

**Bioenergy-I: From Concept to Commercial Processes** 

### CONTINUOUS CULTURE OF FLOCCULENT YEA FOR ETHANOL PRODUCTION

José Cardoso Duarte Vera Lourenço, Belina Ribeiro, M<sup>a</sup> do Céu Sàágua, Lina Baeta-Hall

INETI Instituto Nacional de Engenharia, Tecnologia e Inovação

**Biotechnology Department** 

Portugal

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### >Brief Introduction



Bioethanol production became an important issue because:

>EU has established that 5.75% of transport fuels must be obtained from renewable sources (such as bioethanol) till 2010;

➢ Bioethanol production is a well known and developed technology but gains in productivity are still possible;

➤The interest for the use of new raw materials (lignocellulosic based) was raised by the recent oil crisis.





How to increase productivity of the bioethanol production process?



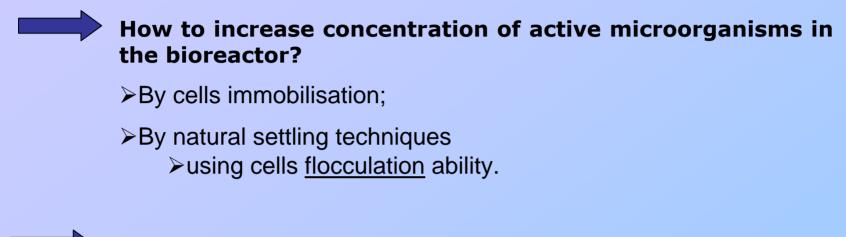
1. use of high concentration of active microorganisms in the bioreactor.

2. increase of dilution rate.

$$r_P = P. D = Y. X. D$$







Answer: many yeast and Zymomonas strains have good flocculation ability.



**Bioreactors using cells flocculation techniques could present some advantages:** 

Low associated capital and operational costs;

➤Design simplicity.





#### >The importance of Bioreactor Configuration

Bioreacto	r <sub>P</sub> (g <sub>EtOH</sub> /L h)	
Batch a	2-5	
Continuous Stirred Tank Reactor	Free Cells	6-8
	Immobilized Cells	10-16
	Entrapm. by Membranes	10-30
Packed	16-40	
Fluidized Bed Reactor	Free Cells	50-170
	Recycled Cells	190*

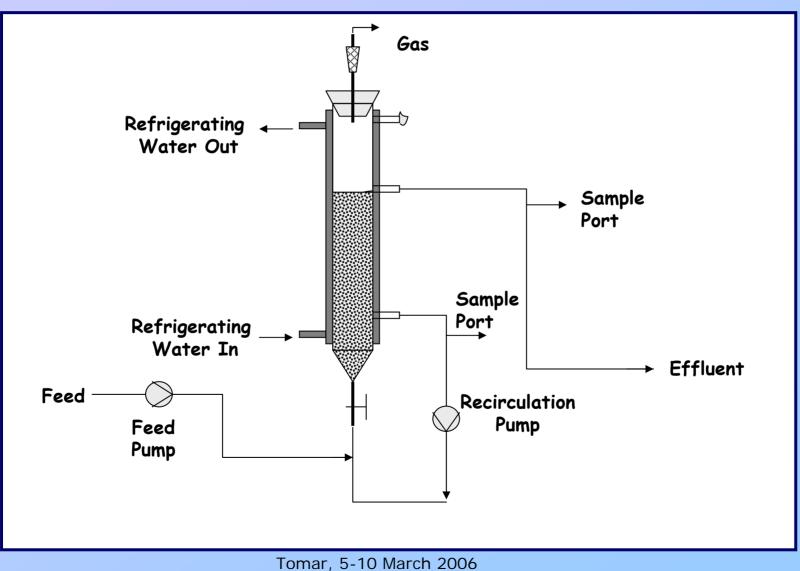
\* Productivity achieved at  $D=2,5 h^{-1}$ .





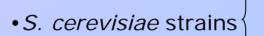
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# The Fluidized Bed Reactor (FBR)with Flocculated Cells Recycling











Relevant parameters

 Maximum specific growth rate (μ<sub>m</sub>)
 Ethanol yield (Y<sub>P/S</sub>)
 Ethanol tolerance





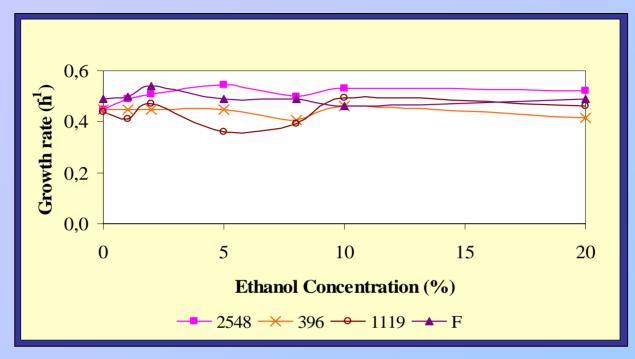


Strain	μ <sub>m</sub> (h <sup>-1</sup> )	X <sub>f</sub> (g/L)	P <sub>f</sub> (g/L)	Y <sub>P/S</sub> (g/g)	<r<sub>P&gt; (g/Lh)</r<sub>
ССМІ 396	0.366	9.87	31.50	0.26	1.31
DER 24	0.366	11.08	41.50	0.35	1.73
DSMZ 2548	0.396	13.60	46.60	0.23	1.94
NCYC 1119	0.440	nd	49.59	0.39	2.07
F	0.490	12.73	43.50	0.40	1.81

#### Substrate: sucrose 120 g/L Temperature: 28°C







#### **Batch Conditions:**

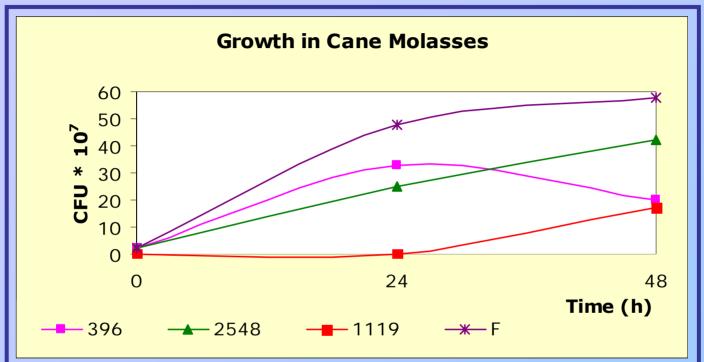
- -Ethanol addition in concentrations ranging from 0 to 20%
- -Initial sucrose: 120 g/L
- -Temperature: 28°C
- -Shake flasks



#### all strains showed good tolerance to ethanol.







#### **Batch Conditions:**

-Initial sugar: 120 g/L -Temperature: 28°C -Shake flasks **strain F showed a better performance** in growing in cane molasses medium. Strain **1119 cell concentration is the lowest** but that is due to the high sedimentation velocity, which difficults a representative sampling.







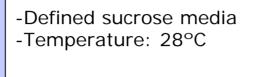
Only strains F and 1119 showed good flocculation ability, which is very important in FBR fermentations with no solid suport to retain biomass; 1119 has more dense flocs than F.

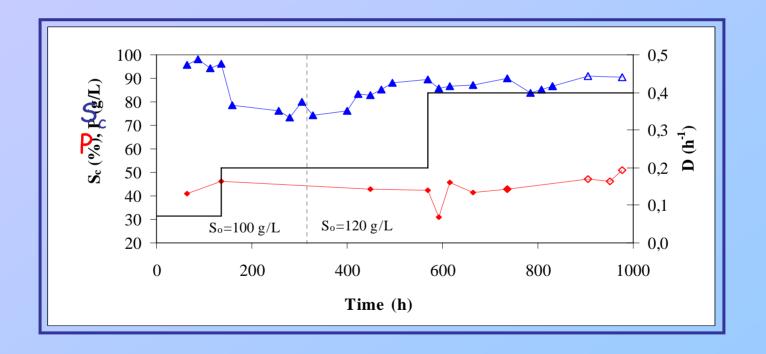


Due to higher maximum specific growth rates, ethanol yields and flocculation ability, strains F and 1119 are suitable for ethanol production in FBR



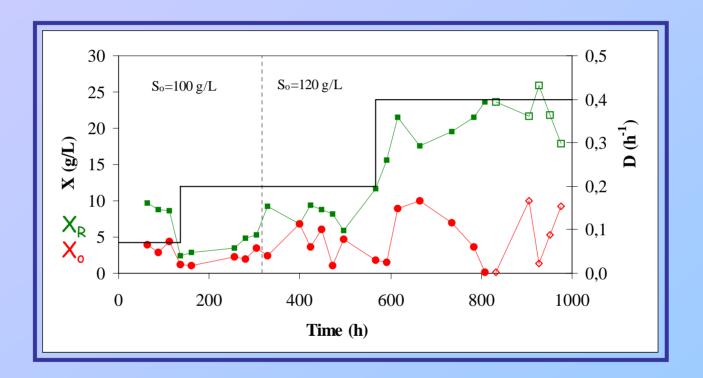








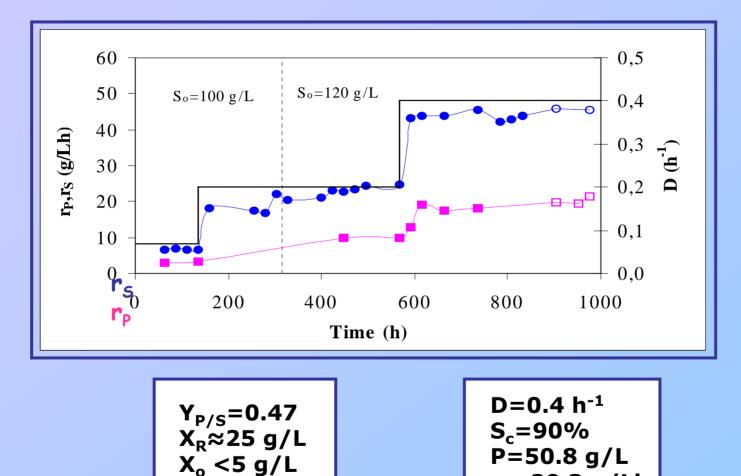
ΠΕΤΙ



#### There is a good capacity for retention of biomass



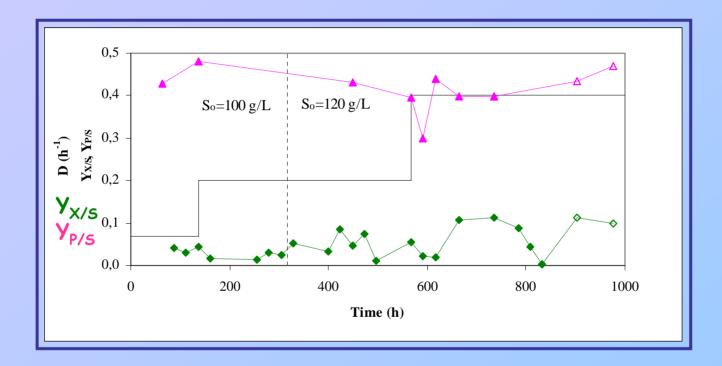




 $r_{P} = 20.3 \text{ g/Lh}$ 



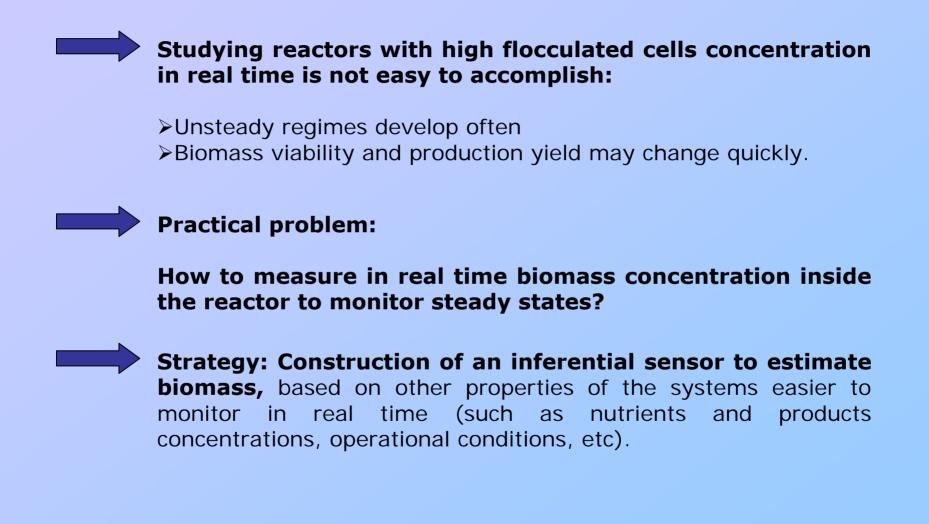




Handicap?: Strain F forms small flocs with low sedimentation velocity.











### Construction of a biomass inferential sensor:

 $\checkmark$  For statistical fitting of data, only steady-state values were used concerning experiments with strain F.

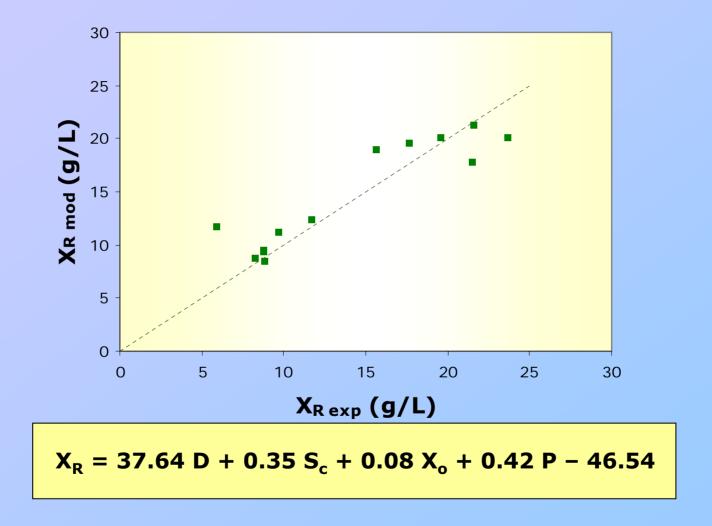
✓ Biomass concentration within the broth was related to ethanol, biomass concentration in the outflow, dilution rate and percentage of consumed sugar.

✓Only linear correlations were tested and the best one was obtained through an Ordinary Linear Square (OLS) Regression.





#### Construction of a biomass inferential sensor:







#### Construction of a biomass inferential sensor:

 $\checkmark$ The constructed biomass inferential sensor permits to predict biomass concentration within the bioreactor by the quicker measurement of other variables.

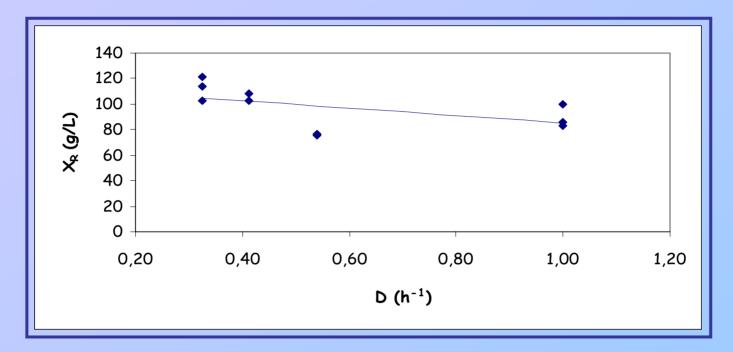
 $\checkmark$ The correlation fitted well the used data, expressed by the Root Mean Square Error (RMSQ=0.68).

✓The correlation is capable of predicting biomass concentration in a quite reasonable way, as the  $R^2_{cv}$  parameter expresses ( $R^2_{cv}$  =0.67).

 $\checkmark$ The most important predictor is dilution rate (37.64) and the least important is biomass concentration in the outflow (0.08).



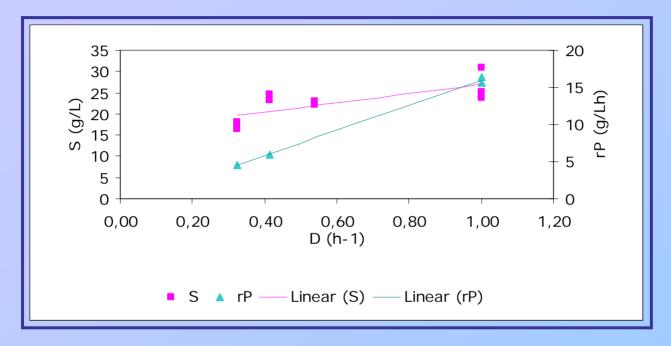




Strain 1119 has bigger flocs and higher sedimentation velocity, which allows it to stay inside the FBR at higher dilution rates ( $D > \mu_m$ ). At all tested dilution rates cells always showed viability ab-ove 90%.







Ethanol productivity of more than **15 g/Lh** was attained, but the yield  $Y_{S/P}$  was of 0.2 (aprox.). If we can raise  $Y_{S/P}$  till near the theoretical one (0.5) ethanol productivity will be near to **40 g/Lh**.





#### **>**Some pictures of FBR in operation



Strain F



#### Strain 1119





### Continuous FBR Tests: Microbial Resistance

FBR showed a natural resistance concerning microbial contamination, mainly due to:

✓ high dilution rates✓ high cells density

This means:

✓longer fermentations✓reduced need for sterility



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#### Continuous FBR Tests: Scale-Up

Prototypes (4;25;400;4000 L ) were built and tested for long-term operation with good results.

## 25L-prototype: a tubular air-lift reactor with an external settler;







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# Thank you for your attention!

For any enquiries email me:

jose.duarte@ineti.pt

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