Chapter 2

Tensile strength of tablets containing two materials with a different compaction behaviour

Summary

The tensile strength of tablets compressed from binary mixtures is in general not linearly related to the strength of tablets prepared from single materials; in many cases it shows a decreased tensile strength relative to interpolation. This chapter evaluates the compaction behaviour of powder blends consisting of the two model materials used in this study. Sodium chloride and pregelatinised starch are both plastically deforming materials, but have a different densification and relaxation behaviour. The yield pressure of the binary mixtures shows an almost linear relationship. As a result of their lower yield pressure, starch particles yield earlier than sodium chloride particles. The following enclosure prevents some sodium chloride particles from yielding or cracking. The relaxation of the tablets is higher than the relaxation calculated by linear interpolation of the relaxation behaviour of the two pure materials. The difference between the measured porosity expansion and the data obtained by linear interpolation can be considered as a measure for the reduced interparticle bonding. SEM photographs indicate that the reduced interparticle bonding is caused by low adhesive forces. The measured decrease in tensile strength of the tablets is also considered to be the result of reduced interparticle bonding. In this chapter it is shown that a similar relationship exists between the tensile strength reduction and the percentage of starch on the one hand and the extra porosity expansion and the starch percentage on the other hand.

Keywords: tablet strength; binary mixture; densification; interparticle bonding; relaxation.

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2.1 Introduction

Tablets produced in the pharmaceutical industry commonly consist of more than one component. The mechanical strength of a tablet depends on both formulation and processing parameters. Even with constant processing parameters, the strength of tablets compressed from binary mixtures can often not be predicted from the compaction properties of the starting materials, because interactions between the materials may occur during the compaction process (1).

Tablet formation is the result of densification, interparticle bonding and relaxation. Each component has its own compaction behaviour. After rearrangement of the particles and building up of the pressure, the particles of the single materials behave brittle or viscoelastic (2, 3). Additionally, the deformation behaviour depends on other factors like particle size in relation to its critical brittle/ductile transition particle size (4) and compaction velocity (5). After release of the applied load, the tablet shows a certain amount of relaxation before reaching its final porosity.

Several investigators have examined the compaction behaviour of binary mixtures. The effects of different parameters, such as compaction behaviour (6), surface area (7) and compaction speed (8, 9), on the final tensile strength of the tablets compressed from blends have been reported. In many cases, the strength of the tablets compressed from the blend is found to be lower than the strength calculated by interpolation of the strength of tablets prepared from the pure materials.

The composition of the powder mixture is directly related to a number of tablet properties, such as tablet strength. Furthermore, it is known that the tensile strength of tablets correlates with the porosity according to the Ryshkewitch-Duckworth equation (10). To compare the tensile strength of tablets compressed from different blends of two materials, it is necessary to use volume fractions instead of weight fractions, because it may be expected that the three-dimensional structure of the tablet is more important than the weight ratio of the components (11). Moreover, a constant porosity is necessary to compare the tensile strength of the tablets of the mixtures with different volume percentages.

This study was performed to investigate differences between tensile strength of tablets compressed from blends of two materials and the tensile strength of tablets compressed from the individual components. These differences will be related to differences in densification, interparticle bonding and tablet relaxation. The materials chosen (sodium chloride and pregelatinised starch) are two plastically deforming materials, but with different densification and relaxation behaviours (12). The final tablet porosity of sodium chloride is almost completely determined by the
densification stage, whereas relaxation hardly occurs. In contrast, the ultimate tablet porosity of pregelatinised starch is the effect of both densification and relaxation behaviour (12).

### 2.2 Materials and methods

The materials used were sodium chloride (glidant-free material, Akzo Nobel, Hengelo, The Netherlands) and pregelatinised potato starch (Paselli WA4, Avebe, Foxhol, The Netherlands). Before use sodium chloride was milled (Moulinex, Birmingham, England) for a short period of time. From both materials the sieve fraction between 106 and 212 µm was obtained using an Air Jet Sieve (Alpine, Augsburg, Germany) equipped with USA Standard testing sieves (W.B Tyler, Mentor, OH, USA). Before use the materials were stored at a temperature of 20°C and a relative humidity of 60% for a period of at least 4 days.

The true density of the 106-212 µm particle size distribution was measured by helium pycnometry (Quantachrome, Syosset, N.Y., USA) at a temperature of 20°C. The true densities of sodium chloride and pregelatinised potato starch were 2239 kg m⁻³ and 1486 kg m⁻³, respectively.

The binary mixtures were prepared in volume percentages; mixing was performed with a Turbula mixer, model 2P (W.A. Bachofen, Basle, Switzerland) at 90 rpm for a period of 15 minutes.

Tablets (round, flat, diameter 13 mm, weight 500 mg) were compressed on a compaction simulator (ESH, Brierley Hill, UK) with a constant compaction speed of 3 mm s⁻¹. The compaction pressures varied between 10 and 350 MPa. The upper punch displacements were sine waves with different amplitudes in order to obtain the different compaction pressures. The lower punch was stationary during compaction. The ejection time was always 10 s. After ejection the tablets were stored for at least 14 hours in a controlled climate chamber (Heraeus, Hanau, Germany) at a temperature of 20°C and a relative humidity of 60%.

Tablet dimensions were measured with an electronic micrometer (Mitutoyo, Tokyo, Japan) and the tablets were weighed on an analytical balance (Mettler-Toledo, Greifensee, Switzerland).

The crushing strength of the tablets was measured with a Schleuniger 4N strength tester (Dr. Schleuniger Productronic, Soloturn, Switzerland). Crushing strengths over 300 N were measured with the compaction simulator (13). The tensile strength (S) of the tablets was calculated according to Fell and Newton (14). The tensile strength was
calculated for at least 40 tablets with different porosities to determine the Ryshkewitch-Duckworth relation (10) between the tensile strength and the tablet porosity for every binary mixture containing a different percentage pregelatinised starch:

\[ S = S_0 \cdot e^{-k \varepsilon} \]  

(1)

where \( S_0 \) is the tensile strength at zero porosity, \( \varepsilon \) the porosity and \( k \) a constant. \( k \) describes the bonding capacity of a single material (15) and indicates the effect of a change in porosity on the tensile strength. From these Ryskewitch-Duckworth relations the tensile strength at a porosity of 15% was calculated for every blend containing a different percentage pregelatinized starch.

Yield pressure (\( P_y \)) of the two materials and the binary mixtures were derived from the force displacement profiles according to Heckel (16, 17). Linear interpolation was performed in the porosity range between 15% and 35%.

2.3 Results and discussion

2.3.1 Densification of the binary powder mixtures

During powder densification, the porosity of a powder bed decreases. If the rearrangement of particles of the different blends is considered as a non-dominant factor, the densification can be studied by both the porosity under load and the yield pressure. Fig. 2.1a reflects the porosity under load of mixtures of sodium chloride with the different percentages pregelatinised starch. The porosity under load, using compaction pressures to obtain tablets with a final porosity of 15% after relaxation, decreases. Moreover, the slope decreases upon increasing pregelatinised starch concentration. Obviously, the largest influence of the pregelatinised starch particles on the densification of the sodium chloride particles occurs at low starch fractions.

Fig. 2.1b shows the yield pressure of sodium chloride, pregelatinised starch and their binary mixtures. The yield pressure of the two single materials illustrates the difference in densification behaviour. As reported previously, sodium chloride has a significantly higher yield pressure than pregelatinised starch (12, 18). The high yield pressure of sodium chloride illustrates its difficult densification. The figure shows that the second component, pregelatinised starch, with the lowest yield pressure enhances the densification of the powder blends. This results in a lower porosity under pressure for the binary mixtures than for pure sodium chloride.
Fig. 2.1a. Porosity under load of different binary mixtures of sodium chloride and pregelatinised starch for obtaining tablets with a final porosity of 15%.

Fig. 2.1b. Yield pressure of sodium chloride, pregelatinised starch and their mixtures (symbols) and the yield pressure of the binary mixtures calculated by linear interpolation (dotted line).

If the two components densify independently, the yield pressure of a mixture ($P_{y,mix}$) can be described by linear interpolation (dotted line, Fig. 2.1b) of the yield pressures of each of the two materials ($P_{NaCl}, P_{starch}$):

$$P_{y,mix} = P_{NaCl} \cdot \phi_{NaCl} + P_{starch} \cdot \phi_{starch}$$  \hspace{1cm} (2)

with $\phi_{NaCl}$ and $\phi_{starch}$ being the volume fractions of sodium chloride and pregelatinised starch in the blends. The explanation of the small deviations from linearity of the data
in Fig. 2.1b with eq. 2 can be found from SEM-photographs of the cross-sections of the tablets (Fig. 2.2a-2.2c).

**Fig. 2.2a**

**Fig. 2.2b**

**Fig. 2.2c**

**Fig. 2.2. Cross-section of a tablet compressed from a binary mixture of sodium chloride and (a) 10% pregelatinised starch (b) 20% pregelatinised starch and (c) 40% pregelatinised starch (S = starch particles, N = sodium chloride particles).**
All visual pregelatinised starch and sodium chloride particles in binary mixtures containing 10% (Fig. 2.2a) or 20% pregelatinised starch (Fig. 2.2b) have yielded and/or are cracked. The SEM picture of tablets compressed from a blend containing 40% pregelatinised starch (Fig. 2.2c) shows that all the starch particles are deformed. However, a number of the sodium chloride particles still have their original shape and are not cracked, even though the compaction pressure was higher than the yield pressure. Obviously, the starch particles, which are arranged around the sodium chloride particles, prevent that these sodium chloride particles are affected to a large extent by the compaction process. So, the sodium chloride particles do not yield or crack. This explains that the measured yield pressures of the blends are somewhat lower than those described by eq. 2. The difference in yield pressure between the two components is responsible for this phenomenon. In a binary mixture of sodium chloride and pregelatinised starch under the given conditions, a sodium chloride particle will only yield if it is in direct contact with another sodium chloride particle. However, the majority of the particles in the blends do yield during the densification of the powder. Therefore, the deviations from linearity are small.

Although the particle size of sodium chloride is larger than its critical particle diameter of 35 µm (4), the particles possess a low degree of fragmentation at high pressures (19). Under high pressures and close packing of the particles, there is hardly any possibility for the particles to rearrange. It may therefore be assumed that this small fragmentation only has a minor influence on the eventual tensile strength.

### 2.3.2 Relaxation of the tablets

The relaxation of tablets can be quantified by an increase in tablet height or volume after compression (20). Van der Voort Maarschalk et al. (13) have suggested that tablet relaxation is the result of two phenomena: a decrease in material density and an increase in tablet porosity. In general, the compression-induced increase in material density is fully reversible. The increase in tablet porosity occurs during and after release of the pressure from the tablet. The porosity expansion ($\Delta \varepsilon$) is given by:

$$\Delta \varepsilon = \varepsilon - \varepsilon_{\min}$$

where $\varepsilon$ is the tablet porosity after relaxation and $\varepsilon_{\min}$ is the porosity under load. $\varepsilon_{\min}$ is zero if the porosity under load was negative as an effect of the increase of material density. Just as previously found for single materials (15), the porosity expansion for all the different blends of sodium chloride and pregelatinised starch was found to be independent of the compaction pressures used.
Fig. 2.3 depicts the relationship between the percentage pregelatinised starch and the porosity expansion for tablets compressed from blends of sodium chloride and pregelatinised starch. In contrast to tablets compressed from pure pregelatinised starch, tablets compressed from sodium chloride show only a small porosity expansion. Fig. 2.3 also shows that the porosity expansion of tablets compressed from the blends is not linearly related with the volume percentage pregelatinised starch, but shows a sharp increase at low fractions of pregelatinised starch. The effect of the addition of pregelatinised starch particles is higher than expected from linear interpolation of the porosity expansion.

![Graph showing porosity expansion](image)

**Fig. 2.3.** Porosity expansion of tablets compressed from sodium chloride, pregelatinised starch and their mixtures (symbols) and the porosity expansion calculated by linear interpolation (dotted line).

Under load, the particles of sodium chloride and pregelatinised starch are closely packed. During tablet relaxation, the tablet volume increases, resulting in an increase in size and number of pores. The formation of extra pores consequently leads to the formation of more free particle surfaces, which is an effect of partial or complete disconnection of particles. In a tablet compressed from a blend of sodium chloride and pregelatinised starch particles, there are three different types of particle-particle bonding that can, partially or completely, be disconnected:

1) the cohesive bonding between sodium chloride particles
2) the cohesive bonding between pregelatinised starch particles
3) the adhesive bonding between sodium chloride and pregelatinised starch particles.
The SEM-photographs (Fig. 2.2a-2.2c), especially the one of tablets compressed from the blend of sodium chloride and 40% pregelatinised starch (Fig. 2.2c), show that pregelatinised starch particles situated around the individual sodium chloride particles have an imprint of the enclosed sodium chloride particles. This implies that pregelatinised starch particles, which were in contact with sodium chloride particles under load, have been detached. This observation is a strong indication for the fact that the adhesion forces between the sodium chloride and pregelatinised starch particles are smaller than the two different cohesion forces. The observation of smaller adhesion forces can be explained by the fact that sodium chloride and pregelatinised starch have a different type of interparticle bonding (21).

### 2.3.3 Tablet strength

Fig. 2.4 shows the tensile strength of tablets with a porosity of 15% compressed from different binary mixtures of sodium chloride and pregelatinised starch as a function of the volume percentage pregelatinised starch. The figure shows that the tensile strength of tablets compressed from 100% pregelatinised starch is higher than that of tablets containing 100% sodium chloride. Furthermore, the strengths are lower than predicted by interpolation.

![Graph showing tensile strength of tablets](image)

**Fig. 2.4.** Tensile strength of tablets compressed from sodium chloride, pregelatinised starch and their mixtures (symbols) and the tensile strength calculated by linear interpolation (dotted line).

Strength reduction can be considered as the result of reduction in interparticle bonding. In line with our conclusions from the SEM-photographs, the decreased
tensile strength of tablets compressed from the blends indicates that the adhesive forces between the sodium chloride particles and the pregelatinised starch particles are smaller than the cohesive forces between particles of the pure materials. The reduction of interparticle bonding inside tablets compressed from different volume percentages sodium chloride and pregelatinised starch is reflected in Fig. 2.5 as the difference between the measured tensile strength and the calculated tensile strength (dotted line, Fig. 2.4).

Fig. 2.5. Difference between the measured tensile strength and calculated tensile strength of tablets compressed from sodium chloride with different volume.

Fig. 2.6. Extra porosity expansion of tablets compressed from sodium chloride with different volume percentages pregelatinised starch.

Fig. 2.6. Extra porosity expansion of tablets compressed from sodium chloride with different volume percentages pregelatinised starch.
A decreased tensile strength, caused by a reduction of interparticle bonding, is related to a larger relaxation of the tablets (15, 22, 23). Therefore, the extra porosity expansion between the measured porosity expansion and the porosity calculated by linear interpolation (Fig. 2.3) is suggested to be a measure for the reduced interparticle bonding. This extra porosity expansion is depicted in Fig. 2.6.

Both the decrease in tensile strength of tablets compressed from sodium chloride and pregelatinised starch (Fig. 2.5) and the extra porosity expansion (Fig. 2.6) are a reflection of the reduced interparticle bonding inside the tablets. Comparison of the figures shows that an almost similar relationship was found between the decrease in tensile strength and the percentage pregelatinised starch on the one hand and the extra porosity expansion of the tablets and the percentage pregelatinised starch on the other hand. Figures 2.5 and 2.6 have been combined into Fig. 2.7. This figure indeed suggests such an interrelation of the extra porosity expansion, the reduced interparticle bonding and the decrease in tensile strength.

![Graph showing difference in tensile strength as a function of extra porosity expansion.](image)

Fig. 2.7. Difference in tensile strength as a function of the extra porosity expansion.

In conclusion, the yield pressures of blends of sodium chloride with different percentages pregelatinised starch can be calculated by linear interpolation of the yield pressures of the two single materials. The porosity expansion of tablets compressed from the blends is higher than calculated from linear interpolation of the porosity expansion of tablets compressed from the pure materials. Both the extra porosity expansion and the decrease in tensile strength of the tablets compressed from the binary blends are related and caused by a reduction in interparticle bonding. Using the interparticle bonding reduction, caused by adhesion forces that are lower than the
cohesion forces, it is possible to predict the relaxation behaviour of the mixture using the properties of the individual components.

2.4 Nomenclature

\begin{itemize}
  \item \( k \) \hspace{1cm} constant, bonding capacity
  \item \( P_y \) \hspace{1cm} yield pressure
  \item \( P_{y,mix} \) \hspace{1cm} yield pressure of binary mixture
  \item \( P_{NaCl} \) \hspace{1cm} yield pressure of sodium chloride
  \item \( P_{starch} \) \hspace{1cm} yield pressure of pregelatinised starch
  \item \( S \) \hspace{1cm} tensile strength
  \item \( S_0 \) \hspace{1cm} tensile strength at zero porosity
  \item \( \varepsilon \) \hspace{1cm} porosity
  \item \( \varepsilon_{min} \) \hspace{1cm} porosity under load
  \item \( \Delta \varepsilon \) \hspace{1cm} porosity expansion
  \item \( \phi_{NaCl} \) \hspace{1cm} volume fraction of sodium chloride
  \item \( \phi_{starch} \) \hspace{1cm} volume fraction of pregelatinised starch
\end{itemize}

2.5 References


