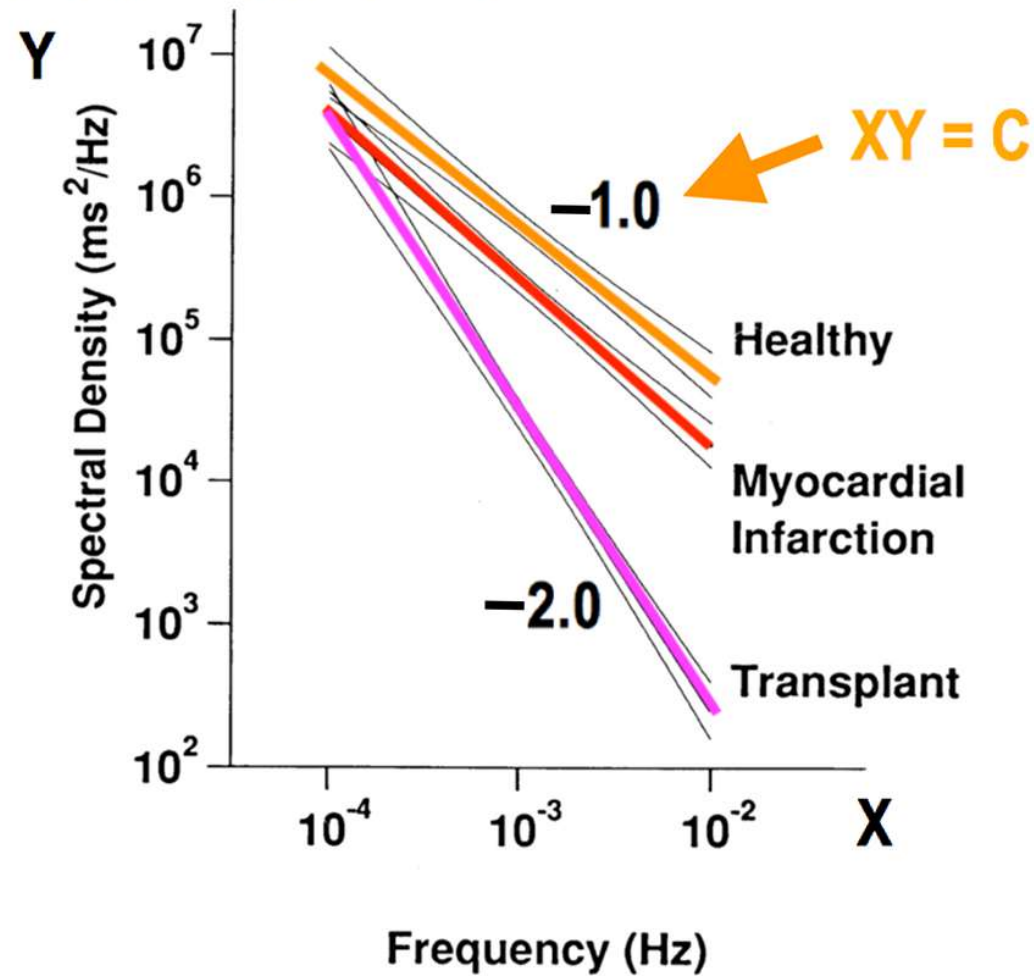




$Y/X = -1$	$\text{Log}(y) = -1 \text{ Log}(x)$	<u>$y = 1/x$ hyperbole</u>	$xy = k$
$Y/X = +1$	$\text{Log}(y) = +1 \text{ Log}(x)$	<u>$y = x$ droite</u>	$y = k.x$
$Y/X = 2$	$\text{Log}(y) = 2 \text{ Log}(x)$	<u>$y = x^2$ parabole</u>	

Regression of log(power) on log(frequency) for healthy subjects (n=274), myocardial infarction patients (n=715), and heart transplant patients (n=19)

Bigger J et al. Circulation 1996;93:2142-2151

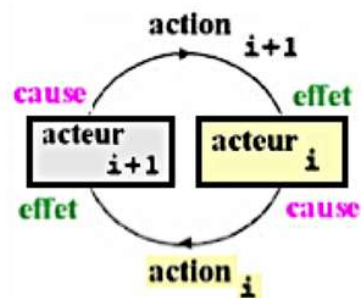
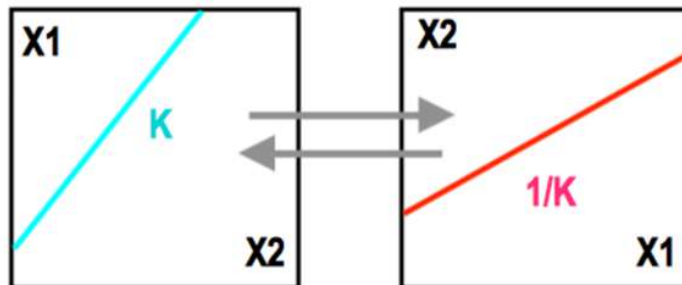


on ne peut pas privilégier,
comme cause ou effet,
un facteur plutôt qu'un autre

$$X1 = K X2 \text{ équivaut à } X2 = 1/K X1$$

$$\text{Log } x1 = K \text{ Log } x2 \quad \text{Log } x2 = 1/K \text{ Log } x1$$

$$x1 = x2^K \quad \text{ou} \quad x2 = x1^{1/K}$$



des interactions antagonistes
permettent la mise en place
d'un processus de rétroaction négative

$$X1 = +K X2$$

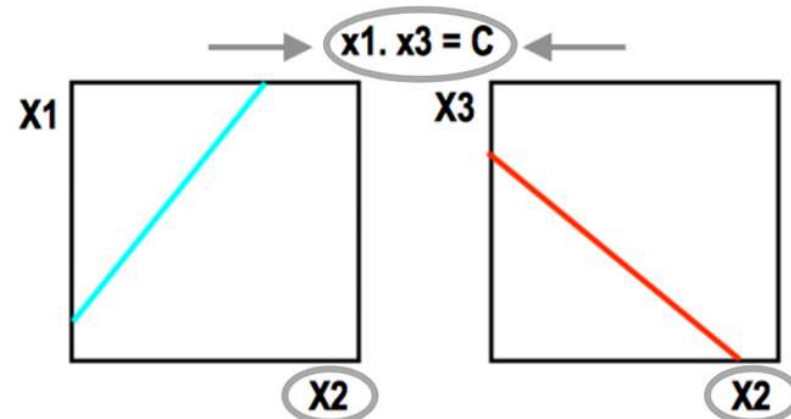
$$X3 = -K X2$$

$$\text{Log } x1 = K \text{ Log } x2$$

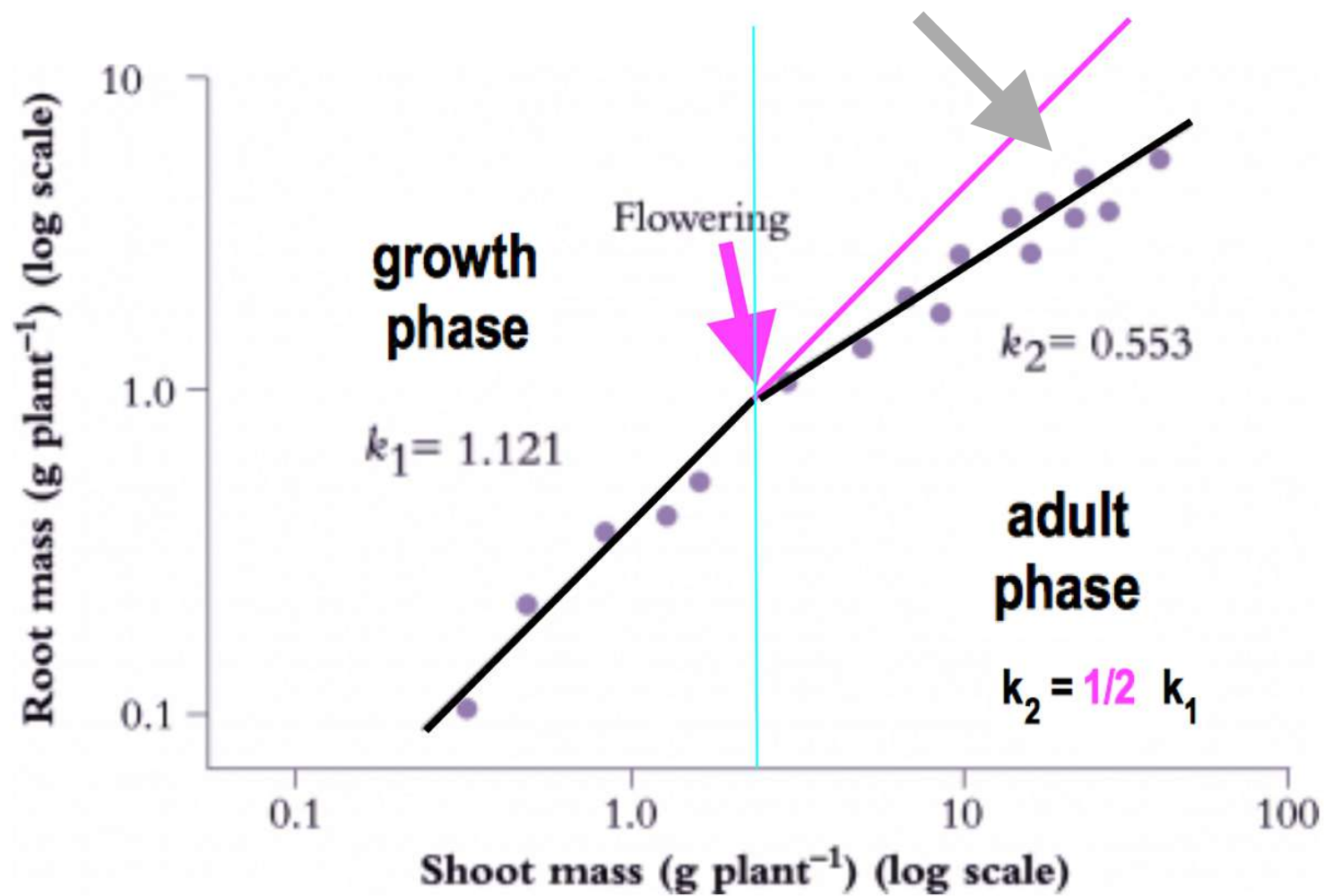
$$\text{Log } x3 = -K \text{ Log } x2$$

$$x1 = C1. x2^K$$

$$x3 = C3. x2^{-K}$$



quand l'un croît,
l'autre décroît



Two-Third power law for curved movements provides a strict constraint on optimal control models

Dongsung B Huh (huh@salk.edu), Terrence J Sejnowski (terry@salk.edu)

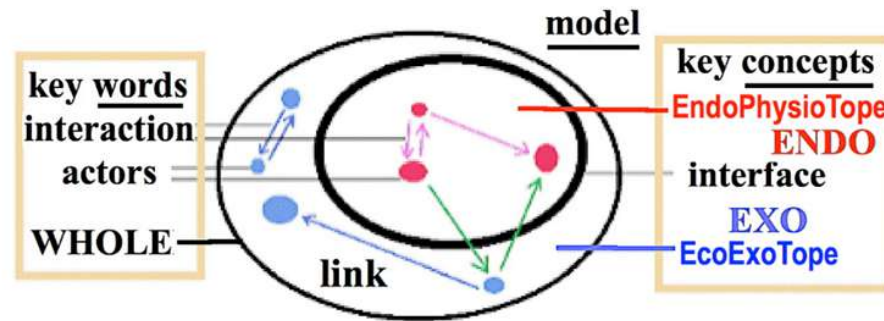
One of the interesting properties of curved hand movements is that the speed v is related to the curvature k through a power law: $v(t) \propto k(t)^{-1/3}$, called the $\frac{2}{3}$ power law (because the angular speed is proportional to $k^{2/3}$).

Several models have been proposed to explain the origin of the $\frac{2}{3}$ power law from optimality principle: the minimum jerk models (Richardson and Flash 2002; Todorov and Jordan 1998; Viviani and Flash 1995; Wann et al. 1988), and the minimum-variance model (Harris and Wolpert 1998), to name a few. It seemed the $\frac{2}{3}$ power law was an easily explainable phenomenon.

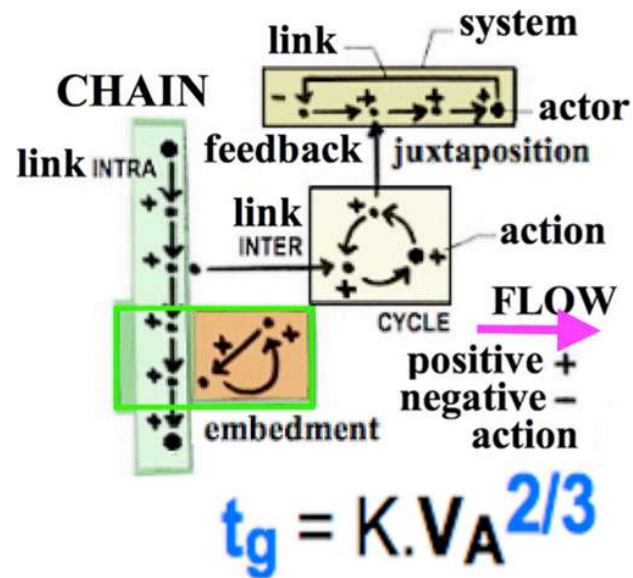
In further analyzing these models, however, we found their predictions to hold only for movement paths that have rather quickly changing curvature (such as ellipses, cloverleaf shapes and figure eights). For paths with slowly changing curvature, such as spirals, the power law is no longer $\frac{2}{3}$.

This failure can be analytically proved for the minimum jerk model. We show that the minimum jerk model in fact predicts a $\frac{1}{3}$ power law ($v(t) \propto k(t)^{-2/3}$) for paths with slowly changing curvature, and that the power law approaches $\frac{2}{3}$ for paths with fast changing curvature.

Our analysis shows that predicting the $\frac{2}{3}$ power law for a wide range of paths is a very strict criterion that narrows down the appropriate cost function. We are currently investigating a family of cost functions that satisfy this criterion.

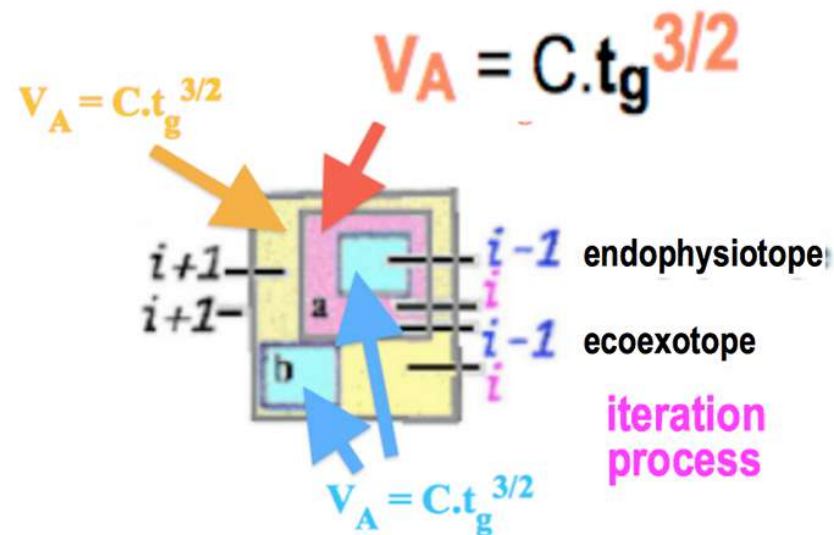


embedment & juxtaposition
of metabolic networks

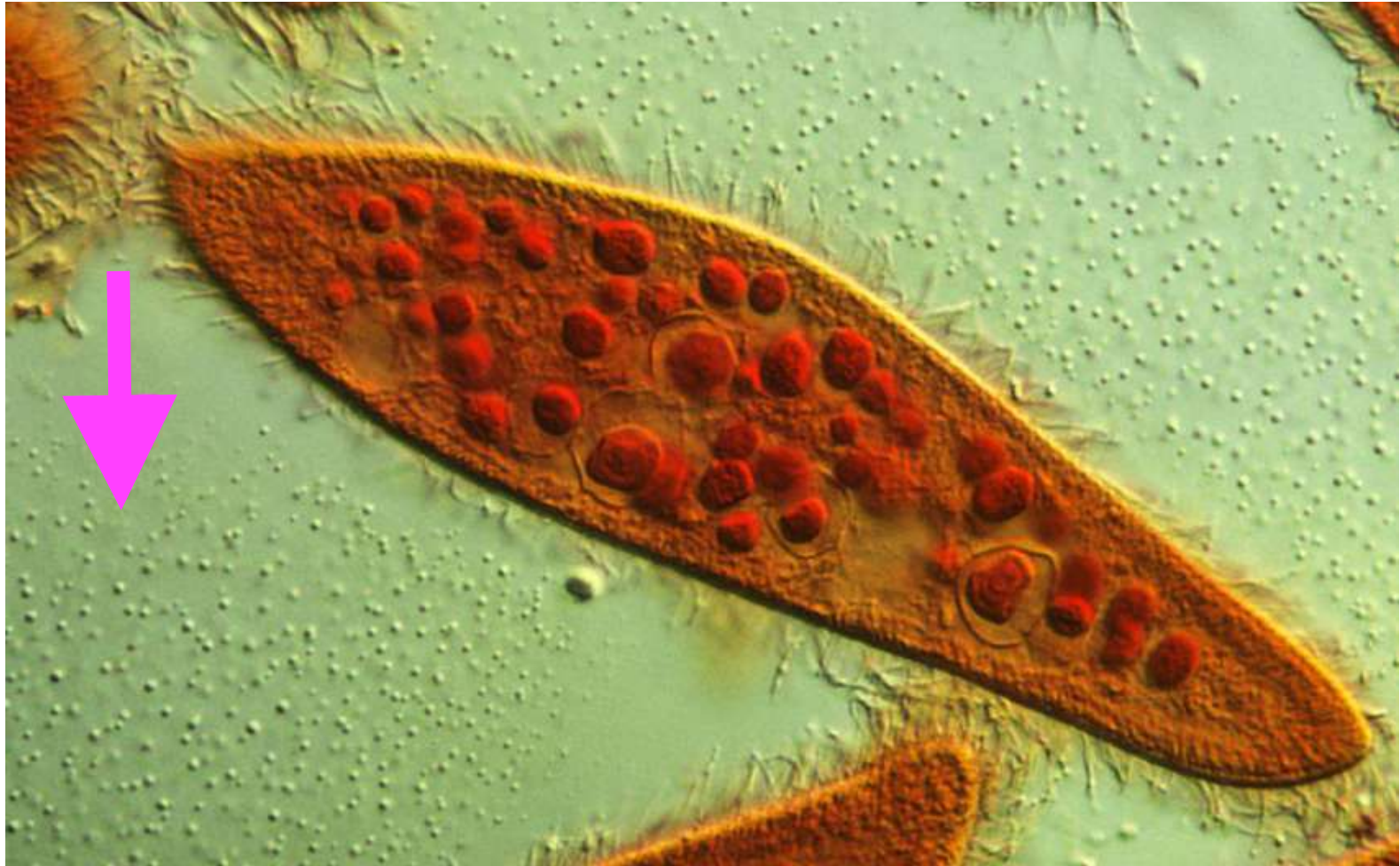


temporal flow

embedment & juxtaposition
of spaces & times



geometric fractal flow

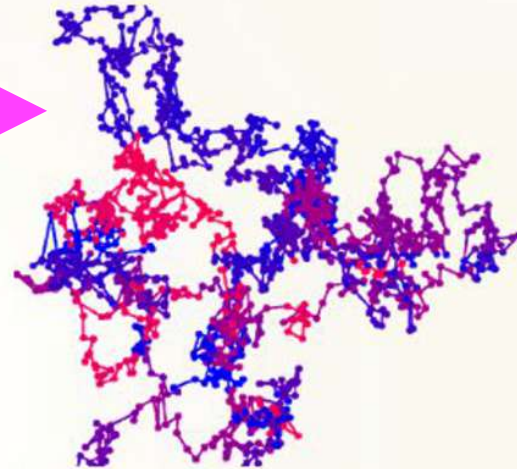


■ Random Walk

- a particle repeatedly moves in all directions

■ Brownian motion

- continuous irregular motion of individual pollen particles
- Robert Brown(1828)



On note $E_x(f(B_s))$ l'espérance de $f(B_s)$
 quand B est un Brownien issu de x . Cette quantité est égale à

$$E(f(x + B_s)) = \frac{1}{\sqrt{2\pi t}} \int_{\mathbb{R}} f(x + y) \exp\left(-\frac{y^2}{2t}\right) dy$$

où B est un Brownien issu de 0.

Soit B un mouvement Brownien, b et σ deux constantes.

Le processus **Brownien géométrique**.

$$X_t = X_0 \exp\left\{\left(b - \frac{1}{2}\sigma^2\right)t + \sigma B_t\right\}$$

est aussi appelé processus "log-normal".

Temps d'atteinte T_a est un temps d'arrêt fini

$$E(\exp -\lambda T_a) = \exp(-|a|\sqrt{2\lambda}).$$

Par inversion de la transformée de Laplace,
 on obtient la densité de T_a qui est, pour $a > 0$

$$\frac{a}{\sqrt{2\pi t^3}} \exp\left(-\frac{a^2}{2t}\right)$$

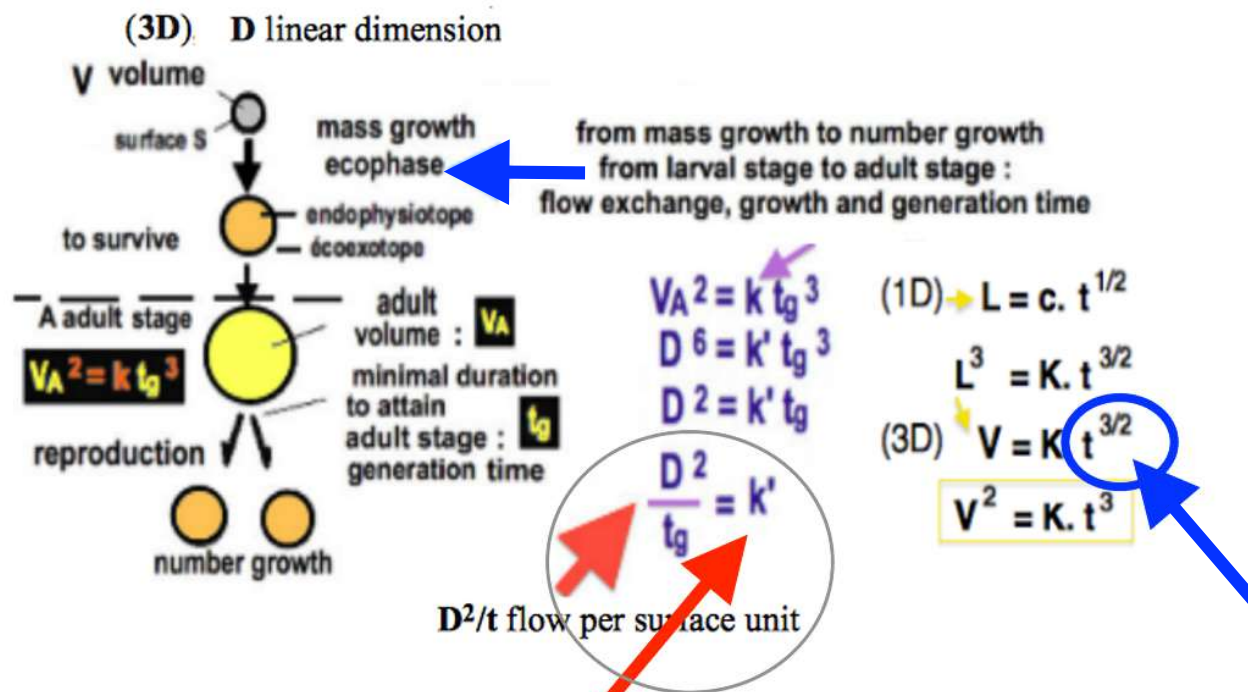


Fig. 9a Exchanges at a constant rate
t time, **V** volume (3D), **A** adult phase
D linear dimension,
 a constant exchange rate **k'**
 To survive with **mass growth** is the prerequisite for
 a life form to itself survive with **number growth**.
Just-in-time exchanges: the exchanges flow rate
 at the surface of exchange is a constant.

Fig. 9b Brownian motion
 The length of the linear movement (1D), **L**, is
 proportional to the square root of the duration of
 observation -time **t**- (Perrin 1908): **1/2** power-law.
 This is the same as: **V** obeys a **3/2** power-law of **t**,
 or $V_A^2 = K t_g^3$
 and **3/2** is just the true exponent that describes
 “the ratio of volume to surface area in a sphere”.

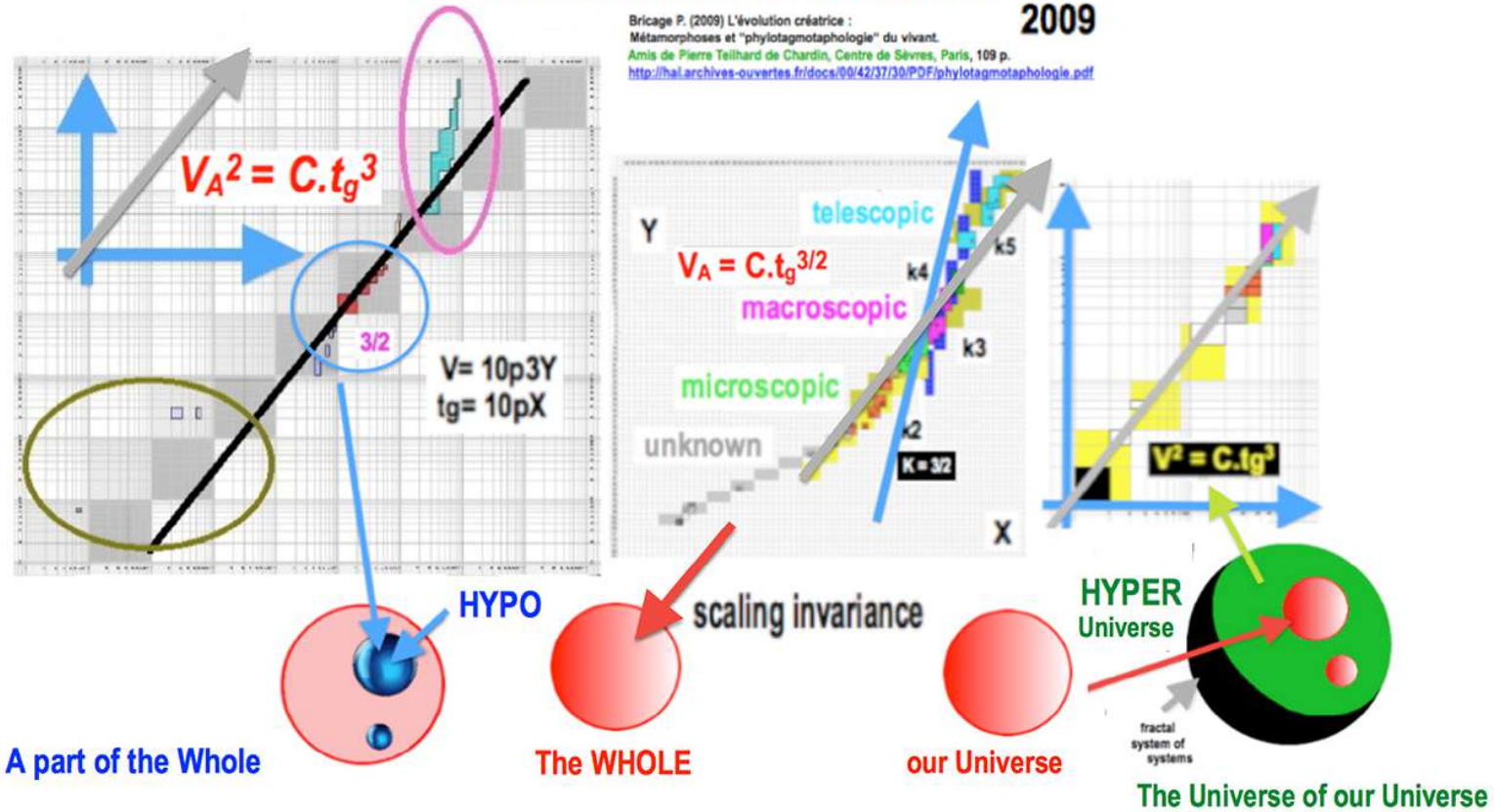
(Reproduced from Bricage 2009a, b, 2014e [Free Share-Alike Creative Commons Licence])

ASSOCIATIONS à AVANTAGES et INCONVÉNIENTS RÉCIPROQUES et PARTAGÉS

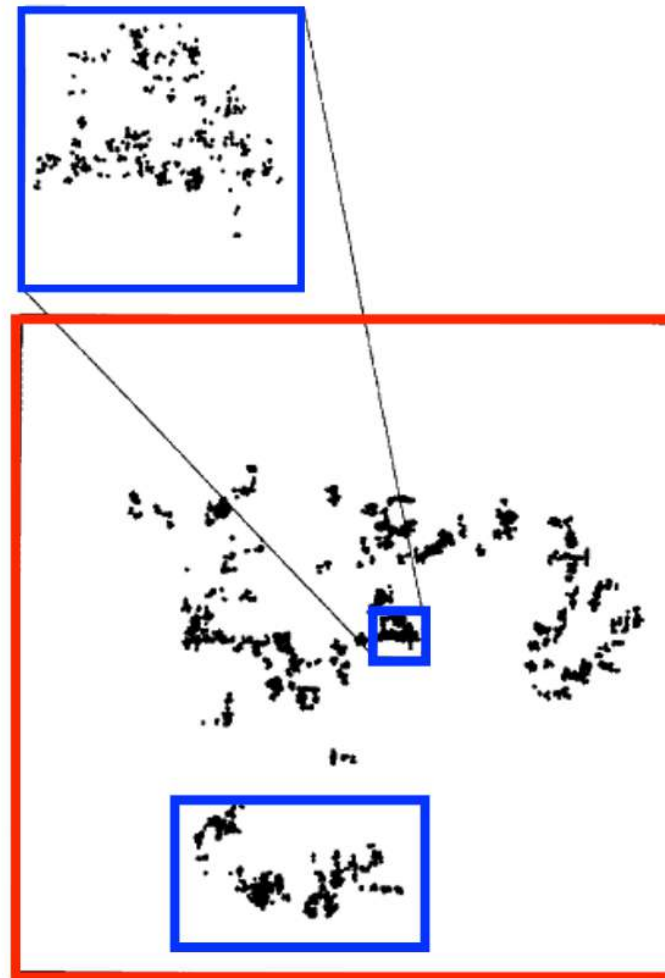
L'INVARIANCE FRACTALE du VIVANT

2009

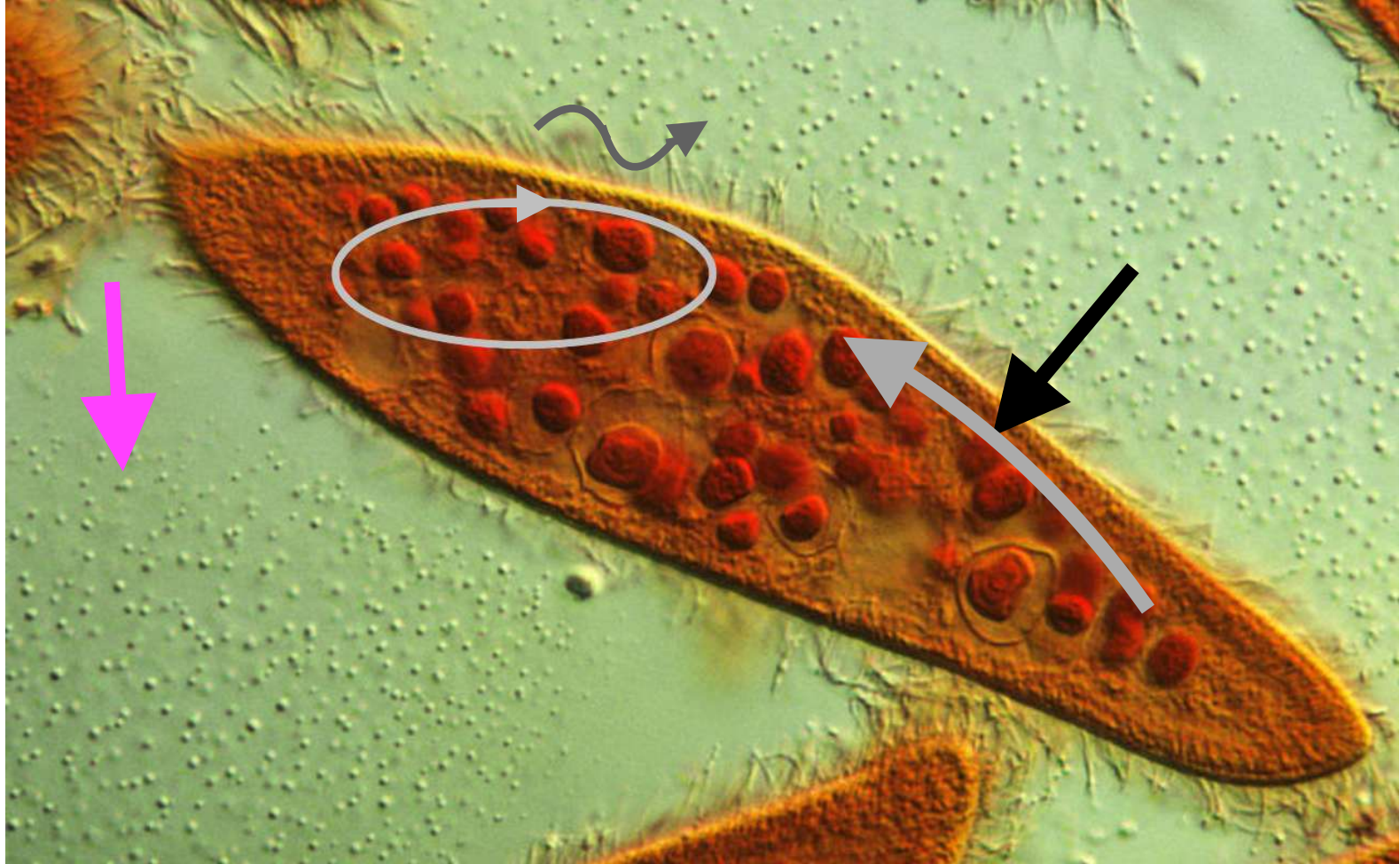
Bricage P. (2009) L'évolution créatrice :
Métamorphoses et "phylotagmotaphologie" du vivant.
Amis de Pierre Teilhard de Chardin, Centre de Sévres, Paris, 109 p.
<http://hal.archives-ouvertes.fr/docs/00/42/37/30/PDF/phylotagmotaphologie.pdf>



ASSOCIATIONS for the RECIPROCAL and MUTUAL SHARING of ADVANTAGES and DISADVANTAGES



The first 10^4 points visited by a two-dimensional random walk generated by a power-law jump distribution with exponent $g = 1.5$, starting at the center of the main frame. The amplification illustrates the self-similar properties of this process.



La conception traditionnelle chinoise de l'éthique

Du rapport entre l'homme et la nature et de leurs expériences agricoles, les ancêtres chinois ont tiré deux idées fondamentales : "L'harmonie et l'ordre".

Il faut maintenir l'harmonie entre l'homme et la nature et ***les hommes doivent respecter les lois de la nature***. L'harmonie est la pièce maîtresse de cette éthique.

Le **confucianisme** et le **taoïsme** sont les deux grands ***courants ago-antagonistes*** qui ont animé l'histoire chinoise avec le **bouddhisme** arrivé en Chine à l'époque des Han (vers le III^{ème} siècle), probablement par des commerçants indiens, via la Route de la Soie.

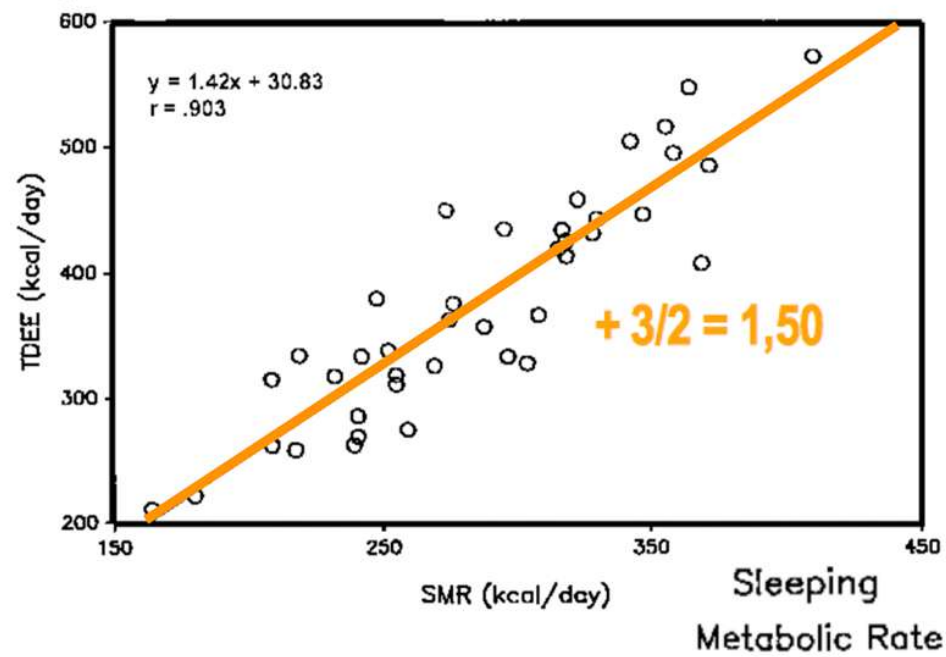
Confucius a dit : "L'harmonie prime en tout".

Le taoïsme est considéré par certains comme étant la seule "vraie" **religion chinoise**. C'est un mélange du culte des esprits, de la nature et des ancêtres, ***une quête mystique des lois qui gouvernent notre vie***, en quelque sorte une quête de l'immortalité.

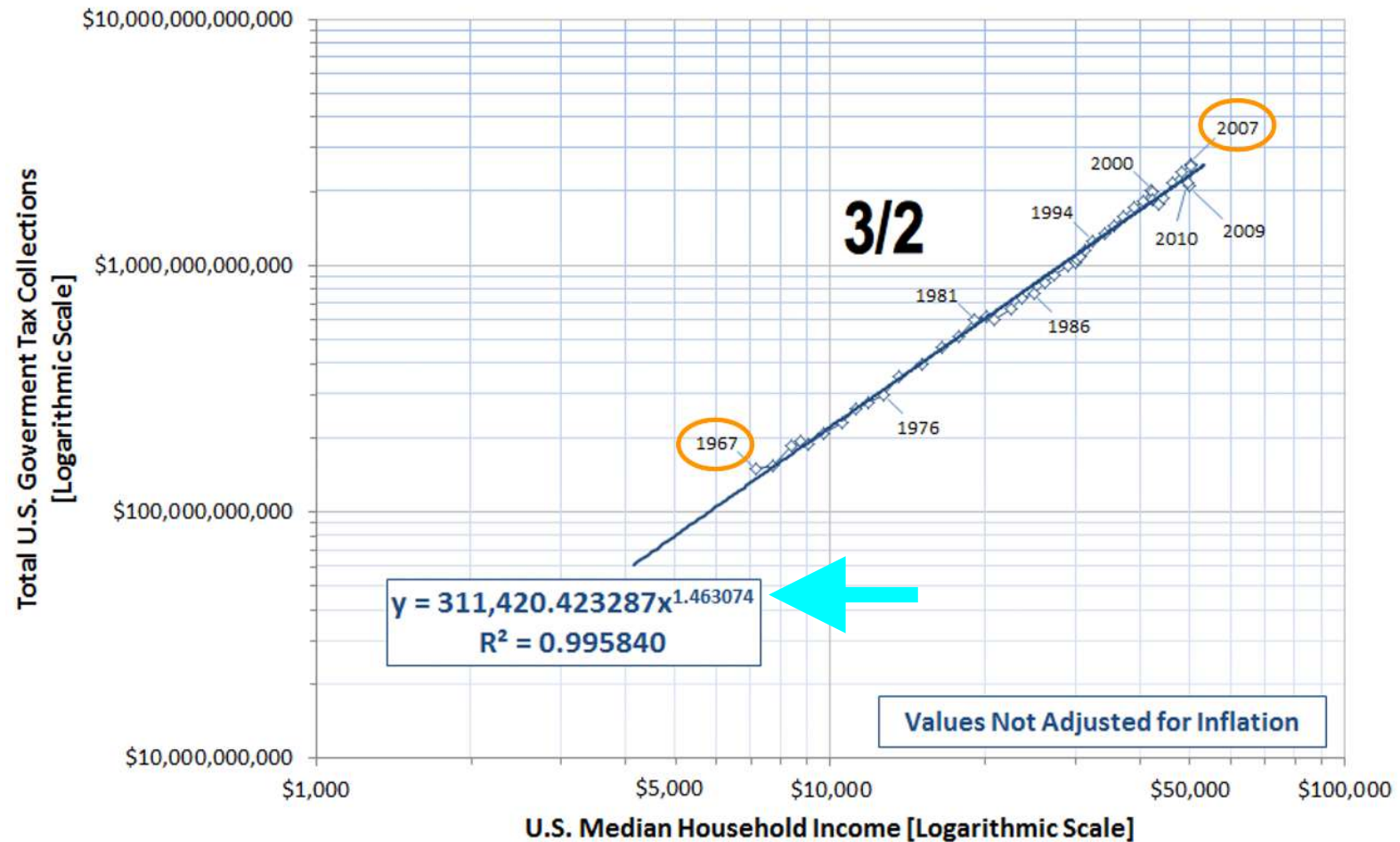
Les valeurs confucéennes se retrouvent dans toutes les sociétés qui ont adopté l'écriture chinoise. Le respect des enfants envers leurs parents, l'épouse envers son mari, conduisant à l'obéissance des travailleurs à leurs chefs, explique la discipline qui règne dans les entreprises chinoises.

Le maintien de l'harmonie se fait à l'aide de la loi et de l'ordre, une société doit être régie par l'ordre, ***ce qui lui permet d'établir une hiérarchie, un système, des lois, des relations humaines et des règles morales.***

Total
Daily Energy
Expenditure



Total U.S. Government Tax Collections vs U.S. Median Household Income, 1967-2010

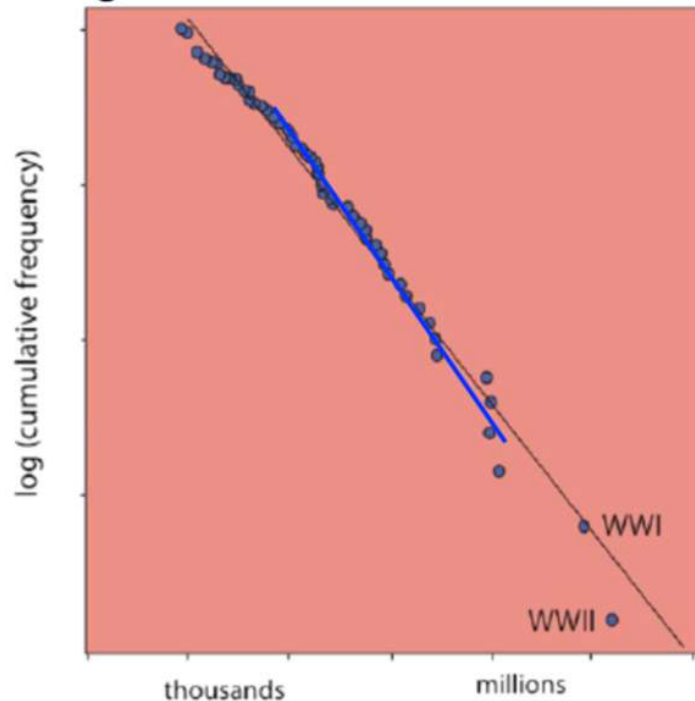


Data Sources: White House Office of Management and Budget and U.S. Census

© Political Calculations 2011

X1= Log x1

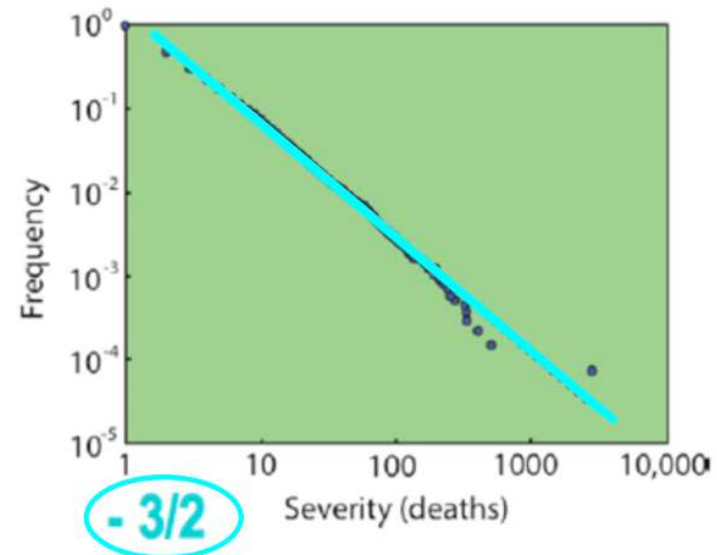
-1



X2 = Log x2 log (severity)

How the number of deaths in a war between states (from 1820 to 1997) varies with the frequency (equivalently, the probability) of such a conflict. The relationship is a mathematically simple one, called a power law. The last two data points are, respectively, World Wars I and II.

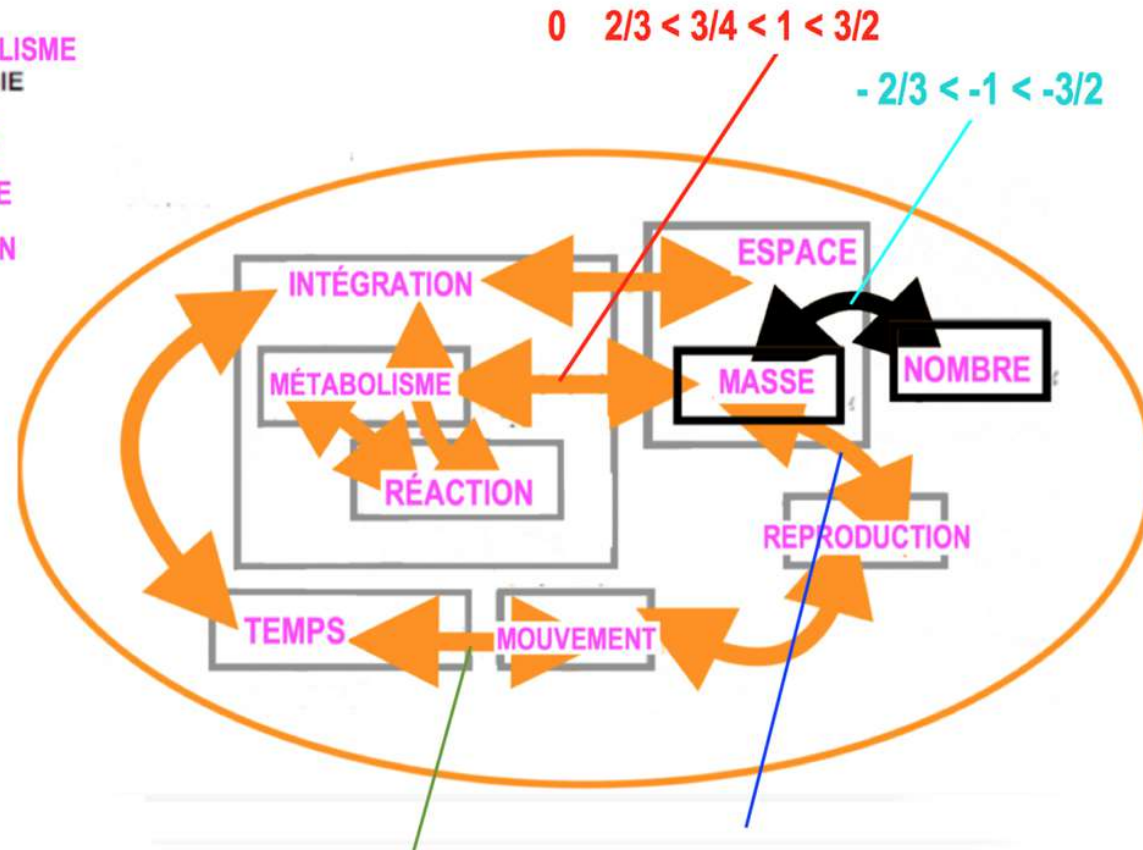
$$x1 \cdot x2 = C$$

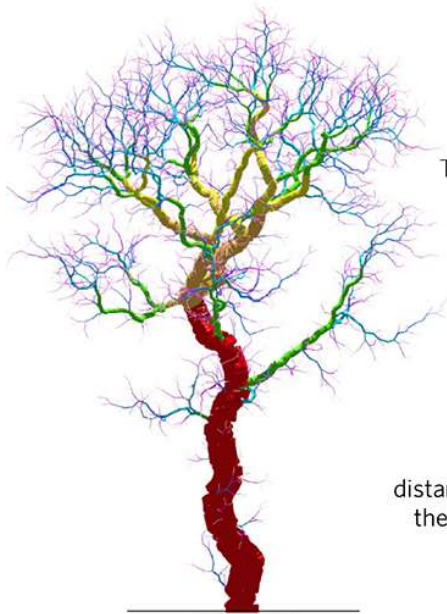


How the number of deaths in a terrorist attack (between 1968 and 2008) varies with the frequency of such an attack. The relationship is another power law, but with a different exponent from that of inter-state wars.

- SURVIVRE**
1. La MOBILISATION de la MATIÈRE et de l'ÉNERGIE **MÉTABOLISME**
 2. La CROISSANCE en masse **MASSE**
 2. La CROISSANCE en nombre **NOMBRE**
 3. La RÉPONSE à des STIMULATIONS **RÉACTION**
 4. L'ORGANISATION interne dans l'espace et dans le temps **ESPACE TEMPS**
 5. L'**INTÉGRATION** externe
 6. Le **MOUVEMENT**
- pour **SE SURVIVRE**
7. La **REPRODUCTION**

systémique

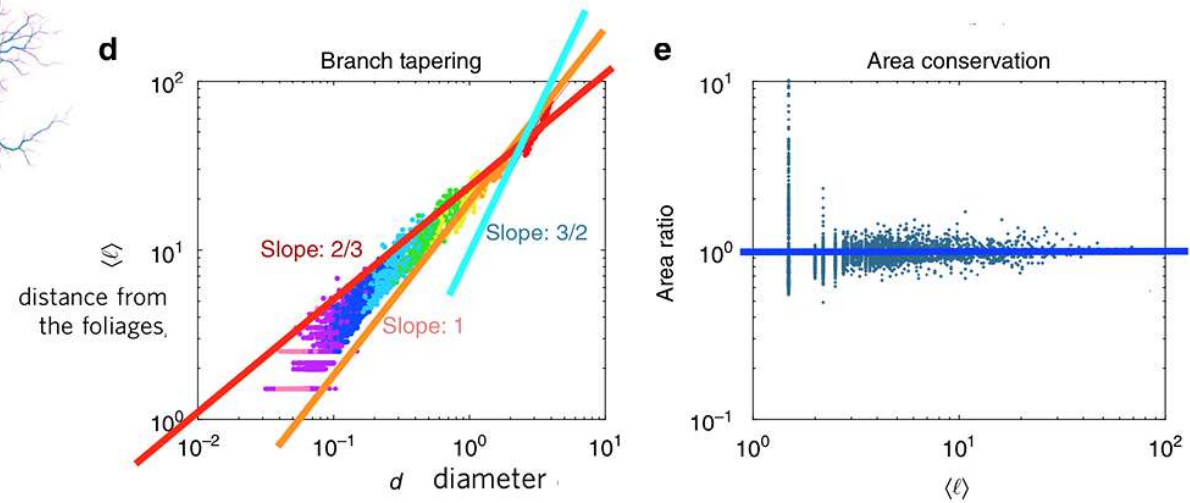


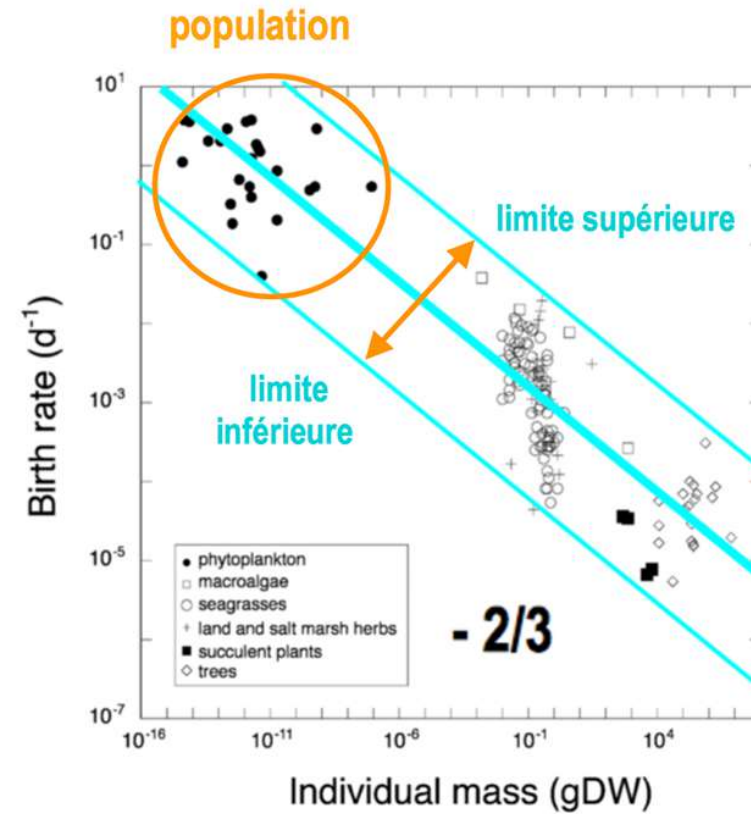
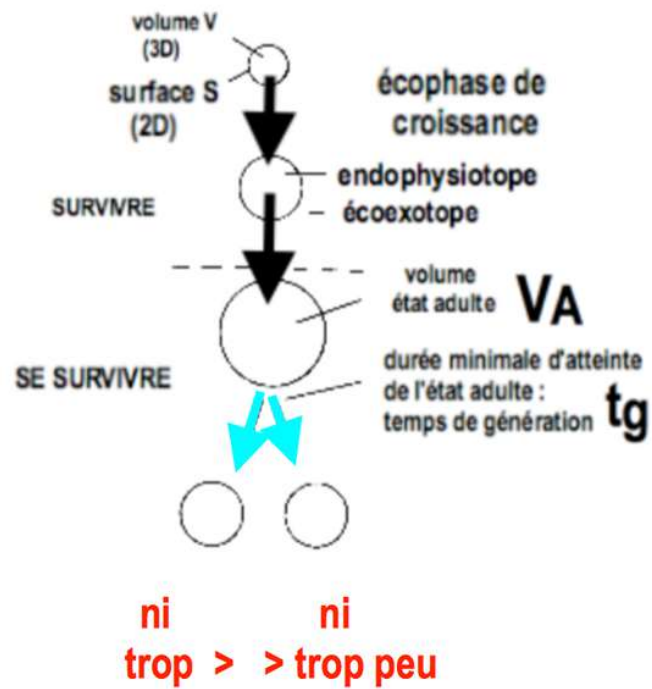


Self-similarity.

length (unit L), area (unit L^2) and diameter (unit L)

The Strahler order of each segment is represented with the same colour





CyberSystemics

How to capture the comprehensive nature of complex systems and processes as a whole?

CyberSystemics abounds of celebrities ...

If you know these authors or these words (concepts), you have made already cybersystemics without knowing it!

R. ASHBY (**requisite variety**), Gregory BATESON (**inter-personal communication**), Henri BERGSON (**creative changing & evolution**), Claude BERNARD (**homeostasis**), Elie BERNARD-WEIL (**agoantagonism, paradoxical strategies**), BERTIN (diagrams, networks, maps: **graphic semiology**), BOLTZMANN (**behavior statistical laws**), BOOLE (**algebra**), BOULDING (**economic typology of systems**), BUNGE (**general typology of systems**), CANNON, Noam CHOMSKY (**language syntax**), L. COUFFIGNAL, Michel CROZIER (**multiple agents decision making systems**), Joël DE ROSNAY (*the macro-scope*), DE SAUSSURE (**linguistic structuralism**), Gérard DONNADIEU & Michel KARSKY (**systemic approach**), J.W. FORRESTER (**systems dynamics**), Michel FOUCAULT (**analogy**), Arthur KOESTLER (**wholeness**), LAVOISIER (**isomorphism**), Jean-Louis LE MOIGNE, LEIBNIZ, J. LESOURNE (**fate systems**), Claude LEVY-STRAUSS (**anthropologic structuralism**), Edward LORENZ (**butterfly effect**), James LOVELOCK (**Gaia system**), MAC CULLOCH (**biomimicry**), Benoît MANDELROT (**fractals, systems scaling invariance, theory of catastrophes**), Edgar MORIN (*the method*), Blaise PASCAL (**the Whole and the parts**), Louis PASTEUR, Jean PIAGET (**psychological structuralism**), PRIGOGINE (**autonomy & auto-organization, dissipation structures**), PYTHAGORE, Michel SERRES (**communication interference**), Claude SHANNON (**communication, quantitative theory of information**), SKINNER (**behavior**), Herbert SPENCER (**progress**), TARSKY (**theory of models**), Pierre TEILHARD DE CHARDIN (**man cosmic system**), Paul VALERY (**models & modeling**), Ludwig VON BERTALANFFY (**general system theory**), VON FOERSTER, (**information**), VON NEUMANN & MORGENTHAU (**games theory**), Norbert WIENER (**cybernetics**).

Les échelles des **systèmes** vivants :

- **espace** <http://www.afscet.asso.fr/Ande10/pbETA vivant10.pdf>
- **temps** <http://www.afscet.asso.fr/Ande12/pbAnde2012.pdf>
- **espace-temps** <http://www.armsada.eu/pb/bernardins/phylogtagmotaphologie.pdf>



LES LOIS SYSTÉMIQUES de l'ÉVOLUTION du VIVANT

- **LOI SYSTÉMIQUE D'INVARIANCE DE JAUGE** :
écoexotopie et endophysiotopie, les 7 caractéristiques fonctionnelles du vivant "vivant".
- **LOI SYSTÉMIQUE ERGODIQUE D'ÉMERGENCE CONSTITUTIVE** :
niveaux d'organisation et échelles d'emboîtements hiérarchiques, règles d'émergence, ARMSADA et confinement.
- **LOI SYSTÉMIQUE ERGODIQUE D'INDIVIDUATION ET D'ÉLAN VITAL** :
hypothèse de linéarité et principe de moindre action, loi puissance $V_A^2 = k.t.g^3$, non-linéarité.
- **LOI SYSTÉMIQUE CONSTRUCTALE** :
global et local, capacité d'accueil et capacité d'être accueilli, action et rétroaction.
- **LOI SYSTÉMIQUE ERGODIQUE DE CAUSALITÉ ONTOGÉNÉTIQUE** :
tableau de la classification périodique du vivant.

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- Bricage P. (2017) **Education for sustainability: lessons from living systems governance**. <https://halshs.archives-ouvertes.fr/halshs-01705968/document>

Online reference to cite this work:

- Bricage P. (2018) **Déterminisme et indéterminisme multi-échelle, invariants et variance des systèmes vivants**. -présentation- Journées annuelles de l'AFSCET à Andé, France, 05 mai 2018, 84 p. http://afscet.asso.fr/Ande18/pbricage_Ande2018_interactions-PowerLaws_presentation.pdf