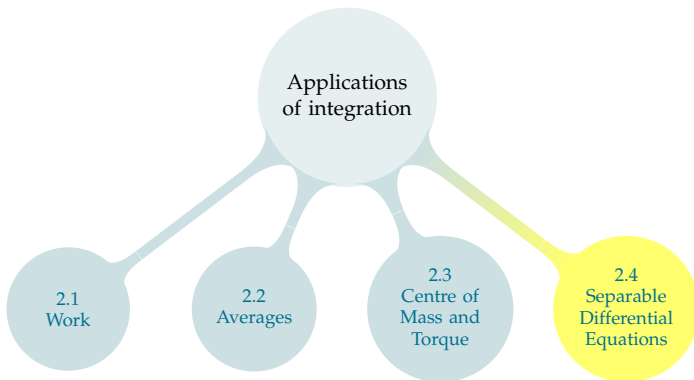
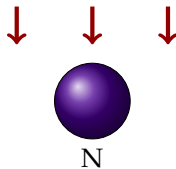


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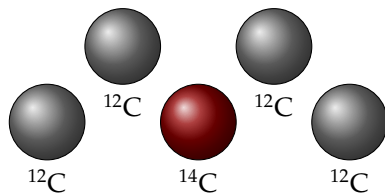


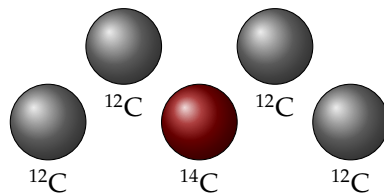
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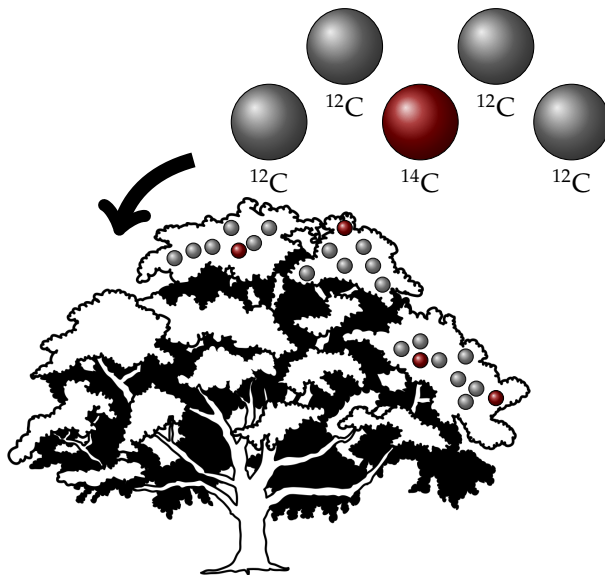


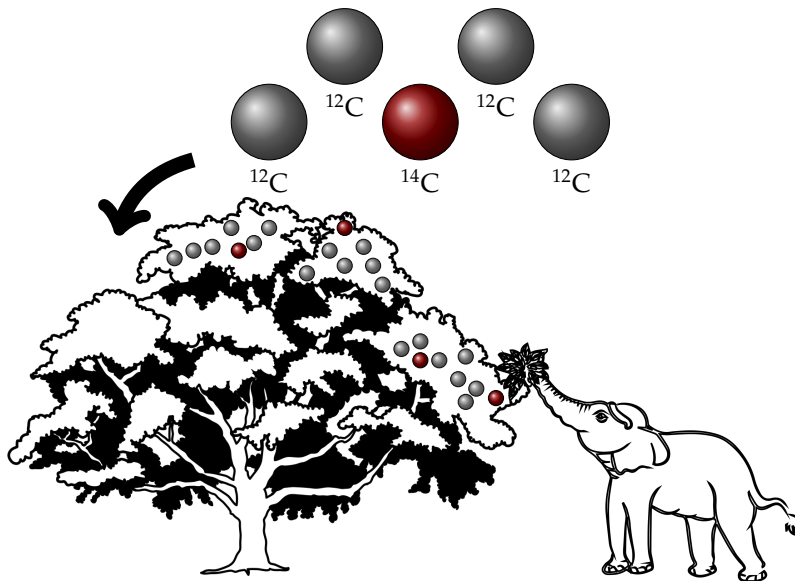


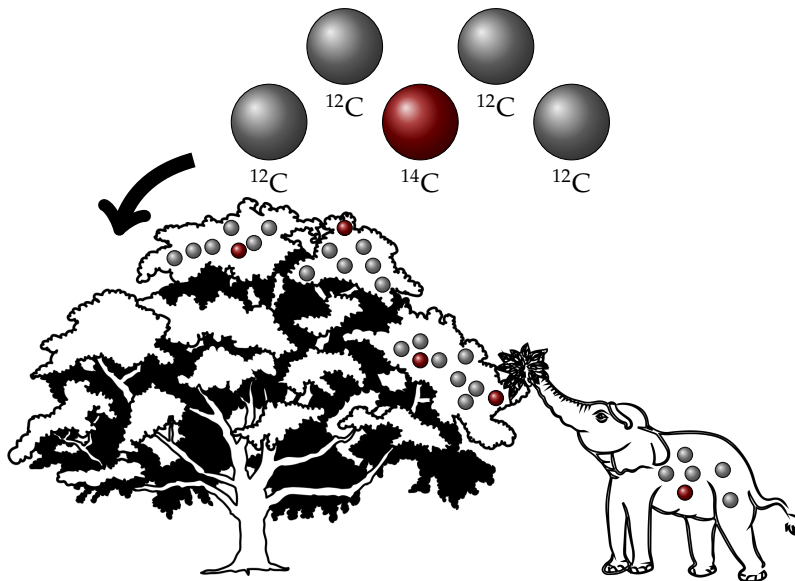
^{14}C

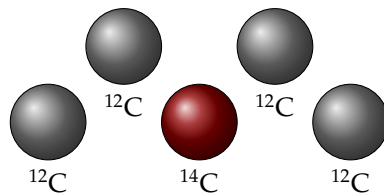


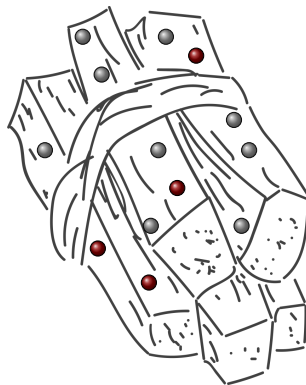
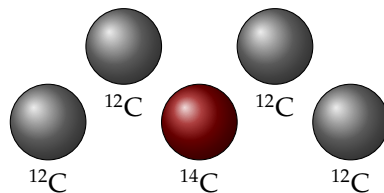


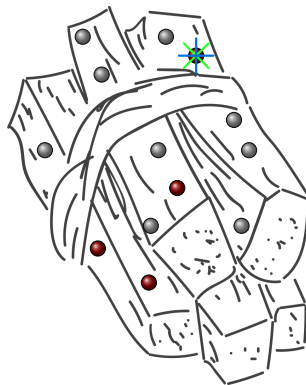
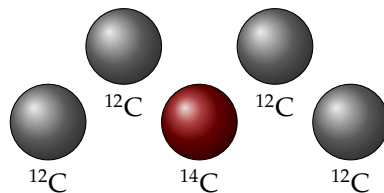


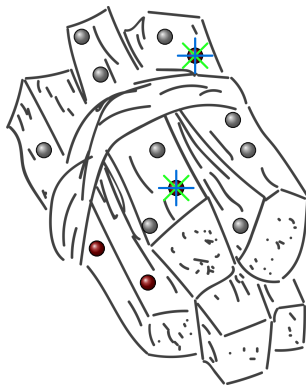
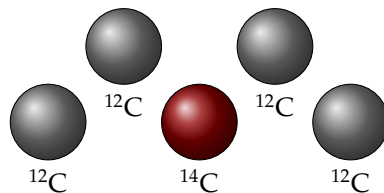


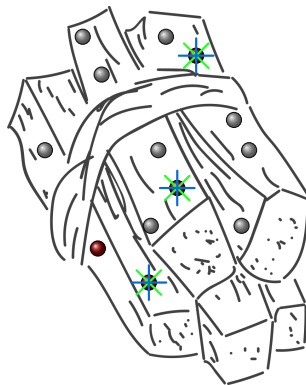
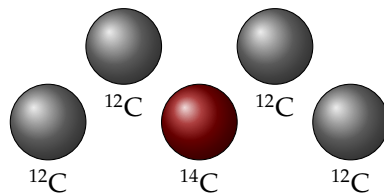












RADIOACTIVE DECAY

One model for radioactive decay says that the rate at which an isotope decays is proportional to the amount present. So if $Q(t)$ is the amount of a radioactive substance, then

$$\frac{dQ}{dt} = -kQ(t)$$

for some constant¹ k .

This is a first-order linear differential equation. Its explicit solutions have the form:

¹By including the negative sign, we ensure k will be positive, but of course we could also write " $\frac{dQ}{dt} = KQ(t)$ for some [negative] constant K ".

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$$Q(t) = Ce^{-kt}$$

where $C = Q(0)$.

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HALF-LIFE

The **half-life** of an isotope is the time required for half of that isotope to decay. If we know the half-life of a substance is $t_{1/2}$, and its quantity at time t is given by $Q(0)e^{-kt}$ we can find the constant k :

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$$\frac{1}{2}Q(0) = Q(t_{1/2}) = Q(0)e^{-kt_{1/2}}$$

$$\frac{1}{2} = e^{-kt_{1/2}}$$

$$2 = e^{kt_{1/2}}$$

$$\log 2 = kt_{1/2}$$

$$\frac{\log 2}{t_{1/2}} = k$$

Plugging this back in gives us a more intuitive equation for the

quantity of a radioactive substance over time:

$$\begin{aligned} Q(t) &= Q(0)e^{-\frac{\log 2}{t_{1/2}}t} \\ &= Q(0) \cdot \left(\frac{1}{2}\right)^{\frac{t}{t_{1/2}}} \end{aligned}$$

So if $t = t_{1/2}$, the initial amount is cut in half; if $t = 2t_{1/2}$, the initial amount is cut in half twice (i.e. quartered), etc.

Radioactive Decay

The function $Q(t)$ satisfies the equation $\frac{dQ}{dt} = -kQ(t)$ if and only if

$$Q(t) = Q(0) e^{-kt}$$

The half-life is defined to be the time $t_{1/2}$ which obeys $Q(t_{1/2}) = \frac{1}{2} Q(0)$. The half-life is related to the constant k by $t_{1/2} = \frac{\log 2}{k}$. Then

$$Q(t) = Q(0) e^{-\frac{\log 2}{t_{1/2}} t} = Q(0) \cdot \left(\frac{1}{2}\right)^{\frac{t}{t_{1/2}}}$$

If the half-life of ^{14}C is $t_{1/2} = 5730$ years, then the quantity of carbon-14 present in a sample after t years is:

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$$Q(t) = Q(0) e^{-\frac{\log 2}{5730} t} = Q(0) \left(\frac{1}{2}\right)^{\frac{t}{5730}}$$

A particular piece of flax parchment contains about 64% as much ^{14}C as flax plants do today. We will estimate the age of the parchment, using 5730 years as the half-life of ^{14}C .

First, a rough estimate: is the parchment older or younger than 5730 years?

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First, a rough estimate: is the parchment older or younger than 5730 years?

Younger: it has *more* than half its ^{14}C left, so it has been decaying for *less* than one half-life.

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Let $Q(t)$ denote the amount of ^{14}C in the parchment t years after it was first created.

$$Q(t) = Q(0) \left(\frac{1}{2} \right)^{\frac{t}{5730}}$$

$$0.64 = \left(\frac{1}{2} \right)^{\frac{t}{5730}}$$

$$\log(0.64) = \frac{t}{5730} \log \frac{1}{2} = -\frac{\log 2}{5730} t$$

$$t = -\frac{5730 \log(0.64)}{\log 2} \approx 3689$$

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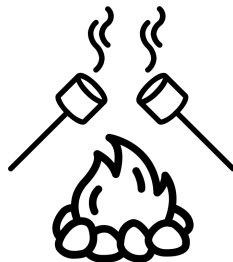
$$t = -\frac{5730 \log(0.64)}{\log 2} \approx 3689$$

So the parchment was made of plants that died about 3700 years ago.

Newton's law of cooling

The **rate of change** of temperature of an object is proportional to the difference in temperature between the object and its surroundings.

The temperature of the surroundings is sometimes called the ambient temperature.

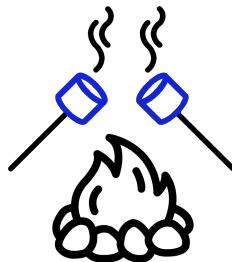


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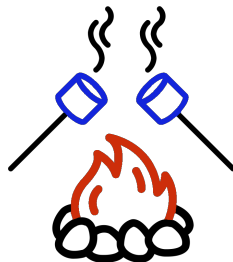


$$\frac{dT}{dt} = K (T(t) - T_a)$$

Newton's law of cooling

The rate of change of temperature of an object is proportional to the difference in temperature between the object and its **surroundings**.

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$$\frac{dT}{dt} = K(T(t) - A)$$

Linear First-Order Differential Equations

Let a and b be constants. The differentiable function $y(x)$ obeys the differential equation

$$\frac{dy}{dx} = a(y - b)$$

if and only if

$$y(x) = \{y(0) - b\} e^{ax} + b$$

Find an explicit formula for functions $T(t)$ solving the differential equation $\frac{dT}{dt} = K(T(t) - A)$ for some constants K and A .

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Find an explicit formula for functions $T(t)$ solving the differential equation $\frac{dT}{dt} = K(T(t) - A)$ for some constants K and A .

$$T(t) = (T(0) - A) e^{Kt} + A$$

The temperature of a glass of iced tea is initially 5° . After 5 minutes, the tea has heated to 10° in a room where the air temperature is 30° . Assume the temperature of the tea as it cools follows Newton's law of cooling,

$$T(t) = (T(0) - A)e^{Kt} + A$$

- (a) Determine the temperature as a function of time.
- (b) When the tea will reach a temperature of 14° ?

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The ambient temperature is $A = 30$ and $T(0) = 5$, so we only have to determine K . (Or, more neatly, e^K .)

$$T(t) = (5 - 30)e^{Kt} + 30$$

$$= 30 - 25e^{Kt}$$

$$10 = T(5) = 30 - 25e^{5K}$$

$$25e^{5K} = 20$$

$$e^{5K} = \frac{4}{5}$$

$$e^K = \left(\frac{4}{5}\right)^{1/5}$$

$$T(t) = 30 - 25 \left(\frac{4}{5}\right)^{\frac{t}{5}}$$

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$$14 = T(t) = 30 - 25 \left(\frac{4}{5} \right)^{\frac{t}{5}}$$

$$\frac{t}{5} = 2$$

$$t = 10$$

$$25 \left(\frac{4}{5} \right)^{\frac{t}{5}} = 16$$

After 10 minutes

$$\left(\frac{4}{5} \right)^{\frac{t}{5}} = \frac{16}{25} = \left(\frac{4}{5} \right)^2$$

A glass of room-temperature water is carried out onto a balcony from an apartment where the temperature is 22°C . After one minute the water has temperature 26°C and after two minutes it has temperature 28°C . Assuming the water warms according to Newton's law of cooling, what is the outdoor temperature?
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$$\begin{aligned}T(t) &= A + (T(0) - A)e^{Kt} \\ &= A + (22 - A)e^{Kt}\end{aligned}$$

$$\begin{aligned}\text{Given: } 26 &= A + (22 - A)e^K \\ \implies e^K &= \frac{26 - A}{22 - A}\end{aligned}$$

$$\begin{aligned}\text{Given: } 28 &= A + (22 - A)e^{2K} \\ \implies e^{2K} &= \frac{28 - A}{22 - A}\end{aligned}$$

$$\begin{aligned}\frac{28 - A}{22 - A} &= (e^K)^2 = \left(\frac{26 - A}{22 - A}\right)^2 \\ (28 - A)(22 - A) &= (26 - A)^2 \\ 28 \cdot 22 - 50A + A^2 &= 26^2 - 52A + A^2 \\ 2A &= 26^2 - 28 \cdot 22 \\ A &= (26)(13) - (22)(14) \\ &= (26)(13) - (22)(13) - 22 \\ &= 4 \cdot 13 - 22 = 30\end{aligned}$$



Let P be the size of a population, and let K be the carrying capacity of its environment (i.e. the population size that can be sustainably supported).

When P is much less than K , our population has...

- A. not enough resources
- B. just enough resources
- C. extra resources

So when the P is much less than K , we expect the population to...

- A. shrink
- B. stay the same
- C. grow

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Malthusian growth

The Malthusian growth model relates population growth to population size:

$$\frac{dP}{dt} = bP(t)$$

where b is a constant representing net birthrate per member of the population.

Let P be the size of a population, and let K be the carrying capacity of its environment (i.e. the population size that can be sustainably supported).

When P is **greater** than K , our population has...

- A. not enough resources
- B. just enough resources
- C. extra resources

So when the P is greater than K , we expect the population to...

- A. shrink
- B. stay the same
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Let P be the size of a population, and let K be the carrying capacity of its environment (i.e. the population size that can be sustainably supported).

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The Malthusian growth model

$$\frac{dP}{dt} = bP(t)$$

is **not** a good model for populations with strained resources

Logistic growth models population growth as:

$$\frac{dP}{dt} = b_0 \left(1 - \frac{P(t)}{K} \right) P(t)$$

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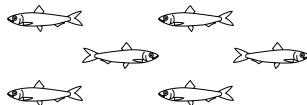


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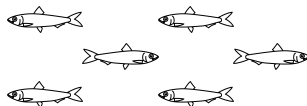
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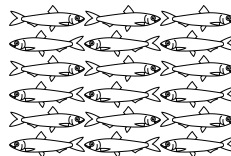
- ▶ If $P \ll K$, then $\frac{dP}{dt} \approx b_0 P(t)$
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- ▶ If $P \ll K$, then $\frac{dP}{dt} \approx b_0 P(t)$
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- ▶ If $P > K$, then $\frac{dP}{dt}$



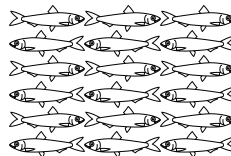
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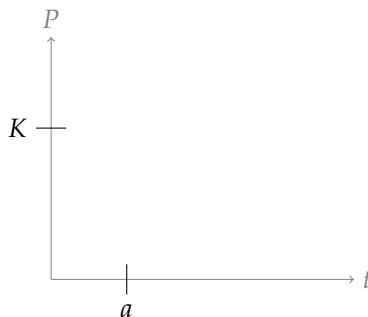
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Before we solve explicitly, let's sketch some solutions to

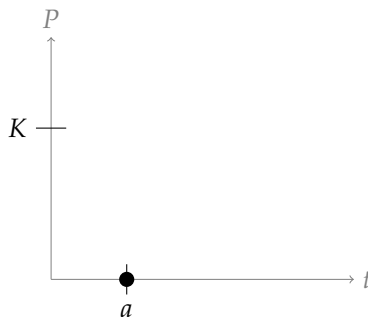
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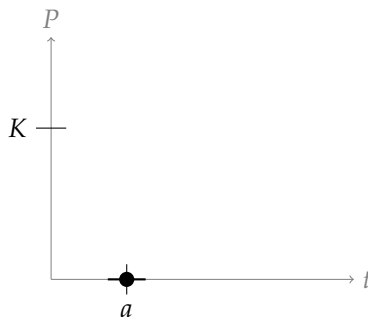
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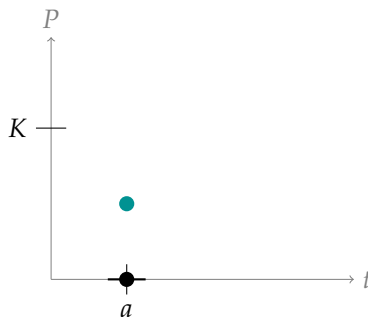
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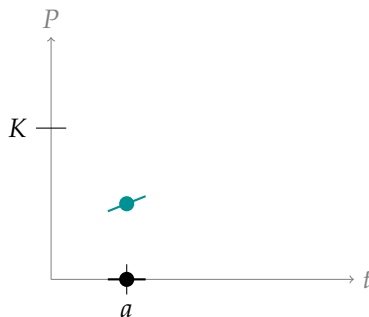
- ▶ If $P(a) = 0$: $\frac{dP}{dt} = 0$
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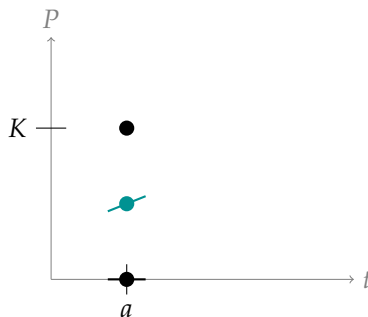
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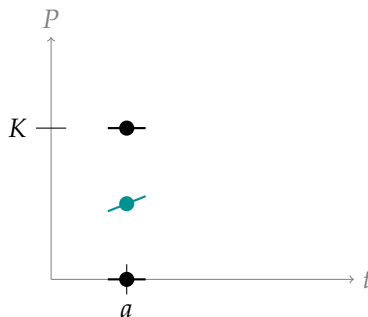
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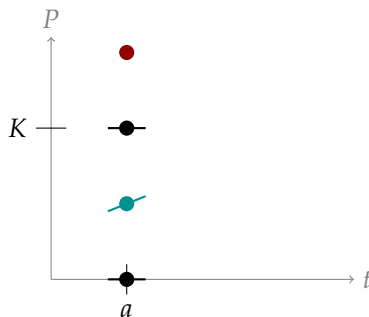
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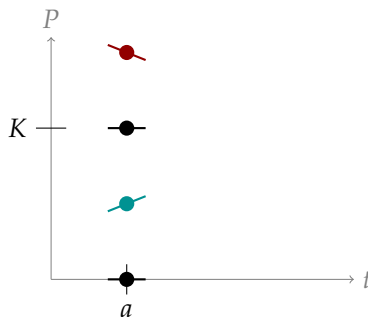
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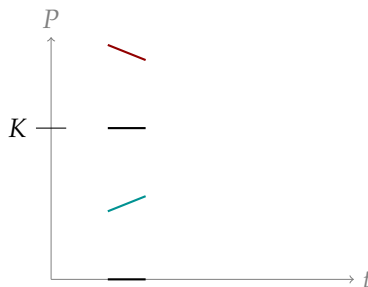
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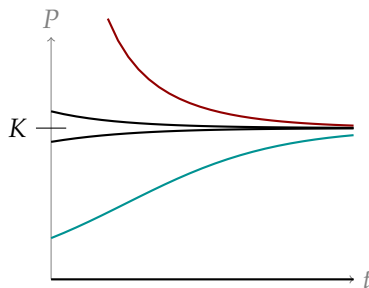
- ▶ If $P(a) = 0$: $\frac{dP}{dt} = 0$
- ▶ If $P(a) = K$: $\frac{dP}{dt}(a) = 0$
- ▶ If $0 < P(a) < K$: $\frac{dP}{dt}(a) > 0$
- ▶ If $K < P(0)$: $\frac{dP}{dt}(a) < 0$



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Find the explicit solutions to

$$\frac{dP}{dt} = b \left(1 - \frac{P(t)}{K} \right) P(t)$$

when b and K are constants.

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$$\int b \, dt = \int \frac{1}{P(1 - \frac{1}{K}P)} \, dP$$

Using partial fractions, we can find $\frac{1}{P(1 - \frac{1}{K}P)} = \frac{1}{P} + \frac{1/K}{1 - \frac{1}{K}P}$

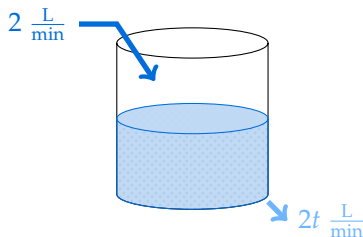
$$\begin{aligned} bt + D &= \int \left(\frac{1}{P} - \frac{1/K}{1 - \frac{1}{K}P} \right) dP \\ &= \log |P| - \log \left| 1 - \frac{1}{K}P \right| = \log \left| \frac{P}{1 - \frac{1}{K}P} \right| \end{aligned}$$

let $C = \pm e^D$:

$$Ce^{bt} = \frac{P}{1 - \frac{1}{K}P} \implies P(t) = \frac{Ce^{bt}}{1 + \frac{C}{K}e^{bt}}$$

At time $t = 0$, where t is measured in minutes, a large tank contains 3 litres of water in which 1 kg of salt is dissolved. Fresh water enters the tank at a rate of 2 litres per minute and the fully mixed solution leaks out of the tank at the varying rate of $2t$ litres per minute.

- (a) Determine the volume of solution $V(t)$ in the tank at time t .
- (b) Determine the amount of salt $Q(t)$ in solution when the amount of water in the tank is at maximum.



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We're given information about the rate of change of V : $\frac{dV}{dt} = 2 - 2t$. Then $V(t) = 2t - t^2 + C$. From the initial value $V(0) = 3$, we see

$$V(t) = 2t - t^2 + 3$$

The maximum value of a downwards-facing parabola occurs at its critical point, so the water in the tank is at its highest level when $t = 1$.

At time $t = 0$, where t is measured in minutes, a large tank contains 3 litres of water in which 1 kg of salt is dissolved. Fresh water enters the tank at a rate of 2 litres per minute and the fully mixed solution leaks out of the tank at the varying rate of $2t$ litres per minute.

- (a) Determine the volume of solution $V(t)$ in the tank at time t .
- (b) Determine the amount of salt $Q(t)$ in solution when the amount of water in the tank is at maximum.

The amount of salt is decreasing as it leaks out. The concentration of salt in the tank water at time t is $\frac{Q(t)}{V(t)} \frac{\text{kg}}{\text{L}}$. If the saltwater is leaking out at a rate of $2t \frac{\text{L}}{\text{min}}$, then:

$$\begin{aligned}\frac{dQ}{dt} &= -\frac{Q(t)}{V(t)} \cdot 2t \frac{\text{kg}}{\text{min}} = \frac{-2t}{2t - t^2 + 3} Q \\ \int \frac{1}{Q} dQ &= \int \frac{2t}{(t+1)(t-3)} dt = \int \left(\frac{1/2}{t+1} + \frac{3/2}{t-3} \right) dt \\ \log |Q| &= \frac{1}{2} \log |t+1| + \frac{3}{2} \log |t-3| + C\end{aligned}$$

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So, we have the following relationship between Q and t :

$$\log |Q| = \frac{1}{2} \log |t + 1| + \frac{3}{2} \log |t - 3| + C$$

To find C , we use the initial condition $Q(0) = 1$.

$$\log |Q(0)| = \frac{1}{2} \log 1 + \frac{3}{2} \log 3 + C$$

$$\implies 0 = C + \frac{3}{2} \log 3$$

$$\implies C = -\frac{3}{2} \log 3$$

At time $t = 0$, where t is measured in minutes, a large tank contains 3 litres of water in which 1 kg of salt is dissolved. Fresh water enters the tank at a rate of 2 litres per minute and the fully mixed solution leaks out of the tank at the varying rate of $2t$ litres per minute.

- (a) Determine the volume of solution $V(t)$ in the tank at time t .
- (b) Determine the amount of salt $Q(t)$ in solution when the amount of water in the tank is at maximum.

Finally, we use our relationship between Q and t to find $Q(1)$.

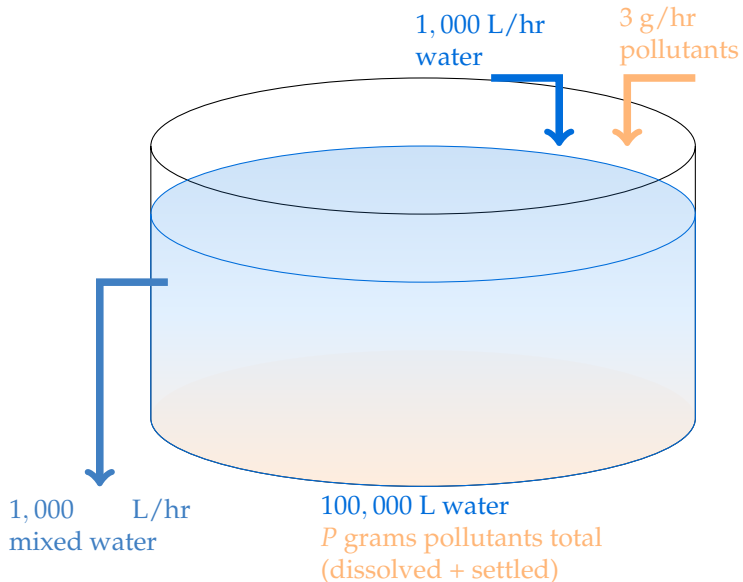
$$\begin{aligned}\log |Q(t)| &= \frac{1}{2} \log |t + 1| + \frac{3}{2} \log |t - 3| - \frac{3}{2} \log 3 \\ \Rightarrow \log |Q(1)| &= \frac{1}{2} \log 2 + \frac{3}{2} \log 2 - \frac{3}{2} \log 3 = 2 \log 2 - \frac{3}{2} \log 3 \\ &= \log 4 - \log \left(3^{3/2} \right) = \log \frac{4}{3^{3/2}} \\ \Rightarrow Q(1) &= \frac{4}{3^{3/2}}\end{aligned}$$

SETTLING TANK

A settling tank is filled with 100,000 litres of pure water. Every hour, 1,000 litres of water, containing 3 grams of pollutants, enters the tank.

90% of the pollutants in the settling tank sink to the bottom, with the remaining 10% well-mixed into the water. The tank drains 1,000 litres of this mixed water into the sewer every hour.

In order to drain the water into the local sewer, the concentration of pollutants cannot be more than 1 gram per 1,000 litres. How long can the settling tank take dirty water until the process must be stopped?



Let $P(t)$ be the total amount (in grams) of pollutants in the tank. Pollutants are entering at a rate of 3 grams per hour. How fast are they leaving?

So, the quantity of pollutants in the tank satisfies the differential equation:

Let $P(t)$ be the total amount (in grams) of pollutants in the tank. Pollutants are entering at a rate of 3 grams per hour. How fast are they leaving?

Every hour, the tank drains 1,000 of its 100,000 litres. That is, every hour, it drains $\frac{1}{100}$ of its total volume. So, every hour, it disgorges $\frac{1}{100}$ of its *dissolved* pollutants. The amount of dissolved pollutants in the tank is $\frac{1}{10}P(t)$. So, the rate the tank leaks pollutants is

$$\frac{1}{100} \cdot \frac{1}{10}P = \frac{1}{1,000}P$$

So, the quantity of pollutants in the tank satisfies the differential equation:

$$\frac{dP}{dt} = (\text{rate in}) - (\text{rate out}) = 3 - \frac{1}{1000}P$$

$$\frac{dP}{dt} = 3 - \frac{1}{1000}P$$

is a linear first-order differential equation, so we know its solution:

$$P(t) = Ce^{-t/1000} + 3000$$

We also know $P(0) = 0$:

$$0 = C + 3000$$

$$P(t) = 3000 \left[1 - e^{-t/1000} \right]$$

If the concentration of **dissolved** pollutants in the tank is 1 gram per 1,000 litres, that means:

- ▶ in the whole 100,000 litre tank, there are 100 grams **dissolved**, so
- ▶ in the whole tank, $\frac{1}{10}P = 100$, so $P = 1000$ grams.

To find when that happens, we solve $P(t) = 1000$ for t .

$$1000 = 3000 \left[1 - e^{-t/1000} \right]$$

$$\frac{1}{3} = 1 - e^{-t/1000}$$

$$e^{-t/1000} = \frac{2}{3}$$

$$-\frac{t}{1000} = \log \left(\frac{2}{3} \right)$$

$$t = -1000 \log \left(\frac{2}{3} \right) \approx 405$$

So the tank can be used for about 405 hours (24 days)

You deposit $\$P$ in a bank account at time $t = 0$, and the account pays $r\%$ interest per year, compounded n times per year. Your balance at time t is $B(t)$.



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If one interest payment comes at time t , then the next interest payment will be made at time $t + \frac{1}{n}$ and will be:

$$\frac{1}{n} \times \frac{r}{100} \times B(t) = \frac{r}{100n} B(t)$$

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So, calling $\frac{1}{n} = h$,

$$B(t+h) = B(t) + \frac{r}{100} B(t)h \quad \text{or} \quad \frac{B(t+h) - B(t)}{h} = \frac{r}{100} B(t)$$

If the interest is compounded continuously,

$$\begin{aligned} \frac{dB}{dt}(t) &= \lim_{h \rightarrow 0} \frac{B(t+h) - B(t)}{h} = \frac{r}{100} B(t) \\ \implies B(t) &= B(0) \cdot e^{rt/100} = P \cdot e^{rt/100} \end{aligned}$$

Continuously compounding interest

If an account with balance $B(t)$ pays a continuously compounding rate of $r\%$ per year, then:

$$\frac{dB}{dt} = \frac{r}{100}B$$
$$B(t) = B(0) \cdot e^{rt/100}$$

You invest \$200 000 into an account with continuously compounded interest of 5% annually. You want to withdraw from the account continuously at a rate of $\$W$ per year, for the next 20 years. How big can W be?

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Let $A(t)$ be the balance in the account t years after the initial deposit.

$$\frac{dA}{dt} = \frac{5}{100}A - W = \frac{1}{20}(A - 20W)$$

$$A(t) = (200\,000 - 20W)e^{t/20} + 20W$$


$$\begin{aligned} 0 = A(20) &= (200\,000 - 20W)e + 20W \\ &= 200\,000e + 20W(1 - e) \end{aligned}$$


$$W = \frac{200\,000e}{20(e - 1)} = 10\,000 \frac{e}{e - 1} \approx 15\,819.77$$

That is, you can withdraw $10\,000 \frac{e}{e-1} \approx 15\,819.77$ each year.


Included Work


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