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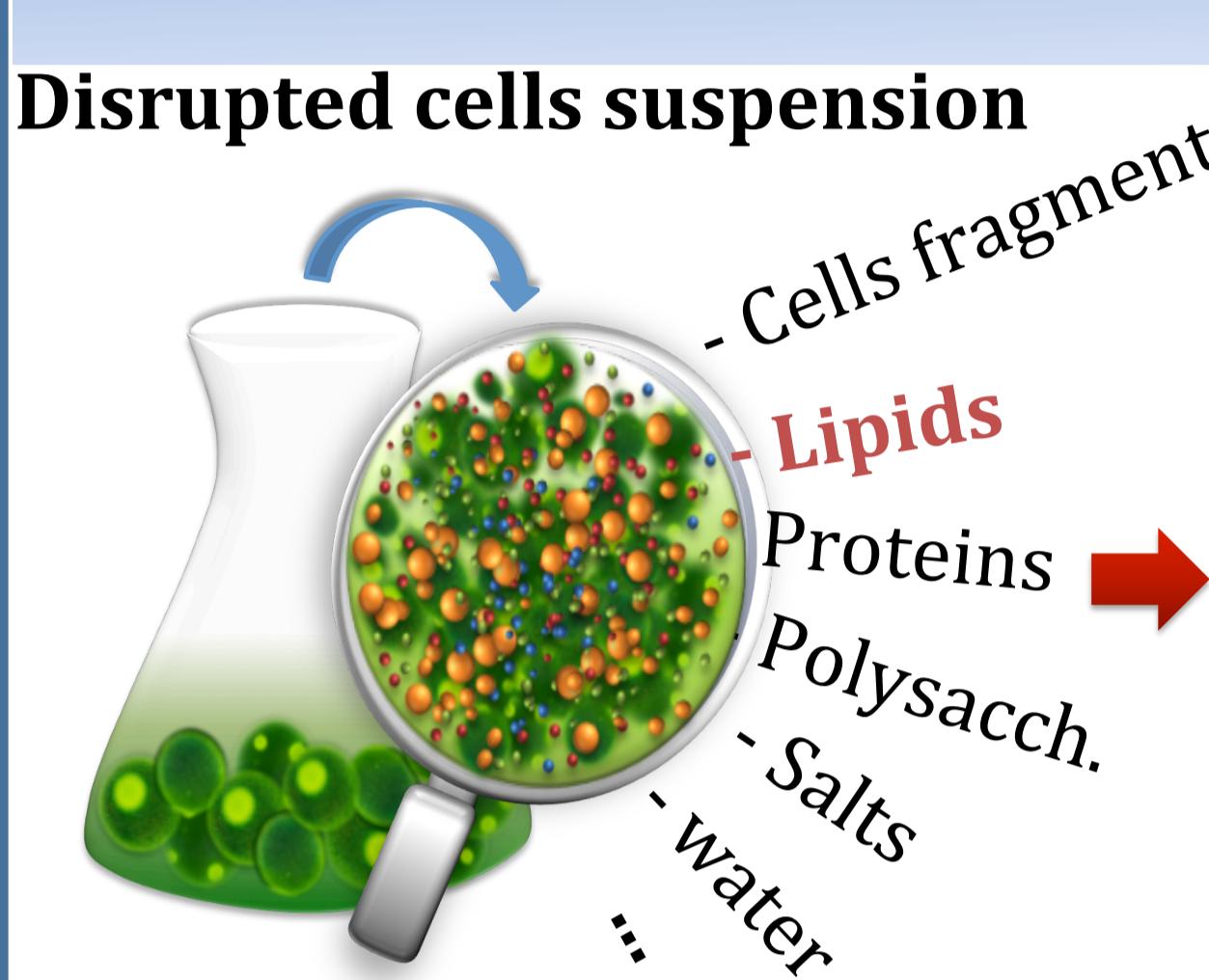


# MEMBRANE FILTRATION FOR THE RECOVERY OF LIPIDS FROM MICROALGAE EXTRACTS

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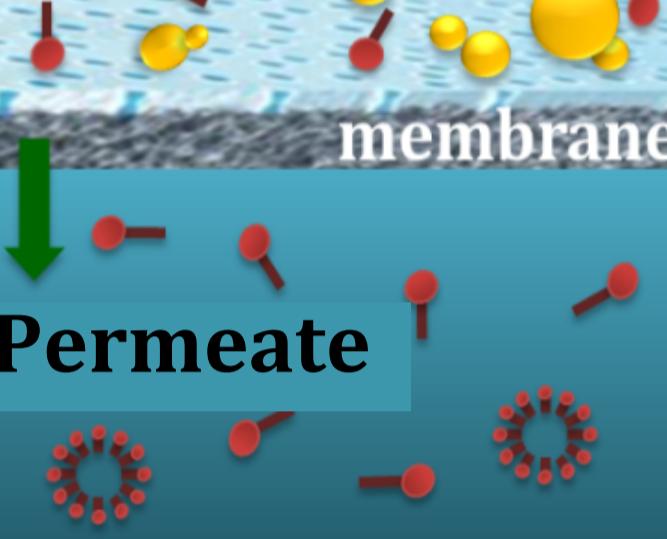
## Clarification

## Feed

## Ultrafiltration

## Retentate

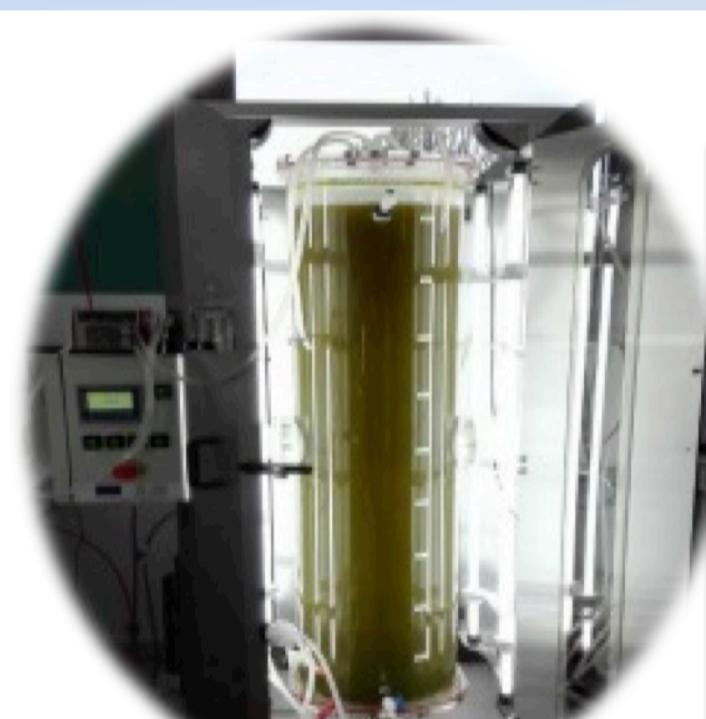
membrane



## Introduction

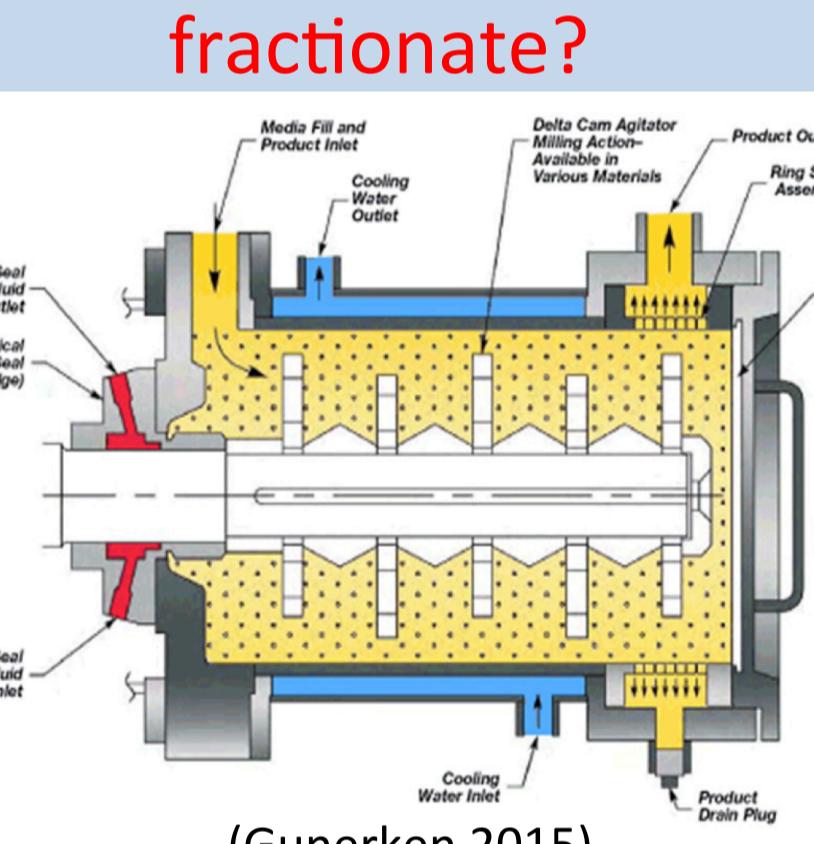
In this work we focus on the recovery of lipids from aqueous extracts of microalgae. After the culture of *Parachlorella Kessleri*, in starving conditions to enhance their lipid production, the micro-organisms were ground and the resulting mixture were centrifuged to extract aqueous fractions containing a large quantity of dispersed lipids.

## Challenge



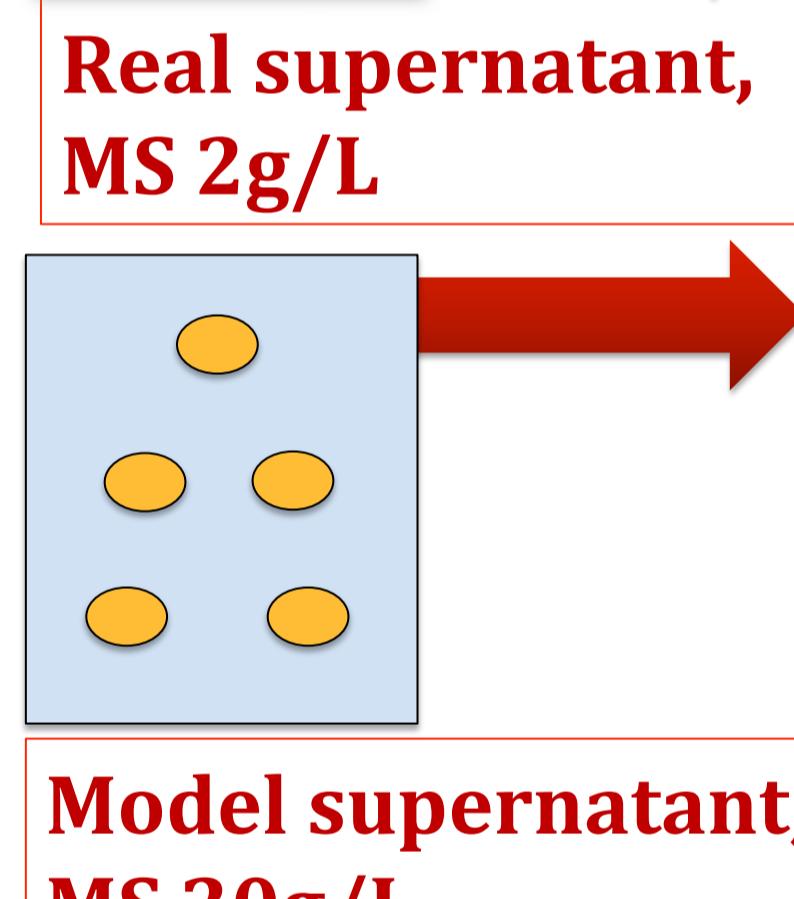
*P-chlorella kessleri*  
Cultured  
in starving conditions

Cell disruption:  
Target molecules are released  
in the aqueous phase.  
+ risk of degradation of their properties  
+risk of reorganisation: how to  
fractionate?

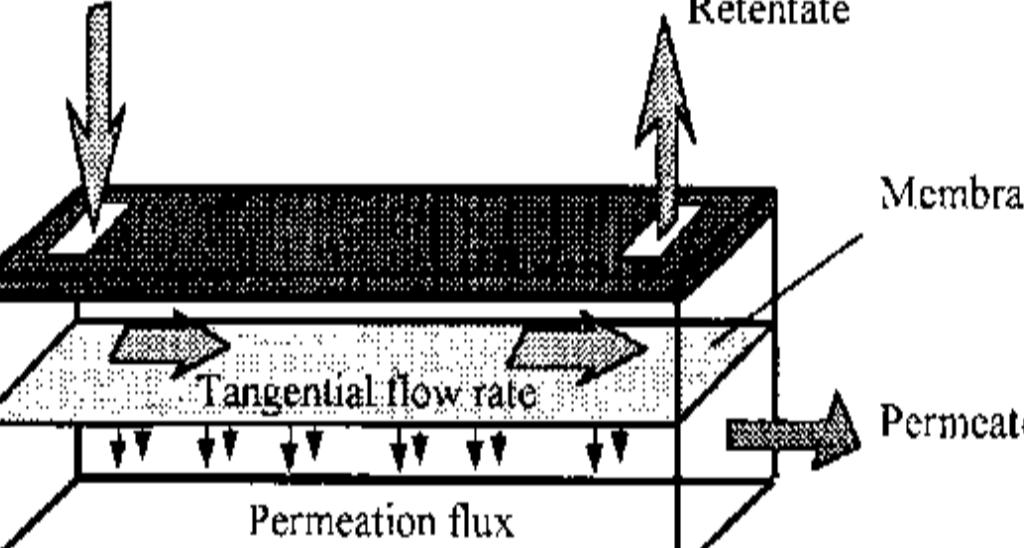


## Ultrafiltration

Crossflow (CF) module  
(Orelis Rayflow Pleiade)



Model supernatant,  
MS 20g/L



Shear rate :  
10,000 – 50,000 (s⁻¹)

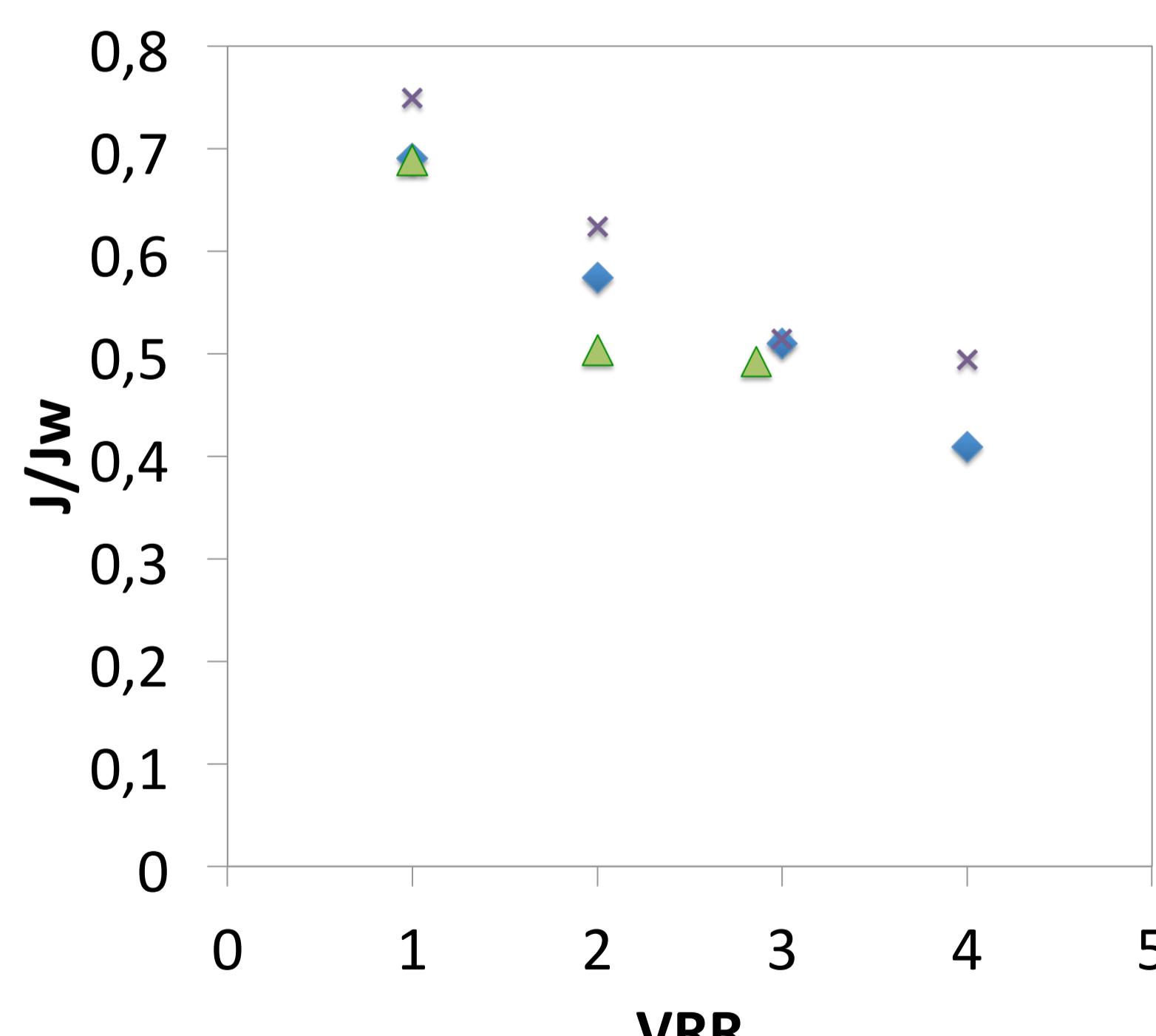
Dynamic filtration (DF) module  
(UTC Rotating disk)

Shear rate:  
10,000 – 500,000 (s⁻¹)

Model mixture: available in large quantities; controlled composition and organisation;  
Tests at high concentration

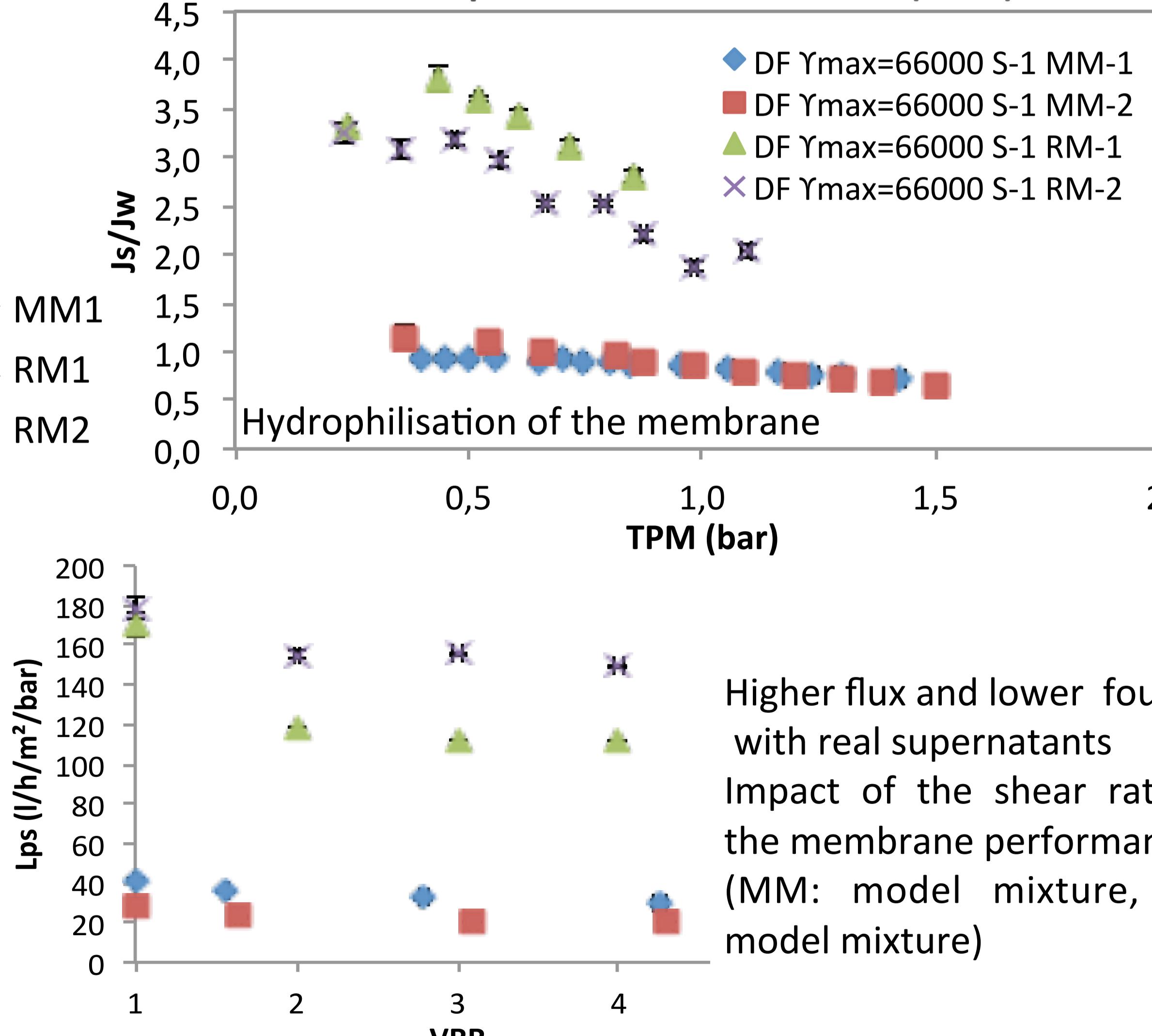
## Filtration results

### Cross flow filtration



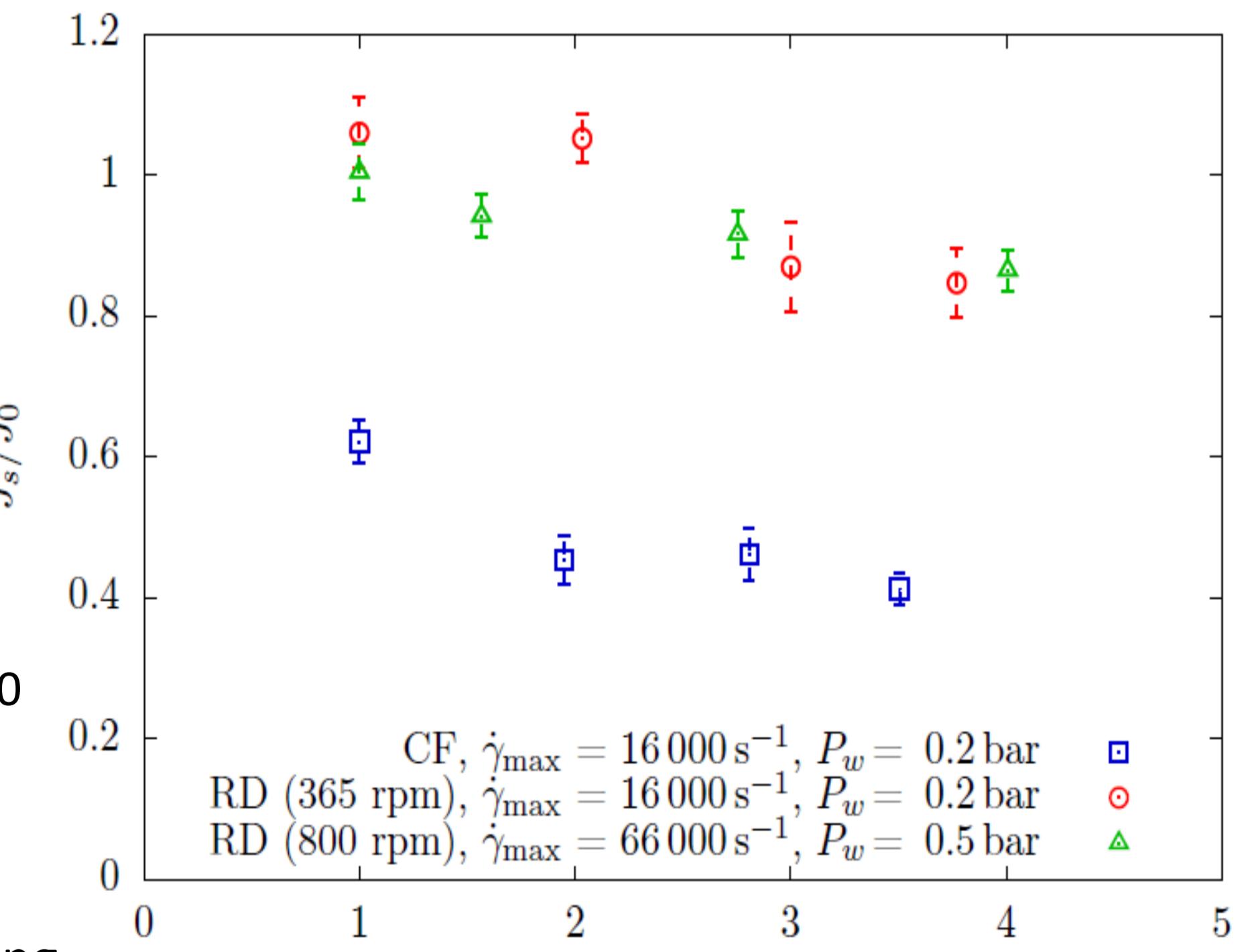
Similar flux and fouling  
with model and real supernatants  
Despite the concentration difference  
(MM: model mixture, RM: model mixture)

### Dynamic filtration



Higher flux and lower fouling  
with real supernatants  
Impact of the shear rate on  
the membrane performances  
(MM: model mixture, RM:  
model mixture)

### Comparison



Higher flux of dynamic filtration compared to  
cross flow filtration using the model mixture

## Conclusions and Perspectives

The membrane filtration allows the concentration of lipids from the aqueous extracts. The dynamic filtration offers interesting flux. The model mixtures allows the prediction of the fouling with concentrated extracts. The retention of the other compounds will depend on the cell disruption conditions: this will be deepened in a future study (PhD Shuli LIU). The membrane modification through hydrophilisation or fouling will be characterised through innovative technics in a futrure work (post-doctoral study, collaboration with ISCR Rennes and INRA Nantes).