



# THE POTENTIAL OF THE MICROWAVE-VACUUM TECHNIQUE IN DRYING OF PROBIOTIC AND STARTER CULTURES

IDS 2012, Xiamen, China

12<sup>th</sup> November 2012

Sabine Ambros, S.A.W. Bauer, Ulrich Kulozik, Petra Först



Chair for Food Process Engineering and Dairy Technology  
Technische Universität München, Freising-Weihenstephan, Germany

# Microbial cultures

- Fields of application of starter cultures and probiotics:

dairy products, pastries, pickles, raw sausages

probiotic foods

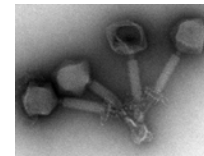


- Advantages of direct starter cultures:

- no contamination by bacteriophages

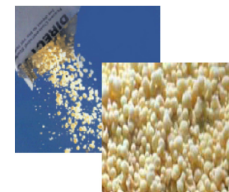
- reproducible product quality

- little know-how and personnel expenditure



- Stabilization by: - freezing

- drying



# Motivation

Current standard procedures in drying of microbial cultures:



High time and energy effort

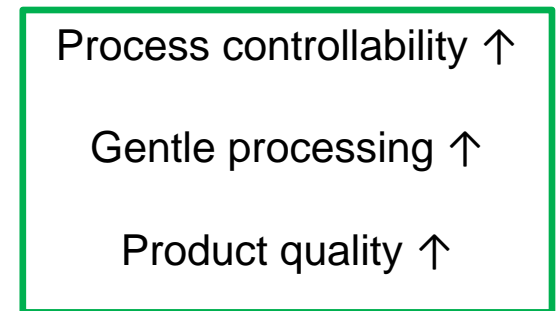
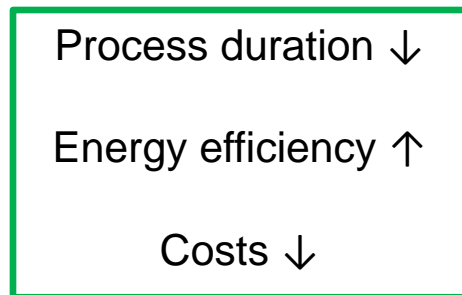


Low product quality



Time consuming

➔ *Microwave vacuum-drying as an alternative?*



# Quality aspects of microbial cultures

- Viability
  - Metabolic activity
  - Storage stability
  - Handling
- Rehydratibility
- Proper dosability

→ Aim: Maintenance of technological and biological functionality for direct inoculation without preculturing

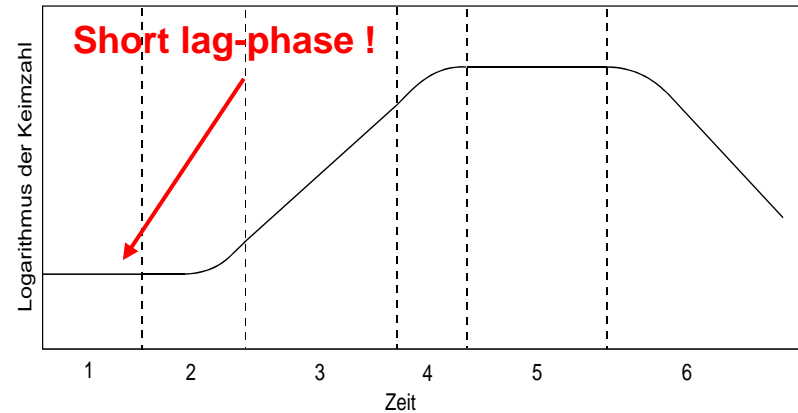
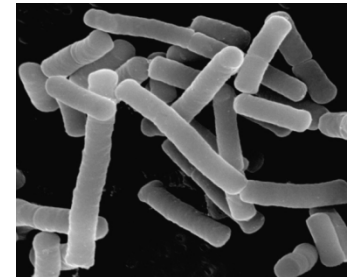
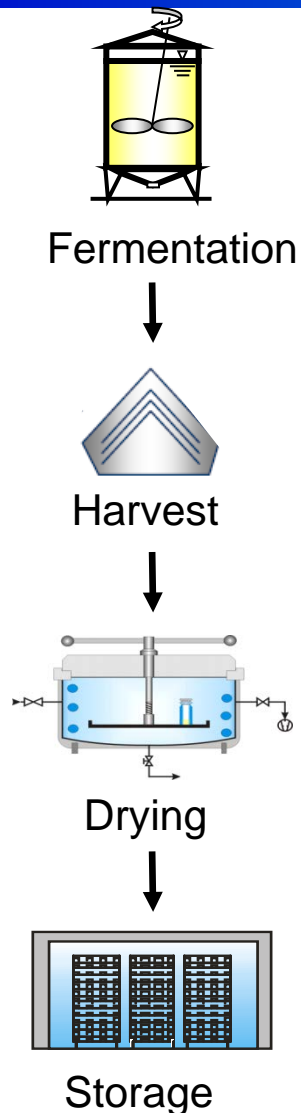


Abb.6.1: Vermehrungsphasen einer Bakterienkultur: 1. Latenzphase, 2. Beschleunigungsphase, 3. exponentielle Vermehrungsphase, 4. Verzögerungsphase, 5. stationäre Phase, 6. letale Phase



# Culture preparation/ Test set-up



## Drying parameters:

- Model strain: *Lactobacillus paracasei* spp. *paracasei* (F19)
- 100 g cell suspension
- $h_{\text{sample}} = 0.5 \text{ cm}$
- Drying jar: Duran glass
- Residual water content = 6-7%

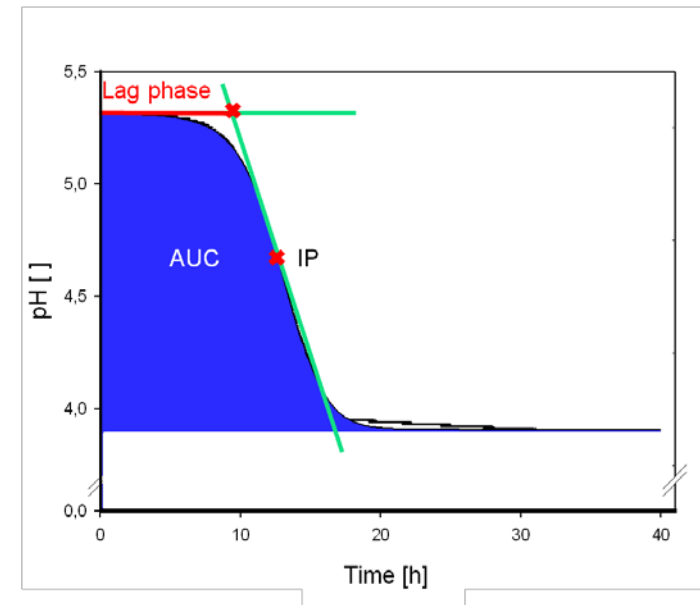


## Analytics:

- Survival rate by plate count method and flow cytometry
- Acidification power test

$$pH(t) = pH_0 + \frac{a}{(1 + e^{\frac{t-t_0}{b}})^c}$$

$$\int_0^{40} pH(t) dt - pH_{\text{end}} \cdot t_{\text{end}} = AUC$$

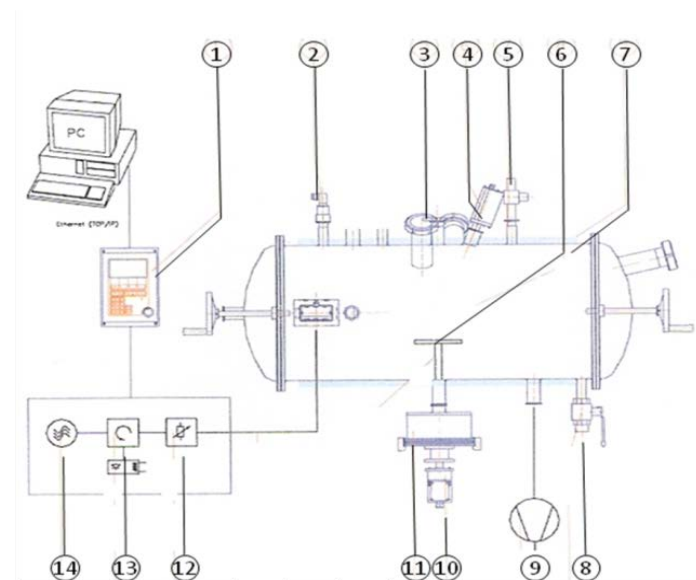
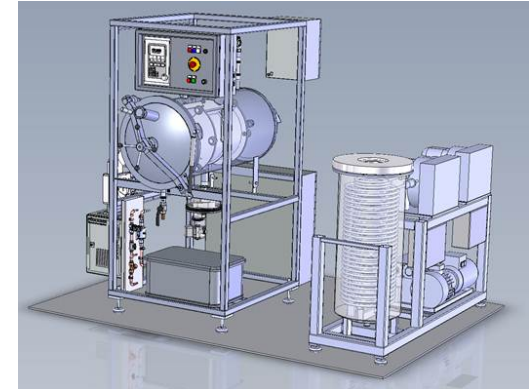


# Microwave vacuum dryer

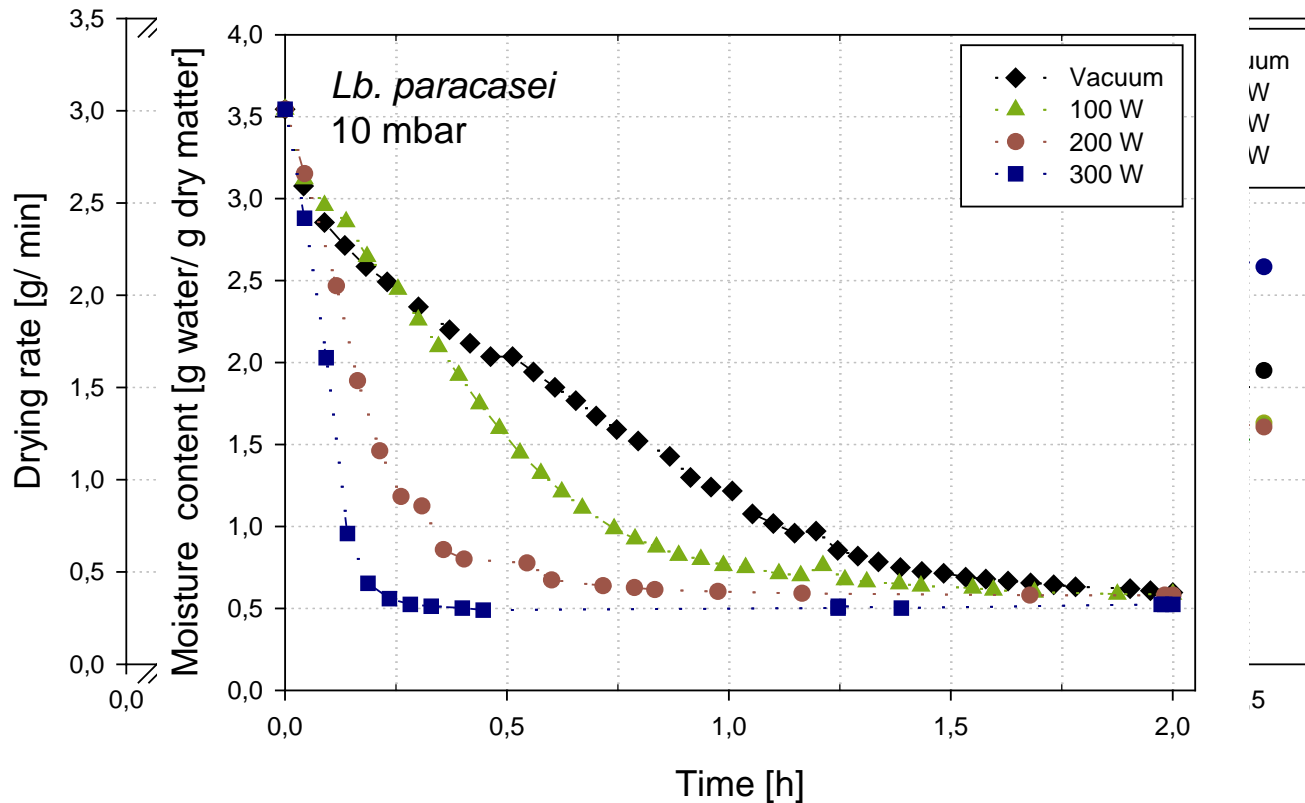
- Manufacturer: Püschner MicrowavePowerSystems, Schwanewede-Bremen, Germany
- Microwave power: max. 1 kW/ 2,450 MHz
- Integrated load cell and IR-camera
- Prozess visualization: online-measuring of  $\rho$ ,  $m$ ,  $T$ ,  $P$
- Influencing factors:
  - Microwave power
  - Pressure
  - Product characteristics  
(dielectric loss factor)

$$P_r = 2 \pi f E^2 \varepsilon_0 \varepsilon''$$

$P_r$ : Relative absorbed power  
 $f$ : Frequency  
 $E$ : Electric field strength  
 $\varepsilon_0$ : Absolute dielectric constant  
 $\varepsilon''$ : Dielectric loss factor

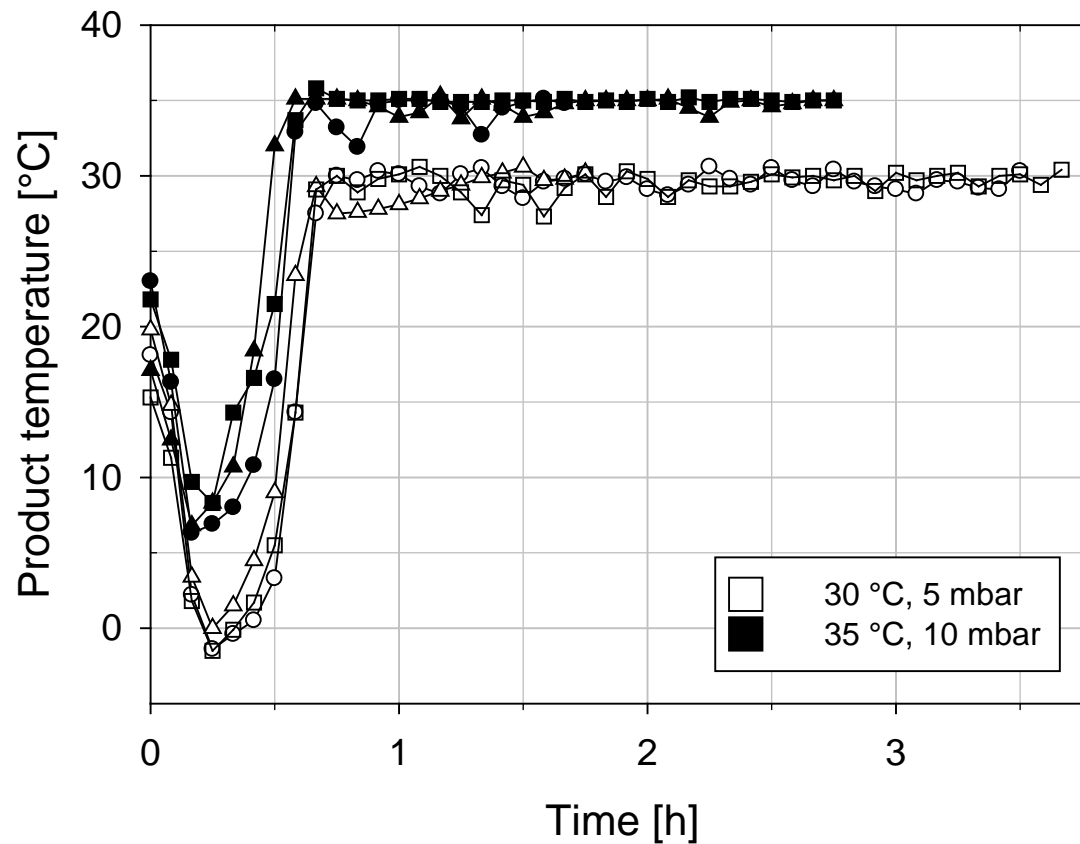


# Drying rates



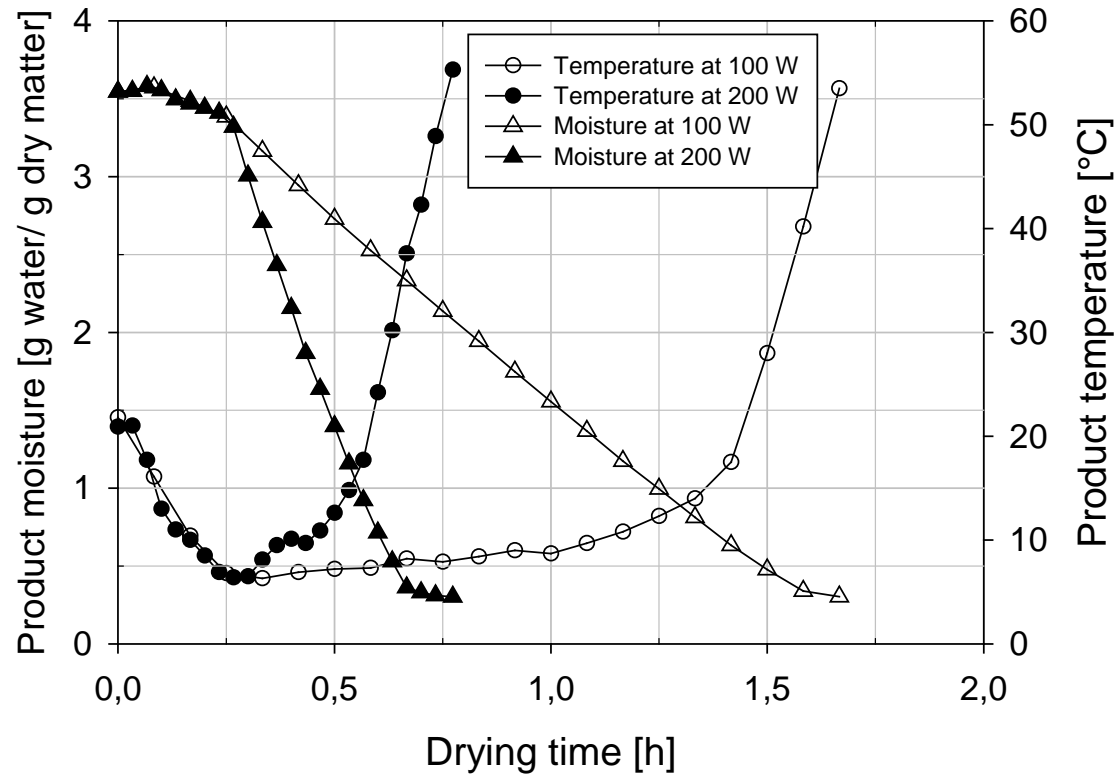
- Clear dependence of the drying rate on the microwave power input
- Process can be controlled via microwave energy input

# Performance and replicability of drying process





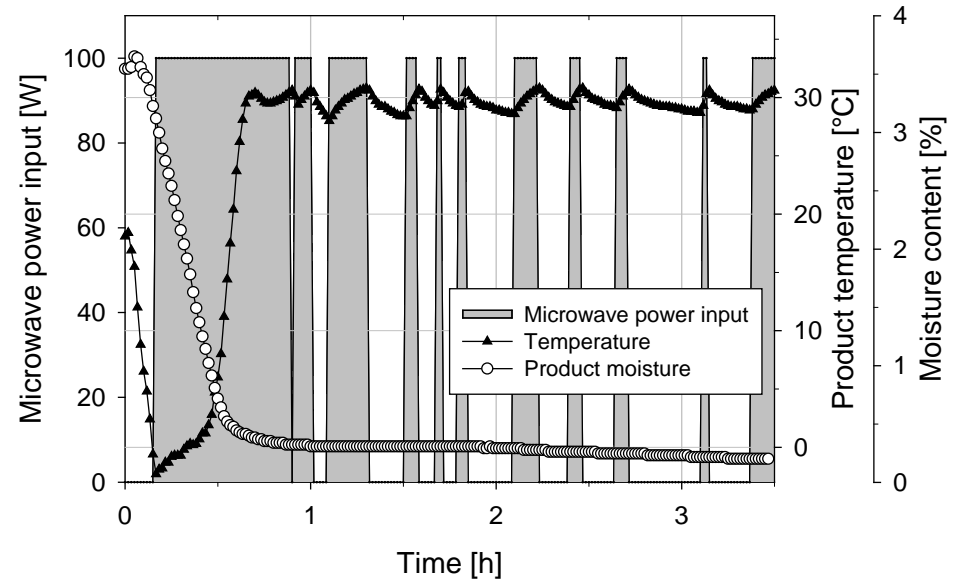
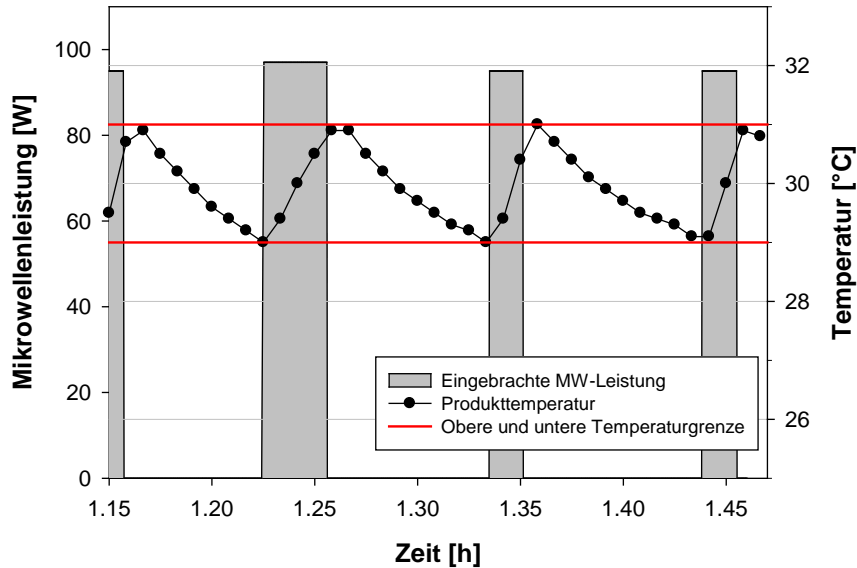
# Drying profiles for continuous microwave input



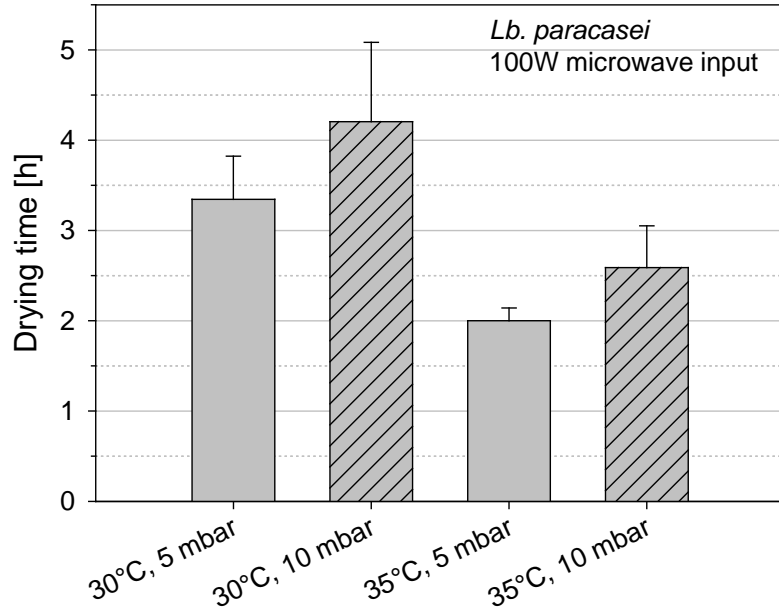
→ Drying below critical residual water content leads to detrimental product temperatures

# Pulsed microwave input

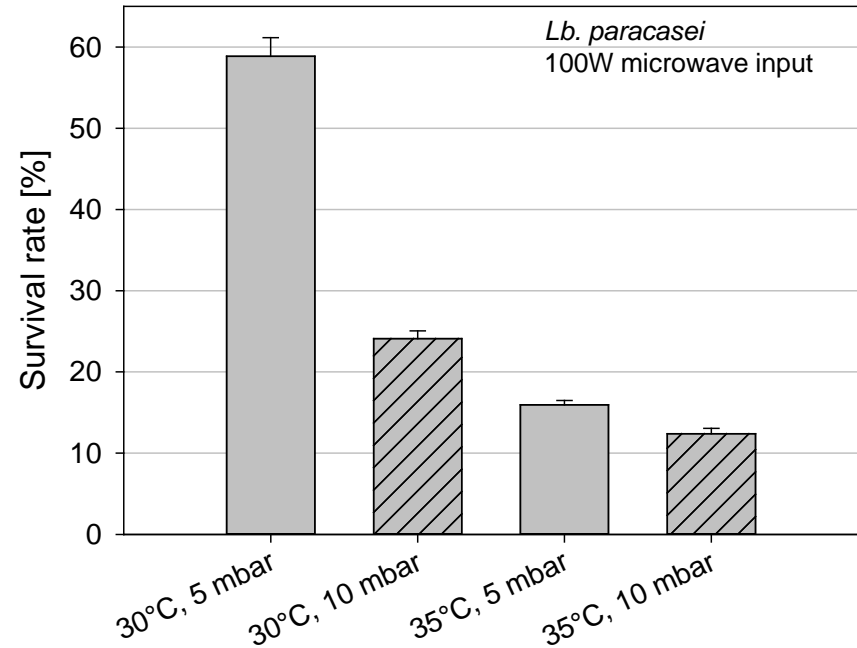
Pulsed energy input: Temperature regulation according to  $T_{max}$



# Influence of chamber pressure and product temperature on drying time and survival

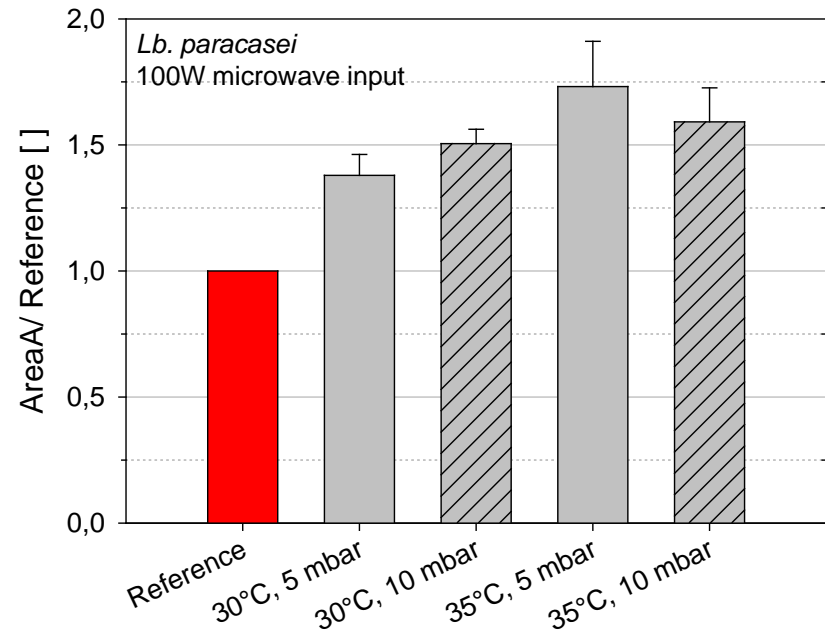
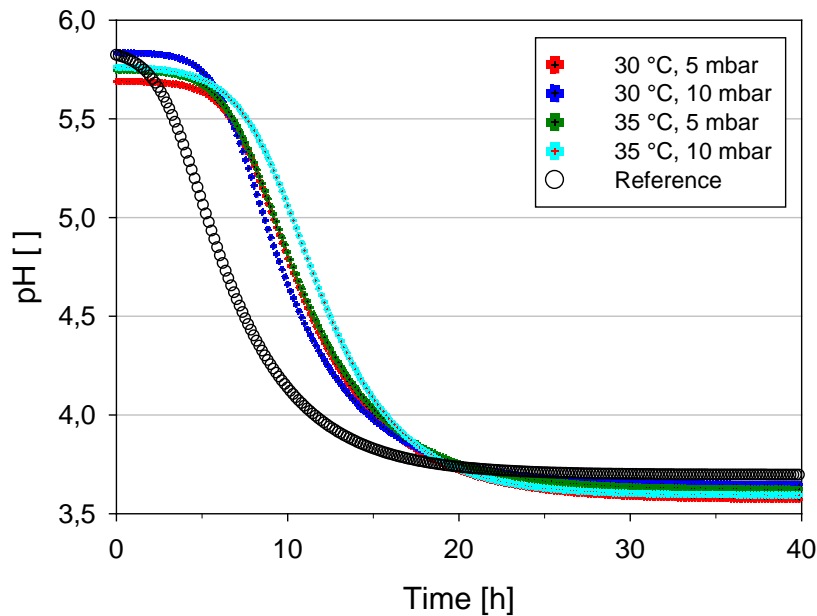


→ Besides microwave power input  $T_{\max}$  determines drying time



→ High survival at low  $T_{\max}$  and low chamber pressures

# Influence of chamber pressure und product temperature on acidification activity



→ Low  $T_{\max}$  and low chamber pressures are favourable for vitality

# Summary

---

## **Microwave vacuum-drying of lactic acid bacteria is generally possible!**

- Microwave enables **shortening of drying time up to 90%** in comparison to freeze and vacuum drying
- **Pulsed microwave input** enables **gentle drying** and thus **high survival rates**



→ *Microwave vacuum-drying has got the potential to overcome many inconveniences in the drying of lactic acid bacteria and so could replace or at least complement the established drying techniques*