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

Laureline Marc, Sabrina Guillemer, Jean-Marie Schneider, Gérard Coquerel

► **To cite this version:**

Laureline Marc, Sabrina Guillemer, Jean-Marie Schneider, Gérard Coquerel. Continuous chiral resolution by diastereomeric salt formation of racemic Ibuprofen in a Couette-Taylor crystallizer. ISIC 2021, Aug 2021, Online event, France. hal-03359396

HAL Id: hal-03359396

<https://normandie-univ.hal.science/hal-03359396>

Submitted on 29 Nov 2021

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Introduction

Batch production mode [1]

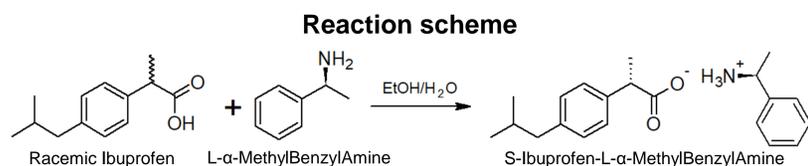
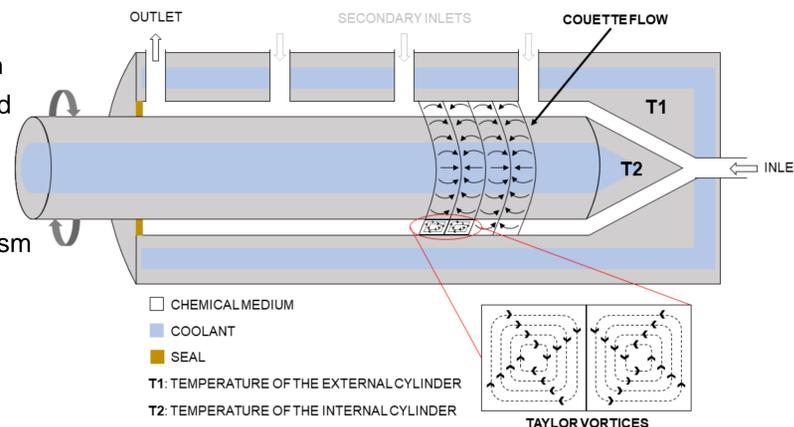
- legacy in pharmaceutical industry
- weaknesses such as batch-to-batch quality variation

Continuous production mode [2]

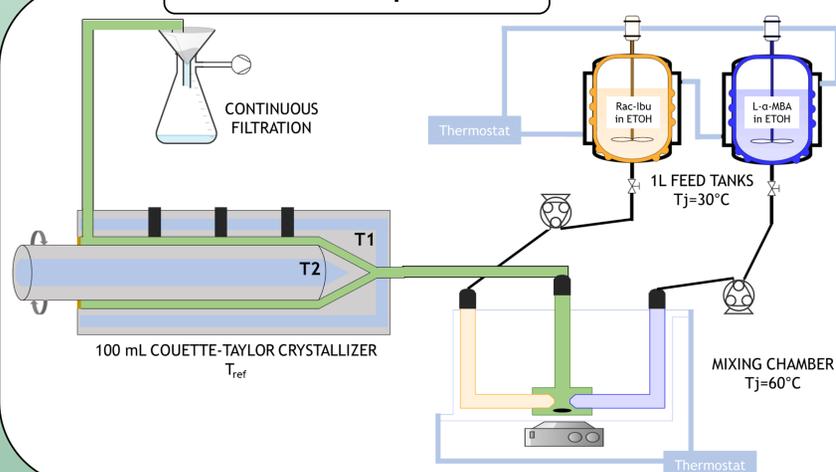
- steady-state functioning, i.e., more constant quality product
- better process control

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]



Continuous process



7 parameters to investigate

- (1) Absolute temperature difference $|\Delta T|$
- (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/H₂O 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 (T ₁ >T ₂)	0 (T ₁ =T ₂)	+10 (T ₁ <T ₂)
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**

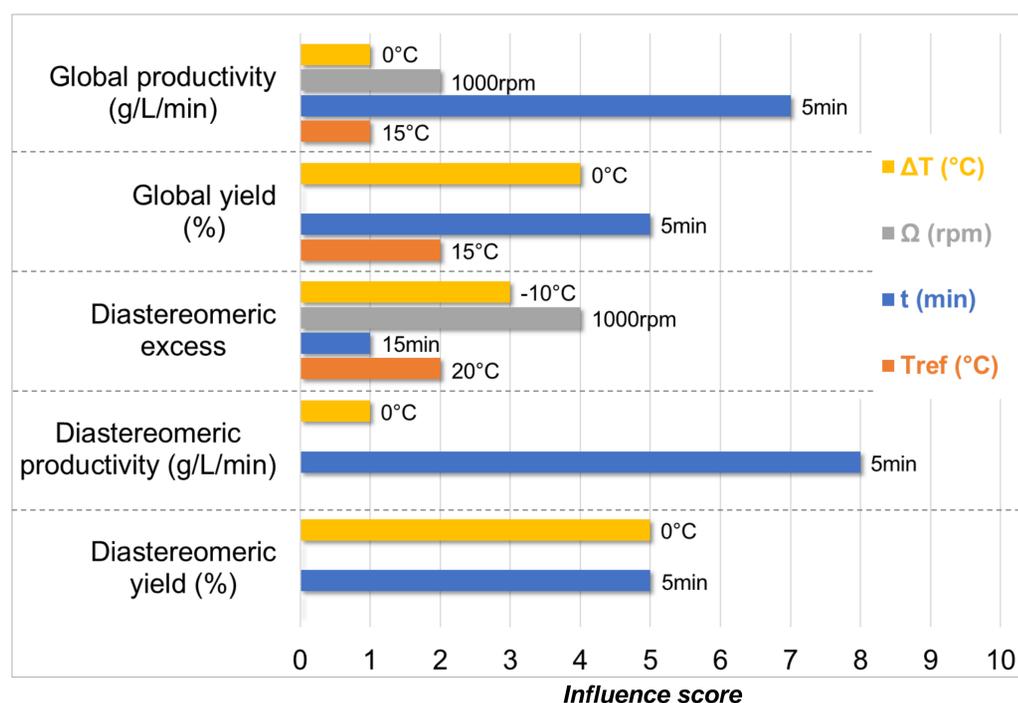
Conclusions and perspectives

- ⇒ 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].

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)
⇒ $\Delta T > 0$ seems to enhance chiral purity	⇒ 1000rpm seems to be the best factor level	⇒ Favor lower residence times	⇒ Apparent poor influence
⇒ Avoid T ₁ <T ₂	⇒ 200 and 500rpm eliminated	⇒ Avoid 30min	⇒ Favor lower temperatures

[References]

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