

Environmental Quality: Monitoring and Analysis
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Lecture – 51
Air-Water Exchange

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Tutorial

1. Emission factor from different chosen source.
2. Estimation of Mixing length.
3. Dispersion $\rightarrow \rho_g(x, y, z, t) = \frac{Q}{2\pi\sigma_y\sigma_z} \exp\left(-\frac{y^2}{2\sigma_y^2} - \frac{z^2}{2\sigma_z^2}\right)$

$\rho_g(x, y, z, t)$ σ_y σ_z

Tutorial is to get emission factor. So, there are different categories I have already given in that. So, you can pick any 3 and go to the website that we talked about, pick the emission factor for any 3 different pollutants and then for those sources we do dispersion problem exercise. So, in the dispersion problem, I think before the dispersion problem, I have one exercise on estimation of mixing length. So, here I think mixing length you have to determine graphically, for example you have to determine what is the temperature at the ground T at $z = 0$ for the environment and T at $z = 0$ for the pollutant.

So, the environmental environment temperature and there is a corresponding environmental lapse rate and for the pollutant whatever temperature is there that is adiabatic lapse rate. So, depending on so in the problem itself, I think you have to decide which is the environment starting point, lapse rate starting point and which is the adiabatic lapse rate starting point, that I think we discussed in class. So, based on whatever the discussion we had, you can recall that, if you if you don't recall this, just go back to the notes and read it, otherwise sent me an email and I will help you at that time.

So, this is a calculation of the mixing length estimate. So, sometimes if the mixing length is infinite, then you say it is infinity, but in this case it will be a finite intersection somewhere. So these will intersect somewhere and that will be mixing height. So you can do it either graphically or arithmetically either way by equations and this is a dry adiabatic lapse rate and this is the environmental lapse rate, the mixing height based on that. The next is a dispersion exercise. So, you start with the full form of the Gaussian dispersion equation.

So, the full form that we have is x , y , z , and H . we have this full equation with reflection. This example we did in class, you start with that and then reduce it. So, one of the things that I have asked in the problem is for a given set of conditions and for a certain stability class, so you have to estimate the stability class for this particular problem, you have to calculate the what we are asking is ρA_1 at some height, I think height I'm asking is z equals to 1 or 2 or 2 meters or some such thing as a function of x , which means that you have to calculate.

You have to calculate ρA_1 x , y , z , H , so along the center line which means you have to calculate at 0, 2 meters and whatever is the height for different x values, which means you have to run this model for different x , running for x , so the dispersion coefficient will change with x , okay. So, you can do it graphically and then you will get you will get some relationship like this. Typically, you will see something like this yeah. The reason you do this, you see this is, what happens is if there is a stack is very high, the plume is going like this, you will not see anything at ground level for some distance.

Once you once the plume reaches the ground is when you start seeing some concentration. So, at this site nothing seen here, then it starts increasing. So, I want you to plot that. So, that is one exercise. This is a manual exercise. When you are doing AERMOD, this is done automatically by the software. Software will do 4000 points at one shot within 1 second. You can set it up like that to do in Excel also provided you know equations for σ_y and σ_z . There are equations, mathematical equations.

It is a little painful but if you know for that stability class, you can take the points from the graph and fix a curve and then use that curve. So automatically you do for one, do for all of them okay. I think you can figure it out how to do it automatically, so okay. So that I would like you to try this on your own and then after that if you have doubts, questions, how to do it,

you are unable to get any of these things. So the emission factor that you get here you, I do not remember if the dispersion problem has a link to this first problem.

The first problem, the emission factor is meant to be used in AERMOD. So, whatever you do here, you will do it and use that source in AERMOD, one of the sources in AERMOD, but you are free to choose this, but I think I have given you some number here for the q . This is q by some equation right $ux \sigma_y \sigma_z$ exponential whatever. So, this q is I think given, I have given this to you in the problem, I do not remember, you check that okay. So, I would like you to try this on your own, sometime next week we can discuss this.

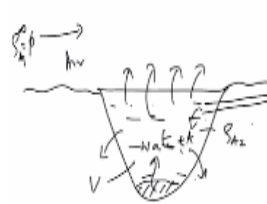
So, before the quiz, you have enough time to revise. The quiz will be in class next to next Wednesday, again this 30th, either in 355 or 356, one of those rooms okay, most likely in 355. We will assemble there and we will this discuss that, decide that day and previous day I will tell you.

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Mass Transfer across interface

Exchange between Air-Water

Case 1 \rightarrow Evaporation of a pollutant from a lake.



(i) What happens to A in the lake. \rightarrow Volume of lake is the system.

$$\text{Rate of Accumulation of } A = \text{Rate in} - \text{Rate out (Evaporation)}$$

$$V \frac{dC_A}{dt} = Q_1 C_{A1} - Q_2 C_{A2}$$

Assume Rate out \Rightarrow Evaporation.

Rate of change of A in lake = Rate of A entering lake via disposal - Rate of A exiting lake via Evap.

$$V \frac{dC_A}{dt} = Q_1 C_{A1} - ()$$

(Ma)

So I want to continue our discussion. This is an application of what we were doing. So, so far the recap, what we did in mass transfer at the interface. We first take the exchange between air and water. So, this is a simplest case for which we even derived the interfacial expression air and water. So, the conditions for air and water for example this is a box model example we started with originally when we started doing the modeling, the description of the model. So, we are looking at exchange between for a simple case, so

case 1:

We look at evaporation of a pollutant from a lake okay. So, there is a lake and there is air. There are some volume and we are talking about this thing okay. So, this problem can be posed differently. One the one way in which it can be posed is let us say that this background concentration of A in the air ρ_{A1} infinite is 0, which means there is no chemical A in the background air and so there is a chemical. This scenario can be that somebody has spilled a lot of chemical here A and this A is entering the water from the sediment or somebody has dumped a lot of chemical into the lake and this lake is contaminated and so on, different scenarios. So, the simplest scenario here is there is a concentration of A in the water and so what are the implications of this? The questions that you can ask are the following. The first one, first possibility is what happens to A in the lake? First question. Is the is the concentration of A in the lake, how is it changing? So, this problem if you want to solve, we can write the mass balance, rate of accumulation.

So, automatically when we ask this question in the lake our system becomes a lake, the lake becomes a system. So, this volume of the lake is the system. So, for this system, rate of accumulation of A in the system equals rate in minus rate out. So, how can A come in, rate of A coming in is let us say somebody is constantly dumping stuff every day continuously, there is a pipeline coming in and disposing material here okay. So let us say there is a pipeline coming in and disposing stuff here.

So you have Q into some concentration of ρ_{A2in} , let us call it as in minus rate out, what is this rate out? So in this case, several processes, possibilities we can discuss. One rate out could be it is going into the sediment, we have not yet considered that, let us for the time being ignore that part say it as 1. So, we assume rate out is evaporation only, which means this is evaporation, rate out by evaporation only. So, you have to write this this statement in English first.

This is very important because then from each term here, so this conversion of the statement will be

$$\begin{aligned} & \text{rate of change of A in the lake} \\ &= \text{rate of A entering lake via disposal} \\ &- \text{rate of A exiting lake via evaporation} \end{aligned}$$

rate of change of A in the lake is a result of rate of A entering lake via disposal minus rate of A exiting the lake via evaporation.

Student: Sir what if already some person has dumped it just for once, yeah, like one time. After that he did not, nothing had happened. Then, how do we consider rate in that situation.

Professor: Yeah, so that there will be no rate in, it will already be there, your initial condition already be will exist.

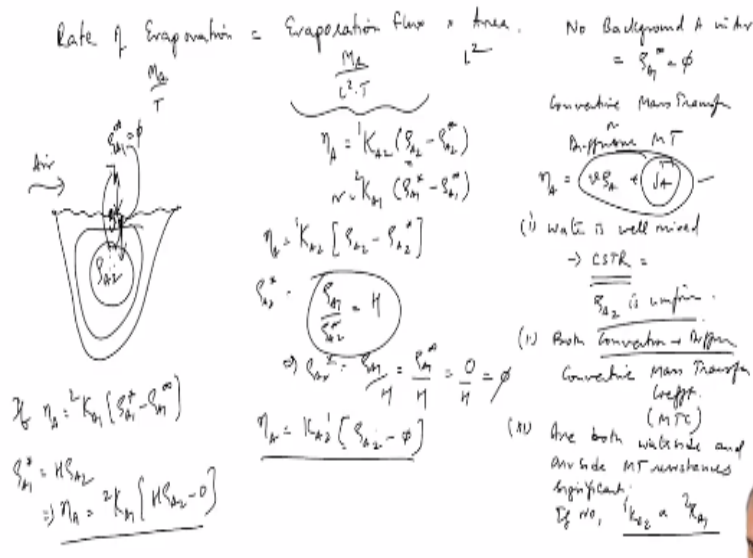
The initial condition exists, even here that initial condition exist. So, I am going to remove one of these terms now. So, we are going to be assuming this first. There is no rate of entering, which means somebody has already dumped it okay. So which means that there is an initial mass that is sitting there already and that we have estimated it somehow, either by measurement or by estimation that you know that 4400 kilograms of some chemical has been dumped in.

So we have to link it back to our first problem that we did in this course, that is mass balance, a 400 kilograms of dumped in what is the concentration at equilibrium in the water, the worst case scenario. That is the highest concentration of A in the water. That is your starting point. From there now, that problem was an equilibrium problem. So now if that is happening, then after that, you take that as starting point and emission starts from there, the release start from there. So, we will take one by one.

This this problem can be as complicated as you want and then corresponding to that the equation will change and mathematics form will change. So so rate of change is so now we are saying that rate of change is equal to rate of evaporation okay. So you see a negative sign there, this is zero, nothing is entering. So, because somebody has dumped already everything and it is a finite volume of dumping and nothing is entering, exiting. So, this becomes the rate of A exiting the lake. So, rate of change of this thing is $\frac{d}{dt} \rho A_2$ into V of the lake of the water.

So, every term in the mass balance has units of mass of A by T. So, therefore, $V \frac{d\rho A}{dt}$. It is the rate of change of liquids that is in unsteady state system, nothing is coming in okay. So, if somebody is putting material into it. So if this is second term, this term is not 0 there will be there will be a term here, so we will have it as ρA in constantly minus rate of evaporation, rate of evaporation will be what here, this is the term now we have been discussing in the last few classes, to fill up this term okay, the rate of evaporation term.

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So, that evaporation is flux into area. So evaporation flux has units of MA by L square into T , area is L square, MA by T . Now, so this is evaporation flux. The evaporation flux is nA okay. This nA has to be now computed. So, now we look at the system. There is water, there is air and there is a concentration here. Now it is going out from here to here. Concentration here is ρ_{A2} infinity which we will assume to be 0 for the time being, no background concentration. When you say no background A in air, we are essentially saying this ρ_{A1} , sorry this is ρ_{A1} infinity is 0.

This is the assumption we are making. Now what do we write this term here? Now, first of all, you have to make this judgment whether it is a convective mass transfer or diffusive mass transfer. We have

$$n_A = v\rho_A + j_A,$$

we make a decision right here, which one do you consider? If you are considering only j_A , how will you write the equation? If you are considering both, how will you write? So what is the likelihood of this in this system, the information given to you have it is not sometimes enough for you to, so you have to make some assumptions here, ok.

So if no information is given, you can do problems in multiple ways. So let us make it easy on you for the time being. Let us say that the lake is very shallow and it is well mixed, one assumption. The water is well mixed. What will this immediately do, what is well mix means? The concentration wherever in this region, everywhere the concentration is the same. So, well

mixed essentially will make this into what is called as a CSTR, completely stirred tank reactor, which means that ρ_{A2} is uniform throughout the volume.

Which also allows you to do this on the left hand side accumulation term, you can write it like that because it is uniform, otherwise you cannot write it like that. You have to integrate that thing over the entire volume, otherwise you cannot write v into $d\rho_{A2} dt$ cannot be written okay. The water is well mixed. Air is anyway moving. So, what is well mix, there is convection in the water and there is convection in the air. So, now we are assuming both and we will say both convection plus diffusion.

When we have conduction and diffusion as a condition, then we use a convective mass transfer coefficient, we use a convective, we give term by MTC, mass transfer coefficient write as MTC, I will write MTC henceforth okay. Now, we have decided that which means there is a k mass transfer coefficient term multiplied by a driving force that needs to come here. So we will start with the driving force. So, yesterday's last class, we had discussed how we can write this flux term, you have various options to doing it.

So, in this particular scenario, we are taking ρ_{A2} which is a solution in the aqueous phase going into the vapor phase in the air. So many resistances are there. This resistance are both water side and air side mass transfer resistances significant, which means can you neglect any of them? By this point of time, you have no reason to neglect either of them? You do not have any information that can tell you that neglecting of it because it is not pure chemical, you cannot neglect one of the phases.

You have to consider the solution here and it is going across. So, there is an interface resistance here and so both. The answer is no, both k_{A21} and k_{A12} are to be considered, okay, both are to be considered. Then, this term here will become, you can have two choices, you either have,

$$\begin{aligned} n_A &= {}^1K_{A2}(\rho_{A2} - \rho_{A2}^*) \\ &= {}^2K_{A1}(\rho_{A1}^* - \rho_{A1}^\infty) \end{aligned}$$

either way. So if you do this, we will do k_{A21} , when you do this, we do k_{A12} okay. Is it clear? Anybody has a question on this how we do this? So we further write this.

Let us take the first one, K_{A2} because we already know ρ_{A2} . We need to do this in ρ_{A2} , why? The other term that is there sitting in the accumulation, we can do it either way. So, let us take this first, how we can do this, ρ_{A2}^* . What is ρ_{A2}^* ? ρ_{A2} is connected to ρ_{A1} . This is the equilibrium concentration, an equivalent concentration of ρ_{A2} that is in equilibrium with whatever is there in the air. What is there in the air? Zero. So it is ρ_{A1} by ρ_{A2}^* equals Henry's constant, which means

$$\frac{\rho_{A1}}{\rho_{A2}^*} = H$$

$$\rho_{A2}^* = \frac{\rho_{A1}}{H}$$

In this case, this ρ_{A1} is ρ_{A1}^∞ by Henry's constant. ρ_{A1}^∞ is 0, 0 by H is 0.

$$\rho_{A2}^* = \frac{\rho_{A1}}{H} = \frac{\rho_{A1}^\infty}{H} = \frac{0}{H} = 0$$

Therefore, this equation becomes

$$n_A = {}^1K_{A2}(\rho_{A2} - 0)$$

This is capital KA. Alternatively, you write the other term. If you write KA, if you take

$$n_A = {}^2K_{A1}(\rho_{A1}^* - \rho_{A1}^\infty)$$

$$\rho_{A1}^* = H * \rho_{A2}$$

based on this. In this case it is ρ_{A1}^* divided by $\rho_{A2} = H$, we are now taking the equivalent concentration of A in the air which is in equilibrium with whatever of A is there in the water and using the water concentration to find an equivalent equilibrium concentration in air.

So what we are doing is we are taking this entire thing and finding an imaginary concentration here that is in equilibrium with this mass, this concentration and so we are calculating this one. In the other case, we did we take this concentration and find out an equivalent concentration here and we are taking this section okay. So which means that

$$n_A = {}^2K_{A1}(H * \rho_{A2} - 0)$$

When you calculate, these 2 expressions are different, so eventually become they should give you the same value if you compute either way okay.