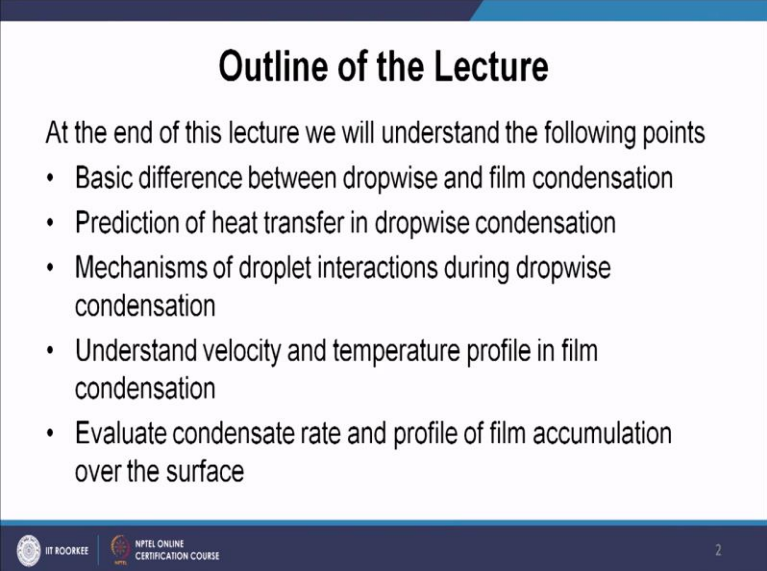


Two phase flow and heat transfer
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Lecture No: 18
Condensation

Hello, welcome in the eighteenth lecture of Two Phase Flow and Heat Transfer. Here, today we will be discussing about condensation heat transfer. So, let us see at the end of this lecture you will be understanding basic difference between drop wise and film condensation. We will be predicting the heat transfer coefficient in drop wise condensation. We will be discussing elaborately the mechanisms of droplet interactions during drop wise condensation.

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Outline of the Lecture

At the end of this lecture we will understand the following points

- Basic difference between dropwise and film condensation
- Prediction of heat transfer in dropwise condensation
- Mechanisms of droplet interactions during dropwise condensation
- Understand velocity and temperature profile in film condensation
- Evaluate condensate rate and profile of film accumulation over the surface

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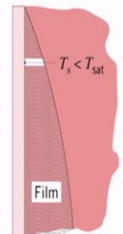
We will be finding out velocity and temperature profile when film condensation is happening over a vertical surface. We will be also seeing how to evaluate condensate rate and the profile of the film okay. Over the vertical surface, so let us start with the definition of condensation. So, condensation is 1 mode of heat transfer in which actually over the surface we will find out due to heat transfer gas will be converting into liquid okay. So here you will find out the surface will be actually having lower temperature than the saturation temperature okay.

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Condensation:

Heat transfer to a surface occurs by condensation when the surface temperature is less than the saturation temperature of an adjoining vapor.

- **Film Condensation**
 - ☐ Entire surface is covered by the condensate, which flows continuously from the surface and provides a resistance to heat transfer between the vapor and the surface.
 - ☐ Thermal resistance is reduced through use of short vertical surfaces and horizontal cylinders.
 - ☐ Characteristic of clean, uncontaminated surfaces



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Now if we try to classify broadly, this condensation will be finding out 2 modes are there. One is called film wise condensation another one is called drop wise condensation. So first let us try to understand what is film condensation? So in film condensation majority of this type of film condensation we see over vertical surface. So here I have given you 1 schematic of the vertical surface. So let us say this is the surface, whose temperature is lower than the saturation temperature of a given pressure okay.

Now as this temperature surface temperature $T_s < T_{\text{sat}}$ then you will be finding out that a liquid film is being formed okay due to the condensation. Condensation means, it will be forming liquid from vapor. So, you will find out that this liquid is forming a film over this vertical surface okay. So in case of film condensation, you will find out that entire surface is actually covered by condensate. It is not like that locally the surface is covered by the liquid, it is entirely covered which flows continuously from the surface and provides a resistance to the heat transfer between the vapor and the surface.

Because this film will be acting as some sort of resistance between you know vapor to come in contact with the cold surface. So you will be finding out this is giving some sort of resistance. So this thermal resistance okay is reduced okay through the use of short vertical surfaces and


horizontal cylinders. If you want to know, remove this film boiling. So what we need to do on the surface, we need to give some sort of horizontal cylinders okay as some sort of appendage.

So that the film will be breaking down and you can once again go back to your drop wise condensation mode, where gas can come direct contact with your surface okay. And this type of film boiling you can see in the surface is where you will finding out surface is clean and you are having no contamination okay. Then let us see another version. So drop wise condensation so you see in drop wise condensation, here I have given you 1 figure in drop wise condensation.

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• Dropwise Condensation


- ❑ Surface is covered by drops ranging from a few micrometers to agglomerations visible to the naked eye.
- ❑ Thermal resistance is greatly reduced due to absence of a continuous film
- ❑ Surface coatings may be applied to inhibit *wetting* and stimulate dropwise condensation





Steam condensation on copper surfaces:

$$q = \bar{h}_{dc} A (T_{sat} - T_s)$$

$\bar{h}_{dc} = 51,100 + 2044 T'_{sat}$	$22^\circ\text{C} < T'_{sat} < 100^\circ\text{C}$
$\bar{h}_{dc} = 255,500$	$T_{sat} > 100^\circ\text{C}$





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You can find out over the surface you are having lots of droplets of liquid being formed okay. Now this drop wise condensation majority of the time you will be finding out over a horizontal surface okay. But here I have given you 1 example, this to give you some sort of comparison between the previous 1 and this 1 that we are having a vertical surface once again. And you can find out lots of droplets are actually being formed due to condensation. So gas is converting into liquid.

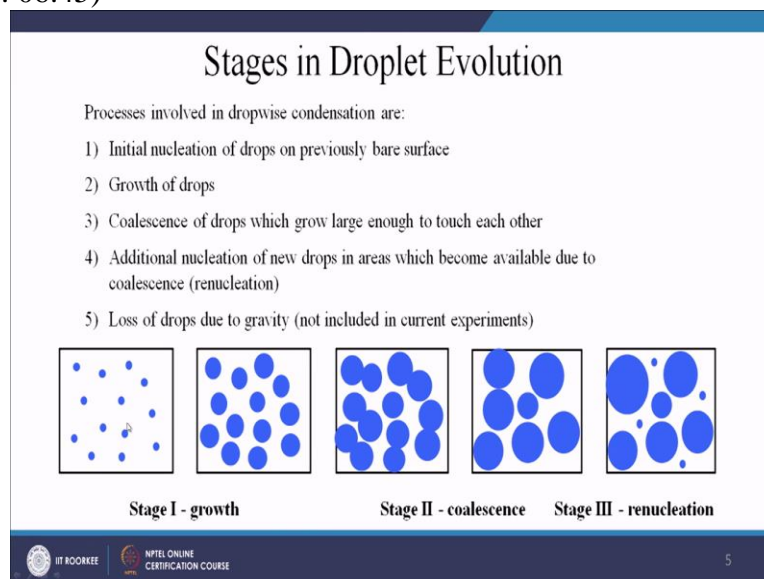
Now in this type of drop wise condensation you will be finding out that surface is covered by drops and the size of drop will be ranging from a wide variety okay. So it will be starting from few micro meters okay. And you know agglomerations of this drops will be making this drop visible even with a naked eyes okay. Here I have shown you 1 typical example. Now this thermal resistance is greatly reduced due to this absence of continuous film whatever we have seen in the

previous case. In case of film condensation and surface coating, you can apply okay to inhibit, to wetting and stimulate the drop wise condensation.

If the surface is having very poor wetting condition that means it is asking the drop to squeeze on the surface then you will be finding out the drop wise condensation is being preferred. Now let us see that how in case of drop wise condensation heat transfer can be calculated. So here I have shown you a situation where condensations, drop wise condensation is being predicted for steam over copper surface. So that can be written as q equals to heat transfer coefficient for drop wise condensation. $h_{dc} \cdot a (T_{sat} - T_s)$ okay.

Now this h_{dc} is very important heat transfer coefficient in drop wise condensation. It depends on the surface temperature. So if saturation temperature is in between you know 22 degree centigrade to 100 degree centigrade, we can write down the coefficient as $51,000, 100 + 2044 \cdot T_{sat}$ and if the saturation temperature is higher than 100 degree then you can write down that is actually 255,500 a constant value okay. Once you know the heat transfer coefficient once again you can go back to this equation to get the drop wise condensation heat flux okay.

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Next let us see that how these droplets in drop wise condensation will be behaving over a surface. So difference stages of droplet evolution. So you will be finding out at the beginning the droplets will be nucleating on the surface at this time the droplet will be having very smaller size in the range of a microns and then you will be finding out the initially those droplets will be

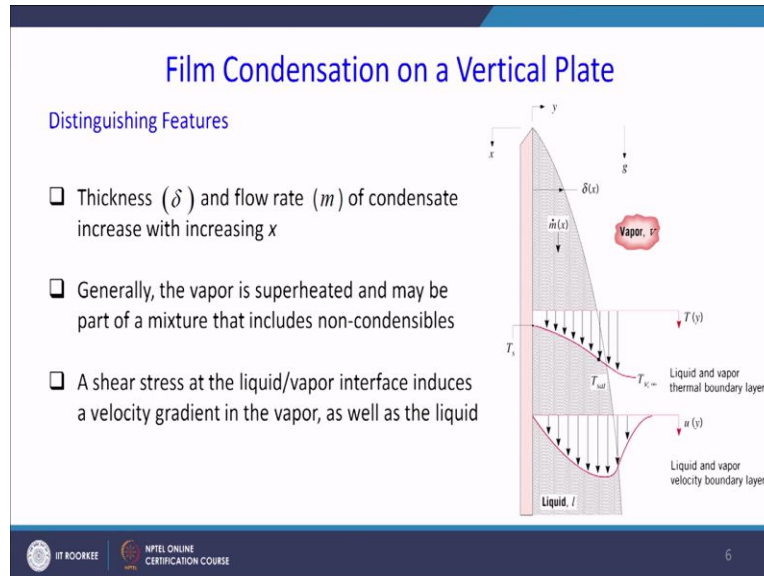
growing in size due to more and more condensation around that. So it is actually increasing its size due to the condensation for that side okay.

And then after some time you will find out whenever this droplet size will be becoming considerably, we will be finding out 2 droplets side by side which are staying side by side they are actually merging or coalescences among themselves okay. So you will be finding out coalescences happens like this. Here you see the coalescences happen and the drop gets bigger and bigger in size, once again so bigger size droplets you can find out okay.

But at the same time if it is on a vertical surface then you will be finding out with the increase of size the droplet is also facing the gravitational pool. So you can find out at some time droplet is also being dislodged from the surface okay. So you will be finding out at some point of time whenever it is increasing in size some droplet you know can also dislodged from the surface okay. So loss of drop due to gravity is also very, very important.

Apart from that simultaneously you will be finding out over the surface more smaller nucleations are still continuing okay. Those are different stages of droplet evaluation and you will be finding out if you continue in this side further. Then once you will once up on a time you will be finding out that all those droplets are actually forming a film over the surface and it is turned into film wise condensation okay. Next lets us see the next subsection which already I have told as film wise condensation.

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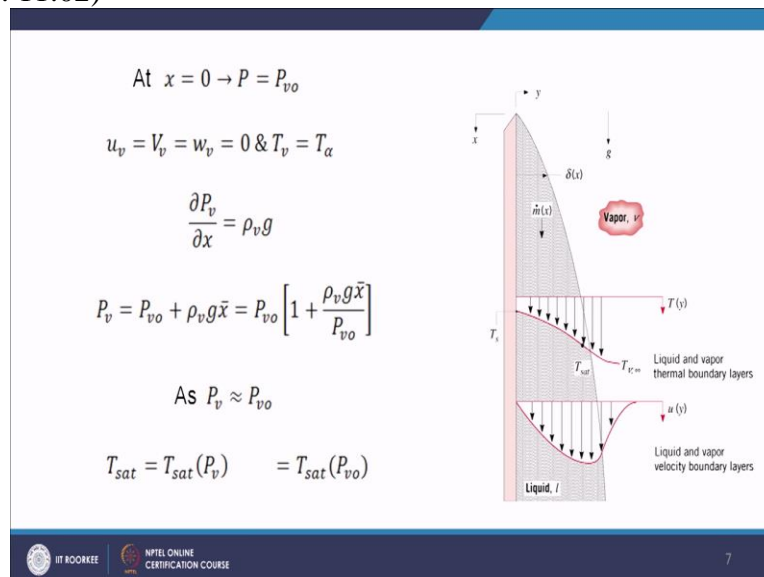
So let us try to see that how film boiling is distinguished from the drop wise condensation. So here I have shown you that vertical surface over which the film is being formed this is your liquid film being formed due to your condensation. So here in the periphery you are having vapor which is suppose to come in contact with the surface and form film okay. So this vapor I have considered as v and liquid I have considered as l . So now first if you try to find out the temperature pattern and velocity pattern over this surface then let us try to see what happens to the temperature.

All of us know that the vapor temperature will be at higher side and obviously surface temperature will be at the lower side which is necessary for condensation. So you will be finding out the temperature will be a smooth fall from the vapor side to the solid surface, where it will be attaining the T_s which is nothing but surface temperature and here it will be attaining T_v which is the T_v infinity you can say which is the free steam vapor temperature okay.

Now at the interface you can find out that there is continuous nature of the temperature also okay. Now if you think about the velocity, so obviously at the wall you will be finding out there is mostly condition. So there is no velocity over here. Obviously on the other hand you will be having no velocity when at the bulk of the gas also. So here you see once again the velocity is 0. So in between whatever film has being it has been formed that will be going in the downward direction. So you can find out a downward velocity in the film okay.

And at the interface you will find out due to inertia you will be finding out the liquid which is falling down will be carrying some amount of gas also along with that in the downward direction. And velocity continuation is there so there is no discontinuity of the velocity so you will be finding out something like this for the velocity profile. Interestingly the highest velocity you can find out somewhere inside the liquid film domain okay.

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So let us now try to understand that how this condensate rate can be predicted okay. \dot{m} can be predicted as well as how this film nature okay, δ or film thickness nature okay. δ is the film thickness so how this film thickness varies with respect to your rate okay. So, in order to do, so we will be doing some sort of mathematical calculation for the vapor as well as the liquid. Let us start with the vapor side. So what will be doing at the leading edge that means where the film starts at this point? Let us call that 1 as $x=0$ where x varies in this side okay.

That means along the plate by it varies and y is actually the perpendicular direction of the wall. So at $x = 0$, you can find out pressure will be equivalent to your vapor pressure okay. So here we

are having no liquid. So obviously the pressure will be equivalence to your P_v okay. Next, in this position as you are not having any film liquid film. So obviously the velocities will be also 0. So we can write down that u_v , v_v and w_v all are equals to 0 okay. And the temperature at this zone obviously will be the equivalent to your infinite temperature or bulk temperature.

So we can call T_v is $= T_{\infty}$ right. If you try to write down the momentum equations for the vapor now as we are having all the u 's $= 0$. So you will be finding out only the pressure term and buoyancy term remain. So we can write down $\frac{dP_v}{dx} = \rho_v g$ okay. So once you integrate this one we find out that $P_v = P_{v,not} + \rho_v g x$. Here we have used the integration criteria as $x = 0$, $P = P_v$. So boundary condition is at $x = 0 \rightarrow P = P_{v,not}$.

So if we take $P_{v,not}$ common form, here we get $1 + (\rho_v g x / P_{v,not})$. Interestingly you see over here ρ_v is there. So which is a very, very small value in case of at least for the water, so what we can do, we can neglect this term. So we can write down vapor pressure will be more or less equivalent at this zone. We can write down the vapor pressure is more or less equivalent to your $P_{v,not}$ okay. So we can write down your corresponding saturation pressure saturation temperature T_{sat} , which is actually your saturation temperature corresponding to P_v that is once again equivalent to your T_{sat} at $P_{v,not}$ because $P_v = P_{v,not}$. We have just now found out okay.

So and the limiting conditions for considering this ρ_v very small. We can write down $T_{sat} = T_{sat, P_{v,not}}$ okay. Next let us try to see from the liquid side but before going to the liquid side let us see the interface okay. So here we are having the interface between the liquid film and the vapor. So what we have taken, we have taken a small sector across the interface. So this is the small sector you see l and g okay. Liquid is there in this side in the film and gas is there in the bulk okay. So if we try to first find out the mass flow rate okay.

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Mass flow rate of liquid film, $\dot{m} = (\rho A V_n)_l = (\rho A V_n)_v$ where, $V_n = \bar{V} \hat{n}$

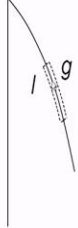
Energy balance, $\dot{m} \left(h_l + \frac{V_l^2}{2} \right) = \dot{m} \left(h_v + \frac{V_v^2}{2} \right) + \dot{Q} - \dot{W}$



$\dot{Q} = - \left(k \frac{\partial T}{\partial n} A \right)_l + \left(k \frac{\partial T}{\partial n} A \right)_v \quad \forall \quad \dot{W} = 0$

As, $k_v \ll k_l$ Therefore, $\left(k \frac{\partial T}{\partial n} A \right)_v = 0 \rightarrow \dot{Q} = - \left(k \frac{\partial T}{\partial n} A \right)_l$

$(\rho A \bar{V} \hat{n})_{int} \left[h_f + \frac{V_l^2}{2} \right] = (\rho A \bar{V} \hat{n})_{int} \left[h_g + \frac{V_g^2}{2} \right] - k_l \left(\frac{\partial T}{\partial n} \right)_{int} A$

$k \left(\frac{\partial T}{\partial n} \right)_{int} = \rho (\bar{V} \hat{n})_{int} h_{fg}$



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So mass flow rate for the liquid film. So mass flow rate \dot{m} can be written as $\rho a v_n$. Here this v_n is the actually normal directional velocity of the interface. So here if you find out a normal of this one. So that directional velocity will be coming into picture over here okay. So this is actually equivalent to 1 and this is actually for the vapor. So because this 2 will be same because we know that whatever amount of mass will be being lost from the gas that will be accumulated in the liquid side okay.

So after doing the mass balance, let us go for the energy balance side. So if you go for the energy balance side in this across this interface then we can write down for the liquid side $\dot{m} * h_l + v_l^2 / 2$. So this h_l is nothing but your enthalpy for the liquid side. v_l is nothing but the velocity for the liquid okay. So here you have added the kinetic energy portion over here okay and on the other hand side for the vapor, we can write down $\dot{m} * h_v$, which is the enthalpy for the vapor + $v_v^2 / 2$.

Along with that we will be having the heat transfer and work done also as for the first law of thermodynamics. So you are having $\dot{q} - \dot{w}$ but obviously in this case we are not having any \dot{w} . So we can make it 0 okay. This can be cancelled out and easily this \dot{q} can be written as now this \dot{q} . \dot{q} will be the heat transfer. Now the heat transfer will be dependent on the conduction to the liquid and conduction to the gas okay.

So conduction to the liquid if you write down then that will be $-k (\frac{\Delta T}{\Delta n}) A$. Now this n is once again the normal direction okay. All the properties I will be taking for l that means liquid. So that means this will be $k_l A$ okay. And here for the vapor side we will be getting $k_v \frac{\Delta T}{\Delta n}$ of a right. Now you see they are opposite in sign because one will be actually accepting heat other one will be actually dumping heat okay. So q we can write down at the summation of both the conduction heats in liquid and vapor side okay.

Now if we put that this q and w in this equation then ultimately we will be getting the equation like this. Here I have also put the value of \dot{m} from the continuity equation. So I have written $\rho A v_n$. So this is v_n is nothing but actually your normal directional velocity. What I have written in place of this v_n okay multiplied by $h_f + \frac{v_l^2}{2}$ equals to once again this is for the gaseous side mass flow rate. Remember what we have done over here. As this l and v are for the liquid side and vapor side in the limiting condition, we can write down there is on the interface.

So here both these l and v , we have written as int , this symbolizes the mass transfer across the liquid okay. So this will be for the gaseous side mass transfer from the gaseous side multiplied by $h_g + \frac{v_g^2}{2}$ and the other hand side we are having over here the conduction part. Now see in the conduction part also. As I have shown you this conduction is actually summation of 2 conductions. So liquid side and vapor side but we know for the vapor side the thermal conduction conductivity will be very low compared to the liquid side.

So this term actually we can cancel out and this q can be written as $-k \frac{\Delta T}{\Delta n} A$ for the liquid side only. So that term we have kept over here as q right. So if we see this equation. So and take this first term in the right hand side to the left hand side. So I can write down that will be coming as $(h_f - h_g)$ okay. Why because this is the limiting condition at the interface and at the interface already we have shown that there is no discontinuity of the velocity. So you will be finding out v_l will be equivalent to your v_g . So that part will be canceling out and will be having $(h_f - h_g)$ which is nothing but h_{fg} okay. So we get in this side the conduction and here in this side we get your condensation heat transfer okay involved with the h_{fg} okay.

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For falling liquid film:

$$\text{Continuity: } \frac{\partial u}{\partial x} + \frac{\partial v}{\partial y} = 0$$

$$\text{x-momentum: } \rho \left(u \frac{\partial u}{\partial x} + v \frac{\partial u}{\partial y} \right) = -\frac{\partial P}{\partial x} + \rho g + \mu \left(\frac{\partial^2 u}{\partial x^2} + \frac{\partial^2 u}{\partial y^2} \right)$$

$$\text{y-momentum: } \frac{\partial P}{\partial y} = 0$$

$$\frac{\partial P}{\partial x} = \frac{\partial P_v}{\partial x} = \rho_v g$$

$$\text{Energy: } \rho C_p \left(u \frac{\partial T}{\partial x} + v \frac{\partial T}{\partial y} \right) = k \left(\frac{\partial^2 T}{\partial x^2} + \frac{\partial^2 T}{\partial y^2} \right)$$



Next let us see after this 1 interfacial condition let us see what happens for the liquid side. So we have taken falling liquid film assumptions. So in falling liquid film assumptions we have written down over here the continuity, x momentum and y momentum equations respectively okay. So these are coming from your fluid mechanics already you know and here as we have already shown that $\frac{\partial p}{\partial x}$ for the vapor side will be $\rho_v \cdot g$ that also you will be using into consideration, because this we have already shown for the gaseous phase momentum equation side.

And energy equation in the liquid if you write down, so in the left hand side, you are having convection and the right hand side you are having conduction okay. Next to reduce this number of equations you see over here we are having at least 1, 2, 3, 4 equations. So these 4 equations if you try to reduce the number of dependence then what will be doing, we have taken some approximations. So first approximation what you take is called Nusselts analysis. So he has, Nusselt, he has proposed this one what he has consider the assumptions are laminar flow in the liquid.

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Nusselts's analysis:

- Laminar
- Steady
- Constant property
- Negligible stream-wise diffusion of momentum
- Negligible stream-wise diffusion of heat
- Negligible inertia and convection term

Using these assumptions x-momentum and energy equation can be written as,

$$0 = (\rho - \rho_v)g + \mu \frac{\partial^2 u}{\partial y^2} \qquad 0 = \frac{\partial^2 T}{\partial y^2}$$



He has considered flow of liquid steady; he has considered the liquid is having constant property negligible stream wise diffusion of momentum. So negligible stream wise diffusion of momentum means in your equation $\frac{\partial^2 u}{\partial x^2}$ will be equals to 0 because x is the stream direction. So that portion is 0 okay diffusion stream wise diffusion is 0. Similarly, negligible stream wise diffusion of heat. So your energy equation you will be finding out this $\frac{\partial^2 T}{\partial x^2}$ will be also going to 0 okay.

And negligible inertia and convection term so this is very important so convection, inertia if you neglect so all these left hand side and in the energy x momentum equation and energy equation will be canceling to 0 okay. So our equation sets of equation becomes very, very simple. So this comes from x momentum equation. 0, already we have cancelled this inertia. So $0 = (\rho - \rho_v) * g$. Where from this $(\rho - \rho_v)$ is coming? Actually you see you are having $\frac{\partial p}{\partial x}$ over here. So $\frac{\partial p}{\partial x} = \rho * g$ we have got over here.

So we put this from this equation over here. $\frac{\partial p}{\partial x}$ value will be getting a $(\rho - \rho_v) * g$. So that term will be coming over here $(\rho - \rho_v) * g$ +. Only the cross wise diffusion is term is remaining in the momentum equation. And from the energy equation we get very simplified form $\frac{\partial^2 T}{\partial y^2} = 0$ because stream wise diffusion, a stream wise diffusion in energy equation and your convection has been neglected as per the Nusslet analysis okay. Let us see the boundary conditions.

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Boundary conditions:

$$y = 0 \rightarrow u = 0 \text{ and } T = T_w$$

$$y = \delta(x) \rightarrow \frac{\partial u}{\partial y} = 0 \text{ and } T = T_{sat}$$

From momentum equation, $\frac{\partial^2 u}{\partial y^2} = -\frac{(\rho - \rho_v)g}{\mu}$

From energy equation, $\frac{\partial^2 T}{\partial y^2} = 0$

After integrating under given set of BC's,

Liquid film velocity, $u = -\frac{(\rho - \rho_v)g}{\mu} \left(\frac{y^2}{2} - \delta y \right)$

Temperature profile, $\frac{T_{sat} - T}{T_{sat} - T_w} = 1 - \frac{y}{\delta}$

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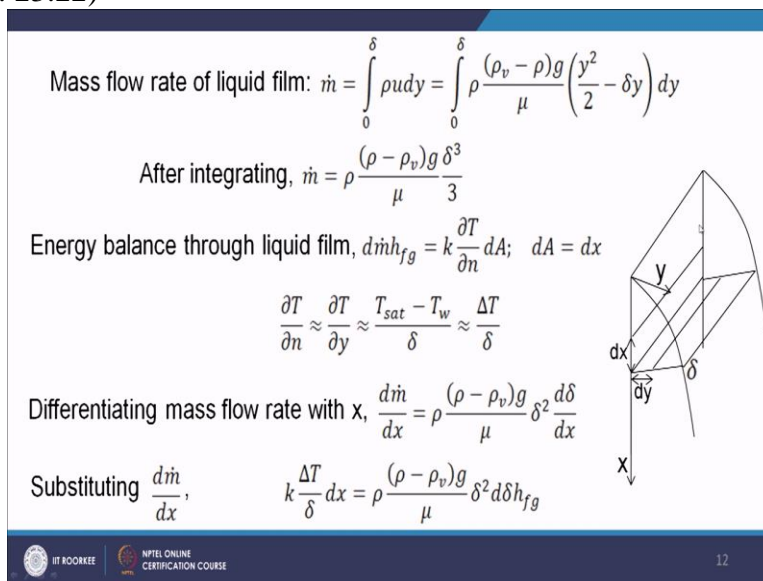
Now so at $y = 0$ that means on the surface obviously the velocity will be 0 and the temperature will be T_w which is the wall temperature or you can call that one as surface temperature T_s okay. Now at interface which is nothing but $y = \delta$ will be finding out that there is no gradient of velocity because the velocity continues over that. So $\partial u / \partial y = 0$ as well as we find out that there the phase change is happening, so the temperature will be actually equivalent to your saturation temperature. So $t = T_{sat}$, so if we now integrate your momentum equation and energy equation which already we have discussed based on the Nusselts analysis.

If we integrate along with this boundary conditions then you will be finding out u from the integration of momentum equation comes out to be $-(\rho - \rho_v)g / \mu * y^2 / 2 - (\delta y)$ very simplified integration we are having and we have to put the boundary conditions as $y = 0, u = 0, y = \delta, \partial u / \partial y = 0$ okay. So we get the liquid film velocity in this fashion and energy equation is very simple. So 2 times integration if we do for this equation and along with the boundary condition $y = 0, T = T_w, y = \delta, T = T_{sat}$.

We get straight forward solution for the temperature profile $T_{sat} - T_w = 1 - y / \delta$ okay. So you see this nature if you see already we have shown you over here. So this y verses u that is parabolic nature. So this is a parabolic nature and for t verses y that is actually a linear nature. So you see that is the linear nature over here. We have seen okay. Next, let us try to find out that

how condensate rate can be achieved. So here I have shown you that condensate rate $\dot{m} = 0$ to δ $\rho u dy$ okay. Already we have found what is y .

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Mass flow rate of liquid film: $\dot{m} = \int_0^\delta \rho u dy = \int_0^\delta \rho \frac{(\rho_v - \rho)g}{\mu} \left(\frac{y^2}{2} - \delta y \right) dy$

After integrating, $\dot{m} = \rho \frac{(\rho - \rho_v)g}{\mu} \frac{\delta^3}{3}$

Energy balance through liquid film, $d\dot{m}h_{fg} = k \frac{\partial T}{\partial n} dA$; $dA = dx$

$\frac{\partial T}{\partial n} \approx \frac{\partial T}{\partial y} \approx \frac{T_{sat} - T_w}{\delta} \approx \frac{\Delta T}{\delta}$

Differentiating mass flow rate with x , $\frac{d\dot{m}}{dx} = \rho \frac{(\rho - \rho_v)g}{\mu} \delta^2 \frac{d\delta}{dx}$

Substituting $\frac{d\dot{m}}{dx}$, $k \frac{\Delta T}{\delta} dx = \rho \frac{(\rho - \rho_v)g}{\mu} \delta^2 d\delta h_{fg}$

So let us put that y over here or y I have shown in the u . I have shown you in the previous slide liquid film velocity. So let us put this expression over there in integration. So if I put this over here and perform the integration then I get \dot{m} is equals to with the obviously with the limits 0 to δ . Because, film condensation will be inside the film thickness only. So \dot{m} becomes $\rho \cdot (\rho - \rho_v) \cdot g \cdot \delta^3 / 3 / \mu$ okay. So this is the condensate rate very, very important term for condensation.

And if you try to find out that the mass balance energy balance across the liquid film then I will be writing down $d\dot{m}h_{fg} = k \cdot \frac{dt}{dn} \cdot dA$. If you remember this you have already explained across the interface earlier. We have got that $d\dot{m} \cdot h_{fg}$ $d\dot{m}$ this was $d\dot{m}$ actually $d\dot{m}$ into h_{fg} was actually $k \cdot \frac{dt}{dn}$ and the interface okay. So same equation you are using over here $d\dot{m}h_{fg} = k \cdot \frac{dt}{dn} \cdot dA$ okay. Now here interestingly if you see here dA is actually dx . So this is the area so this is rectangle.

If you see this rectangle is actually giving you the heat flux to the film. So you will be finding out that dA is actually $dx \cdot 1$. We have considered unitary directions in this way in this side okay z direction. So you can write down $da = dx$. So if I put this $\frac{dt}{dn}$ over here. $\frac{dt}{dn}$ will be once again the normal direction is y direction. Already I have showed you so $\frac{dt}{dn}$ will

be $\delta t / \delta y$ and that can be reduced to $T_{sat} - T_w / \delta$ okay. So if I put all these things over here $\delta t / \delta n$ will be $\delta t / d$ and $dA = \delta x$ will be getting $\dot{m} dx = \rho \delta x \delta y v$ okay.

So what we can do once again in we have got over here. So if I make the derivative of m and put it over here in this equation then I will be getting this equation you see in this equation the left hand side we are having derivative of x and in the right hand side we are having derivative of δ . So we can easily integrate this one okay. So if we integrate then you find out if we integrate from you know 0 to x okay. Then we will be finding out that this is becoming δ^4 to the power 4 and in the right hand side we are having $\mu k \delta t / \rho \delta x \delta y v g h_{fg} \delta y$ plus A okay.

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Rearranging and integrating δ from 0 to x : $\int \delta^3 d\delta = \int \frac{\mu k \Delta T}{\rho(\rho - \rho_v) g h_{fg}} dx$

After integrating, $\frac{\delta^4}{4} = \frac{\mu k \Delta T}{\rho(\rho - \rho_v) g h_{fg}} x + A$

At, $x = 0 \rightarrow \delta = 0 \rightarrow A = 0$

Therefore, $\delta = \left[\frac{4 \mu k \Delta T}{\rho(\rho - \rho_v) g h_{fg}} x \right]^{1/4}$

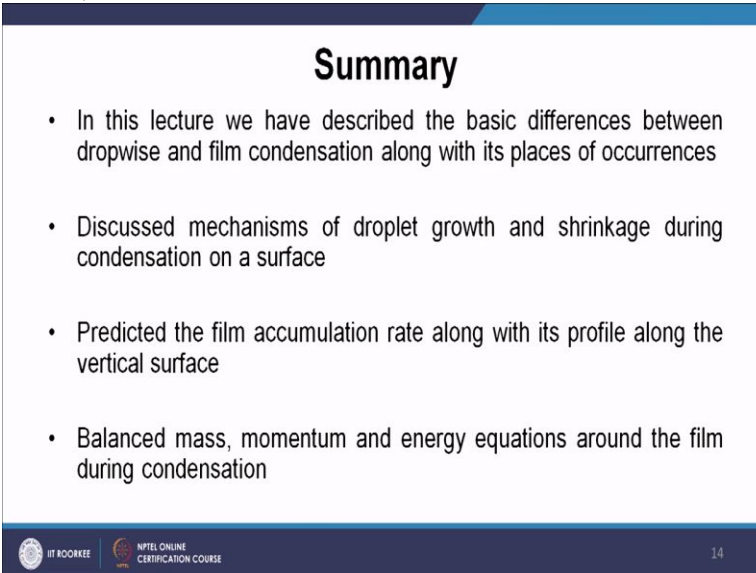
Rearranging, $\frac{\delta}{x} = \left[\frac{4 \theta \alpha \Delta T}{(\rho - \rho_v) g x^3} \times \frac{C_p \Delta T}{h_{fg}} \right]^{1/4}$

Further, $\frac{\delta}{x} = \left[\frac{4}{Ra_x} Ja \right]^{1/4} = \left[\frac{4}{Ra_{fx}} \right]^{1/4}$ where, $Ra_{fx} = \frac{Ra_x}{Ja}$

Now we need the boundary condition for finding out A . So we need the boundary condition means we know that $x = 0$ $\delta = 0$ at the leading edge. So we get that value of $A = 0$ okay. Finally the δ turns out to be $4 \mu k \delta t / \rho \delta x \delta y v g h_{fg} \delta y$ * x to the power $1 / 4$ okay. So if you rearrange this a term a little bit then you will be finding out this is this will be coming out as ratio between 2 non-dimensional numbers 1 is called Jacob number, another 1 is called Rayleigh number and we define the film Rayleigh number as Rayleigh number / Jacob number.

Then we can get this δ / x will be coming out to be $4 / \text{Rayleigh number}$, film Rayleigh number to the power $1/4$. So this is very, very important relationship. If I can find out the film Rayleigh number then easily the δ / x can be obtained okay. So to summarize this lecture what we have done, we have described the basic differences between drop wise and film wise condensation and where those occurs those things we have discussed. We have discussed the mechanisms of droplet growth and reduction okay in case of drop wise condensation.

(Refer Slide Time: 27:23)



The slide is titled "Summary" in a bold, black font. It contains four bullet points summarizing the lecture content. The slide has a white background with a blue header and footer. The footer includes the IIT Roorkee logo, the text "IIT ROORKEE", the NPTEL logo, the text "NPTEL ONLINE CERTIFICATION COURSE", and the page number "14".

Summary

- In this lecture we have described the basic differences between dropwise and film condensation along with its places of occurrences
- Discussed mechanisms of droplet growth and shrinkage during condensation on a surface
- Predicted the film accumulation rate along with its profile along the vertical surface
- Balanced mass, momentum and energy equations around the film during condensation

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We predicted the film accumulation rate okay in case of film wise condensation over vertical surface and we have balanced mass momentum and energy equations to find out the film profile over a vertical surface okay. So let us test your understanding. So we are having once again 3 questions over here.

(Refer Slide Time: 27:48)

Test your understanding ?

1. Necessary condition for condensation
 - a. $T_w < T_{sat}$
 - b. $T_w > T_{sat}$
 - c. $T_w = 0\text{ K}$
 - d. $T_w = T_{\text{triple point}}$
2. Necessary assumption for Nusselt analysis
 - a. Laminar
 - b. Neglect inertia and convection
 - c. Neglect buoyancy momentum
 - d. Neglect crosswise diffusion of momentum
3. Condensate rate is proportional to
 - a. $1/\delta$
 - b. δ^2
 - c. δ
 - d. δ^3



First question, necessary condition for condensation. You are having 4 answers part a $T_w < T_{sat}$, part b $T_w > T_{sat}$, part c $T_w = 0\text{ kelvin}$ and part d $T_w = t\text{ triple point}$ okay. Answer I think all of us know answer will be part a, where surface temperature requires to be lesser than your saturation temperature okay.

Second question goes like this necessary assumption for Nusselt Analysis. We have given you the assumptions today. So the assumptions the answers you are having as first assumption can be Laminar, second assumption is neglect inertia and convection, third assumption neglect buoyancy and fourth assumption neglect crosswise diffusion of momentum okay. So the correct answer will be obviously part a and part b because we neglect the stream wise diffusion and we and we will not be neglecting buoyancy otherwise g will not be coming into picture okay.

Let us see the last question condensate rate is proportional to very simple in the previous slide. I have shown you. So condensate rate \dot{m} will be proportional to 4 answers you are having $1/\delta$, δ^2 , δ and δ^3 . δ is the film thickness here and part c is δ and part d is δ^3 . So the correct answer is δ^3 . In the last slide we have seen. So with this we will be ending this lecture, thank you.