

The continuous production of bioethanol: One, two and three tank reactor designs

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Talk Outline

- Brief background.
- Model description.
- Results — dynamical & optimal performance.
- Conclusion and future work.

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Background — bioethanol

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- Reduced emissions — greenhouse gas emissions reduced by up to 19%, tailpipe carbon monoxide by as much as 30%
- It takes only six months to harvest a substantial crop for fuel.

Model — biochemistry

A.B. Jarzebski. (1992). “Modelling of oscillatory behaviour in continuous ethanol fermentation”. *Biotechnology Letters*, **14**(2), 137–142.

- Substrate (S).
- Product - ethanol - (P)
- Biomass (*Zymomonos mobilis*)
 - Viable cells (X_v)
 - Non-viable cells (X_{nv}) — non-growing, but still retain the ability to produce ethanol.
 - Dead cells (X_d) — eqn uncouples.

Model — single reactor

$$V \frac{dS}{dt} = F(S_0 - S) - V \left(\frac{\mu_v X_v}{Y_{x|s}} + m_s X_{nv} \right),$$

$$V \frac{dX_v}{dt} = -F X_v + V(\mu_v - \mu_{nv} - \mu_d) X_v,$$

$$V \frac{dX_{nv}}{dt} = -F X_{nv} + V(\mu_{nv} X_v - \mu_d X_{nv}),$$

$$V \frac{dX_d}{dt} = -F X_d + V \mu_d (X_v + X_{nv}),$$

$$V \frac{dP}{dt} = -F P + V \left(\frac{\mu_v X_v}{Y_{x|p}} + m_p X_{nv} \right).$$

Model — rate expressions

$$\mu_v = \mu_{\max} \frac{S}{K_1 + S} \left(1 - \frac{P}{P_c} \frac{S}{K_2 + S} \right),$$

$$\mu_d = -\mu_{\max} \frac{S}{K_1 + S} \left(1 - \frac{P}{P_c} \frac{S}{K_2 + S} \right),$$

$$\mu_{nv} = \mu'_{\max} \frac{S}{K_1 + S} \left(1 - \frac{P}{P'_c} \frac{S}{K_2 + S} \right) - \mu_v.$$

- Reaction rates can not be negative.
- Substrate limitation.
- Product inhibition.

Results — ethanol concentration

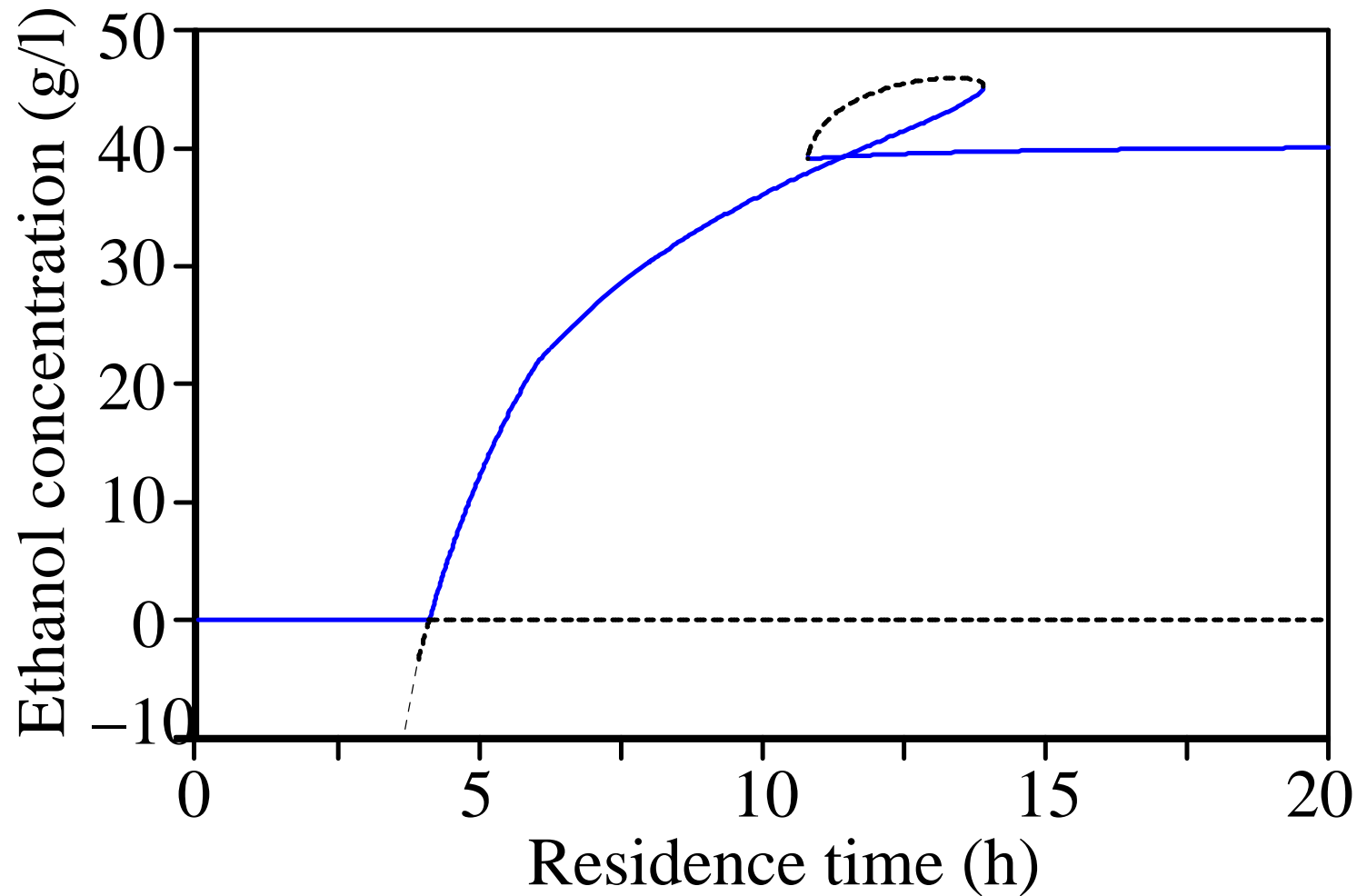


Figure 0: $S_0 = 100 \text{ g l}^{-1}$. $\tau_W = 4.12 \text{ hr}$.

Results — productivity ($\text{Pr} = P/\tau$)

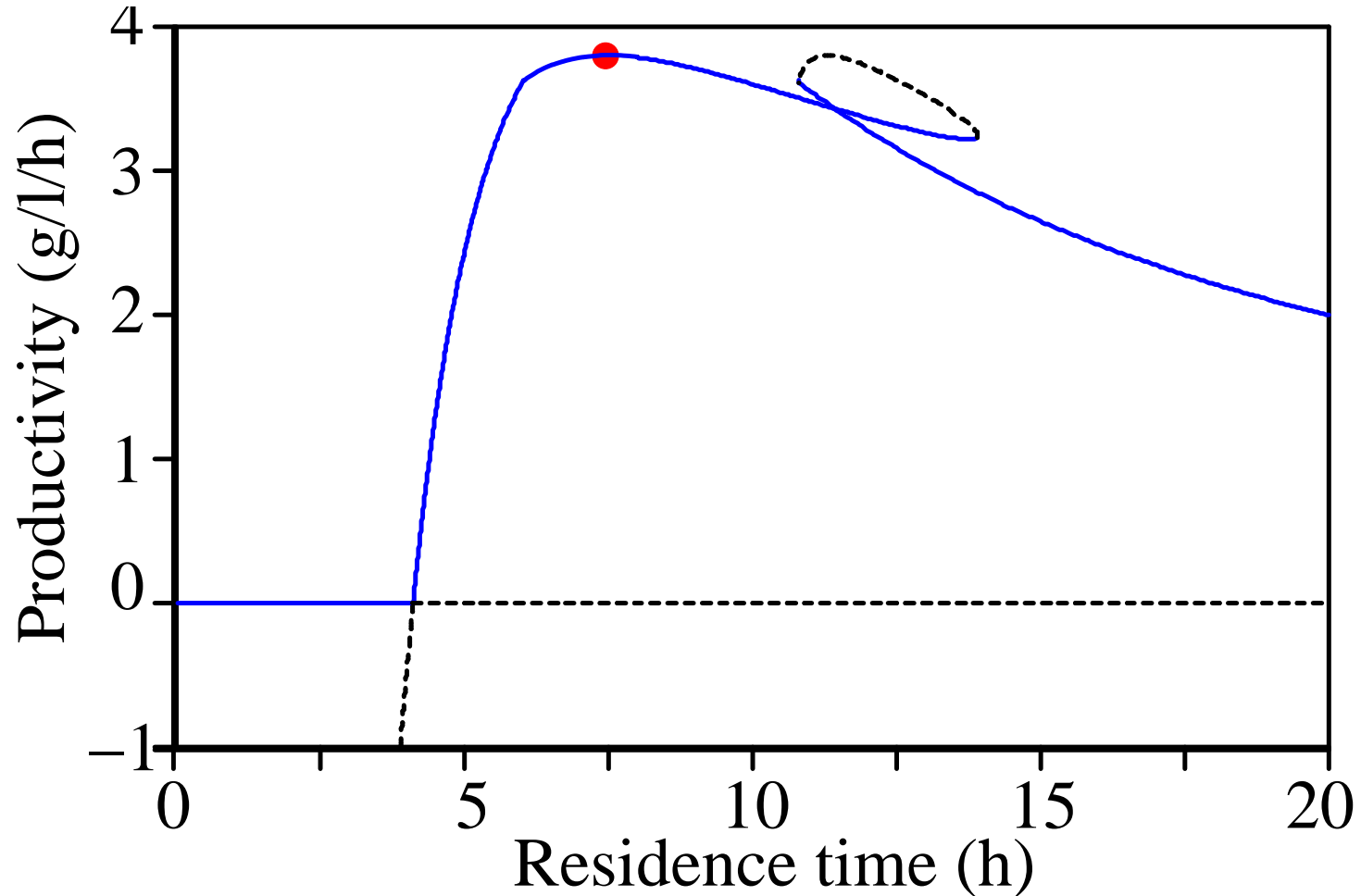


Figure 0: $S_0 = 100 \text{ g l}^{-1}$. $\text{Pr}_{\max} = 3.8 \text{ g l}^{-1} \text{ hr}^{-1}$ at $\tau = 7.5 \text{ hr}$.

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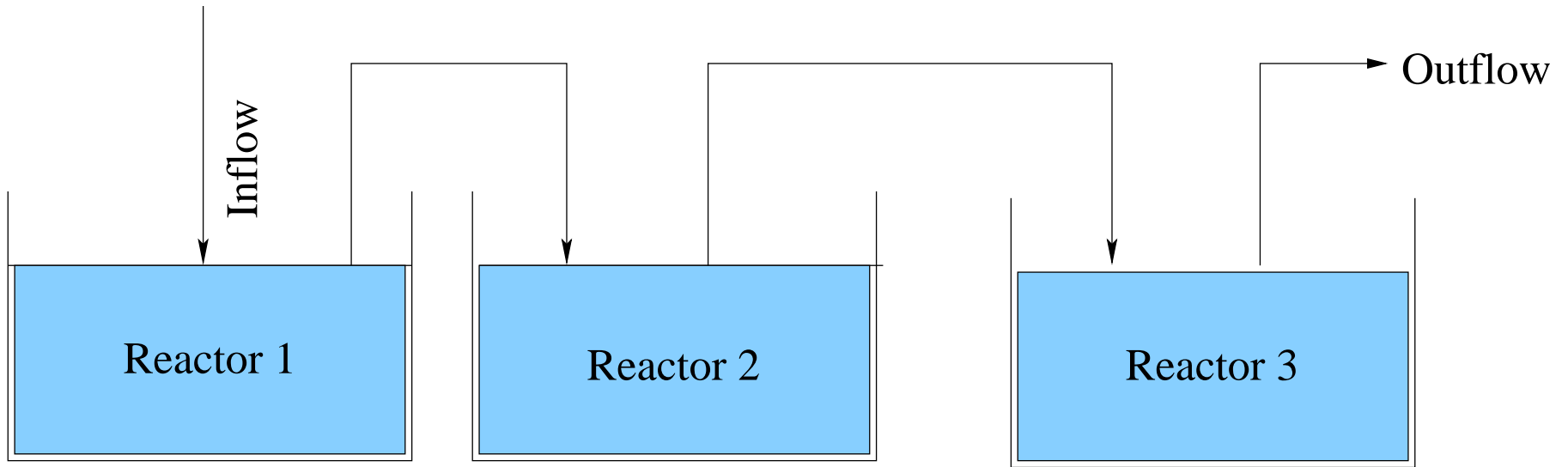
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- Maximum productivity: system operated at a static steady-state.
- If $100 \leq S_0 \text{ (g l}^{-1}\text{)} \leq 160$ then

$$Pr_{\max} = 3.8039 \text{ (g l}^{-1}\text{hr}^{-1}\text{)} \pm 9.8738 \times 10^{-5}$$

Results — reactor cascade



$$Pr_2 = \frac{P_2}{\tau_1 + \tau_2}$$

$$Pr_3 = \frac{P_3}{\tau_1 + \tau_2 + \tau_3}$$

Results — double reactor cascade

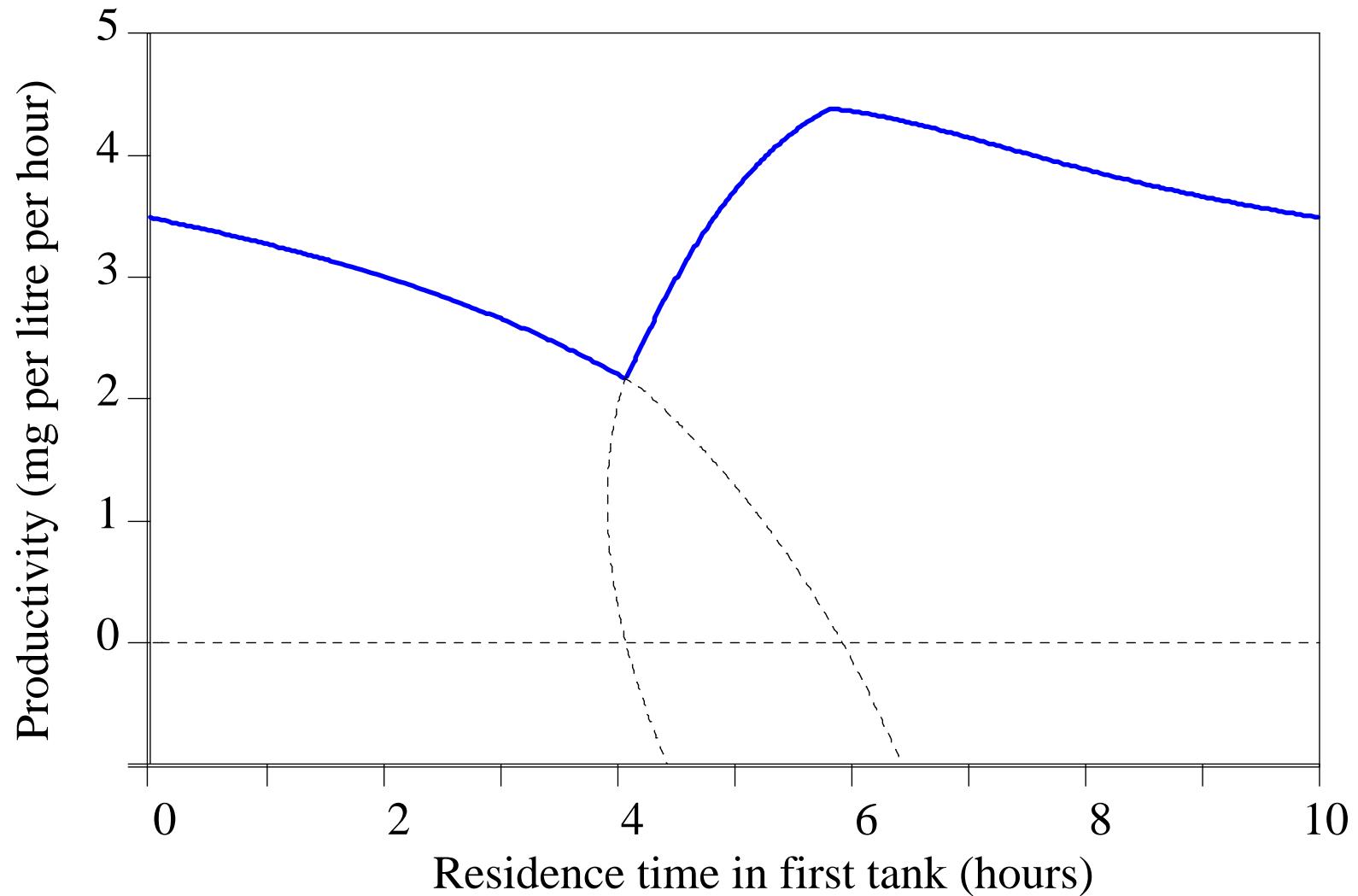


Figure 0: Total residence $\tau_1 + \tau_2 = 10$ (h). $S_0 = 100 \text{ g l}^{-1}$

Results - optimal double reactor cascade

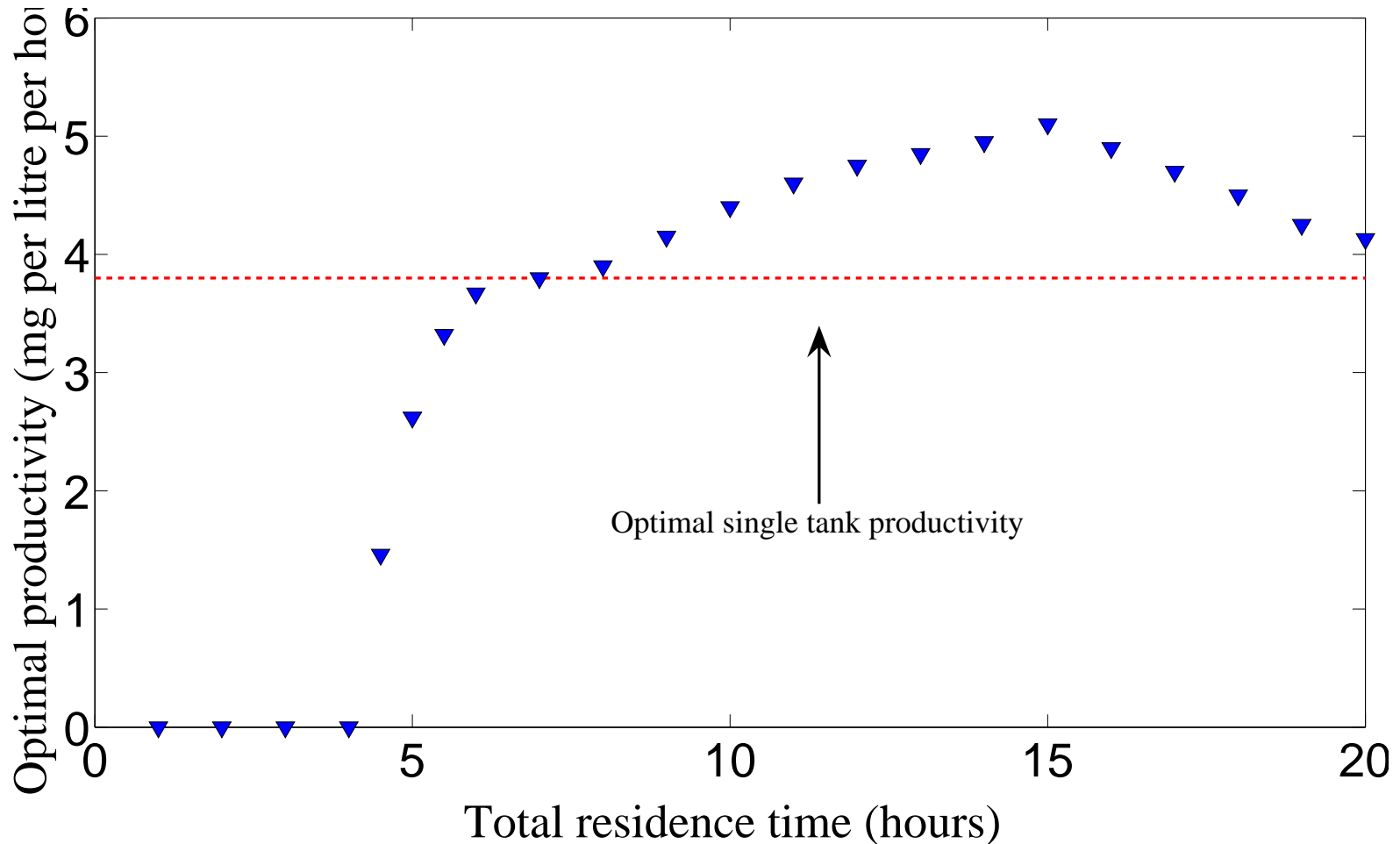


Figure 0: Optimal performance of a cascade of two (un)equal reactors.

$$S_0 = 100 \text{ g l}^{-1}.$$

Results — scatter plot

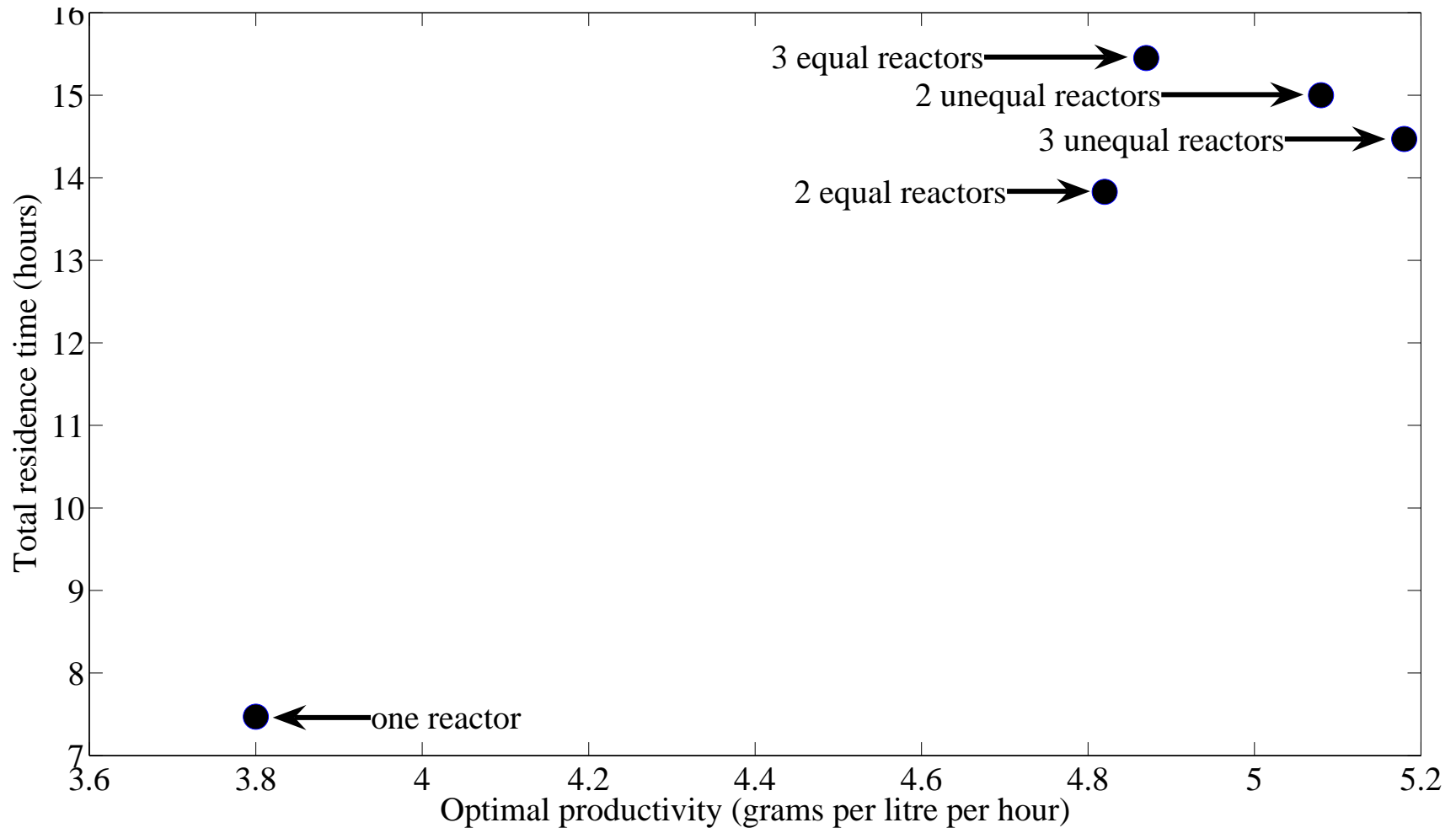


Figure 0: Optimal performance of a cascade of various reactor configurations. $S_0 = 100 \text{ g l}^{-1}$.

What's next?

Feed concentration : Extensive investigation.

Reactor costs : Can we afford to optimise production?

Recycle : Improves performance.

Performance : Maximise product concentration.

Undergraduate lab : Department of Chemical Engineering,
Monash University.

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6. Equal double reactor cascade can outperform single reactor by 27%.

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7. Equal triple reactor cascade can outperform single reactor by 28%.