

Fibonacci Phyllotaxis: Models, Data and Alan Turing

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Novel Fibonacci and non-Fibonacci structure in the sunflower: results of a citizen science experiment

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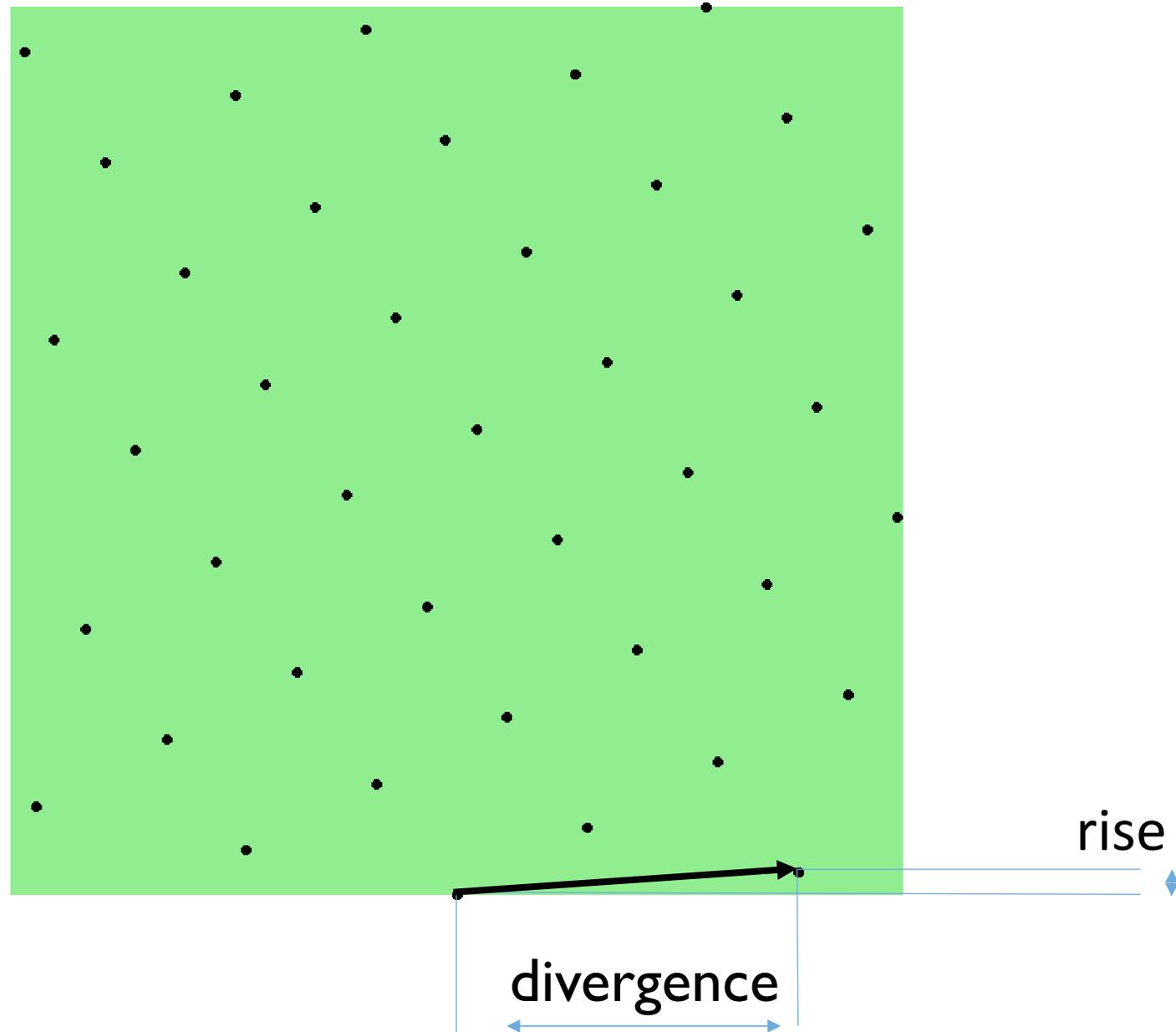
D'Arcy Thompson

The Fibonacci series...and the hypothesis of its introduction into plant-structure through natural selection, are all matters which deserve no place in the plain study of botanical phenomena...all such speculations as these hark back to a school of mystical idealism

I'm going to talk about

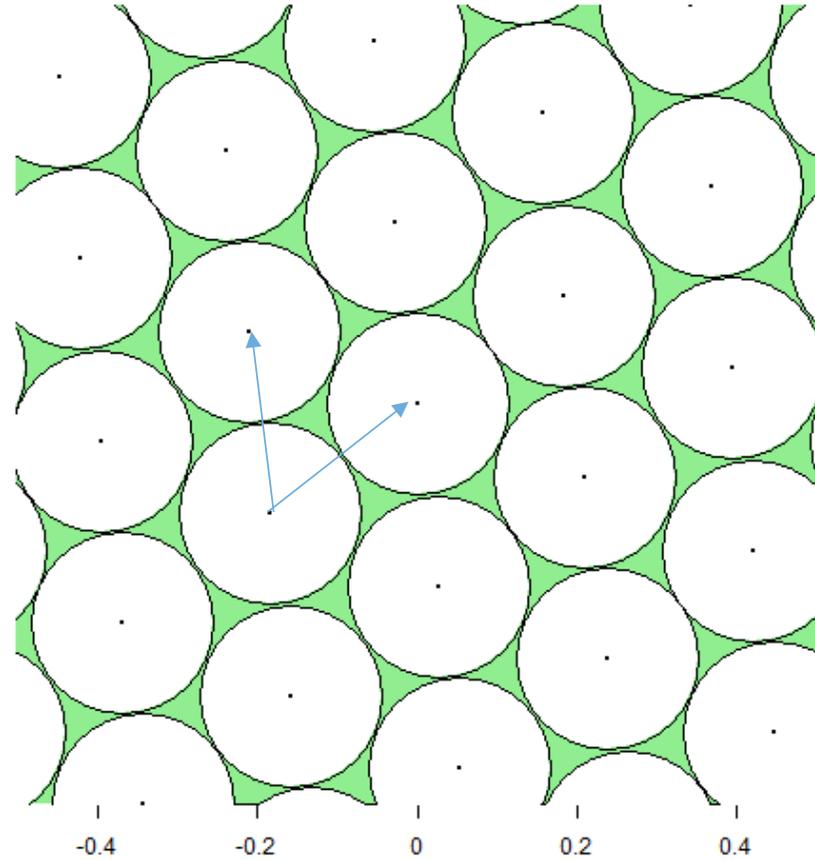
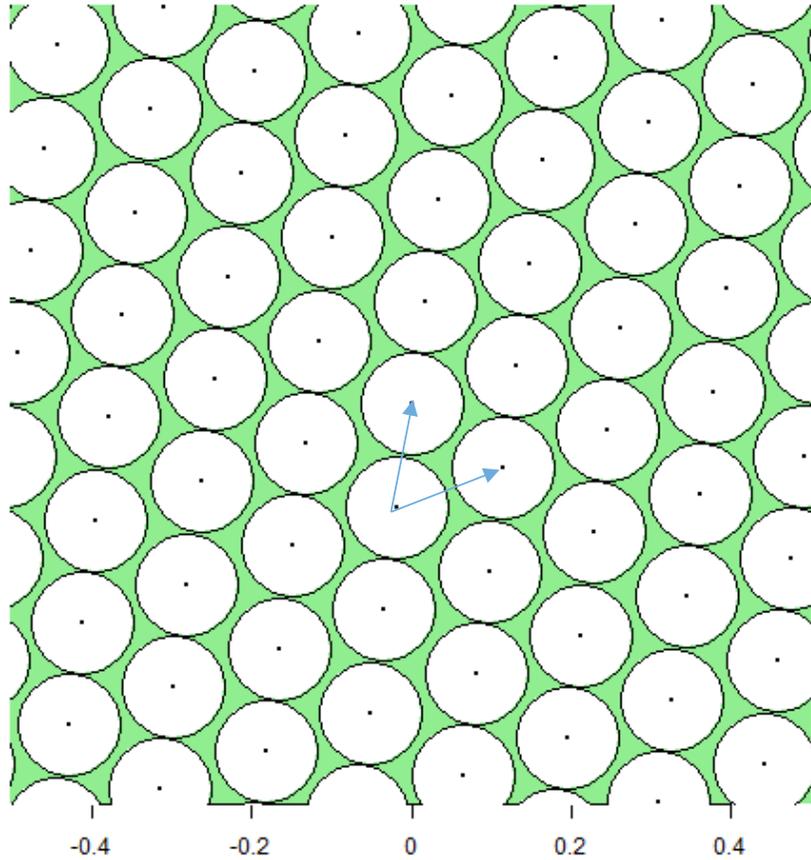
- Mathematics of lattices
- Citizen science & Alan Turing: first computational biologist
- Opportunities for today's mathematical biologists

Lattice is on
a cylinder
has two
parameters

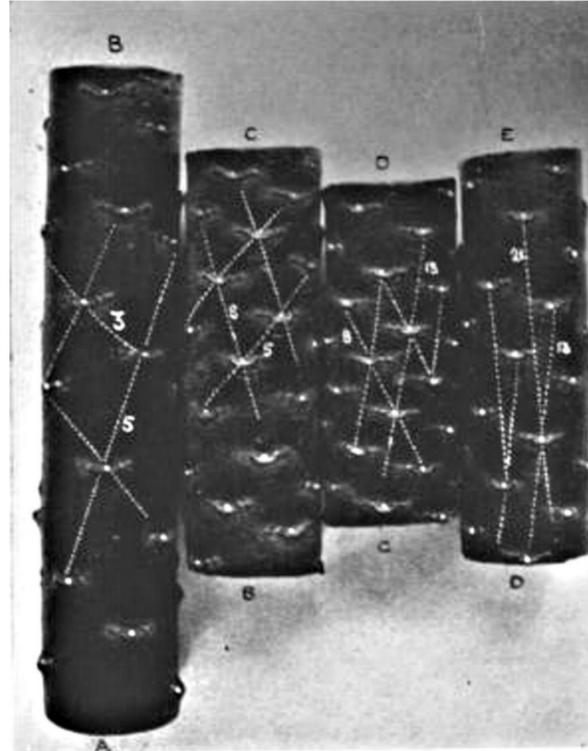


Lattices on cylinders

- can define 'obvious' parastichies
- only some lattices are disk packings

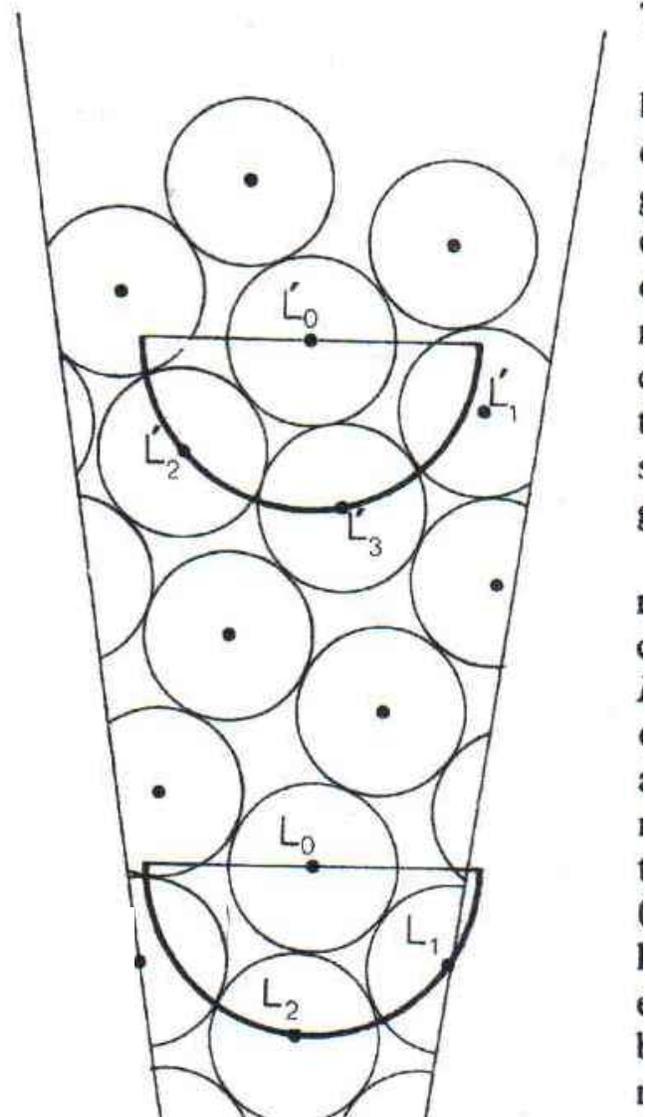


Model: lattice at each height on the stem is a quasi steady state solution of repeated node-placement decisions

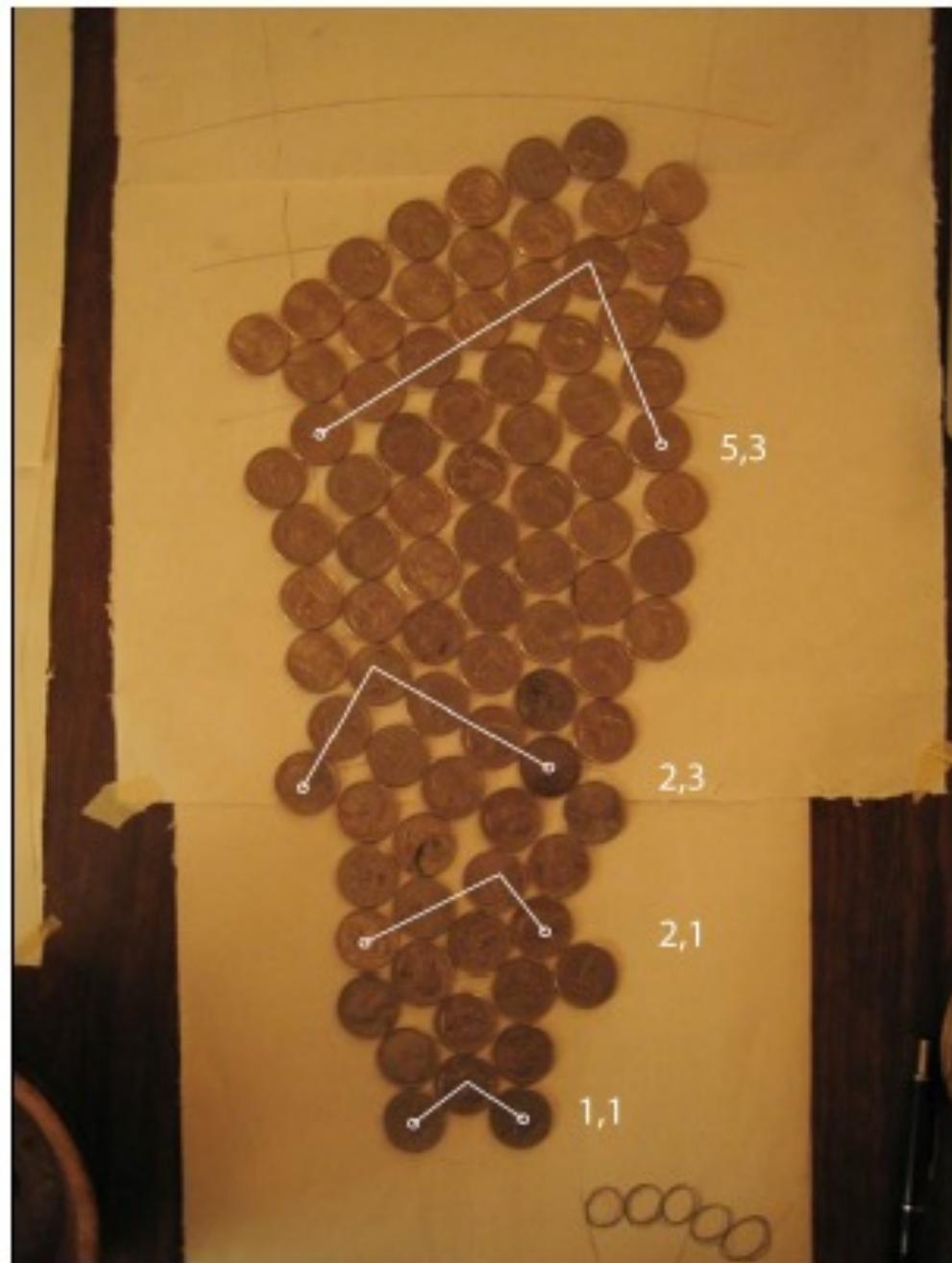


Church 1904

Node placement models

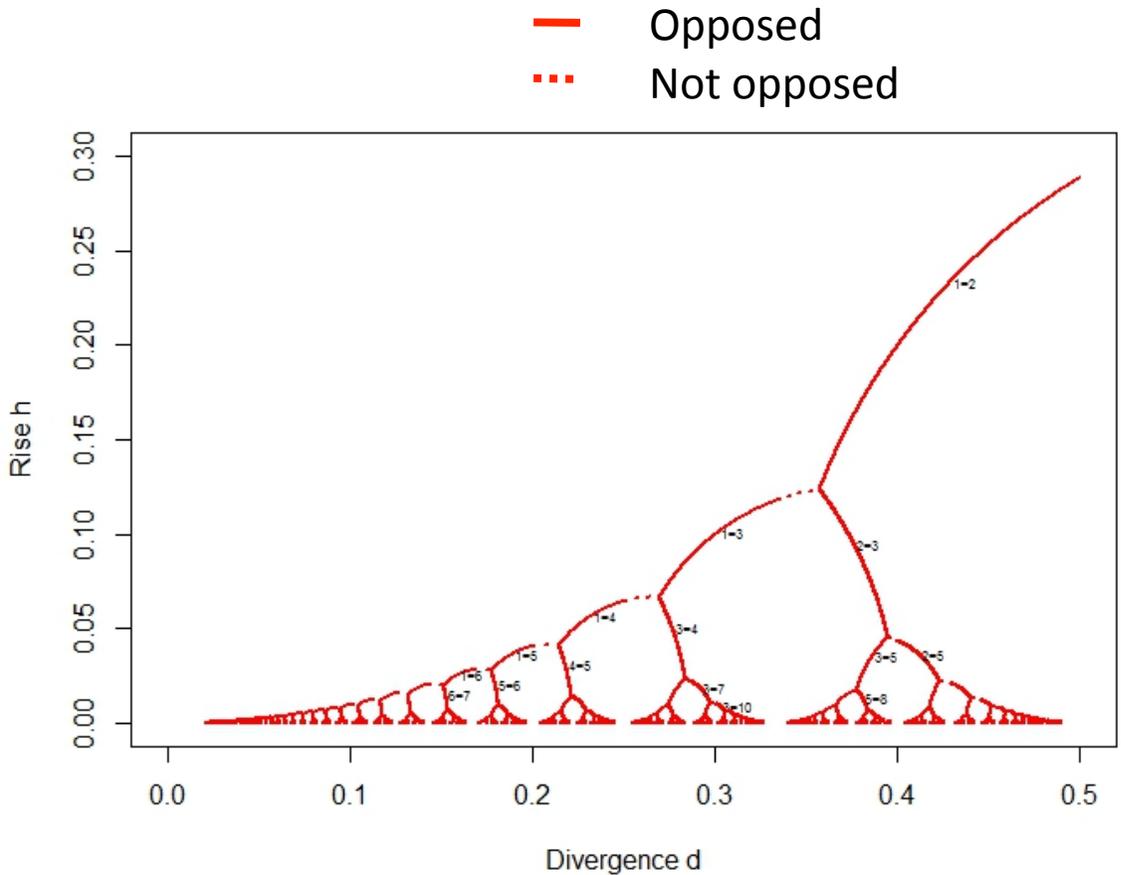
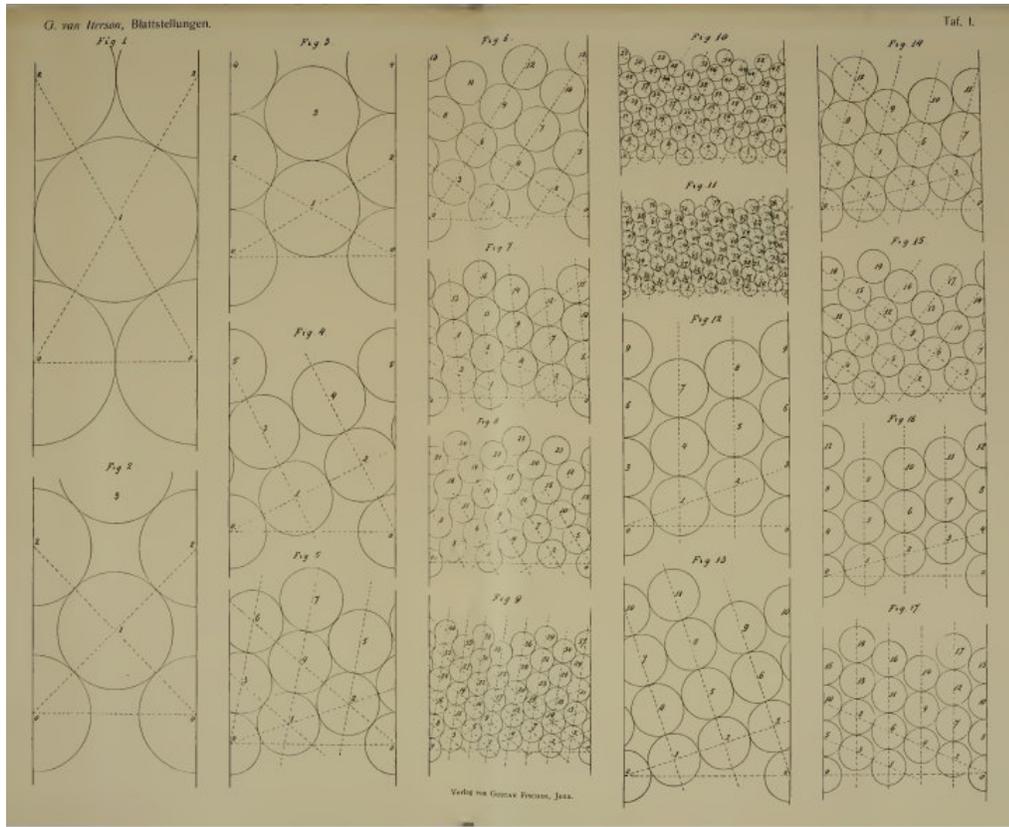


Mitchison 1977



Atela 2011

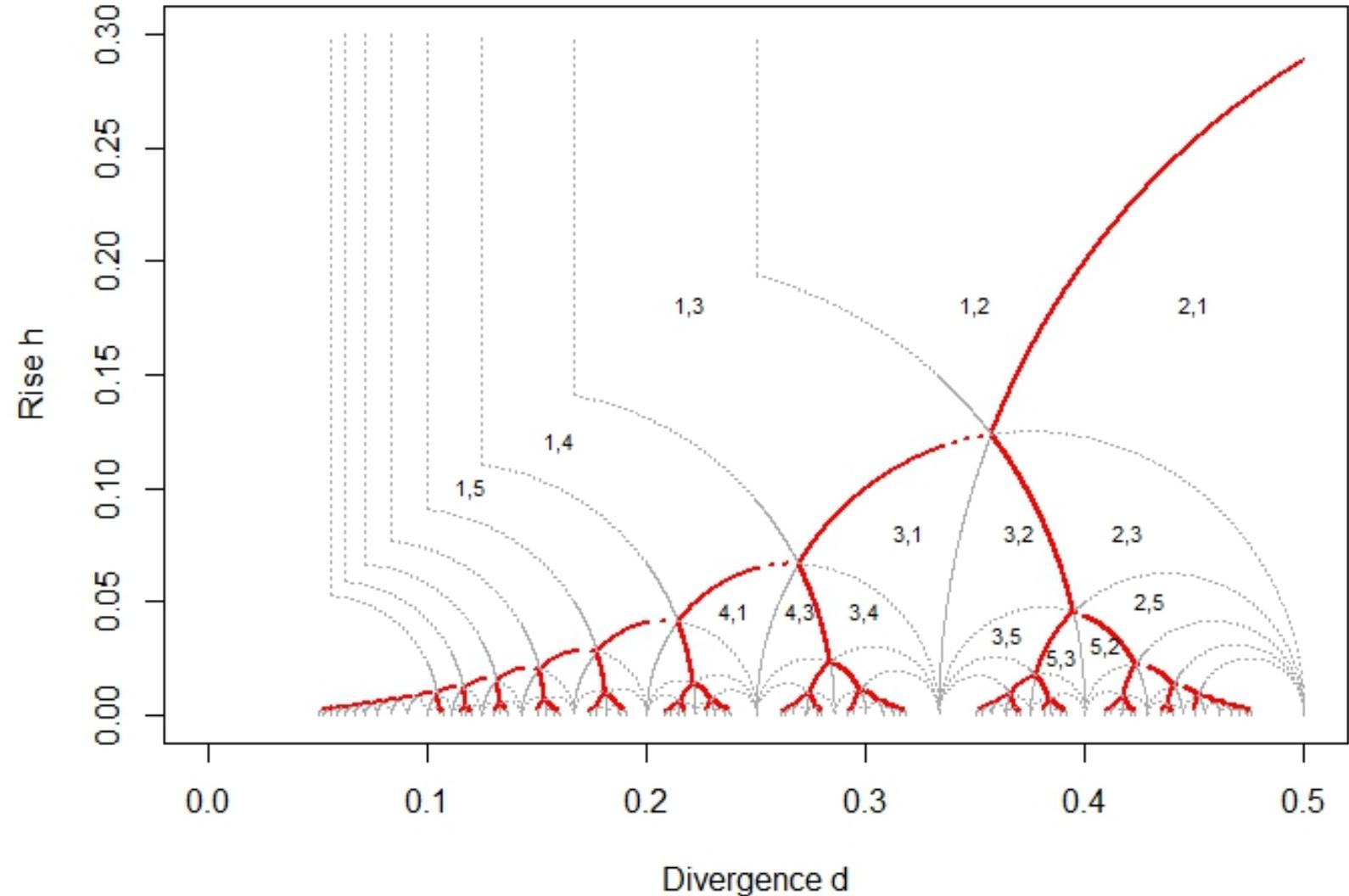
The van Iterson diagram – packed disk lattices



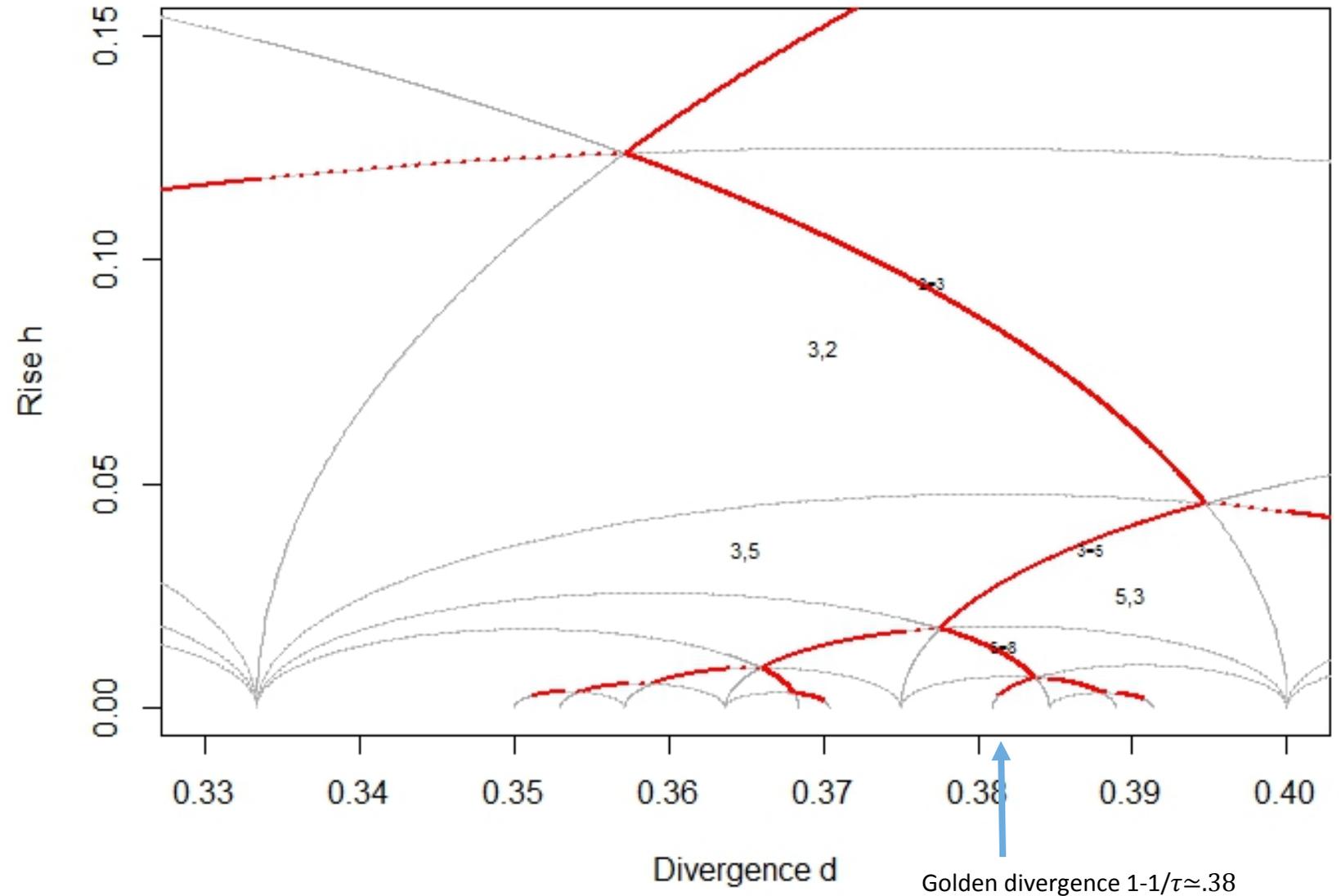
van Iterson 1907

Lattice space categorised by primary and secondary parastichies

- Every relatively prime pair of integers appears once
- Observation of parastichy pairs constrains divergence

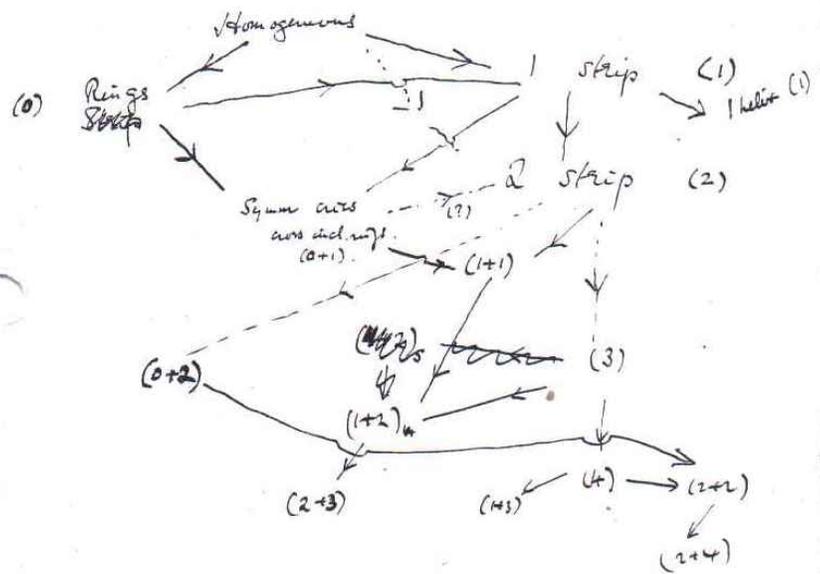


Branch points: (m,n) splits into $(n,m+n)$, and $(m, m+n)$



A path to Fibonacci phyllotaxis

- A model for new node placement that locally produces regular lattices
- A constraint that causes these lattices to be disk-packing
 - Keeps us on the van Iterson tree
- A smoothly changing parameter that gradually increases the complexity of the lattice
- A constraint that causes the Fibonacci property to be preserved at each bifurcation
 - Turing: 'The hypothesis of geometrical phyllotaxis'



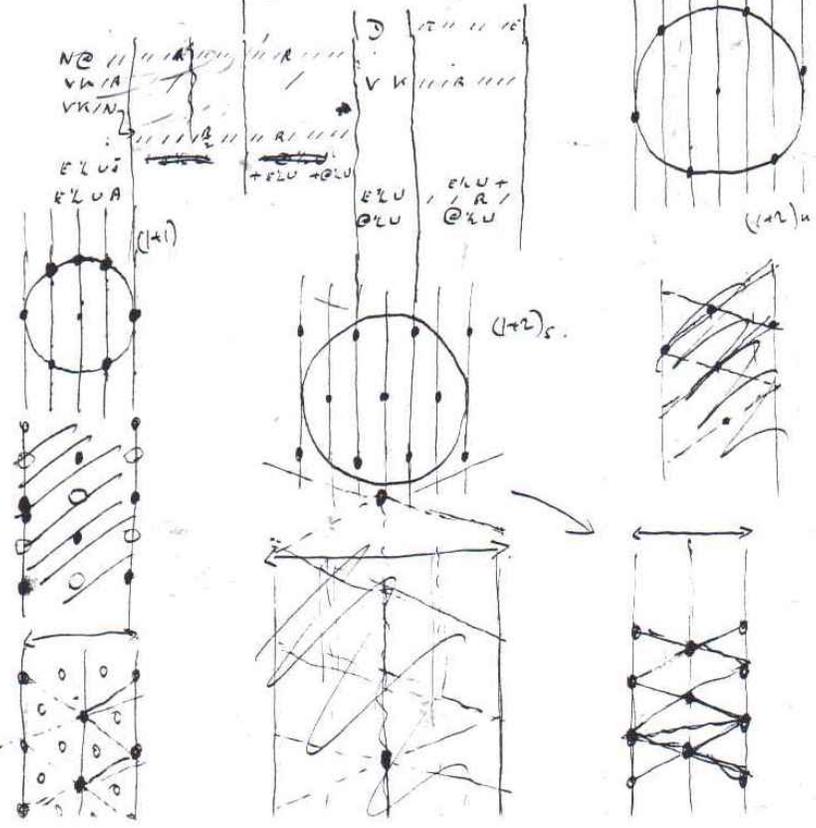
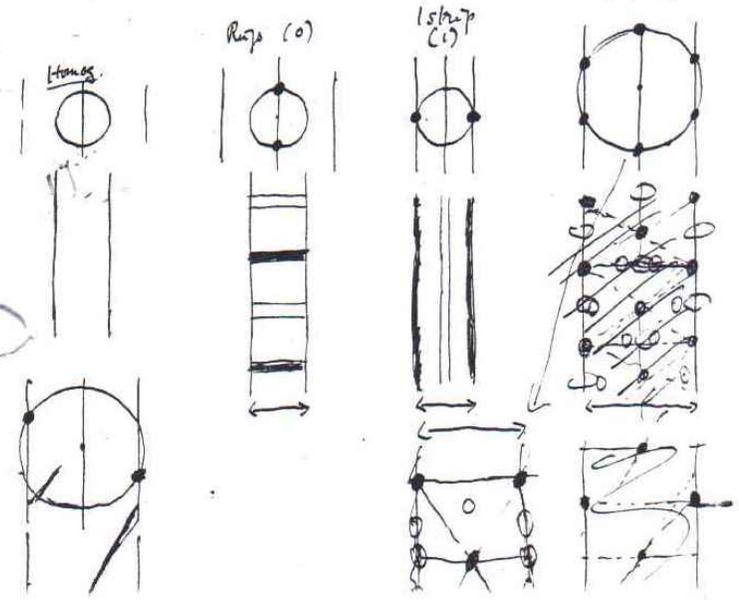
7	N	0	1	1	F
V	N	A	A	C	
V	N	N	N	K	
E	Z	U	J	T	
E	Z	U	A	Z	
F	:	B	S	C	
F	@	:	T	H	

/// R /

(U//N//E) R /

VK/R /

(E//@//@) E//@//@



At present I am not working on the problem at all, but on my mathematical theory of embryology, which I think I described to you at one time. This is yielding to treatment, and it will so far as I can see, give satisfactory explanations of -

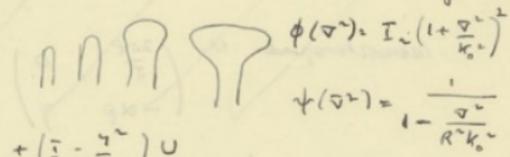
- i) Gastrulation.
- ii) Polygonally symmetrical structures, e.g., starfish, flowers.
- iii) Leaf arrangement, in particular the way the Fibonacci series (0, 1, 1 = 2, 3, 5, 8, 13,.....) comes to be involved.
- iv) Colour patterns on animals, e.g., stripes, spots and dappling.
- v) Patterns on nearly spherical structures such as some Radiolaria, but this is more difficult and doubtful.

I am really doing this now because it is yielding more easily to treatment. I think it is not altogether unconnected with the other problem. The brain structure has to be one which can be

$$\frac{\partial U}{\partial t} = \phi(\nabla^2)U + aU^2 - U + U(\psi(\nabla^2)U^2) + \Sigma(\tau, \gamma)U$$

$$(\bar{I}_2)(k_0)(G) \bar{I}_0 \sigma + R e$$

$$\frac{\partial U}{\partial t} = (1 + \nabla^2)^{-1}U + U^2 - U + U \frac{U^2}{1 - \nabla^2/R^2} + (\bar{I}_0 - \frac{\gamma^2}{\epsilon^2})U$$



$$U = \sum_{\nu=0,2} \gamma_\nu e^{i(\nu, z)} \quad U^2 = \sum_{\nu+\mu=\nu} (\sum_{\nu+\mu=\nu} \gamma_\nu \gamma_\mu) e^{i(\nu, z)}$$

As an approx. $\frac{U^2}{1 - \nabla^2/R^2} = \sum (\gamma_\nu \gamma_\mu) = V$ and $\sigma = \infty$ (R small)

$$\frac{d\gamma_\nu}{dt} = (1 - \nu^2)^{-1} \gamma_\nu + (\bar{I}_0 - 4V) \gamma_\nu + \sum_{\nu+\mu=\nu} \gamma_\mu \gamma_\nu$$

A letter pattern remains a letter.

Suppose $\gamma_0, \gamma_2, \dots, \gamma_{2k}$ different from zero

$$\frac{d\gamma}{dt} = (\bar{I}_0 - 4V)\gamma + 2\gamma^2 \quad V = 6\gamma^2$$

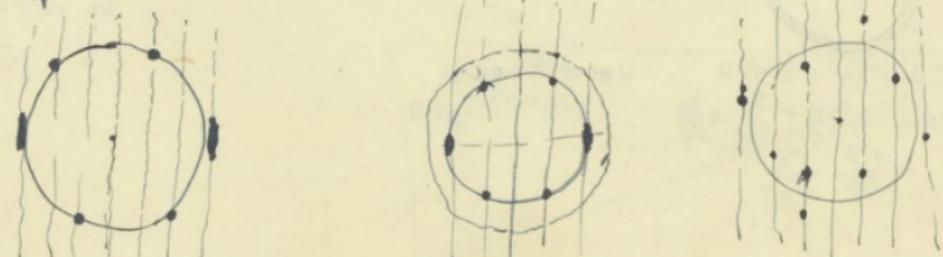
$$= \gamma(\bar{I}_0 + 2\gamma - 6\gamma) = \gamma(\bar{I}_0 - 4\gamma)$$

Solve $\gamma^{(0)}$ $F(\gamma^{(0)}) = 0$
 Stability of $F(\gamma^{(0)}) < 0$. Roots alternately stable + unstable.

One root at $\gamma=0$, also at $\gamma = \frac{1}{2H}(\sqrt{1+6H\bar{I}_0} + 1)$

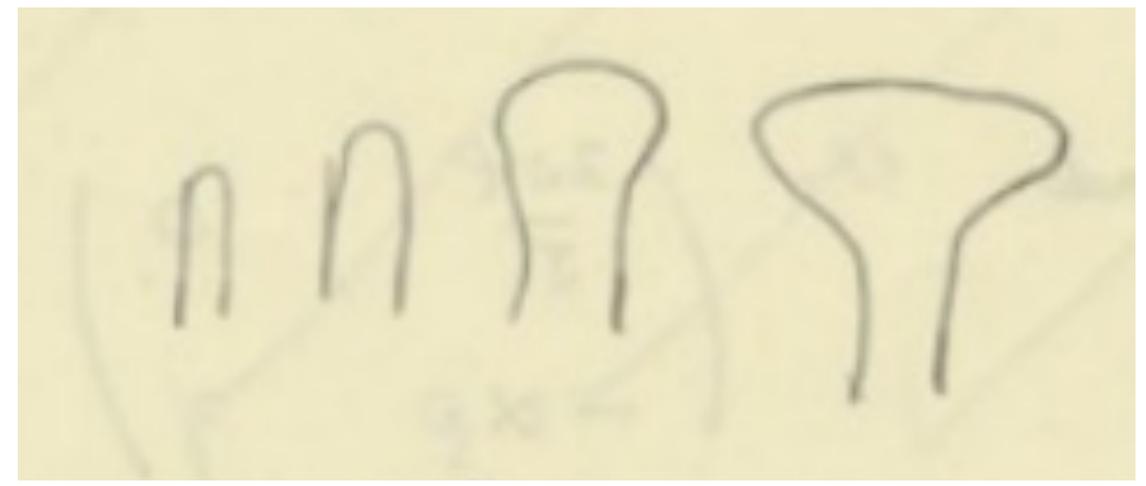
$F'(\gamma) \rightarrow -\infty$ at $\gamma \rightarrow \pm\infty$. Derivatives are stable

Transition from unstable to stable if $\bar{I}_0 > 0$
 If $1+6H\bar{I}_0 < 0$ then only root is $\gamma=0$ and stable
 If $-\frac{1}{6H} < \bar{I}_0 < 0$ then $\gamma=0$ is stable root and also $\frac{1}{2H}(1 + \sqrt{1+6H\bar{I}_0})$
 If $\bar{I}_0 > 0$ then $\gamma=0$ is unstable and $\frac{1}{2H}(1 \pm \sqrt{1+6H\bar{I}_0})$ are stable.



Outline of development of the Daisy.

XXXXXXXXXXXXXXXXX The theory developed in this paper is limited by a number of assumptions which are by no means always satisfied. Two are of special importance





Dynamic node placement models

- Airy (1873): spheres glued to an elastic band
- Hofmeister's rule: new primordia appears 'periodically' in largest available space
- Snows' rule: new primordia whenever 'enough' space
- Veen & Lindenmayer (1977): Turing-like RD model
- Douady and Couder (1992-) repulsive particles (soft disks)
- Levitov (1991), Kunz (1995) – soft disks
- Mitchison (1977) , Atela et al – coin-dropping models
- All obey the HofGP – why?

HofGP-type models say:

- Placement in older parts of the plant influences placement in younger (later) parts but not vice versa
 - Increases in parastichy number in later parts of the plant
 - Parastichy numbers remain adjacent members of (usually) the Fibonacci sequence
 - If not strict Fibonacci, then adjacent members of one particular Fibonacci-structure sequence
 - Ratio of parastichy numbers close to the golden ratio
 - Specific models will predict how development responds to noise
-
- Hard to imagine any other model structure that could predict high Fibonacci numbers such as the sunflower head
 - Intrinsically mathematical (or at least computational) – by contrast with eg current developmental biology

New generations of models with noise/defects

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M.F. Pennybacker et al. / I

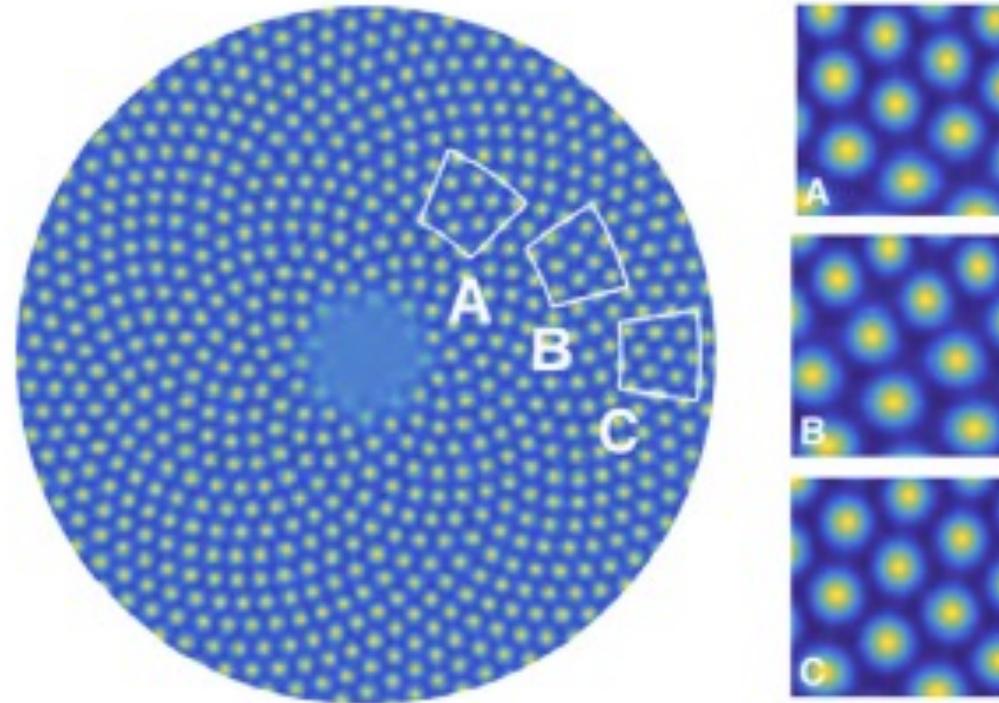


Fig. 13. Simulation of (1) showing a phyllotactic pattern that has propagated as a front from the outer edge towards the center of a disk, a model of pattern formation on a sunflower inflorescence meristem. The simulation is described in detail in Section 3. Cutouts of the pattern centered at A $r = 75/\phi$, B $r = 75/\sqrt{\phi}$, and C $r = 75$, where ϕ is the golden number, transformed as described in the text, appear as panels A–C.

Literature before 2012

Table 7.7. *Patterns for 319 Helianthus (sunflower) capituli*

Sequence	Frequency	%
$\langle 1, 2, 3, 5, 8, \dots \rangle$	262	82.13
$2\langle 1, 2, 3, 5, 8, \dots \rangle$	9	2.82
$\langle 1, 3, 4, 7, 11, \dots \rangle$	46	14.42
$\langle 1, 4, 5, 9, 14, \dots \rangle$	2	0.63

Source: Schoute (1938).

Table 7.8. *Data for 140 Helianthus capituli*

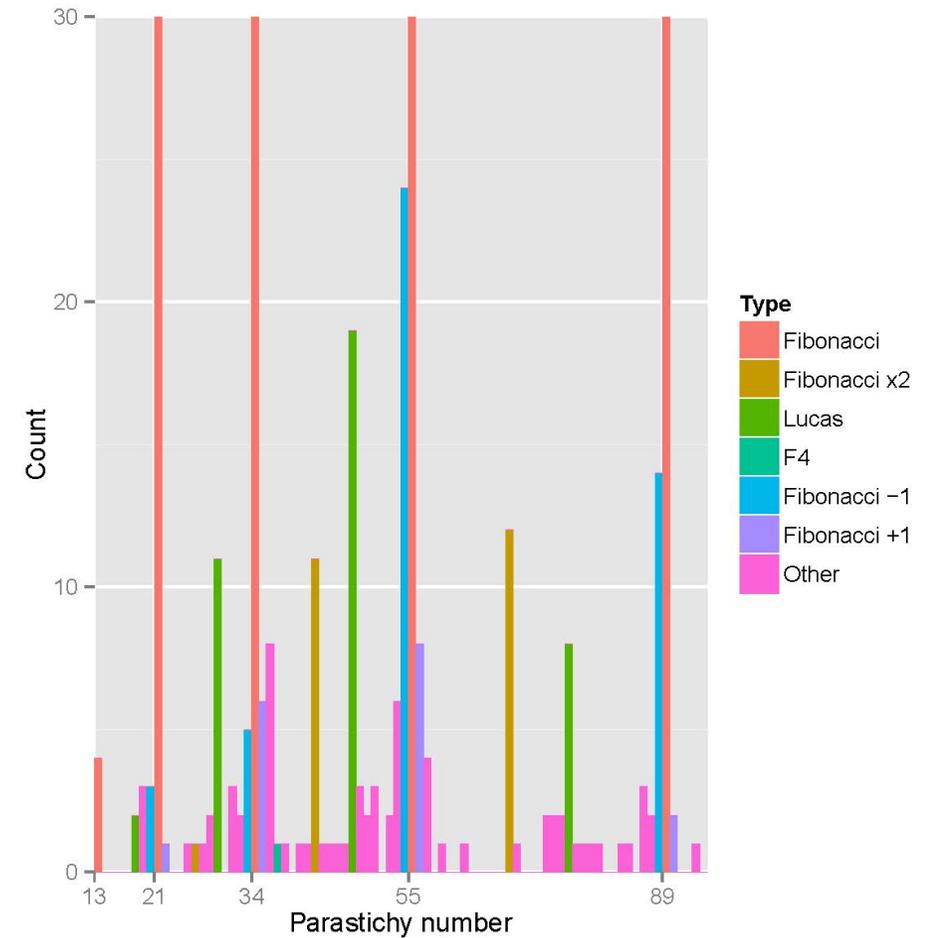
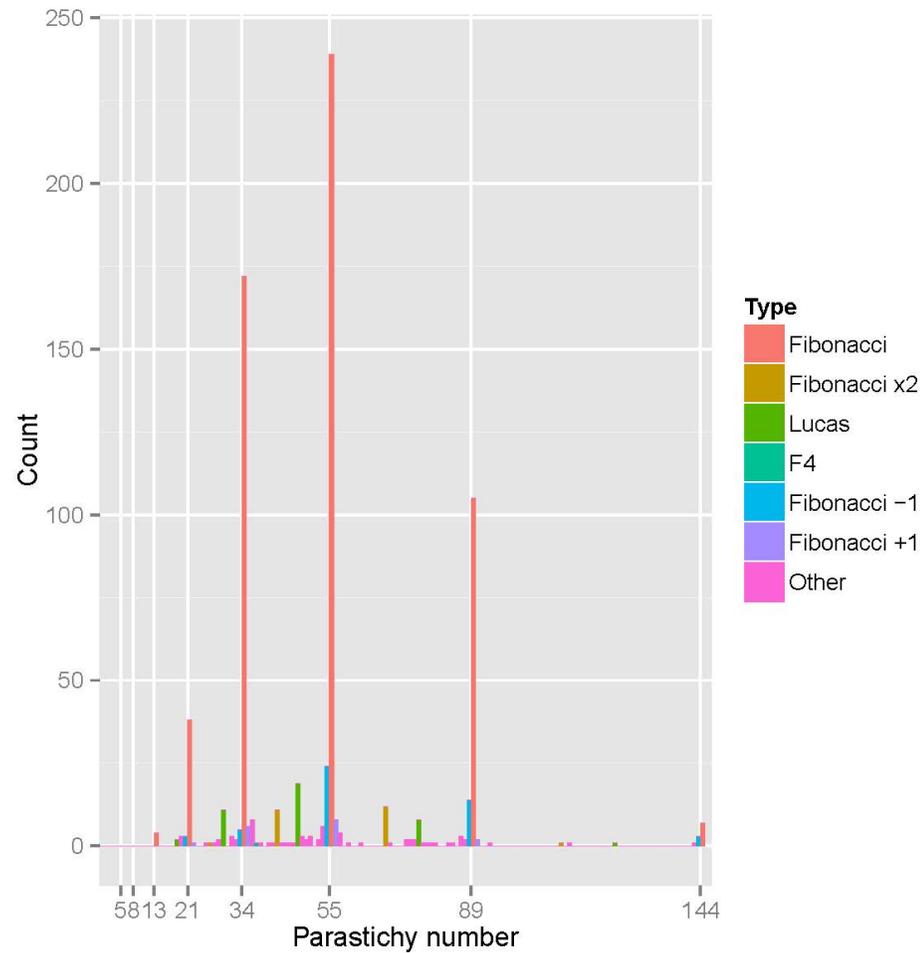
Sequence	Frequency	%
$\langle 1, 2, 3, 5, 8, \dots \rangle$	133	95.0
$2\langle 1, 2, 3, 5, 8, \dots \rangle$	1	0.71
$\langle 1, 3, 4, 7, 11, \dots \rangle$	6	4.29
$\langle 1, 4, 5, 9, 14, \dots \rangle$	0	0

Source: Weisse (1897).

Fibonacci structure

- Fibonacci rule $F_n = F_{n-1} + F_{n-2}$
 - Any sequence obeying this rule has $F_n / F_{n-1} \rightarrow \tau$
 - $\tau \simeq 1.618$ is the golden ratio $\tau^2 = \tau + 1$
- Fibonacci sequence 1, 2, 3, 5, 8, 13, 21, 34, 55, 89, 144 ...
- Lucas sequence 1, 3, 4, 7, 11, 18, 29, 47, 76, 123, ...
- F4: 1, 4, 5, 9, 14, 23, 37, 60, 97, ...
- F5: 1, 5, 6, 11, 17, 28, 45, 73, ...
- Double Fibonacci 2, 2, 4, 6, 10, 16, 26, 42, 68, 110, ...

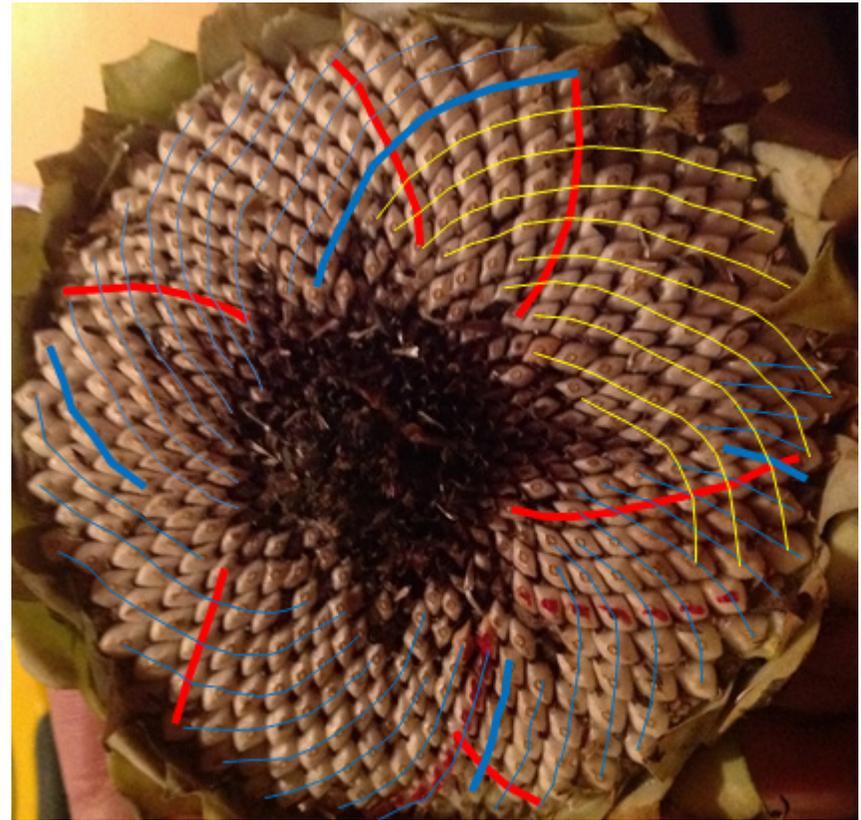
Manchester experiment



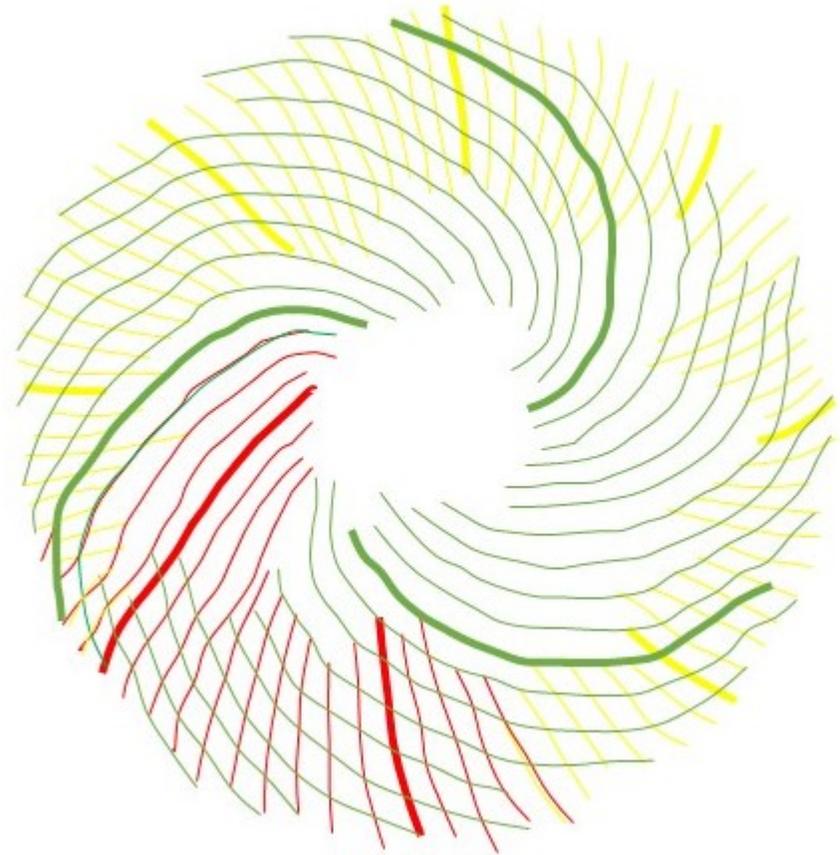
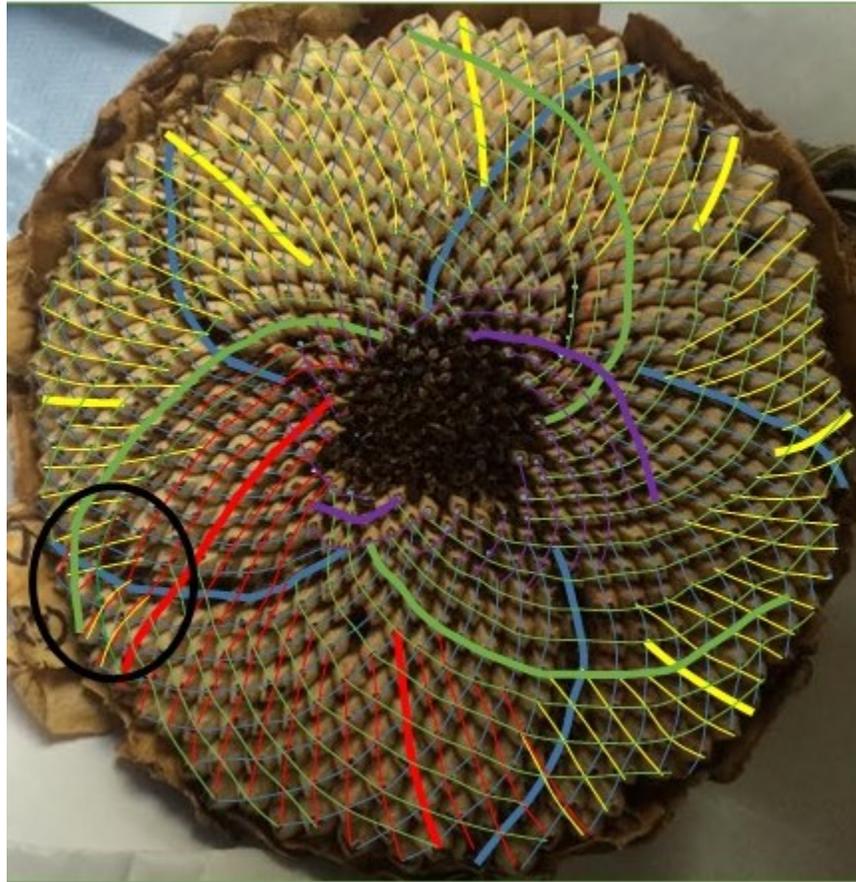
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(55,-)



Overlapping parastichies (50,81/31/20)



? (77,56)



? (62,31)



Manchester results summary

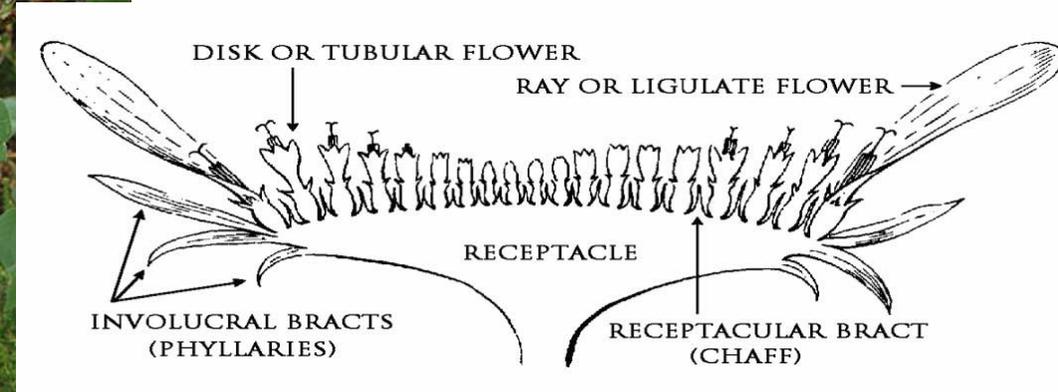
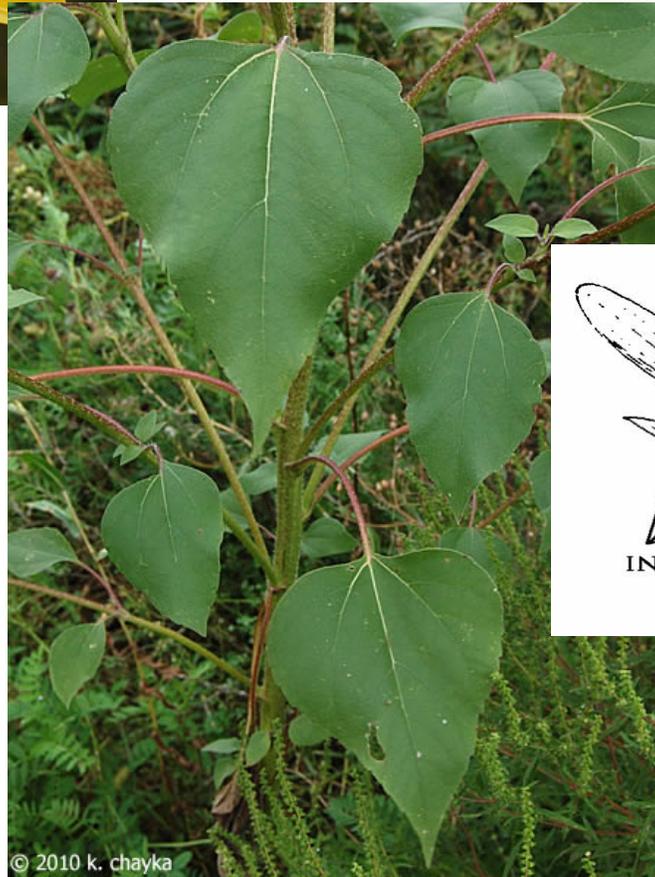
- Fibonacci is commonest; Lucas more common than double Fibonacci
- Approximately Fibonacci is common (54 more common than Lucas 47) mainly $F-I$, $F+I$
- $F-I$ (significantly) more common than $F+I$
- Common departures from rotational symmetry making parastichies uncountable
- Count in one direction often much more ordered than in the other
- Some seedheads possess completely disordered regions

Turing's sunflowers: a citizen science experiment



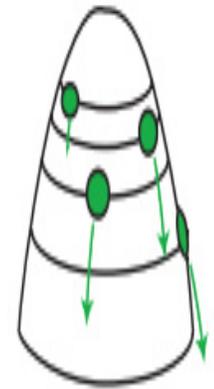
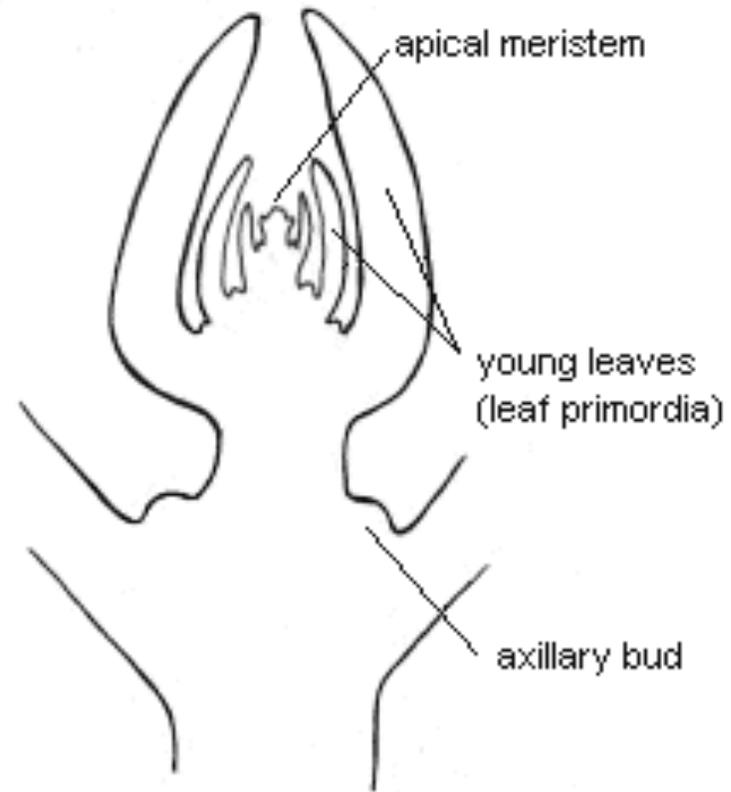
Summary

- Fibonacci phyllotaxis is a real phenomenon whose observation deserved replication
- We have plausible mathematical reasons why it should exist – although some more work on why the HofGP is often true would be worthwhile
- The assumptions and predictions of these models are not well connected to empirical biology
- If they were, they would likely make a powerful argument for systems biology
- The Manchester dataset should be useful for this – let's make some more!





garden.org



Atela 2002

Comparison with previous studies

