

Introduction

• Using in tablet formulations microcrystalline cellulose (MCC) with plastic behaviour and calcium phosphate anhydrous (**CaHPO**_₄) with brittle behaviour under compaction is very popular in the pharmaceutical industry for achieving desirable structural-mechanical properties of tablets.

Compaction and Structural-Mechanical Properties of Tablets as a Function of Volume Ratios of Excipients

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Materials

- MCC (CEOLUS UF-711; Asahi Kasei, Japan)
- CaHPO4 (DI-CAFOS A60; Budenheim KG, Germany)
- Silica (SYLOID® 244FP; Grace GmbH, Germany)
- Sodium stearyl fumarate (PRUV®; JRS Pharma, Germany)

Methods

- Powder mixtures (Table 1) were tableted with 11.28 mm flat punches to obtain 500 mg tablets using a compaction simulator (STYL'One Nano, Medelpharm, France).
- Compression cycles simulated small rotary press at tableting speed of 70 rpm with





• The aim of this study was to investigate the compaction properties of mixtures of MCC and CaHPO₄ in different volume ratios at the range of compaction pressure to observe and explain the influence of mixture composition on the structuralmechanical properties.

Table 1 50-50 25-75 -100 -25 100-(5 0 mg/mm³ w/w Ingredients CEOLUS[™] UF-711 0.608 0.346 0.151 0.977 0.000 1.586 DI-CAFOS[®] A60 0.369 0.631 0.826 0.977 2.890 0.000 PRUV® 0.020 0.020 0.020 0.020 1.110 0.020

pre-compaction force of 5 MPa and compaction force of 10-50 MPa.

- The **powder feeding** into the die was performed automatically via the feed shoe.
- The tablet height (t), diameter (d), and hardness (F) were measured (n=10) by a tablet tester (ST50 WTDH; SOTAX AG, Switzerland) immediately after the compaction and converted into tensile strength (MPa).
- The calculated true density of composition was obtained based on the true density (pt) of components and their shares (x, w/w) using the additive methodology:

 $\rho_t = (\rho_{exc\,1} \cdot x_{exc\,1}) + (\rho_{exc\,2} \cdot x_{exc\,2}) + \dots + (\rho_{exc\,i} \cdot x_{exc\,i})$

- For **in-die Heckel plot**, the **relative density In(1/ε)** was calculated with Alix software (Medelpharm).
- X-ray Micro-Computed Tomography (µCT) of samples has been done with a 3D Micro X-ray CT (CT Lab HX; Rigaku Corp., Japan) at 70 kV with a current of 50 µA, focus set on S, and with no filter. CT scan were exported and processed with Dragonfly soft (Object Research Systems Inc., Canada).







50 100 2 00 100 40	~
Compaction pressure, MPa	

Fig. 9

Fig. 10B: F 25-75 @ 450 MPa

Compression pressure, MPa

Compression pressure, MPa

Fig. 12

Fig. 11

Results

- Lubricant amount was justified by the tablet ejection force profile (Fig. 1)
- Tensile strength increased along with MCC fraction and compaction pressure (Fig. 2)

Fig. 10A: F 25-75 @ 100 MPa

- Proportion-tensile strength profiles in the 25-75 vol.% MCC range is almost linear (Fig. 3)
- Proportion–porosity profiles in the same 25-75 %vol. MCC range is almost linear (Fig. 4)
- Punches' distance is smaller, tablets are thinner because of the true CaHPO₄ density (Fig. 5)
- Increasing CaHPO₄ portion has increased the in-die yield pressure Py (Fig. 6)
- Increase in the MCC portion increased the value of elastic recovery (Fig. 7)
- Out-of-die was in line with in-die Heckel plot except for F25-75 (Fig. 8)
- Out-of-die Heckel plot of F25-75 at 100-150 and 150-500 MPa cause of MCC and CaHPO₄ (Fig. 8-9) μCT 100 vs 450 MPa (Fig. 10); Comparison of processed μCT data (Fig. 11) vs. calculated (Fig. 12)

Discussion & Conclusion

- F100-0, F75-25 and F50-50 were in the Successful Formulation Window (**Fig. 1-2**)
- Non-linear segments can be related to the percolation threshold (Fig. 3-4)
- The plasticity and elastic recovery increase with increasing MCC (**Fig. 6-8**)
- F25-75 has the two-stage out-of-die Heckel plot (Fig. 8-9)
- 1st stage controlled by MCC and 2nd by CaHPO₄ (Fig. 8-9)
- µCT can be useful for de-formulation (Fig. 10-12)

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