

**Introduction to Process Modeling in Membrane Separation Process**  
**Prof. Sirshendu De**  
**Department of Chemical Engineering**  
**Indian Institute of Technology, Kharagpur**

**Lecture - 09**  
**Osmotic Pressure Controlling Filtration (Contd.)**

Welcome to the session. We will be now we are looking into the some simplified versions of osmotic pressure control filtration. We have looked into the first simplified case there is the no concentration polarization and we have looked into the system how the system can behave and prediction can be obtained.

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(ii) Low polarization:  $\frac{v_w}{K} \ll 1$   $e^{\frac{v_w}{K}} = 1 + \frac{v_w}{K} + \frac{(\frac{v_w}{K})^2}{2} + \dots$   
 $= 1 + \frac{v_w}{K}$

Film Theory:  $\frac{c_m - c_p}{c_0 - c_p} = e^{v_w/K} = 1 + \frac{v_w}{K}$

$c_p = c_m (1 - R_r)$

$v_w = L_p [\Delta P - \Delta \pi]$

$v_w = L_p \left[ \Delta P - b R_r c_0 \frac{(1 + v_w/K)}{R_r + (1 - R_r)(1 + v_w/K)} \right]$

$\Rightarrow$  quadratic in  $v_w$ ;

The second simplified case will be talking about is low polarization and low polarization can be realized by exponential  $v_w$  by  $K$  is going to be realized when  $v_w$  by  $K$  is much lesser than 1 that means, again in this case turbulence is quite high. So, Reynolds number will be quite high very high and therefore, the mass transfer co efficient will be high compared to the permeate flux. So,  $v_w$  by  $K$  will be much less than 1 and that will allow us to have an exponential expansion of  $e$  to the power  $v_w$  by  $K$  as 1 plus  $v_w$  by  $K$  plus  $v_w$  by kernel square of that 1 over factorial 2 here plus other higher order terms. We

neglect the since  $V_w$  by  $K$  itself is much less than 1  $V_w$  by  $k$  square on the higher terms will be neglected and this can be approximated as  $1 + V_w$  by  $K$ .

Now what we will be doing next? We will be writing the film theory the film theory. If you remembered this is  $C_m$  minus  $C_p$  divided by  $C$  not minus  $C_p$  e to the power  $V_w$  by  $K$  and this will be replaced by  $1 + V_w$  by  $K$ .

So, these a simplified fraction of the film theory and then will be writing  $C_p$  as  $C_m$  into 1 minus real retention and then will be writing the osmotic pressure model as  $V_w$  is equal to  $L_p$   $\Delta \pi$  minus  $\Delta \pi_i$  and we have seen earlier how  $\Delta \pi_i$  can be replaced in terms of real retention and membrane surface concentration  $C_m$ . Now combining these 3 equation will be getting a simplified version as  $V_w$  is equal to  $L_p$   $\Delta \pi$  minus  $b R_r C$  not  $1 + V_w$  by  $K$  divided by  $R_r$  plus  $1 - R_r$  into  $1 + V_w$  by  $K$ . Now in this case what will be ultimately getting? So, ultimately you will be getting a quadratic in  $V_w$ . So, all the other quantities are known to us  $L_p$  is membrane permeability  $\Delta \pi$  is the trans membrane pressure drop  $b$  is the osmotic co efficient known to us  $R_r$  is real retention known to us  $C$  not is the feed concentration  $K$  is the mass transfer co efficient that will be calculate and we have seen how to calculate the mass transfer co efficient.

So, we can calculate the  $V_w$  and from the  $V_w$  one can estimate. So, one can directly get the expression of  $C_m$  because  $C_m$  minus  $C_p$  divided by  $C$  not minus  $C_p$  can be expressed. So,  $V_w$  can be calculated independently. Let us see now how  $C_m$  will be calculated;  $C_p$  will be calculated.

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Handwritten derivation on a whiteboard:

$$\frac{C_m - C_p}{C_0 - C_p} = 1 + \frac{V_w}{K}$$

$$\Rightarrow \frac{C_m R_r}{C_0 - C_m(1 - R_r)} = 1 + \frac{V_w}{K} \quad R_r = 1 - \frac{C_p}{C_0}$$

$$\Rightarrow C_m R_r = (1 + \frac{V_w}{K}) C_0 - (1 - R_r)(1 + \frac{V_w}{K}) C_m$$

$$\Rightarrow C_m [R_r + (1 - R_r)(1 + \frac{V_w}{K})] = (1 + \frac{V_w}{K}) C_0$$

$$\Rightarrow C_m = \frac{(1 + \frac{V_w}{K}) C_0}{R_r + (1 - R_r)(1 + \frac{V_w}{K})}$$

Final boxed equation:

$$C_p = C_m (1 - R_r) \quad \checkmark$$

So, if you look into the osmotic expression film theory  $C_m$  minus  $C_p$  divided by  $C_0$  not minus  $C_p$  is equal to one plus  $V_w$  by  $K$ . We have already got an explicit expression in quadratic form of  $V_w$ . So, therefore, these  $C_m$  minus  $C_p$  can be replaced in terms of  $C_m$ . So, this will be nothing, but  $C_m$  times  $R_r$  because 1 of the definition of real retention 1 minus  $C_p$  by  $C_m$ . So,  $C_m$  minus  $C_p$  is nothing, but  $C_m$  time  $R_r$  and  $C_p$  will be the denominator  $C_p$  can be replaced  $C_m$  into 1 minus  $R_r$  is equal to 1 plus  $V_w$  by  $K$ .

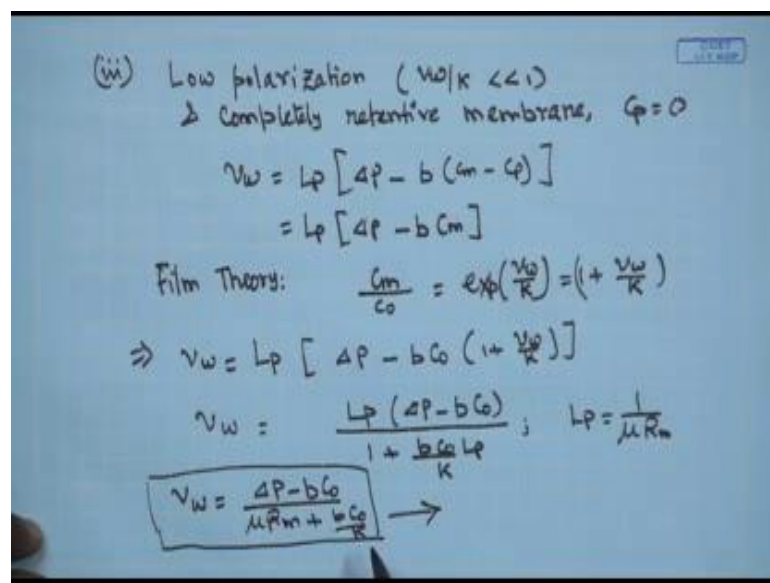
So, just rearrange  $C_m$ . So, this become  $C_m R_r$  is equal to 1 plus  $V_w$  by  $K$  times  $C_0$  not minus 1 minus  $R_r$  1 plus  $V_w$  by  $K$  times  $C_m$ . So,  $C_m$  will be  $R_r$  plus 1 minus  $R_r$  1 plus  $V_w$  by  $K$  and here it will be 1 plus  $V_w$  by  $K$  times  $C_0$  not or one can estimate the value of  $C_m$  as 1 plus  $V_w$  by  $K$  time  $C_0$  not divided by the denominator  $R_r$  plus 1 minus  $R_r$  into 1 plus  $V_w$  by  $K$ . So,  $V_w$  you have estimated already as a quadratic in the only few minutes back and  $K$  is known to as  $C_0$  not is known to us  $R_r$  is known to us and one can estimate the value of  $C_m$ . Once the  $C_m$  is estimated then permeate concentration can be estimated as well through their definition of real retention.

So, we can have an independent system prediction in terms of permeate concentration and permeate flux for a low polarization case. Where in the polarization is not low whether 1 it is significant we have already seen that we have to solve 2 equations and 2

unknown 3 equations in 3 unknown system by using Newton Raphson after combining all the equations we will be getting will be landing of with a single non-linear algebraic equation which will be solved iteratively by using the Newton Raphson method.

So, we will be doing another simplification, that the things will be physically quite significant and apparent to us.

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(ii) Low polarization ( $v_w/K \ll 1$ )  
 & completely retentive membrane,  $C_p = 0$

$$v_w = L_p [\Delta P - b(C_m - C_p)]$$

$$= L_p [\Delta P - bC_m]$$

Film Theory:  $\frac{C_m}{C_0} = \exp\left(\frac{v_w}{K}\right) = \left(1 + \frac{v_w}{K}\right)$

$$\Rightarrow v_w = L_p \left[ \Delta P - bC_0 \left(1 + \frac{v_w}{K}\right) \right]$$

$$v_w = \frac{L_p (\Delta P - bC_0)}{1 + \frac{bC_0 L_p}{K}} ; L_p = \frac{1}{\mu R_m}$$

$$v_w = \frac{\Delta P - bC_0}{\mu R_m + \frac{bC_0}{K}} \rightarrow$$

The third simplified case will be the low polarization and completely retentive membrane. Low polarization that is  $V_w$  by  $K$  is much much less than 1 and completely retentive membrane; that means,  $C_p$  is equal to 0. So, in this case permeate flux will be  $L_p \Delta P$  minus  $bC_m$ ,  $bC_m$  will be  $bC_0$  minus  $C_p$  and  $C_p$  will be equal to 0. So, you will be having  $L_p \Delta P$  minus  $bC_0$  and from the film theory what you will be getting? For film theory you will be getting  $C_m$  by  $C_0$  not because all  $C_p$ s are 0 it is a completely retentive membrane exponential  $V_w$  by  $K$ . So, it will be 1 plus  $V_w$  by  $K$ .

So, you will be getting permeate flux is equal to  $L_p \Delta P$  minus  $bC_0$  not 1 plus  $V_w$  by  $K$  now if you bring  $V_w$  to the other side and write everything in terms of that. So, you will be getting  $L_p \Delta P$  minus  $bC_0$  not divided by 1 plus  $bC_0$  not into  $L_p$  divided by mass transfer coefficient  $K$  and now if I replace  $L_p$  the membrane permeability in terms of

membrane hydraulic resistance  $R_m$  then I will be getting a concrete expression of permeate flux as  $\Delta P - \pi C$  is equal to  $\mu R_m + \pi C$  not divided by  $K$ .

So, these expressions clearly show that, this is the effective driving force. This is the applied trans membrane pressure drop minus osmotic pressure divided by there will be 2 resistances in series one is membrane hydraulic resistance other is the resistance due to the mass transfer in the mass transfer boundary layer. So, two resistance in series that will be appearing in the denominator and in the numerator you will be having the effective trans membrane pressure drop which will be nothing, but the actual trans membrane pressure drop minus the osmotic pressure that is developed in to the system. So, this clearly shows that mass transfer resistance and membrane resistance they will be acting in series in an actual system. So, before going into the other variance of solution diffusion model I would like to now in at this point of time I would like to include or introduce velocity variation technique or one more technique.

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Velocity variation Technique to estimate  $R_r$  of membrane-solute-solvent system

Film Theory:

$$\frac{C_m - C_p}{C_o - C_p} = \exp\left(-\frac{v_w}{K}\right) \leftarrow$$

$$R_r = 1 - \frac{C_p}{C_m} \Rightarrow \frac{C_p}{C_m} = 1 - R_r$$

$$\frac{C_m}{C_p} = \frac{1}{1 - R_r}$$

$$\Rightarrow \boxed{C_o = \frac{C_p}{1 - R_r}}$$

$$R_o = 1 - \frac{C_p}{C_o}$$

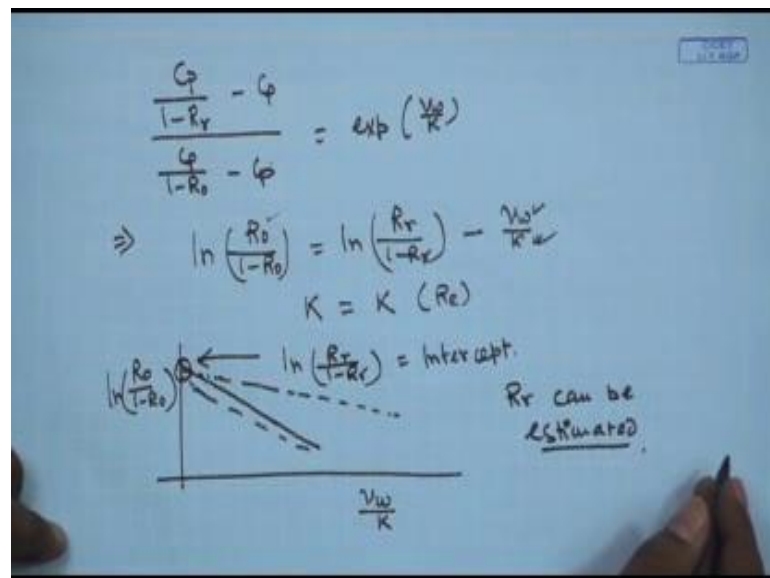
$$\Rightarrow C_o = \frac{C_p}{1 - R_o}$$

If you remember in an earlier class we have said that there are two ways or two experimental methods to estimate the real retention of the system. Now will be really looking into how to we have seen 1 method already how to conduct a separate set of experiments. So, that the real retention of the system can be estimated by under low

polarization condition, that is low operating pressure low feed concentration high turbulence. Now in this will be looking into 1 more method to estimate the velocity variation technique which will be utilizing the film theory model to estimate the real retention of the membrane; technique to estimate  $R_r$  of membrane solute solvent system.

So, we take request to the film theory model and it is equation. So,  $C_m$  minus  $C_p$  divided by  $C$  not minus  $C_p$  is equal to exponential  $V_w$  by  $K$ . Next what we will be doing will be writing the definition of real retention and observe retention. So, real retention is  $1$  minus  $C_p$  by  $C_m$ . So,  $C_m$  can be written as.  $C_p$  by  $C_m$  will be  $1$  minus  $R_r$   $C_m$  by  $C_p$  will be  $1$  over  $1$  minus  $R_r$ . So, therefore,  $C_m$  is nothing, but  $C_p$  divided by  $1$  minus  $R_r$ . Similarly from the definition of observe retention  $R_{not}$  is  $1$  minus  $C_p$  by  $C$  not and we can estimate the value of  $C$  not in terms of  $C_p$ . So,  $C$  not will be nothing, but  $C_p$  divided by  $1$  minus  $R_{not}$ . So, now, we are going to replace the expression of the  $C_m$  in favor of  $C_p$  and  $R_r$  and  $C$  not in favors of  $C_p$  and  $R_{not}$  in this equation and see what we get.

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So, if you really do that you will be getting  $C_p$   $1$  minus  $C_p$  divided  $1$  minus  $R_r$  minus  $C_p$  is equal to  $C_p$  divided by  $1$  minus  $R_{not}$  minus  $C_p$  exponential  $V_w$  by  $K$ . So, this can be. So,  $C_p$  will be canceling out from both the numerator and denominator and you will be getting an expression of observe retention and real retention and permeate flux and mass

transfer coefficient. If you simplify this equation I am just writing the final form  $\ln R_{not}$  divided by  $1 - R_{not}$  is equal to  $\ln R_r$  by  $1 - R_r$  minus  $V_w$  by  $k$ .

Now in this expression if we have the real retention as constant and let us see how if you change your operating conditions operating conditions means here we cannot change the operating conditions in terms of pressure, because the mass transfer coefficient is appearing here and as you have already seen that mass transfer coefficient is a strong function of Reynolds number and Reynolds number will be basically will be a strong function of the cross flow velocity.

So, this method is known as velocity variation technique. What will be doing? Will be applied different cross flow velocity in the system that will alter the mass transfer coefficient; once the mass transfer coefficient will be altered the permeate flux will be altered and once the permeate flux will be altered as well as it will be affecting the permeate quality. So, observe retention because permeate concentration also be altered. So, observe retention will be altered. So, that is therefore, we change the cross flow velocity and get different type different values of permeate mass transfer coefficient and permeate flux and then we plot  $\ln R_{not}$  by  $1 - R_{not}$  versus  $V_w$  by  $K$ .

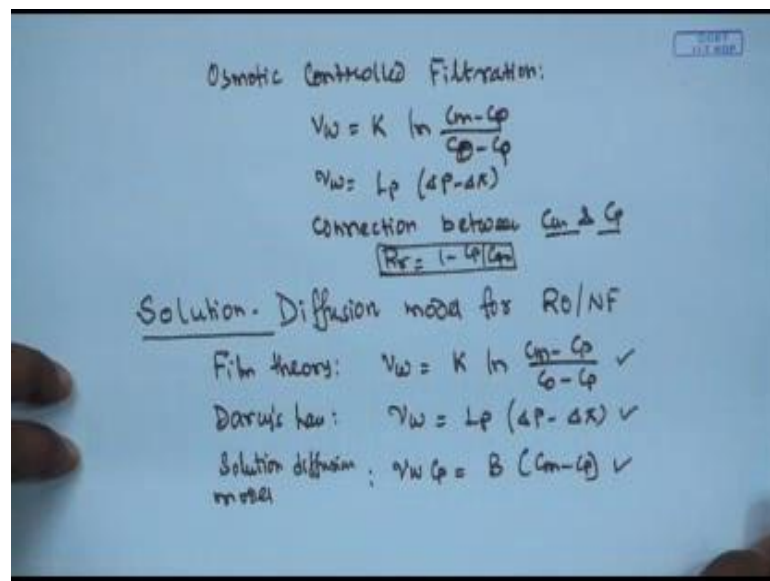
So, now, if a plot  $\ln R_{not}$  divided by  $1 - R_{not}$  versus  $V_w$  by  $K$  you will be getting a curve something like this. So, similarly actually these will be hitting the y axis the curve will be something like this and similarly it will be for the other values of mass transfer coefficient will be getting this. So, ultimately all these points will be going to the intercept  $R_{not}$  by  $1 - R_{not}$ . So, this is for a particular  $K$  this is that is for another particular  $K$ .

So, you will be getting from the intercept. What is the intercept? The intercept is  $\ln R_r$  by  $1 - R_r$ . So, from the intercept one can estimate the value of real retention. So,  $R_r$  can be estimated. So, what is the drawback of this system? The major drawback of this system is that, the experiments should be very very accurate. If there is slight in accuracy in the experiments then in the log scale these will be even magnified and therefore, it will giving a very wrong estimation. So, there will be number of more number of experiment one has to conduct not only that not only the more number of experiments the

experiments has to be very very accurate then only 1 can get an estimation of real retention appropriate estimation of real retention by this velocity variation method, OK.

Next, we will be looking into as you have discussed that the third equation let us look into once again the modeling equation for the osmotic pressure control filtration one dimensional model.

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So, we have at the film theory  $V_w$  is equal to  $K \ln C_m$  minus  $C$  not  $C_p$  minus  $C$  not we have the Darcy's law  $V_w$  is equal to  $L_p \Delta P$  minus  $\Delta \pi$  and we have 2 equations 2 unknown  $C_m$   $C_p$   $V_w$  these are the 3 unknown and 2 equations though other equation is the connection between the  $C_m$  and  $C_p$ .  $C_m$  and  $C_p$  and we have already seen that 1 way of connecting  $C_m$  and  $C_p$  is the real retention that is nothing, but a partition coefficient between across the membrane of the solute partition coefficient across the membrane. Now we have already seen the all the formulations and the method for solving these third equation to get a system prediction of the system performance. Now we are what will be doing? Will be replacing if the expression of real retention that will be valid more valid for the ultra filtration system and various cut off ultra filtration system by the solution diffusion model for the for the reverse osmosis nano filtration system.



So, real retention is a concept which will be more valid for the ultra filtration system, but for the reverse osmosis and nano filtration will not beyond be able to use the expression of real retention as partition coefficient  $1 - \frac{C_p}{C_m}$ . You have to use the other variance of the connection between the  $C_p$  and  $C_m$  that is the solution diffusion model and it is various variations or modifications. So, first will be talking about the solution diffusion model for reverse osmosis and nano filtration system; for RO-NF system, now, let us look in to the 1 of the various governing equations. So, 1 equation will be the film theory is  $V_w$  is equal to  $k \ln \frac{C_m}{C_p}$ . So, is not  $C_p$   $C$  not minus  $C_p$   $C_m$  minus  $C_p$  divided by  $C$  not minus  $C_p$  the other 1 is the Darcy's law which is nothing, but the solvent flux through the membrane.  $V_w$  is equal to  $L_p \Delta P - \Delta \pi$  and the third 1 is the solute flux through the membrane, which is the solution diffusion model.

So, these will be giving you the solvent flux through the mass transfer boundary layer, solvent flux to the porous membrane and the solute flux to the porous membrane  $V_w C_p$  is equal to  $b (C_m - C_p)$ . So, we will be having 3 equations and 3 unknowns  $V_w$   $C_m$  and  $C_p$  and can get the system prediction. So, I will just give a small algebraic manipulation in order to solve this set of equations.

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Handwritten mathematical derivation on a blue background:

$$V_w = L_p (\Delta P - \Delta \pi) \quad R = \alpha C$$

$$V_w = L_p \Delta P \left[ 1 - \frac{\alpha}{\Delta P} (C_m - C_p) \right]$$

$$V_w = V_w^* [1 - \alpha (C_m - C_p)] \quad \dots (1)$$

$V_w^* = L_p \Delta P \rightarrow$  Pure water flux

$$\alpha = \frac{R}{\Delta P} v$$

$$V_w^* [1 - \alpha (C_m - C_p)] = K \ln \left( \frac{C_m - C_p}{C_0 - C_p} \right) \dots (2)$$

Solution-Diffusion  $\Rightarrow V_w C_p = B (C_m - C_p) \dots (3)$   
 Combine Eq. (3) and Eq. (1)

So,  $V_w$  can be the solvent flux to the membrane can be written as  $V_w$  is equal to  $L_p \Delta P - \Delta \pi$ . So, this will be, if you take  $L_p$  out of  $\Delta P$ . So, it will be giving  $1 - \alpha$   $\Delta \pi$  and  $\Delta \pi$  is let us say  $\pi$  is equal to  $\alpha C_m - C_p$ . So,  $\Delta \pi$  will be basically  $\alpha \Delta P C_m - C_p$ . So, I write  $V_w$  is equal to  $V_w \text{ not } 1 - \alpha C_m - C_p$ . So, this is my equation number 1 and what is  $V_w \text{ not}$ ?  $V_w \text{ not}$  is basically  $L_p \Delta P$  it is a pure water flux. Pure water flux and what is the parameter  $\alpha$ ? So, these will be  $V_w \text{ not}$  will be known to us because  $L_p$  is known to us and  $\Delta P$  is the trans membrane pressure drop. So, what is  $\alpha$ ?  $\alpha$  is the parameter  $\alpha$  by  $\Delta P$ .

So, it is non dimensional. So, there will be equation number 1 then will be equating these with the film theory equation. So,  $V_w \text{ not } 1 - \alpha C_m - C_p$  will be equal to  $K \ln \frac{C_m - C_p}{C \text{ not } - C_p}$ . So, these will be the combination of the film theory and the osmotic pressure model Darcy's law. So, I write it as; this is an equation number 2, then we will be having the solution diffusion model for the solute flux through membrane. That will be  $V_w C_p$  is equal to  $b C_m - C_p$ . So, now, we can combine these two equations and finally, will be getting into these. So, we will be combining the equation number 3 solution diffusion model and Darcy's law that is equation 1 you can combine these 2 equations and see what we will get? If you combine these two equations you will be getting  $V_w \text{ not } 1 - \alpha C_m - C_p$  is equal to  $b C_m - C_p$  divided by  $C_p$ .

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$$V_w^0 [1 - \alpha (C_m - C_p)] = B \left( \frac{C_m - C_p}{C_p} \right)$$

$$\Rightarrow 1 - \alpha C_m + \alpha C_p = \beta \frac{C_m - C_p}{C_p}$$

$$\beta = B / V_w^0$$

$$C_m = C_p \left[ 1 + \frac{1}{\beta + \alpha C_p} \right]$$

Eq. (2)

$$\frac{\beta V_w^0}{\alpha C_p + \beta} - K \ln \left[ \frac{C_p}{(\alpha C_p + \beta)(C_p - C_p)} \right] = 0$$

$$f(C_p) = 0 \Rightarrow \text{Iterative Solution of } C_p$$

$\leftarrow$  N-R method

So, these equations can be written in this form 1 minus alpha times  $C_m$  plus alpha times  $C_p$  is equal to beta  $C_m$  minus  $C_p$  divided by  $C_p$ . Where alpha we have already defined and beta is  $b$  by  $V_w$  not. So, from these equations the membrane surface concentration is obtained in terms of permeate concentration. So, from these we can get  $C_m$  membrane surface concentration as function of permeate concentration  $C_p$  into 1 plus 1 plus beta plus alpha  $C_p$ .

Now, this expression of  $C_m$  can be put into the equation that we have already talked about. Equation number 2. We are going to put the value of  $C_m$  that we are obtained in the in these equation - equation number 2 if you really do that put into equation number 2 you will be getting 1 non-linear algebraic equation in the form of I will be giving the final expression beta  $V_w$  not alpha  $C_p$  plus beta minus  $K \ln C_p$  divided by alpha  $C_p$  plus beta into  $C$  not minus  $C_p$  is equal to 0. Again these expression is nothing, but a non-linear algebraic equation in  $C_p$  and again 1 has to take care go by a Newton-Raphson method iteratively. So, one has to have an iterative solution. iterative solution of  $C_p$  and the method can be used as Newton Raphson method. once you get the value of  $C_p$  you will be getting the value of  $C_m$   $C_p$  is equal will be getting because from this equation you will be getting the value of  $C_m$  because  $C_p$  you have already estimated. once you

know the value of  $C_p$  and  $C_m$  then 1 can go through the film theory equation and can get the expression of permeate flux.

So, again in this method also, we can land up with 3 equations only difference is the definition of real retention which basically acts as a partition co-efficient between the membrane surface concentration solute concentration in the membrane surface and in the permeate will be replaced by the solution diffusion equation which is basically nothing, but the solute transport equation for the solute flux through the membrane. Then again these 3 equations can be solved and can be recast in the form of single equation which will be algebraic non-linear equation and that can be solved by using an iterative method. By that 1 can get the permeate concentration membrane surface concentration and permeate flux ultimately.

So, I will stopping in this class in the next class next session will be looking into the other variance of solution diffusion model and how to use these solution diffusion model, what there will be the there will be a modifications of we need in order to get the system prediction perfectly.

Thank you very much.