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Comprehensive study of dynamic curing effect on tablet coating structure

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The dissolution method is still widely used to determine curing end-points to ensure long-term stability of film coatings. Nevertheless, the process of curing has not yet been fully investigated. For the first time, joint techniques were used to elucidate the mechanisms of dynamic curing over time from ethylcellulose (Aquacoat[®])-based coated tablets. X-ray micro-computed tomography (X μ CT), Near Infrared (NIR), and Raman spectroscopies as well as X-ray microdiffraction were employed as non-destructive techniques to perform direct measurements on tablets. All techniques indicated that after a dynamic curing period of 4 h, reproducible drug release can be achieved and no changes in the microstructure of the coating were any longer detected. X μ CT analysis highlighted the reduced internal porosity, while both NIR and Raman measurements showed that spectral information remained unaltered after further curing. X-ray microdiffraction revealed densification of the coating layer with a decrease in the overall coating thickness of about 10 μ m as a result of curing. In addition, coating heterogeneity attributed to cetyl alcohol was observed from microscopic images and Raman analysis. This observation was confirmed by X-ray microdiffraction that showed that crystalline cetyl alcohol melted and spread over the coating surface with curing. Prior to curing, X-ray microdiffraction also revealed the existence of two coating zones differing in crystalline cetyl alcohol and sodium lauryl sulfate concentrations which could be explained by migration of these constituents within the coating layer. Therefore, the use of non-destructive techniques allowed new insights into tablet coating structures and provided precise determination of the curing end-point compared to traditional dissolution testing. This thorough study may open up new possibilities for process and formulation control.

1. Introduction

Functional aqueous polymer coatings are frequently used to control drug release rates [1]. The film formation process is induced from the coalescence of polymer particles for which a post-thermal treatment, known as curing, must generally be conducted to ensure long-term stability [2,3]. Currently, *in vitro* dissolution testing is commonly used to determine curing end-points and evaluate the long-term stability of film coatings [4–6]. A comprehensive understanding of the mechanisms involved during film formation is lacking as are methods of characterizing film coatings. Some authors reported studies on isolated films to assess the influence of formulation and process parameters on film coating quality [7,8]. However, it was shown that film properties differ from those

of film coatings when applied to substrates [9]. The use of fast non-destructive techniques, allowing direct characterization of solid dosage forms during the curing process, is a welcome addition for the understanding of this crucial step.

In the present work, various techniques were investigated to detect subtle variations and elucidate changes in the tablet coating layer during curing. New insights into coating structures were gained using X-ray micro-computed tomography (X μ CT), X-ray microdiffraction, and Near Infrared (NIR) and Raman spectroscopies. Compared to scanning electron microscopy or mercury porosimetry, X μ CT enables non-destructive 3D visualization of the internal structure of samples [10,11], whereas X-ray diffraction allows a non-invasive means of probing information on crystallinity [12,13]. In this study, X-ray techniques were coupled to high-intensity synchrotron radiation, giving access to faster data acquisition and more accurate information relative to conventional X-ray sources [14]. NIR and Raman analyses are frequently employed to characterize solid dosage forms [15,16] and monitor

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real-time pharmaceutical processes [17]; very few NIR or Raman studies, however, describe spectral changes induced upon curing [18–20].

For this study, a functional polymer combination of ethylcellulose/poly(vinyl-alcohol)-poly(ethylene-glycol) graft copolymer, using triethylcitrate as plasticizer, was chosen as the coating material to modulate drug release rate from matrix tablets and thus acting as a diffusion barrier. Dissolution studies previously demonstrated stable controlled drug release profiles from pellets coated using this polymer combination, which were subjected to either static or dynamic curing conditions [21,22]. Curing is often performed under static conditions in an oven, but can be carried out preferably within the coating equipment under dynamic conditions to eliminate additional tablet transfer steps [22]. In the present work, dynamic curing was carried out at 60 °C and ambient relative humidity. Drug release properties from cured coated tablets were evaluated to confirm dynamic curing performances.

In accordance with current pharmaceutical development regulations based on ICH Q8 recommendations [23], the aim of this work was to determine curing end-point with accuracy to ensure film coating stability and thus improve the overall understanding of formulation and processing steps. A new insight into the coating structure of tablets subjected to different dynamic curing times was provided by combining results from the aforementioned techniques.

2. Materials and methods

2.1. Materials

Hydrophilic matrix tablets of biconvex shape (with a diameter of 8 mm, a radius of curvature of 7 mm, and an average mass of 200 mg), made from hypromellose and calcium hydrogenophosphate as main excipients, containing a freely soluble drug substance were supplied by Les Laboratoires Servier Industrie (Gidy, France).

Aqueous ethylcellulose dispersion (Aquacoat ECD 30[®], FMC Biopolymer, Philadelphia, USA), poly(vinyl-alcohol)-poly(ethylene-glycol) graft copolymer (Kollicoat IR[®], BASF, Ludwigshafen, Germany), and triethylcitrate (Citroflex 2[®], Morflex, Greensboro, USA) were purchased as coating materials.

The commercially available aqueous ethylcellulose dispersion also contains cetyl alcohol as emulsifier (concentration range: 1.7–3.3% w/w of total solids) and sodium lauryl sulfate as surfactant (concentration range: 0.9–1.7% w/w of total solids) [24]. Ethylcellulose powder (Ethocel[®] Standard 10, Dow Chemical Company, Midland, USA), cetyl alcohol (Sigma-Aldrich, Buchs, Switzerland), and sodium lauryl sulfate (Texapon[®] K 12 G PH, Cognis, Düsseldorf, Germany) were used as crystallographic and spectroscopic references.

2.2. Coating operation

The coating suspension was prepared with an Aquacoat ECD 30[®]:Kollicoat IR[®] ratio of 90:10 w/w (expressed as dry masses). Aquacoat ECD 30[®] was plasticized overnight with Citroflex 2[®] (1:4 w/w, based on ethylcellulose content), under continuous agitation (Rayneri[®] apparatus, 200 rpm). An aqueous solution of Kollicoat IR[®] was then added to adjust the final solid content at 20% (w/w). The actual coating suspension was mixed for an additional period of 30 min prior to coating and stirred throughout the coating operation (Rayneri[®] apparatus, 100 rpm). Approximately 4 kg of preheated tablets (20 min at 40 °C) were coated inside a partially perforated pan coater (Driacoater 500[®], Driam, Eriskirch, Germany) using a spray nozzle with a diameter of

0.8 mm at a rotation speed of 20 rpm. The inlet temperature was set at 63 °C to maintain a product temperature of 36 °C during coating. Tablet bed temperature was controlled using a hand-held infrared temperature probe, regularly placed at the surface of the tablet bed of the pan coater (Fluke 574, Fluke Corporation, Everett, USA).

The process was stopped after spraying polymer blend equivalent to 10% theoretical weight gain (w/w expressed as percentage of core tablet mass). At the end of the coating operation, a fraction of uncured tablets was collected, together with a sample of dried coating suspension removed from the inner surface of the pan coater.

2.3. Curing operation

Coated tablets remaining inside the pan coater were subjected to dynamic curing. The rotation speed was reduced to 10 rpm to prevent tablet damage. The inlet temperature was gradually increased to 65 °C to attain a steady product temperature of 60 °C throughout the curing operation. Tablet bed temperature was controlled using an infrared thermometer (Fluke 574, Fluke Corporation, Everett, USA). No additional water was sprayed. Tablets were collected at different time intervals over 6 h (0.5, 1, 2, 4, and 6 h).

2.4. Drug release studies

In vitro drug release from uncured and cured tablets was evaluated using the USP apparatus 2 dissolution system (AT7 Smart Off-line, Sotax, Allschwiwz, Switzerland), in 0.05 M phosphate buffer at pH 6.8 (1000 ml dissolution medium, 37 °C, paddle rotational speed = 50 rpm, 3 tablets analyzed per condition). At appropriate time intervals, 10 ml samples were withdrawn over a 16 h period and analyzed by UV spectrophotometry at 230 nm (Agilent 8453 UV-visible spectroscopy system, Hewlett Packard, Waldbronn, Germany).

To assess long-term storage stability, cured tablets were stored in closed polypropylene vials at 25 °C and 60% RH (ambient conditions) and at 40 °C and 75% RH (stress conditions). Drug release was measured before and after 6 months storage for stress conditions and after 18 months storage for ambient conditions.

Comparison between drug release profiles was carried out using the similarity factor (f_2) [25], given by the following equation:

$$f_2 = 50 \cdot \log \left(\left[1 + (1/n) \sum_{t=1}^n (R_t - T_t)^2 \right]^{-0.5} \times 100 \right) \quad (1)$$

where n is the number of time points, R_t denotes the percentage of drug released of the reference condition at time t , and T_t is the percentage of drug released from the test condition at time t .

Dissolution profiles were considered similar for f_2 values between 50 and 100 [26].

2.5. X-ray micro-computed tomography ($X\mu$ CT) measurements

High-resolution X-ray micro-tomography acquisitions [27] were performed on beamline ID19 at the European Synchrotron Radiation Facility (Grenoble, France). Each data collection consisted in a set of 1200 radiographies acquired in 0.15 degree increment over 180 degrees of rotation of the sample specimen. The detector resolution was 0.28 μ m, leading to a reconstruction with an isotropic voxel size of 0.28 μ m. The 17.6 keV incident X-ray parallel beam was truncated to the dimensions of 600 μ m (wide) \times 300 μ m (high). A Fast Readout Low Noise – FreLoN – 14 bit CCD camera [28] was set at 12 mm from the sample to increase the propagation distance and, therefore, to enhance the

phase contrast. 3D images were reconstructed using a standard filtered back projection algorithm. No further image treatment was necessary. The reconstructed volume was a cylinder of 600 μm in diameter and 300 μm in height. Measurements were carried out in a non-destructive and non-invasive way, that is, from intact tablets without any sample preparation. However, due to the limited size of the X-ray beam, only part of the coating was analyzed.

Both uncured and cured tablets were analyzed (2 tablets per sample). Quantitative analysis of coating porosity, expressed as the ratio (Volume of pores)/(Total Volume), was performed from a basic segmentation of the reconstructed volume using threshold adjustment and manual refinement. A representative coating volume of $2048 \times 2048 \times 200$ (pixels³) was selected from all samples and then analyzed with VGStudio MAX 2.1 software. The 200 pixels corresponded to a thickness of 56 μm chosen in the middle of the coating in order to avoid artifacts from the surface or from the interface zone with the tablet core.

2.6. Spectroscopic measurements

2.6.1. Near infrared measurements

NIR spectral acquisitions were performed with a BUCHI NIRFlex N500 spectrometer (BUCHI Labortechnik AG, Flawil, Switzerland) in the diffuse reflectance mode. The illumination spot diameter was 4 mm. Spectra were acquired over the wavelength range of 12,200–4000 cm^{-1} , at a spectral resolution of 16 cm^{-1} and accumulated over 128 scans. NIR acquisitions were carried out on the top of one tablet face, at approximately the same location for all samples. Tablets were positioned on a sample plate allowing accurate and repeatable NIR acquisitions. Fifteen tablets, including uncured tablets, were analyzed for each dynamic curing time. Spectral acquisition was performed using BUCHI NIRCal[®] 5.2.3000 software (BUCHI, Flawil, Switzerland).

2.6.2. Raman measurements

Raman spectral acquisitions were performed with a HR Labram microspectrometer (Horiba Jobin Yvon, Lille, France). The excitation source was a 785 nm single-mode diode laser generating approximately 50 mW on the sample. The microspectrometer was equipped with an Olympus microscope, and measurements were recorded using a 50 \times MPlan objective (spot size laser \sim 10 μm). Light scattered by the sample was collected through the same objective. Using a Notch filter, the elastically scattered photons were separated from the Raman signal. A multichannel CCD detector (1024×256 pixels²) detected the Raman Stokes signal dispersed with a spectral resolution of 2 cm^{-1} by a holographic grating (600 grooves/mm). The spectral range was from 500 to 1800 cm^{-1} . All spectra were acquired at a total laser light exposure time of 2×30 s per collected spectrum. Each component of the coating suspension and a sample of dried coating suspension were analyzed to assign Raman bands in film coating spectral information. A total of 9 spectra were then acquired from both uncured and cured tablets (3 tablets per condition and 3 acquisitions per tablet at different positions on the surface). Spectral acquisition was carried out using Labspec5 software (Horiba Jobin Yvon SAS, Lille, France).

2.6.3. Spectral data analysis

All Raman and NIR data were exported from Labspec5 software and BUCHI NIRCal[®] software, respectively, and analyzed using Matlab[®] 7.8, R2009a software (The MathWorks Inc., Natick, USA) and PLS Toolbox 5.8.3 (Eigenvector Research Inc., Wenatchee, USA).

Raman spectra were pre-processed using baseline correction [29] and normalization based on the highest peak intensity [30].

NIR spectra were pre-processed using Savitzky–Golay smoothing (filter width of 9 points and a second-order polynomial fitting) [31] and second derivative. A multivariate qualitative analysis was

performed from pre-processed NIR spectra using Principal Component Analysis (PCA). This method defines the significant sources of variance within the samples as a series of principal components (PC) [32]. The study of PCA scores based on the relative contribution of each spectrum on the PC construction highlights the relationship between samples.

2.7. X-ray microdiffraction measurements

Experiments were performed at the European Synchrotron Radiation Facility (Grenoble, France) on the microfocus beamline ID13 [33]. The high intensity monochromatic beam (energy 12.41 keV), obtained from an in-vacuum undulator and a Si(1 1 1) double-crystal monochromator, was focused with an ellipsoidal mirror (focal spot 20 (wide) \times 40 (high) μm^2) and a size-limited down to a 1.2 μm diameter section by Kirpatrick–Baez optics. A two-pieces guard aperture (Pb, about 10 μm square aperture) was used to reduce the diffuse scattering from the exit of the collimator. Samples were mounted on a motorized gantry coupled with a microscope allowing sample positioning with a resolution close to 0.1 μm . The experiments were carried out in transmission geometry with a sample-detector distance of 164.9 mm, calibrated using silver behenate (first order spacing 58.38 Å). 2D X-ray scattering patterns were recorded from 0.01 to 0.5 \AA^{-1} on a MAR-CCD camera (16 bit readout; 130 mm entrance window; 2048×2048 pixels²; pixel size of $78.94 \times 78.94 \mu\text{m}^2$) with a beam-stop diameter of approximately 200 μm . The data collection procedure consisted of a series of 50 shots collected along a 150- μm long line perpendicular to the coating surface at the level of the tablet band, penetrating progressively into the coating with a 3 μm step size. Time exposure for recording one shot was 1 s. No radiation damage effect was detected in the scattering pattern for this exposure time. The diffraction experiments were performed on 4 tablets prior to curing and following 4 h of dynamic curing (2 tablets per condition). To ensure the representativeness of the results obtained along a single scan, a series of 20 scans from a distance of 50 μm were also collected, leading to a 2D image of the crystalline structure of the coating layer.

To identify the diffraction peaks, additional experiments were carried out on pure crystalline components present in the coating formulation inserted in a 1-mm diameter glass capillaries using a conventional laboratory set-up: Rigaku rotating Cu anode (50 kV, 100 mA), beam energy 8.04 keV, beam size $0.6 \times 0.6 \text{ mm}^2$, distance sample-detector 250 mm, MAR-CCD 2300 image plate detector, resolution $0.15 \times 0.15 \text{ mm}^2$, exposure time 100 s.

3. Results and discussion

3.1. Dynamic curing effect on drug release and storage stability

Fig. 1 represents drug release profiles obtained from tablets collected at the end of the coating operation and at regular time intervals during dynamic curing. Approximately 10% of theoretical polymer blend (w/w expressed as percentage of core tablet mass) was applied to obtain controlled release of the drug studied over a 16-h period. As expected, a significant decrease in drug release was observed between uncured and cured coated tablets regardless of curing time ($38 < f_2 < 45$). This confirmed that film formation was incomplete immediately after coating. However, identical dissolution profiles were obtained irrespective of curing time ($75 < f_2 < 97$). Therefore, even short dynamic curing periods at 60 $^\circ\text{C}$ and ambient relative humidity could be efficient to obtain fully coalesced film coatings.

Storage stability was then conducted to confirm complete film formation after dynamic curing. Recently, long-term stable film

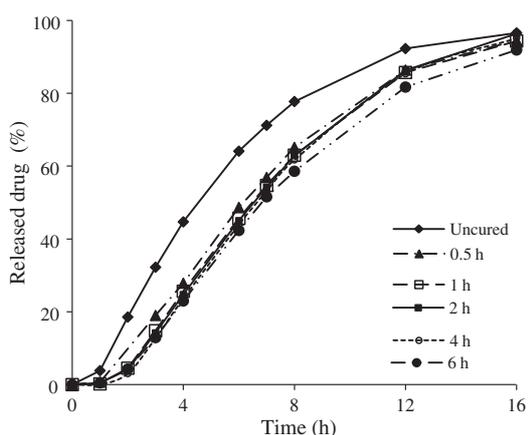


Fig. 1. Effect of curing time on drug release from tablets coated with ethylcellulose:Kollicoat IR[®] 90:10 (10% coating level) and cured inside a pan coater at 60 °C and ambient RH. To not overload the figure, standard deviations were not represented but were included between 2% and 4% of released drug.

coatings prepared from similar coating suspension (15% ethylcellulose:Kollicoat IR[®] 85:15) were obtained after 2 h of dynamic curing in a fluidized bed at 57 °C/15% RH [22]. In the present study, long-term storage stability of tablets coated with 10% ethylcellulose:Kollicoat IR[®] 90:10 and dynamically cured for 2 h at 60 °C and ambient RH was verified. As it can be seen in Fig. 2, drug release rates from cured tablets stored for 6 months under stress conditions and 18 months under ambient conditions were constant ($f_2 = 85$ and 84, respectively). In addition, stable film coatings for 4 h dynamically cured tablets were also obtained (Fig. 2), under both ambient ($f_2 = 77$) and stress conditions ($f_2 = 95$).

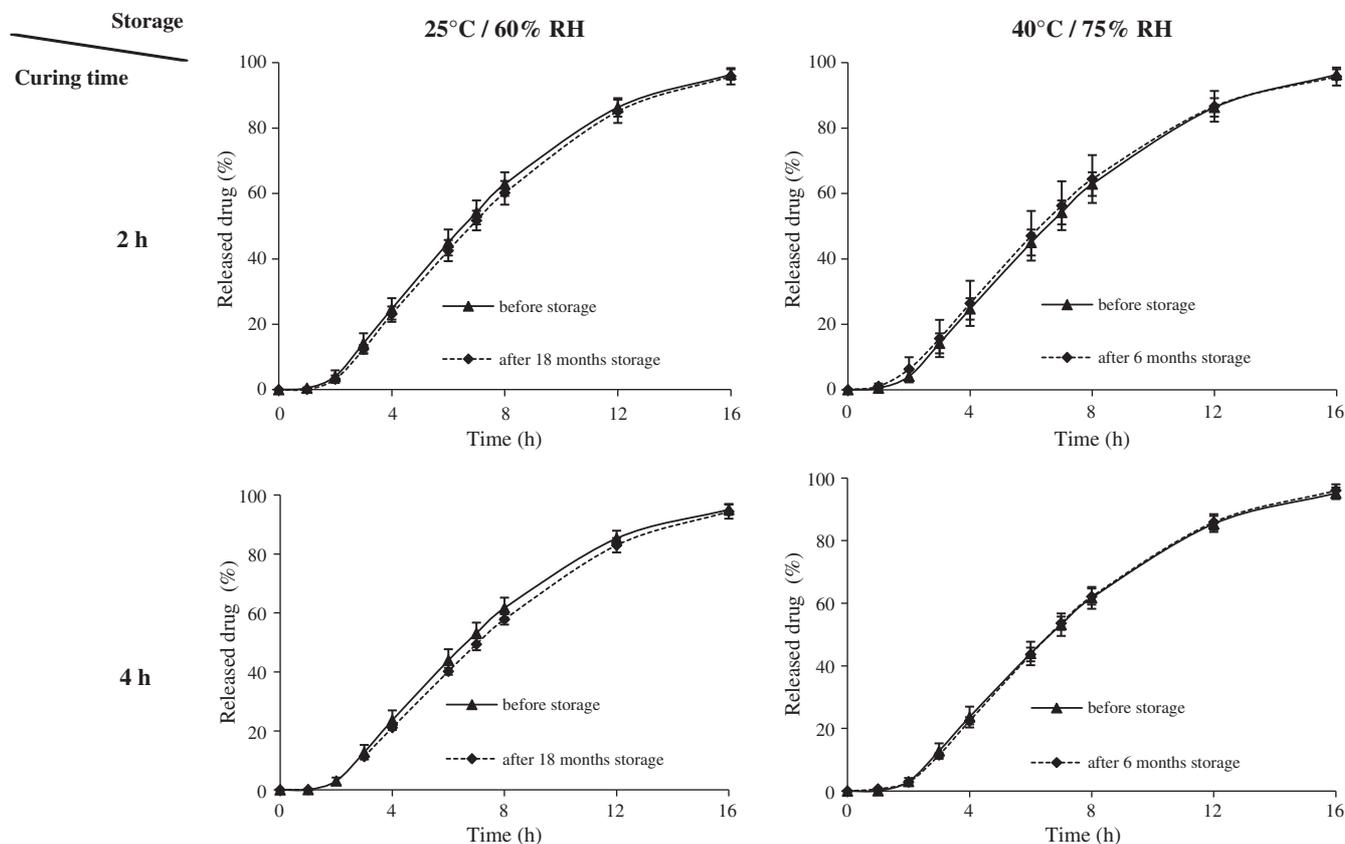


Fig. 2. Storage stability of dynamically cured tablets before and after 18 months under ambient conditions and 6 months under stress conditions (as indicated in the figures). The storage conditions are shown at the top, the curing time on the left.

Drug release profiles were stable even under elevated conditions of temperature and humidity. The high stability of the film coating and the efficiency of the dynamic curing operation were, therefore, confirmed.

Dissolution testing is currently the most widely used method to evaluate curing performances directly from coated solid dosage forms. However, it remains a destructive and time-consuming operation giving access to overall information on both the substrate and the coating layer. For a better understanding of film formation, new techniques were investigated to overcome dissolution drawbacks and provide a further insight into film coating structure irrespective of tablet core.

3.2. Non-destructive determination of internal coating structure using X μ CT

X-ray micro-computed tomography was used to provide information about the internal coating porosity of uncured and cured tablets. Currently, this information can be assessed using conventional but destructive microscopic techniques associated with image analysis. Whilst mercury porosimetry can determine the presence of pores within a solid dosage form directly, it cannot detect closed pores. In contrast, both open and closed pore populations can be obtained by X μ CT measurements [10].

Typical reconstructed X μ CT cross-sectional images are shown in Fig. 3. These 2D images are representative of all recorded acquisitions for the two tablets analyzed per condition and of all cross-sectional images recorded for the 3D imaging reconstruction (1200 X-ray radiographies). The use of monochromatic synchrotron X-ray radiation provided a relevant contrast between the tablet core and the coating layer thanks to the phase contrast mode, thus allowing an accurate insight into the coating layer.

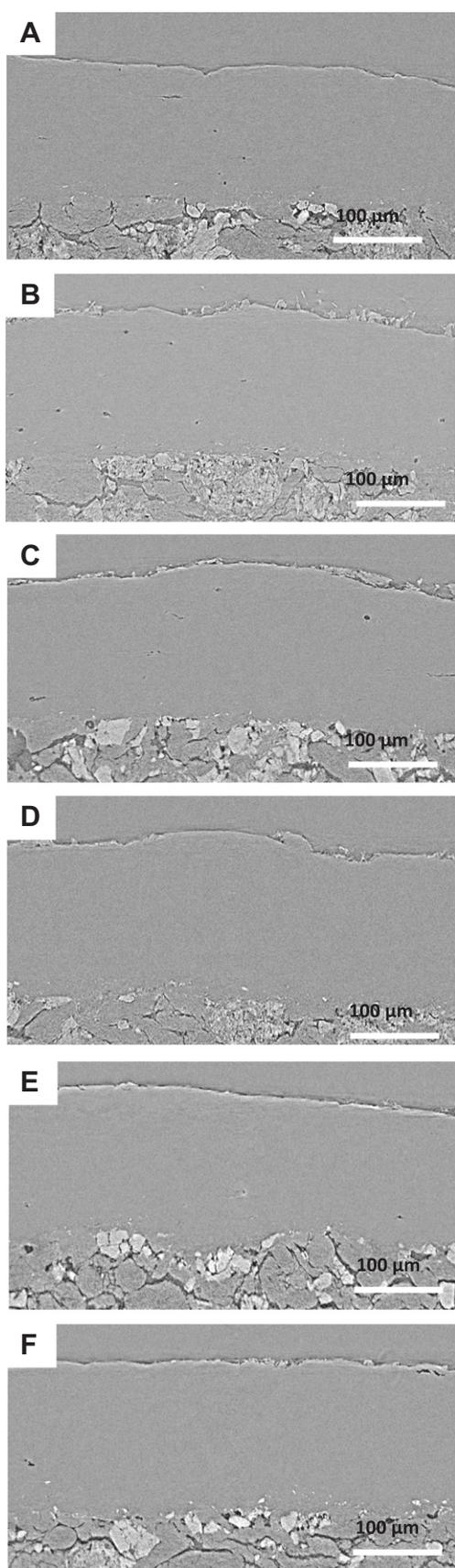


Fig. 3. Representative micro-tomographic images obtained from coated tablets (A) prior to curing and following various dynamic curing times: (B) 0.5 h, (C) 1 h, (D) 2 h, (E) 4 h, and (F) 6 h.

The non-destructive X μ CT analysis highlighted the presence of micropores that gradually decreased as curing time continued. Uncured tablets and tablets cured for as long as 2 h (Fig. 3A–D) were characterized by the presence of numerous micropores within the coating layer and by a coating surface with a high roughness. A qualitative analysis revealed the existence of two types of micropores presenting either (i) a spherical shape with a characteristic diameter of around 5 μ m or (ii) a broader and flatter shape. Quantitative volume analysis demonstrated that uncured and 0.5 h dynamically cured tablets presented similar porosities (\sim 0.20% v/v), while a 2-h curing period led to lower porosity (\sim 0.14% v/v). In contrast, tablets cured for longer times (Fig. 3E and F) presented coating layers with a much lower porosities (\sim 0.03% v/v) and smoother surfaces.

These results suggested that 4 h of dynamic curing could lead to a more reduced coating permeability than the coating permeability obtained after 2 h of curing. A slower drug release could thus be expected. However, the observed variations in coating porosity were not correlated with dissolution profiles, which exhibited similar release patterns from all cured tablets. Therefore, X μ CT allowed the detection of more subtle variations within film coating during dynamic curing in comparison with dissolution. This X-ray technique can successfully be carried out to provide a deeper insight into film coating structures, independently of substrate properties. Interestingly, more accurate determination of curing end-point could be assessed by combining results from X μ CT coating porosity and long-term drug release.

3.3. Monitoring of dynamic curing by near infrared and Raman spectroscopies

3.3.1. Near infrared analysis

NIR spectra were recorded from uncured and cured tablets. Pre-processed NIR spectra are represented in Fig. 4. Spectral evolution was observed with respect to curing times. Variations in absorbance were observed in two characteristic NIR regions over the ranges 5060–5380 cm^{-1} (region A, Fig. 4) and 6970–7190 cm^{-1} (region B, Fig. 4). Both NIR regions identified are characteristics of combination and overtone bands of water due to stretching and deformation vibrations of the OH groups [34]. Observed

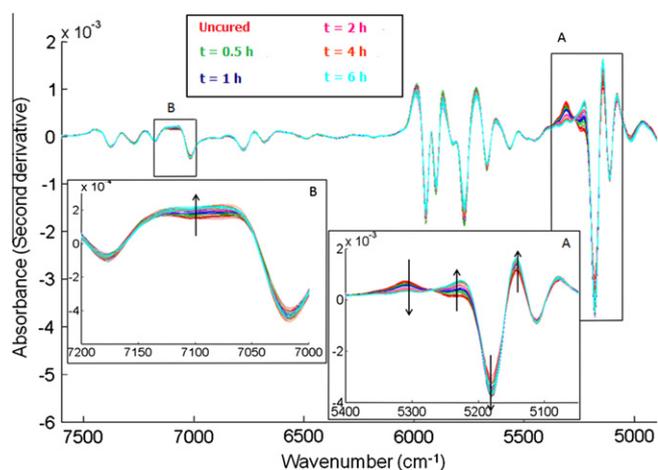


Fig. 4. Pre-processed NIR spectra of coated tablets prior to curing and undergoing dynamic curing until 6 h (as indicated in the figure). Enlarged areas A and B correspond to NIR spectral regions of interest where absorbance varies in a progressive way according to curing time. Arrows indicate the increase in curing time. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

changes could, therefore, be related to the presence of water in film coating. No other NIR spectral variations were detected over the entire spectral range that could be linked to curing times.

PCA was performed to highlight the relationship between samples exhibiting spectral variations observed within both of the selected NIR regions. As seen in Fig. 5, an obvious discrimination between samples was obtained on PC2, related to relevant information from uncured and cured tablets. Uncured tablets were clearly separated from cured tablets. Spectra corresponding to curing times ranging from 0.5 to 4 h were also separated in distinct groups, while spectra for longer curing times were clustered in a same area. This result confirmed that sample classification based on curing times could be correlated with the removal of water trapped within the coating layer during coating. In addition, this NIR spectral analysis demonstrated the efficiency of a 4-h dynamic curing period to achieve constant spectral information.

Nevertheless, NIR results obtained both from spectral observations and PCA were not directly related to dissolution results that did not appear significantly different from all cured tablets. Similarly to $\chi\mu$ CT, NIR spectroscopy allowed the detection of finer variations in coating structure than dissolution. This vibrational technique could, therefore, be used in association with dissolution testing to determine optimal curing end-point in order to strengthen the control strategies of final product quality.

Furthermore, previous studies also reported NIR changes during the curing of acrylic coatings. However, these changes, observed for a different coating formulation, were either attributed to the addition of an acidic polymer (Eudragit L[®]), resulting in ionic interactions between functional groups of Eudragit[®] RS:RL 30D [18], or caused by the N-R swinging of quaternary ammonium groups promoted by high humidity and temperature, required for acrylic coatings, during curing [19]. In the present study, NIR spectroscopy was efficiently used to monitor dynamic curing operation from tablets coated with an ethylcellulose based-coating suspension. Its ability to highlight changes in water content between uncured and cured tablets was demonstrated. It can be noticed that curing was performed at ambient relative humidity. Nevertheless, in some situations, high humidity during curing may be required to obtain long-term stable film coatings [22]. Therefore, in such a case, the use of NIR spectroscopy would have to be evaluated independently and it may not be suitable to monitor changes in water content as the high humidity would dominate the spectrum.

NIR spectroscopy is a well-known technique to monitor water content and provides detection limit as low as 0.1% (w/w) [35]. Nonetheless, this high sensitivity to moisture could mask other more subtle changes in the polymer formulation that might take place in parallel. Raman spectroscopy was, therefore, used as a

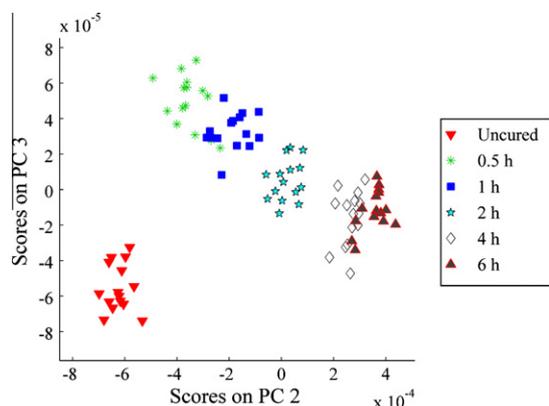


Fig. 5. PCA scores plot obtained from pre-processed NIR spectra of uncured tablets and tablets dynamically cured until 6 h. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

complementary vibrational technique to characterize the polymer structure prior to and following dynamic curing.

3.3.2. Raman analysis

Raman spectra of each excipient in the coating formulation, that is, cetyl alcohol (Fig. 6a), sodium lauryl sulfate (Fig. 6b), graft copolymer PVA-PEG (Fig. 6c), triethylcitrate (Fig. 6d), and ethylcellulose (Fig. 6e), were initially acquired. Dried coating suspension was then analyzed to identify Raman bands from each coating component (Fig. 6f).

Spectral analysis revealed ethylcellulose to be the main component in film coating; nonetheless, specific Raman bands arising from the presence of other components were also observed, as indicated in Fig. 6f. The peak at 870 cm^{-1} was assigned to $\nu(\text{C}-\text{O}-\text{C})$ stretching vibration from triethylcitrate or graft copolymer PVA-PEG [34,36]. Peaks at 1063 cm^{-1} , 1080 cm^{-1} , 1100 cm^{-1} , and 1297 cm^{-1} were attributed to $\nu(\text{C}-\text{C})$ stretching vibration from cetyl alcohol and sodium lauryl sulfate [36–39], while the peak at 1738 cm^{-1} was attributed to $\nu(\text{C}=\text{O})$ stretching vibration from triethylcitrate [36].

Raman spectra of uncured and cured tablets after various dynamic curing times were acquired and compared. The intensity of the peak at 1080 cm^{-1} decreased as curing proceeded, to disappear entirely after 4 h. This can be seen quite clearly in Fig. 7 representing the peak area, calculated between 1075 and 1085 cm^{-1} , as a function of curing time. This specific Raman band at 1080 cm^{-1} was assigned to the $\nu(\text{C}-\text{C})$ stretching mode in the *gauche* conformation [37] and may be attributed to either cetyl alcohol or sodium lauryl sulfate. As expected, it can be concluded that no chemical reaction occurred during curing as no new chemical entity was identified. Curing led only to a structural rearrangement showing a decrease in the *gauche* conformers along the alkyl chains, resulting in denser and more organized coating structure with lower permeability for water and drug. Consequently, a

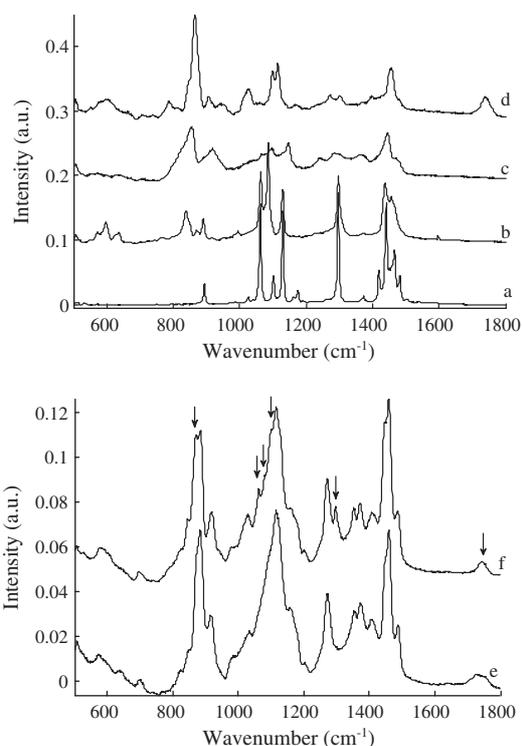


Fig. 6. Raman spectra of (a) cetyl alcohol, (b) sodium lauryl sulfate, (c) graft copolymer PVA-PEG, (d) triethylcitrate, (e) ethylcellulose, and (f) dried powdered sample of film coating. Arrows indicate Raman bands related to the presence of other components in coating suspension in addition to ethylcellulose.

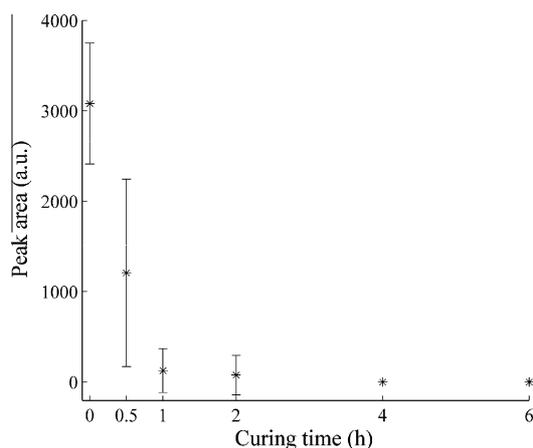


Fig. 7. Evolution of peak area, calculated between 1075 and 1085 cm^{-1} from Raman spectra as a function of dynamic curing time, ranging from 0 (uncured) to 6 h. Mean area values obtained from 9 spectra, associated with its standard deviation, are represented per curing time.

slower drug release rate could be expected for long curing times. However, dissolution results did not show significant changes in drug release with increased curing times. Raman results, therefore, revealed more subtle variations in film coating structure during dynamic curing and indicated a stable coating structure after a 4-h curing period.

Interestingly, real-time and in-line spectroscopic measurements could be successfully performed inside a coating apparatus to monitor dynamic curing, in accordance with current pharmaceutical development regulations and Process Analytical Technology control strategies [23,40]. NIR and Raman techniques are commonly employed to control and monitor various real-time pharmaceutical processes [17]. To our knowledge, no study up until now has reported in-line monitoring of dynamic curing with the selected coating formulation. Therefore, these results highlighted the potential of both NIR and Raman spectroscopies for the control and the monitoring of this pharmaceutical unit operation.

In addition, optical microscope images were recorded of the surface of uncured tablets (Fig. 8), showing some coating heterogeneity. Specific Raman bands from darker areas, as observed in Fig. 8b, were identified and attributed to higher concentrations of cetyl alcohol (Fig. 8a), while these bands were attenuated for spectra acquired from lighter areas (Fig. 8c). As curing proceeded, a quantitative decrease in darker areas was observed. The melting of cetyl alcohol (melting point between 45 and $52\text{ }^{\circ}\text{C}$ [41]) is likely to occur during curing, thus leading to its partial diffusion at the surface or inside the coating layer. This would favor the improvement of coating homogeneity with curing.

Recently, Yang described similar coating heterogeneity from Aquacoat[®]-based free films, prepared by a spraying technique, after a static curing of 1 h at $60\text{ }^{\circ}\text{C}$ [20]. The author suggested the existence of crystalline cetyl alcohol agglomerates to explain non-homogeneous distribution within the coating. On the other hand, Schmid et al. [42] using X-ray investigations observed cetyl alcohol in Aquacoat[®] to be in the amorphous state.

Based on these studies, X-ray microdiffraction measurements were conducted to elucidate the non-homogeneous distribution of cetyl alcohol and to identify its potential crystalline evolution during the curing operation.

3.4. Crystalline composition of coating layer using X-ray microdiffraction

In this study, X-ray diffraction using synchrotron radiation was performed at the microscale to identify the crystalline composition

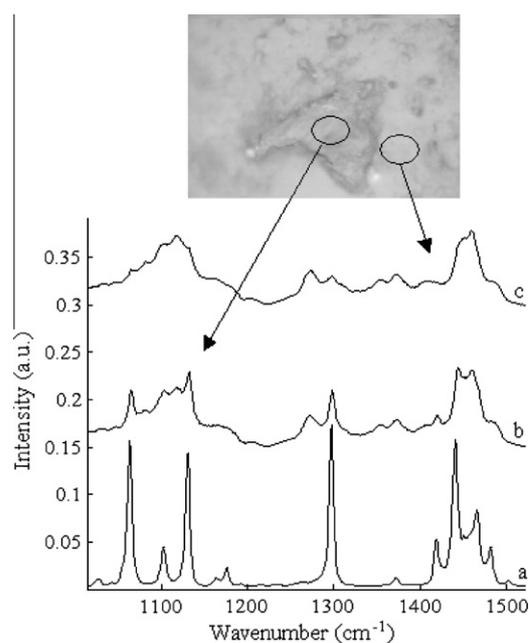


Fig. 8. Raman spectra of (a) cetyl alcohol, (b) darker areas, and (c) lighter areas corresponding to distinct regions observed on optical microscopic images from uncured tablets (size: 614×460 pixels²).

of the coating layer and to gain a better understanding of coating heterogeneity.

3.4.1. Identification and distribution of crystalline components within the film coating layer

Fig. 9 illustrates representative diffraction patterns obtained from uncured tablets. The series of patterns collected during a scan perpendicular to the coating clearly showed a progressive change of scattering features from the coating surface up to the interface with the tablet core. In fact, each pattern consisted of the sum of the scattering contributions from all the matter located along the beam path. Therefore, close to the coating surface, the signal only arose from the surface. However, due to the tablet curvature, the contribution of the internal coating structure became more and more important when going deeper into the coating. The depth resolution was thus not perfect. In this work, measurements were carried out in a non-destructive way without microtoming the tablets. Such data collection could have complicated considerably the data analysis; nevertheless, the changes of the scattering features were well marked, leading to the straightforward interpretation

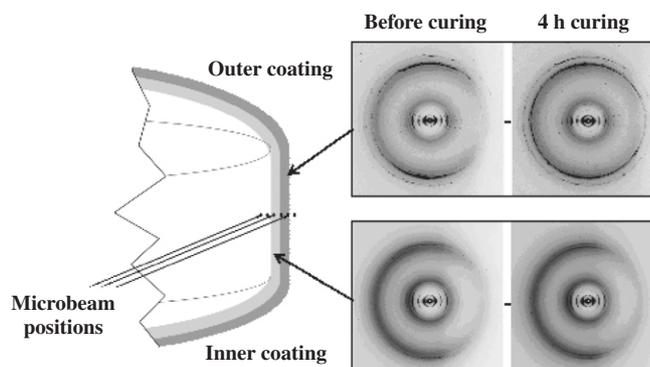


Fig. 9. X-ray microdiffraction patterns corresponding to inner and outer coating zones, as indicated on the schematic representation of uncured and 4 h dynamically cured coated tablets.

of the pattern evolution. The patterns exhibited mainly broad diffuse rings, corresponding to amorphous components, plus a series of sharp diffraction arcs/rings corresponding to crystallized components. The progressive change of these sharp diffraction features can be interpreted by the existence of two zones: an outer zone close to the tablet surface and an inner zone in contact with the tablet core (Fig. 9), which clearly indicated a difference in coating compositions.

The comparison between the outer and inner intensity profiles for uncured samples, obtained by circular integration, is given in Fig. 10. The intensity profiles are also shown of the three main crystallized or partly crystallized components in the initial coating composition: ethylcellulose, sodium lauryl sulfate, and cetyl alcohol.

It can be noticed that the relative intensities of the peaks in the coating were different from those of the pure components, firstly because of the non-uniform spatial orientation of the crystallites in the coating, and secondly and most importantly because of different concentrations of the components. Nevertheless, the positions of most diffraction peaks of the coating corresponded to the peaks of these three components. In addition, the slight shift in position that was observed for a few peaks and the presence of a few extra peaks revealed the existence of a structural change during film coating formation, such as polymorphism and/or phase mixing.

Furthermore, main diffraction peaks in the outer zone of the coating corresponded to a lamellar thickness packing of 45 Å, and the two most intense WAXS peaks at 4.15 Å and 3.68 Å were typical of a rectangular organization in the plane of the layers. The comparison with the patterns of the pure components demonstrated that these diffraction features were characteristic of cetyl alcohol. The inner zone also showed diffraction peaks from a lamellar system with a layer thickness of 41.5 Å and two main WAXS peaks at 4.32 Å and 4.10 Å. They corresponded to the features of sodium lauryl sulfate. With regards to ethylcellulose, which only displayed a sharp but weak peak (about 7.8 Å), it was evenly distributed though the coating except in the 10- μm thick outer surface where it was missing.

An opposite behavior of sodium lauryl sulfate and cetyl alcohol emerged from these observations. Crystalline cetyl alcohol was mainly found in the outer part of the coating and its concentration decreased progressively when going toward the deeper parts of the coating. On the contrary, the concentration of sodium lauryl sulfate was low close to the surface of the coating and increased in the deeper parts. Therefore, the existence of two zones was clearly highlighted and could be inferred: an outer zone with high cetyl

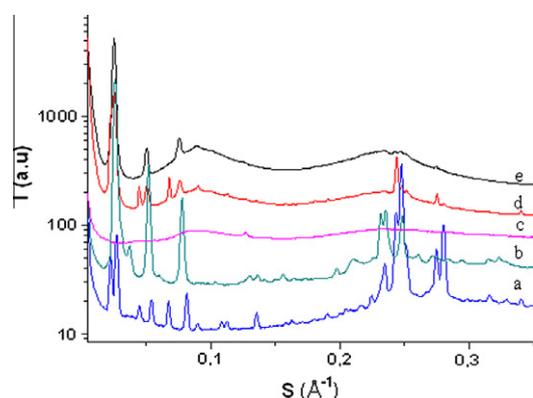


Fig. 10. X-ray microdiffraction patterns of (a) cetyl alcohol, (b) sodium lauryl sulfate, (c) ethylcellulose, (d) outer, and (e) inner coating layers obtained from uncured coated tablets. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

alcohol content, low sodium lauryl sulfate content, and no ethylcellulose on the extreme surface, and an inner ethylcellulose zone, rich in sodium lauryl sulfate with a low cetyl alcohol content.

In agreement with Yang's suggestions [20], X-microdiffraction results confirmed a non-homogenous distribution of crystalline cetyl alcohol observed by microscopic imaging and Raman measurements. In addition, an unexpected distribution of crystalline sodium lauryl sulfate within the coating layer was also highlighted. The homogeneity of coating suspension was checked prior to and during coating. Therefore, the main reason may be related to an upward migration of cetyl alcohol associated with a downward migration of sodium lauryl sulfate in coated tablets during coating.

Taking into account the observations of Schmid et al. [42] who reported that cetyl alcohol and sodium lauryl sulfate were molecularly dispersed in Aquacoat[®] and thus in the amorphous state, it could be assumed that these two components crystallized during coating.

3.4.2. Influence of curing on crystalline coating structure

Analysis of 4-h dynamically cured samples did not reveal any significant changes in the scattering intensity profiles, except for the thickness of the lamellar system attributed to cetyl alcohol layers which decreased by 1 Å. The two-zone coating was still present with the same characteristics. This result confirmed that melted cetyl alcohol present at the coating surface was spread over the coating surface during curing and did not diffuse deeper within the coating layer, as initially suggested by microscopic and Raman observations.

As previously mentioned, conducting diffraction experiments without sectioning tablets limits precise determination of concentration profiles due to the superposition of diffraction contributions along the X-ray beam path. However, one can roughly estimate that the outer and inner zone thicknesses are, respectively, 30 μm and 70 μm . These values were fairly constant along the 1-mm long zone which was analyzed in 2D. Nevertheless, the overall coating thickness decreased by about 10 μm during curing, indicating a denser coating layer.

In addition to this result, differences in the orientation of cetyl alcohol crystallites in the outer coating were observed in cured tablets. Without curing the equatorial position of the SAXS arcs (perpendicular to the coating plane), combined with the meridional position of the WAXS arcs clearly indicated that the cetyl alcohol layers lie mainly parallel to the coating surface (Fig. 9). After 4-h dynamic curing, the arcs tend to transform into circles, indicating a random orientation of the layers. This observation can be interpreted as a melting/recrystallization phenomenon induced by the heat-cooling cycle of cured tablets.

A new insight into the internal coating structure was provided prior to and following curing, revealing the crystalline distribution of coating components without requiring destruction of samples. Interestingly, results highlighted the existence of two coating zones presenting different concentrations in crystalline cetyl alcohol and sodium lauryl sulfate, irrespective of curing.

4. Conclusions

Deeper insights into film coating structures from dynamically cured tablets were highlighted using non-destructive techniques. X μ CT, NIR, and Raman analyses revealed more subtle variations occurring in coating layers during curing compared with usual dissolution testing. It was demonstrated that stable film coating structures could be obtained after a 4-h dynamic curing period. Interestingly, and even prior to curing, unexpected distributions of two crystalline components, cetyl alcohol and sodium lauryl sulfate, within coating layers were observed from X-ray microdiffraction.

tion measurements. Further, quantitative investigations would be required to fully elucidate these observations and determine their impact on the properties of film coating structures. Nonetheless, the aforementioned techniques can efficiently be used to provide a better understanding of curing process, leading to relevant determinations of curing end-points.

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