

CELL MECHANICS

LECTURE 2

2. Physical principles

2.1. Forces at molecular and cell level

- Physical forces and their magnitudes at the single-molecule level
- Modeling complex mechanical devices as protein machines by using three elements:
case study: Mass, Stiffness and Damping of Proteins

2.2. Thermal forces, diffusion, and chemical forces

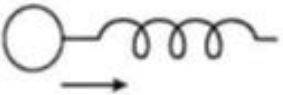
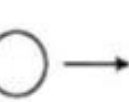
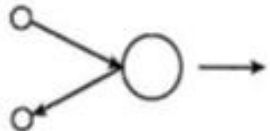

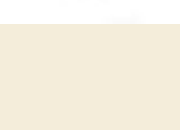
- Boltzmann's law and the Principle of Equipartition of Energy
- Diffusion equation - Einstein relation – Stokes law
- Autocorrelation function and Power Spectrum
- The effect of force on the equilibria and rate of chemical reactions
- Example of single molecule force spectroscopy experiments – unbinding, unfolding

Outline:

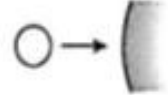

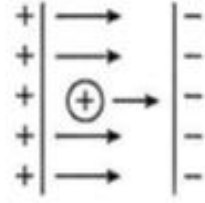
- Physical forces and their magnitudes at the single-molecule level
- Modeling complex mechanical devices as protein machines by using three elements:
Spring, Dashpot, Mass
- Mass, Stiffness and Damping of Proteins

- The **force** drives change and motion. E.g. motor proteins and other molecular machines are able to move and do work because they generate force.
- What types of interactions and forces occur in cells ?
- Where these forces come from ?
- Which is the magnitude of forces acting on molecules ?

Table 2.1 Examples of forces acting on molecules

Type of force	Diagram	Approximate magnitude
Elastic		1-100 pN
Covalent		10,000 pN
Viscous		1-1000 pN
Collisional		10^{-12} to 10^{-9} pN for 1 collision/s
Thermal		100-1000 pN

Range:
pN - nN

Gravity		10^{-9} pN
Centrifugal		$< 10^{-3}$ pN
Electrostatic and van der Waals		1-1000 pN
Magnetic		$\ll 10^{-6}$ pN

Elastic

$F = k \cdot x$, where: k – spring constant (stiffness), x – displacement

Example: motor protein $k = 1 \text{ pN/1nm}$, spring strained through distance $x = 1 \text{ nm} \rightarrow F = 1 \text{ pN}$

Viscous

$F = \gamma \cdot v$, where: γ – drag coefficient, v – relative velocity between object and liquid

$\gamma = 6\pi\eta r$, with η – liquid viscosity, r – radius of a spherical particle

Example: for a globular protein (radius $r = 3 \text{ nm}$, molecular mass $MM = 100 \text{ kDa}$) in water ($\eta \sim 1 \text{ mPa s}$) $\rightarrow \gamma \sim 60 \text{ pN s/m}$

the average thermal speed

$$v_{rms} = \sqrt{\frac{3KT}{m}}$$

$1\text{Da} \sim 1.66 \times 10^{-27} \text{ kg}$

$KT \sim 4.1 \text{ pN nm}$

$\rightarrow v_{rms} \sim 8 \text{ m/s}$

$F \sim 480 \text{ pN}$

Collisional and thermal

Example: Protein – water molecule collision / s: $F = \Delta p / \Delta t$

Water molecule: mass $m \sim 18 \text{ Da}$, average thermal speed $v_{rms} \sim 600 \text{ m/s}$, momentum: $p \sim 18 \times 10^{-24} \text{ kg m/s}$.

Assuming the interaction is perfectly elastic: $F = \Delta p / \Delta t = \Delta(mv) / \Delta t = 2 p / \Delta t \sim 36 \times 10^{-12} \text{ pN}$ - very small

However, the number of collisions / s is much much bigger ($> 10^{13}$), such that the instantaneous thermal force acting on a the protein is on the order of the viscous force: **$F \sim 500 \text{ pN}$** , and drives diffusion.

Optical forces

Optical pressure due to the momentum of light (photon's linear momentum : $p=hv/c$)

Example: if an object absorbs one green photon / second, the corresponding force is:

$$F= \Delta p/ \Delta t \sim \mathbf{1.3 \cdot 10^{-27} \text{ N}}$$
 - very small

$$h= 6.63 \cdot 10^{-34} \text{ m}^2 \text{ kg} / \text{ s} , \nu = 6 \cdot 10^{14} \text{ Hz} , c= 3 \cdot 10^8 \text{ m/s} , E= h \nu = 4 \cdot 10^{-19} \text{ J}$$

A laser beam of power $P=1 \text{ mW}$ has about $N \sim 2.5 \cdot 10^{15}$ photons ! $\rightarrow F (1\text{mW}) \sim \mathbf{3.25 \text{ pN}}$

still small but enough to make an effect on small objects (see optical tweezers)

Gravity

Example: protein 100 kDa = $166 \cdot 10^{-21} \text{ g}$, the gravitational force $F= 1.7 \cdot 10^{-9} \text{ pN}$ very small

$$F= mg, m - \text{mass}, g - \text{gravitational acceleration}$$

Centrifugal

Ultracentrifuges \rightarrow acceleration $a_c \sim 10^5 \text{ g}$, associated force on protein 100 kDa is still modest: $F= 1.7 \cdot 10^{-3} \text{ pN}$, but this is large enough to cause the protein to drift at an average speed of $\sim 3 \mu\text{m/s}$ \rightarrow protein sedimentation through a distance of 100 mm (typical length of centrifuge tube) in about 10 h.

Electrostatic

$$F = qE$$

Example: force experienced by a potassium ion K^+ , traveling through an ion channel of the plasma membrane.

The charge of the ion $q = 1.60 \cdot 10^{-19}$ C; the electric field across a typical plasma membrane: $E = 15 \cdot 10^6$ V/m

(60 mV potential across the 4 nm thick membrane) -- **> $F = 2.4$ pN**

Similar force exists between two monovalent ions in water that are separated by 1 nm (*homework*).

Van der Waals forces are also electrostatic – they arise from the charge separation induced by nearby atoms.

These forces can be as high as 100 pN / nm² of protein-protein interface

Magnetic

Very small at the molecular level because molecules interact very weakly with magnetic fields.

Example: max force on a proton in the strongest nuclear magnetic resonance (NMR) machines is only of the order of 10^{-12} pN.

Thus even with a huge protein with 3000 amino acids and 60000 atoms subject to a very strong magnetic field the magnetic force is **< 10^{-6} pN**.

Example 2.1 Physical forces and their magnitudes at the single-molecule level

ELASTIC FORCES. If an object is connected to a spring of stiffness κ that is stretched a distance x beyond its resting length, then the object will experience a force of $F = \kappa x$. For a motor protein, the stiffness might be about $1 \text{ mN/m} = 1 \text{ pN/nm}$. If the spring is strained through a distance of $1 \text{ nm} = 10^{-9} \text{ m}$, a distance appropriate to the size of proteins, then the force exerted on the object is 1 pN .

VISCOUS FORCES. If an object is held fixed in a moving liquid or is moving through a stationary fluid, then it will experience a viscous, or drag, force from the liquid. The force is proportional to the relative velocity, v , between the object and the fluid according to $F = \gamma v$. The constant of proportionality, γ , is called the drag coefficient. The drag coefficient is related to the size and the shape of the object as well as the viscosity. For example, for a sphere of radius r moving through a liquid of viscosity η , the drag coefficient is $6\pi\eta r$ (Stokes' law, Chapter 3). The viscous forces on proteins are large. For a globular protein of diameter 6 nm , corresponding to a molecular mass of $\sim 100 \text{ kDa}$ (see Table 2.2), the drag coefficient measured by centrifugation studies at 20°C is $\sim 60 \text{ pN}\cdot\text{s/m}$ (Creighton, 1993), in good agreement with Stokes' law. The average instantaneous thermal speed of such a protein in solution at standard temperatures is $\sim 8 \text{ m/s}$ (this is a consequence of thermally driven collisions from the surrounding solvent molecules, Chapter 4). The corresponding viscous force is therefore $\sim 480 \text{ pN}$.

COLLISIONAL AND THERMAL FORCES. If an object is struck by another, it experiences a force equal to the rate of change in momentum (mv) of the striking particle, $F = d(mv)/dt$. For example, the mass of a water molecule is $\sim 30 \times 10^{-27} \text{ kg}$, the average speed associated with its kinetic energy is $\sim 600 \text{ m/s}$ (Chapter 4), and therefore its momentum is $\sim 18 \times 10^{-24} \text{ kg}\cdot\text{m/s}$. If a protein were struck head-on every second by a water molecule that bounced straight back, then the average force would be equal to $36 \times 10^{-12} \text{ pN}$ (twice the momentum for an elastic collision). This is a very small force. However, in solution a huge number of collisions take place per second. The collisions come from all directions, and the resulting randomly directed force, called the thermal force, drives diffusion. The average instantaneous thermal force acting on a 100 kDa protein is on the order of the viscous force, or $\sim 500 \text{ pN}$ (Chapter 4).

OPTICAL FORCES. Another example of a collisional force is optical pressure. Because photons have momentum, they exert a force when they are diffracted by an object. The momentum of a photon is $h\nu/c = h/n\lambda$, where h is Planck's constant, ν is the frequency of the light, c is the speed of light, n is the refractive index, and λ is the wavelength (in a vacuum). If an

object in water ($n = 1.33$) absorbs one green photon ($\lambda = 500 \text{ nm}$) per second, the corresponding optical force on it is $1.0 \times 10^{-15} \text{ pN}$ (the values for the physical constants can be found in the table on the endpapers). This is a very small force. Even if a molecule adsorbs 10^9 photons per second, which would require very bright laser illumination, the optical force would still be only 10^{-6} pN .

GRAVITY. An object of mass m experiences a gravitational force of magnitude mg , where g is the acceleration due to gravity, equal to $\sim 9.8 \text{ m/s}^2$ at the Earth's surface. With a mass of only $166 \times 10^{-24} \text{ kg}$, a 100 kDa protein experiences a gravitational force of only $1.6 \times 10^{-9} \text{ pN}$. At the single-molecule level, gravitational forces are very small and can be ignored.

CENTRIFUGAL FORCES. An object spinning in a centrifuge experiences a centrifugal force equal to ma_c . Ultracentrifuges are capable of generating centrifugal accelerations, a_c , in excess of 100,000 times that of gravity. The associated centrifugal forces on molecules are still quite modest, $\sim 160 \times 10^{-18} \text{ N} = \sim 160 \times 10^{-6} \text{ pN}$ for our 100 kDa protein, but this is large enough to cause the protein to drift at an average speed of $\sim 3 \mu\text{m/s}$ (using the drag coefficient from Table 2.2). The slow drift is superimposed on the rapid, randomly directed thermal motion. At this speed the protein will sediment through a distance of 100 mm , a typical length of a centrifuge tube, in about 10 hours.

ELECTROSTATIC FORCES. A particle with charge q , in an electric field of strength E , will experience a force $F = qE$. An ion such as sodium experiences an electrostatic force when it moves through an ion channel in the plasma membrane. The charge on the ion is $160 \times 10^{-21} \text{ coulombs}$ (see the table of physical constants on the rear endpapers), and the electric field across a typical plasma membrane is $15 \times 10^6 \text{ V/m}$ (60 mV potential across the 4-nm -thick membrane). The corresponding force is 2.4 pN . A similar-sized force exists between two monovalent ions in water that are separated by 1 nm (Problem 2.7): The force will be smaller in a salt solution due to charge screening, but will be larger in the interior of proteins where the dielectric constant is low.

Van der Waals forces are also electrostatic: They arise from the charge separation induced by nearby atoms. Van der Waals forces can be as high as 100 pN per nm^2 of protein-protein interface (Appendix 3.1).

MAGNETIC FORCES. Magnetic forces are very small at the molecular level because molecules interact only very weakly with magnetic fields. For example, the maximum force on a proton, the nucleus with the largest magnetic moment, in the strongest nuclear magnetic resonance (NMR) machines is only on the order of 10^{-12} pN . Thus even for a huge protein with 3000 amino acids and 60,000 atoms, subject to a very strong magnetic field, the magnetic force is less than 10^{-6} pN .

Table 2.2 Physical properties of a globular protein of molecular mass 100 kDa

Property	Value	Comment
Mass	166×10^{-24} kg	Mass of 1 mole/Avogadro constant
Density	1.38×10^3 kg/m ³	1.38 times the density of water
Volume	120 nm ³	Mass/density
Radius	3 nm	Assuming it is spherical
Drag coefficient ^a	60 pN·s/m	From Stokes' law (Chapter 3)
Diffusion coefficient ^a	67 μm ² /s	From the Einstein relation (Chapter 4)
Average speed ^b	8.6 m/s	From the Equipartition principle (Chapter 4)

Note: 1 nm = 10⁻⁹ m, but 1 nm³ = (1 nm)³ = 10⁻²⁷ m³.

^aIn water at 20°C

^bRoot-mean-square (the square root of the average value of the square of the velocity)

In a simplified approach, a protein can be thought as a mechanical device composed of

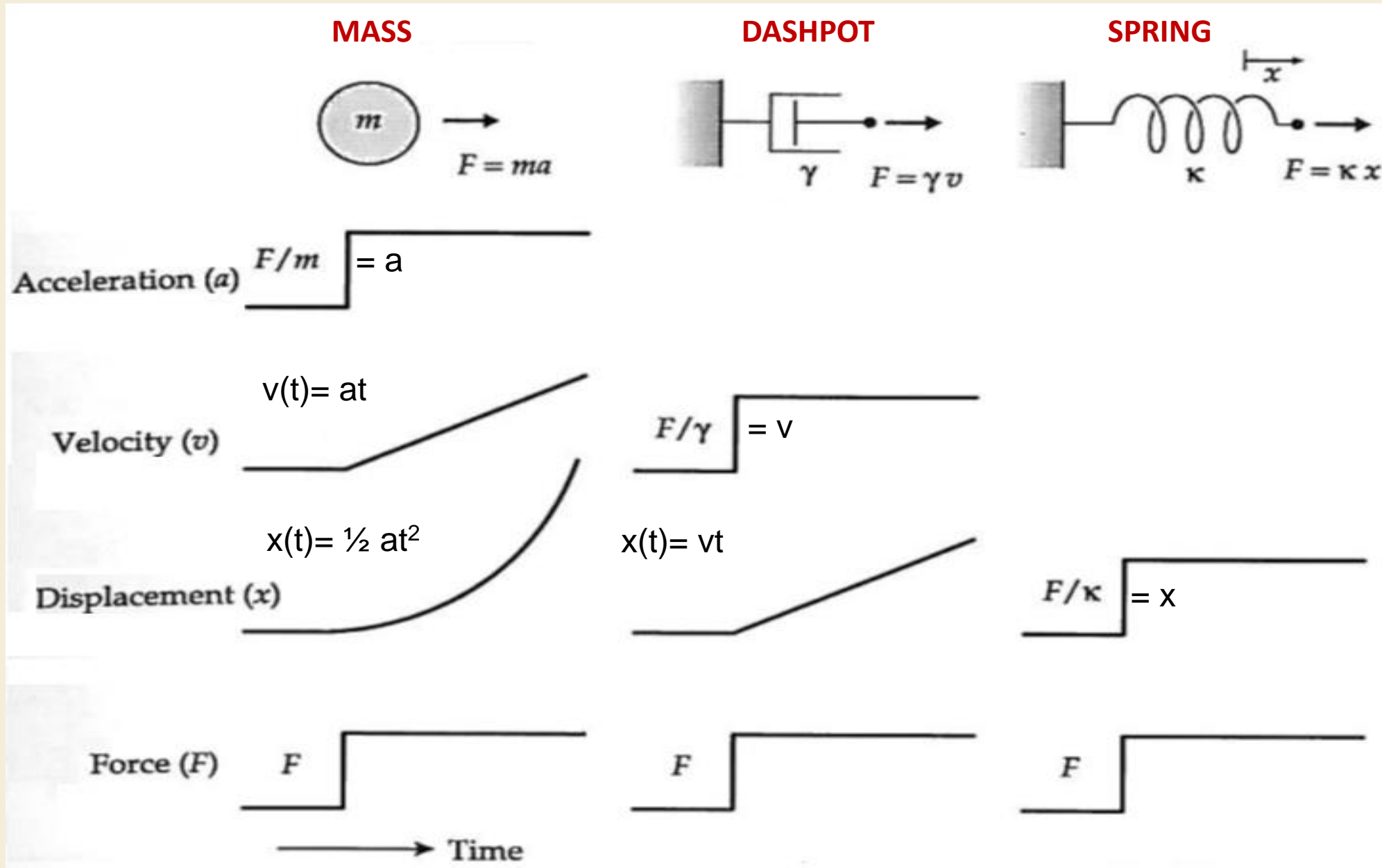
atoms that have mass,

connected by **bonds that have elasticity**, like springs,

and moves **in liquid environment**, facing **viscosity** like dashpots.

All mechanical devices can be built with three fundamental mechanical elements:

SPRING, DASHPOT, MASS.



Example. The force generated by the bacterial motor.

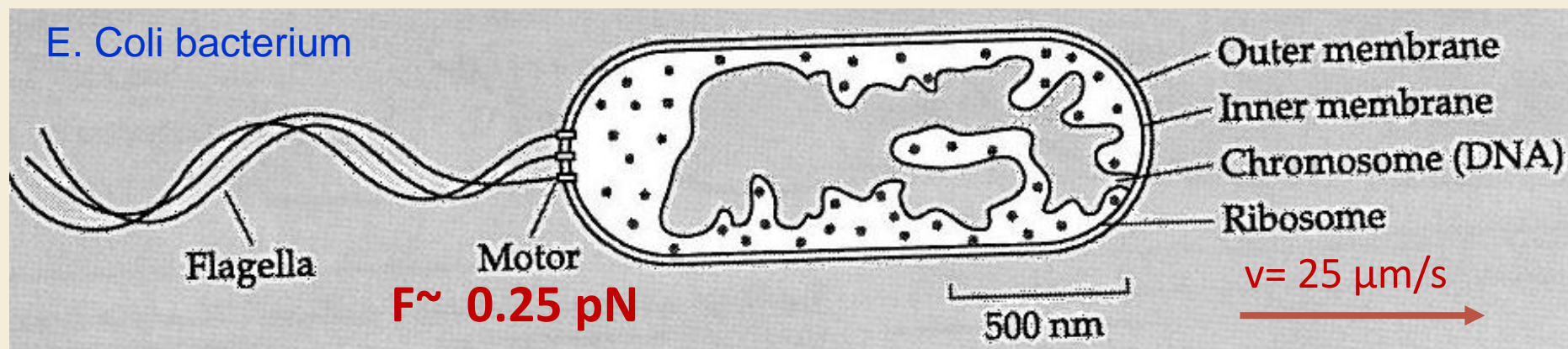
The bacterial flagellar motor should generate a force to move an E. Coli bacterium through water at a constant velocity $v = 25 \mu\text{m/s}$, which is the force to do this ?

$\eta \sim 1 \text{ mPa s}$ – water viscosity, $D \sim 1 \mu\text{m}$ (diam of E. Coli)

$$F = \gamma \cdot v = 3\pi\eta Dv$$

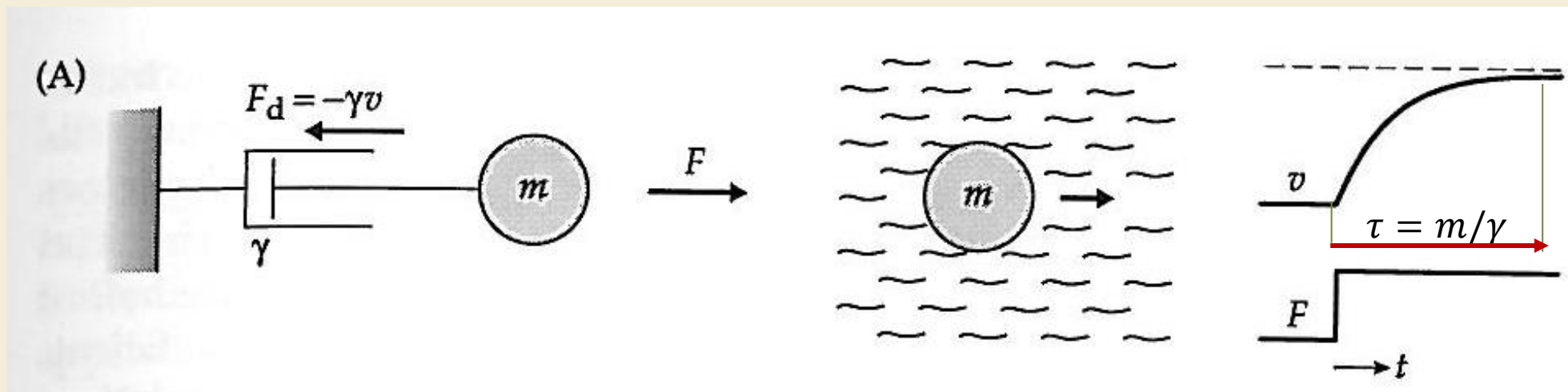
$$\gamma = 3\pi D \eta \sim 10 \text{ mPa s } \mu\text{m} = 10^{-8} \text{ N s/m}$$

$$F \sim 250 \text{ mPa } \mu\text{m}^2 = \mathbf{0.25 \text{ pN}}$$



Motion of Combinations of Mechanical Elements

A) **DASHPOT and MASS**. Model for the movement of a cell / bacterium or a protein through a liquid



Eq of motion

$$m \frac{dv}{dt} + \gamma v = F$$

Solution

$$v(t) = \frac{F}{\gamma} \left[1 - \exp\left(-\frac{t}{\tau}\right) \right]$$

Time constant

$$\tau = \frac{m}{\gamma}$$

Example. Inertia of a bacterium vs protein

Consider a bacterium swimming through water at a constant velocity $v(0) = 25 \mu\text{m/s}$.

How long will it continue to coast after its motor has stopped ?

$$m \frac{dv}{dt} + \gamma v = 0 \quad v(t) = v(0) \cdot \exp\left(-\frac{t}{\tau}\right) \quad \tau = \frac{m}{\gamma}$$

$$m \approx 0.33 \cdot 10^{-15} \text{ kg} \quad \tau_b = 3.3 \mu\text{s} !$$

$$\gamma_b \approx 10 \text{ mPa s } \mu\text{m} = 10^{-8} \text{ N s/m}$$

$$x_{stop} = \int_0^{\tau} v(0) \cdot \exp\left(-\frac{t}{\tau}\right) dt = v(0) \cdot \tau \approx 1 \text{ \AA}$$

Less than the diameter of a water molecule ($\sim 2.7 \text{ \AA}$)

For a globular protein of 100 kDa the average instantaneous speed of such a protein is $v = 8.6 \text{ m/s}$

After the protein gains speed due to molecular collisions with solvent molecules, the velocity persists for only a very short time as other collisions rapidly randomize the protein's direction of travel.

$$m_p \approx 0.166 \cdot 10^{-21} \text{ kg} \quad \tau_p = 2.8 \text{ ps} !!!$$

$$\gamma_p \approx 60 \cdot 10^{-3} \text{ mPa s } \mu\text{m} = 0.6 \cdot 10^{-10} \text{ N s/m}$$

$x_{stop} = 0.24 \text{ \AA}$ the distance that the protein moves before its speed is randomized by molecular collisions

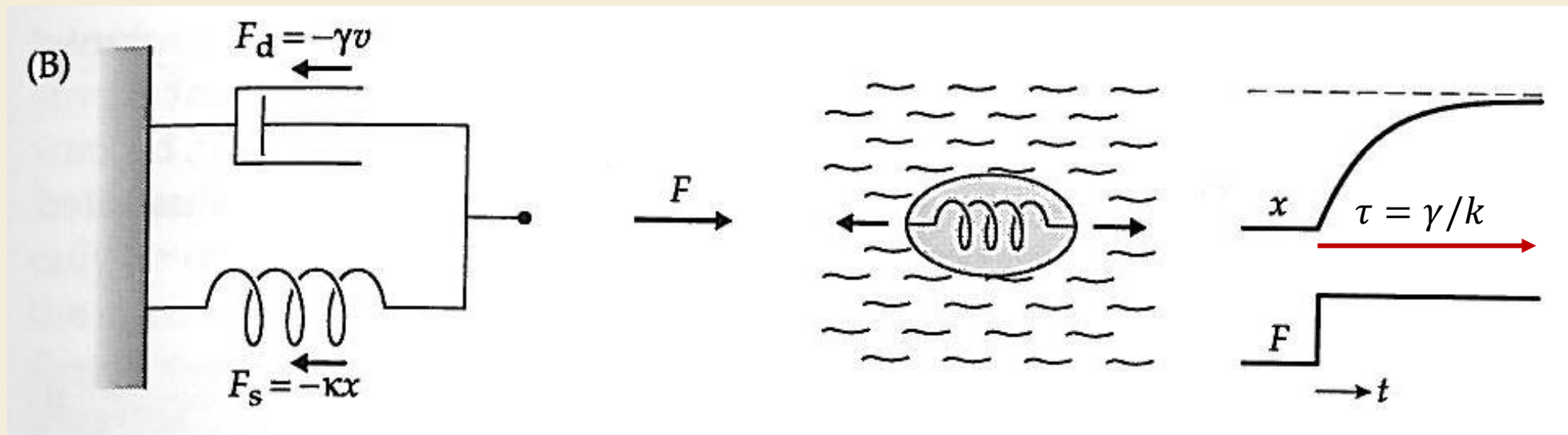
Both time constants are small, but there are 6 orders of difference between the two !

Motion of Combinations of Mechanical Elements

B) SPRING and DASHPOT in parallel.

Model for a compliant low- mass object that is deformed in a liquid, such as a **protein that undergoes a global conformational change**.

It can be used also to model a viscoelastic material, such as **skin, that takes finite time to adopt a new shape**.



Time constant

$$\gamma \frac{dx}{dt} + kx = F$$

$$x(t) = \frac{F}{k} \left[1 - \exp\left(-\frac{t}{\tau}\right) \right]$$

$$\tau = \frac{\gamma}{k}$$

Example. The timescale of protein conformational changes.

Globular protein 100 kDa, the global conformational changes of the whole protein are relatively slow

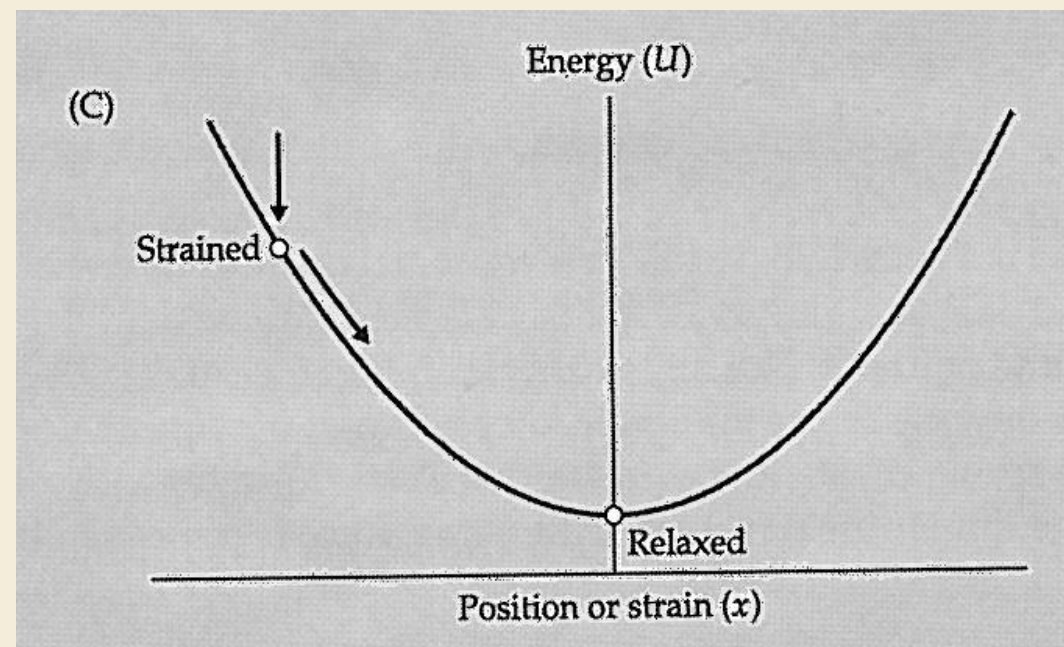
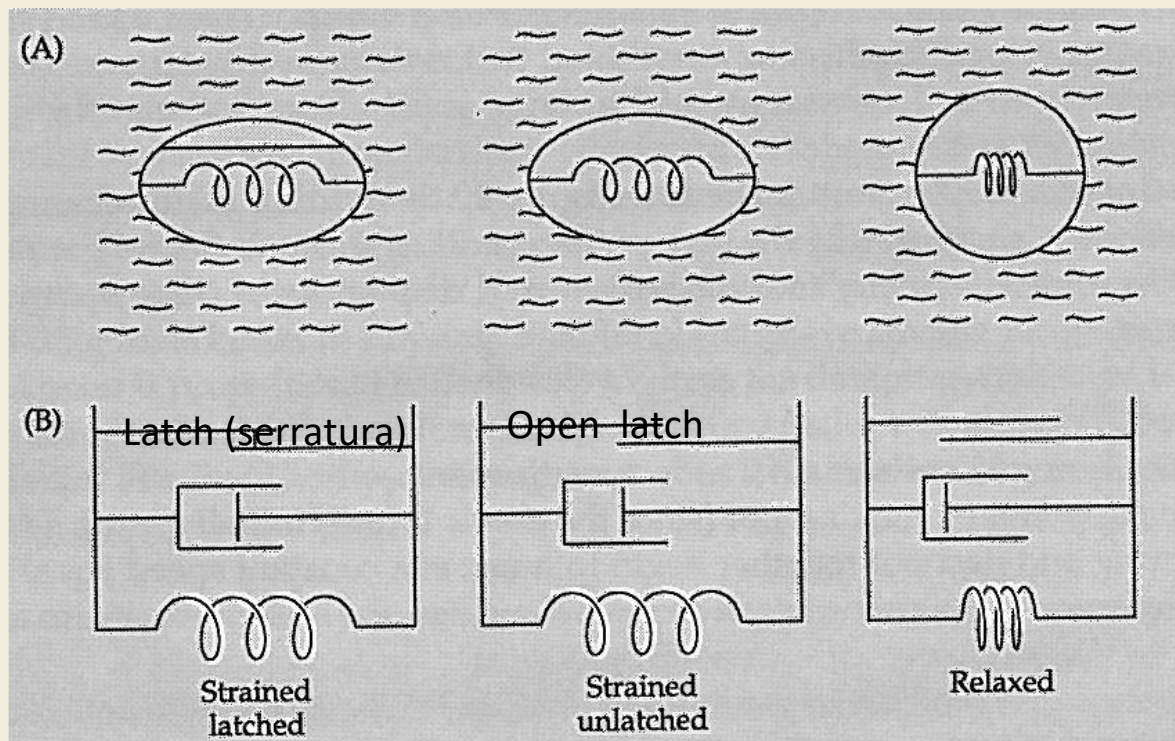
Protein held in a strained conformation due to an internal strut (montante)

Protein relax, changes shape \rightarrow unstrained conformation

$$\gamma = 60 \text{ pN s / m} \quad k = 4 \text{ pN/nm}$$

$$\tau = \frac{\gamma}{k}$$

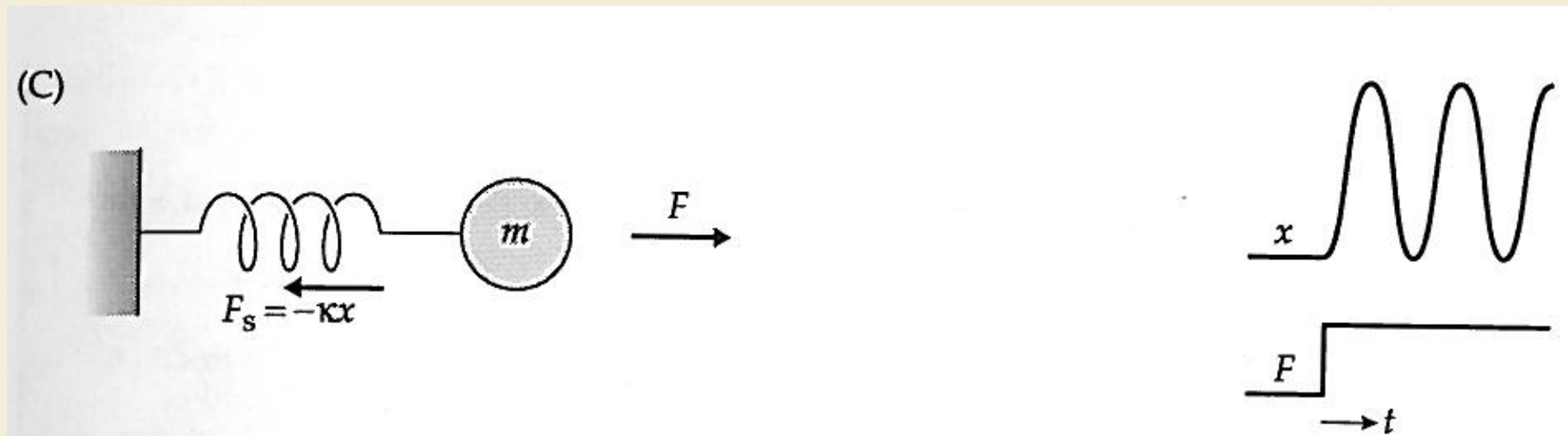
$$\tau = 15 \text{ ns}$$



The global conformation changes in protein occur in nanoseconds

Motion of Combinations of Mechanical Elements

C) **MASS and SPRING** in serie. Model to describe the **vibrations of the atomic bonds**.



$$m \frac{d^2 x}{dt^2} + kx = F$$

$$x(t) = \frac{F}{k} [1 - \cos(\omega t)]$$

Harmonic motion

$$\omega = \sqrt{\frac{k}{m}}$$

Example. **Vibration of chemical bonds.**

Chemical bonds can be thought as having stiffness (chemical bonds vibrate at frequency $\omega=2\pi f$, which can be detected spectroscopically when the molecule absorbs light of the same frequency as the molecular vibration).

Ex: the fundamental vibration frequency of the **H-Cl** bond in HCl is $f= 89.6 \cdot 10^{12}$ Hz (2990 cm^{-1})

The corresponding wavelength is $\lambda= c / \nu= 3.53 \text{ }\mu\text{m}$

The appropriate mass $m \sim 1.63 \cdot 10^{-27}$ kg (approx mass of the hydrogen nucleus)

$$\omega = \sqrt{\frac{k}{m}}$$

Stiffness $k= m \omega^2= 517 \text{ N/m}$ – **very stiff !!!**

Example. **Protein vibrations.**

Consider the motor protein myosin.

Motor domain has a mass $m \sim 0.166 \cdot 10^{-21}$ kg and stiffness $k \sim 4 \text{ pN/nm}$.

The vibration frequency is calculated to be: $f \sim 10^9$ Hz, which means a period of oscillation $T= 1 \text{ ns}$.

By contrast, the relaxation time is 15 ns

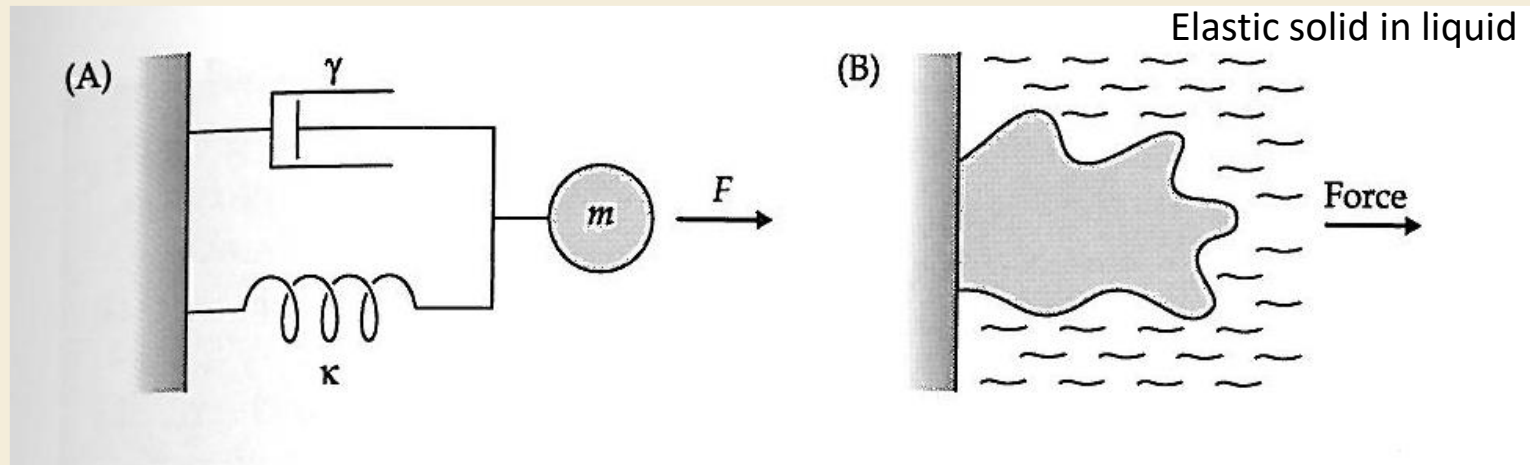
Does the protein oscillate when it detaches from the actin filament or does it creep exponentially into its relaxed state ?

The answer requires solution of the full model, with mass, spring, and dashpot, and it shows that the protein creeps rather than rings.

MASS and SPRING with DAMPING.

Simple mechanical model of a protein undergoing a large scale conformational change that is damped by the surrounding fluid, and possibly by internal viscosity.

This model captures the main qualitative features of more complex models in that it can display oscillatory or monotonic motions depending on the strength of the damping.



$$m \frac{d^2 x}{dt^2} + \gamma \frac{dx}{dt} + kx = F$$

The solution depends on whether the **damping** is:

small

$$\frac{\gamma^2}{4mk} < 1$$

or

large

$$\frac{\gamma^2}{4mk} > 1$$

Underdamped Motion ($\gamma^2 < 4m\kappa$)

$$x(t) = \frac{F}{\kappa} \left[1 - \exp\left(-\frac{t}{\tau}\right) \frac{\sin(\omega t + \phi)}{\sin \phi} \right] \quad (\text{A2.1})$$

where

$$\tau = \frac{2m}{\gamma}, \quad \omega^2 = \omega_0^2 - \frac{1}{\tau^2}, \quad \omega_0^2 = \frac{\kappa}{m}, \quad \tan \phi = \omega\tau$$

Overdamped Motion ($\gamma^2 > 4m\kappa$)

$$x(t) = \frac{F}{\kappa} \left[1 - \frac{\tau_1}{\tau_1 - \tau_2} \exp\left(-\frac{t}{\tau_1}\right) + \frac{\tau_2}{\tau_1 - \tau_2} \exp\left(-\frac{t}{\tau_2}\right) \right] \quad (\text{A2.2})$$

where

$$\tau_1 = \frac{\gamma + \sqrt{\gamma^2 - 4m\kappa}}{2\kappa} \quad \text{and} \quad \tau_2 = \frac{\gamma - \sqrt{\gamma^2 - 4m\kappa}}{2\kappa}$$

Both τ_1 and τ_2 satisfy $(m/\tau) + \kappa\tau = \gamma$. When the motion is highly overdamped ($\gamma^2 \gg 4m\kappa$), the time constants become $\tau_1 = \frac{\gamma}{\kappa}$ and $\tau_2 = \frac{m}{\gamma}$, where $\tau_1 \gg \tau_2$.

Critically Damped Motion ($\gamma^2 = 4m\kappa$)

$$x(t) = \frac{F}{\kappa} \left[1 - \left(1 + \frac{t}{\tau} \right) \exp\left(-\frac{t}{\tau}\right) \right] \quad (\text{A2.3})$$

where

$$\tau = \frac{2m}{\gamma} = \frac{\gamma}{2\kappa} = \sqrt{\frac{m}{\kappa}}$$

This solution is monotonic, like that in the overdamped case. Note that there is a lag, of duration $\sim \tau/2$, before the displacement starts to rise quickly.

Unrealistic case:

Globular protein – 16 MDa (hypothetical)

Stiffness $k = 30 \text{ N/m}$ (very rigid)

Little damping $\gamma = 150 \text{ pN s/m}$ (unrealistic)

More realistic case:

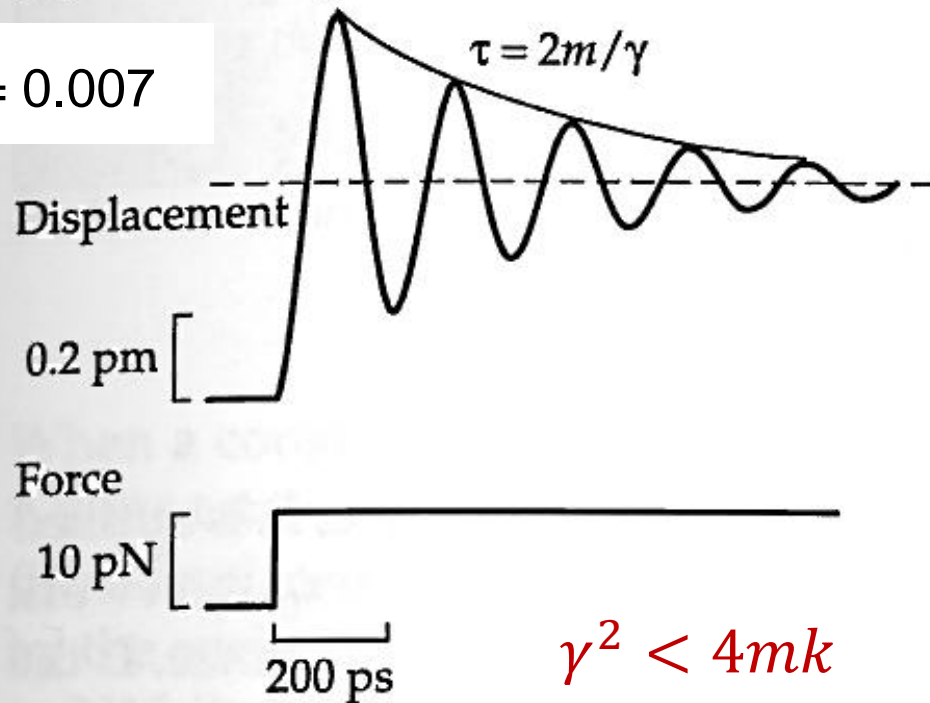
protein undergoing a large scale conformational change that is damped by the surrounding fluid, and by internal viscosity.

Globular protein MM=100 kDa ; Stiffness $k = 4 \text{ pN/nm}$;

damping $\gamma = 60 \text{ pN s/m}$

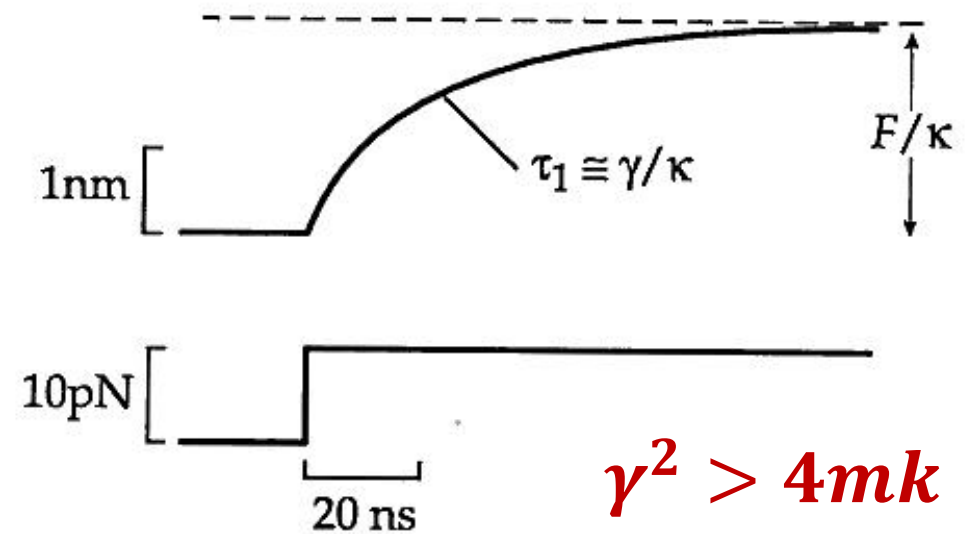
(C) Underdamped motion

$$\frac{\gamma^2}{4mk} = 0.007$$



(D) Overdamped motion

$$\frac{\gamma^2}{4mk} \sim 1400$$



The inertial forces are usually very small at the microscopic and molecular levels, so that the overdamped case usually applies.

Energy of chemical bonds:

the dissociation energy is seen as being approximately equal with the potential energy in the bond:

$$U = \frac{1}{2}kd^2, \text{ where } d \text{ is the extension required to break the bond, } d \sim 0.05 \text{ nm.}$$

For H-Cl, the stiffness $k \sim 517 \text{ N/m} \rightarrow U \sim 650 \times 10^{-21} \text{ J} = 650 \text{ pN} \cdot \text{nm} \rightarrow \mathbf{U \sim 161 K_B T}$

(K_B Boltzmann ct: $K_B = 1.38 \cdot 10^{-23} \text{ J/K}$; T- temperatue, e.g. T=300 K ; $1 K_B T \sim 4 \cdot 10^{-21} \text{ J} = 4 \text{ pN} \cdot \text{nm}$)

Energy stored in protein conformational changes:

Myosin molecule. The stiffness is about $k \sim 4 \cdot 10^{-3} \text{ N/m}$ (or 4 pN/nm)

For a conformational change of $d=5 \text{ nm}$ the total energy $U = \frac{1}{2}kd^2 = 50 \text{ pN nm} = 50 \cdot 10^{-21} \text{ J}$, $\mathbf{U \sim 12.5 K_B T}$

This energy is approximately half of the chemical energy derived from hydrolysis of the gamma phosphate bond of ATP.

We can generalize this argument to global conformational changes of other protein machines:

The energies are on the order of 10 to 100 x 10⁻²¹ J (2.5 to 25 K_BT), conformational changes are on the order of 1 to 10 nm.

Therefore the stiffnesses are on the order of 0.2 to 200 pN/nm.

Get a feeling for what proteins are like mechanical devices

Questions:

- How rigid the proteins are ? Density, viscosity ?
- How quickly do they move and change shape ?
- What happens when a protein is struck by a force: does it ring like a fork (underdamped motion), or does it creep monotonically into a new shape (overdamped motion ?).

$$m = \rho V$$

Proteins are composed of relatively light **components**:
carbon, oxygen, nitrogen, and hydrogen

Proteins are about **40 % denser than water**, with different proteins having slightly different densities.

The **average density** of proteins is consider to be:

$$\rho = 1.38 \times 10^3 \text{ kg/m}^3$$

Rule of thumb:

The density of proteins is such that each kDa of protein occupies a volume of about 1.2 nm^3 .

The **SI of mass** is **kg**, but in biochemistry the mass of proteins and other biomolecules is usually expressed as **molecular mass**, defined as the mass in grams of a mole of the molecules.

The unit is the **Dalton** : $1 \text{ Da} = 1.66 \times 10^{-24} \text{ g}$

Ex: A protein of 100 kDa has a mass, $m = 166 \times 10^{-21} \text{ g}$

The volume V , occupied by such a protein is: $V \sim 120 \text{ nm}^3$.

Table 3.1 Densities of molecules, proteins, organelles, and cells relative to water

Substance	Density (relative to water ^a)
Water	1.00
Glycerol	1.26
Glycine	1.16 (solid)
Alanine	1.40 (solid)
Glutamic acid	1.46 (solid)
Hemoglobin	1.33 (in solution)
Trypsin	1.38 (in solution)
Lysosyme	1.42 (in solution)
Chromosome	1.36
Virus	1.15
Mitochondrion	1.18
Synaptic vesicle	1.05
Erythrocyte	1.10
Fibroblast	1.05

For a homogenous and isotropic solid:

$$\boxed{\frac{F}{A}} = E \boxed{\frac{\Delta L}{L}} \quad E : \text{constant [N/m}^2\text{] [Pa]}$$

pressure strain

E - Young's modulus or elastic modulus

- Young's modulus E is a material property:
it does not depend on the object size or shape

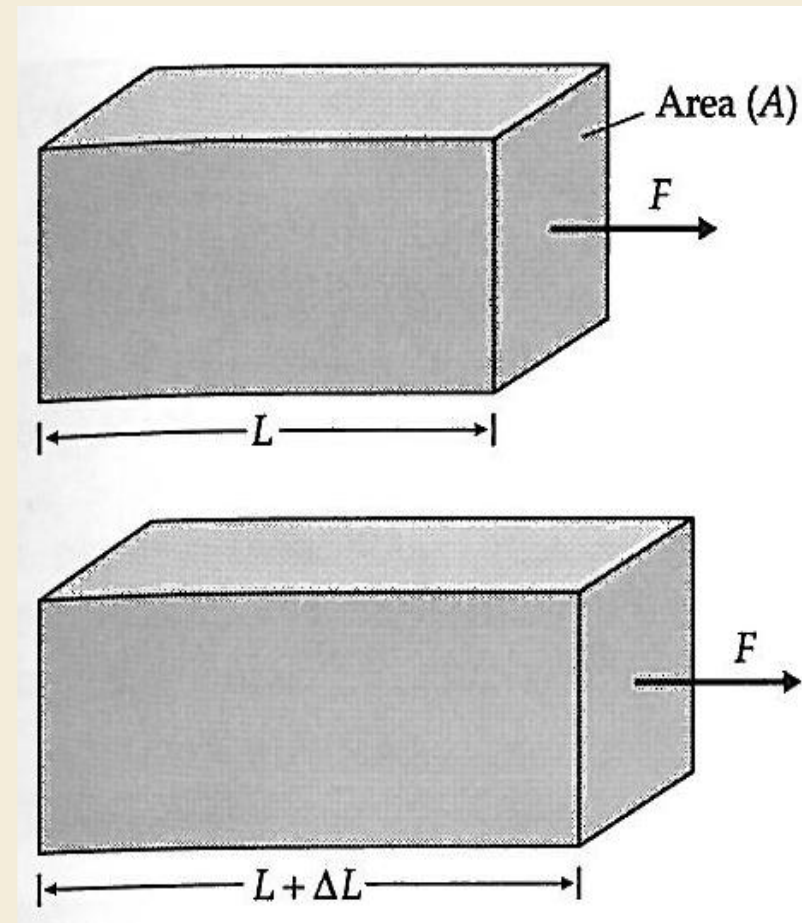
$$\boxed{F} = k \boxed{\Delta L} = \frac{AE}{L} \Delta L \quad k : \text{constant [N/m]}$$

force extension $F \sim \Delta L$; Hooke's law

K – stiffness

- The stiffness, k , of an object does depend on the size and shape.

A solid strained by a tensile force.



Material	Young's modulus, E (GPa)
Carbon nanotube	1300
Diamond	1200
Steel (stainless)	211
Glass (quartz)	73
Wood (fir, along grain)	16
Plexiglas	3
Plastic (polypropylene)	2.4
Teflon (PTFE)	0.34
Rubber (polyisoprene)	0.02
Silk (<i>Bombyx mori</i>)	5–10
Keratin (hair)	2.4
Actin	2.3
Collagen	2
Tubulin	1.9
Elastin	0.002

For many materials (e.g. metals, plastics and structural proteins) the Hooke's law $F = k \Delta L$ applies only for forces that cause strains up to:

$$\frac{\Delta L}{L} = 0.1 - 1 \%$$

At higher forces the material yields and the yield pressure is called **tensile strength**.

Other materials such as rubber and proteins like elastin and titin can be strained up to 100 % or more.

The behavior / motion of an object in response to mechanical force - oscillatory (underdamped) or monotonic (overdamped) - depends on the relative magnitudes of the inertial and viscous forces.

These in turn depend on the material properties: mass, stiffness and damping.

The scaling argument:

as the dimension of an object gets smaller, the viscous forces increase relative to the inertial forces, and as a result, the global motions of small, comparatively soft objects such as proteins in aqueous solution are expected to be overdamped.

Considering a crude mechanical model of a globular protein as a homogeneous and isotropic cube with side L , density ρ , and Young's modulus E , damped by fluid viscosity η .

The mass: $m = \rho V = \rho L^3$. The stiffness: $k = EL$

The drag force associated with a global conformational change that alters the shape of a protein: $F = -\gamma v$, with $\gamma = 3\pi \eta L$.

Overdamped condition:

$$\frac{\gamma^2}{4mk} > 1$$



$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

Overdamped:

$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

How small must a protein be to ensure that its motion is overdamped and that it does not oscillate when subject to an external force ?

For the middle rigid proteins the Young's moduli, $E \sim 1 \text{ GPa}$; the density, $\rho \sim 10^3 \text{ kg/m}^3$, viscosity of water $\eta \sim 1 \text{ mPa s}$.

$$\frac{\eta^2}{\rho E} \approx 1 \text{ nm}^2$$



$$L < 5 \text{ nm}$$

This length corresponds to a medium-sized globular protein of ~ 1000 amino acids.

Average MM of an amino acid is $\sim 100 \text{ Da}$.

Thus the model predicts that global motions of rigid globular proteins or protein domains of molecular weight less than **100 kDa** should be overdamped.

$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

The motion of larger proteins is also overdamped because:

The rigidity of energy-transducing proteins, such as motor proteins, and the ribosomes is likely to be much less than that of rigid proteins like those of the cytoskeleton. Considering a protein undergoing a $x=2$ nm (modest) conformational change and assuming this is associated to a large amount of mechanical work, say $W=25 K_bT$ (equal to the free energy of hydrolysis of the gamma phosphate bond of one molecule of ATP) from the energy $W= \frac{1}{2} kx^2$ we get the stiffness $k= 2W/x= \mathbf{50 pN/nm}$, which is much smaller than the stiffness of a rigid protein of length 10 nm and Young's modulus $E= 2$ GPa.

This value of stiffness leads to a much greater **characteristic length $L= 50$ nm**, implying that even the motion of a ribosome, one of the largest protein machines, would be overdamped.

Moreover, since we consider a small value for the conformational change and a large value for the work, even this low stiffness is likely to be an overestimate; indeed the stiffness of motor proteins is on the order of pN/nm (0.001 N/m) \rightarrow arguing once more that that protein motions are overdamped.

Example 3.6 Ribosome If a large protein were to oscillate, how fast and how large might these oscillations be? Consider the ribosome, a globular protein–RNA enzyme complex of diameter ~30 nm (Ban et al., 1999, Clemons et al., 1999). The ribosome is the molecular machine that synthesizes proteins. If the ribosome were very rigid ($E = 1$ GPa), and the only damping came from the surrounding fluid, then it would oscillate at a frequency of $\sim(\kappa/m)^{0.5}/2\pi$ Hz = $(E/\rho)^{0.5}/2\pi L \sim$ 5 GHz, corresponding to a period of 200 ps. The oscillation would decay quickly, with a time constant of $2m/\gamma \sim (2/3\pi)\rho L^2/\eta \sim$ 200 ps (Equation A2.1 in the Appendix). In other words, the oscillations would die out after only a few cycles. The magnitude of the oscillations would depend on the size of the force. Suppose that the force did work on the protein equal to 100×10^{-21} J (= 25 kT), the free energy associated with the hydrolysis of one molecule of ATP (Chapter 14). If we think of this chemical energy as being converted into mechanical potential energy within the protein during the protein synthesis reaction, then the amplitude, x , of the deformation would be only ~ 0.8 Å (energy = $\frac{1}{2}\kappa x^2$, and we assume that ribosome is as rigid as a cytoskeletal protein with $\kappa = EL = 30$ N/m). The oscillations, if they occurred, would be very small indeed. Considering that the lifetimes of different chemical states are in the order of microseconds to milliseconds, it is unlikely that such small oscillations, even if they were to occur, would play important roles in the chemistry of protein synthesis.

$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

Based on the scaling argument, since cells have linear dimensions about 1000 larger than those of proteins, **one might expect that cells undergo underdamped motions.**

Experimentally it is shown that **this is not the case**: the motions of the cells are very highly damped.

For example, the cytoplasm of macrophages that have ingested 1 μm diameter magnetic particles can be perturbed using a weak external magnetic field. The particles reorient extremely slowly, with time constant of minutes.

The apparent **intracellular viscosity is very high, approx 1000 Pa s** and the motion is highly overdamped. Because actin gels crosslinked with the actin binding protein ABP have similar viscoelastic properties to cells, it is likely that the viscoelasticity of cells arises from the stiffness and damping on cytoskeletal filaments. Since the long cytoskeletal filaments are highly damped, so too are cells.

The cytoskeletal filaments **form a gel with a mesh size of about 50 nm**. Small solutes and proteins can readily diffuse through the pores, but the motion of larger particles, such as ribosomes and organelles is severely restricted.

Summary 1

- By considering three mechanical elements – mass, damping, spring - we introduced some of the mechanical concepts required to understand how forces influence protein and cells.

$$\tau = \frac{m}{\gamma}$$

movement of a cell / bacterium or a protein through a liquid, inertia of a protein (100 kDa), measured as time to stop ($\tau = 3 \text{ ps}$)

$$\tau = \frac{\gamma}{k}$$

protein that undergoes a global conformational change ($\tau = 15 \text{ ns}$)

$$\omega = \sqrt{\frac{k}{m}}$$

Vibration of chemical bonds ex. H-Cl , $f = 89.6 \cdot 10^{12} \text{ Hz}$; mass $m \sim 1.63 \cdot 10^{-27} \text{ kg}$ $\rightarrow k = m \omega^2 = 517 \text{ N/m}$ – very stiff

- The mass and spring with damping illustrate that system can respond to mechanical forces in two ways:

they can oscillate or they can move monotonically.

$$\frac{\gamma^2}{4mk}$$

Summary 2

- The mechanical models discussed in this lecture can be generalized in two ways:
 1. Increase the number of mechanical elements to include several masses, springs and dashpots, and even other elements as latches and stops; the equations of motion are solved by balancing the forces across each element.

Ex: Molecular dynamics: each atom in protein and surrounding fluid is represented by a point mass, each bond is represented by a spring with constant stiffness and the damping is dropped from the equation. The ensuing motion is complex and best solved numerically by computer.
 2. Consider the mechanical behavior of «continuum» solids that have material properties such as elasticity, density and viscosity (our approach in next lectures)

Summary 3

- The rigidity of cytoskeleton proteins as actin, tubulin, keratin is similar to that of hard plastics but less than that of other materials such as glass or metal, because proteins are held together by weak Van der Waals bonds
- The rigidity of protein machines/motors undergoing large conformational changes as they transduce chemical energy into mechanical work is expected to be much less than that of structural proteins
- As proteins move and change shape, they experience damping forces from the surrounding fluid as well from the internal friction. These forces arise from the rapid making and breaking of bonds.
- Due to the small size of proteins, the viscous forces are preponderant over the inertial forces → global motions of proteins, especially less rigid ones, are overdamped → they creep rather than oscillate when subject to forces

$$\frac{\gamma^2}{4mk} \cong 25 \frac{\eta^2}{\rho EL^2} > 1$$

$$k = \frac{AE}{L}$$

- The motion of long, thin cytoskeletal filaments are also overdamped, due to their large aspect ratios, this in turn causes the motion of the cells to be overdamped.