

$$1) \quad (a) \quad (T+a)v = b(T_0 - T)$$

$$v = \frac{b(T_0 - T)}{T+a}$$

$$\text{Power } vT = \frac{bT(T_0 - T)}{T+a}$$

$$\text{max power } \frac{d}{dT}(vT) = 0, \quad P = vT$$

$$\frac{dP}{dT} = 0 = \left(\frac{1}{a} + \frac{1}{T}\right) = \left(\frac{T_0}{T^2} - \frac{1}{T}\right)$$

$$T = \frac{-2a \pm \sqrt{4a^2 + 4aT_0}}{2}$$

$$v_{\text{opt}} = \frac{b \left[ 1 + \frac{a}{T_0} - \sqrt{\frac{a^2}{T_0^2} - \frac{a}{T_0}} \right]}{\sqrt{\frac{a^2}{T_0^2} + \frac{a}{T_0}}}$$

A cyclist chooses a gear so that muscle velocity =  $v_{\text{opt}}$   
so as to maximise power o/p.

(b) Thermal activation of molecules leads to progressive loss of correlation along the length of the molecule. The persistence length  $\zeta$  is the length over which correlation is lost by thermal activation

$$kT \sim \frac{1}{2} EI \phi^2 / \zeta$$

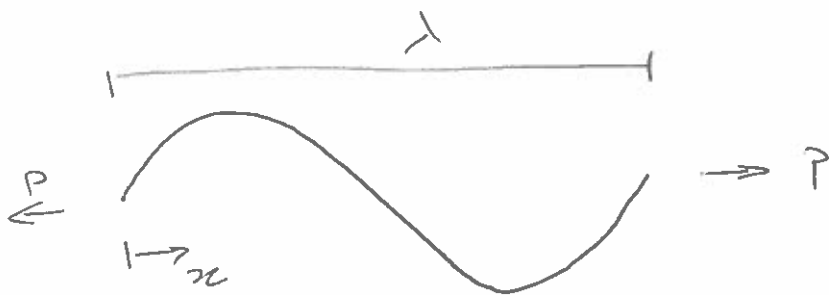
where  $\phi$  is end-to-end rotation

$$\Rightarrow \zeta = \frac{EI}{2kT} \quad \text{taking } \phi \sim 1.$$

Stiff molecules like microtubules have long persistence lengths & remain straight over the length of the cell while intermediate filaments with small  $\zeta$  coil randomly

(c) The cellulose cell wall in plants gives it structural stiffness. Animal cells do not have this & rely on the cytoskeleton for structural support.

2  
(a)



we assume  $w(z) = a \sin \frac{2\pi z}{\lambda}$

$$M(z) = Pw = EI \frac{d^2 u}{dz^2}$$

$$\frac{d^2 u}{dz^2} = \frac{Pa}{EI} \sin \frac{2\pi z}{\lambda}$$

$$u = - \left( \frac{\lambda}{2\pi} \right)^2 \frac{Pa}{EI} \sin \frac{2\pi z}{\lambda}$$

assume  $P \ll \frac{EI, \lambda^2}{4\pi^2} \Rightarrow |u| \ll |w|$

$$e = - \frac{1}{\lambda} \int_0^{\lambda} \left\{ \left[ 1 + \left( \frac{\partial u}{\partial z} + \frac{\partial w}{\partial z} \right)^2 \right]^{\frac{1}{2}} - \left[ 1 + \left( \frac{\partial w}{\partial z} \right)^2 \right]^{\frac{1}{2}} \right\} dz$$

$$\frac{\partial u}{\partial z} = - \frac{\lambda}{2\pi} \frac{Pa}{EI} \cos \frac{2\pi z}{\lambda}$$

$$\frac{\partial w}{\partial z} = \frac{2\pi a}{\lambda} \cos \frac{2\pi z}{\lambda}$$

$$\left[ 1 + \left( \frac{\partial u}{\partial z} + \frac{\partial w}{\partial z} \right)^2 \right]^{\frac{1}{2}} \sim 1 + \frac{1}{2} \left[ \left( \frac{\partial u}{\partial z} \right)^2 + 2 \frac{\partial u}{\partial z} \frac{\partial w}{\partial z} + \left( \frac{\partial w}{\partial z} \right)^2 \right]$$

$$\left[ 1 + \left( \frac{\partial w}{\partial z} \right)^2 \right]^{\frac{1}{2}} \sim 1 + \frac{1}{2} \left( \frac{\partial w}{\partial z} \right)^2$$

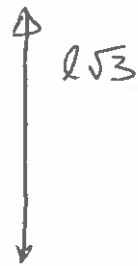
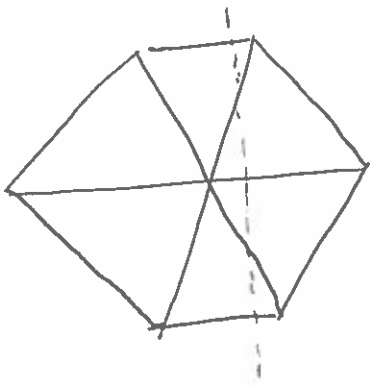
$$e_1 \approx \int_0^l dx \left( \frac{\partial u}{\partial x} \frac{\partial w}{\partial x} \right)$$

$$= \frac{1}{2} \frac{Pa^2}{EI}$$

$$e = \frac{Pa^2}{2EI} \quad ; \quad EI = \frac{E_s}{3} \frac{\pi}{4} \left( \frac{d}{2} \right)^4$$

$$\Rightarrow e = \frac{32}{\pi} \frac{a^2 P}{E_s d^4}$$

(b) By symmetry  $T_B = T_F = T_E = T_C$  &  $T_A = T_D$



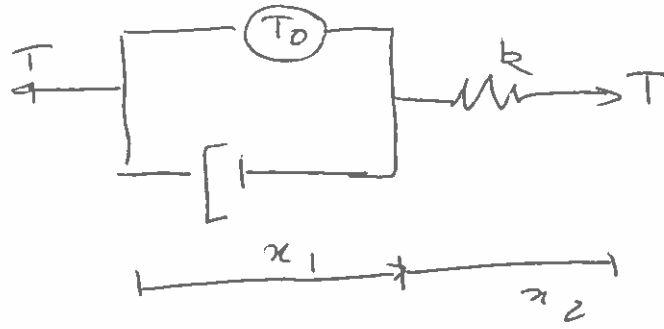
$$\& T_B = T_F = T_E = T_C = 0$$

$$\Rightarrow T_A = T_D = \sqrt{3} l d \Sigma_{11}$$

$$(c) \frac{F_1}{e} = \frac{\Sigma_{11}}{e} = \frac{\pi}{32\sqrt{3}} \frac{E_s d^3}{a^2 l}$$

(3)

(a)



$$k \Delta x_2 = T \quad \& \quad B \dot{x}_1 + T_0 = T$$

Since muscle under isometric tension  $\dot{x}_1 = -\dot{x}_2 = -\frac{\dot{T}}{k}$

$$T + \frac{B}{k} \dot{T} = T_0$$

with  $T(0) = 0$

$$\Rightarrow T(t) = T_0 \left[ 1 - e^{-\frac{kt}{B}} \right] \quad 0 \leq t \leq c$$

~~II~~

$$(b) \quad T(c) = T_0 \left[ 1 - e^{-\frac{kc}{B}} \right]$$

$$\& \quad T + \frac{B}{k} \dot{T} = 0$$

$$\Rightarrow T(t) = \left[ 1 - e^{-\frac{kc}{B}} \right] e^{-\frac{k(t-c)}{B}} \quad c \leq t \leq c+A$$

(c) ~~Problem~~ Problem is linear  $\Rightarrow$  use superposition  
 $\Rightarrow$  between  $c+A$  &  $2c+A$  the response due to 2nd input is

$$T_0 \left[ 1 - e^{-\frac{k(t-c-A)}{B}} \right]$$

$$\& T(t) = T(c) e^{-k \frac{(t-c)}{B}} + T_0 \left[ 1 - e^{-k \frac{(t-c-A)}{B}} \right]$$

(d) The superposition nature of solution suggests that it will be able to capture the differences between a twitch & tetanus.

(4)

(a) Myoglobin stores  $O_2$  & releases it when the environmental  $O_2$  is low. This mechanism gives the enhanced diffusion rate rather than the actual faster transport of  $O_2$

(b) Large molecules such as glucose are transported across the cell membrane via a carrier mediated transport mechanism such as uniports, symports etc.

(c)  
(i) In its E1 conformation the  $Na^+/K^+$  ATPase has 3  $Na^+$  binding sites

4  
(c)  
(i)

In its E1 conformation, the Na<sup>+</sup>/K<sup>+</sup> ATPase has three high-affinity Na-binding sites and two low-affinity K-binding sites accessible to the cytosolic surface of the protein. The K<sub>m</sub> for binding of Na<sup>+</sup> to these cytosolic sites is 0.6 mM, a value considerably lower than the intracellular Na concentration of approx. 12mM; as a result, Na<sup>+</sup> ions normally will fully occupy these sites. Conversely, the affinity of the cytosolic K-binding sites is low enough that K<sup>+</sup> ions, transported inward through the protein, dissociate from E1 into the cytosol despite the high intracellular K concentration. During the E1 → E2 transition, the three bound Na<sup>+</sup> ions become accessible to the exoplasmic face, and simultaneously the affinity of the three Na-binding sites becomes reduced. The three Na<sup>+</sup> ions, transported outward through the protein and now bound to the low-affinity Na<sup>+</sup> sites exposed to the exoplasmic face, dissociate one at a time into the extracellular medium despite the high extracellular Na concentration. Transition to the E2 conformation also generates two high-affinity K<sup>+</sup> sites accessible to the exoplasmic face. Because the K<sub>m</sub> for K<sup>+</sup> binding to these sites (0.2 mM) is lower than the extracellular K<sup>+</sup> concentration (4 mM), these sites will fill with K<sup>+</sup> ions. Similarly, during the E2 → E1 transition, the two bound K<sup>+</sup> ions are transported inward and then released into the cytosol.

(ii)

Overall per ATP molecule hydrolyzed the pump moves 3 Na<sup>+</sup> ions & 2 K<sup>+</sup> ions. Increasing the ATP concentration will increase the rate of the pump.



**Q1 Muscle power + qualitative reasoning of biological fibres**

15 attempts, Average mark 73%

A question that was well-attempted. Most students calculated muscle power correctly but gave poor explanation of persistence length and the cell walls of plant and animal cells

**Q2 Modulus of a network of wavy struts**

5 attempts, Average mark 60%

Generally poorly attempted and few attempts too. The students struggled to calculate the effective modulus of a wavy strut and even in calculating strut forces in the network by method of sections

**Q3 Hill muscle model**

11 attempts, Average mark 67%

Generally well attempted but they struggled on using superposition to calculate the effect of multiple stimuli.

**Q4 Qualitative question on ion pumps and transport mechanisms**

14 attempts, Average mark 70%

Generally well attempted although the explanation of ATP pumps was generally not adequate.