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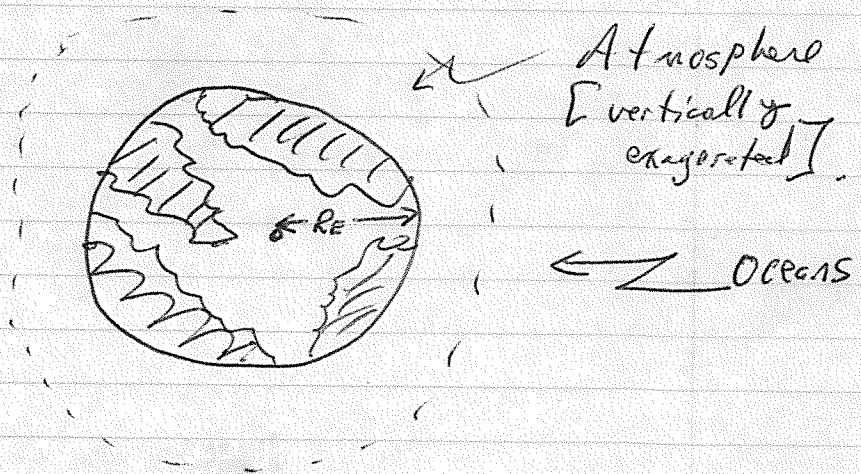
Example Problem

Pg 1

Problem Statement:

Given  $\text{CO}_2$  forcing is about  $3 \text{ W/m}^2$ ; how long would it take to raise the temperature of the atmosphere by  $1^\circ\text{C}$ ? IF this power was instead dropped into the ocean, how long would it take to raise ocean temperature by  $1^\circ\text{C}$ ?

Note:  $3 \text{ W/m}^2$  is the "extra" radiative forcing caused by anthropogenic  $\text{CO}_2$  emitted to the atmosphere since the industrial revolution.

Solution:

Let's treat the problem in a general way.

Q heat in in Joules  $\Rightarrow$   $\begin{matrix} C & T \\ & M \end{matrix}$  object

"Specific heat"

$C = \frac{\text{Heat capacity}}{\text{mass}} \left[ \frac{\text{J}}{\text{K kg}} \right]$

$M = \text{Object mass}$

$\Delta T = \text{temperature change}$

$$Q = \cancel{\Delta T} = MC\Delta T$$

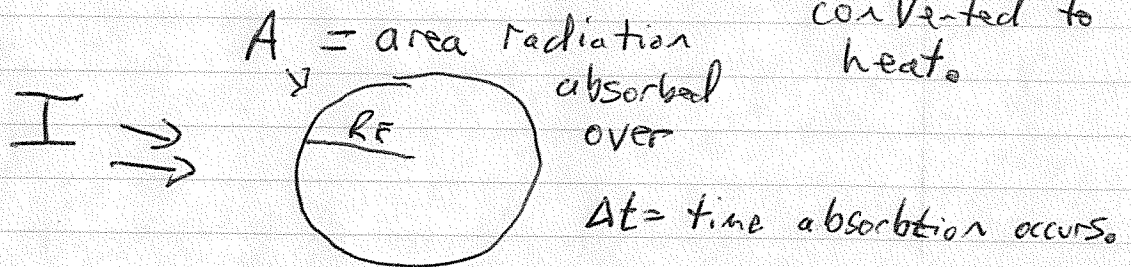
(1)

$$\boxed{\Delta T = Q / MC}$$

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We need to convert radiative forcing to  $Q$ .

Let  $I \equiv 3 \frac{W}{m^2}$  = irradiance to be converted to heat.



(2) Then  $Q = I \left[ \frac{W}{m^2} \right] * A [m^2] * \Delta t$

Since  $W = \frac{J}{s}$ .

Now all we need to do is to find the mass of the atmosphere and ocean and the area  $A$ .

Combining (1) and (2) we can get the time needed for the temperature change by 1 K.

$$\Delta T = \frac{I A \Delta t}{m C} \quad \text{So the}$$

(3) time needed is

$$\Delta t = \frac{m C \Delta T}{I A}$$

A high heat capacity  $C$  and mass  $M$  makes for a long time to increase the temperature by 1 C.

Aside: Heat capacity per unit mass for air and water.

Pg 3

Air is composed of  $N_2$  and  $O_2$ , mostly.



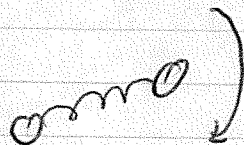
From the equipartition theorem of physics, the energy of air is, for  $N$  molecules,

(4)

$$E = N n \frac{1}{2} kT$$

$n =$  "# degrees of freedom, energy pathways for a molecule"

$\frac{1}{2} kT$  energy for every mode of vibration or rotation or translation.



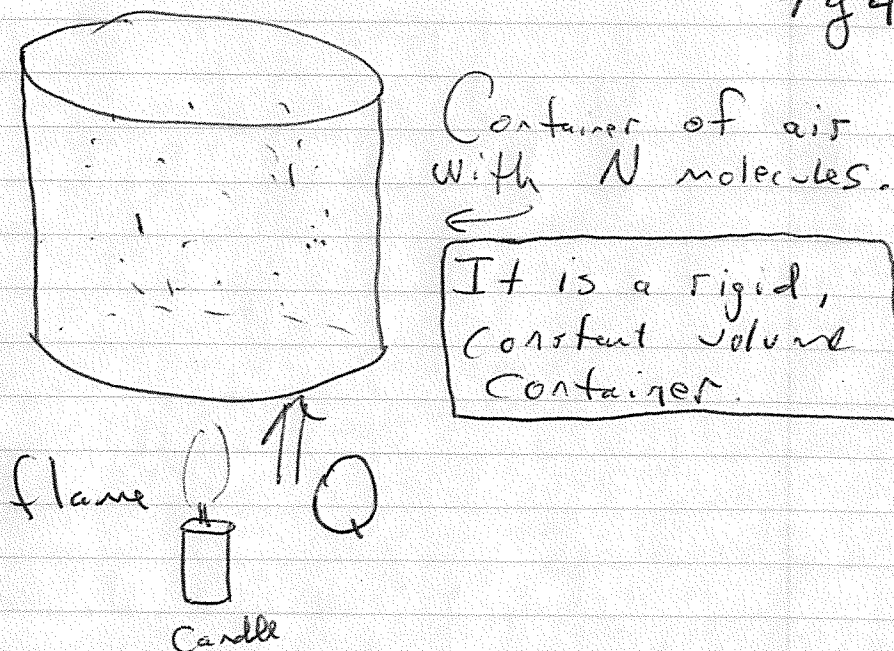
Can move in 3 directions,

$n = 5$  for  $N_2$  and  $O_2$  molecules

Can rotate in 2 directions.

Vibration is "frozen out" at atmospheric temperatures.

In (4),  $k = 1.38 \times 10^{-23} \frac{J}{\text{molecule } ^\circ K}$  is Boltzmann's constant.

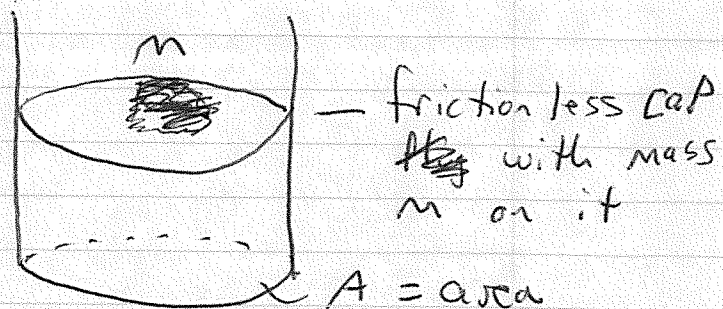


When we add  $Q$  heat into our air sample the energy change will be

(5)

$$\Delta E = Q = \frac{5}{2} N k_B T.$$

That's fine, but the atmosphere is not a sealed container. It is more like this



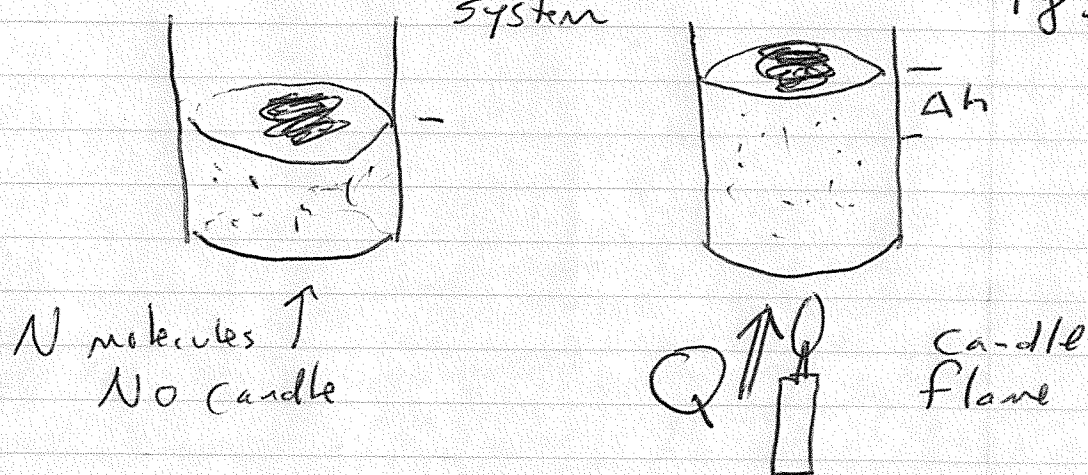
When we apply the candle, the cap can move up.

(6)

The pressure in the chamber is  $\frac{mg}{A} = P$

Constant Pressure system

Pg 5



The candle both causes the molecules to move around faster, and the cap to move up. The increase in potential energy of the cap is

(7)

$$P.E. = mg \Delta h.$$

This increase in P.E. is provided by the action of the faster moving molecules pushing the cap up; they were made to move faster by the heat provided by the candle flame. We can play with  $E(7)$  to get

(8)

$$P.E. = \frac{mg}{A} A \Delta h$$

Pressure  
in the  
gas  
(constant)

$\Delta V =$  volume  
change of the  
gas



The effect of  $Q$  is

(9)

$$Q = \frac{5}{2} N k \Delta T + P \Delta V$$

internal energy of the gas molecules has gone up, but some of  $Q$  goes into the work done by the gas,  $P \Delta V$ , to move the cap up.

In the constant volume system on page 4 and Eq (5), all of  $Q$  goes into changing the temperature, while in the constant pressure system of page 5 and Eq 9, some of  $Q$  goes into doing work, changing the gas volume, so the temperature change in the constant pressure system is less than in the constant volume system.

Our ideal gas air equation is

(10)

$PV = NkT$  so if we hold pressure constant, and ask, how much do we need to change the temperature of the gas by to ~~increase~~ do the work  $P \Delta V$  in Eq (9), we find out quickly that

(11)

$$P \Delta V = Nk \Delta T.$$

So for the same amount of heat  $Q$  from the candle, we will have a different temperature change for our constant volume and constant pressure systems.

$$(9) \quad Q = \frac{7}{2} Nk \Delta T \quad \text{constant pressure}$$

$$(5) \quad Q = \frac{5}{2} Nk \Delta T \quad \text{constant volume}$$

We define :

$$(12) \quad C_p = \frac{7}{2} Nk$$

$$(13) \quad C_v = \frac{5}{2} Nk$$

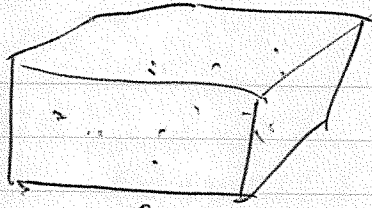
as the heat capacity for constant pressure and constant volume processes.

Then

$$(9) \quad Q = C_p \Delta T \quad \text{constant pressure}$$

$$(5) \quad Q = C_v \Delta T \quad \text{constant volume}$$

Next we want to work of the specific heat capacity.  $C_p = C_p / \text{mass}$ , the heat capacity per unit mass of air.



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← Box of air with  $N$  molecules.

$M =$  mass of air in box. what is the mass?

$$M = N \text{ molecules} \times \underbrace{\mu}_{\text{Average molecular weight of air}} \frac{\text{kg}}{\text{mole}} \times \underbrace{N_A}_{\text{Avogadro's \#}} \frac{\text{mole}}{\text{molecule}}$$

Rearranging,

$$(14) \quad N = \frac{M N_A}{\mu} \text{ molecules}$$

Using (14) in (12) and (13),  $R^*$

$$C_p = \frac{7}{2} N k = \frac{7}{2} \frac{M N_A}{\mu} k$$

$$(15) \quad C_p = \frac{7}{2} \frac{M}{\mu} R^* = \frac{7}{2} m \frac{R_{\text{air}}^*}{\mu}$$

We used a universal gas constant definition,

$$R^* = N_A k$$

and

$$R_{\text{air}} = R^* / \mu$$

We get  $C_p = \frac{C_p}{M} = \frac{7}{2} R_{\text{air}}$



The value of  $R_{air}$  is

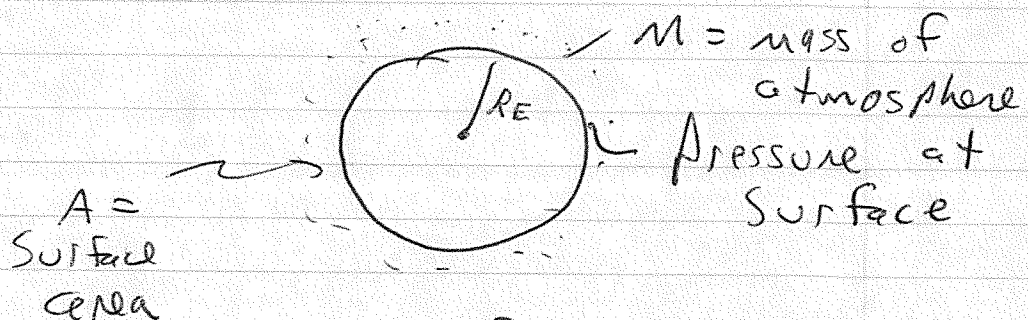
$$R_{air} = \frac{8.3143 \text{ J}}{\text{mole K}} \cdot \frac{1000 \text{ mole}}{29 \text{ Kg}}$$

$$(16) \quad R_{air} = 286.7 \frac{\text{Joules}}{\text{Kg} \cdot \text{K}}$$

Now we return to our problem, calculating  $\Delta t$  in Eq 3, the time it takes for the atmosphere to increase temperature by 1 K due to absorption of radiation  $I_0$

$$(3) \quad \Delta t = \frac{M}{A} \frac{C_p \Delta T}{I}$$

Referring back to Eq (6), the surface pressure on the planet is



$$(6) \quad P = \frac{mg}{A} \quad \text{So}$$

$$(17) \quad \Delta t = \frac{P}{g} \frac{C_p \Delta T}{I}$$

For the atmosphere then,

Pg 10

$$\Delta t = \frac{10^5 \text{ Pa}}{9.8 \text{ m/s}^2} \cdot \frac{7}{2} \cdot \frac{286.7 \text{ J}}{\text{kg K}} \frac{\Delta T}{1 \text{ K}}$$

$$3 \text{ W/m}^2$$

$$\Delta t = 3.4 \times 10^6 \text{ Seconds}$$

$$\Delta t = 39.5 \text{ days} \quad \text{Atmosphere.}$$

Given about 40 days and an excess radiative forcing of  $3 \text{ W/m}^2$ , the atmosphere will warm up, on average, by  $1^\circ \text{C}$ .

For the oceans, we use

$$M = 1.38 \times 10^{21} \text{ Kg} \quad (\text{mass of oceans})$$

$$A = 4\pi R_e^2 = 4\pi \times (6.37 \times 10^6 \text{ m})^2$$

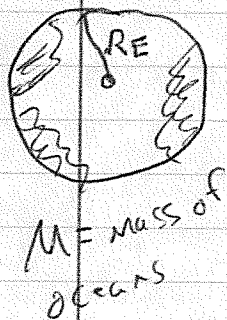
$$A = 5.1 \times 10^{14} \text{ m}^2$$

$$C = \frac{1 \text{ calorie}}{\text{gram}^\circ \text{K}} = \frac{4180 \text{ J}}{\text{kg K}}$$

$$\Delta t = \frac{M}{A} \frac{C \Delta T}{I} = \frac{1.38 \times 10^{21}}{5.1 \times 10^{14}} \times \frac{4180 \text{ J/K}}{3}$$

$$\Delta t = 3.8 \times 10^9 \text{ seconds} \\ = 43,636 \text{ days}$$

$$\Delta t = 120 \text{ years}$$



In summary, the atmosphere takes about 40 days to warm by 1 K due to excess radiation absorbed of  $3 \frac{W}{m^2}$

While the oceans take about 120 years to warm by 1 K.

Roughly speaking, the atmosphere responds quickly to radiative imbalances, though oceans take much longer; about a factor of

$$\frac{43636 \text{ days ocean}}{40 \text{ dys atmosphere}}$$

~ factor of 1000 times as long for ocean to heat up as atmosphere.

The bad news is, eventually the ocean does warm up too.

Fun  
thoughts  
in  
closing

For air,  $C_p = \frac{7}{2} R_{\text{air}} = 1004 \frac{J}{kg K}$

For water  $C = 4180 \frac{J}{kg K}$

Per mole,  
Air

$$C_p = 1004 \frac{J}{kg K} \times \frac{29 \text{ kg}}{1000 \text{ mole}} \sim 29 \frac{J}{\text{mole K}}$$

$$C = 4180 \frac{J}{kg K} \times \frac{18 \text{ kg}}{1000 \text{ mole}} = 75 \frac{J}{\text{mole K}}$$

The heat capacity of water  $\approx 2.6$   
<sub>air</sub>  
 So <sup>Liquid</sup> water molecules have about 2.6  
 equivalent degrees of freedom  
 more than the  $N_2$  and  $O_2$  molecules  
 of air.

For water vapor in the atmosphere  
 $n = 6$  (3 rotation + 3 translation)

degrees of freedom so

$$C_p^{\text{water vapor}} = \frac{5}{2} R \frac{\text{J}}{\text{mole}^\circ\text{K}} \approx 33 \frac{\text{J}}{\text{mole}^\circ\text{K}}$$

which is considerably less than  
 the heat capacity of water molecules in  
<sub>mole</sub>  
 the liquid state.