

CE407 SEPARATIONS

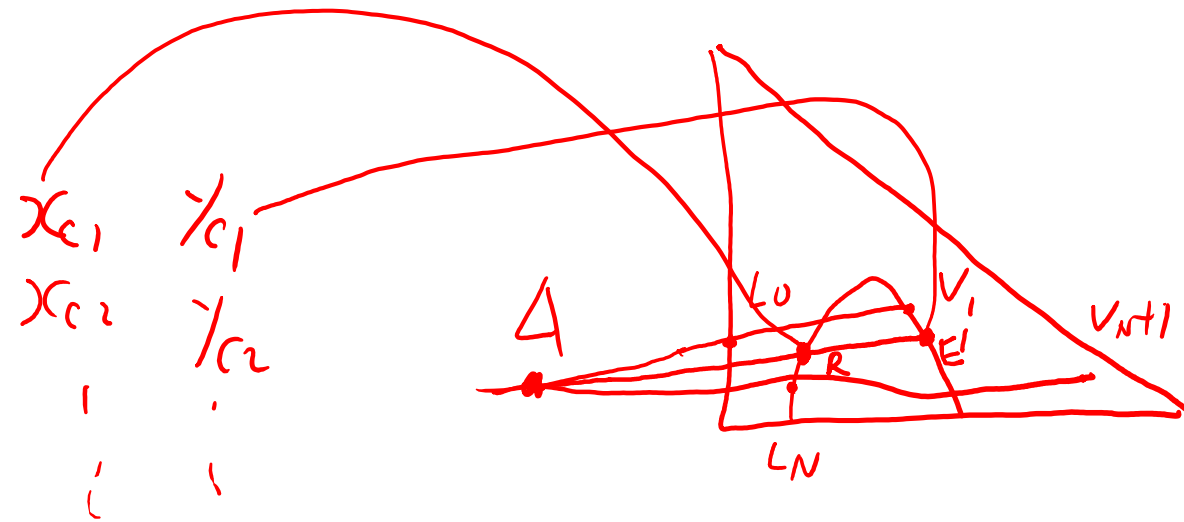
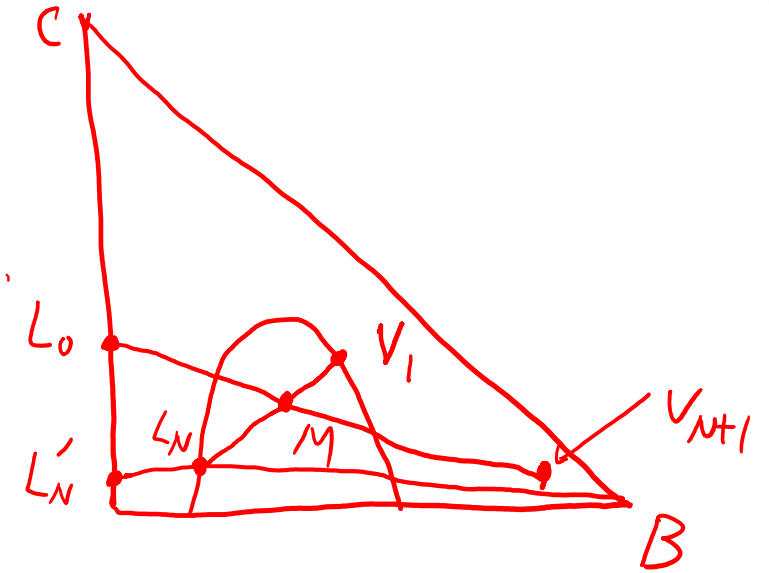
Lecture 17

Instructor: Miao Yu



No. of ideal stages

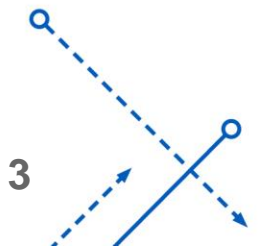
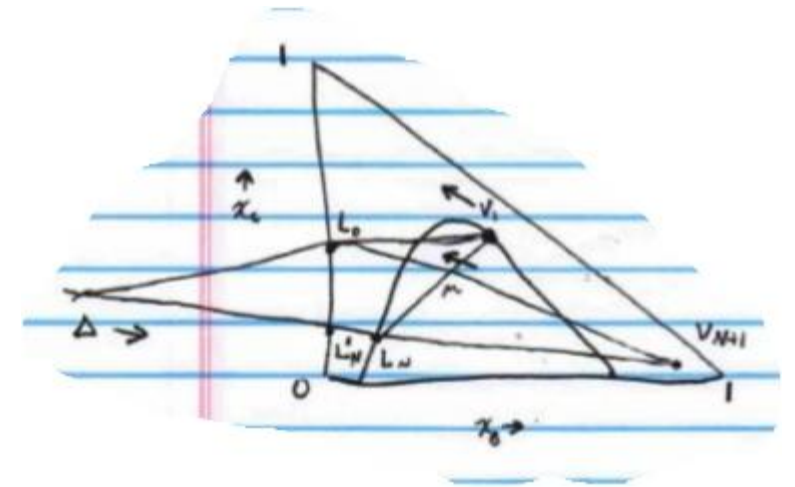
- ① Locate given points L_0, V_{N+1}, L'_N
- ② Find M, L_N
- ③ Find V_1
- ④ Find Δ
- ⑤ Find points on OP
- ⑥ Plot EQ, OP
- ⑦ No. of stages



Multi-Stage Countercurrent Extraction Treybal pp. 452 Fig 1040

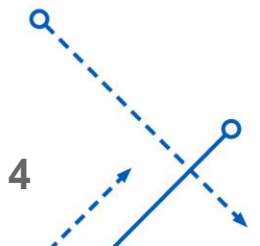
Minimum Entering Solvent Flow

- So far we have started with a given solvent flow, now we will see how to determine a reasonable flow
- **Point #1**
- Revisit the diagram for locating mixing point, **M**
- As the amount of solvent DECREASES
 - “**M**” moves toward L_0
 - V_1 moves to the left
 - Δ will move to the right
 - The line $\overline{V_1 L_0}$ becomes steeper



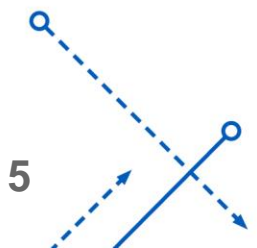
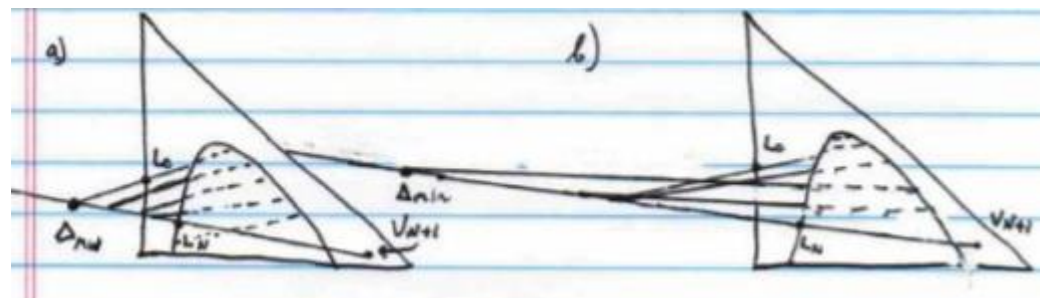
Minimum Entering Solvent Flow

- **Point #2: Review Hunter-Nash method**
- The # of steps are determined by alternating between:
 - Using Δ lines to do mass balances
 - Using tie lines to establish EQ relationships
- When the slopes of the Δ lines and tie lines are very different we make a lot of progress with each step
 - Similar to when OP lines and EQ curve are far apart
- When the slope of a Δ line is the same as the slope of a tie line we stop making progress
 - This is a pinch point
- The infinite number of steps corresponds to minimum solvent flow



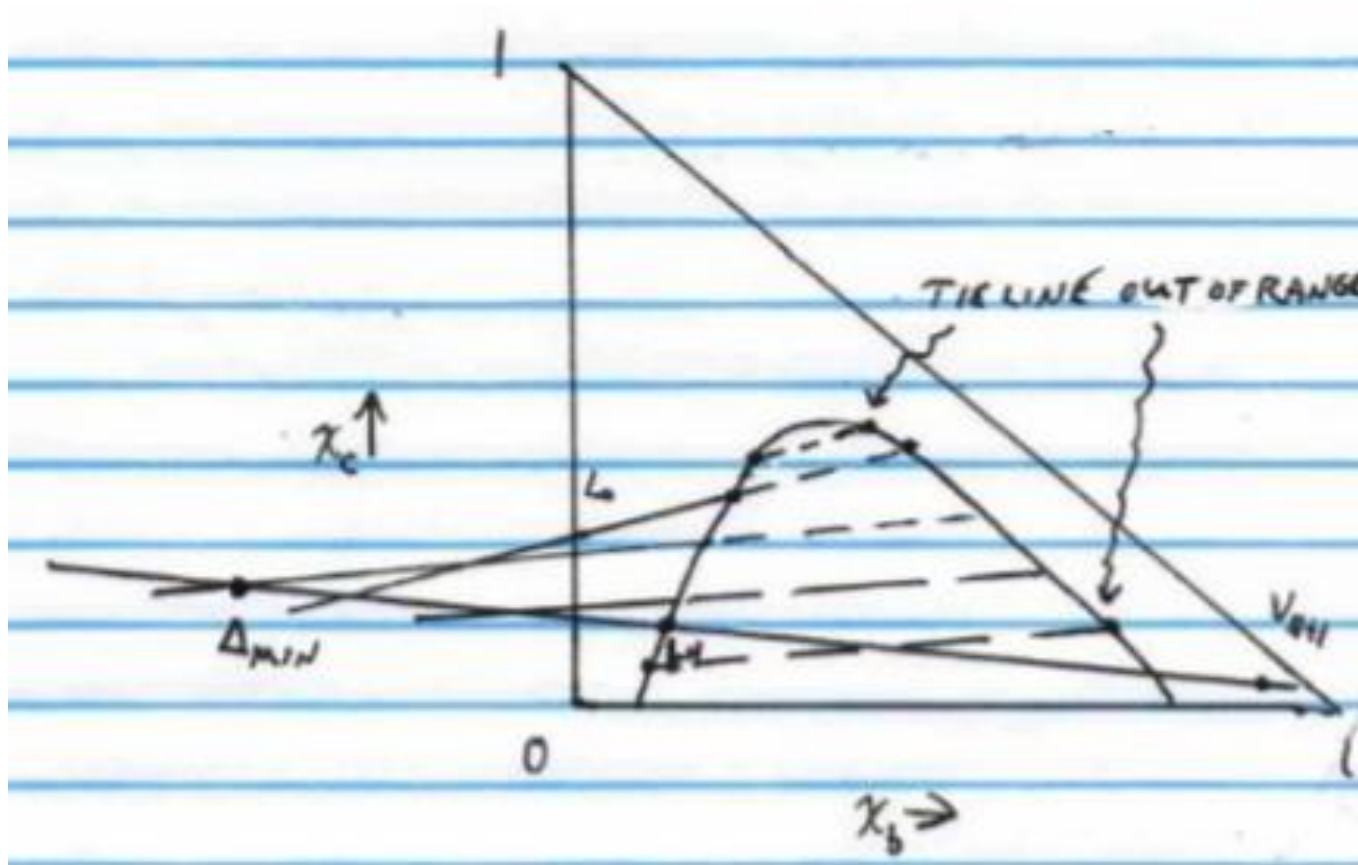
Minimum Entering Solvent Flow

- If we extend all of the relevant tie lines we see which leads to the furthest Δ location
 - Relevant tie lines are the those located between the tie line that passes through L_0 and the one that passes through L_N
- The Δ location furthest left corresponds to the largest flow that leads to a pinch point – this is the Minimum Solvent Flow
 - Note that all smaller flows will have a pinch point, we are looking for one where you reach the point where there are no more pinch points
 - When Δ lies to left of triangle it is furthest out, when Δ lies to right of triangle it is closest
- If the tie lines all have similar slopes this will be the tie line that crosses at L_0 (Fig a)
- If the slopes vary, it could be a different tie line (Fig b)



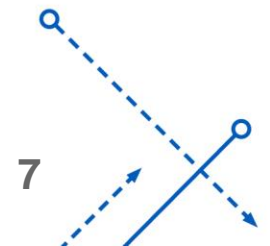
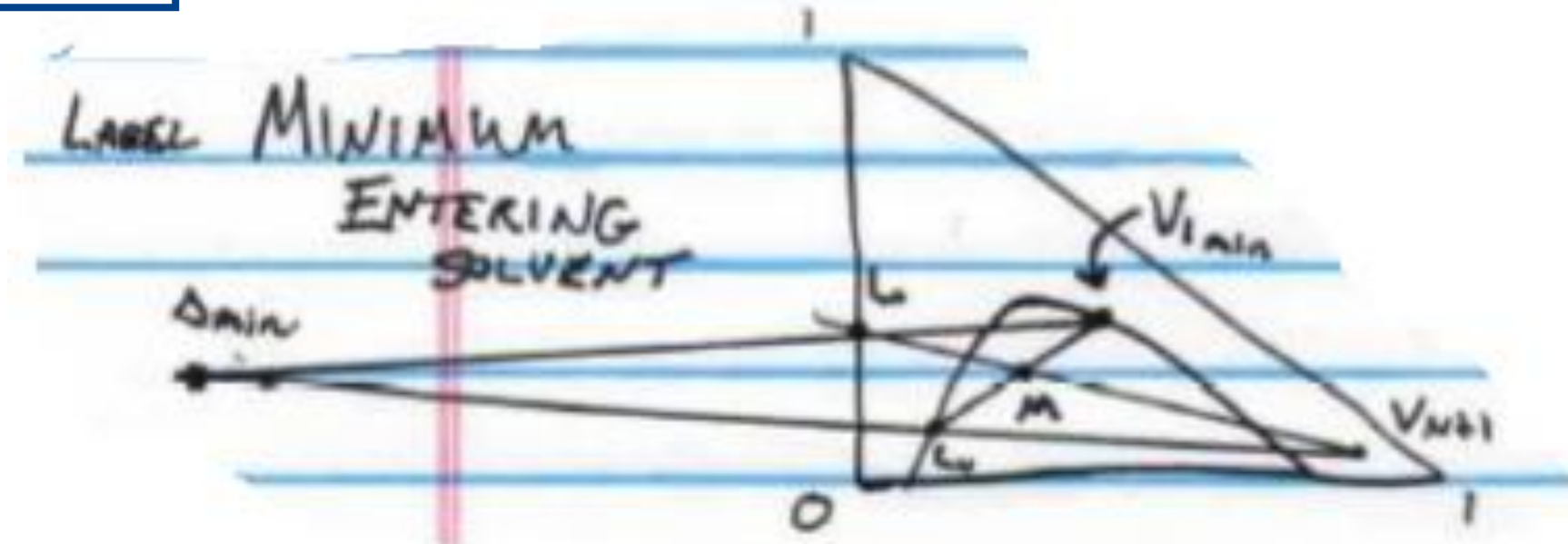
Minimum Entering Solvent Flow

- Label Leftmost intersection as Δ_{\min}
- Notice the tie lines that are out of range are not used



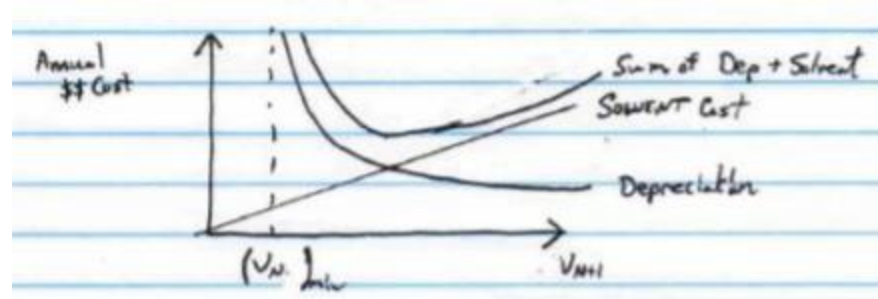
Minimum Entering Solvent Flow

- Now draw a line from Δ_{\min} to L_0 and extend it to right hand side of phase boundary
- This determines $V_{1,\min}$
- Note that this line is NOT necessarily a tie line
- Draw in $\overline{L_N V_{1,\min}}$ and $\overline{L_0 V_{N+1}}$, their intersection determines **M**
- $\frac{(V_{N+1})_{\min}}{L_0} = \frac{x_0 - x_M}{x_M - y_{N+1}}$ this gives the ratio of minimum solvent flow to feed flow



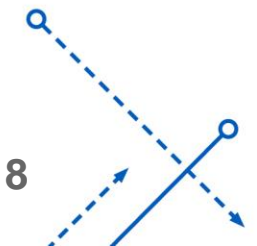
What Flow Should We Use?

- Same optimization as we did for other Unit Operations...
- Annual Cost = Depreciation + Solvent Cost

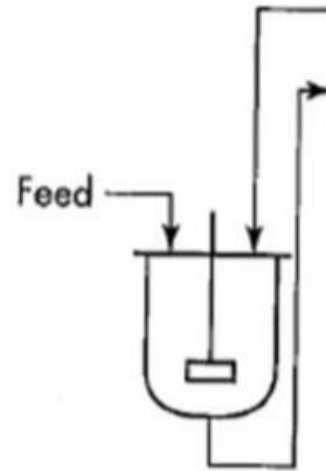


- Once again it turns out that it typically reaches a minimum at

$$(V_{N+1})_{opt} = 1.3(V_{N+1})_{min}$$

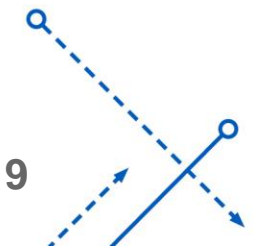


Batch Operation of a Stage



Think in terms of “before” and “after” the mixing and settling

- Charge Feed and Solvent to Vessel
- Mix thoroughly – need proper hold time
- Stop agitation and let phase settle
- Aqueous phase is more dense and will be on bottom
- Drain material from bottom of vessel
 - First material is aqueous phase
 - Switch to another receiver when organic phase starts to come out



Continuous Operation

Mixer-settlers

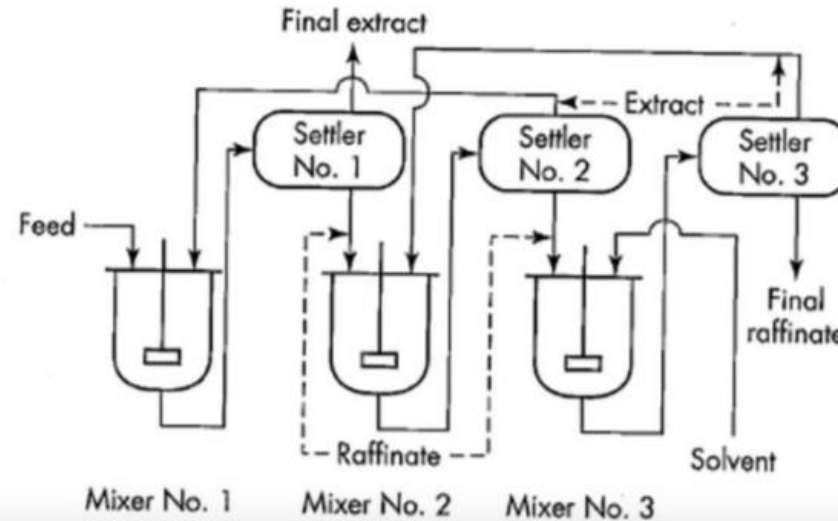
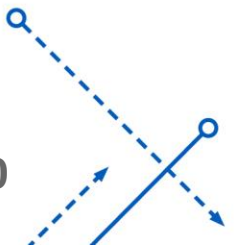


FIGURE 23.4

Think in terms of “flow in” and “flow out” each mixing and settling stage

- Continuous flow of Feed and Solvent to Mixing Vessel
- Mix thoroughly – need proper residence time
- Mixture is continuously flowing to settler
- The two phases separate in settler and exit as two streams



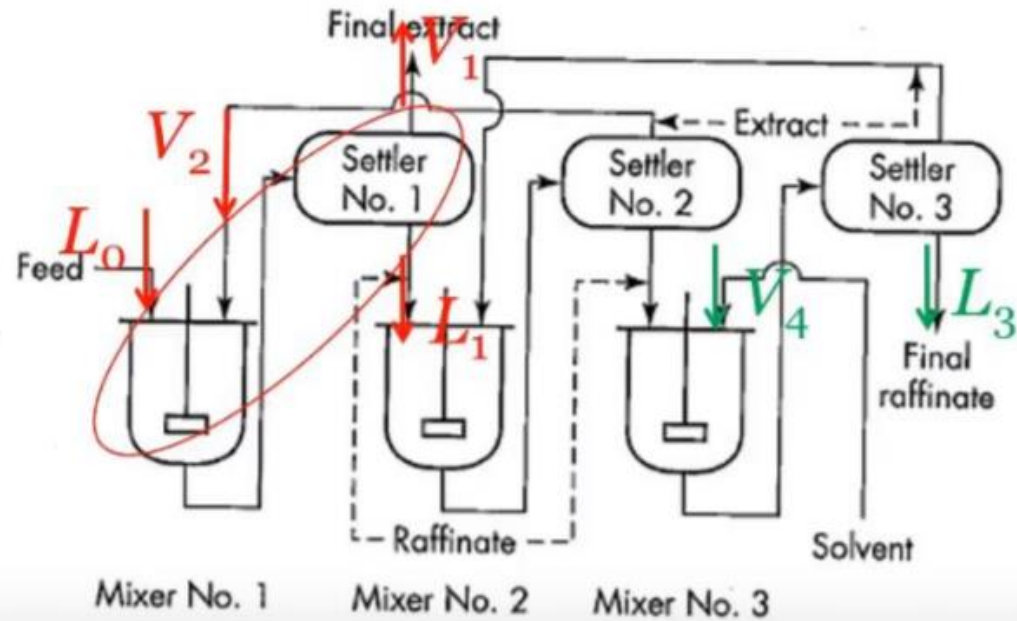
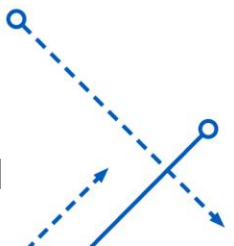


FIGURE 23.4

- Ellipse represents Stage 1
- L_0 is Feed into Stage 1
- V_2 is extract from Stage 2 feeding into Stage 1
- L_1 is raffinate flow leaving Stage 1
- V_1 is extract flow leaving Stage 1
- V_4 is solvent flow entering Stage 3, ie V_{N+1}
- L_3 is final raffinate flow exiting Stage 3, ie L_N



Packed and plate towers

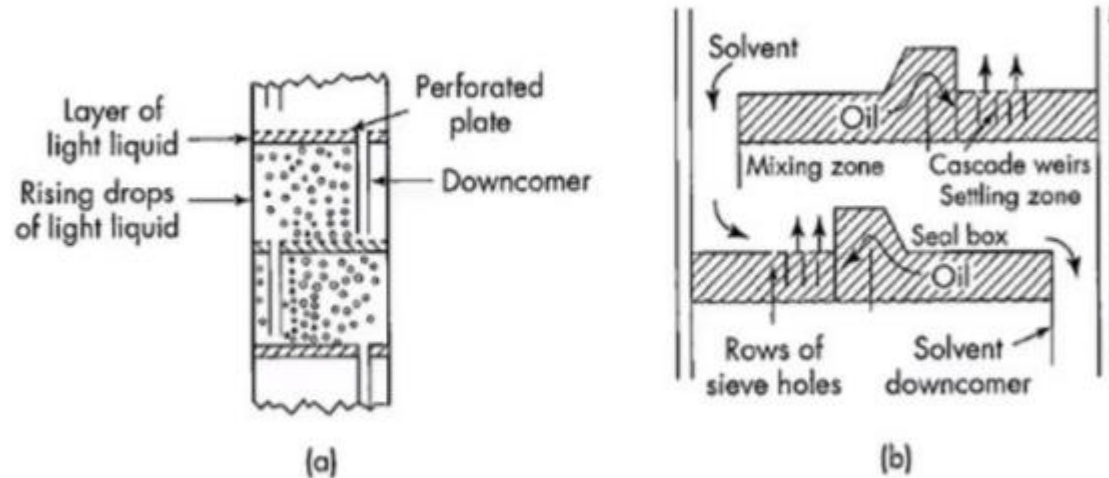


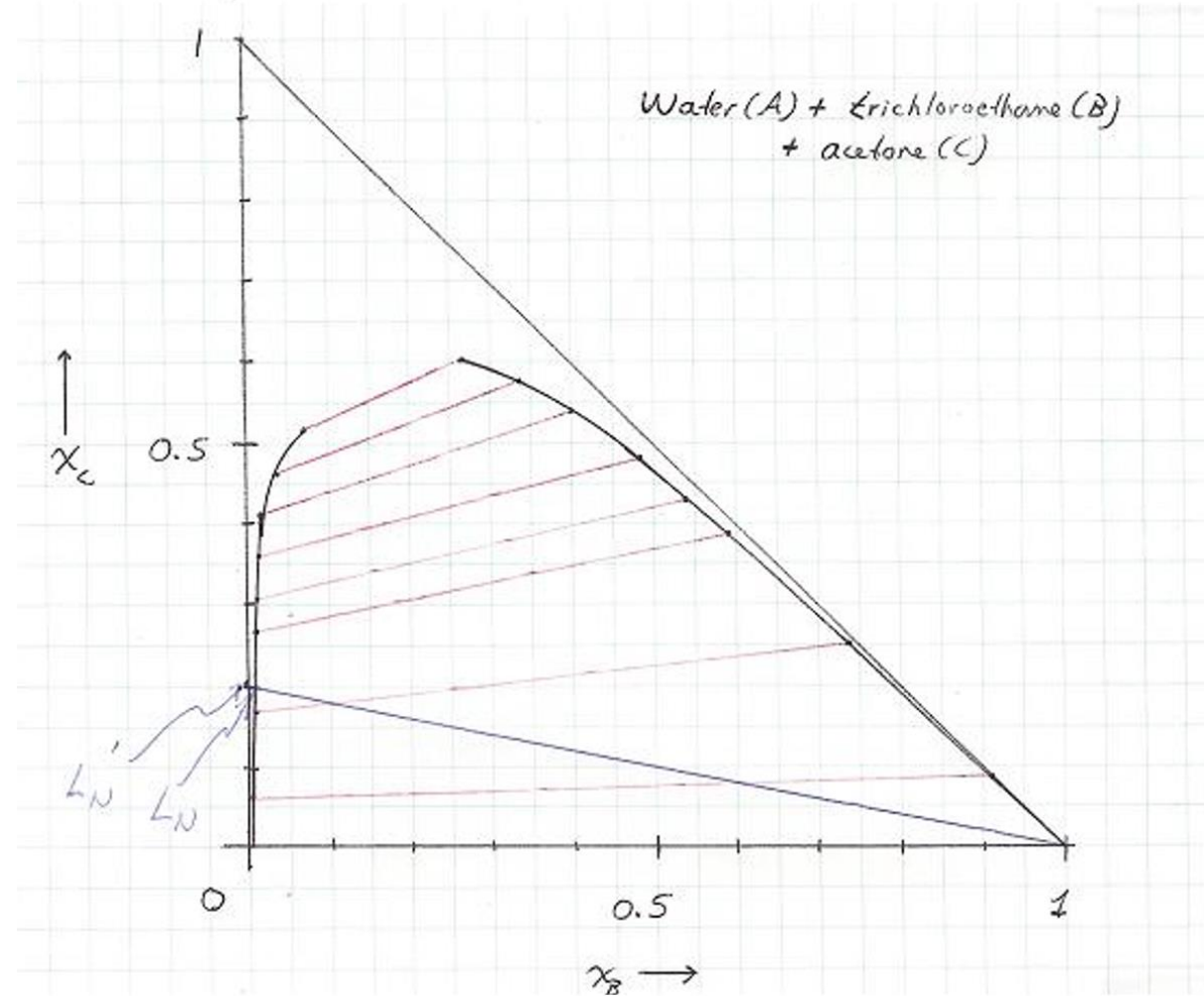
FIGURE 23.5
 Perforated-plate extraction towers: (a) perforations in horizontal plates;
 (b) cascade weir tray with mixing and settling zones. (After Bushell and
 Fiocco.⁴)

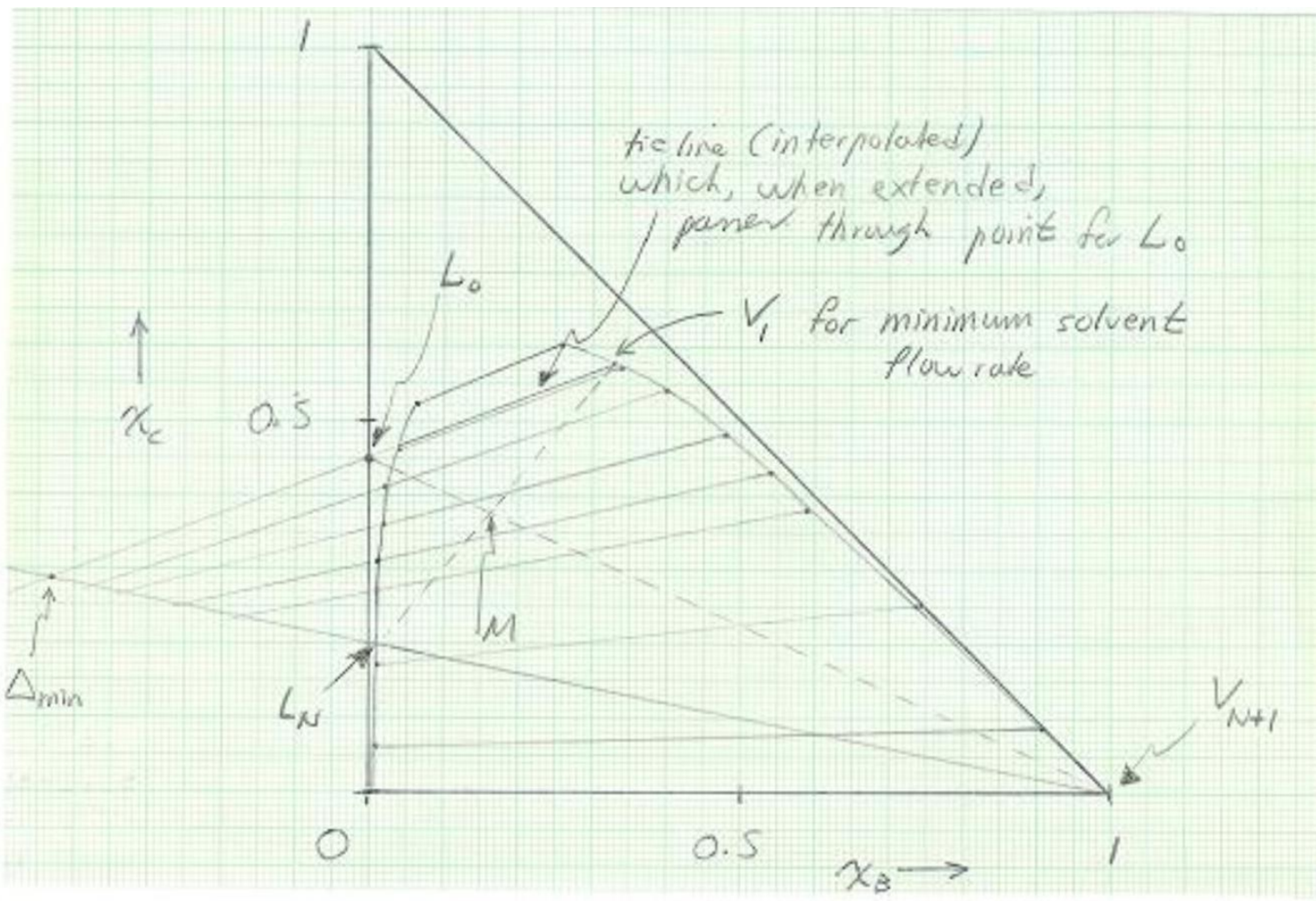
Very similar to the trays we have discussed in a Distillation Column

- Density difference is orders of magnitude lower than in a rectifying gas/liquid column (sp gr of 1 for aqueous and around 0.7 for organic)
- Both phases will be relatively high viscosity as opposed to the low viscosity vapor phase in distillation (velocities will be lower than in distillation column)
- Aqueous phase is more dense and will travel downward, organic phase will travel upward
- This means to location of the extract leaving the column depends on whether the extract is the aqueous phase or whether it is the organic phase

3. A 500 kg/h feed stream with composition 45 mass % acetone (solute, C) and 55 mass % water (diluent, A) is to be contacted with trichloroethane (solvent, B) in a countercurrent liquid extraction battery. Entering trichloroethane is pure. The exiting raffinate should contain 20.2 mass % acetone on a trichloroethane-free basis. Our very good friend Elroy poses the following two questions:

- (a) What is the minimum flow rate of trichloroethane required to achieve the desired composition of the exiting raffinate (corresponding to an infinite number of stages)?





$$\begin{aligned}
 \frac{(V_{N+1})_{\min}}{L_0} &= \frac{x_0 - y_m}{y_m - y_{N+1}} \\
 &= \frac{0.45 - 0.38}{0.38 - 0} = 0.194
 \end{aligned}$$

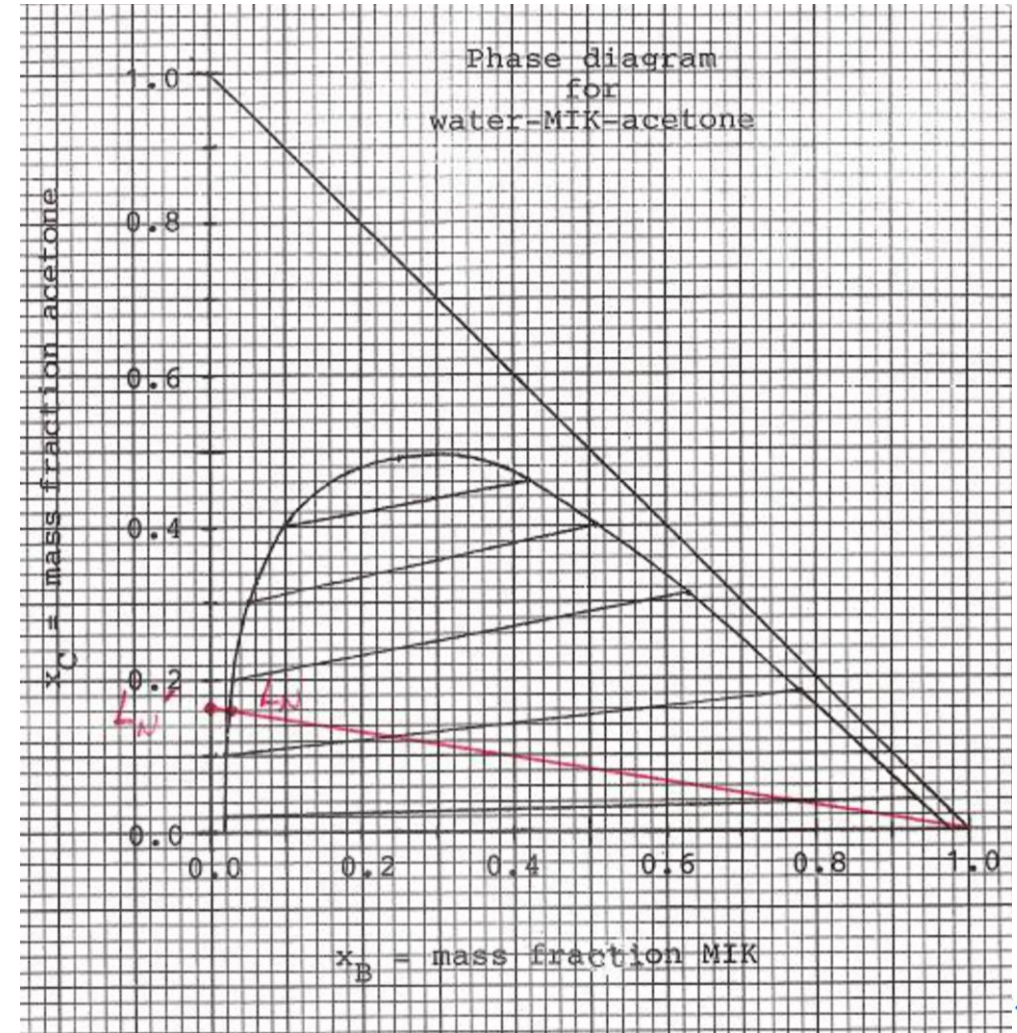
$$L_0 = 500$$

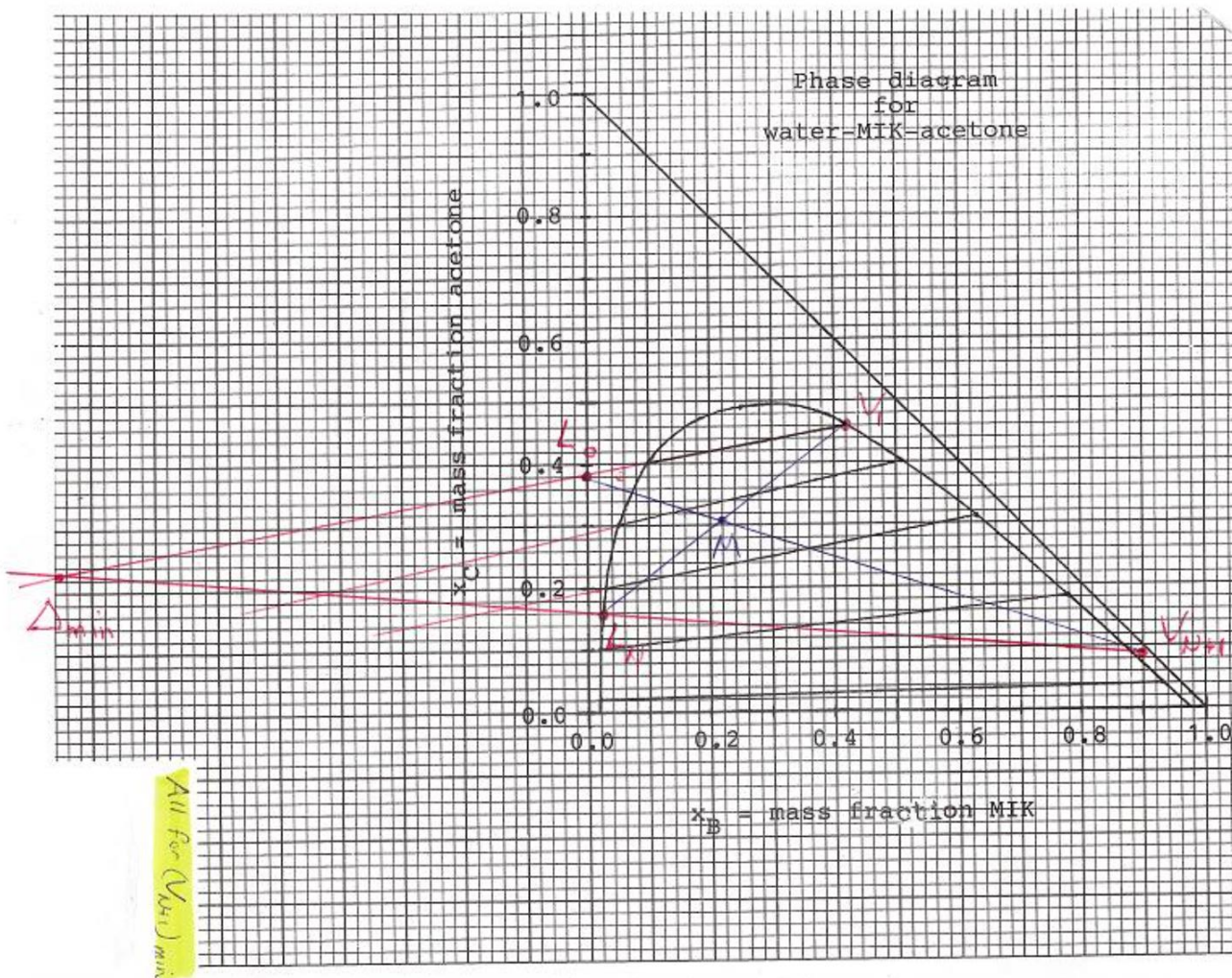
$$(V_{N+1})_{\min} = 96 \text{ kg/h}$$

$$\begin{aligned}
 V_{N+1} &= 1.3 \times (V_{N+1})_{\min} \\
 &= 125
 \end{aligned}$$

5. A 450 kg/h feed stream with composition 38 mass % acetone (solute, C) and 62 mass % water (diluent, A) is to be contacted countercurrently with an MIK(solvent B)-rich solution of which the precise composition is 90 mass % MIK, 9 mass % acetone and the balance water. The exiting raffinate should contain 16.5 mass % acetone (C) and 83.5 mass % water (A) on an MIK(B)-free basis.

(a) What is the minimum flow rate $(V_{N+1})_{\min}$ of the entering MIK-rich solvent stream required to achieve the desired separation (corresponding to an infinite number of stages)?





$$\frac{(V_{N+1})_{min}}{L_0} = \frac{x_{L_0} - x_M}{x_M - y_{N+1}}$$

$$= \frac{0.38 - 0.31}{0.31 - 0.09} = 0.33$$

$$(V_{N+1})_{min} = 450 \times 0.33 \approx 150$$

$$V_{N+1} = 1.3 \times 150 = 195$$