FAT-BASED ENCAPSULATION: PHYSICOCHEMICAL CHARACTERISATION AND RELEASE KINETICS



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1. Introduction

Fat-based cold extrusion represents a high potential but largely unexplored encapsulation technique [1,2]. Understanding the functioning and efficiency of encapsulated systems the determination of physicochemical characteristics of materials and their interactions is crucial [3].

Approach/Aim

- matrix design of fat mixtures consisting of high-melting hydrogenated rapeseed oil + variable types and contents of lowmelting fats/oil for encapsulation of 10% beetroot powder (RBP) by cold extrusion
- physicochemical characterization of fat matrices and encapsulants
- in-vitro release kinetics of encapsulated RBP

2. Methods

Cold extrusion

- Matrices: high-melting 1) Fat hydrogenated rapeseed oil (HMF) + 3% to low-melting fats (LMF): 16% of rapeseed oil (ROE)/palm kernal fat (PKF)/two coconut fats (COC; COP)
- encapsulants: matrix material 2) RBP = fat mixtures with 12% and 16% LMF; 10% encapsulated RBP

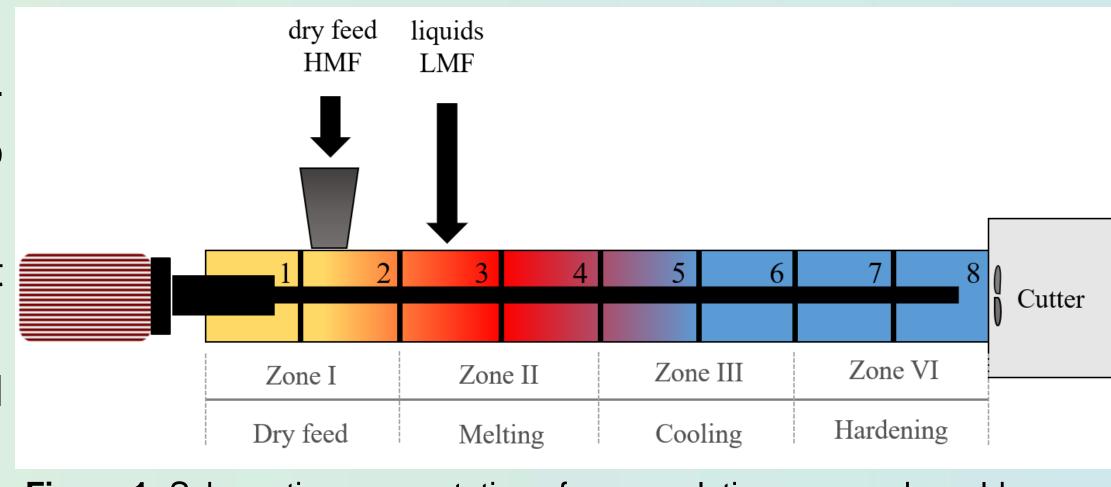


Figure 1: Schematic representation of encapsulation process by cold extrusion.

Physicochemical Analyses

- Particle size distribution (sieve analysis)
- Melting behaviour (DSC)
- Solid fat content (NMR)
- Microstructure (CLSM)

Release kinetics of RBP encapsulants

Encapsulation efficiency, release profile, release behaviour (UV-visspectrophotometry)

3. Results and Discussion

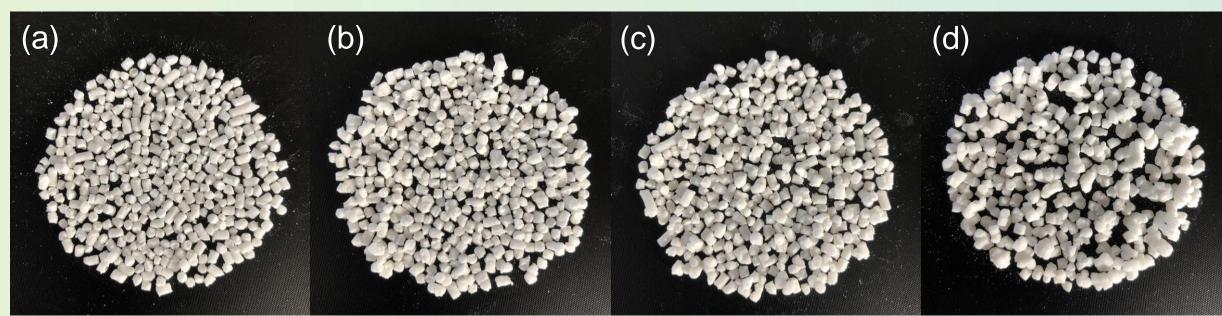


Figure 3: Extruded fat matrices consisting of fat mixtures with (a) 3%, (b) 6%, (c) 12% and (d) 16% ROE and HMF as a base.

22,04

23,97

Cococnut fat (Walther Rau) + hydrogenated rapeseed oil

21,36

Cococnut fat (Palmin) + hydrogenated rapeseed oil

■ 3 min (oral)

13,07

20,28

27,38

Rapseed oil + hydrogenated rapeseed oil

Palm kernal fat + hydrogenated rapeseed oil

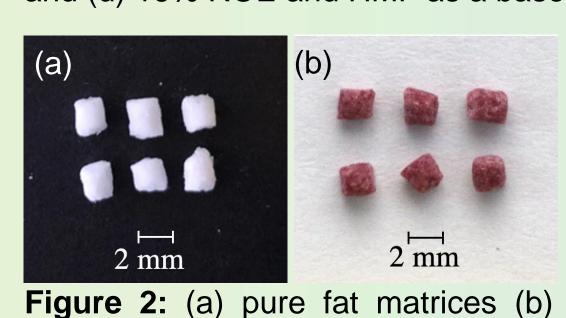
6,26 3,74

4,34 3,41

3,14 2,46

2,96 5,14

16% 2,22 3,31



RBP

Total release of

during digestion

12% and 16% of

The

with

material

and HMF

encapsulated

Figure 5:

RBP

phases.

Matrix

consist

LMF

mixtures

as a base.

extrusion and cutting.

encapsulants

12% 2,48 4,01

16% 2,352,16

Increasing share of LMF:

- increasing proportion of particles within size fraction > 2.8mm → agglomerates of a number of pellets
- softer consistency

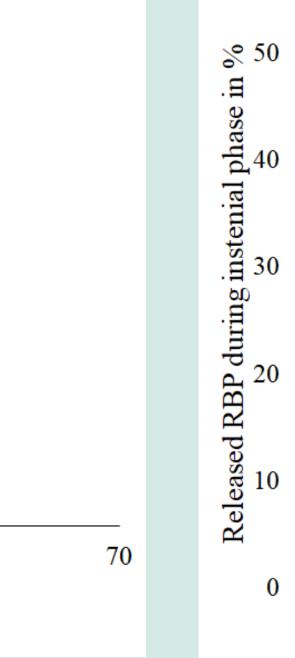
Influencing factors on melting behaviour: temperature and time

 increased storage duration led to increasing melt of fat crystals; homogeneouse RBP distribution

Figure 7: CLSM images of RBP encapsulants with 16% PKF in the matrix coloured with Nile Red and FITC measured at (a) 37°C after 5h, (b) 37°C after 1d.

Figure 6: CLSM images of fat matrix with 16% PKF coloured with Nile Red and

measured at (a) 25°C, (b) 37°C, (c) 37°C after 5h, (d) 37°C after 1d.



phase

Release mechanisms

• 12% LMF: ≈ 92.16%

Encapsulation efficiency

- Melting (temperature & time dependent)
- 16% LMF: ≈ 90.85%
- Enzymatic lipid digestion

Released RBP in each digestion phase in %

■ 120 min (gastric) ■ 240 min (intestinal)

48,14

• Matrix with 16% rapeseed oil (ROE) + 10% encapsulated RBP ■ Matrix with 16% palm kernal fat (PKF) + 10% encapsulated RBP between ◆ Matrix with 16% coconut fat (COC) + 10% encapsulated RBP Matrix with 16% coconut fat (COP) + 10% encapsulated RBP matrices Difference in SFC at 20.0 °C and 36.5 °C in %

deviation

→ 16% PKF in the matrix peformed best

HMF as the base 8 and 16% of LMF. Correlation between released proportion of RBP during intesinal 36.5°C 20.0°C and

Figure 8:

20.0°C

released

during

phase

RBP).

material:

mixtures

consisting

Comparison

36.5°C and

encapsulated

deviations SFC at

and

the

fat

(10%

Matrix

fat

of

intestinal

Thermal properties strongly influence release behaviour deviations in solid fat content as a measure of structural

changes in a temperature range of 20.0°C to 36.5°C are crucial

SFC

at

4. Conclusion

- Good extrudability of fat mixtures up to a share of 16% LMF
- Matrix composition and encapsulated RBP impact physicochemical properties

References

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