

Studies of the properties of tablets from directly compressible isomalt

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SUMMARY

Studies of the properties of tablets from directly compressible isomalt

The paper deals with the study of the properties of tablets from two types of directly compressible isomalt – namely galenIQ™ 720 and galenIQ™ 721. The study focuses on the dependence of tensile strength and disintegration time of the tablets, on both the compression force, the addition of lubricants, namely magnesium stearate and sodium stearyl fumarate (Pruv) with the concentrations of 0.5% and 1%, and a 50% addition of model active ingredients, namely ascorbic acid and acetylsalicylic acid. The used compression forces were 6, 8 and 10 kN, tablets containing drugs were compressed only with the force of 10 kN. The tensile strength of tablets from both substances increased with increasing compression force, the presence of lubricants did not result in any decrease in strength. The disintegration time of tablets was longer with the use of substance galenIQ™ 720, it was prolonged by lubricants and increased with increasing compression force for both substances. The presence of drugs decreased tensile strength, and the tablets containing acetylsalicylic acid showed higher tensile strength. For both drugs, differences within the type of isomalt or lubricants were not significant. The disintegration time of tablets was shorter when ascorbic acid was used; it was the shortest in the case of galenIQ™ 721 with Pruv, but in the case of acetylsalicylic acid the disintegration time was the shortest with galenIQ™ 721 with magnesium stearate.

Key words: directly compressible isomalt – lubricants – tensile strength of tablets – disintegration time of tablets

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SOUHRN

Studium vlastností tablet z přímo lisovatelného isomaltu

Práce se zabývá studiem vlastností tablet ze dvou typů přímo lisovatelného isomaltu – galenuIQ™ 720 a galenuIQ™ 721. Studovala se pevnost tablet v tahu a doba rozpadu tablet, obojí v závislosti na lisovací síle, přídavku stearanu hořečnatého a stearylfulmarátu sodného (Pruvu) jako mazadel v koncentracích 0,5% a 1% a 50% přídavku modelových léčivých látek kyseliny askorbové a kyseliny acetylsalicylové. Tablety byly lisovány lisovacími silami 6, 8 a 10 kN, výlisky s léčivou pouze lisovací silou 10 kN. Pevnost tablet z obou látek rostla s lisovací silou, vlivem mazadel se nesnížila. Doba rozpadu byla delší s látkou galenIQ™ 720, u obou látek byla prodloužena mazadly a rostla s lisovací silou. Přítomnost léčiv snížila pevnost tablet, pevnější byly výlisky s kyselinou acetylsalicylovou. U obou léčiv nebyl výrazný rozdíl v rámci typu použitého isomaltu ani mazadla. Doba rozpadu byla kratší u tablet s kyselinou askorbovou, kdy byla nejkratší s látkou galenIQ™ 721 s Pruvem, kdežto v případě kyseliny acetylsalicylové byla nejkratší s látkou galenIQ™ 721 se stearanem hořečnatým.

Klíčová slova: přímo lisovatelný isomalt – mazadla – pevnost tablet v tahu – doba rozpadu tablet

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Introduction

The interest in the use of sugar alcohols in pharmaceutical formulations has recently increased due to their sweetening capacity, a lower content of calories, and a non-cariogenic character. In addition, they can be administered to diabetic patients. Sugar alcohols may be used as fillers in normal as well as chewing tablets. After a suitable adjustment of tablets, e.g. by granulation or spray drying, they can serve as dry binders in directly compressible tablets¹⁾. The sugar alcohols employed for these purposes include also isomalt. Isomalt is a mixture of hydrogenated monosaccharides and disaccharides and its principal components are the disaccharidic alcohols 1-O- α -D-glucopyranosyl-D-mannitol dihydrate (GPM) and 6-O- α -D-glucopyranosyl-D-sorbitol (GPS). Isomalt possesses a half of the sweetening capacity of saccharose and low negative heat of dissolution, it is non-cariogenic and a small source of energy, as it is not absorbed in the small intestine, but fermented in the large intestine.¹⁾ Due to stable glycosidic bonds, it is resistant to chemical degradations and it does not produce a cooling effect in the mouth. Its agglomerated product, composed of small primary particles, was developed for direct compression. Isomalt adjusted in this way possesses excellent flowability, a high dilution potential, and low hygroscopicity²⁾. Ndindayno et al.³⁾ have demonstrated that the directly compressible properties of isomalt can be improved by melting. After melting, the crystalline form is converted into the amorphous one. The disintegration time of the tablet compressed from molten isomalt and the release of the active ingredient from the tablet are very quick. A problem is agglomeration caused

by recrystallization, where the amorphous phase passes to the stable crystalline phase in the presence of aerial humidity. This problem must be solved in further research.

The agglomerated isomalt product is manufactured by the German firm Palatinit GmbH under the name of galenIQ™, types 720 and 721 being intended for direct compression. The individual types differ in the ratios of basic disaccharidic alcohols; in type 720 the ratio of GPM and GPS is 1 : 1 and in type 721 this ratio is 1 : 3, and due to this fact it is more soluble in water²⁾. The present paper studies the properties of tablets, in particular the tensile strength and disintegration time in the above-mentioned types of directly compressible isomalt in dependence on compression force, addition of two types of lubricants in two concentrations, and two types of active ingredients in one concentration.

EXPERIMENTAL PART

Materials

galenIQ™ 720 – agglomerated isomalt (1-O- α -D-glucopyranosyl-D-mannitol dihydrate and 6-O- α -D-glucopyranosyl-D-sorbitol in the ratio of 1 : 1) (Palatinit GmbH, Germany); galenIQ™ 721 – agglomerated isomalt (1-O- α -D-glucopyranosyl-D-mannitol dihydrate and 6-O- α -D-glucopyranosyl-D-sorbitol in the ratio of 1 : 3) (Palatinit GmbH, Germany); Pruv® – sodium stearyl fumarate (J. Rettenmaier & Söhne GmbH + Co,

Table 1. Tableting materials without active ingredients

Raw material	Tableting material									
	1	2	3	4	5	6	7	8	9	10
galenIQ™720	+		+	+	+	+				
galenIQ™721		+					+	+	+	+
0.5% Mg st.			+				+			
0.5% Pruv				+				+		
1% Mg st.					+				+	
1% Pruv						+				+

Mg st. – magnesium stearate

Table 2. Tableting materials with active ingredients

Raw material	Tableting material							
	11	12	13	14	15	16	17	18
galenIQ™720	+	+	+	+				
galenIQ™721					+	+	+	+
1 % Mg st.	+		+		+		+	
1 % Pruv		+		+		+		+
50% Ac. ac.			+	+			+	+
50% Asc. ac.	+	+			+	+		

Mg st. – magnesium stearate, Ac. ac. – acetylsalicylic acid, Asc. ac. – ascorbic acid

Rosenberg, Germany); magnesium stearate (Acros organics, New Jersey, USA); ascorbic acid (Northeast General Pharmaceutical Factory, China); acetylsalicylic acid (Merck KGaA, Darmstadt, Germany).

Preparation of tableting materials and tablets

The tests included tableting materials from dry binders without lubricants, mixtures with lubricants, and mixtures with active ingredients. Altogether 18 tableting materials of the following composition were used (Table 1 and 2).

Dry binders with lubricants were mixed for 5 minutes in a stainless steel cube KB 15S (Erweka GmbH, Hausenstamm, Germany). Mixtures with active ingredients were prepared by mixing the dry binder and the active ingredient for 5 minutes, then the lubricant was added and the material was mixed for 5 minutes again. The rotation rate of the mixing cube was always 17 revolutions/minute. The amount of prepared tableting materials without active ingredients was always 30 g, with active ingredients, 20 g.

All tableting materials were used to produce 16 tablets compressed with the use of a special die with an upper and a lower punch on a material testing equipment T1 – FRO 50 TH.A1K Zwick/Roell (Zwick GmbH & Co, Ulm, Germany). Proper compaction took place by applying the pressure on the upper punch. The tablets were of a cylindrical shape without facets with a diameter of 13 mm and weight of 0.5 ± 0.0010 g. Compression velocity was 1 mm/s and compression forces 6, 8 and 10 kN, in the case of pure dry binders, 8 kN. Mixtures including active ingredients were compressed using only a compression force of 10 kN.

Measurement of tensile strength of tablets and evaluation of the lubricant sensitivity of tableting materials

Tensile strength was always evaluated in 10 tablets, first no sooner than 24 hours after compaction. Measurements were performed on a Schleuniger apparatus (Dr. Schleuniger Pharmatron AG, Solothurn,

Table 3. Values of LSR for galen IQ 720 and 721 at the compression force of 8 kN

Lubricant	galen IQ 720	galen IQ 721
0.5 % Mg st.	-0.1469	0.0151
1% Mg st.	-0.0563	0.0016
0.5% Pruv	-0.0649	0.0601
1% Pruv	-0.0134	0.0277

Mg st. – magnesium stearate

Switzerland), which measured tablet sizes accurate to 0.01 mm and destruction force in N. Tensile strength of tablets was calculated according to Eq. [1]:

$$P = \frac{2F}{\pi \cdot d \cdot h}, \quad [1]$$

where P is the tensile strength of tablets (MPa), F is the destruction force (N), d is the tablet diameter [mm], and h is the thickness of the tablet (mm)⁴⁾.

LSR (lubricant sensitivity ratio) values, which make it possible to quantify and mutually compare the lubricant sensitivity of tableting materials, were calculated according to Eq. [2]:

$$LSR = \frac{(C_{su} - C_{sl})}{C_{su}}, \quad [2]$$

where C_{su} is the crushing strength of tablets without an added lubricant and C_{sl} is the crushing strength with a lubricant. The more this value approaches 1, the more the dry binder is sensitive to an added lubricant from the viewpoint of decreased strength of tablets⁵⁾. In the present paper, the values of tensile strength, not those of crushing strength, are used in the equation.

Measurement of disintegration time of tablets

Disintegration times of tablets were evaluated earliest 24 hours after compaction always in 6 tablets. Measurements were performed on an apparatus for the determination of disintegration time of tablets Erweka ZT 301 (Erweka GmbH, Hausenstamm, Germany) following the method described in the chapter *Pharmaceutical Technical Procedures* in the *Ph. Eur. 2005*. The test was carried out without discs in the medium of purified water tempered to $37 \text{ }^\circ\text{C} \pm 1 \text{ }^\circ\text{C}$. The tablet was considered disintegrated at the moment when there was no remainder on the net.

The results of strengths and disintegration times were statistically processed by means of the computer programmes Excel and Qcexpert. Elementary data analysis yielded the mean values with standard deviations, which were plotted into dependences on compression force. In the cases of unclear significance of differences in the values, unpaired t-test at a level of significance of 0.05 was employed.

RESULTS AND DISCUSSION

The paper examined the properties of tablets, tensile strength and disintegration time, from the directly compressible isomalt galenIQ 720 and 721 in dependence on compression force. The impact factors were additions of two types of lubricants in two concentrations and two types of active ingredients in one concentration. The compression forces were 6, 8 and 10 kN and they were selected in such a way to make the tensile strength of tablets made of isomalt with lubricants oscillate within the range of the

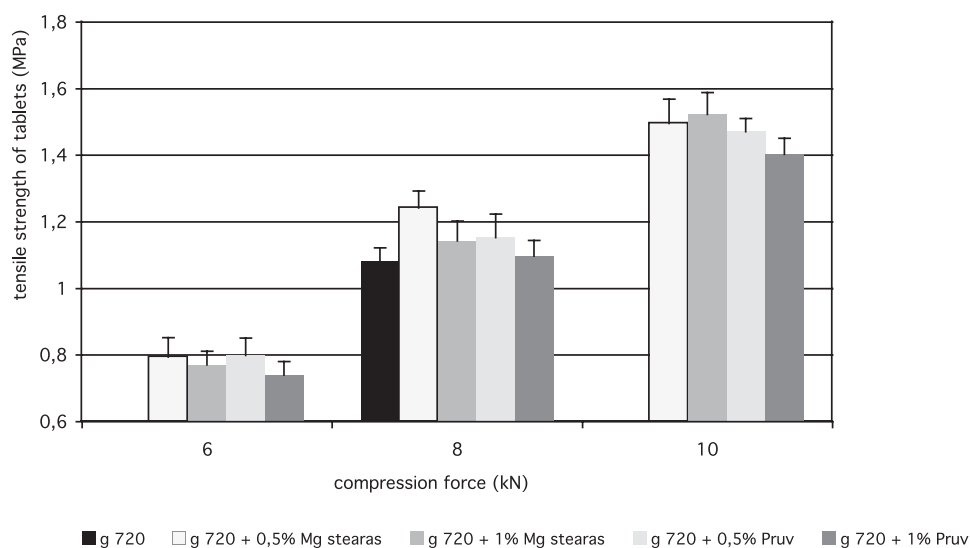


Fig. 1. Tensile strength of tablets in function of compression force: galen IQ 720
g 720 – galenIQ 720, Mg stearas – magnesi stearas

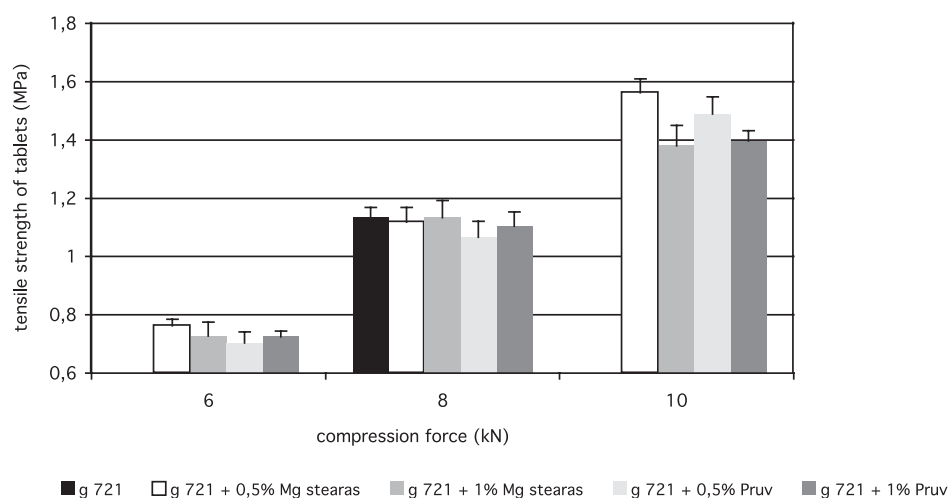


Fig. 2. Tensile strength of tablets in function of compression force: galen IQ 721
g 721 – galenIQ 721, Mg stearas – magnesi stearas

optimal strength of tablets (0.56–1.11 MPa)⁶⁾. The mixtures with active ingredients were compressed using only the compression force of 10 kN. The lubricants used were magnesium stearate and sodium stearyl fumarate in the concentrations of 0.5 and 1%. The model active ingredients were ascorbic acid and acetylsalicylic acid.

Figure 1 shows the dependence of tensile strength of tablets on compression force for galenIQ 720. In the compression force of 8 kN, it also states the value of strength for pure substance, which with the use of this compression force could be still compacted without a lubricant, though there was a problem of its sticking to the matrix. This value was intended to serve for the evaluation of the influence of lubricants on the strength of compacts, which should be theoretically decreased due to the plastic behaviour of isomalt¹⁾. However, there was no decrease in strength due to the lubricants. The tablets with a 0.5% addition of lubricants were stronger than those with a 1% addition. The lowest strength was

found in the tablets containing 1% of Pruv, the value of strength for the compression force of 8 kN not being statistically significantly different from the value for the pure substance. The values of strength of tablets increased with compression force. Luhn⁷⁾ examined also the influence of the concentration of magnesium stearate and the length of the period of mixing on tablet strength in the substance galenIQ 720. He found that the length of the period of mixing longer than 15 minutes resulted in lower tablet strength, but with increased concentration of the stearate from 0.5 to 2% there was no influence on the strength.

Figure 2 represents the identical dependence for galenIQ 721. Neither in this case the strength of tablets was influenced by lubricants, only the value for the mixture with 0.5% of Pruv was statistically significantly lower. In the compression forces of 6 and 10 kN, there is a statistically significant difference only between the strengths of compacts with 0.5% of lubricants, the compacts containing 0.5% of

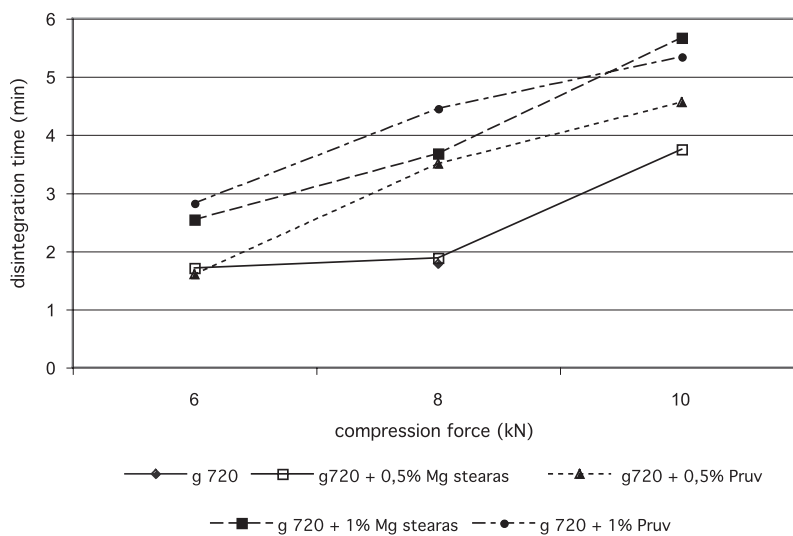


Fig. 3. Disintegration time in function of compression force: galen IQ 720
g 720 – galenIQ 720, g 721 – galenIQ 721, Mg stearas – magnesiū stearas

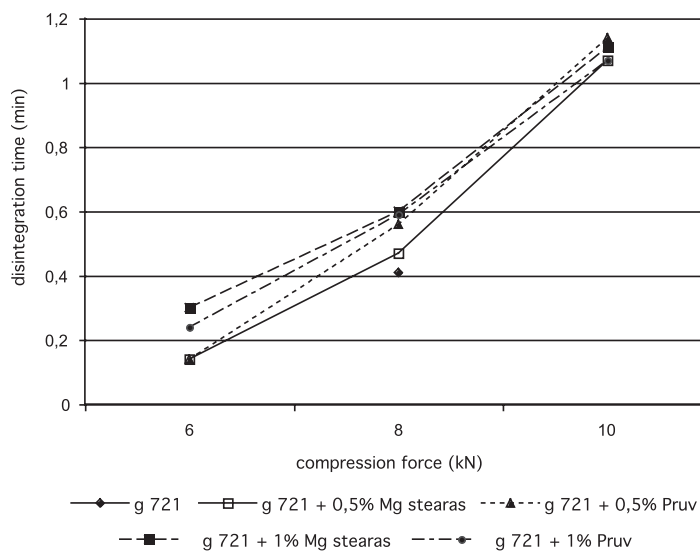


Fig. 4. Disintegration time in function of compression force: galen IQ 721
g 720 – galenIQ 720, Mg stearas – magnesiū stearas

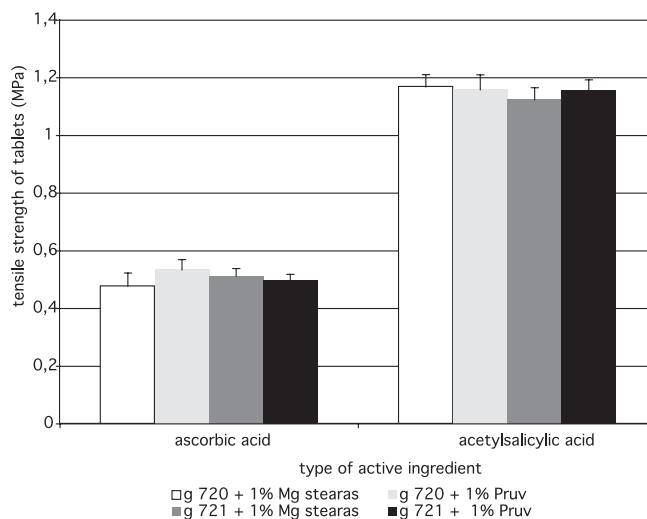


Fig. 5. Tensile strength of tablets at the compression force of 10kN: galen IQ 720 and 721 with active ingredients
g 721 – galenIQ 721, Mg stearas – magnesiū stearas

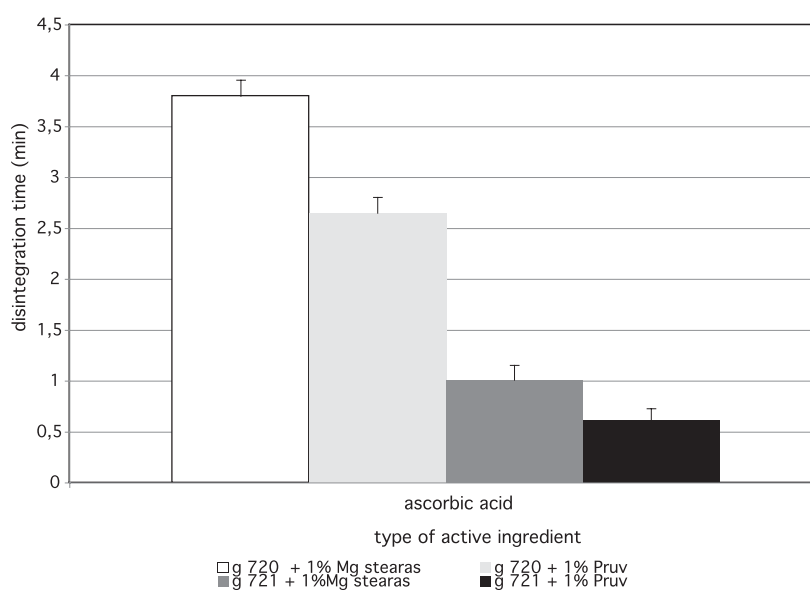


Fig. 6. Disintegration time of tablets at the compression force of 10 kN: galenIQ 720 and 721 with ascorbic acid
g 720 – galenIQ 720, g 721 – galenIQ 721, Mg stearas – magnesií stearas

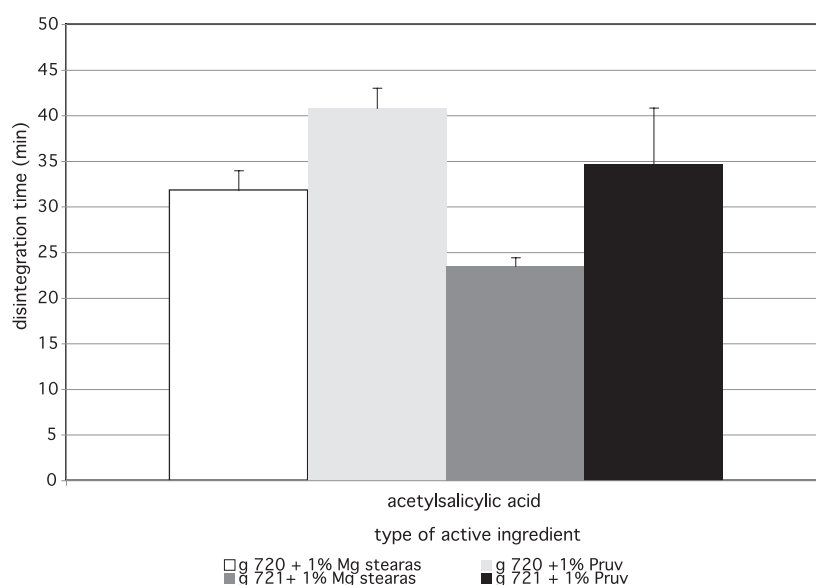


Fig. 7. Disintegration time of tablets at the compression force of 10 kN: galenIQ 720 and 721 with acetylsalicylic acid
g 720 – galenIQ 720, g 721 – galenIQ 721, Mg stearas – magnesií stearas

magnesium stearate being stronger. In tablets containing 1% of lubricants, there is no statistically significant difference in the values of strength within the type of the lubricant used. The strength of tablets was increased with compression force.

As follows from the comparison of both graphs, there is no marked difference between the strengths of tablets made from both types of isomalt. No effect of the lubricant on tablet strength is confirmed by the LSR (lubricant sensitivity ratio) values presented in Table 3. The more the value approaches 1, the higher sensitivity of the dry binder to the lubricant is observed. In the substance galenIQ 720, the LSR values were minus, the strength of tablets with lubricants was even marginally higher.

Figure 3 shows the dependence of disintegration times of tablets on compression force for galenIQ 720. A longer disintegration time was observed in the tablets with 1% of lubricants, the shortest disintegration time was found in the compacts with 0.5% of magnesium stearate, the value in the compression force of 8 kN not being statistically significantly different from the value for the pure substance.

Figure 4 describes the identical dependence for galenIQ 721. Disintegration time was prolonged by the action of lubricants.

The comparison of the values of disintegration times presented in the Graphs shows a marked difference in the disintegration times of the compacts made from both substances. The compacts containing the substance

galenIQ 721 unambiguously possessed a shorter disintegration time due to a higher content of GPS, which is more soluble in water than GPM¹⁾.

Figure 5 lists the values of the strength of compacts containing active ingredients at the compression force of 10 kN. The presence of active ingredients decreased the strength of tablets. The strength of tablets containing acetylsalicylic acid was higher, which could be theoretically assumed, due to better compressibility of the active ingredient resulting from the prevailing mechanism of compaction by plastic deformation. Ascorbic acid is compressed by fragmentation of its particles and this mechanism does not guarantee good compressibility and strength of compacts⁸⁾. Within the framework of the employed dry binder and lubricant, there were no marked differences between the values of strength of tablets.

Figures 6 and 7 show the values of disintegration times of compacts with ingredients. The Graphs are divided into two, as the values of disintegration times of compacts containing ascorbic acid are markedly lower due to its good solubility in water. In the compacts with ascorbic acid, the tablets containing the substance galen IQ 720 possessed a longer disintegration time due to its lower solubility. In both cases, the compacts with stearate disintegrated longer than those with Pruv. The tablets containing acetylsalicylic acid possessed long disintegration times, high above the pharmacopoeial limit of classic tablets (15 min) because of bad solubility of acetylsalicylic acid. A shorter disintegration time was observed in the compacts containing galen IQ 721 with the use of the same lubricant. The disintegration time was longer in the case of both substances with less hydrophobic Pruv, though there were no marked

differences in the strengths of tablets. A longer disintegration time would be theoretically assumed in the tablets containing magnesium stearate, which is more hydrophobic.

The compressibility of both types of directly compressible isomalt is the same, but when we need prepare the tablets with a shorter disintegration time, then dry binder galen IQ 721 is more suitable use.

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ZPRÁVY

XLIX. symposium z historie farmacie a veterinární medicíny

Sekce dějin farmacie České farmaceutické společnosti a Společnosti pro dějiny věd a techniky, České farmaceutické muzeum, středisko Univerzity Karlovy v Praze, Farmaceutické fakulty v Hradci Králové, Česká lékárnická komora a Veterinární a farmaceutická univerzita Brno pořádají dne 4. listopadu 2009 od 10 hodin do 16 hodin XLIX. symposium z historie farmacie a veterinární medicíny na téma **Významné farmaceutické a veterinární osobnosti u nás i ve světě.** Symposium se koná v Kabinetu dějin Ústavu cizích jazyků a dějin veterinárního lékařství v budově č. 32 kampusu VFU Brno, Palackého 1-3 – podkroví.

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