



# ME-446: Liquid-gas interfacial heat and mass transfer

## Boiling: Critical Heat Flux

Zhengmao Lu

Energy Transport Advances  
Laboratory

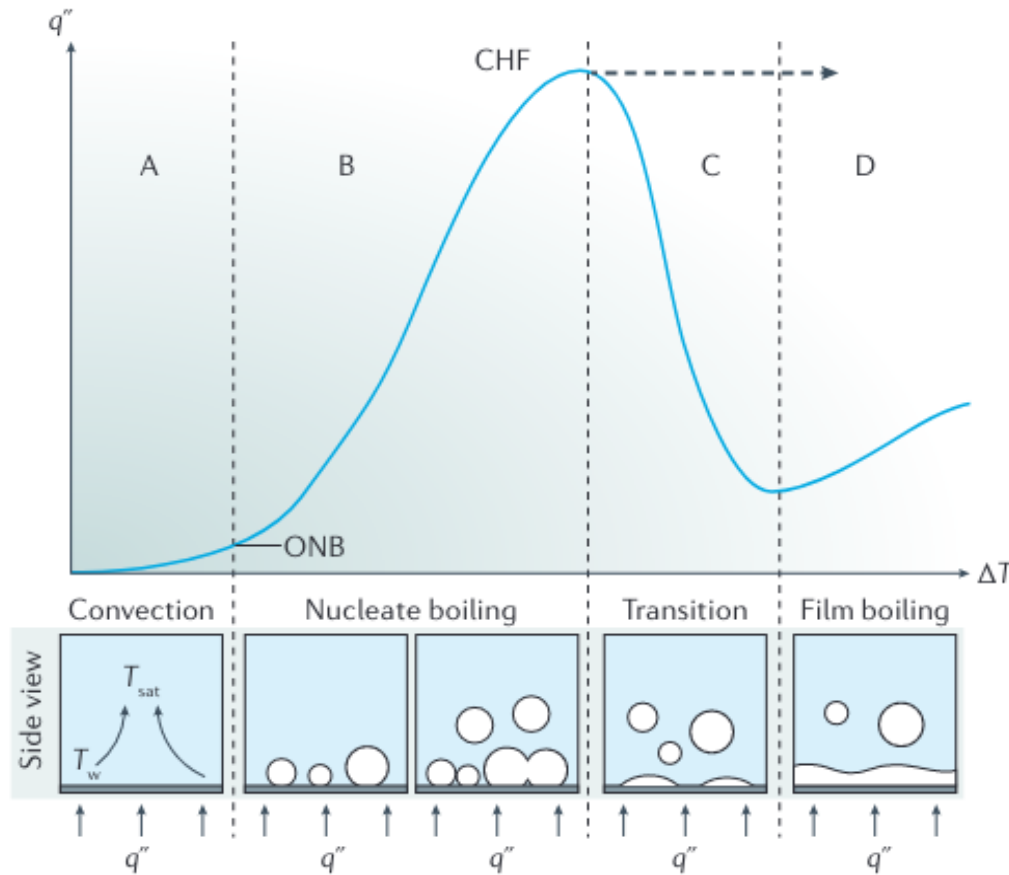
EPFL Mechanical Engineering

2025 Fall Semester

Photo Credit: Trougnouf

- Pool boiling curve
- Rohsenow's microconvection model for nucleate boiling
- Helmholtz and Taylor instabilities

# Pool Boiling Curve



A. At very low superheats, heat transfer is mostly due to natural convection

B. After superheat is large enough to form vapor bubbles, nucleate boiling dominates, promoting bubble-motion-induced convection

C-D. After vapor generation becomes too much, passing the critical heat flux (CHF), insulating vapor film will start to form, decreasing the HTC

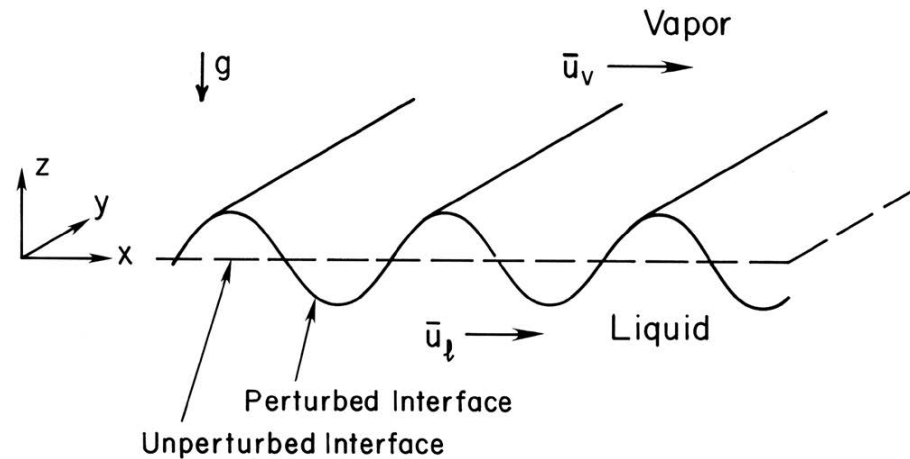
Convective transport facilitated by bubbles

$$\text{Nu}_b = \frac{hL_b}{k_l} \propto \text{Re}_b^{1-r} \text{Pr}_l^{1-s}$$

$$\text{Re}_b = \frac{\rho_v U L_b}{\mu_L} \quad U = \frac{q''}{\rho_v h_{lv}} = \frac{h \Delta T}{\rho_v h_{lv}}$$

$$\frac{q''}{\mu_l h_{lv}} \left[ \frac{\sigma}{g(\rho_l - \rho_v)} \right]^{1/2} = \left( \frac{1}{C_{sf}} \right)^{1/r} \text{Pr}_l^{-s/r} \left[ \frac{c_{pl} \Delta T}{h_{lv}} \right]^{1/r}$$

# Helmholtz Instability

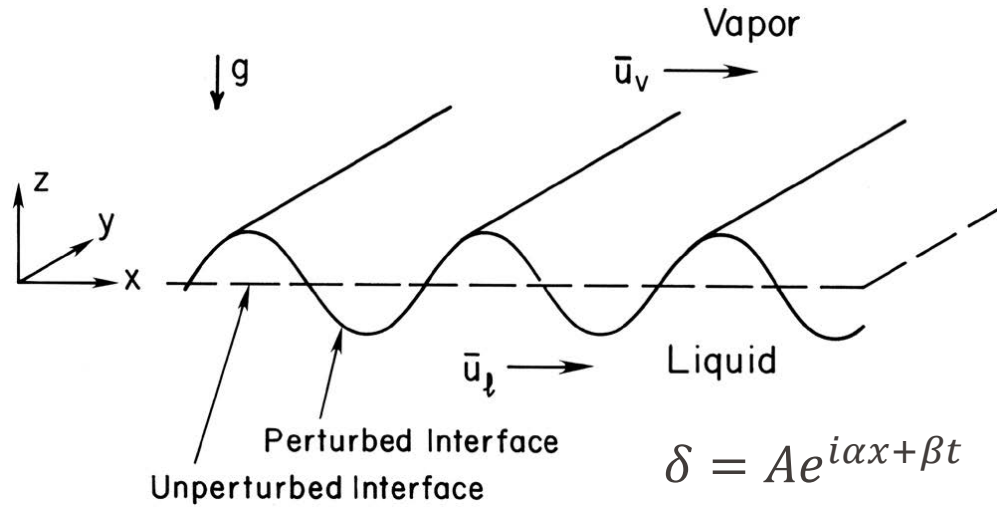


Perturbation  $\delta(x, t = 0) = Ae^{i\alpha x}$   
 Response:  $\delta = Ae^{i\alpha x + \beta t}$

$$\beta = \pm \frac{\sqrt{\alpha^2 \rho_v \rho_l (\bar{u}_v - \bar{u}_l)^2 - (\sigma \alpha^3 + \Delta \rho g \alpha)}}{\rho_v + \rho_l} - i\alpha \frac{\rho_l \bar{u}_l + \rho_v \bar{u}_v}{\rho_v + \rho_l}$$

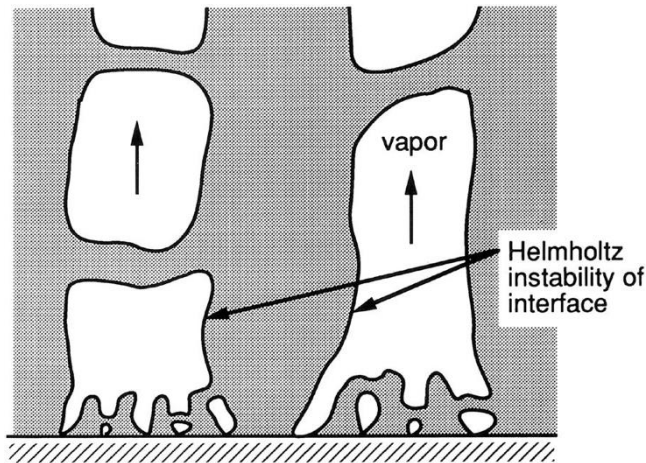
- Instability condition:  $|\bar{u}_v - \bar{u}_l| > \sqrt{\frac{(\sigma \alpha + \frac{\Delta \rho g}{\alpha})(\rho_l + \rho_v)}{\rho_l \rho_v}}$
- When  $|\bar{u}_v - \bar{u}_l| > \left[ \frac{2(\rho_l + \rho_v)}{\rho_l} \right]^{\frac{1}{2}} \left( \frac{\sigma \Delta \rho g}{\rho_v^2} \right)^{\frac{1}{4}}$ , there will be some wave number  $\alpha$  growing exponentially
- $\bar{u}_v - \bar{u}_l$  promotes instability while gravity and surface tension suppressing instability, we can adjust the value of  $g$  based on the orientation of the system.

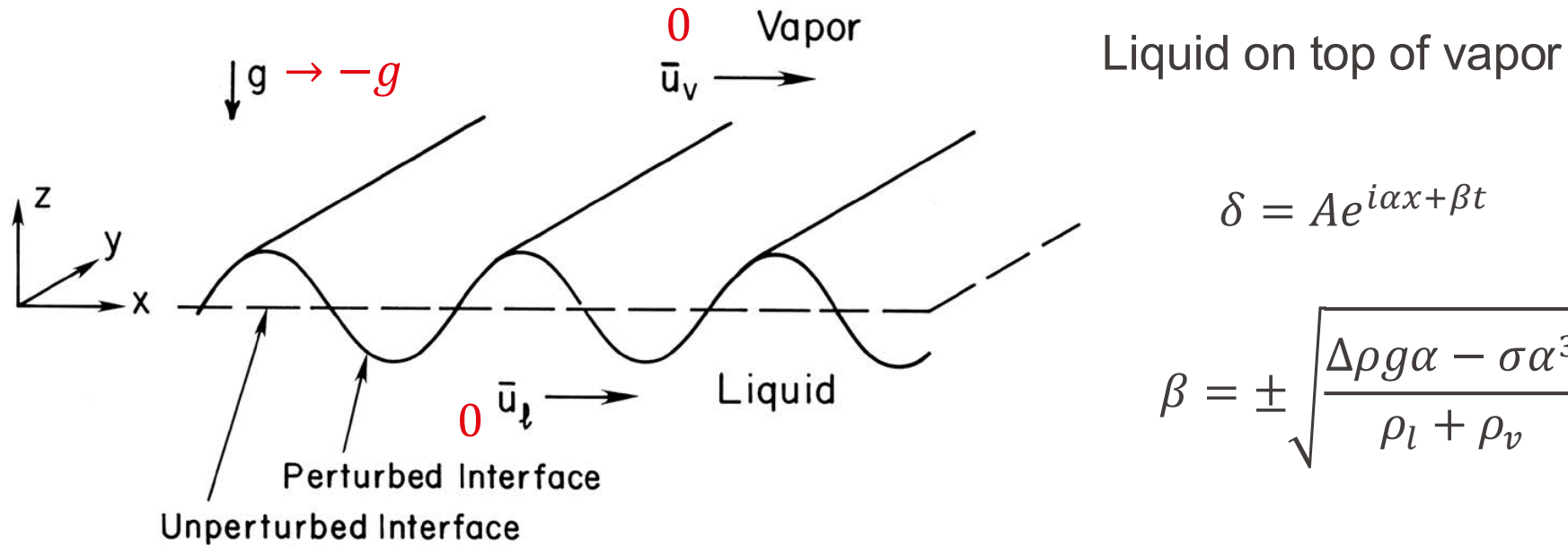
# Vapor Column Instability



Setting  $g = 0$  for in Helmholtz instability for vertical surfaces

$$|\bar{u}_v - \bar{u}_l| > \sqrt{\frac{\sigma \alpha (\rho_l + \rho_v)}{\rho_l \rho_v}} = \sqrt{\frac{2\pi\sigma (\rho_l + \rho_v)}{\rho_l \rho_v \lambda_H}}$$





$$\delta = Ae^{i\alpha x + \beta t}$$

$$\beta = \pm \sqrt{\frac{\Delta\rho g\alpha - \sigma\alpha^3}{\rho_l + \rho_v}}$$

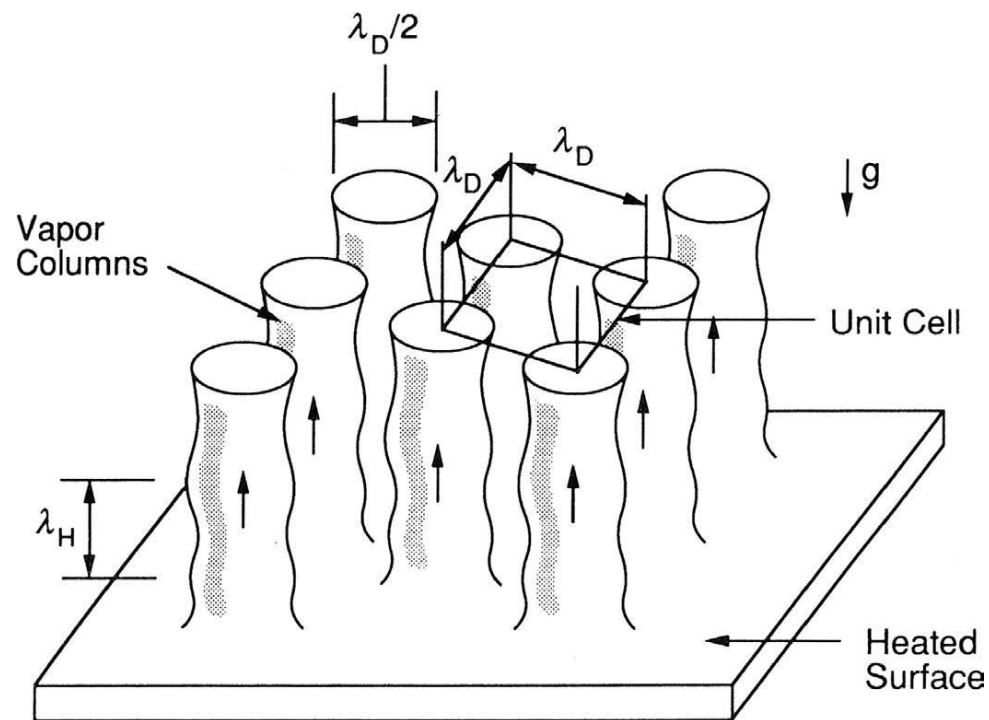
The fastest growing perturbation ( $\alpha_{max}$ ) in this case can be found by setting  $\frac{d\beta}{d\alpha} = 0$

The corresponding most dangerous wavelength  $\lambda_D = \frac{2\pi}{\alpha_{max}} = 2\pi \sqrt{\frac{3\sigma}{\Delta\rho g}}$

- Zuber' CHF model based on hydrodynamic instability
- Force balance model for CHF
- Statistical approach for CHF

# Zuber's Model Assumptions

$$|\bar{u}_v - \bar{u}_l| > \sqrt{\frac{\sigma \alpha (\rho_l + \rho_v)}{\rho_l \rho_v}} = \sqrt{\frac{2\pi\sigma (\rho_l + \rho_v)}{\rho_l \rho_v \lambda_H}}$$



- Vapor columns are formed near CHF
- CHF is reached when interface of vapor columns becomes Helmholtz unstable ( $\lambda_H$ )
- The pitch of the vapor columns coincides with the most dangerous wavelength in Taylor instability

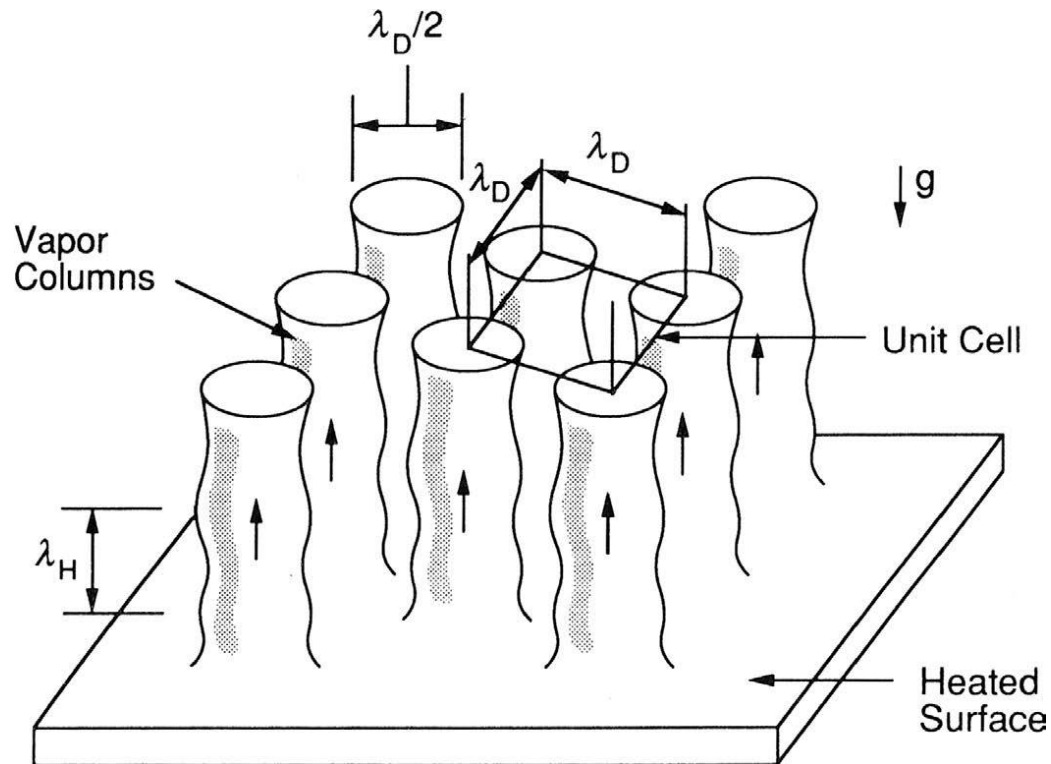
$$\lambda_D = 2\pi\sqrt{3\sigma/\Delta\rho g}$$

- The diameter of vapor column is  $\lambda_D/2$

- Assuming  $\lambda_H = \lambda_D \Rightarrow u_c = \sqrt{\frac{2\pi\sigma (\rho_l + \rho_v)}{\rho_l \rho_v \lambda_H}} \approx \sqrt{\frac{2\pi\sigma}{\rho_v \lambda_D}}$

# Zuber's CHF Prediction

$$u_c = \sqrt{\frac{2\pi\sigma}{\rho_v\lambda_D}} \quad \lambda_H = \lambda_D = 2\pi\sqrt{\frac{3\sigma}{\Delta\rho g}}$$



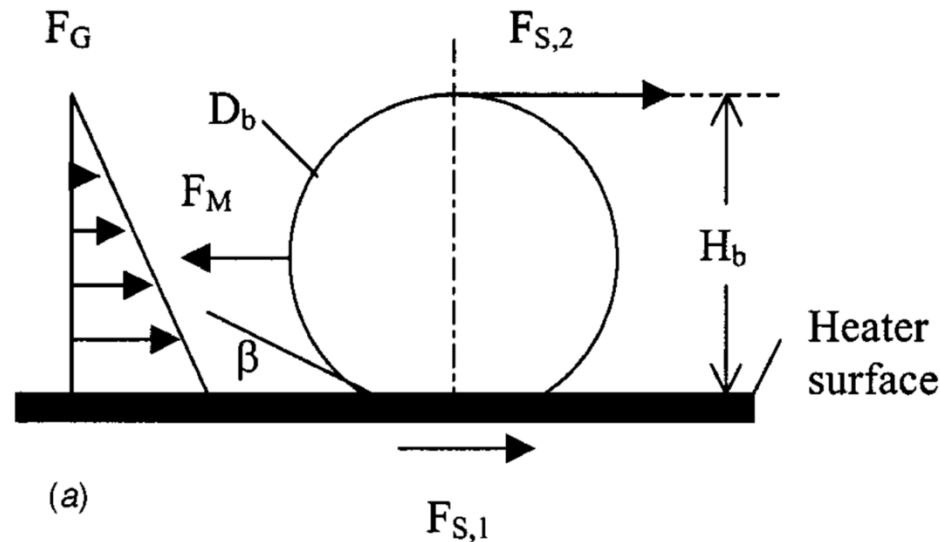
$$u_c = \frac{q''_{max}}{\rho_v h_{lv}} \left( \frac{A_{surf}}{A_{col}} \right) = \frac{16 q''_{max}}{\pi \rho_v h_{lv}}$$

$$q''_{max} = 0.149 \rho_v h_{lv} \left( \frac{\sigma \Delta \rho g}{\rho_v^2} \right)^{1/4}$$

# Comments on Zuber's Model

- No way to accommodate effects from geometry and surface wettability
- No clear justification for the choice of vapor column diameter as  $\lambda_D/2$
- No visual observation of Helmholtz instability during boiling to date
- Still widely used as a reference model for all subsequent CHF models

Kandlikar 2001

<https://doi.org/10.1115/1.1409265>


Considering the lateral direction  
on half of a bubble

Surface tension force :  $F_{S,1}, F_{S,2}$

Hydrostatic force:  $F_G$

Momentum force:  $F_M$  (due to phase change)

Simple force balance requires  $F_{S,1} + F_{S,2} + F_G = F_M$

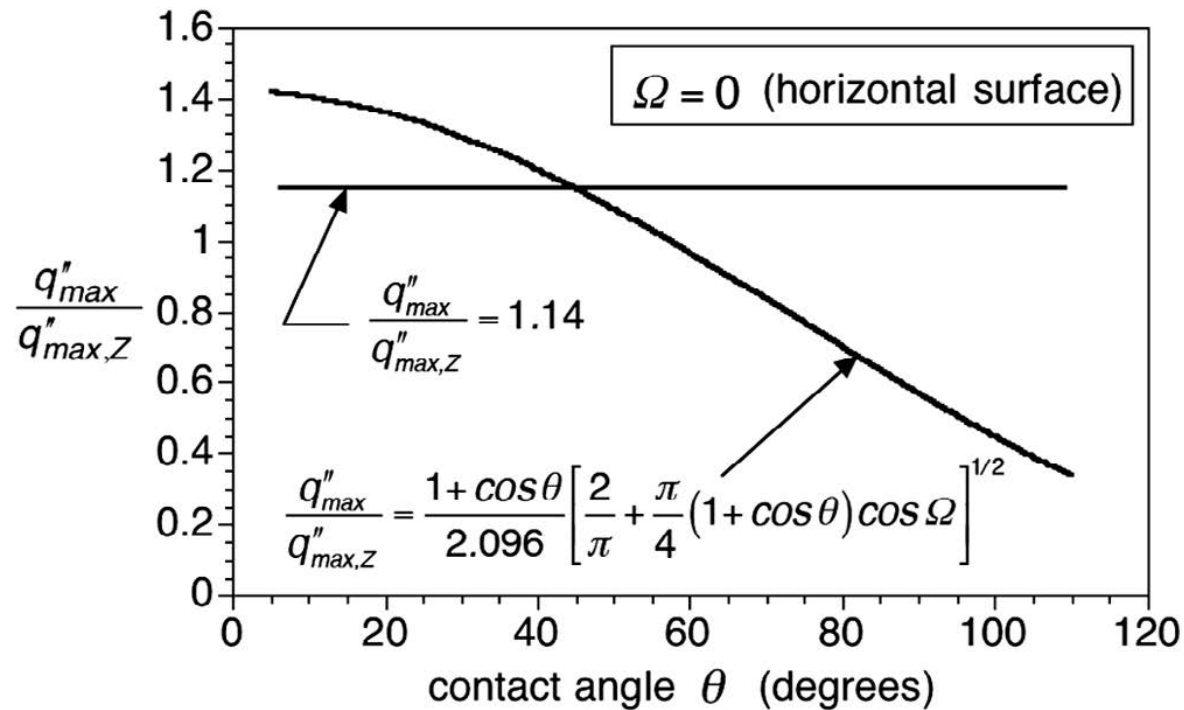
Choose a bubble diameter at CHF  $D_b = \lambda_D/2$

$\lambda_D$ : the most dangerous  
wavelength in Taylor instability  $\lambda_D = 2\pi \sqrt{\frac{3\sigma}{\Delta\rho g}}$

and set a bubble influence area  $\pi D_b^2$

# Contact Angle Dependence

$$q_K'' = \rho_v h_{fg} \left( \frac{1 + \cos \beta}{16} \right) \left[ \frac{2}{\pi} + \frac{\pi}{4} (1 + \cos \beta) \right]^{\frac{1}{2}} \left( \frac{\sigma \Delta \rho g}{\rho_v^2} \right)^{1/4} \quad (\text{Horizontal surface})$$

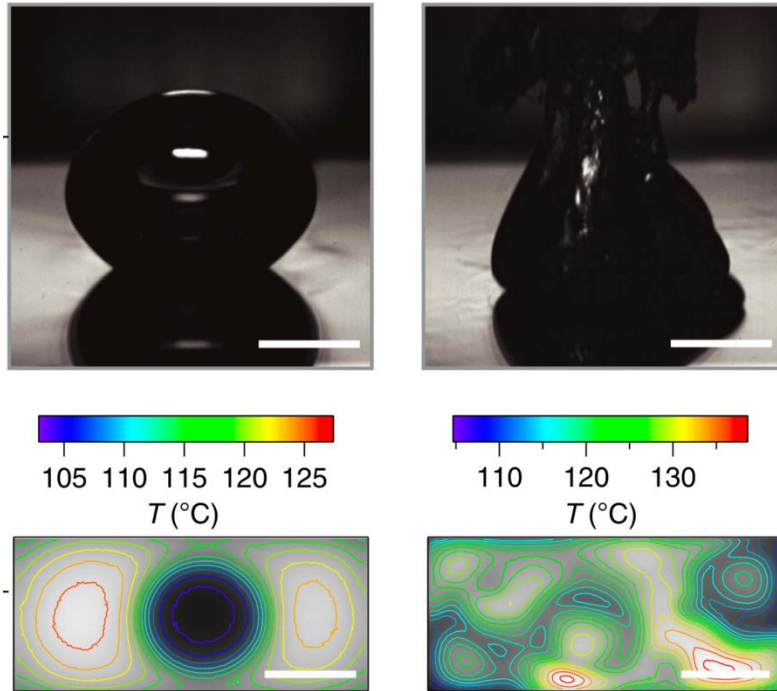


$$q''_{max,Z} = 0.149 \rho_v h_{lv} \left( \frac{\sigma \Delta \rho g}{\rho_v^2} \right)^{1/4}$$

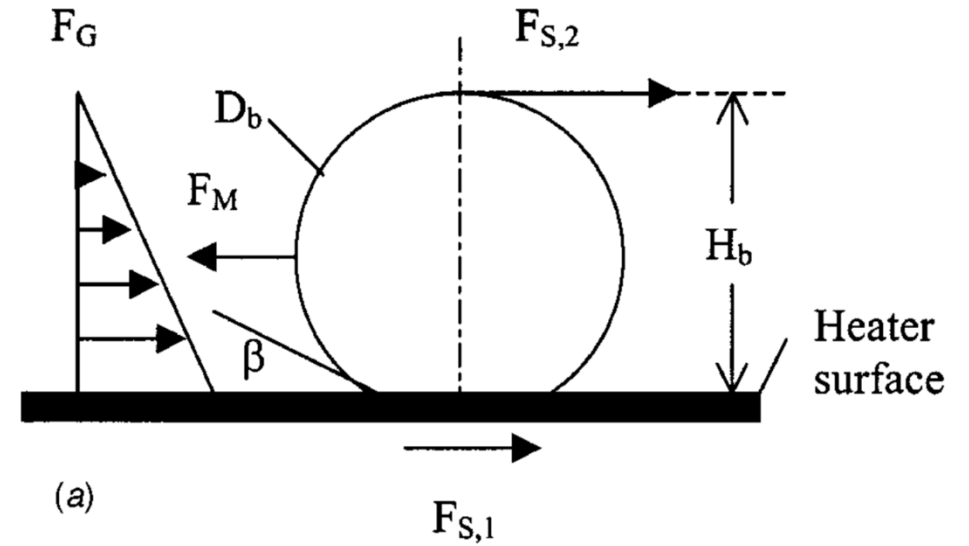
$$\frac{q_K''}{q''_{max,Z}} = \frac{1 + \cos \theta}{2.096} \left[ \frac{2}{\pi} + \frac{\pi}{4} (1 + \cos \theta) \right]^{\frac{1}{2}}$$

FIGURE 7.19 in Carey

Dhillon *et al.*, *Nat Commun* 2015



Choice of geometric parameters not quite justified at CHF



$$F_{S,1} + F_{S,2} + F_G = F_M$$

Liquid-vapor pressure difference not accounted for in momentum balance

# Statistical Approach for Flat Surface Boiling

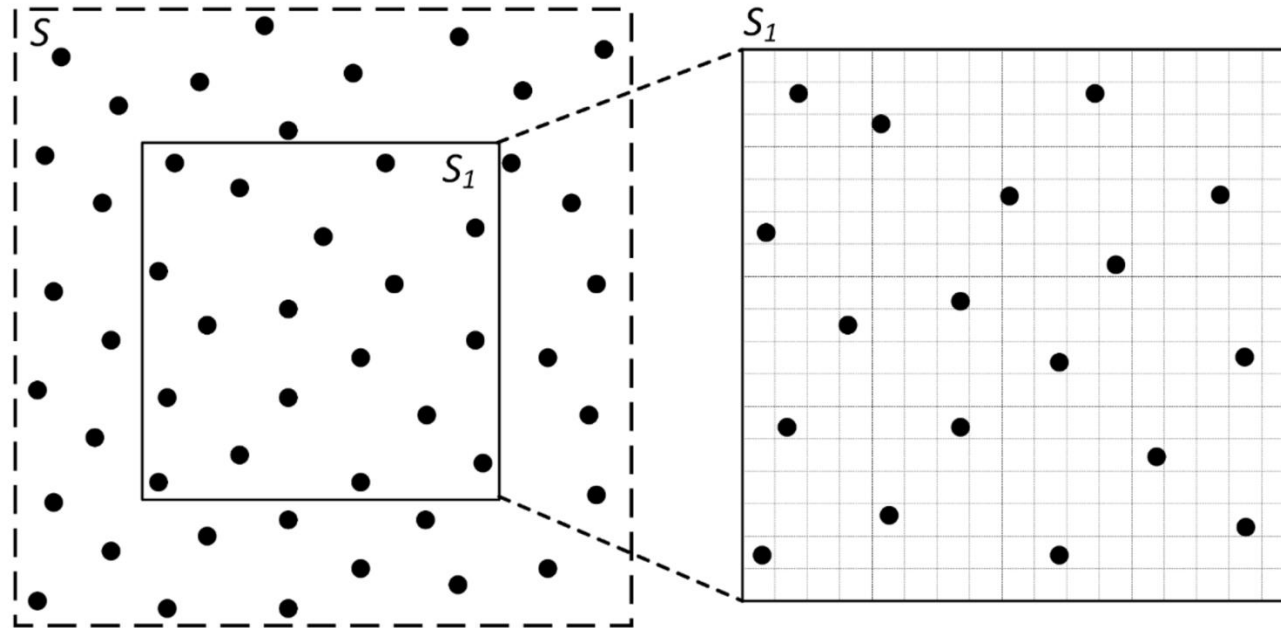


Working fluid: R134a

Heat flux:  
from 1.5 W/cm<sup>2</sup> to 38 W/cm<sup>2</sup>

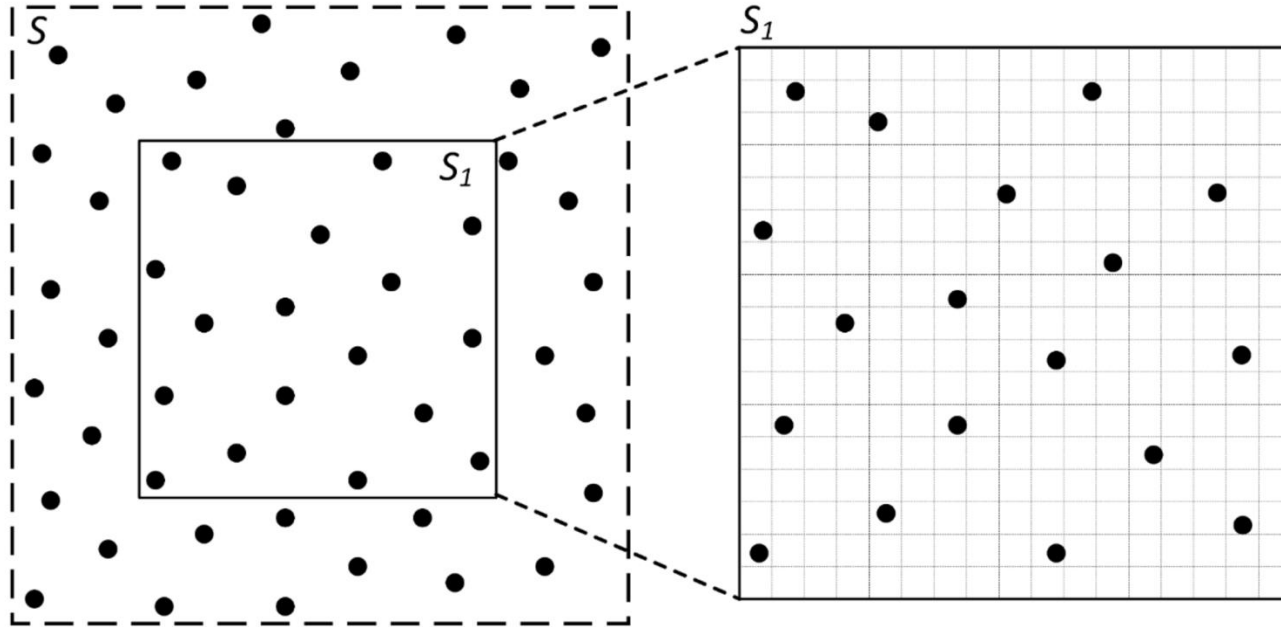
Yazdani, 2016  
<https://doi.org/10.1063/1.4940042>

# Population Distribution of Nucleation Sites

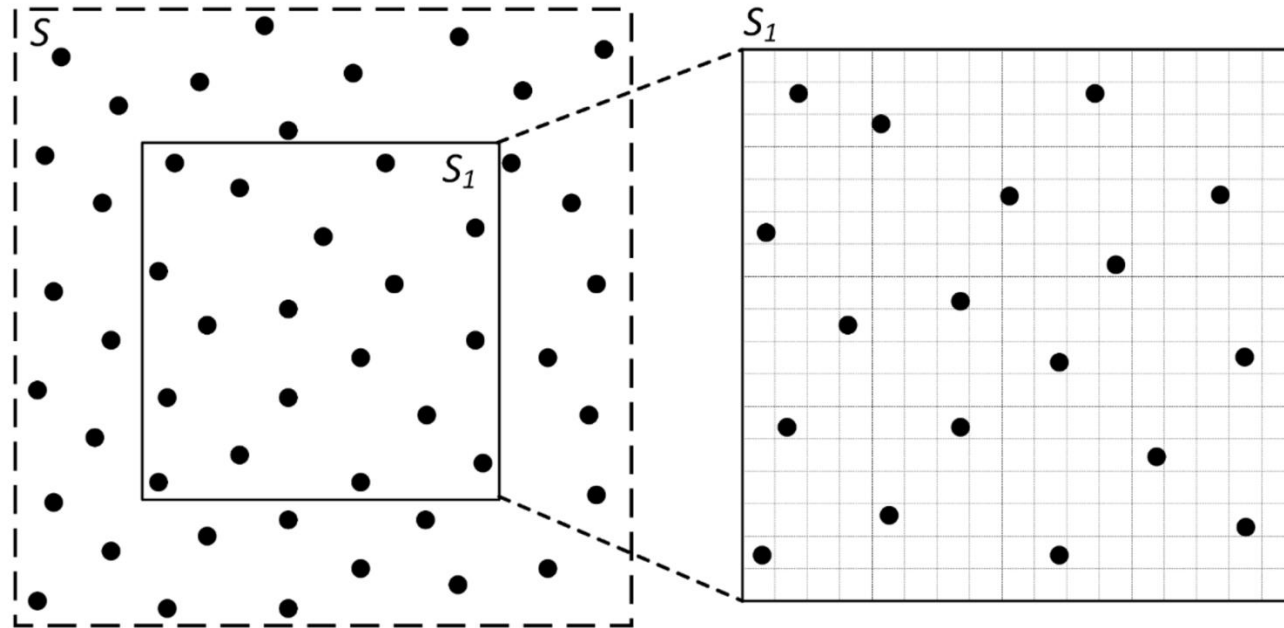


<https://doi.org/10.1016/j.ijheatmasstransfer.2021.121904>

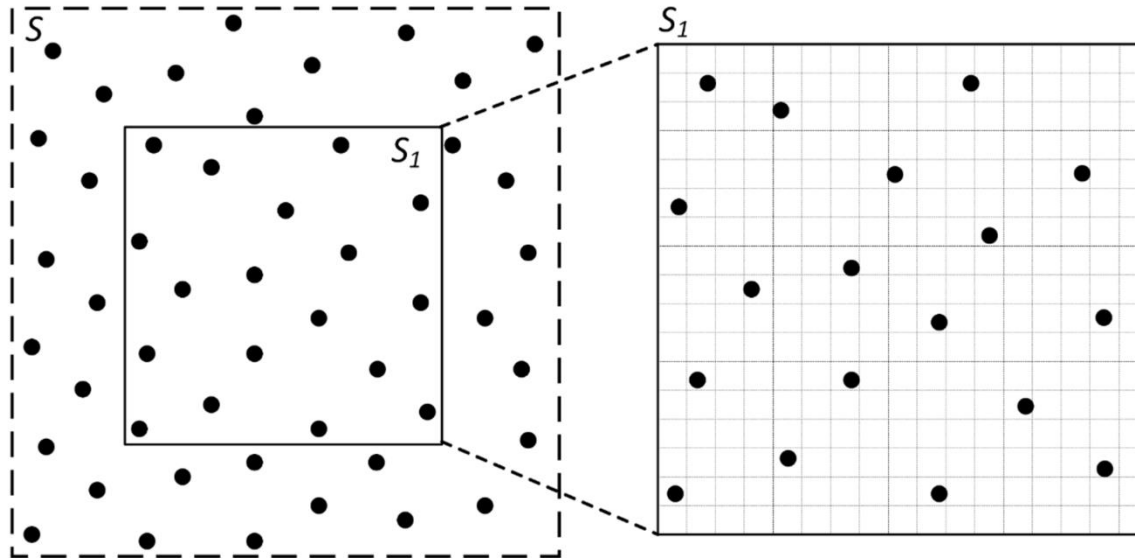
- Consider a large surface  $S$  (large enough to ignore edge effects)
- Assume that probability of each point on the surface becoming an active nucleation sites is equal



- The actual number of active nucleation sites in  $S_1$ ,  $N$ , can be considered as a random variable with an expectation value  $N_0$
- Average nucleation density  $n_0$  [ $\text{m}^{-2}$ ], can be defined as  $n_0 = N_0/A$
- $n_0$  can be assumed as a surface property



- Divide  $S_1$  into  $M$  segments, each with the same area  $A/M$
- Make  $M$  large enough such that the number of nucleation sites in each square is 0 or 1.
- Probability of finding a nucleation site in each square  $p=N_0/M$

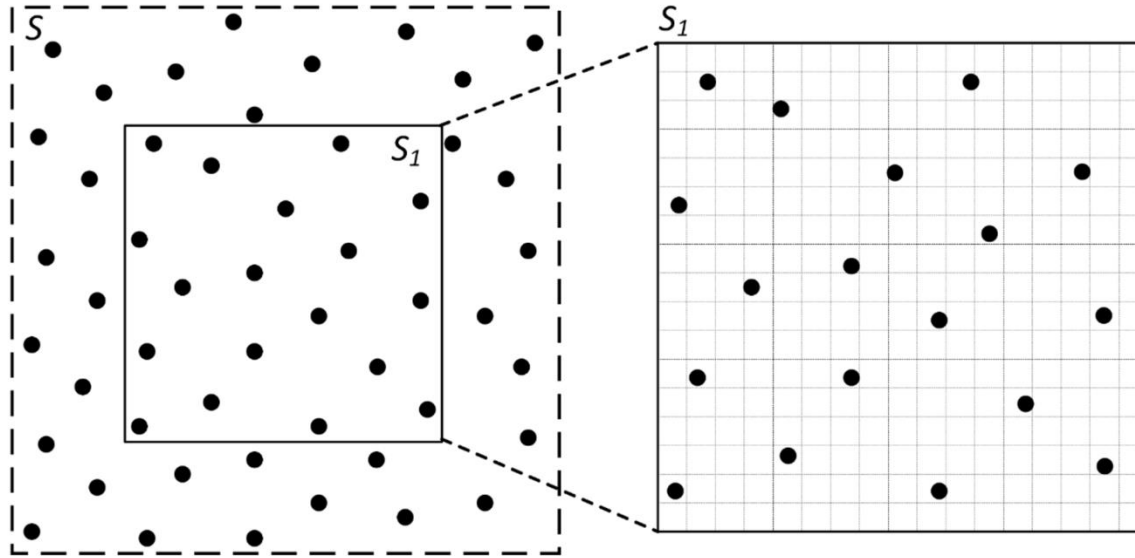


$$P(N, N_0, M) = \frac{1}{N!} \cdot \frac{M!}{(M - N)! M^N} \cdot N_0^N \left(1 - \frac{N_0}{M}\right)^{M-N}$$

Stirling's approximation:  $\ln(n!) = n \ln n - n + O(\ln n)$  for  $n \rightarrow \infty$

$$\lim_{M \rightarrow \infty} \ln \left[ \frac{M!}{(M - N)! M^N} \right] = 0 \Rightarrow \lim_{M \rightarrow \infty} \frac{M!}{(M - N)! M^N} = 1$$

$$\lim_{M \rightarrow \infty} P(N, N_0, M) = \frac{N_0^N}{N!} e^{-N_0}$$



- Probability of finding  $N$  squares that contain one nucleation site is a binomial distribution

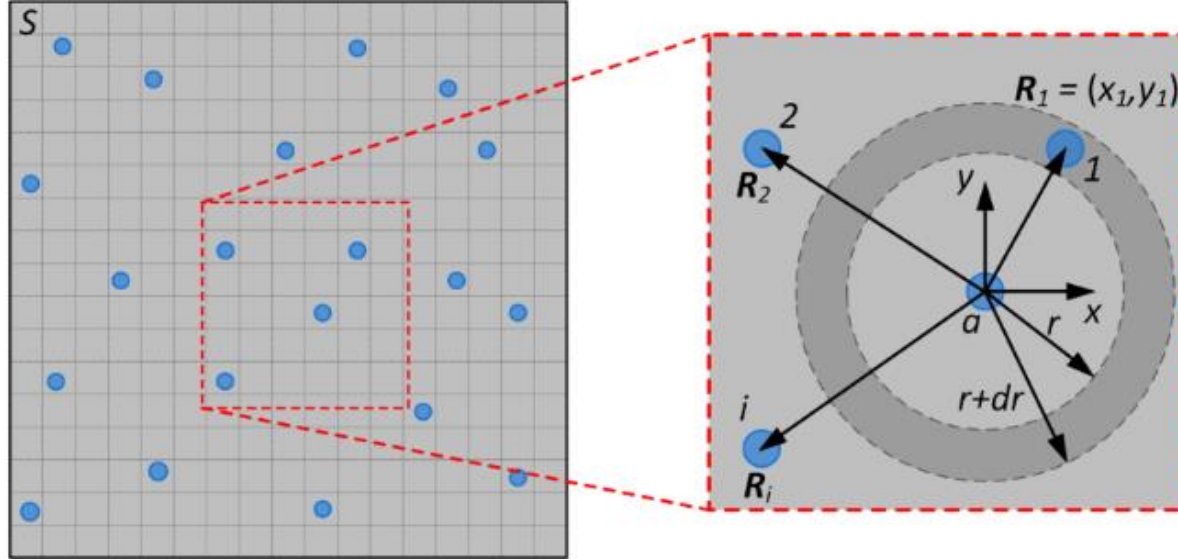
$$P = \frac{M!}{N! (M - N)!} p^N (1 - p)^{M - N} = \frac{1}{N!} \cdot \frac{M!}{(M - N)! M^N} \cdot N_0^N \left(1 - \frac{N_0}{M}\right)^{M - N}$$

$$\text{Po}(N, N_0) = \frac{N_0^N}{N!} e^{-N_0}$$

$$\sum_{N=0}^{\infty} P(N, N_0) = \sum_{N=0}^{\infty} \frac{N_0^N}{N!} e^{-N_0} = e^{N_0} \cdot e^{-N_0} = 1$$

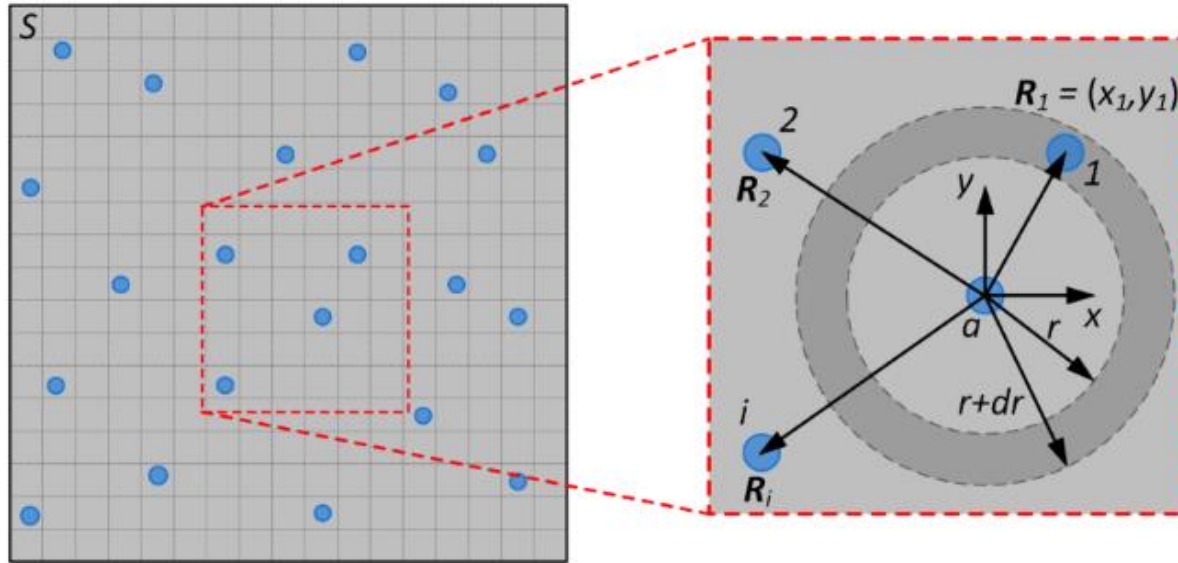
- Isolated bubbles dissipate heat better than merged bubbles
- CHF is reached when you have the **maximum number** of isolated bubbles
- With elevated temperature, 1) more nucleation sites become activated while 2) more bubbles are likely to merge into each other.

# Nearest Neighbor (NN) Distance Between Nucleation Sites



- We would like to know the probability distribution function for distance between two nearest activated nucleation sites
- “NN distance  $> s$ ”  $\Rightarrow$  none of the other  $N - 1$  points inside the disk of radius  $s$

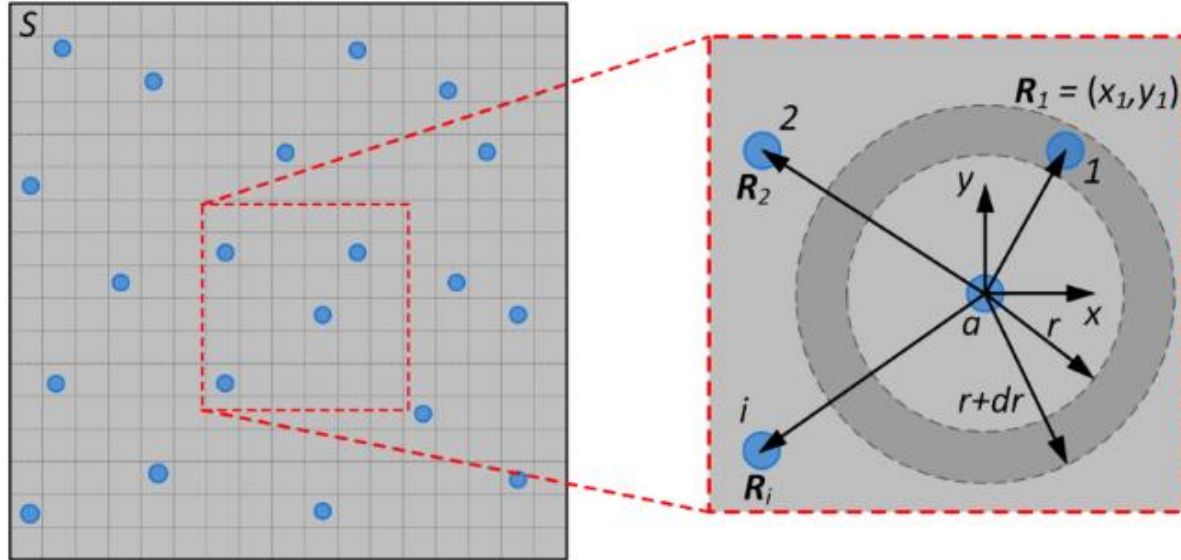
# Nearest Neighbor (NN) Distance Between Nucleation Sites



- Assuming the surface is sufficiently large (no edge effects)

$$\Pr(r > s) = \left(1 - \frac{\pi s^2}{A}\right)^{N-1} \Rightarrow \Pr(r \leq s) = 1 - \left(1 - \frac{\pi s^2}{A}\right)^{N-1}$$

# Nearest Neighbor (NN) Distance Between Nucleation Sites



$$\Pr(r \leq s) = 1 - \left(1 - \frac{\pi s^2}{A}\right)^{N-1}$$

The probability distribution function for NN distance  $f(s)$

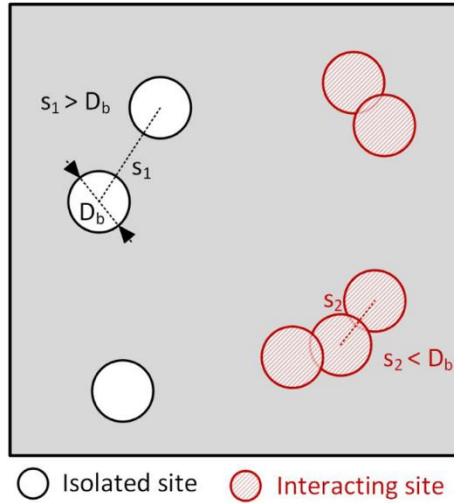
$$f(s) = \frac{d}{ds} (\Pr(r \leq s)) = (N - 1) \frac{2\pi s}{A} \left(1 - \frac{\pi s^2}{A}\right)^{N-2}$$

For  $N \rightarrow \infty$  and  $A \rightarrow \infty$

$$f(s) \approx \frac{2\pi N}{A} s \cdot \exp\left(-\frac{\pi N}{A} s^2\right)$$

Rayleigh  
distribution

# Number of Isolated Bubbles



$$P_{iso} = P(s > D_b) = \int_{D_b}^{\infty} f(s) ds = \int_{D_b}^{\infty} \frac{2\pi N s}{A} e^{-\frac{\pi N s^2}{A}} ds = e^{-\frac{\pi N D_b^2}{A}}$$

$$N_{iso} = \sum_{N=1}^{\infty} N P_{iso} \text{Po}(N, N_0) = \sum_{N=1}^{\infty} \frac{N_0^N}{(N-1)!} \exp\left(-N_0 - \frac{\pi N D_b^2}{A}\right)$$

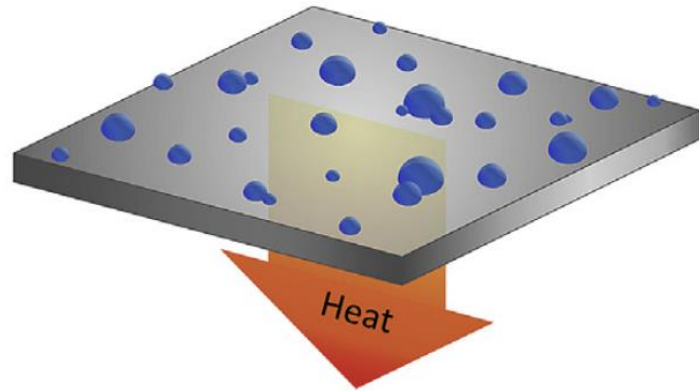
$$\frac{\partial N_{iso}}{\partial T} = 0$$

$$\frac{\partial}{\partial T} \left[ \sum_{N=1}^{\infty} \frac{N_0^N}{(N-1)!} \exp \left( -N_0 - \frac{\pi N D_b^2}{A} \right) \right] = 0$$

$$\Rightarrow n_0 \pi D_b^2 = 1$$

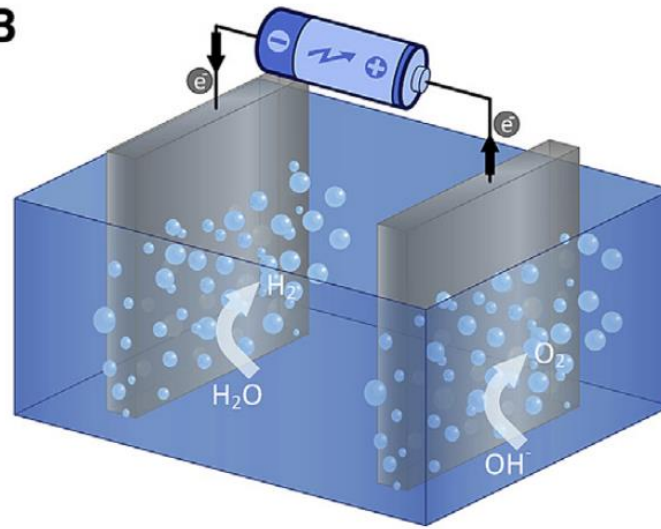
A unified relationship between the nucleation density at CHF and bubble diameter

A



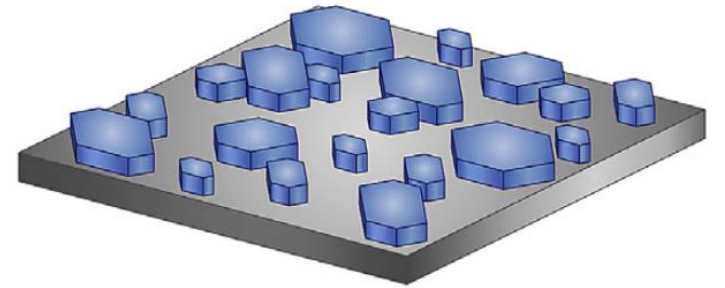
Condensation

B



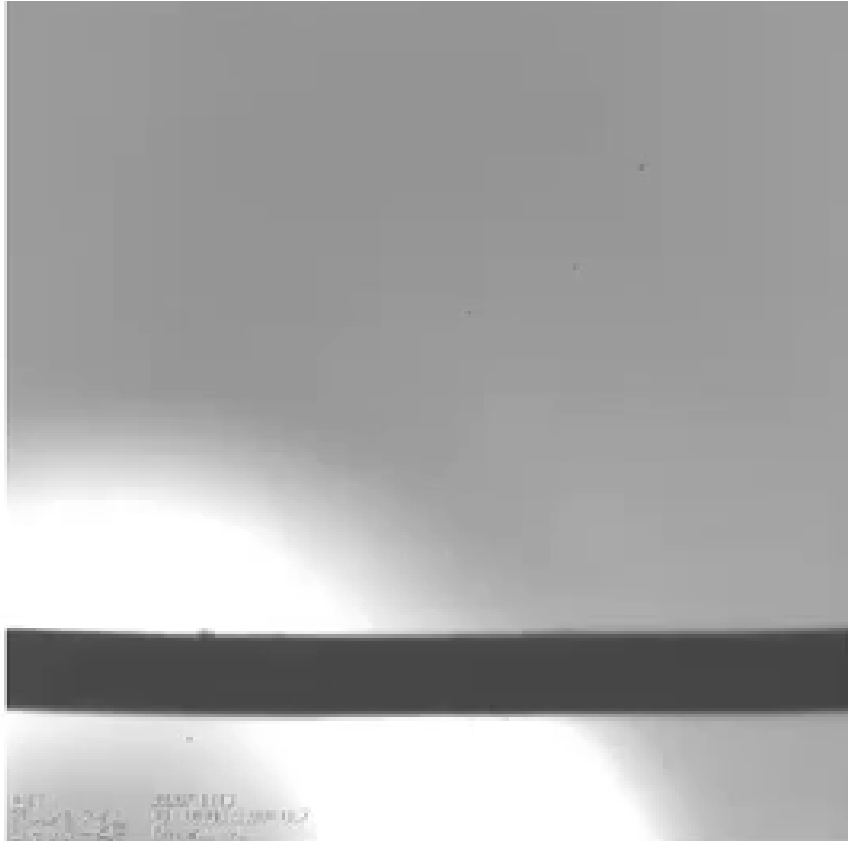
Water splitting

C



Material growth

# Gas Evolving Reaction (GER)



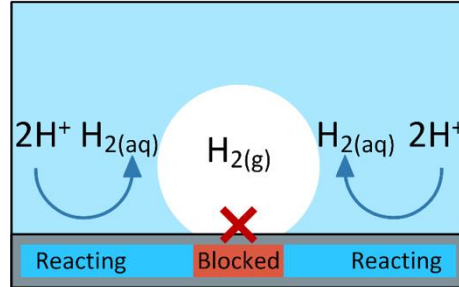
Industrial electrochemical processes that involves bubbles

Water splitting

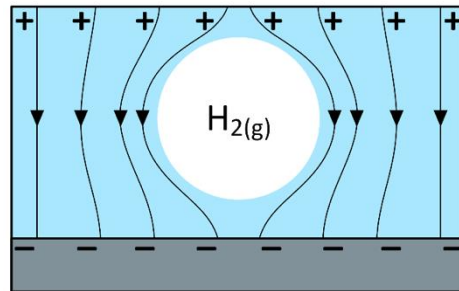
Aluminum production

Sodium chlorate production

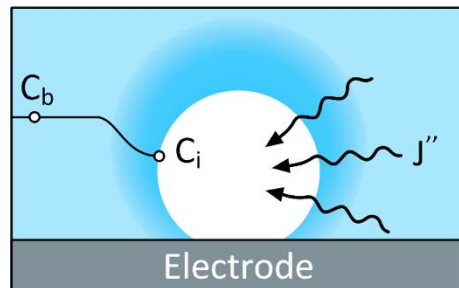
Chlorine and sodium hydroxide production



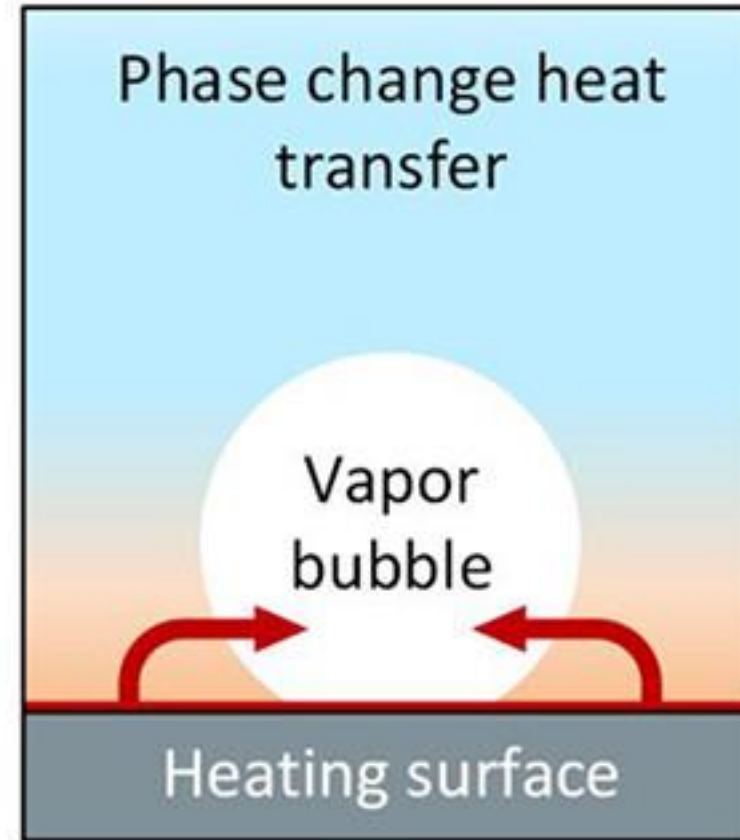
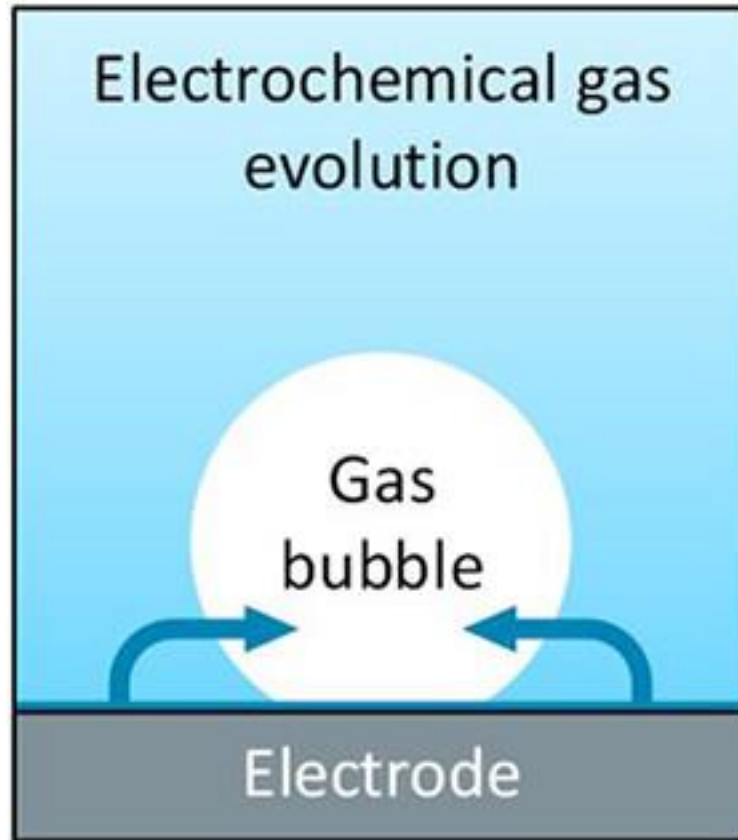
Blocking reaction area

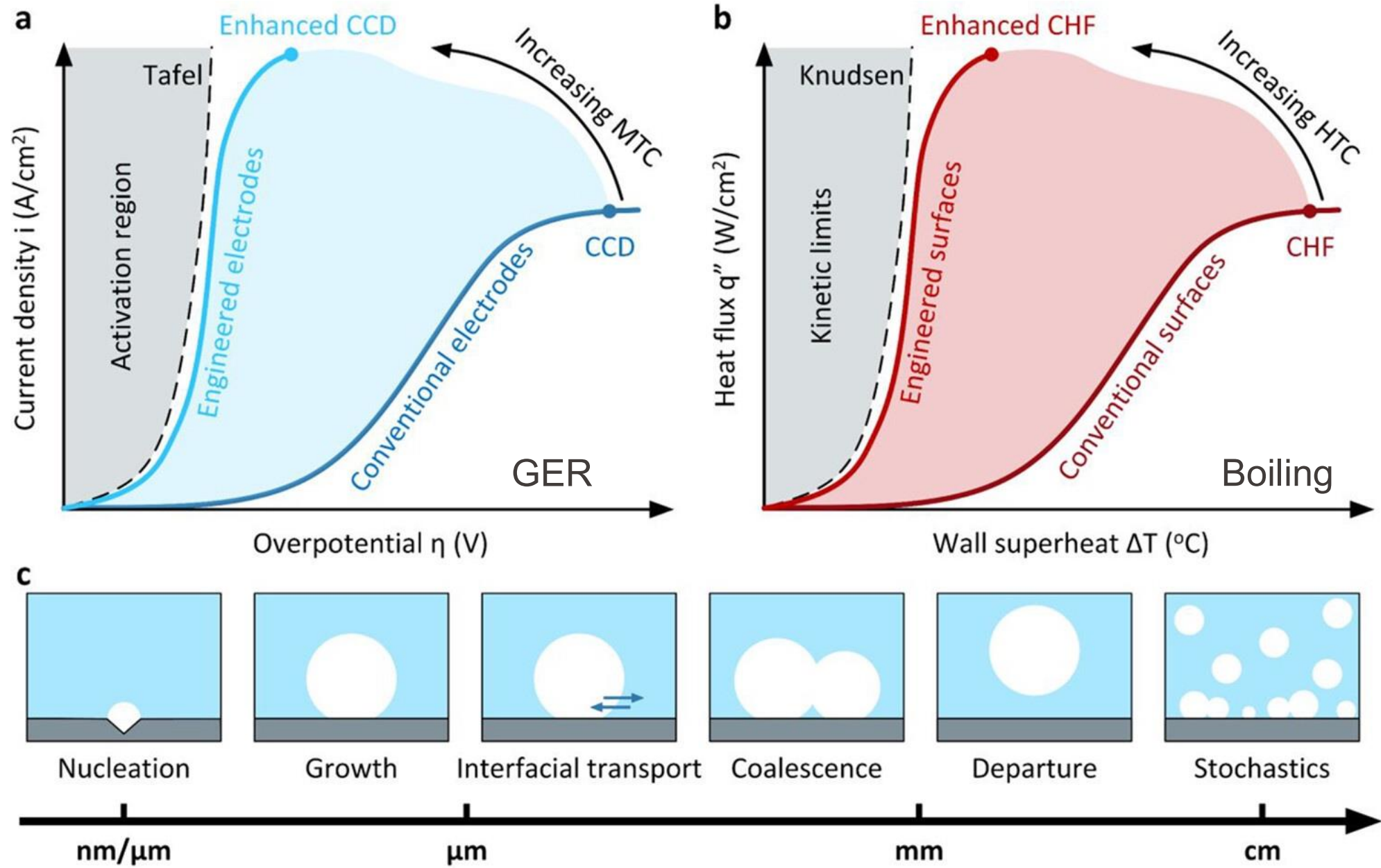


Additional resistance for ion transport

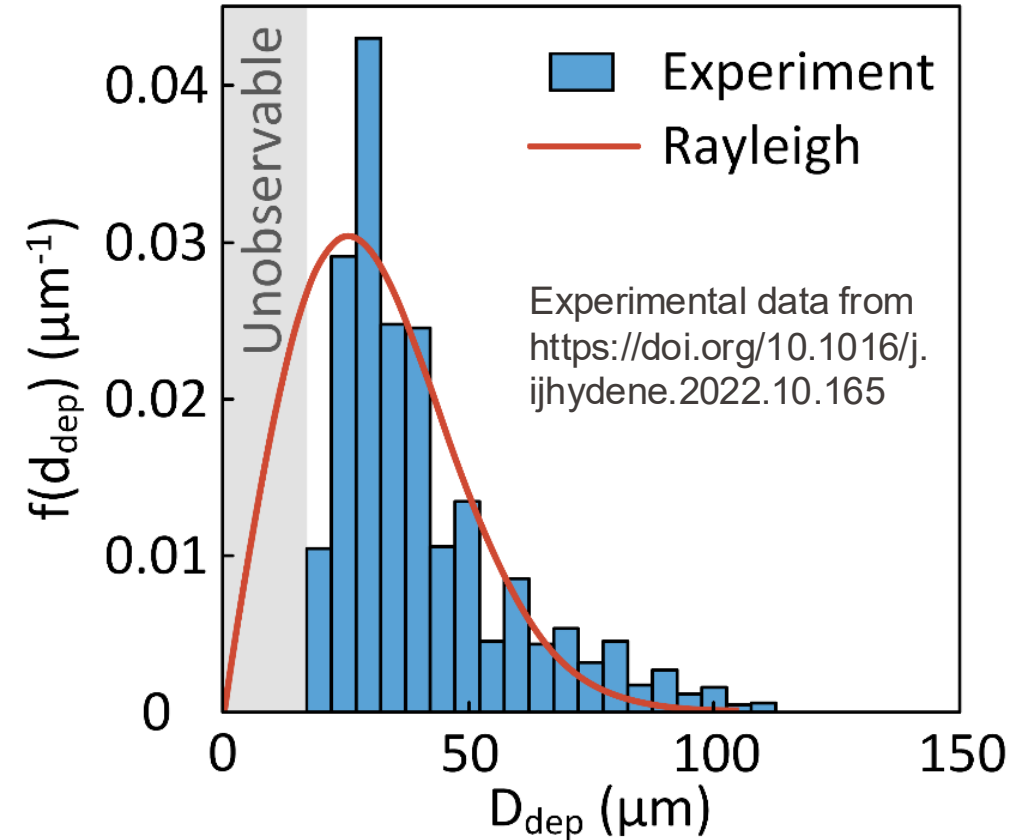
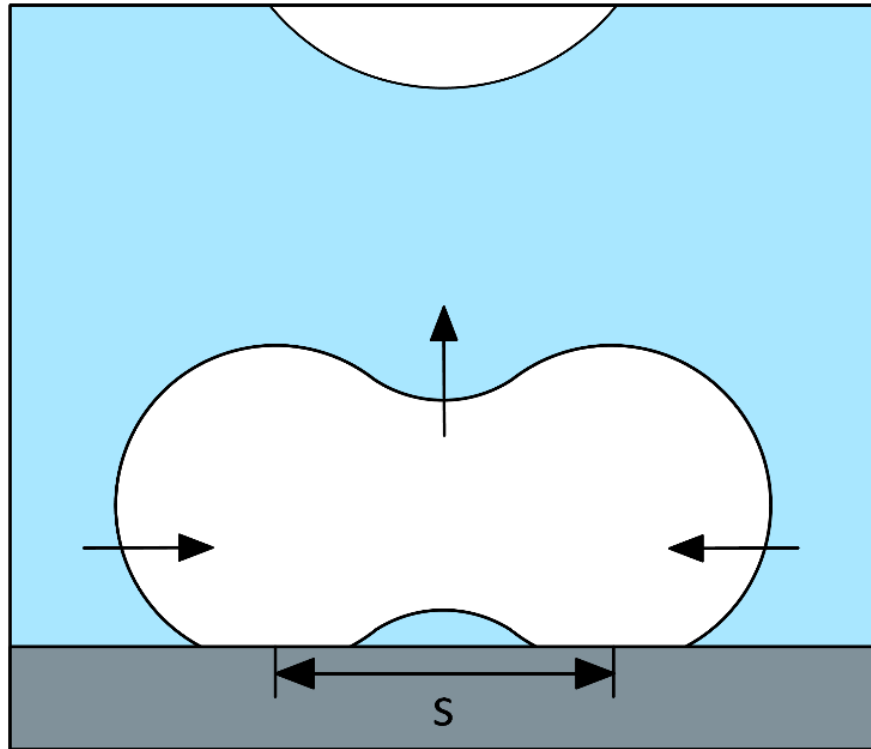


Additional diffusion resistance  
Bubble-induced convection





# Coalescence-Driven Bubble Departure



- When two bubbles coalesce into one, surface area becomes smaller. Excess surface energy needs to be released.
- This may cause bubble to depart immediately after contacting its nearest neighbor