Philosophy of Biology

Stavros Ioannidis / MA Cont. Phil.

General information

Day/time: Thursday 15.15 - 18.00, Library

Office hours: Monday 18.00 - 19.00 (old building, ground floor, first office on your left)

Email: stavros.ioannidis.phil@gmail.com

Main Bibliography

- **Godfrey-Smith** (2016) *Philosophy of Biology*, Princeton University Press.
- Hull & Ruse (2017) The Cambridge Companion to the Philosophy of Biology, Cambridge University Press.
- Rosenberg & McShea (2008) Philosophy of Biology: a contemporary introduction, Routledge.
- **Sober** (ed.) (2006) *Conceptual Issues in Evolutionary Biology*, Cambridge, Massachusetts: MIT Press.
- Sterelny & Griffiths (1999), Sex and Death: An Introduction to the Philosophy of Biology, Chicago: University of Chicago Press.
- **Sterelny** (2003) *Dawkins Vs Gould: Survival of the Fittest, Icon Books.*

Other sources

Griffiths, P.E. & K. Stotz (2013) *Genetics and Philosophy: an introduction*, New York: Cambridge University Press.

Stanford Encyclopedia of Philosophy: http://plato.stanford.edu

Darwin (1859) On the Origin of Species (first edition).

http://darwin-online.org.uk/content/frameset? itemID=F373&viewtype=text&pageseq=1

(intro, ch. 1-4, 6, 14)

Assessment

- Presentation of one or two papers
- Final essay (4,500-5,000 words, without the references)

Emergence of philosophy of biology in the 70s

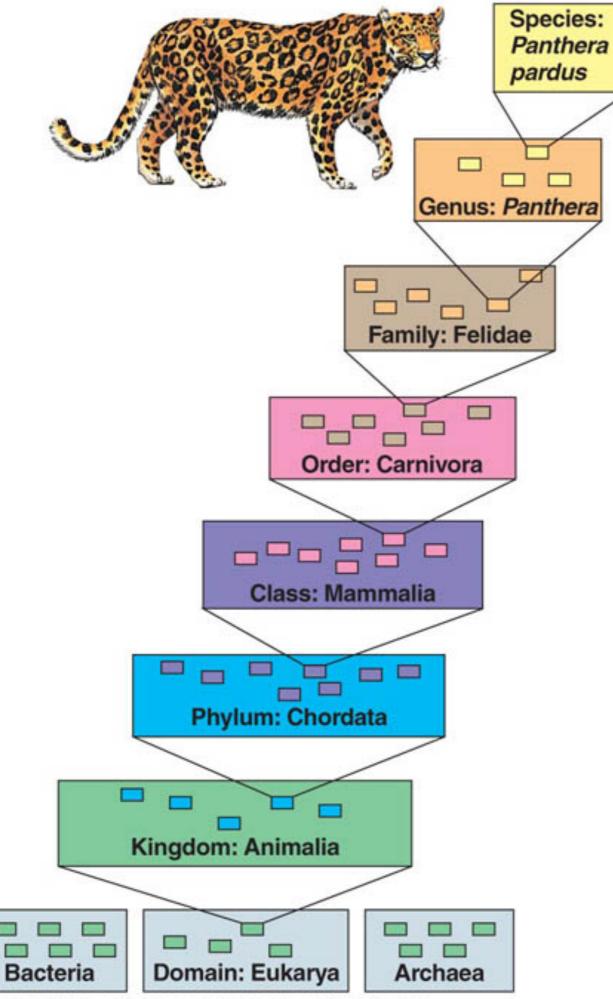
Three main reasons:

physics-centrism of traditional philosophy of science

central place of biology in 20th century science
 conceptual problems within biology

- 'naturalistic turn' in philosophy





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Binomial nomenclature

CAROLI LINNÆI

EQUITIS DE STELLA POLARI, ARCHIATRI REGII, MED. & BOTAN. PROFESS. UPSAL.; ACAD. UPSAL. HOLMENS. PETROPOL. BEROL. IMPER. LOND. MONSPEL. TOLOS. FLORENT. Soc.

SYSTEMA NATURÆ

PER REGNA TRIA NATURA, SECUNDUM

CLASSES, ORDINES, GENERA, SPECIES,

CUM

CHARACTERIBUS, DIFFERENTIIS. SYNONYMIS, LOCIS.

TOMUS I.

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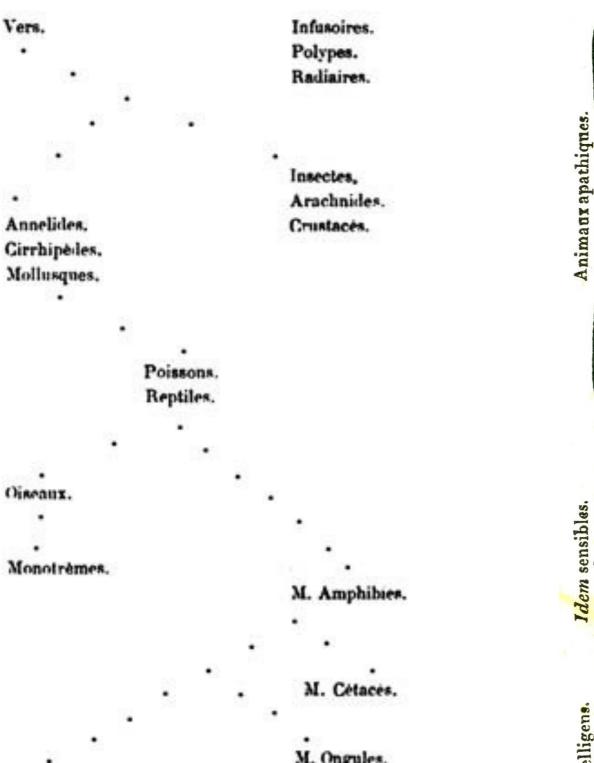
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ANIMAUX



ORDRE présumé de la formation des Animaux, offrant 2 séries séparées, subrameuses. [1] SÉRIE DES ANIMAUX [2] Série des ANIMAUX INARTICULÉS. ARTICULÉS. Infusoires. Polypes. Radiaires. Ascidiens. Vers. Epizoaires.

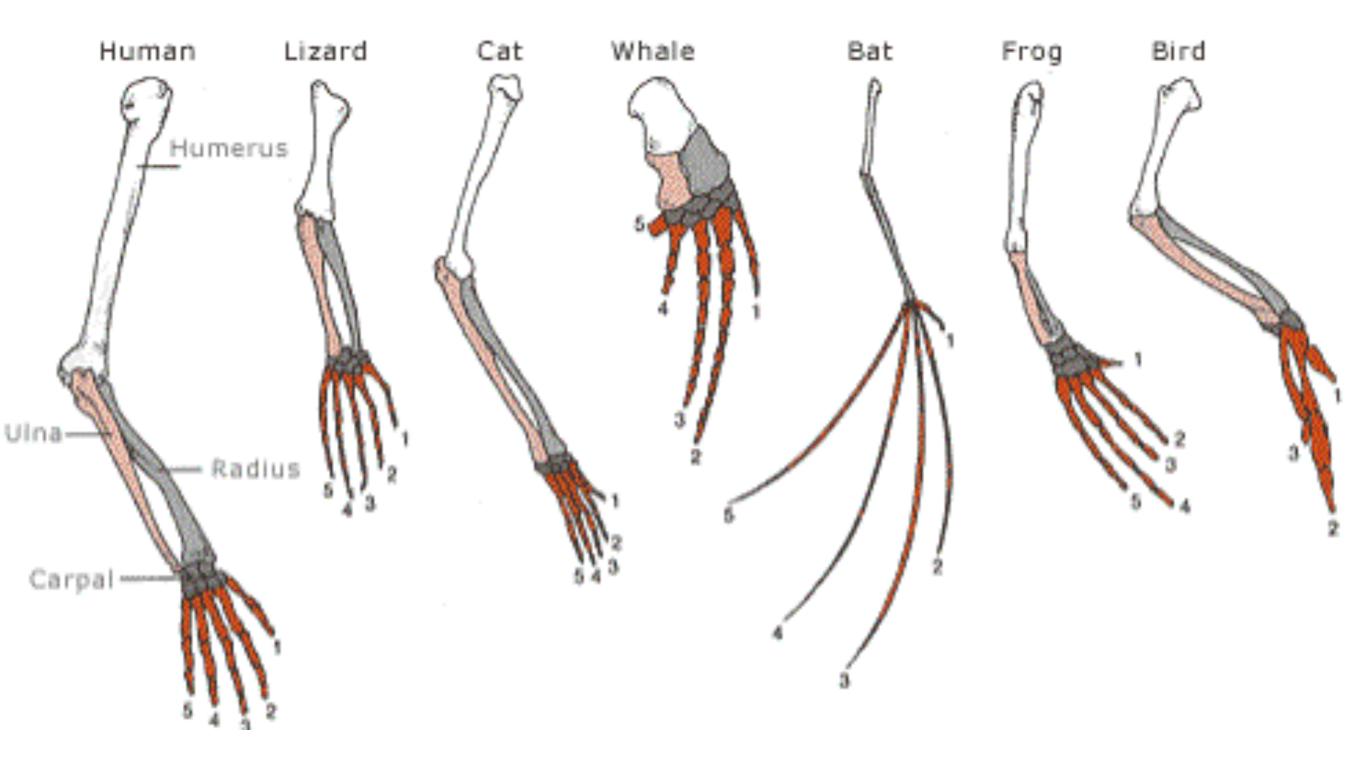
Annelides.

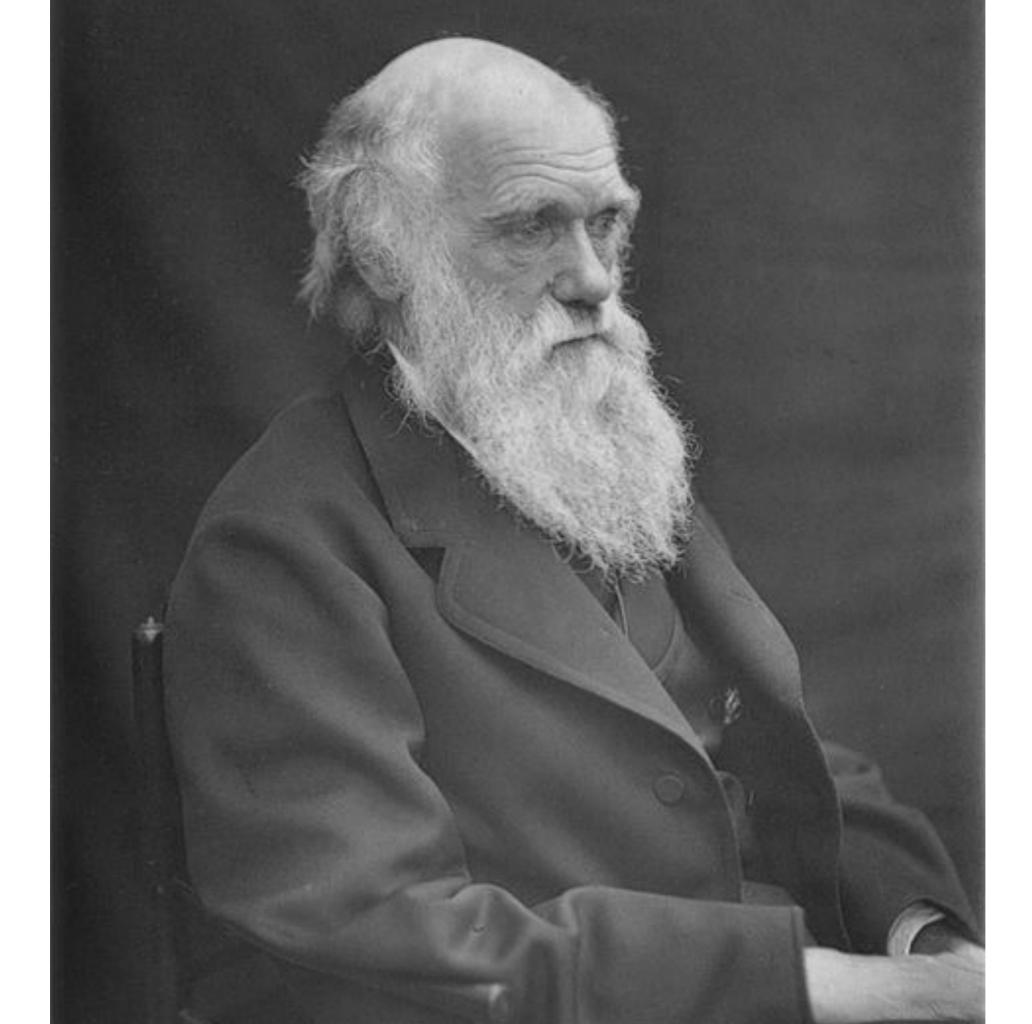
Insectes.

Arachnides. Mollusques. Crustacés. Cirrhipèdes. Poissons. intelligent Reptiles. M. Ongules, Oiseaux. M. Ongaicules. Mammifères. MCH OSTADO d.

Acéphales.

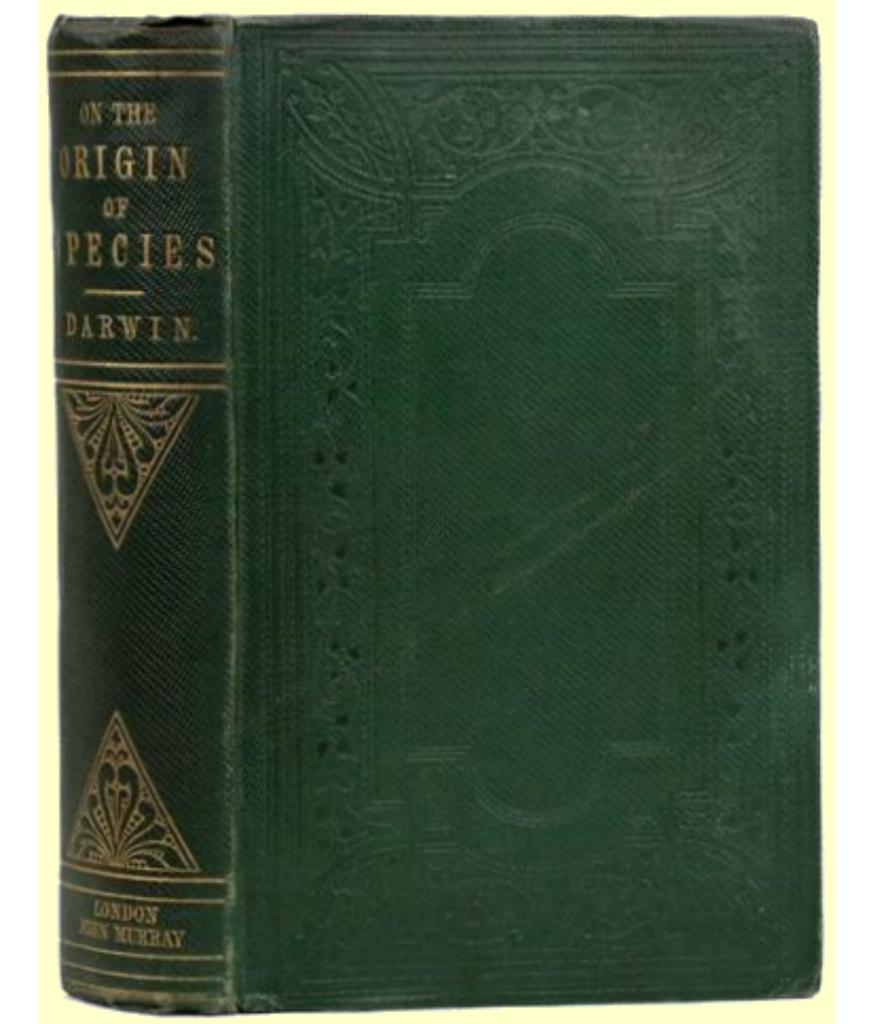
Evolution of animals according to Lamack, as depicted in Zoological Philosophy (1809, left), and in Natural History of Invertebrate Animals (1815, right).





1809 - 1882

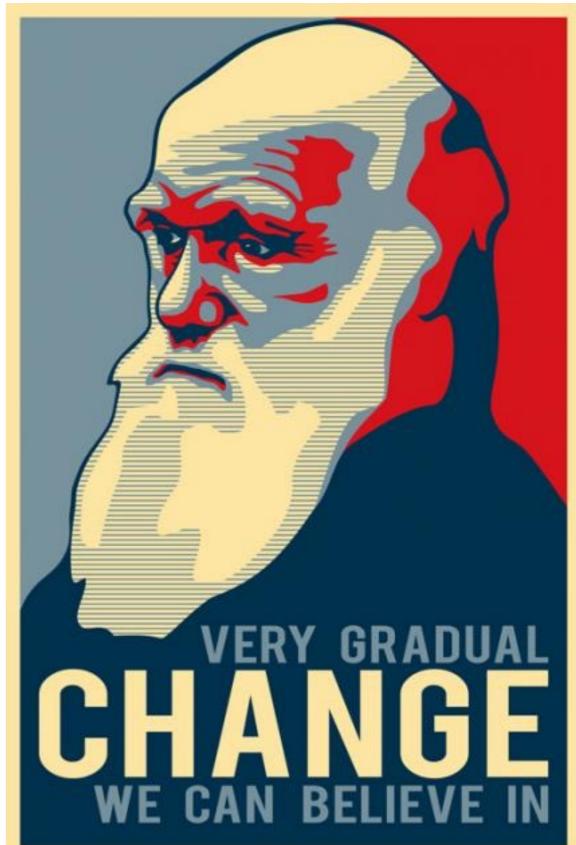
1859 Origin of Species

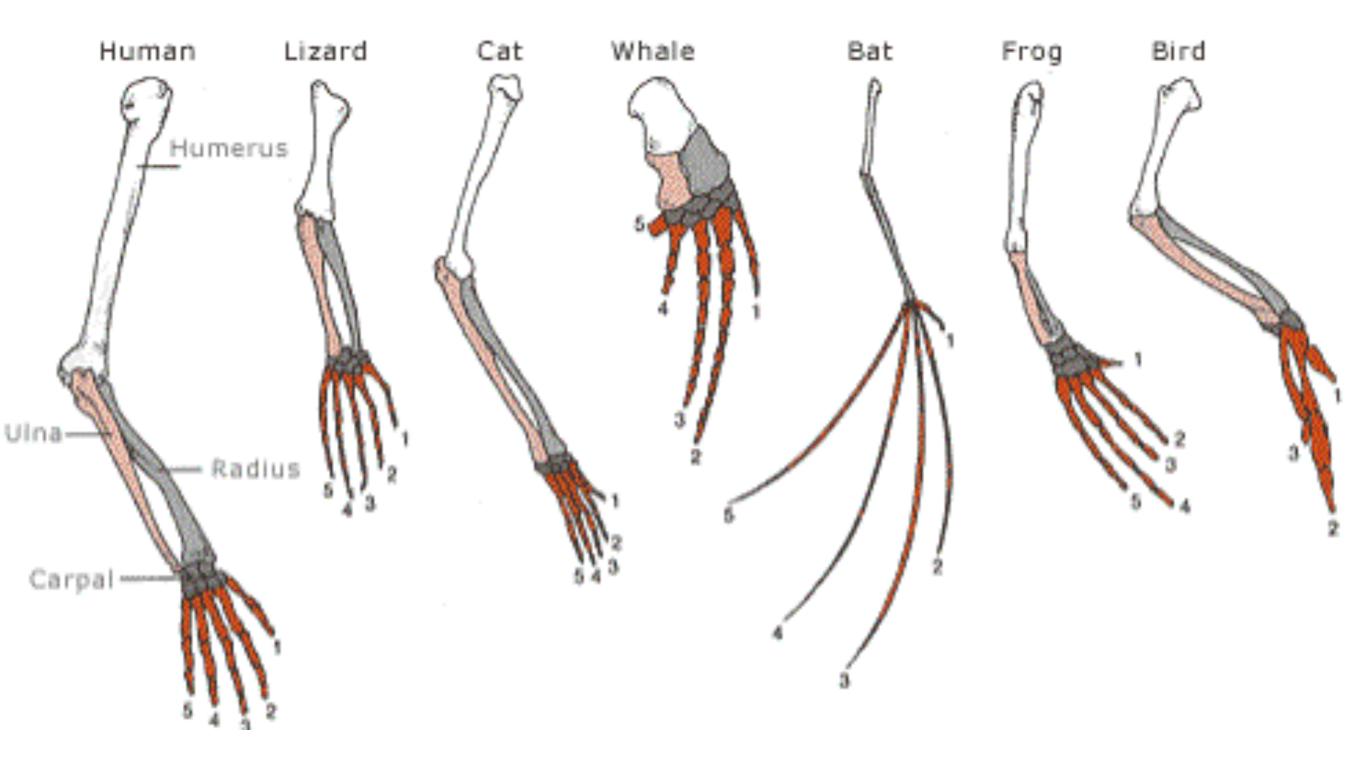


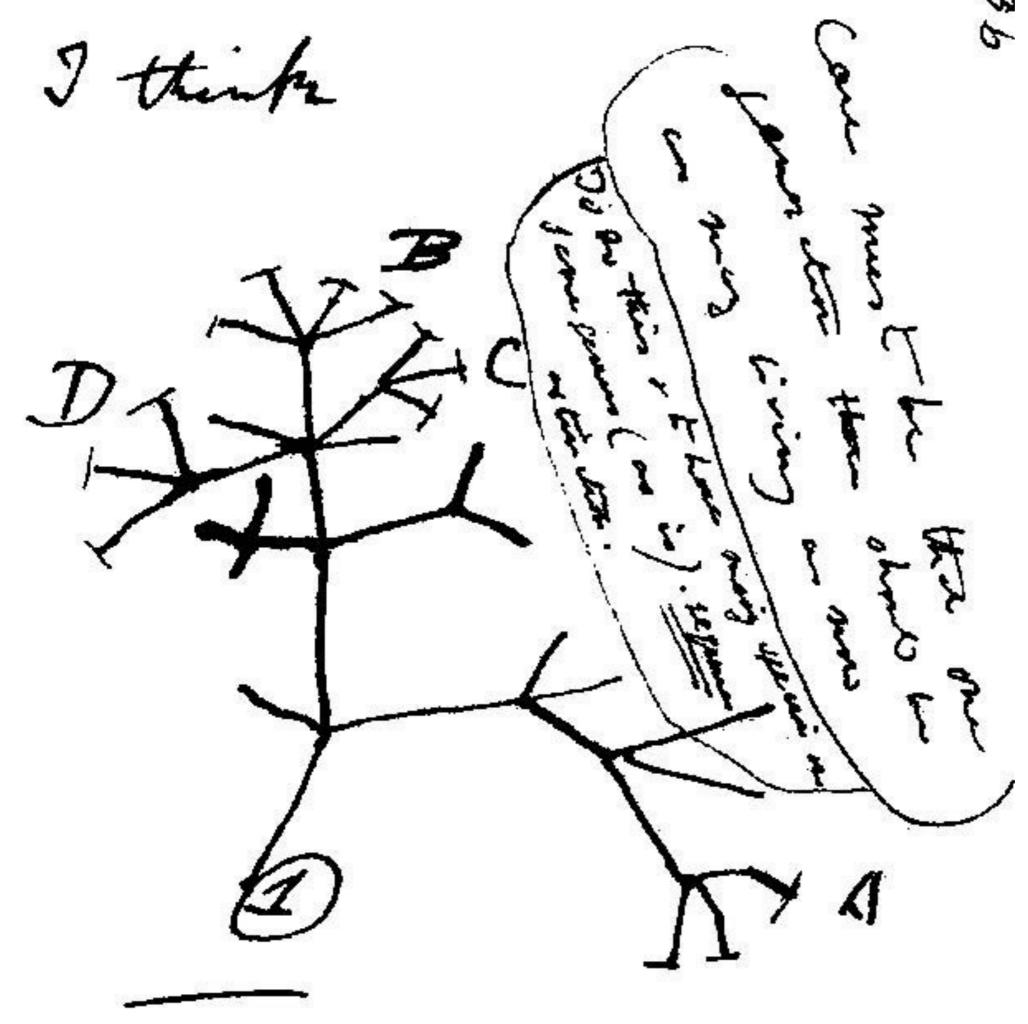


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- 1. Life **evolves**: populations **change** with time
- This change happens gradually (over hundreds or millions of years)



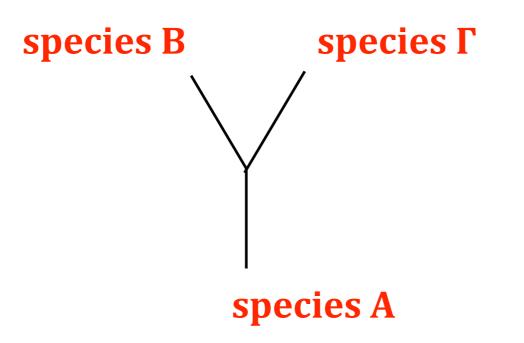


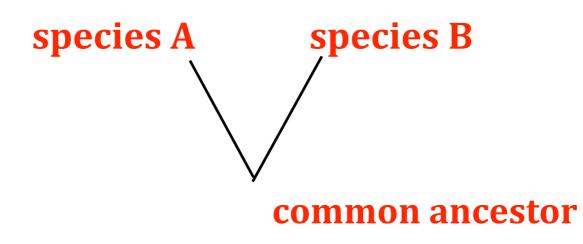


organised beings represent a tree. Irregularly branched some branches far more branched. -Hence Genera. - «as many terminal buds dying, as new ones generated» (1837, Notebook B)

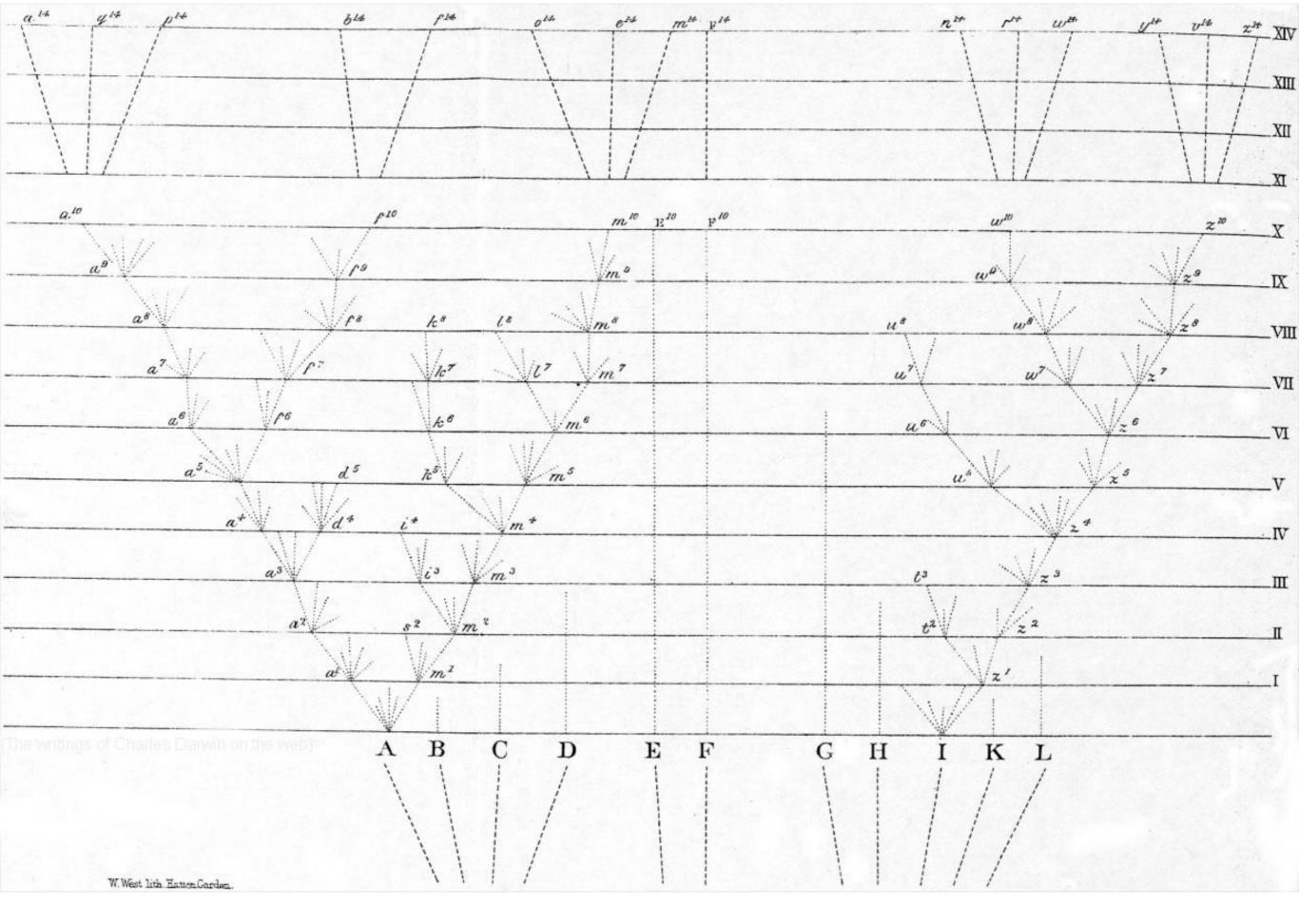
- 1. Life **evolves**: populations **change** with time
- 2. This change happens **gradually** (over hundreds or millions of years)

3. **Speciation** occurs (one species splits into two or more)



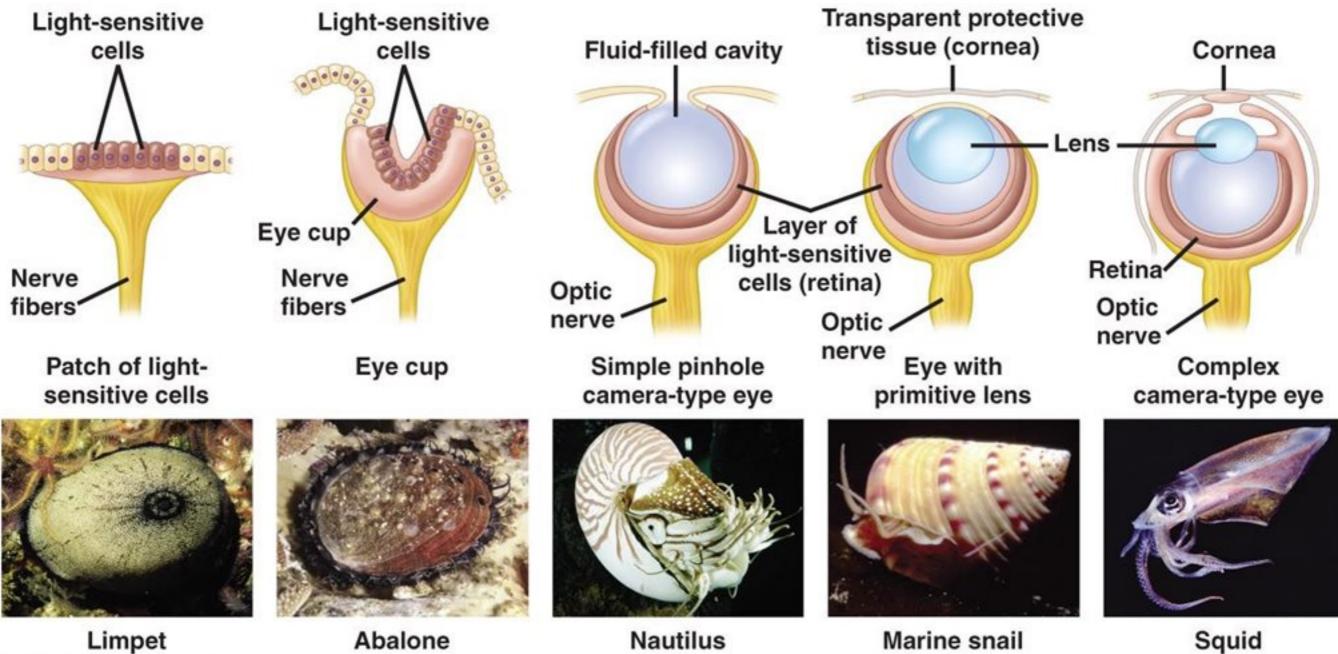


4. All species share a **common ancestor**!



The tree of life, the only picture in Origin

5. Evolution occurs through **natural selection**, which produces adaptations in organisms (organisms appear as if they have been designed)

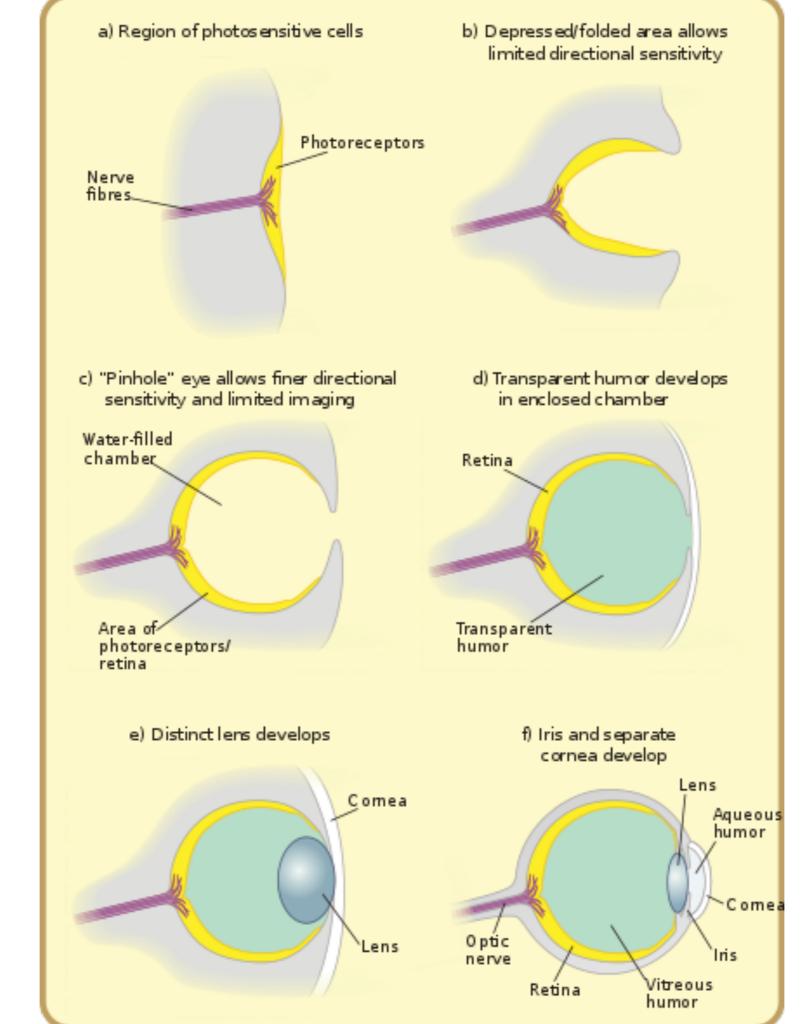


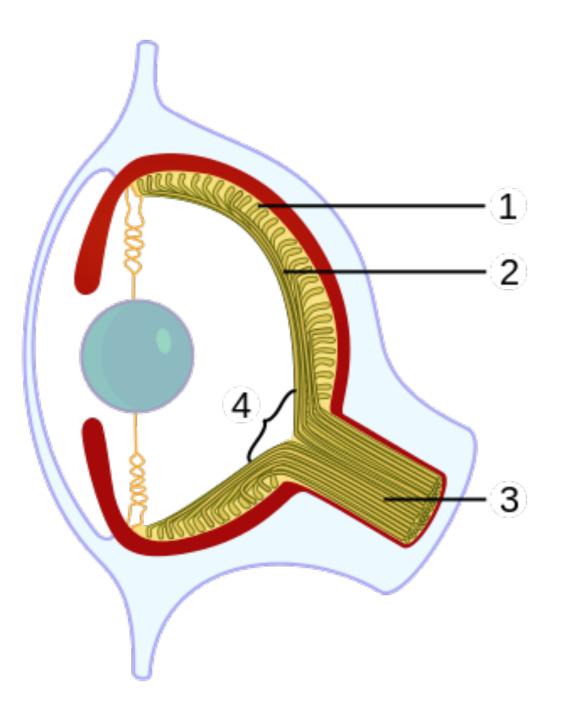
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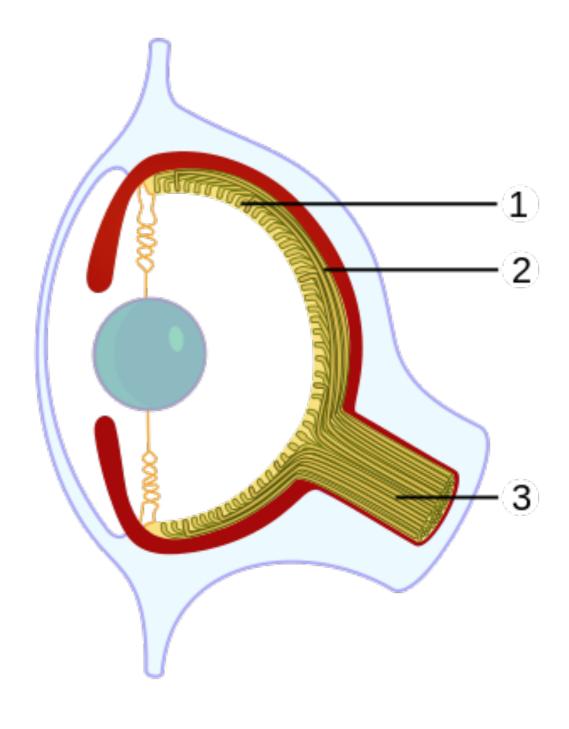
Abalone

Nautilus

Marine snail







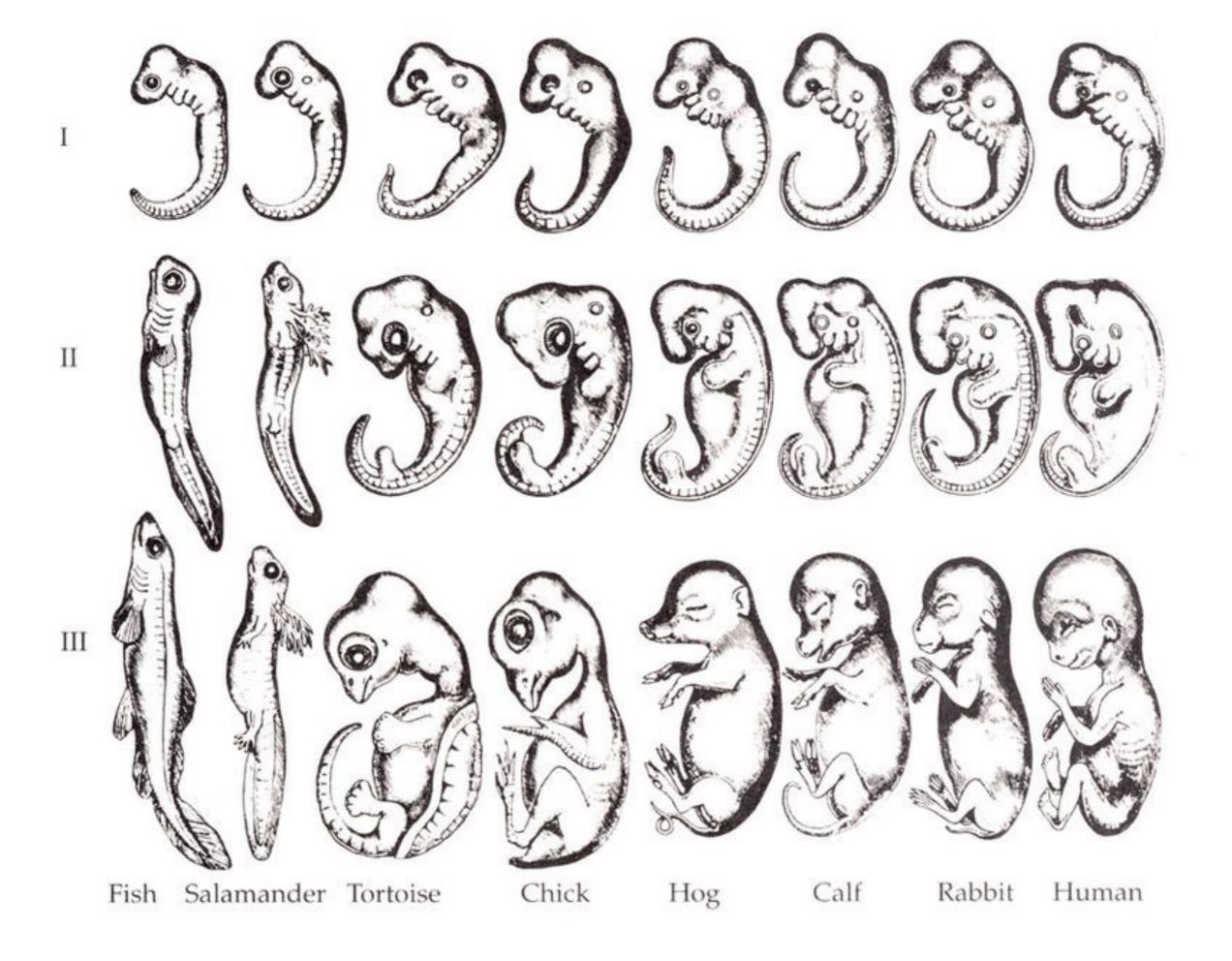
1. Life **evolves**

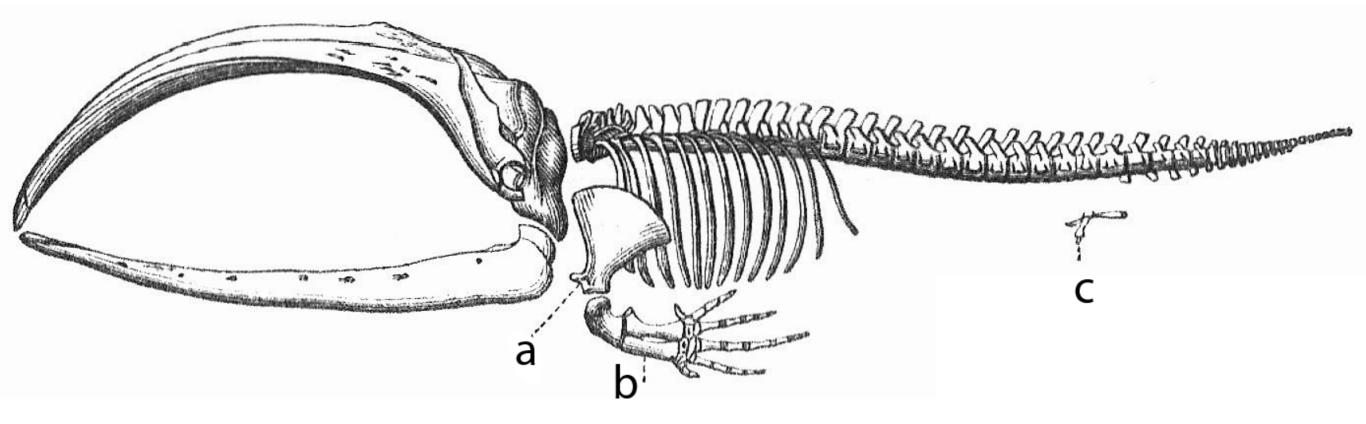
2. This change happens gradually

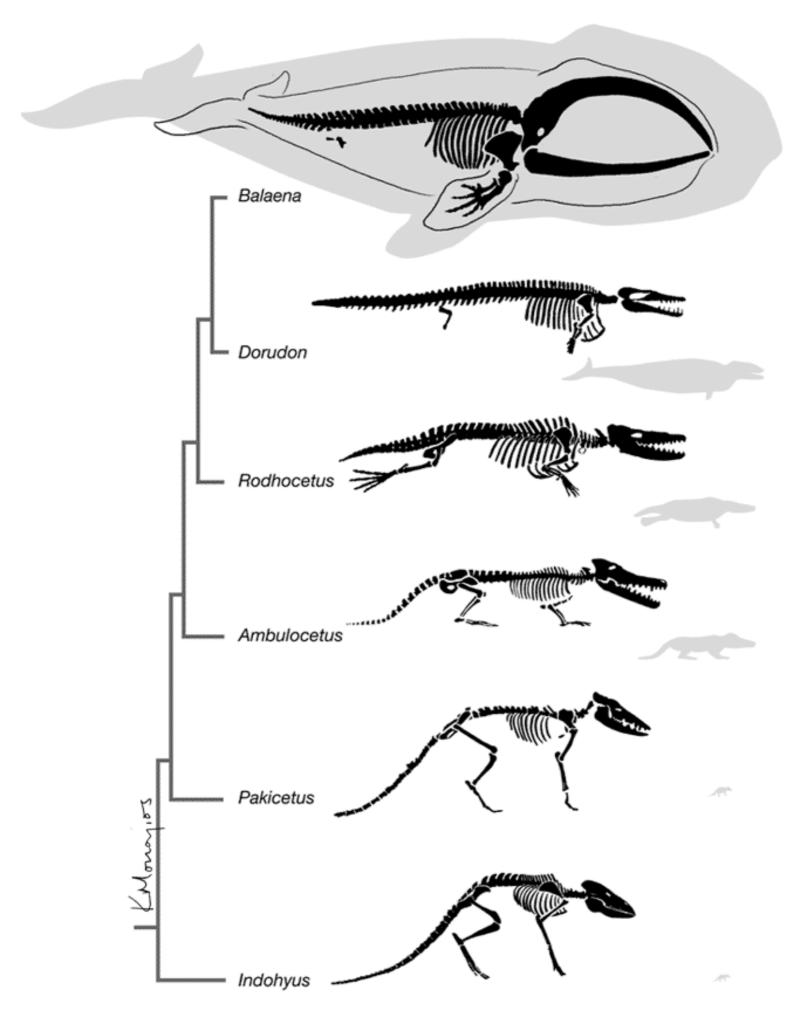
3. Speciation occurs

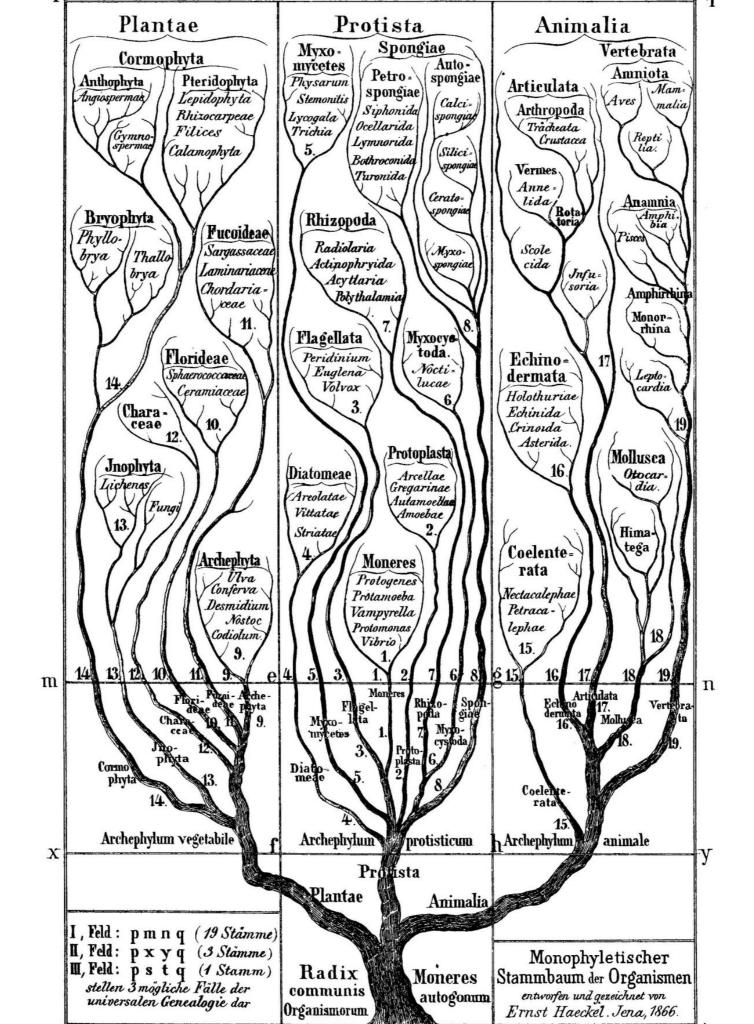
4. All species share a **common ancestor**

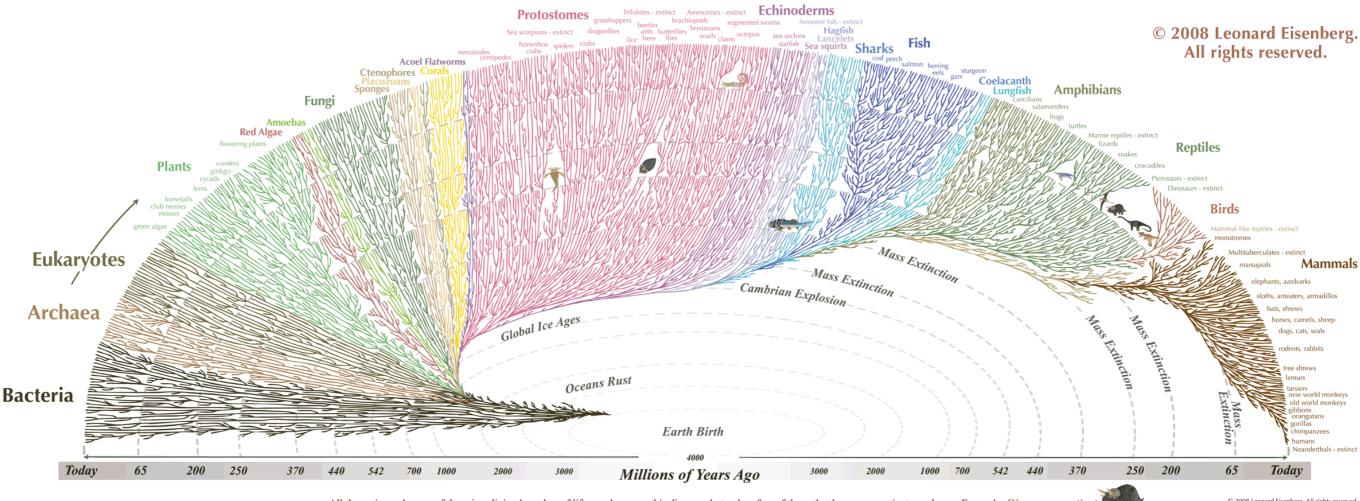
5. Evolution occurs through **natural selection**





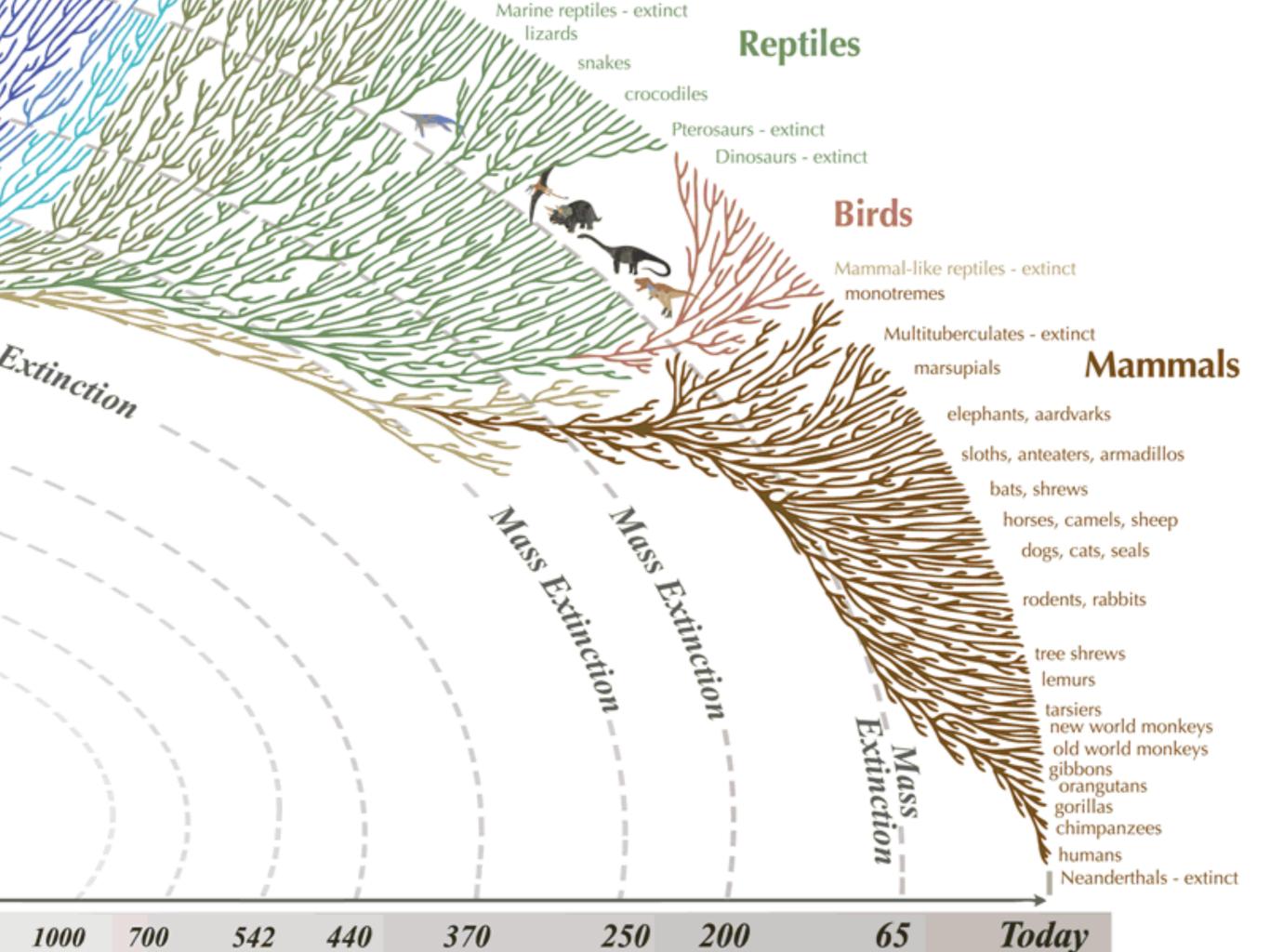


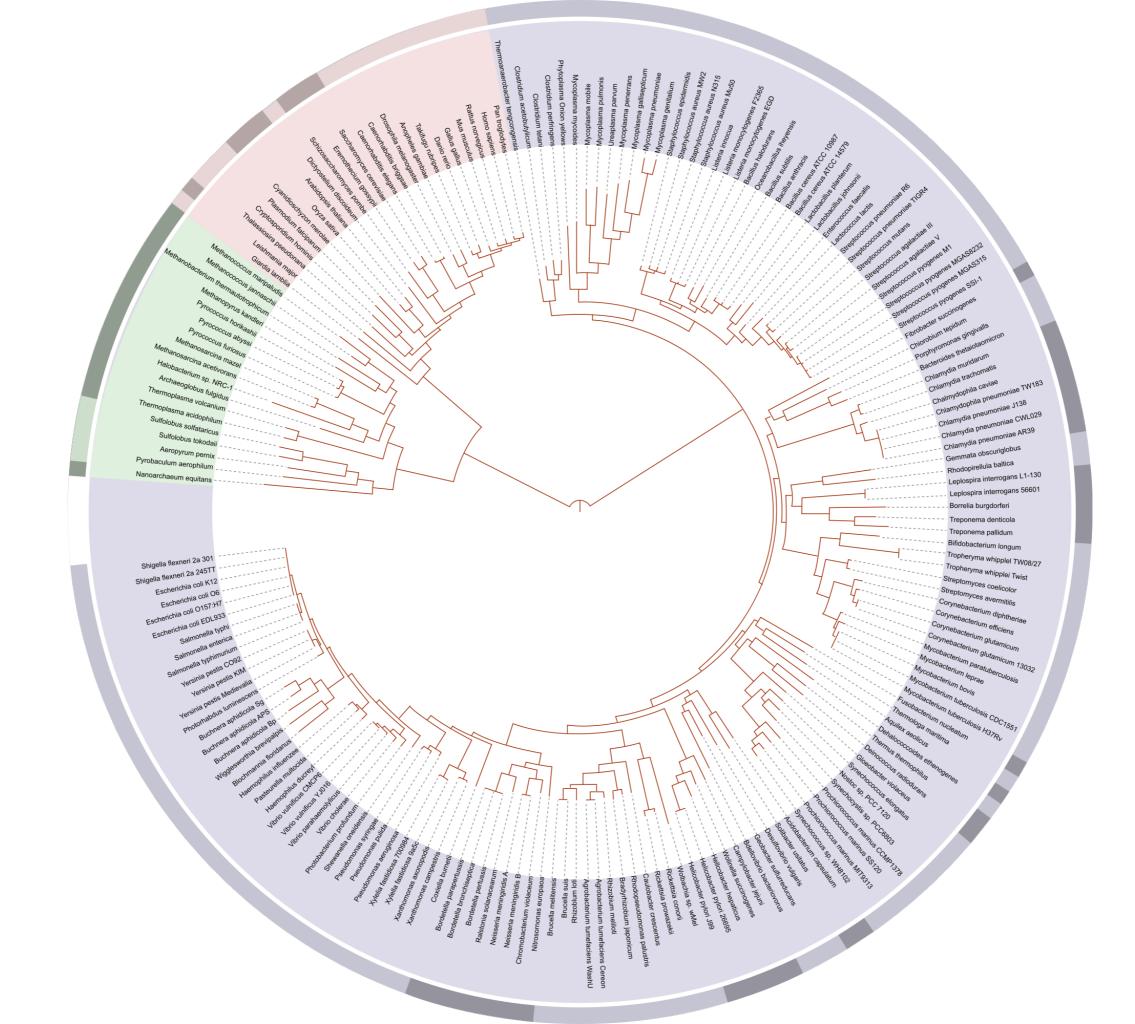




All the major and many of the minor living branches of life are shown on this diagram, but only a few of those that have gone extinct are shown. Example: Dinosaurs - extinct

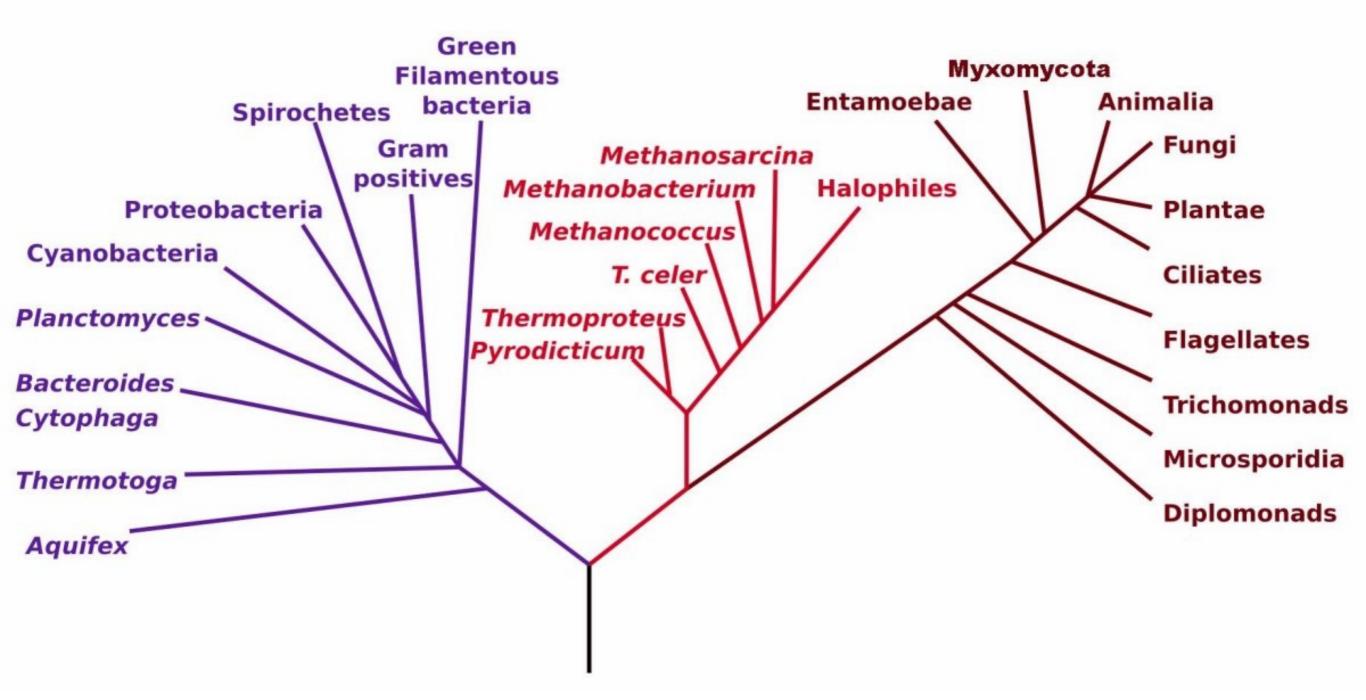
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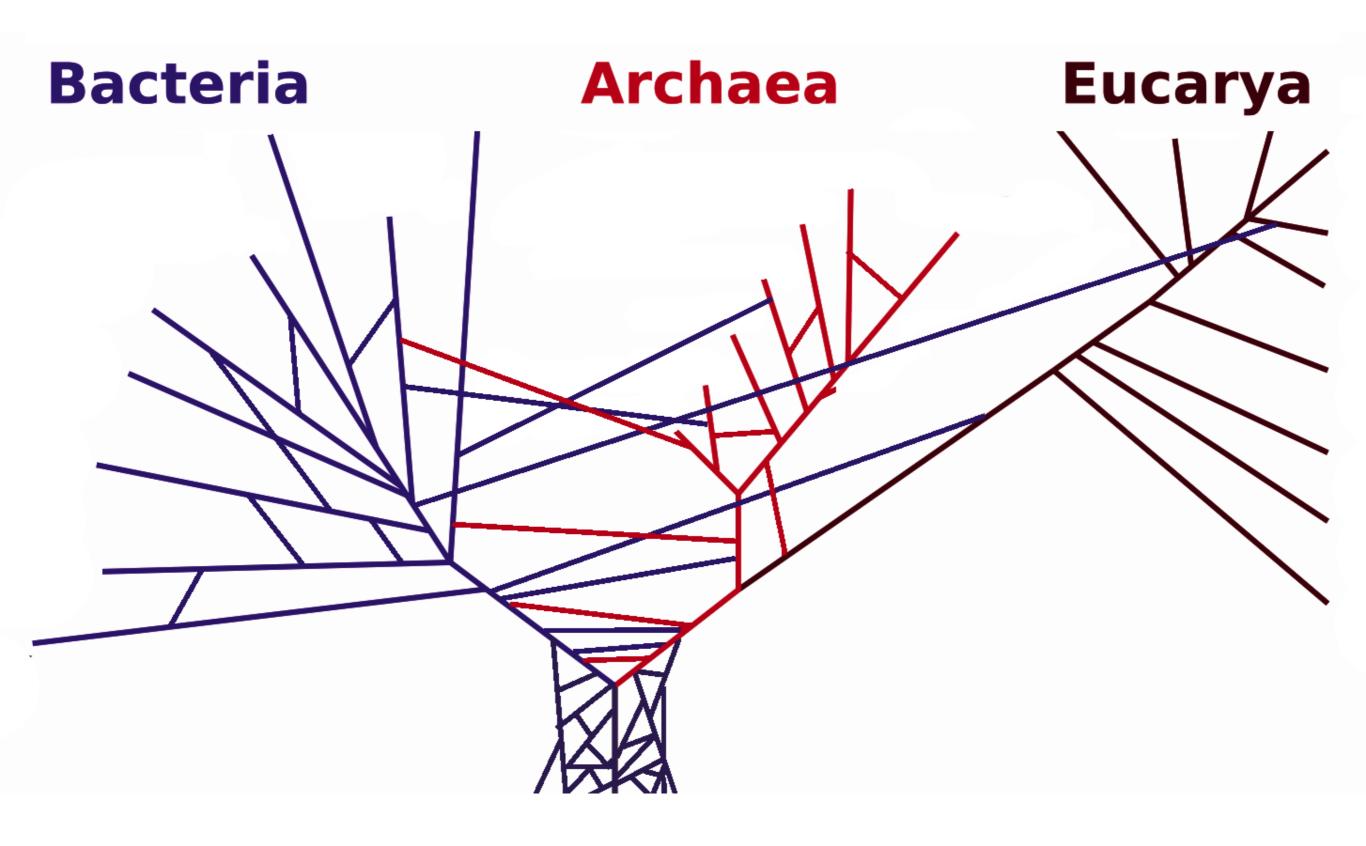


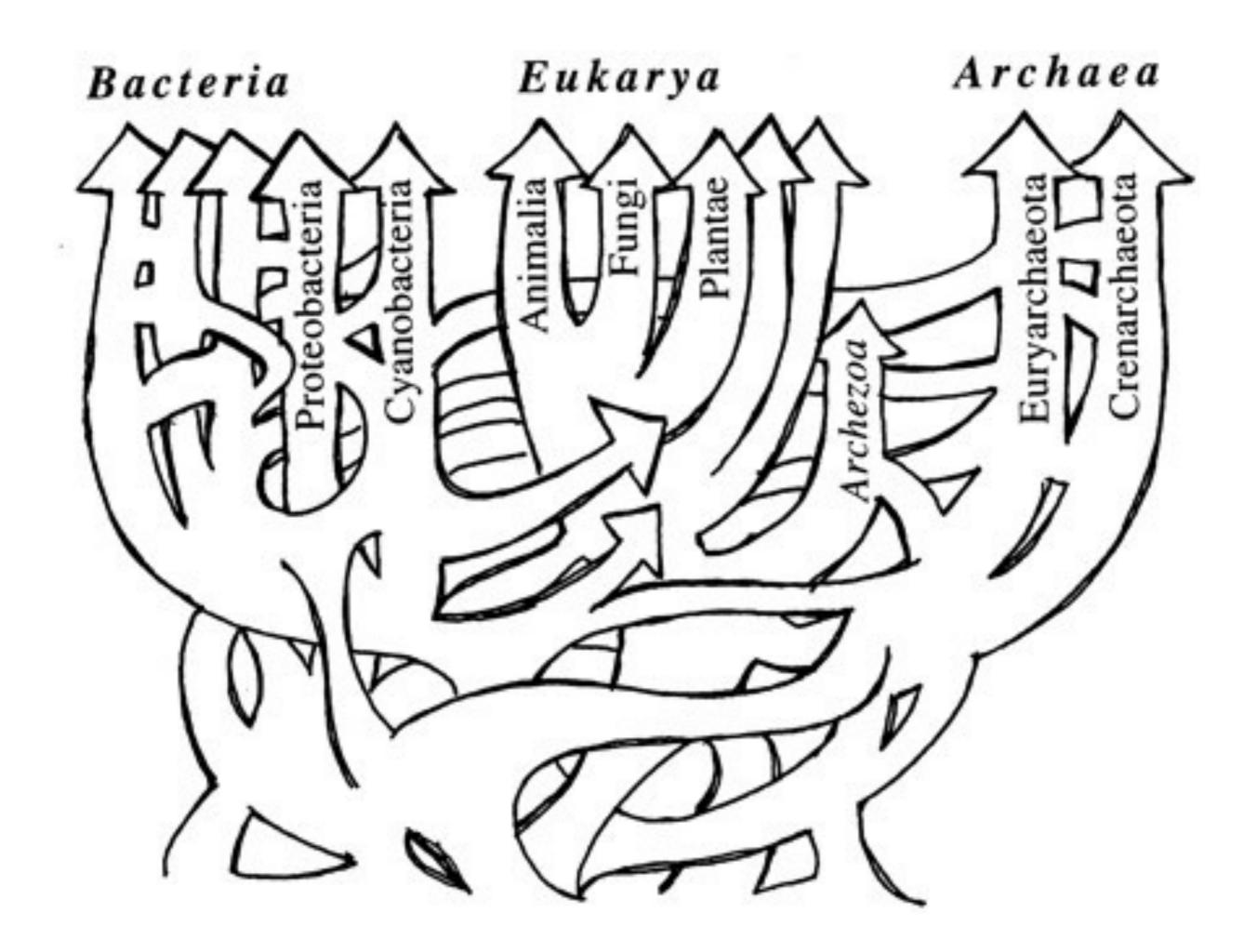


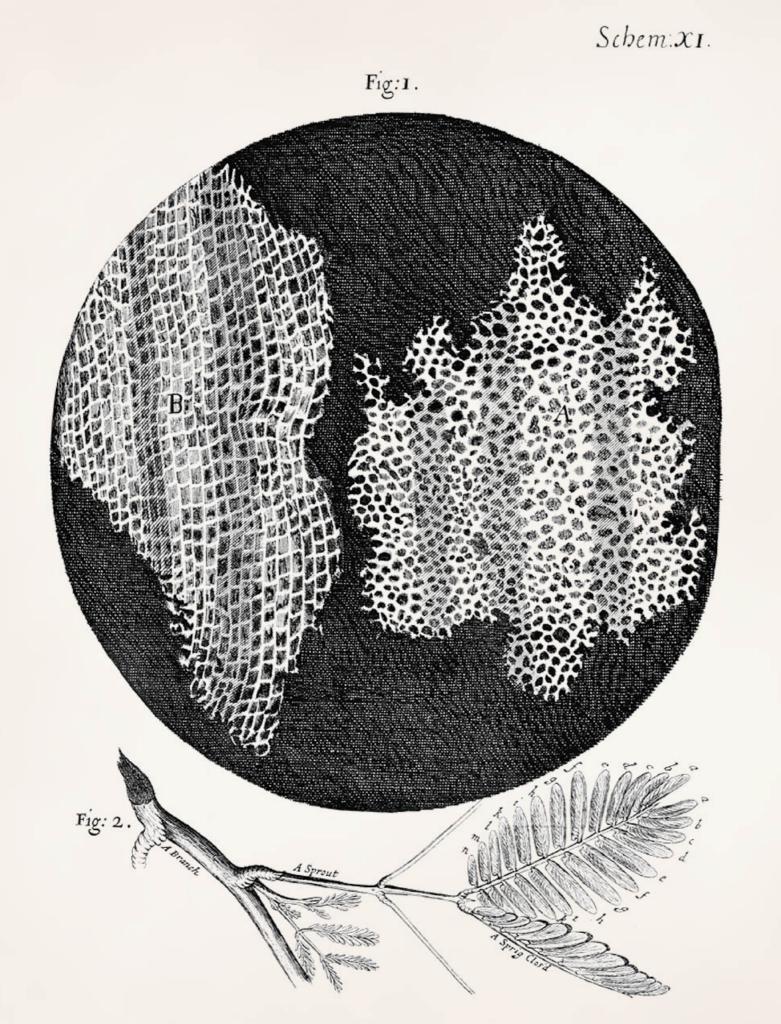
Phylogenetic Tree of Life

Bacteria Archaea Eucarya



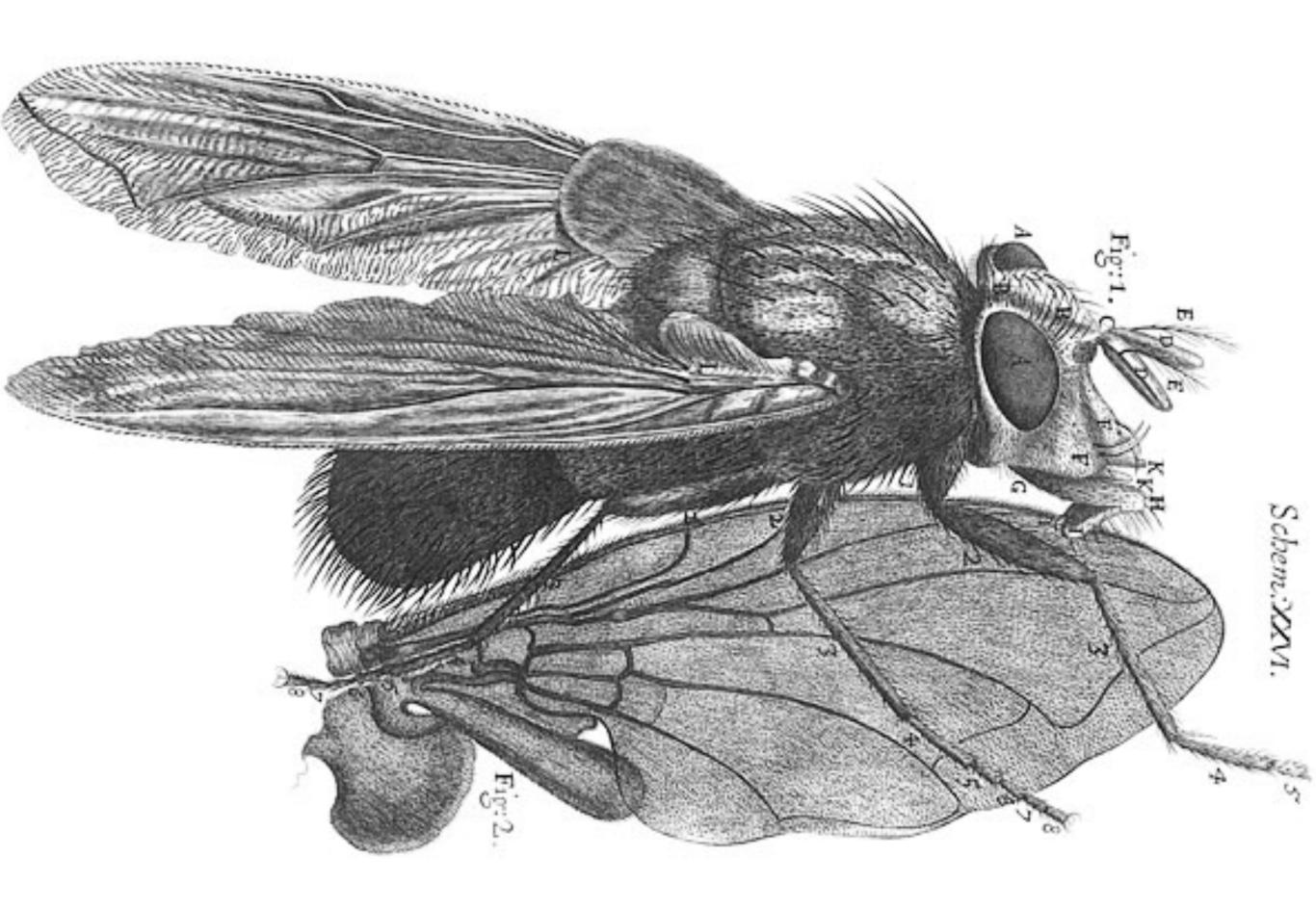








Left: Fig. 1 cork cells Right: Hooke's microscope



Cell theory

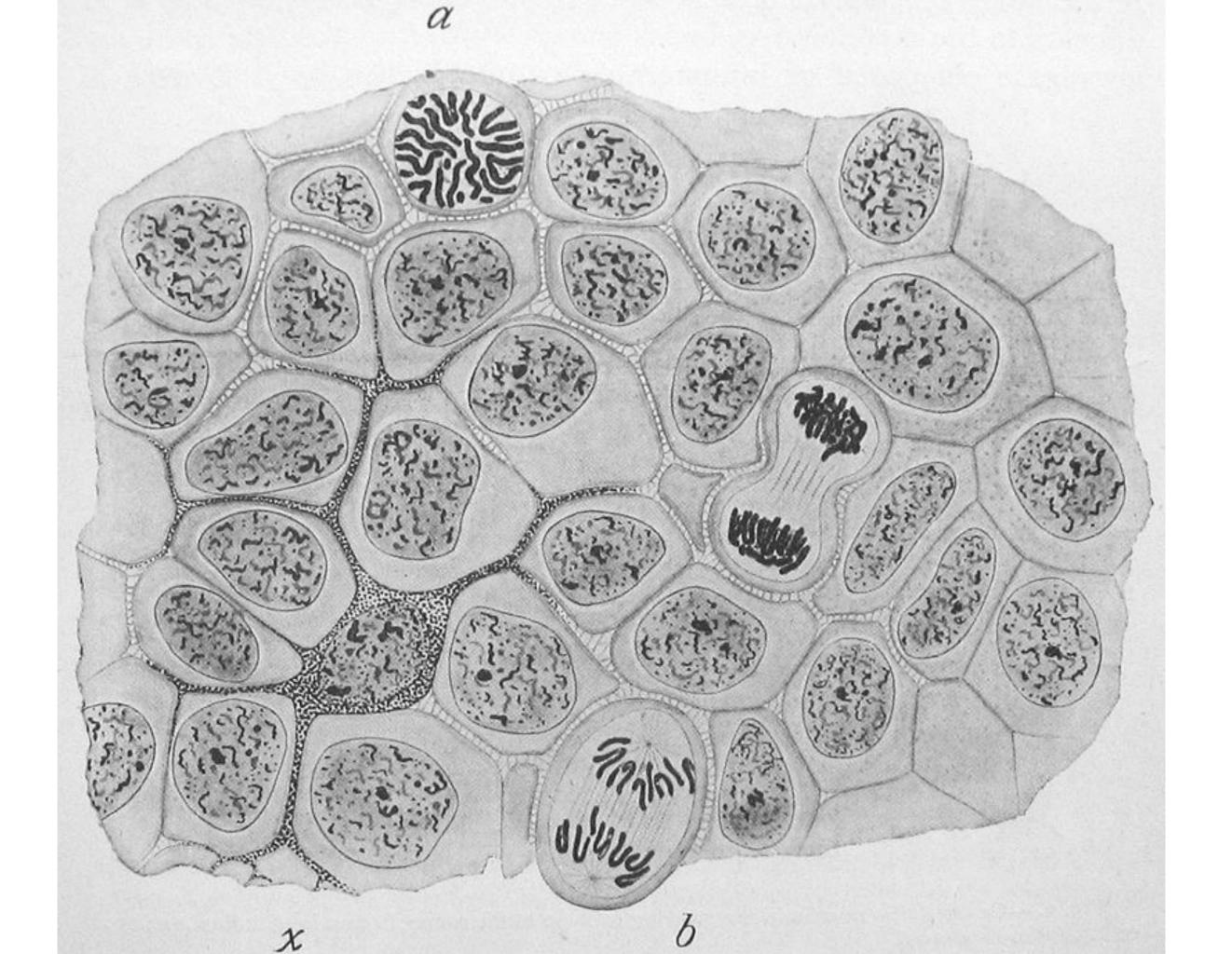
-For the biologists of the **2nd half of the 19th century**, the cell was the **key** to all biological problems.

-> all vital **functions** (**metabolism, development, reproduction, heredity**) could be considered as **cellular activities** - as activities that occur inside cells.

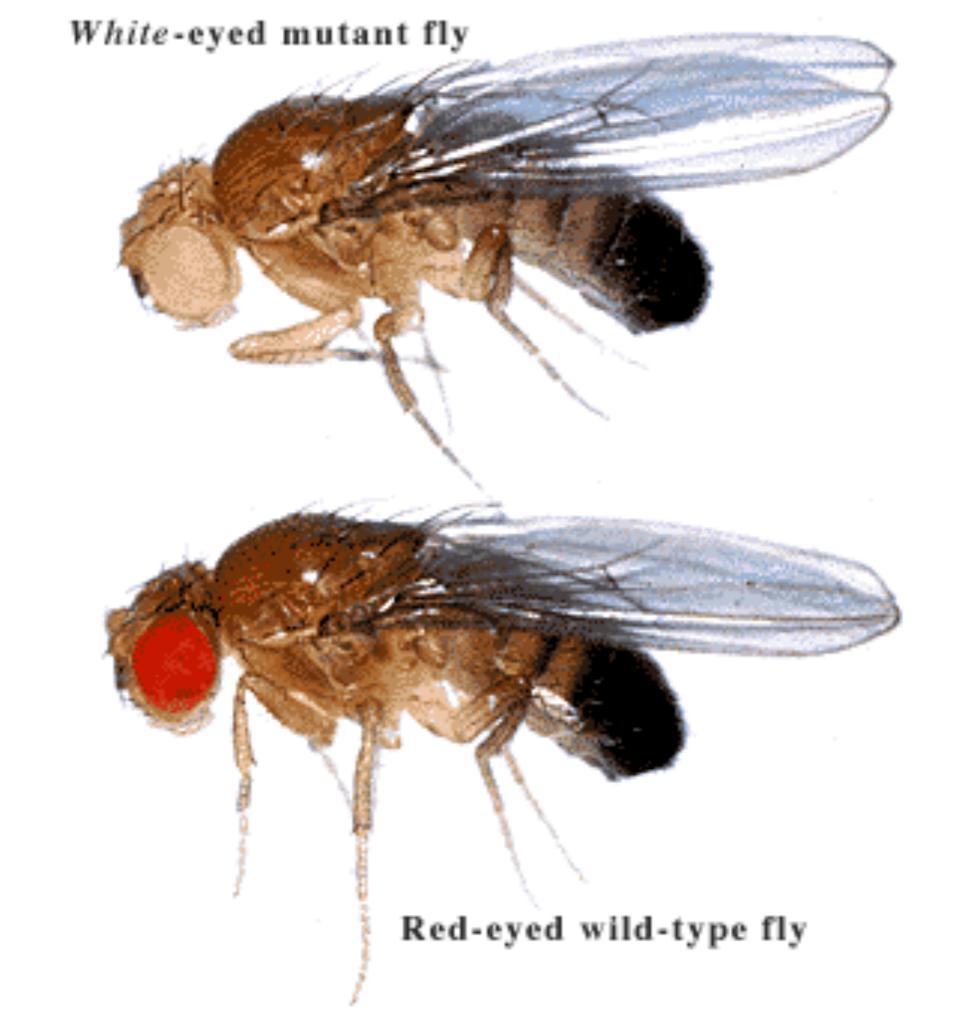
The cell theory consisted of **3 principles**:

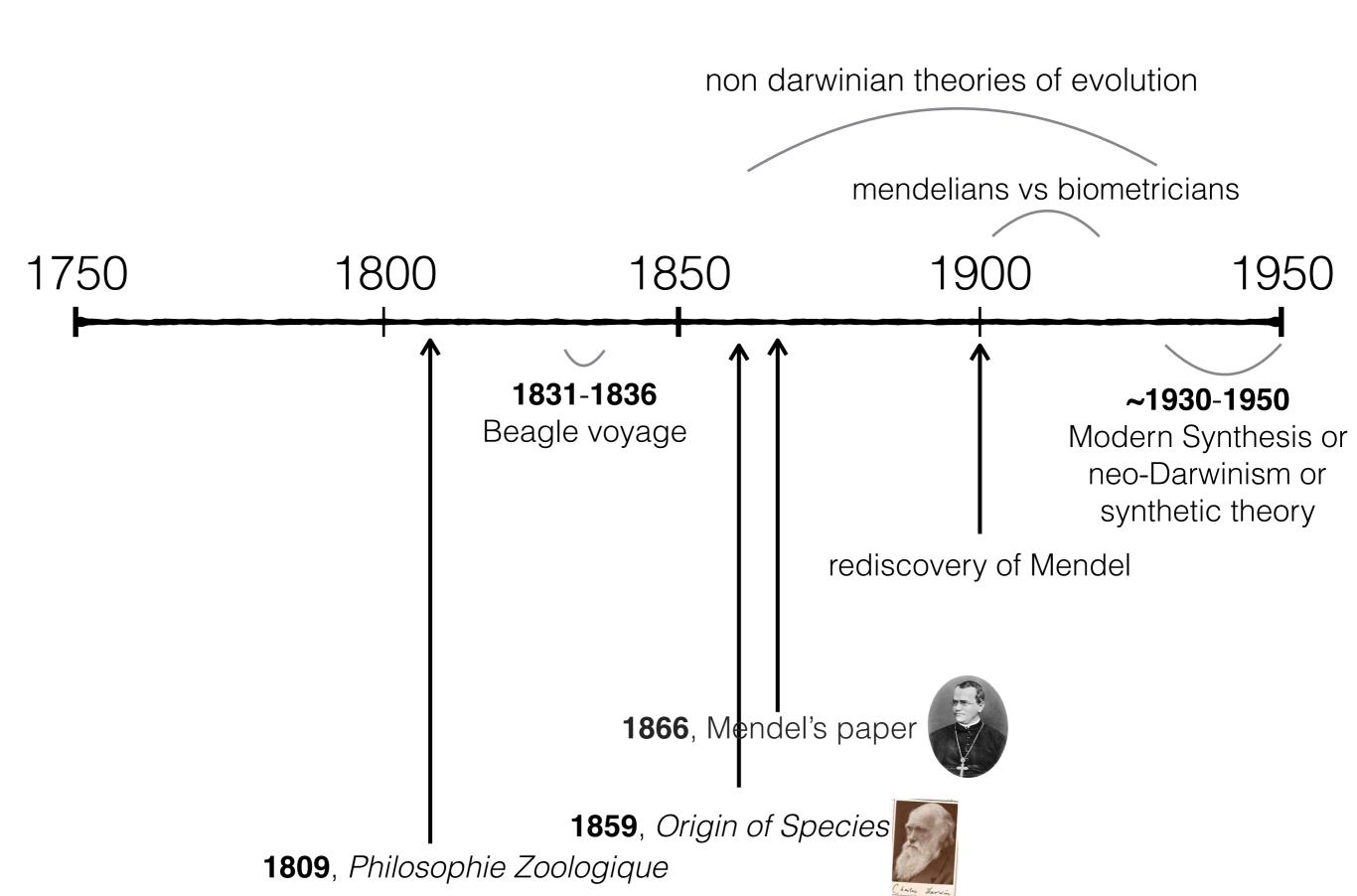
- 1. all plants and animals are made of cells
- 2. cells have **all the characteristics of life** (e.g. metabolism, growth, reproduction)
- 3. all cells arise **by division of preexisting cells**

-> the cell as the universal **structural** and **functional** unit









Modern Synthesis or neo-Darwinism

-evolutionary synthesis or Modern Synthesis or neo-Darwinism or synthetic theory

- -> synthesis of darwinism and mendelism
 -> convergence between naturalists and geneticists
- -> <u>central event</u> in the history of biology of **20th century**
- -the evolutionary synthesis took place during the 30s and 40s

-for many biologists it constitutes even today the **fundamental framework in evolutionary biology**

-the phrase 'evolutionary synthesis' comes from **Julian Huxley's** (1887–1975) book: *Evolution: the Modern Synthesis* (1942)

Modern Synthesis or neo-Darwinism

2 main theses:

-> evolution occurs natural selection that acts on variation that results from mutation and recombination

-> the phenomena observed by **paleontologists**, **systematists**, and **field biologists** can be explained in a manner **compatible** with **already known genetic mechanisms**

-'**merging**' of different specialties (and not a **replacement** of one 'paradigm' with another)

-conditions of the 2nd world war **favourable** for such a synthesis, as **research** in the field and in labs is **hindered**

Modern Synthesis or neo-Darwinism

-a central tenet of the **synthetic theory**:

-> natural populations contain a large gene pool, and thus enough variation for natural selection to act

new mutations are rarely the direct source of variation on which natural selection acts

- variation mostly results from genetic recombination

 there is no relationship between mutation rate and the direction of evolution

-> this view spread during the **40s**

Population Genetics

- -emergence of **population genetic** in the 1920s
- -quantitative models of evolutionary processes
- -evolution as change in gene frequencies in populations
- [Hardy-Weinberg equilibrium]
- -> natural selection as a force that changes gene frequencies
- -> R. A. **Fisher** (1890–1962) & J. B. S. **Haldane** (1889–1988) in UK -> Sewall **Wright** (1892–1964) in USA
- -founders of theoretical **population genetics**

Population Genetics

Haldane:

-series of articles **between 1924 & 1931**, in which he constructs **population genetics models**

-> in **1924** publishes a classic example of evolution, the case of the **peppered moth** (Amphidasys betularia)

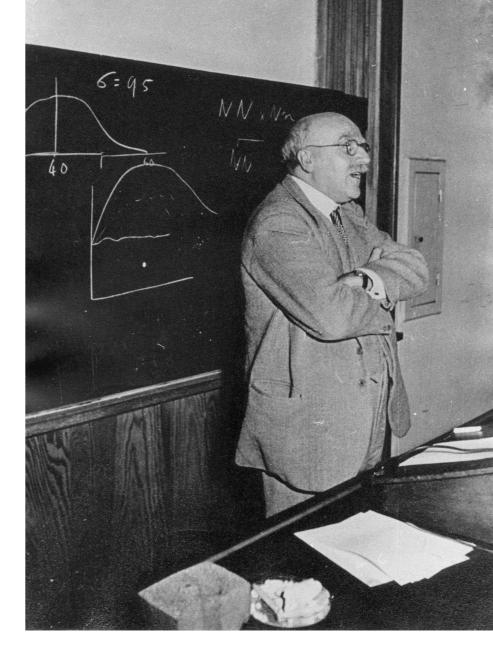
-> shows the **power** of **natural selection**

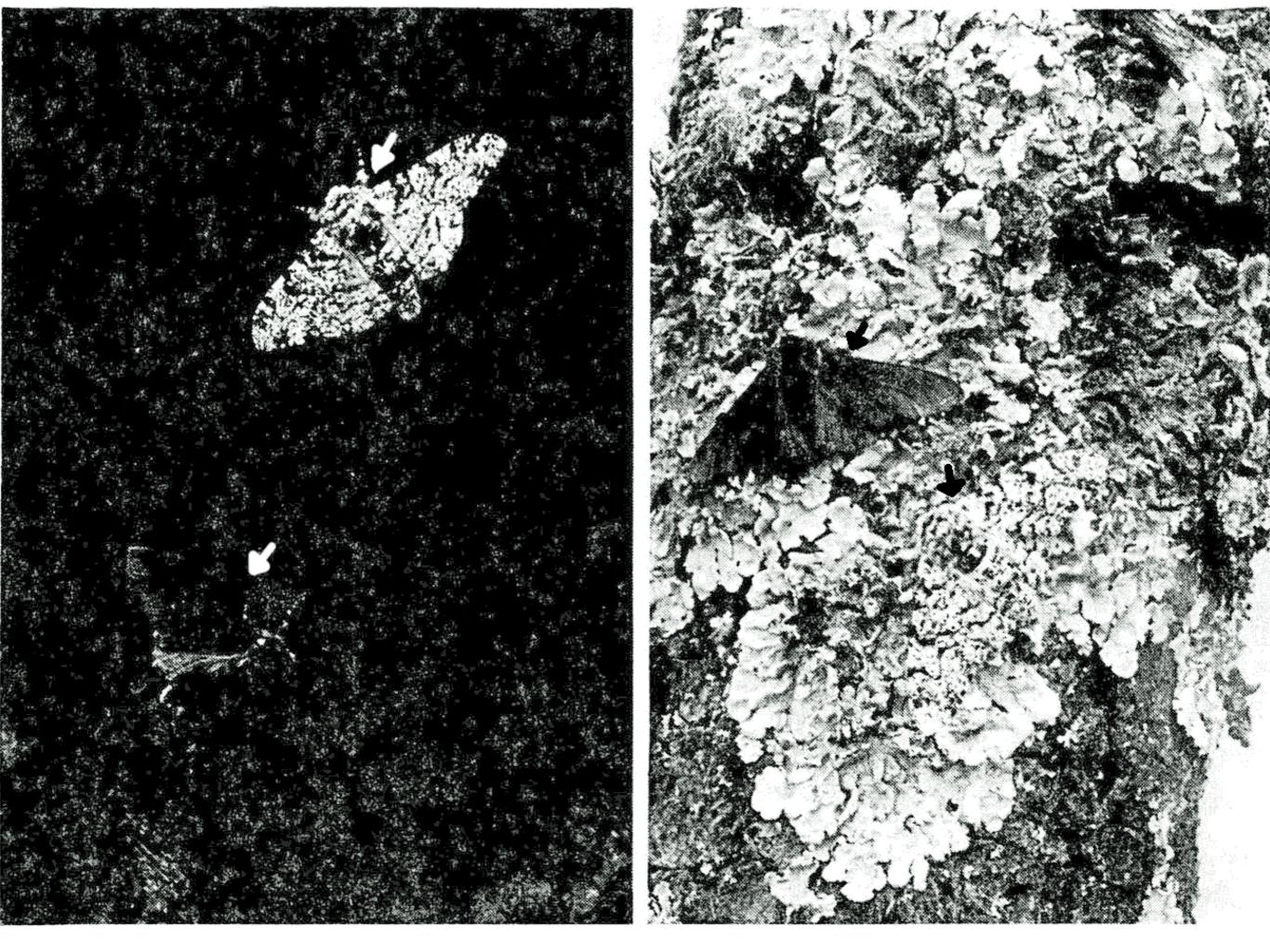
[-> frequency of melanic form < 1% in 1848, until 1901 ~ 99%
-> calculates that melanic form must have had 50% selective advantage]

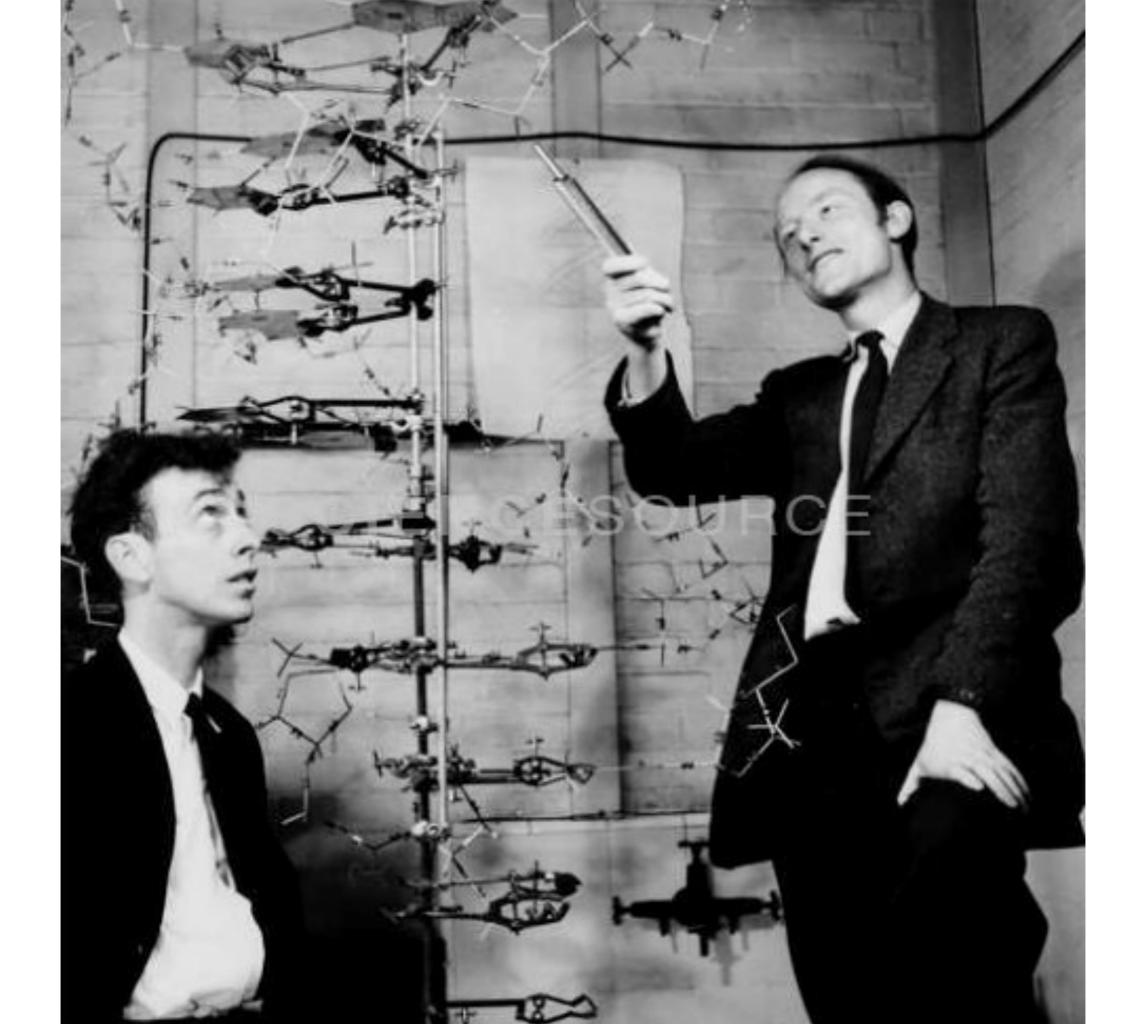
-Main conclusion of population geneticists:

-> even a small genetic difference could spread quickly through the population

-> many neo-Lamarckians are convinced (e.g. Ernst Mayr, Bernhard Rensch)







equipment, and to Dr. G. E. R. Deacon and the captain and officers of R.R.S. Discovery II for their part in making the observations,

NATURE

Young, F. B., Gerrard, H., and Jevons, W., Phil. Mag., 40, 149 Longuet-Higgins, M. S., Mon. Not. Roy. Astro. Soc., Geophys. Supp., 5, 285 (1949).

¹ Von Arx, W. S., Woods Hole Papers in Phys. Ocearog. Meteor., 11 (3) (1950).

*Ekman, V. W., Arkiv. Mat. Astron. Fysik. (Stockholm), 2 (11) (1905).

MOLECULAR STRUCTURE OF NUCLEIC ACIDS

A Structure for Deoxyribose Nucleic Acid

WE wish to suggest a structure for the salt of deoxyribose nucleic acid (D.N.A.). This structure has novel features which are of considerable biological interest.

A structure for nucleic acid has already been proposed by Pauling and Corey¹. They kindly made their manuscript available to us in advance of publication. Their model consists of three intertwined chains, with the phosphates near the fibre axis, and the bases on the outside. In our opinion, this structure is unsatisfactory for two reasons : (1) We believe that the material which gives the X-ray diagrams is the salt, not the free acid. Without the acidic hydrogen atoms it is not clear what forces would hold the structure together, especially as the negatively charged phosphates near the axis will repel each other. (2) Some of the van der Waals distances appear to be too small.

Another three-chain structure has also been suggested by Fraser (in the press). In his model the phosphates are on the outside and the bases on the inside, linked together by hydrogen bonds. This structure as described is rather ill-defined, and for

this reason we shall not comment on it.

We wish to put forward a radically different structure for the salt of deoxyribose nucleic acid. This structure has two helical chains each coiled round the same axis (see diagram). We have made the usual chemical assumptions, namely, that each chain consists of phosphate diester groups joining β-p-deoxyribofuranose residues with 3',5' linkages. The two chains (but not their bases) are related by a dyad perpendicular to the fibre axis. Both chains follow righthanded helices, but owing to the dyad the sequences of the atoms in the two chains run in opposite directions. Each chain loosely resembles Furberg's² model No. 1; that is, the bases are on the inside of the helix and the phosphates on the outside. The configuration of the sugar and the atoms near it is close to Furberg's 'standard configuration', the sugar being roughly perpendicular to the attached base. There

is a residue on each chain every 3.4 A. in the z-direction. We have assumed an angle of 36° between adjacent residues in the same chain, so that the structure repeats after 10 residues on each chain, that is, after 34 A. The distance of a phosphorus atom from the fibre axis is 10 A. As the phosphates are on the outside, cations have easy access to them.

The structure is an open one, and its water content is rather high. At lower water contents we would expect the bases to tilt so that the structure could become more compact.

The novel feature of the structure is the manner in which the two chains are held together by the purine and pyrimidine bases. The planes of the bases are perpendicular to the fibre axis. They are joined together in pairs, a single base from one chain being hydrogen-bonded to a single base from the other chain, so that the two lie side by side with identical z-co-ordinates. One of the pair must be a purine and the other a pyrimidine for bonding to occur. The hydrogen bonds are made as follows : purine position 1 to pyrimidine position 1; purine position 6 to pyrimidine position 6.

If it is assumed that the bases only occur in the structure in the most plausible tautomeric forms (that is, with the keto rather than the enol configurations) it is found that only specific pairs of bases can bond together. These pairs are : adenine (purine) with thymine (pyrimidine), and guanine (purine) with cytosine (pyrimidine).

In other words, if an adenine forms one member of a pair, on either chain, then on these assumptions the other member must be thymine ; similarly for guanine and cytosine. The sequence of bases on a single chain does not appear to be restricted in any way. However, if only specific pairs of bases can be formed, it follows that if the sequence of bases on one chain is given, then the sequence on the other chain is automatically determined.

It has been found experimentally^{3,4} that the ratio of the amounts of adenine to thymine, and the ratio of guanine to cytosine, are always very close to unity for deoxyribose nucleic acid.

It is probably impossible to build this structure with a ribose sugar in place of the deoxyribose, as the extra oxygen atom would make too close a van der Waals contact.

The previously published X-ray data5,6 on deoxyribose nucleic acid are insufficient for a rigorous test of our structure. So far as we can tell, it is roughly compatible with the experimental data, but it must be regarded as unproved until it has been checked against more exact results. Some of these are given in the following communications. We were not aware of the details of the results presented there when we devised our structure, which rests mainly though not entirely on published experimental data and stereochemical arguments.

It has not escaped our notice that the specific pairing we have postulated immediately suggests a possible copying mechanism for the genetic material. Full details of the structure, including the conditions assumed in building it, together with a set of co-ordinates for the atoms, will be published elsewhere.

We are much indebted to Dr. Jerry Donohue for constant advice and criticism, especially on interatomic distances. We have also been stimulated by a knowledge of the general nature of the unpublished experimental results and ideas of Dr. M. H. F. Wilkins, Dr. R. E. Franklin and their co-workers at

Nobel Prize in Physiology or Medicine (1962) - Crick, Watson & Wilkins

"for their discoveries concerning the molecular structure of nucleic acids and its significance for information transfer in living material"

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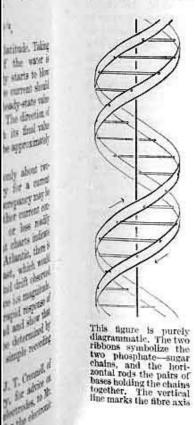
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Structure of the course

The course consists of **5 thematic units**:

1 Philosophical issues in evolutionary biology

Weeks 2-5 (evolution, natural selection, adaptation, adaptationism, niche construction, teleology, units of selection)

2 The ontology of evolution

Weeks 6-7 (organisms, biological individuals, biological species, higher taxa)

3 Philosophical issues in genetic and molecular biology

Weeks 8-10 (reductionism, gene, genetic causation, genetic information)

4 Biology as a science

Week 11 (biological laws and mechanisms, complexity, evolutionary progress, life)

5 Evolution and human nature

Weeks 12-13 (sociobiology, evolutionary psychology, cultural evolution, human nature, evolutionary ethics)

	Торіс	PB	DG
1	Introduction	PB ch. 1	
2	Evolution and natural selection	PB 28-42	
3	Adaptationism and niche construction	PB 50-59	
4	Teleological concepts in biology	PB 59-65	
5	The units of selection and the gene's-eye view of evolution	PB 42-49 93-99	DG part 2
6	Organisms and biological individuals	PB 66-80	
7	Biological species and higher taxa	PB 100-119	
8	The concept of the gene	PB 81-93	
9	Reductionism in biology		
10	Genetic information	PB 144-157	
11	Biological laws and mechanisms	PB 11-27	DG part 3
12	Evolutionary explanations of social behaviour	PB 120-136	
13	Cultural evolution and human nature	PB 45-49 PB 136-143	

- Richard Dawkins , 1nine76 The selfish gene
- Quine Epistemology naturalised
- taxonomy classification
- rank
- naturalists
- Buffon
- Lamarck
- saltationists
- finches / mockingbirds / turtle
- evolutionary lineage
- LUCA
- life experiment genetic code

- patterns process in evolution
- vestiges vestigial traits
- trait = feature / characteristic