Looking into tablets, Characterization of pore structure in tablets using image analysis
Wu, Yu San

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2008

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):
Chapter 2

Location dependent analysis of porosity and pore direction in tablets

Abstract

Several phenomena in tablets indicate that an inhomogeneous pore distribution is formed during the compaction process. Examples are lamination and the capping of corners. In order to gain an understanding of the relation between structure and compact properties, analyzing the structure in a location dependent manner would be extremely useful. Our aim was to visualize and to quantitatively analyze the pore distribution in compacts.

This was done by embedding a cubic (sodium chloride) compact with polymer, allowing the compact to be cut without disrupting the structure. By doing so, it was possible to make scanning electron microscopic images from different angles at different locations in the compact. These images were made binary with a two-means cluster algorithm (Isodata) after which the porosity could be calculated. Counting the number of transitions from the pixels in the pores to the pixels in the sodium chloride particles in two perpendicular directions allows us to construct a measure for the anisotropic connectivity of the particles.

The results show an increase in porosity towards the bottom of the compact and showed a preferred orientation of the pores in the direction of compression. The proposed method is suitable for analyzing the pore distribution quantitatively and for evaluating anisotropy.

2.1 Introduction

Despite the popularity of tablets as a dosage form, problems during tabletting and with the produced tablets occur frequently. Since failures such as lamination and the capping of corners occur in specific locations of the tablet, inhomogeneity in (pore) structure is expected. Inhomogeneity in structure cannot be evaluated with the current methods such as mercury porosimetry and gas adsorption. These methods consider the compact as a whole and discard important local variations. For a better understanding of compact properties, analyzing structure as a function of location within a compact is quite useful.

An important parameter describing the structure is the pore network in a compact. Ryshkwitch described the relation between bulk porosity and tensile strength, stating that there is a linear relation between the bulk porosity and the logarithm of the tensile strength [1]. However, this does not explain all differences in tensile strength. It has been hypothesised that the pore shape also has an important influence on tensile strength [2, 3]. Thus, it is desirable to characterize the pore distribution on a smaller scale than an entire compact. One factor which could then be evaluated is the anisotropy, meaning that a property is dependent on the direction in which it is measured. Anisotropy in structure is suggested by anisotropy in tensile strength, as reported by several authors [4-7]. Ando et al. [4] made pictures of cross sections of the tablets revealing anisotropy in structure, but no quantitative information was obtained [4].

Obtaining detailed information about pore distribution is difficult. Current methods have the disadvantage that they only measure porosity and pore size distribution of an entire compact; no information about the shape or the direction of the pores is obtained. Modern techniques such as X-ray computed tomography enable visualization of the pore morphology as demonstrated in granules [8, 9], but the current resolution of this method (approximately 10 µm) is not sufficient, yet, for the evaluation of pore distribution (shape) in compacts, as the pore diameter is typically in the order of 1 µm or even less [10-12].

In order to gain a fundamental understanding of compact properties in relation to pore distribution, a more detailed characterization is needed. The development of a suitable method for this purpose forms the aim of the research described in this paper. The method presented here describes how pore distribution can be visualized and quantitatively analyzed.
2.2 Materials and methods

2.2.1 Experimental set-up

In order to investigate the pore distribution in the compact, different planes in a cubic sodium chloride tablet of 7x7x7 mm were analyzed. The tablets were made with uni-axial compression. A cube has the advantage that the dimensions of the tablet are the same irrespective of the direction of testing, thus making it easier to examine the tablet structure, for instance the anisotropy, and to check the validity of the imaging method. We examined the structure by making plan and elevation photos.

Plan photos provide a top view and are made when looking down on the tablet at different heights, in the same direction as compression was performed. This direction is defined as the z-direction. Plan photos were made of different planes in the tablets (figure 1a). The first plane is the upper surface of the cube. The other planes lie parallel to this plane at 1.75 mm, 3.5 mm, and 5.25 mm from the upper surface. These distances were arbitrarily chosen, but at equal distance from each other. In the plan photos, an x- and a y-direction were defined. Both directions lie parallel to the cube axes.

![Figure 1: The different planes in the compact of which images are made (a) Plan. (b) Elevation.](image)

Elevation photos provide a side view, observed from the compression direction, of the tablet structure. These photos were made of three different planes. The first plane is a randomly chosen side plane of the cube. The other planes lie parallel to this plane at 2.3 mm and 3.5 mm from the side (see figure 1b). Only one half of the tablet was examined as plane symmetry was expected. In each of these planes an x- and z-direction are defined.
Of each plane 9 photos were taken. These images almost cover the whole plane. Figure 2a depicts which photo number corresponds to which position for the plan photos. Photo position 5 corresponds to the centre of the plane, while photos 1, 3, 7 and 9 correspond to the corner positions. For the elevation photos, the photos 1, 4, 7 are closest to the bottom of the tablet, while the numbers 3, 6 and 9 are closest to the top of the tablet as shown in figure 2b.

2.2.2 Materials
The 212-250 µm fraction of sodium chloride (Chemically pure quality, Akzo Nobel, Hengelo, The Netherlands) was obtained by 30 min. vibratory sieving (Fritsch analysette 3, Germany) followed by 12 min air jet sieving over a sieve of 212 µm (Alpine A200, Augsburg, Germany) to remove the fines. The true density of the 212-250 µm fraction as measured with helium pycnometry (Quantachrome, Syosset, New York, USA) was 2175 kg/m$^3$.

2.2.4 Tablet compaction
A specially manufactured die was used to compress the compacts. The die was square shaped, both sides being 7 mm. Before compaction, the die was lubricated with magnesium stearate using a brush. After filling the die with 530 mg sodium chloride, the powder was compacted with 2.25 kN using a hydraulic press (ESH compaction apparatus, Hydro Mooi, Appingedam, The Netherlands). The rate of compaction was 0.5 kN/s and the maximum pressure was maintained for 0.1 s. Compact dimensions were measured after 24 hours with an electronic micrometer (Mitutoyo, Tokyo, Japan) and the weight of the compacts was measured with an analytical balance (Mettler-Toledo, Greifensee, Switzerland). From these data and the real density the final porosity was calculated. The target porosity of the tablets was 30%. The standard deviation of the porosities of all tablets was 0.3%. This porosity was chosen since it still resulted in manageable tablets with a high porosity assuring that the tablet could be completely infiltrated with polymer.
2.2.5 Embedding and cutting the slices
Compacts were embedded in glycol metacrylate (GMA). This technique is widely used for embedding of biological specimens and subsequent sectioning. A desired volume of Technovit 7100 (Kulzer, Germany) was carefully mixed with hardener I. Shortly before embedding the compact, a second hardener was added. The compacts were immersed with the mixture and completely infiltrated and subsequently stored at room temperature for two days.

To make photos of the designated planes in the compact, the appropriate quantity of sodium chloride/polymer matrix was scraped off the sample with a lathe (Schaublin 125, Esmeijer, Rotterdam, The Netherlands). After scraping off the material with the lathe, a microtome (Reichert-Jung 2050, Vienna, Austria) was used to slice off approximately 1 µm material to obtain a smooth surface. These surfaces were used for scanning electron microscopic (SEM) inspection.

2.2.6 Images
The images were made using back scattered electron imaging. With this technique, also used by Sriamornsk [13], a good contrast is achieved between regions with sodium chloride and regions where the polymer is present. This is due to the fact that elements with a higher atomic number (i.e. the sodium and chloride in our case) deflect more electrons than elements with a lower atomic number (i.e. the polymer). Therefore the elements with a lower atomic number are represented by the darker regions in the image.

SEMs were made using a JEOL scanning electron microscope (JEOL, type JSM-6301F, Japan) operated at an accelerating voltage of 10 kV. The diaphragm was 50 µm, spot size 6 and the working distance was 15 mm. Back scatter was used (standard backscatter detector, JEOL, Japan). Samples were not coated and the magnification was 60x.

2.2.7 Image analysis: local porosity
Each picture used for calculation of the local porosity had a size of 1676 x 1676 pixels. Matlab 6.5.1 (The MathWorks Inc, Natick, USA) and the Dipimage toolbox (Quantitative Imaging Group, Faculty of Applies Sciences, Delft University of Technology, the Netherlands) [14] were used for the image analysis. With this software, the grey-scale images were made binary with the Isodata threshold algorithm [15]. This algorithm calculates the best threshold value out of the grey value histogram of the image.
The binary images consist of two phases. One phase, ‘salt phase’, corresponds to the surface area containing the sodium chloride particles. The other phase, the ‘pore phase’, corresponds to the places where the polymer had replaced the air in the original pores in the compact. In each binary image the total number of pixels in the pore phase was counted. With this number the local porosity was calculated:

\[
\varepsilon_l = \frac{N_{\text{pixel}_\text{pore}}}{N_{\text{pixel}_\text{image}}} \times 100\% \tag{1}
\]

in which:

- \(\varepsilon_l\) Local porosity in percentage
- \(N_{\text{pixel}_\text{pore}}\) Number of pixels in the pore phase
- \(N_{\text{pixel}_\text{image}}\) Number of pixels in the image

The plan photos at a tablet depth of 1.75 mm were made without using the microtome first. Therefore, the calculated local porosities of these images were corrected by adding the mean difference between the local porosity before and after treatment with the microtome. This difference was obtained from the plan photos made at a tablet depth of 3.5 mm of which plane both before and after treatment with the microtome photos were taken (90 photos in total).

### 2.2.8 Image analysis: number of transitions

Each picture used for calculating the number of transitions (\(N_T\)) was a square image of 1676x1676 pixels. A Gaussian filter (\(\sigma = 4\) pixels) was applied to suppress noise and surface artefacts caused by the microtome before making the images binary with the isodata threshold algorithm. Both these operations were carried out with the DIPimage toolbox. After this step the number of transitions from the salt phase to the pore phase and vice versa was calculated. This was done along each pixel line, meaning that for each photo the number of transitions along 3352 lines was counted; in the plan photos this was done along each pixel line in the x direction and in the y direction, and in the elevation photos this was done in the z direction and in the x direction. Counting the number of transitions is similar to the ‘line intercept count’, a method used in stereology [16] that gives information about the structure of a sample. After calculating the number of transitions, a quotient was calculated by dividing the \(N_T\) in one direction by the \(N_T\) in the other direction. For the plan photos this meant:
Chapter 2. Pore distribution in tablets

\[ Q_P = \frac{N_{Ty}}{N_{Tx}} \]  

(2)

And for the elevation photos:

\[ Q_E = \frac{N_{Tz}}{N_{Tx}} \]  

(3)

In which:

- \( Q_{[P,E]} \) Quotient of transitions in [Plan,Elevation] image(s)
- \( N_{Ty} \) Number of transitions in the y-direction
- \( N_{Tx} \) Number of transitions in the x-direction
- \( N_{Tz} \) Number of transitions in the z-direction

The method is illustrated by the pictures in figure 3. In both pictures the total number of squares is \( 8 \times 8 = 64 \). Both pictures have 12 black squares. If the black squares represent the pore phase and the white squares represent the salt phase, the local porosity is in each case 18.8%. However, the quotient of the number of transitions counted from left-to-right and from top-to-bottom, is different for the two pictures: the number of transitions for the picture in figure 3a is 8 counted from left to right and 24 counted from top to bottom. This gives a quotient of 0.33. For the picture in figure 3b, the quotient is \( 20/20 = 1 \). This means that the distribution of black squares is different for the two pictures. It can also be seen that when the quotient deviates from 1, the sample shows anisotropy.

Figure 3: (a) Square divided in 8x8 blocks. Percentage black is 19% and there is a preferential direction. (b) Square divided in 8x8 blocks. Percentage black is 19%, there is no preferential direction for the black squares.
2.3 Results and discussion

2.3.1 Experimental procedure
The chemical formula of glycol methacrylate makes it very unlikely that sodium chloride dissolves in this embedding medium. That the particles do not dissolve is also suggested by the still sharp edges of the sodium chloride particles seen in the SEM images. If dissolution had taken place, smooth edges would have been observed as sharp corners dissolve first. After embedding, tablet dimensions stayed the same, also supporting the assumption that the structure is preserved during embedding.

2.3.2 Photos: visual observation
Figure 4a is an example of a plan photo. There is a good distinction between the salt particles and the spaces where the polymer is present (pores). The observed shape of the particles is mainly rectangular. Some particles lie separate while others are connected to each other. There does not seem to be a preferential direction of either the pores or the particles. Figure 4b shows an elevation photo. Again there is a clear distinction between the salt-phase and the pore-phase and, just as in the plan photo, the particle’s shape is mainly rectangular. However, a difference in structure is discernable. Contrary to the structure in the plan photos, there does seem to be a preferential direction in the elevation photos. Here the particles seem to be connected to each other mainly in the z-direction. However, it is clear that just from the observation the anisotropy is not quantified. Figure 4a and figure 4b are both photos taken from the upper surface of the tablets. Photos taken at the different tablet depths show similar structures.

Figure 4: (a) Plan photo of the upper surface (b) Elevation photo of the upper surface.
2.3.3 Local porosity: plan

Figure 5 shows the local porosities for photo positions 1, 4 and 5. These graphs are representative for the other photo positions. There are hardly differences between the local porosities at the different photo positions. However, differences between the local porosities at different compact depths, compact depth being the distance to the top surface measured in the direction of compression, are considerable. The local porosity increases with increasing compact depth.

The increase in local porosity with increasing compact depth is caused by the fact that the applied compaction pressure is not fully transmitted to lower regions, due to friction between salt particles and the die wall and between the particles themselves [17]. This implies that for uni-axial compression, with a stationary lower punch, a density gradient can be found with a higher density at the top of the compact and a lower density in the bottom. Several authors have reported this [17-20], but this effect has not been visualized so far. Another observation that can be made is that the pattern of increasing local porosity with increasing compact depth is similar for all photo positions. This is in contradiction with results of other authors who found higher densities in the top corners compared to the centre of the top layer [17, 18, 21]. The discrepancy is probably caused by the use of different materials. Train used magnesium carbonate [18], Briscoe et al. used an alumina powder [21] and Ozkan et al. used alumina agglomerates [17]. These materials have different properties compared to the sodium chloride used in this study. The use of a different shape of punch and die, and low compression force, can also be the cause of the deviations from previously reported results.

![Figure 5: Surface porosity of the plan photos. Points are the average of 5 measurements, error bars are standard deviations. Insert shows photo positions of which the results are shown.](image-url)
2.3.4 Local porosity: elevation

Figure 6 shows mean local porosities and standard deviations of the elevation photos. This is shown for the plane in the middle of the compact at 3.5 mm from the side, but the results are representative for the other planes. The line marked ‘side A’ represents the local porosities of the photo position 1, 2 and 3. The line ‘centre’, is the result of the measurements from photo position 4, 5 and 6 and the line ‘side B’ is calculated with positions 7, 8 and 9. The compact depth is the mean tablet depth of the corresponding photos. At all planes the local porosities in the tablet columns side A and side B are not different from the local porosity in the tablet column centre.

Contrary to the plan photos, there does not seem to be a relation between local porosity and compact depth in the elevation photos. This is probably because the compacts are made with uni-axial compression. Since the direction of compression is the z-direction, voids between particles perpendicular to this direction are closed first during compression. It can be argued that pores parallel to the z-direction remain open at this compression force. In the elevation photos these pores are the main pores in the images (see for example figure 4b). Apparently the pores in the z-direction exist over the whole tablet depth, explaining the lack of porosity gradient with increasing tablet depth in the elevation photos. Both plan and elevation photos show no difference in local porosity between corner positions or the more central parts of the compacts.

![Figure 6: Surface porosity at different tablet depths for the elevation photos at 3.5 mm from the side. Points represent the average of 5 measurements, error bars are stand. dev. Insert shows photo positions of which the results are shown.](image-url)
2.3.5 Anisotropy

The quotient of transitions, $Q_{[P,E]}$ (section 2.2.7), has been the basis to quantify the preferential direction of the pores in the tablets. Figure 7 shows the mean values and standard deviations for the plan and elevation planes at 0 mm and 3.5 mm tablet depth. The quotient of transitions in the plan photos does not significantly differ from “unity” in both tablet locations, meaning that there is no preferential direction of the pores in either the x-direction or y-direction. This is in agreement with the image shown in figure 4a. However, in the elevation photos, the quotient of transitions is significantly lower than unity (one sample t-test, $p<0.001$). This implies that the number of transitions in the z-direction is lower than the number of transitions in the x-direction. Hence the pores and particle clusters are preferentially oriented in the z-direction. Particles are pressed together in this direction and it is therefore not surprising that the particles are mainly connected to each other in the z-direction.

Another approach to the same topic is a comparison of the plan photos with the elevation photos. For the upper surface the data show that the quotient number of transitions in the plan photos is significantly higher than in the elevation photos (two-tailed Student’s t-test, $p<0.001$). The same applies to the elevation and plan photos of the planes at 3.5 mm from the tablet surface. This means that the pore distribution as shown in the plan photos is different from the pore distribution as shown in the elevation photos, again proving anisotropy in pore distribution.

Figure 7: Quotient of transitions in plan and elevation images. Error bars represent standard deviations ($n=45$).
2.4 Conclusion

The method presented in this article is suitable for visualization and quantitative analysis of the pore distribution in tablets. By embedding a cubic sodium chloride compact with polymer it was possible to cut the tablets and make images of the inner structure. These images had a high resolution which made it possible to do a quantitative analysis. This analysis clearly showed a porosity gradient in the tablet with higher porosities towards the underside of the compact. In the elevation photos such a gradient was not found. Still, elevation photos proved their use, since those photos in combination with plan photos clearly showed anisotropy in the tablet structure. In addition, it could be determined that the preferential direction of the pores was the z-direction.

In this study sodium chloride was used as an excipient to introduce the method. The choice for this material showed that the density distributions in a compact as found by other authors [17, 18, 21] are not universally applicable.

Finally, the method described here seems to be feasible for pore characterization in standard shaped tablets in order to investigate production problems such as lamination and capping.
2.5 References


