Continuous chiral resolution by diastereomeric salt formation of racemic Ibuprofen in a Couette-Taylor crystallizer

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Introduction

Batch production mode [1]

- legacy in pharmaceutical industry
- weaknesses such as batch-to-batch quality variation

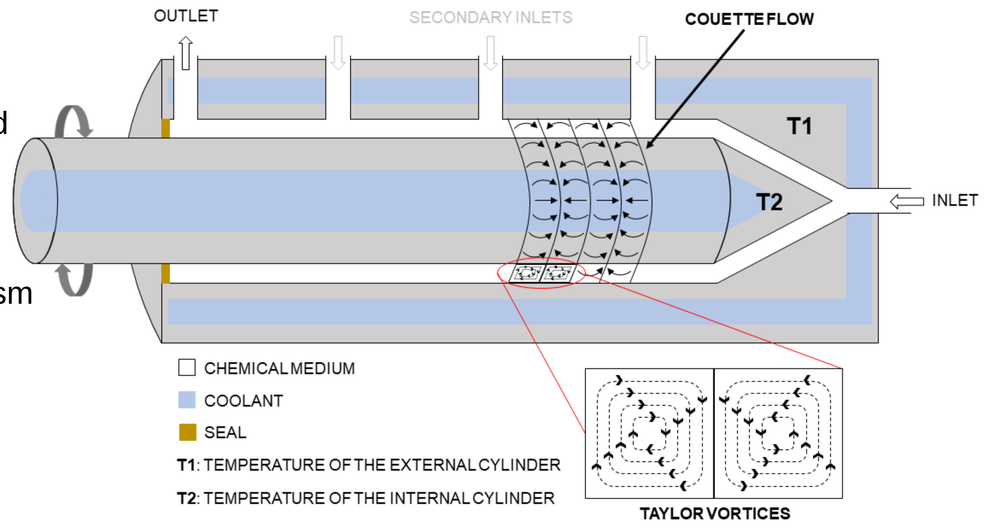
Continuous production mode [2]

- steady-state functioning, i.e., more constant quality product
- better process control

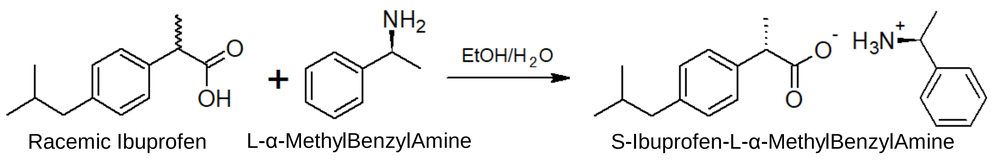
Couette-Taylor crystallizer

- specific flow characteristics at high rotation speed: Couette flow coupled with Taylor vortices [3]

- impact on crystal size distribution (CSD), morphology and polymorphism in crystallization processes [4]

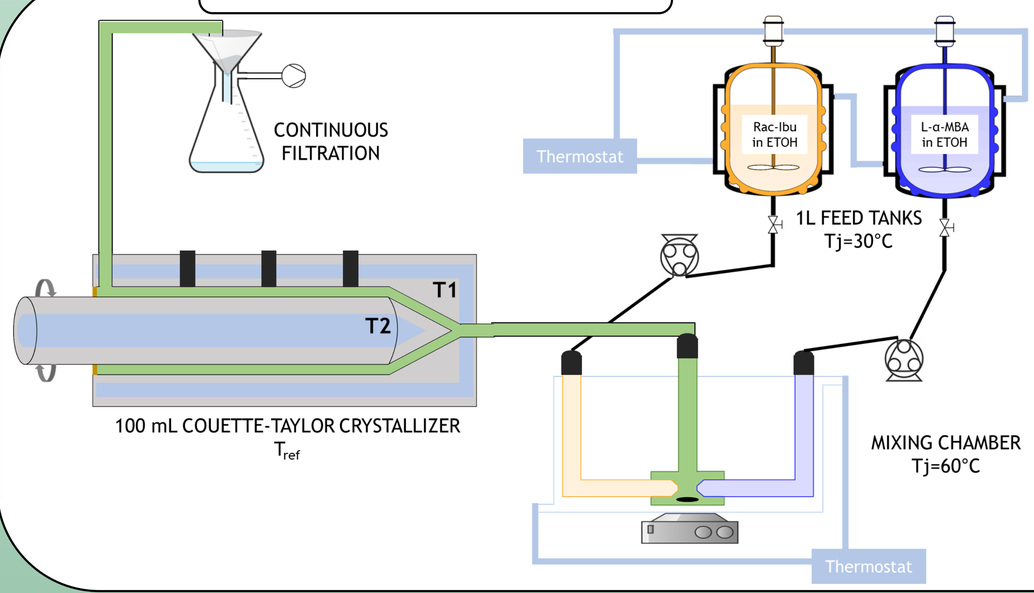


Reaction scheme



Racemic Ibuprofen + L-α-MethylBenzylAmine → S-Ibuprofen-L-α-MethylBenzylAmine

Continuous process



7 parameters to investigate

(1) Absolute temperature difference |ΔT|

(2) Sign of ΔT

(3) Stirring speed Ω

(4) Residence time t (i.e., flowrate)

(5) Temperature inside the CT crystallizer ("central point" of the two coolant set temperatures) T_{ref}

(6) Medium dilution

(7) EtOH/H_2O mass ratio

⇒ **Rationalized study through a Design of Experiments (DoE)**

First screening with 4 factors

(1)(2) Temperature difference ΔT

(3) Stirring speed Ω

(4) Residence time t (i.e., flowrate)

(5) Temperature inside the CT crystallizer ("central point" of the two cryostat set temperatures) T_{ref}

Fixed medium dilution (15V) and EtOH/H2O mass ratio (80/20)

Design of Experiments

Factor	Levels		
ΔT (°C)	-10 (T1>T2)	0 (T1=T2)	+10 (T1<T2)
Ω (rpm)	200	500	1000
t (min)	5	15	30
T_{ref} (°C)	15	20	25

5 main responses to study

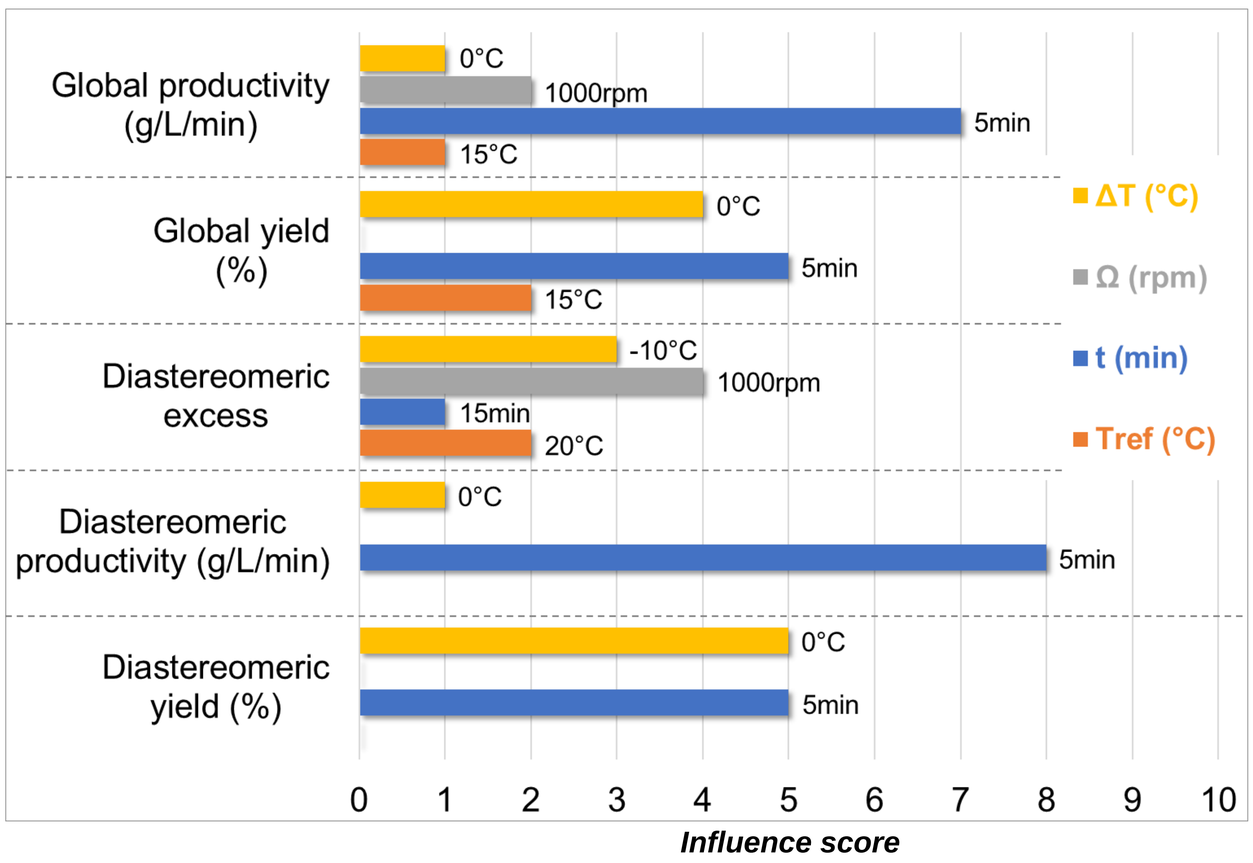
1. Global productivity (g/L/min)
2. Global yield (%)
3. Diastereomeric excess
4. Diastereomeric productivity (g/L/min)
5. Diastereomeric yield (%)

⇒
- **Most impacting factor(s)**
- **Best factor level**

Results

From 16 experiments (including 4 repeated ones) lasting 10 residence times:

Relative influence of each studied factor on the main responses and best factor level for influence score ≥1



Temperature difference ΔT (°C)	Stirring speed Ω (rpm)	Residence time t (min)	Cryostat average set temperature T_{ref} (°C)
⇒ ΔT>0 seems to enhance chiral purity	⇒ 1000rpm seems to be the best factor level	⇒ Favor lower residence times	⇒ Apparent poor influence
⇒ Avoid T1<T2	⇒ 200 and 500rpm eliminated	⇒ Avoid 30min	⇒ Favor lower temperatures

Conclusions and perspectives

⇒ Thanks to the set up of a **Design of Experiments**, **trends** were identified on the 4 studied factors in order to **favor yield, productivity and/or diastereomeric excess**.

⇒ With the **suitable parameter set**, the **chiral purity** of the recovered product is **higher** than that obtained in **batch mode**.

⇒ The **yield** is generally **lower** than that obtained in batch mode. However, experiments performed on a period exceeding **10 residence times** suggest that it **improves after 14 resident times**. **Changes** should be done on the **current set-up** in order to confirm this trend.

⇒ **Further work** should be done on specific ranges determined by this first screening, in order to draw a **response surface** for the **7 parameters** to be studied.

⇒ **Potential interaction(s) between factors** should also be examined, as it has already been seen that the **combination of both Ω and ΔT** can have an influence on chiral purity and crystal size [5].

[References]

[1] Y. Ma et al., Organic process Research & Development, **vol. 24**, n°10, p. 1785-1801 (2020).

[2] D. L. Hughes, Organic Process Research & Development, **vol. 22**, n°1, p. 13-20 (2018).

[3] W.-S. Kim, Journal of Chemical Engineering of Japan, vol. 47, n°2, p. 115-123 (2014).

[4] W.-S. Kim, T. Yu, Z. Wu, U.S. Patent Application 2017/0081188 A1 (2017)

[5] M. Schindler, Thèse de doctorat : physique. Rouen: Université de Rouen Normandie, 2020, 144p.