

Savčí buněčná biologie v číslech

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Vychází z knihy:

- ▶ Ron Milo & Rob Phillips: Cell Biology by the numbers, Garland Sciences, 2016

Matematika a chemie pro biology

Big numbers at your disposal

- Seconds in a year $\approx \pi \times 10^7$ (the approximate value of pi, a nice coincidence and an easy way to remember this value)
- Seconds in a day $\approx 10^5$
- Hours in a year $\approx 10^4$
- Avogadro's constant $\approx 6 \times 10^{23}$
- Cells in the human body $\approx 4 \times 10^{13}$

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Rules of thumb

Just as there are certain arithmetical rules that help us quickly get to our order-of-magnitude estimates, there are also physical rules of thumb that can similarly extend our powers of estimation. We give here some of our favorites and you are most welcome to add your own at the bottom of our list and also send them to us. Several of these estimates are represented pictorially as well. Note that here and throughout the book we try to follow the correct notation where “approximately” is indicated by the symbol \approx , and loosely means accurate to within a factor of two or so. The symbol \sim means “order of magnitude,” so only to within a factor of 10 (or in a different context it means “proportional”). We usually write approximately because we know the property value indeed roughly but to better than a factor of 10, so \approx is the correct notation and not \sim . In the cases where we only know the order of magnitude, we will write the value only as 10^x without extraneous significant digits.

- 1 dalton (Da) = 1 g/mol $\approx 1.6 \times 10^{-24}$ g (as derived in **Estimate 0-1**).
- 1 nM is about 1 molecule per bacterial volume, as derived in **Estimate 0-2**, 10^1 – 10^2 per yeast cell, and 10^3 – 10^4 molecules per characteristic mammalian (HeLa) cell volume. For 1 μ M, multiply by a thousand; for 1 mM, multiply by a million.
- 1 M is about one per 1 nm³.
- There are 2–4 million proteins per 1 μ m³ of cell volume.

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- Concentration of 1 ppm (part per million) of the cell proteome is ≈ 5 nM.
- 1 mg of DNA fragments 1 kb long is ≈ 1 pmol or $\approx 10^{12}$ molecules.
- Under standard conditions, particles at a concentration of 1 M are ≈ 1 nm apart.
- Mass of typical amino acid ≈ 100 Da.
- Protein mass [Da] $\approx 100 \times$ Number of amino acids.
- Density of air ≈ 1 kg/m³.
- Water density ≈ 55 M $\approx \times 1000$ that of air ≈ 1000 kg/m³.
- 50 mM osmolites ≈ 1 atm osmotic pressure (as shown in **Estimate 0-3**).
- Water molecule volume ≈ 0.03 nm³, (≈ 0.3 nm)³.
- A base pair has a volume of ≈ 1 nm³.
- A base pair has a mass of ≈ 600 Da.
- Lipid molecules have a mass of ≈ 500 – 1000 Da.
- $1 k_B T \approx 2.5$ kJ/mol ≈ 0.6 kcal/mol ≈ 25 meV ≈ 4 pN nm $\approx 4 \times 10^{-21}$ J.
- ≈ 6 kJ/mol sustains one order of magnitude concentration difference [$= k_B T \ln(10) \approx 1.4$ kcal/mol].

Velikosti a počty

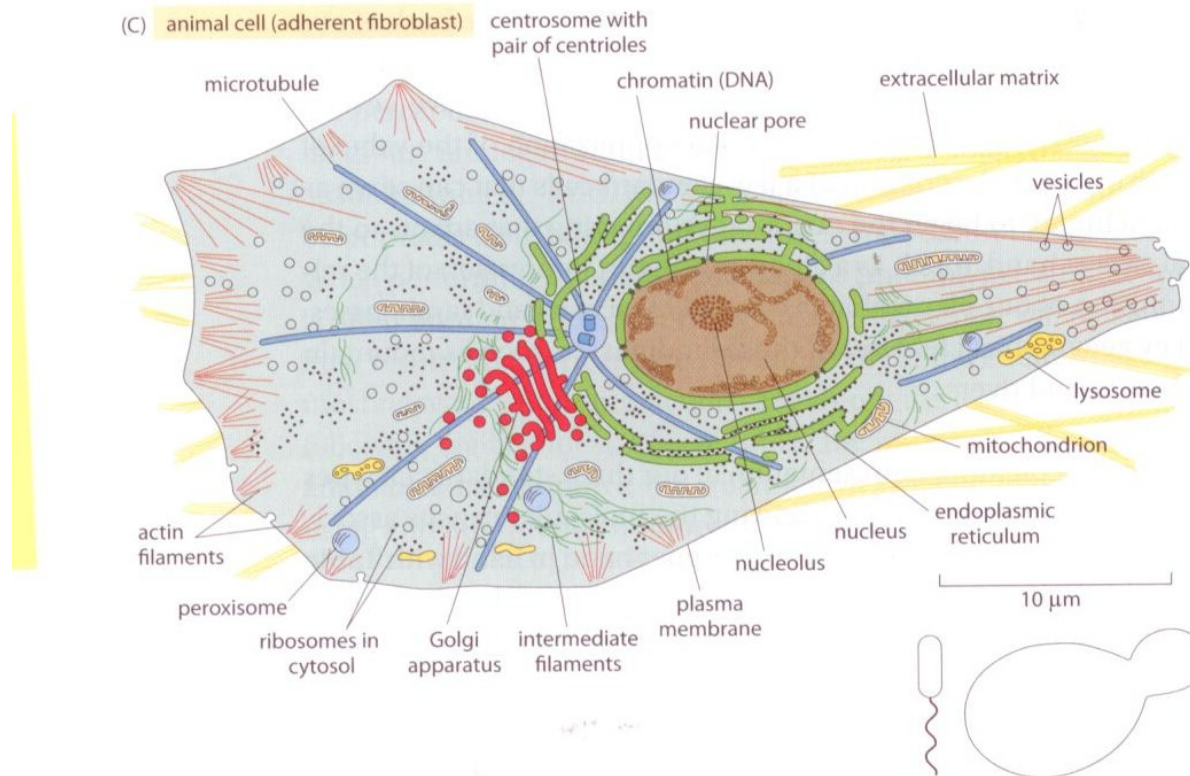


Figure 0-1 The standard cells. (A) A schematic bacterium revealing the characteristic size and components of *E. coli*. (B) A budding yeast cell showing its characteristic size, its organelles, and various classes of molecules present within it. (C) An adherent human cell. We note that these are very simplified schematics. For example, only a small fraction of ribosomes are drawn. Each cell is drawn to a different scale, as indicated by the distinct scale bars in each schematic. The relative sizes of the bacterial and yeast cells at the same scale as the mammalian cell are shown in the bottom right. (A, and C, adapted from Alberts B, Johnson A, Lewis J et al. [2015] *Molecular Biology of the Cell*, 6th ed. Garland Science.)

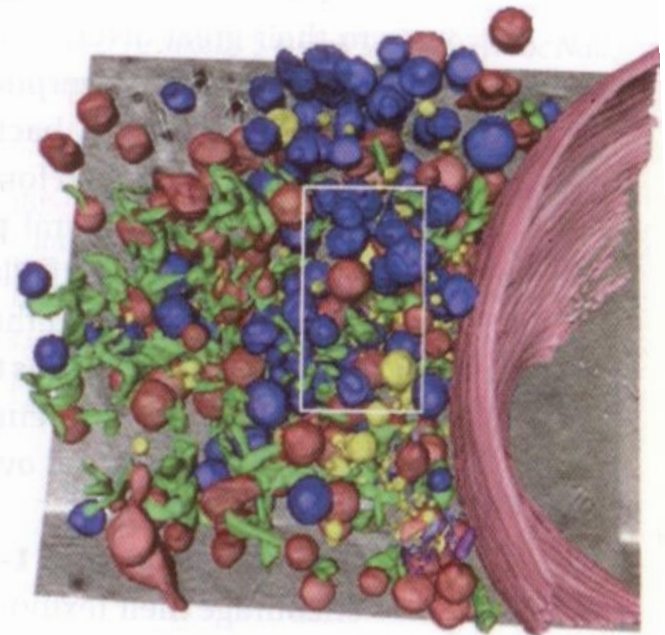
Složení buňky

intracellular compartment	percentage of total cell volume
cytosol	50–60
mitochondria	20
rough ER cisternae	10
smooth ER cisternae plus Golgi cisternae	6
nucleus	6
peroxisomes	1
lysosomes	1
endosomes	1

Table 1-5 The volume fraction occupied by different intracellular compartments in a liver hepatocyte cell. (Adapted from Alberts B, Johnson A, Lewis J et al. [2015] *Molecular Biology of the Cell*, 6th ed. Garland Science.)

a stack of closely spaced confocal images, it is possible to gain insights into the three-dimensional structure of the organelle over time. In these images, we see that during interphase the ER is reticular (netlike). To appreciate the tangled arrangement of organellar membranes even more deeply, **Figure 1-24** provides a reconstructed image using X-ray microscopy of the ER and other ubiquitous membrane systems in the cell. In this cell type and under

Figure 1-24 X-ray microscopy image of cellular ultrastructure highlighting the endoplasmic reticulum. This image is a volumetric rendering of images of a mouse adenocarcinoma cell. The numbers represent percent of the volume occupied by the different compartments. (Adapted from Schneider G, Guttman P, Heim S et al. [2010] *Nature Meth* 7:985–987.)



■ lysosomes	13%
■ mitochondria	17%
■ endoplasmic reticulum	3%
■ vesicles	2%
external	65%

Počty nejdůležitějších biomolekul

(C) mammalian cell (specifically, HeLa: $V \approx 3000 \mu\text{m}^3$; $L \approx 20 \mu\text{m}$; $\tau \approx 1$ day)

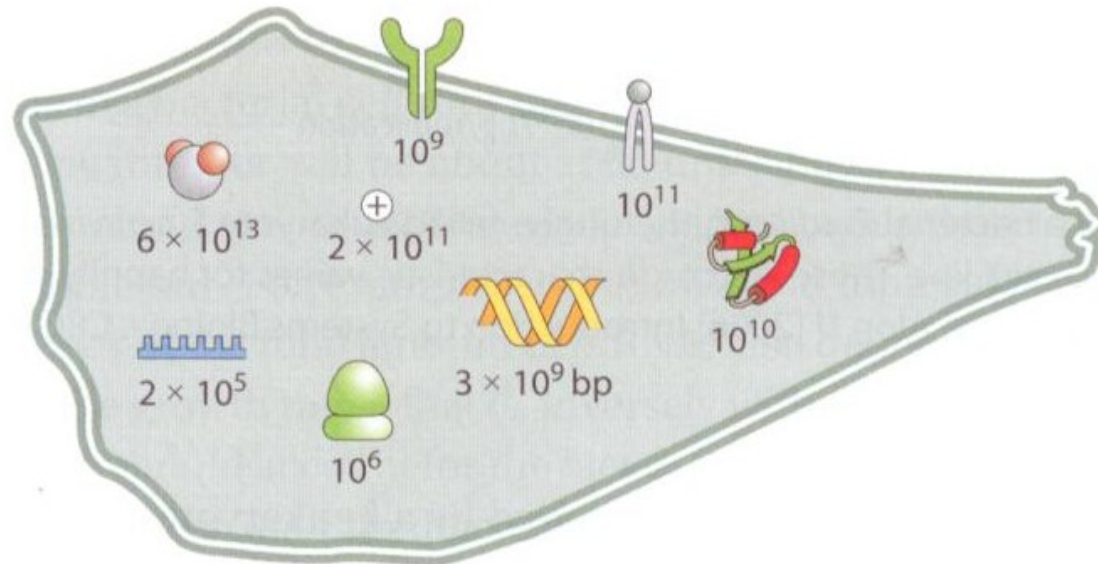


Figure 0-2 An order-of-magnitude census of the major components of the three model cells we employ often in the lab and in this book. A bacterial cell (*E. coli*), a unicellular eukaryote (the budding yeast *S. cerevisiae*), and a mammalian cell line (such as an adherent HeLa cell).

Signální proteiny jsou relativně vzácné

organism	transcription factor	copies per cell order of magnitude	BNID
<i>E. coli</i>	LacI (carbon utilization)	10^1 – 10^2	100734
<i>E. coli</i>	AraC (carbon utilization)	10^2	105139
<i>E. coli</i>	ArcA (general aerobic respiration control)	10^4	102632
<i>S. cerevisiae</i>	Gal4 (carbon utilization)	10^2	109208
<i>S. cerevisiae</i>	Tfb3 (general transcription initiation factor)	10^3	109208
<i>S. cerevisiae</i>	Pho2 (phosphate metabolism)	10^4	109208
<i>D. melanogaster</i> , anterior blastoderm nuclei	bicoid (development)	10^4	106843
<i>D. melanogaster</i> , S2 cells	GAGA zinc finger	10^6	106846
mouse/rat macrophage	glucocorticoid, thyroid and androgen receptors associated zinc fingers	10^4	106899
mouse/rat macrophage	NF-kappaB p65	10^5	106901
<i>H. sapiens</i> cell lines	P53 (growth and apoptosis)	10^4 – 10^5	100420
<i>H. sapiens</i> cell lines	glucocorticoid, estrogen, steroid receptors associated zinc fingers	10^4 – 10^5	106904, 106906, 106911
<i>H. sapiens</i> cell lines	STAT6	10^4 – 10^5	106914
<i>H. sapiens</i> cell lines	NF-kappaB p65	10^5	106909
<i>H. sapiens</i> cell lines	Myc (global chromatin structure regulation)	10^5	106907

Table 2-8 Absolute copy numbers from a number of different organisms. Values are rounded to closest order or magnitude. For more values, see Biggin MD [2011] *Dev Cell* 21:611–626; BNID 106842).

Počty buněk v těle

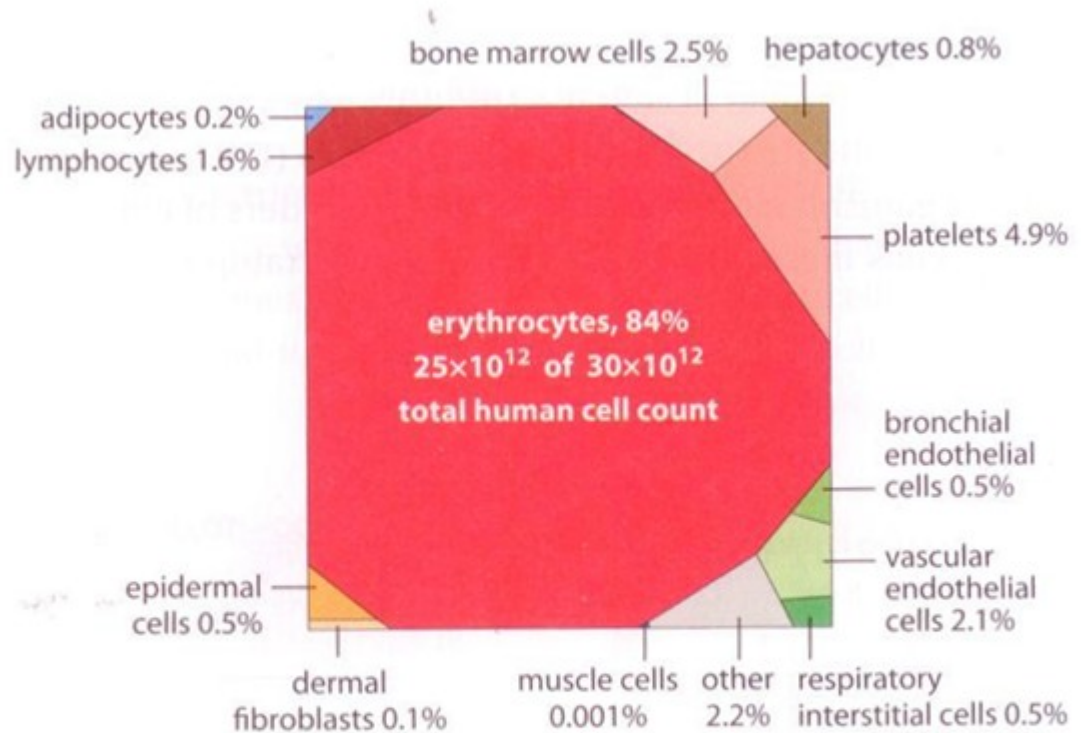


Figure 6-1 Estimate of the number of cells in an adult human partitioned according to cell type. Each cell type in the human body is represented as a polygon with an area proportional to the number of cells. The dominant component is red blood cells. (Based on data from Sender R et al. [2015] in preparation.)

Velikosti

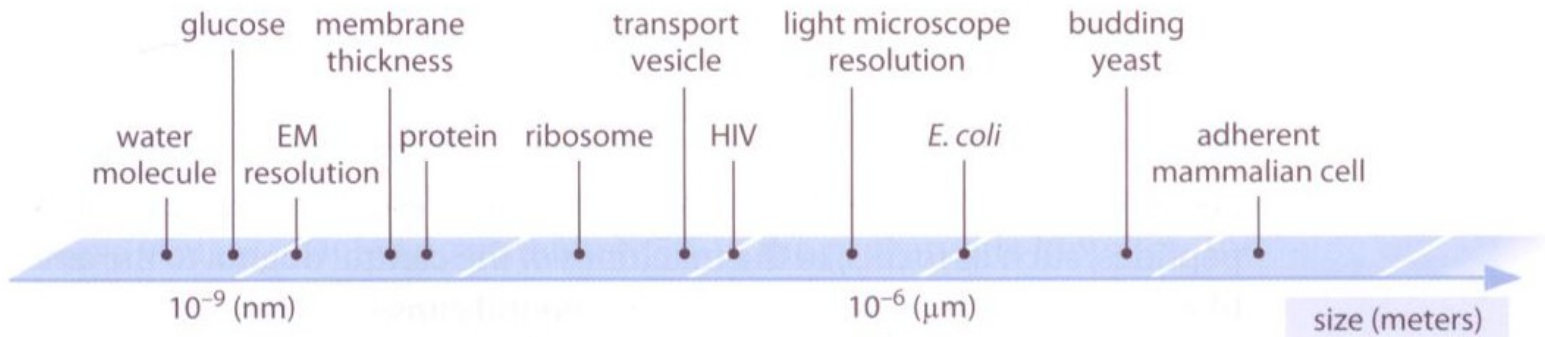


Figure 1-1 Range of characteristic sizes of the main biological entities relevant to cells. On a logarithmic scale we depict the range from single molecules, serving as the nuts and bolts of biochemistry, through molecular machines, to the ensembles that are cells.

Velikosti buněk

cell type	average volume (μm^3)	BNID
sperm cell	30	109891, 109892
red blood cell	100	107600
lymphocyte	200	111788
neutrophil	300	108241
beta cell	1000	109227
enterocyte	1400	111216
fibroblast	2000	108244
HeLa, cervix	3000	103725, 105879
hair cell (ear)	4000	108242
osteoblast	4000	108088
alveolar macrophage	5000	103566
cardiomyocyte	15,000	108243
megakaryocyte	30,000	110129
fat cell	600,000	107668
oocyte	4,000,000	101664

Table 1-3 Characteristic average volumes of human cells of different types. Large cell-cell variation of up to an order of magnitude or more can exist for some cell types, such as neurons or fat cells, whereas for others the volume varies by much less (for example, red blood cells). The value for a beta cell comes from a rat, but we still present it because average cell sizes usually change relatively little among mammals.

Velikosti a tvary proteinů

? Nejpočetnější živočišný protein na Zemi?

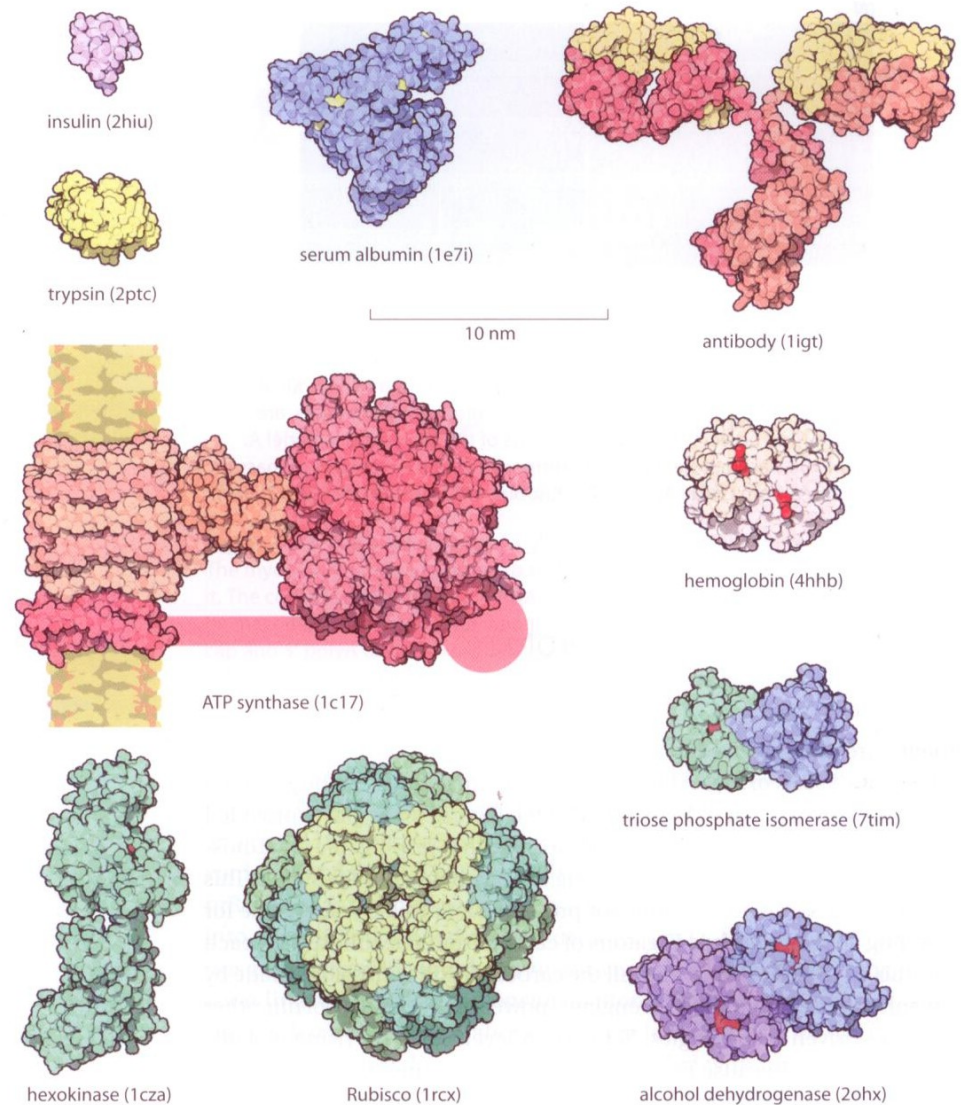


Figure 1-35 Gallery of proteins. Representative examples of protein size are shown with examples drawn from across biology to illustrate some of their key functional roles. Examples range from the antibodies so important to the immune system to Rubisco and photosynthesis. All the proteins in the figure are shown on the same scale to give an impression of their relative sizes. The small red objects shown on some of the molecules are the substrates for the protein of interest. For example, in hexokinase, the substrate is glucose. The handle in ATP synthase is known to exist, but the exact structure was unavailable and thus only schematically drawn. Names in parentheses are the PDB database identifiers. (Courtesy of David Goodsell.)

Velikosti složek cytoskeletu

	actin	microtubules
functions	cell motility, cytokinesis, muscle contraction, hearing	cell division, intracellular transport
subunit	actin monomer	α -tubulin + β -tubulin
subunit weight	≈ 40 kDa	≈ 50 kDa
subunit size	5 nm	4 nm (dimer)
protofilaments number	2	13 (variable)
cross section area	20 nm^2	200 nm^2
filament diameter	6 nm	25 nm
helical repeat period	36 nm	8 nm
persistence length	10 μm	1–10 mm
filament length	from 35 nm in erythrocyte cortex to 10–100 μm in ear hair cells	from $<1 \mu\text{m}$ in <i>S. pombe</i> through 100 μm in rat neurons to >1 mm in insect sperm

Table 1-7 Properties of the main cytoskeleton components: actin and microtubules (BNID 107897).

Velikosti a tvary lipidů

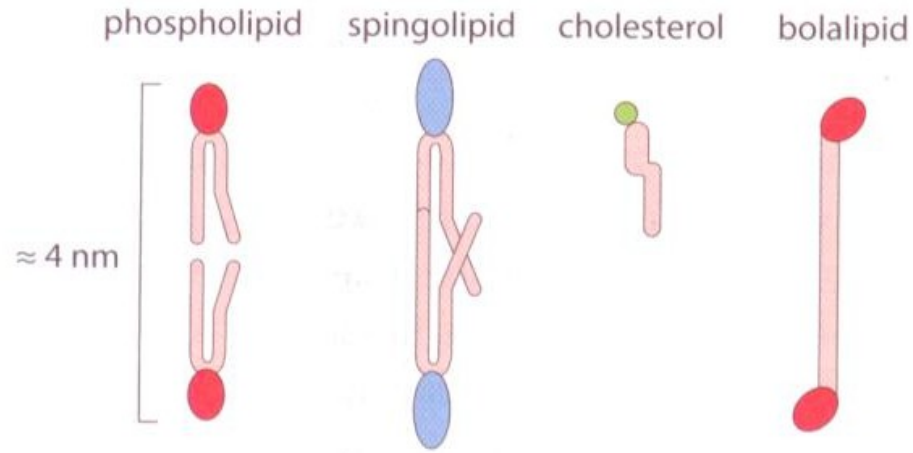


Figure 1-42 Characteristic relative sizes and shapes of the lipid molecules making up biological membranes.

Velikost RNA a proteinu

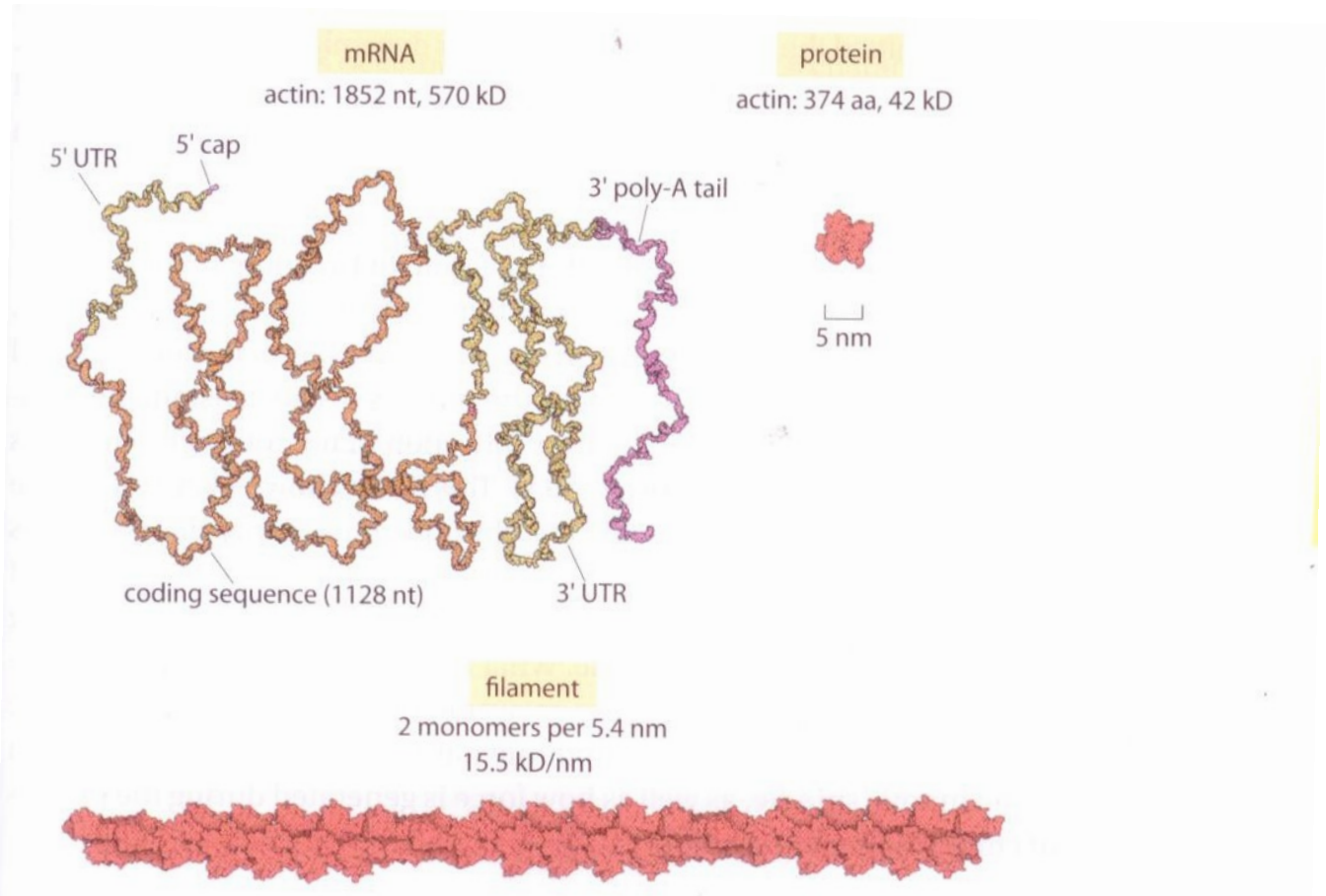


Figure 1-45 Sizes of actin mRNA, protein, and filament. The mRNA molecule is shown next to the corresponding protein monomer that it codes for (based on human actin A). The monomers assemble into actin filaments such as the one indicated schematically at the bottom. For reference, this filament fragment is only 1% of the measured persistence length of these structures. (Courtesy of David Goodsell.)

DNA

- Lidská DNA – 3 mld párů bazí (haploidní) – 1 m
- Buněk – 3×10^{13}
- Buněk s DNA: 5-10 %
- DNA v těle: $2 \times 1.5 \times 10^{12}$ m
- Vzdálenost Země-Slunce: 150 milionů kilometrů = $1,5 \times 10^{11}$

Je lidský genom velký nebo malý?

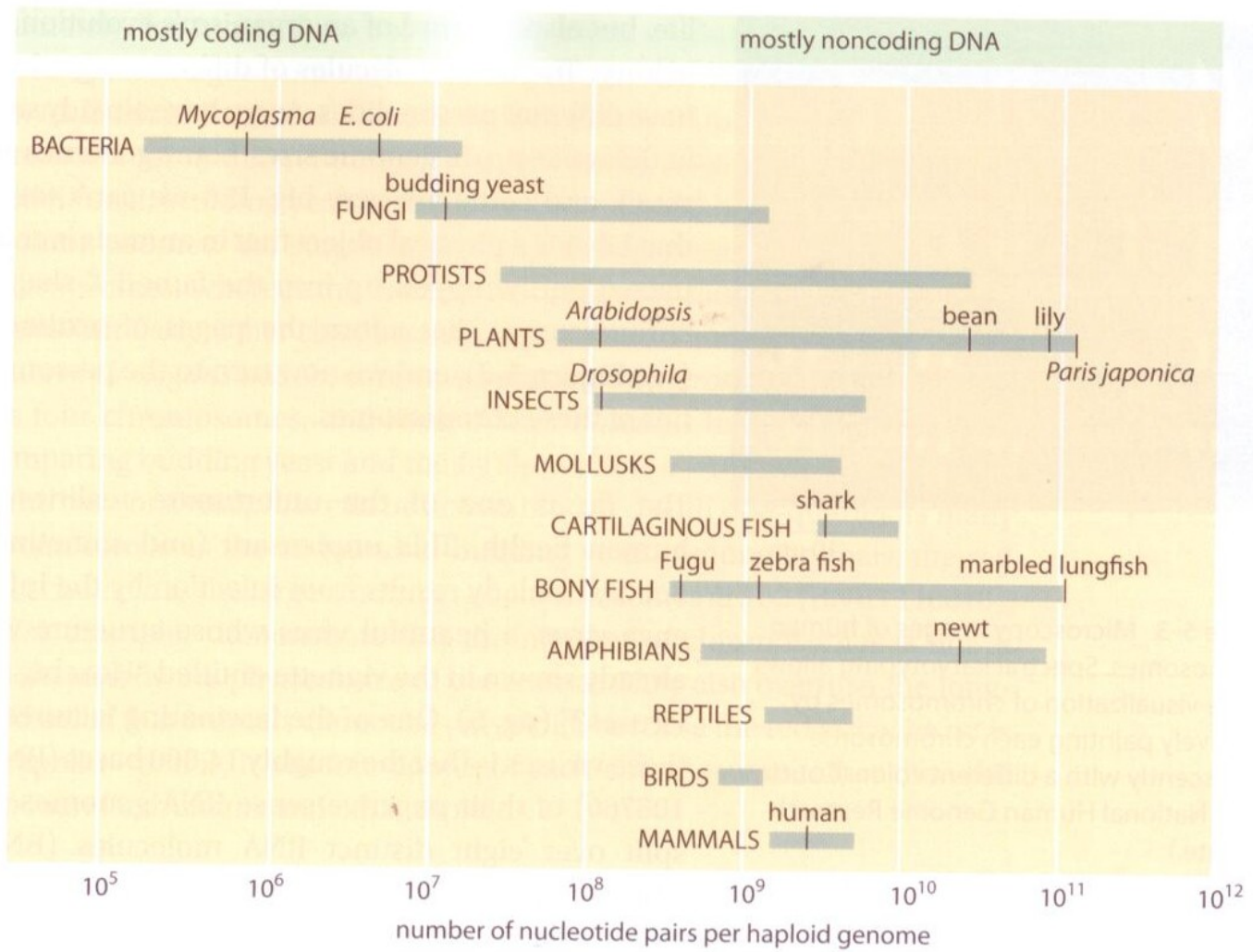


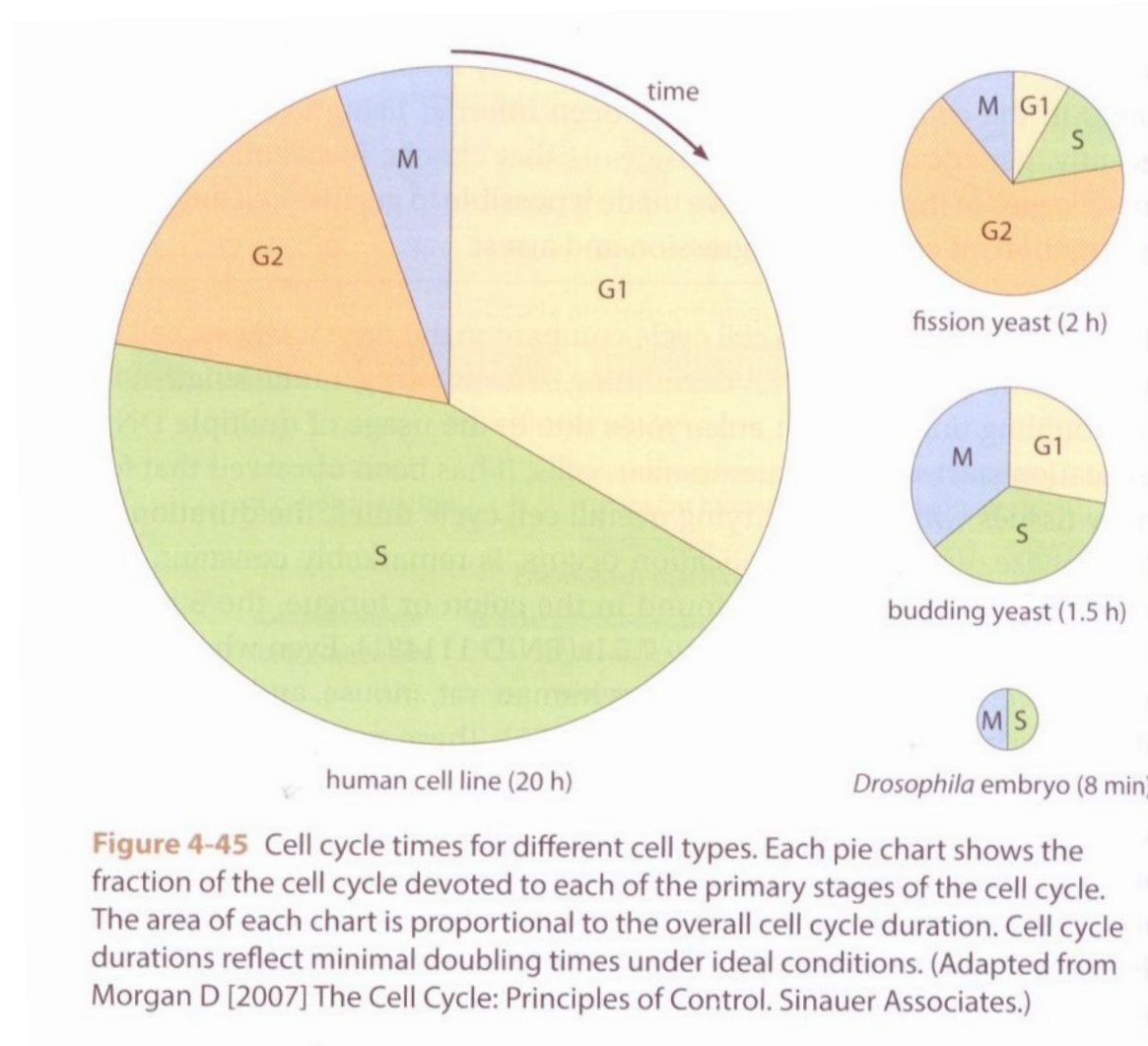
Figure 5-2 Genome sizes of different organisms.

Je lidský genom velký nebo malý?

	organism	# of protein-coding genes	# of genes naïve estimate: (genome size / 1000)	BNID
viruses	HIV 1	9	10	105769
	Influenza A virus	10–11	14	105767
	Bacteriophage λ	66	49	105770
	Epstein-Barr virus	80	170	103246
prokaryotes	<i>Buchnera sp.</i>	610	640	105757
	<i>T. maritima</i>	1900	1900	105766
	<i>S. aureus</i>	2700	2900	105500
	<i>V. cholerae</i>	3900	4000	105760
	<i>B. subtilis</i>	4400	4200	111448
	<i>E. coli</i>	4300	4600	105443
eukaryotes	<i>S. cerevisiae</i>	6600	12,000	105444
	<i>C. elegans</i>	20,000	100,000	101364
	<i>A. thaliana</i>	27,000	140,000	111380
	<i>D. melanogaster</i>	14,000	140,000	111379
	<i>F. rubripes</i>	19,000	400,000	111375
	<i>Z. mays</i>	33,000	2,300,000	110565
	<i>M. musculus</i>	20,000	2,800,000	100308
	<i>H. sapiens</i>	21,000	3,200,000	100399, 111378
<i>T. aestivum</i> (hexaploid)	95,000	16,800,000	105448, 102713	

Table 5-2 A comparison between the number of genes in an organism and a naïve estimate based on the genome size divided by a constant factor of 1000 bp/gene—that is, predicted number of genes = genome size/1000. This crude rule of thumb works surprisingly well for many bacteria and archaea, but fails miserably for multicellular organisms.

Časy – buněčný cyklus



Časy – délka „života“ buněk

cell type	turnover time	BNID
small intestine epithelium	2–4 days	107812, 109231
stomach	2–9 days	101940
blood neutrophils	1–5 days	101940
white blood cells eosinophils	2–5 days	109901, 109902
gastrointestinal colon crypt cells	3–4 days	107812
cervix	6 days	110321
lungs alveoli	8 days	101940
tongue taste buds (rat)	10 days	111427
platelets	10 days	111407, 111408
bone osteoclasts	2 weeks	109906
intestine paneth cells	20 days	107812
skin epidermis cells	10–30 days	109214, 109215
pancreas beta cells (rat)	20–50 days	109228
blood B cells	1 month	111516
trachea	1–2 months	101940
hematopoietic stem cells	2 months	109232
sperm (male gametes)	2 months	110319, 110320
bone osteoblasts	3 months	109907
red blood cells	4 months	101706, 107875
liver hepatocyte cells	0.5–1 year	109233
fat cells	8 years	103455
cardiomyocytes	0.5–10% per year	107076, 107077, 107078
central nervous system	life-time	101940
skeleton	10% per year	109908
lens cells	life-time	109840
oocytes (female gametes)	life-time	111451

Časy – délka života proteinů

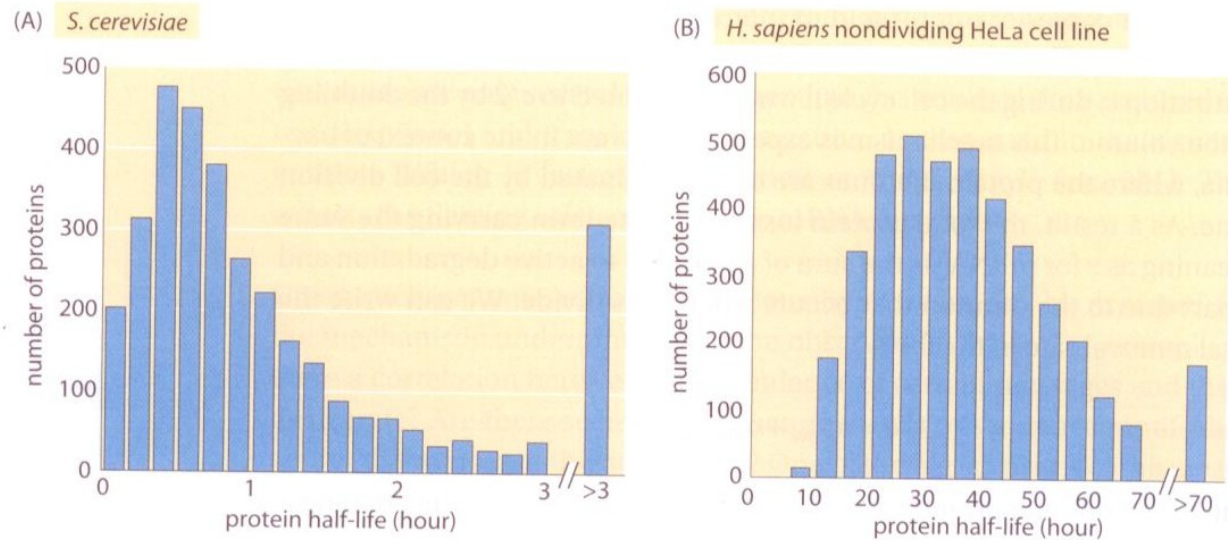
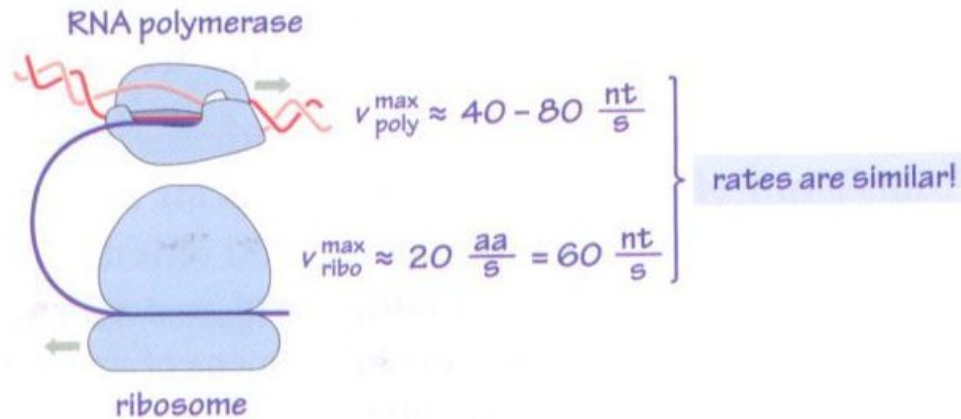


Figure 4-22 Measured half-lives of proteins in budding yeast and a HeLa human cancer cell line. The yeast experiment used the translation inhibitor cycloheximide, which disrupts normal cell physiology. The median half-life of the 4100 proteins measured in the nondividing HeLa cell is 36 h. (A, adapted from Belle A, Tanay A, Bitincka L et al. [2006] *Proc Natl Acad Sci USA* 103:13004–13009. B, adapted from Cambridge S, Gnad F, Nguyen C et al. [2011] *J Proteome Res* 10:5275–5284.)

Rychlosti transkripce a translace

Which is faster: transcription or translation?



Estimate 4-9

” Bakshi S, Siryaporn A, Goulian M et al. (2012) Superresolution imaging of ribosomes and RNA polymerase in live *Escherichia coli* cells. *Mol Microbiol* 85:21–38.

Rychlosti – šíření vzruchu

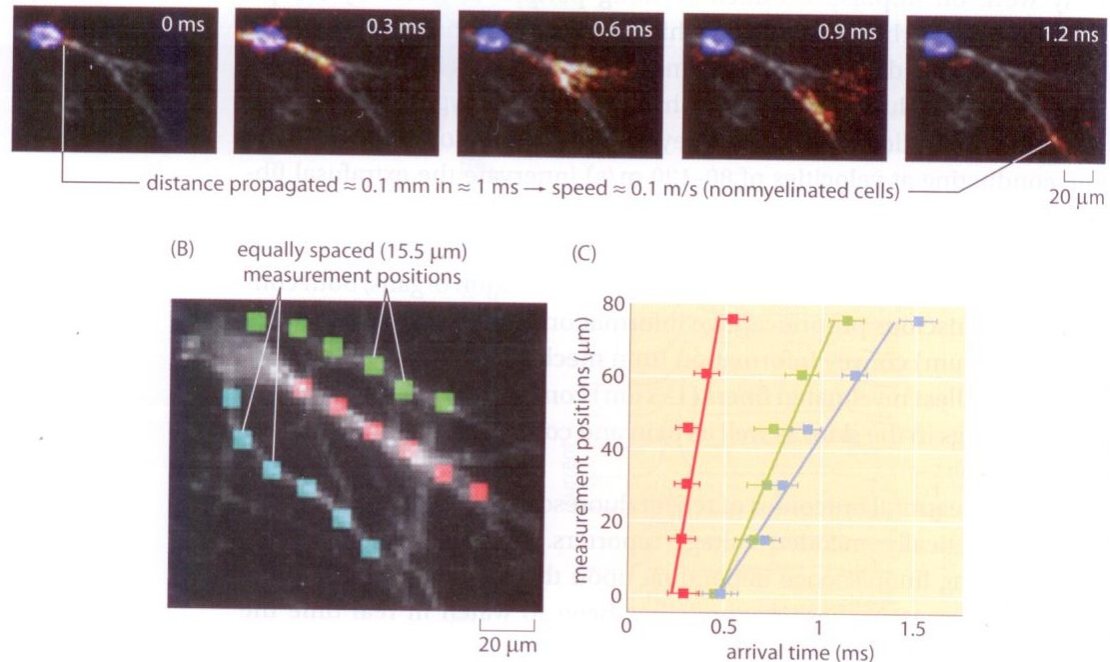


Figure 4-28 Optical measurement of action potential speed. (A) Series of images of the fluorescence in the cell as a function of time. (B) A series of equally spaced ($15.5 \mu\text{m}$) measurement points along three different processes are used to measure the arrival time of a propagating action potential. (C) Arrival times for the three processes shown in (B). For example, for the action potential propagating along the fiber labeled with red boxes, the signal arrives with a time delay of roughly 0.05 ms from one measurement point to the next. The action potential speed can be read off of the graph in the usual way by dividing distance traveled by time elapsed. Due to technical limitations, these are unmyelinated neuronal cells, and thus the propagation speed is much slower than *in vivo*. (Courtesy of Daniel Hochbaum and Adam Cohen.)

Myelinizované vlákno – 100 m/s tj. až $1000\times$ rychlejší

Rychlosti – růstu mikrotubulů

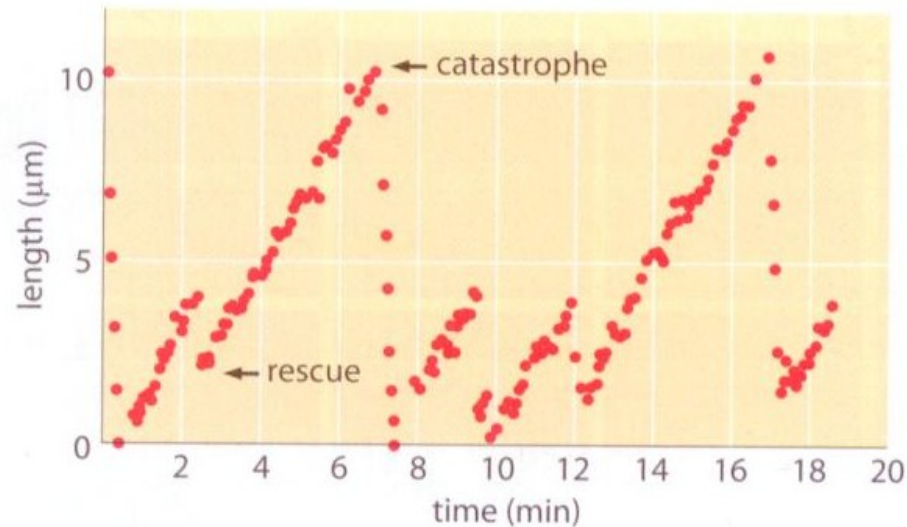


Figure 4-33 Microtubule length vs. time. The length of microtubules as a function of time reveals periods of growth punctuated by catastrophes in which the filaments rapidly depolymerize. (Adapted from Fygenson DK, Braun E & Libchaber A [1994] *Phys Rev E* 50:1579–1588.)

Rychlosti – pohybu motorů po cytoskeletu

motor	function	speed <i>in vivo</i> (nm/s)	rate (ATPase, s ⁻¹ , <i>in vitro</i>)	mode of action
myosins				
myosin XI	cytoplasmic streaming in algae	60,000	not determined	unknown
myosin II	fast skeletal muscle	6000	20	large arrays (10 ⁴ –10 ⁹)
myosin IB	amoeboid motility, hair cell adaptation	200 (<i>in vitro</i>)	6	small arrays (10–10 ³)
myosin II	smooth muscle contraction	200	1.2	large arrays (10 ⁴ –10 ⁹)
myosin V	vesicle transport	200	5	alone or in small numbers (<10)
myosin VI	vesicle transport?	-60 (<i>in vitro</i>)	0.8	unknown
dyeins				
axonemal	sperm and cilia motility	-7000	10	large arrays (10 ⁴ –10 ⁹)
cytoplasmic	retrograde axonal transport, mitosis, transport in flagella	-1000	2	alone or in small numbers (<10)
kinesins				
			1 μm/s	
Fia10/KinII	transport in flagella, axons, melanocytes	2000	not determined	small arrays (10–10 ³)
conventional	anterograde axonal transport	1800	40	alone or in small numbers (<10)
Nkin	secretory vesicle transport	800	80	alone or in small numbers (<10)
Unc104/KIF	transport of synaptic vesicle precursors and mitochondria	700	110	alone or in small numbers (<10)
Bimc/Eg5	mitosis and meiosis	18	2	small arrays (10–10 ³)
Ncd	mitosis and meiosis	-90 (<i>in vitro</i>)	1	small arrays (10–10 ³)

Table 4-7 Summary of experimental data on the dynamics of translational molecular motors. Based on BNID 106501, 101506. Values were rounded to one significant digit. Negative speeds indicate movement towards the minus end of the filament.

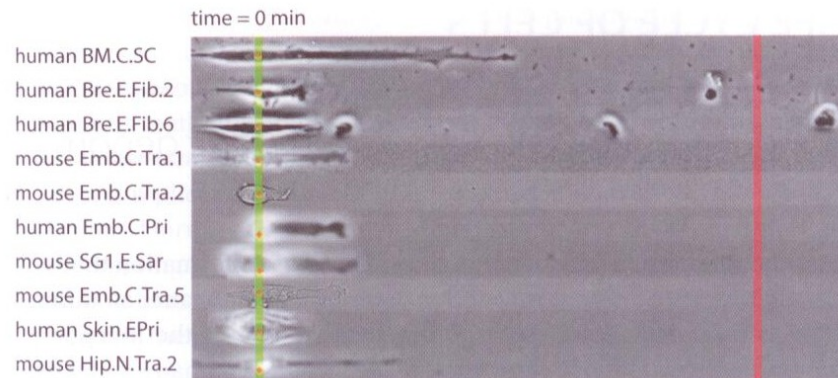
Rychlosti – pohybu buněk

organism	speed	speed in body lengths (bl) per time	BNID and comments
eukaryotes			
<i>Ciliate Paramecium tetraurelia</i>	100–1000 $\mu\text{m/s}$	1–5 bl/s	108087, ciliated, assuming 200 μm cell length
<i>Tetrahymena thermophila</i>	200–400 $\mu\text{m/s}$	4–8 bl/s	111429, 111435, 111436, ciliated
<i>Gyrodinium dorsum</i>	300 $\mu\text{m/s}$	10 bl/s	111432, flagellated
green algae <i>Chlamydomonas Reinhardtii</i>	50–150 $\mu\text{m/s}$	5–15 bl/s	108086, 111430
fish keratocytes - wound healing fibroblasts of the cornea	10–50 $\mu\text{m/min}$	0.7–3 bl/min	106807, 106817
<i>Amoeba Dictyostelium discoideum</i>	10 $\mu\text{m/min}$	≈ 1 bl/min	106825
human neutrophil	9 $\mu\text{m/min}$	≈ 1 bl/min	106809
glioma cells	50 $\mu\text{m/h}$	4 bl/h	106810
mouse fibroblastoid L929 cells	30 $\mu\text{m/h}$	2 bl/h	106808
human H69 small cell lung cancer cell	16 $\mu\text{m/h}$	1 bl/h	106815

Table 4-8 Cell speeds of different cells given in μm per time unit and as body lengths per time unit. Assume a bacterial length of $\approx 2 \mu\text{m}$ and a eukaryotic cell length of $\approx 15 \mu\text{m}$ unless otherwise stated. Speeds depend on temperature, experimental conditions, etc. Values given here are those reported in the literature. Most measurements are based on time-lapse microscopy.

Rychlosti – pohybu buněk

World Cell Race 2011



Rychlosti – pohybu buněk

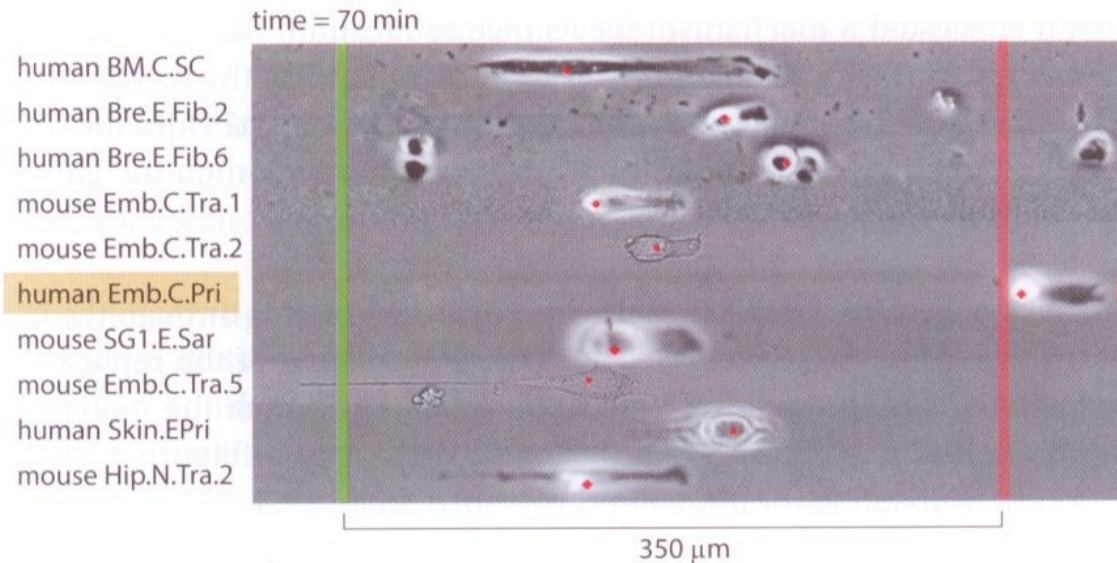
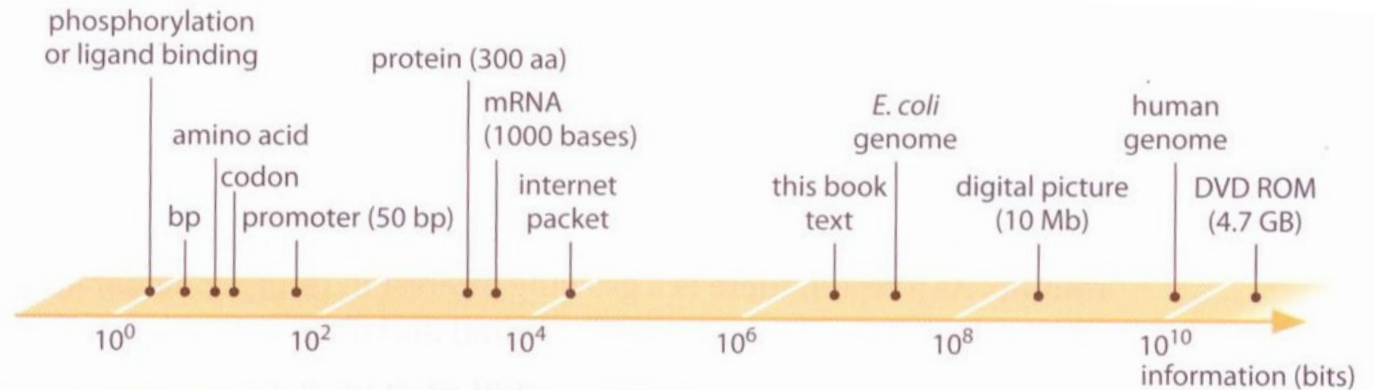


Figure 4-40 Finals of the World Cell Race. The 10 fastest cell lines are displayed competing over a 350 μm micro-fabricated sprinting lane. Each of the cells was found to be the fastest among its cell type. Each cell type was recorded in a separate well, and movies were combined to show one lane per cell type. The time difference between the two images is about an hour. The winner is highlighted in brown. (Courtesy of Matthieu Piel.)

Informace



information, $I = \log_2(\text{\# possible configurations})$

e.g., base pair (bp) has four possibilities $\Rightarrow I = \log_2(4) = 2$

Figure 5-1 Information content of biological entities and some human designed information storage devices. Information is quantified through binary bits, where a base pair that has four possibilities is 2 bits, etc.

Malé shrnutí

property	<i>E. coli</i>	budding yeast	mammalian (HeLa line)
cell volume	0.3–3 μm^3	30–100 μm^3	1000–10,000 μm^3
proteins per μm^3 cell volume	2–4 $\times 10^6$		
mRNA per cell	10^3 – 10^4	10^4 – 10^5	10^5 – 10^6
proteins per cell	$\sim 10^6$	$\sim 10^8$	$\sim 10^{10}$
mean diameter of protein	4–5 nm		
genome size	4.6 Mbp	12 Mbp	3.2 Gbp
number protein coding genes	4300	6600	21,000
regulator binding site length	10–20 bp	5–10 bp	
promoter length	~ 100 bp	~ 1000 bp	$\sim 10^4$ – 10^5 bp
gene length	~ 1000 bp	~ 1000 bp	$\sim 10^4$ – 10^6 bp (with introns)
concentration of one protein per cell	~ 1 nM	~ 10 pM	~ 0.1 – 1 pM
diffusion time of protein across cell ($D \approx 10 \mu\text{m}^2/\text{s}$)	~ 0.01 s	~ 0.2 s	~ 1 – 10 s
diffusion time of small molecule across cell ($D \approx 100 \mu\text{m}^2/\text{s}$)	~ 0.001 s	~ 0.03 s	~ 0.1 – 1 s
time to transcribe a gene	<1 min (80 nts/s)	~ 1 min	~ 30 min (incl. mRNA processing)
time to translate a protein	<1 min (20 aa/s)	~ 1 min	~ 30 min (incl. mRNA export)
typical mRNA lifetime	3 min	30 min	10 h
typical protein lifetime	1 h	0.3–3 h	10–100 h
minimal doubling time	20 min	1 h	20 h
ribosomes/cell	$\sim 10^4$	$\sim 10^5$	$\sim 10^6$
transitions between protein states (active/inactive)	1–100 μs		
time scale for equilibrium binding of small molecule to protein (diffusion limited)	1–1000 ms (1 μM –1 nM affinity)		
time scale of transcription factor binding to DNA site	~ 1 s		
mutation rate	10^{-8} – 10^{-10} /bp/replication		

Table 0-1 Typical parameter values for a bacterial *E. coli* cell, the single-celled eukaryote *S. cerevisiae* (budding yeast), and a mammalian HeLa cell line. These are crude characteristic values for happily dividing cells of the common lab strains. (Adapted from Alon U [2006] Introduction to Systems Biology. CRC Press. See full references at BNID 111494.)

Velikosti genomu

organism	genome size (base pairs)	protein-coding genes	number of chromosomes
model organisms			
eukaryotes – multicellular			
pufferfish <i>Fugu rubripes</i> (smallest known vertebrate genome)	400 Mbp	19,000	22
poplar <i>P. trichocarpa</i> (first tree genome sequenced)	500 Mbp	46,000	19
corn <i>Z. mays</i>	2.3 Gbp	33,000	20 (2n)
dog <i>C. familiaris</i>	2.4 Gbp	19,000	40
chimpanzee <i>P. troglodytes</i>	3.3 Gbp	19,000	48 (2n)
wheat <i>T. aestivum</i> (hexaploid)	16.8 Gbp	95,000	42 (2n = 6x)
marbled lungfish <i>P. aethiopicus</i> (largest known animal genome)	130 Gbp	unknown	34 (2n)
herb plant <i>Paris japonica</i> (largest known genome)	150 Gbp	unknown	40 (2n)

Table 5-1 Genomic census for a variety of selected organisms. The table features the genome size, current best estimate for number of protein-coding genes, and number of chromosomes. Genomes often also include extra chromosomal elements such as plasmids, which might not be indicated in the genome size and number of chromosomes. The number of genes is constantly under revision. The numbers given here reflect the number of protein-coding genes. tRNA and noncoding RNAs are not accounted for since many of them are still to be discovered. Bacterial strains often show significant variations in genome size and number of genes among strains. Values were rounded to two significant digits. See full references in BNID 111493.

Rychlosti – mutací a změn v DNA

organism	mutations/ base pair/ replication	mutations/ base pair/ generation	mutations/ genome/ replication	BNID
multicellular				
human <i>H. sapiens</i>	10^{-10}	$1-4 \times 10^{-8}$ (mitochondria: 3×10^{-5})	0.2–1	105813, 100417, 105095, 108040, 109959, 105813, 110292, 111227, 111228
mouse <i>M. musculus</i>	2×10^{-10}	10^{-8}	0.5	100315, 106792, 100320
<i>D. melanogaster</i>	3×10^{-10}	10^{-8}	0.06	100365, 106793, 100370
<i>C. elegans</i>	10^{-10} – 10^{-9}	10^{-8}	0.02–0.2	100290, 100287, 109959, 103520, 107886
unicellular				
bread mold <i>N. crassa</i>		10^{-10}	0.003	100355, 100359, 106747
budding yeast		10^{-10} – 10^{-9}	0.003	100458, 100457, 109959, 110018
<i>E. coli</i>		10^{-10} – 10^{-9}	0.0005–0.005	106748, 100269, 100263

Table 5-3 Mutation rates of different organisms from different domains of life. RNA virus mutation rates are especially high, partially due to not having a proofreading mechanism. For multicellular organisms, a distinction is made between mutations per replication versus mutations per generation, which includes many replications from gamete to gamete—see vignette entitled “How quickly do different cells in the body replace themselves?” (pg. 278). To arrive at the mutation rate per genome, the rates per base pair are multiplied by the genome length. Mutation rates in the mitochondrial genome are usually an order of magnitude higher (BNID 109959).