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Molecular dynamics and physical stability of amorphous nimesulide drug and its binary drug-polymer systems

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ABSTRACT

In this paper we study the effectiveness of three well known polymers: inulin, Soluplus and PVP in stabilizing amorphous form of nimesulide (NMS) drug. The re-crystallization tendency of pure drug as well as measured drug-polymer systems were examined at isothermal conditions by using broadband dielectric spectroscopy (BDS), and at non-isothermal conditions by differential scanning calorimetry (DSC). Our investigation has shown that the crystallization half-life time of pure NMS at 328 K is equal to 33 minutes. We found that this time can be prolonged to 40 years after adding to NMS 20% of PVP polymer. This polymer proved to be the best NMS's stabilizer, while the worst stabilization effect was found after adding the inulin to NMS. Additionally, our DSC, BDS and FTIR studies indicate that for suppression of NMS's re-crystallization in NMS-PVP system, the two mechanisms are responsible: the polymeric steric hindrances as well as the antiplastization effect exerted by the excipient.

KEY WORDS

amorphous drug, physical stability, molecular dynamics, nimesulide, drug-polymer system, PVP, Soluplus, inulin

INTRODUCTION

Nimesulid (NMS) is a non-steroidal anti-inflammatory drug (NSAIDs) which shows high anti-inflammatory, antipyretic, and analgesic activities in addition to low toxicity, and a high therapeutic index¹. Like many non-steroidal anti-inflammatory drugs, NMS is characterized by low aqueous solubility (≈ 0.01 mg/mL), and poor dissolution rate². The commonly employed strategies to increase solubility and dissolution rate of poorly soluble drugs are based on using an cyclodextrin, macronization, and conversion the Active Pharmaceutical Ingredients (APIs) to their salt forms³. Another, an alternative way is preparing the amorphous form of those compounds. Recently, it has been often reported that pharmaceuticals prepared in the amorphous form are significantly more soluble, have higher dissolution rate, and have greater bioavailability than their crystalline counterparts^{4,5,6}. Unfortunately, it should be highlighted that, amorphous APIs are in general physically unstable and, consequently, during storage or manufacturing they may revert to the crystalline form and, consequently, lose their superior properties.^{7,8,9,10}

It is believed that the main factor affecting physical stability of disordered APIs is molecular mobility^{11,12}. It has been demonstrated many times that the most efficient experimental tool to investigate molecular dynamics of glass-forming systems is broadband dielectric spectroscopy (BDS)^{13,14}. This technique enables the determination of the characteristic time of molecular reorientation over a broad range of frequencies (over 12 decades) and temperatures. Such a wide f and T range allows the detection of various relaxation processes. In the supercooled liquid, the structural (α) relaxation, which originates from cooperative movements of many molecules, is usually the most prominent relaxation process observed in the dielectric spectra. In the glassy state it is possible to observe faster local motions, which have an inter- or intramolecular origin. Additionally, using dielectric spectroscopy, the kinetics of crystallization at isothermal conditions can be monitored. Such studies are extremely valuable, especially for physical stability tests.

The objective of the current work was to convert nimesulid drug to its amorphous form and examine by means of BDS its physical stability. Since, our measurements indicate that amorphous NMS is extremely physically unstable system, a lot of attention has been paid to find an efficient method of stabilization this material. A promising strategy for inhibiting devitrification of amorphous drugs is to disperse them into a polymeric matrix.^{15,16,17} Therefore we decided to check how three well known and commonly used in pharmaceutical industry polymers: inulin (INL), Soluplus (SOP) and poly(vinylpyrrolidone) (PVP) stabilize nimesulid drug. It is worth noting that each of these additives belong to different groups of

polymers. SOP represents a very complex copolymeric system, whereas both INL and PVP have strictly one polymerizable unit. In contrast to inulin, which has a stiff molecular chain, poly(vinylpyrrolidone) belongs to a group of polymers with highly flexible structures. In order to achieve our aim and answer the question of which polymer can better suppress NMS's crystallization, we investigated the molecular dynamics and kinetics of crystallization of pure NMS and the following binary mixtures: NMS+PVP, NMS+SOP and NMS+INL. The crystallization process in nimesulide has been examined at both isothermal and non-isothermal conditions. The latter was measured by using differential scanning calorimetry (DSC). Finally, we also tried to establish which stabilization mechanism is involved in case of the investigated binary mixtures.

EXPERIMENTAL METHODS

Materials

Nimesulide (NMS) (Kemprotec Limited, UK), inulin DP-23 (INL) (Frutafix TEX, Sensus, Netherlands), polyvinyl pyrrolidone 10 kDa (PVP) (Sigma Aldrich, Ireland) and SoluplusTM (SOP) (BASF, Germany).

Preparation of physical mixtures

Physical mixtures of NMS and polymers, respectively; INL, PVP or SOP were prepared by milling at room temperature using mixer mill MM 200 (Retsch, Germany) for 15 min at mixing frequency of 25 1/s.

Differential Scanning Calorimetry (DSC)

Thermodynamic properties of NMS, NMS + 5% wt. INL, NMS + 5% wt. SOP, NMS + 5% wt. PVP, NMS + 20% wt. SOP and NMS + 20% wt. PVP were examined using a Mettler–Toledo DSC 1 STAR^e System. The measuring device was equipped with HSS8 ceramic sensor having 120 thermocouples and liquid nitrogen cooling station. The instrument was calibrated for temperature and enthalpy using zinc and indium standards. Crystallization and melting points were determined as the onset of the peak, whereas the glass transition temperature as the midpoint of the heat capacity increment. The samples were measured in an aluminum crucible (40 μ L). All measurements were carried out in range from 270 K to 430/435 K with a variety – 5 K/min, 10 K/min, 20 K/min and 30 K/min – of heating rates.

Broadband Dielectric Spectroscopy (BDS)

Dielectric measurements of NMS, NMS + 5% wt. INL, NMS + 5% wt. SOP, NMS + 5% wt. PVP, NMS + 20% wt. SOP and NMS + 20% wt. PVP were carried out using a Novo-Control GMBH Alpha dielectric spectrometer, in the frequency range from 10^{-1} Hz to 10^6 Hz, at temperatures from 193 K to 328 K with steps of 2.5 K. The temperature was controlled by a Quattro temperature controller with temperature stability better than 0.1 K. Dielectric studies of all samples were performed immediately after fast cooling of the melt in a parallel-plate cell made of stainless steel (diameter 15 mm and a 0.1 mm gap with Teflon spacers). Crystallization kinetics for all samples were carried out at four different temperatures, as follows: pure NMS at 318 K, 323 K, 328 K and 330.5 K; NMS + 5% INL at 318 K, 323 K, 328 K and 333 K; NMS + 5% SOP at 328 K, 330.5 K, 333 K and 338 K; NMS + 5% PVP

at 328 K, 330.5 K, 333 K and 338 K; NMS + 20% SOP at 343 K, 353 K, 358 K and 363 K; NMS + 20% PVP at 360.5 K, 363 K, 365.5 K and 368 K.

Fourier Transform Infrared Spectroscopy (FTIR)

FTIR was performed using a Spectrum One FT-IR Spectrometer (Perkin Elmer, Connecticut, USA) equipped with Spectrum Software version 6.1. A spectral range of 450-4000 cm^{-1} , resolution of 4 cm^{-1} , scan number of 10 and scan speed of 0.2 cm/s were used. KBr disks were produced by direct compression, using a pressure of approximately 10 bar for 1 minute. A sample loading of 1% was used. The spectra were normalized and background corrected.

MOLECULAR DYNAMICS OF PURE NIMESULIDE ABOVE AND BELOW THE GLASS TRANSITION

Dielectric loss spectra of amorphous nimesulide were measured in both the supercooled liquid and glassy state by using BDS technique. In this experiment, the temperature was increased from 193 K to 328 K in steps of 2.5 K. Representative spectra are shown in Figure 1. For better visualization of obtained dielectric results, the data have been divided into two panels (a and b). Panel a of Figure 1 presents spectra collected above the glass transition temperature. Herein, we can see one well-resolved loss peak connected to the α -relaxation process. This relaxation mode moves toward higher frequencies with increasing temperature and at $T > 323$ K its intensity begins to rapidly decrease. Such a drop in the intensity of α -relaxation is caused by cold crystallization of the sample and it reflects the degree of reduction in amorphous fraction.¹⁸ In order to determine the value of β_{KWW} for NMS, and to check whether temperature affects the shape of the structural relaxation process, we constructed so-called masterplot (see inset Figure 1a) by horizontal shifting of spectra (taken at deferent temperatures from 300.65 to 323.15 K) to superimpose on the reference spectrum at 303.15 K. As can be seen in presented masterplot, the shape of the structural relaxation in NMS is not changing with temperature, and the parameter β_{KWW} is the same for all spectra and equal to 0.72. According to the Shamblin observations, narrower α -relaxation peaks correspond to higher physical stability of amorphous APIs.¹⁹ However, as our studies show, the amorphous NMS reveals quite a strong tendency to recrystallize, despite of fact that the α -relaxation peak is quite narrow. Similar behavior was previously observed in the case of ezetimibe, celecoxib and sildenafil.^{20,21,22,23}

Figure 1. Dielectric loss spectra of NMS obtained on heating. The panel (a) presents spectra collected above the glass transition temperature, whereas panel (b) shows spectra below the T_g . In the inset of panel (a) we present the masterplot that was constructed by superimposed dielectric spectra at nine different temperatures above T_g . The dashed black line represents the KWW fit with the $\beta_{KWW} = 0.72$.

Panel b of Figure 1 presents dielectric spectra collected at temperatures below T_g . In this temperature region the structural relaxation becomes too slow to be observed and only secondary relaxation processes may be visible. As can be easily seen in Figure 1b, NMS has only one secondary (β) process, which, just like the α -mode, moves toward higher frequencies with increasing temperature. However, this process is less sensitive to temperature changes when compared to the behavior of the α -relaxation.

From analysis of dielectric loss spectra collected above and below T_g , we determined the temperature dependence of α - and β -relaxation times (see Figure 2). Firstly, to obtain the values of τ_α and τ_β at various temperatures, the asymmetric α -peak and symmetric β -process have been fitted using the Havriliak–Negami (HN) and the Cole–Cole (CC) function, respectively.²⁴ The empirical HN approach is given by the formula:

$$\varepsilon_{HN}^*(\omega) = \varepsilon'(\omega) - i\varepsilon''(\omega) = \varepsilon_\infty + \frac{\Delta\varepsilon}{[1 + (i\omega\tau_{HN})^a]^b} \quad (1)$$

where ε' and ε'' are real and imaginary parts of complex dielectric permittivity, ε_∞ is high frequency limit permittivity, $\Delta\varepsilon$ is dielectric strength, τ_{HN} is the HN relaxation time, and a and b represent symmetric and asymmetric broadening of the relaxation peak. This function is equivalent to the CC function when the parameter b is equal to 1. In this special case Eq. (1) fits NMS's β -relaxation.²⁵ Next, the fitting parameters were used to calculate the τ_α and τ_β by means of the equation:

$$\tau_{\alpha/\beta} = \tau_{HN} \left[\sin\left(\frac{\pi a}{2 + 2b}\right) \right]^{-1/a} \left[\sin\left(\frac{\pi ab}{2 + 2b}\right) \right]^{1/a} \quad (2)$$

Figure 2. The relaxation map of NMS drug. Temperature dependence of τ_α in the supercooled liquid has been described by the VFT equation. The α -relaxation times in the glassy state were found by horizontal shift of the structural relaxation peak from the region above the glass transition to the region below T_g . The temperature dependence of τ_β was fitted using the Arrhenius equation.

In order to parameterize the $\tau_\alpha(T)$ dependence we employed the Vogel–Fulcher–Tammann (VFT) equation:

$$\tau_\alpha(T) = \tau_\infty \exp\left(\frac{B}{T - T_0}\right) \quad (3)$$

where τ_∞ , T_0 , and B are fitting parameters.^{26,27,28} Their values for NMS are equal to: $\log_{10}(\tau_\infty/\text{s}) = -19.4 \pm 0.7$, $T_0 = 216 \pm 4$, and $B = 3711 \pm 310$. Based on the VFT parameters we estimated T_g (defined as $\tau_\alpha = 100$ s) of NMS as 291 K. The glass transition temperature determined by this method is in good agreement with the T_g obtained from calorimetric studies ($T_{gDSC\ 10K/min} = 294$ K – see Figure 5) as well as with the literature data.²⁹ From VFT fits we also calculated the value of the fragility parameter for NMS as $m_p = 82$. This parameter, also called steepness index, is a measure of deviation from Arrhenius behavior of the $\tau_\alpha(T)$ dependence and is defined as follows:³⁰

$$m_p = \left. \frac{d \log \tau_\alpha}{d(T_g/T)} \right|_{T=T_g} \quad (4)$$

Typical values of steepness index are between 16 and 200. The higher the value of m_p , the more fragile (less strong) liquids are. The fragility is frequently considered as a key parameter for researchers dealing with physical stability of amorphous drugs because it has been implied that strong materials are more stable than fragile ones.³¹ However, very recently a few exceptions to this correlation were found. In the case of NMS we found that its low physical stability correlates to a high value of fragility. Thus NMS satisfies the above mentioned correlation.

ISOTHERMAL AND NON-ISOTHERMAL CRYSTALLIZATION KINETICS STUDIES OF PURE NMS ABOVE THE GLASS TRANSITION TEMPERATURE

In this section we shall concentrate on the crystallization kinetics of NMS drug. To fully characterize the tendency of amorphous NMS to devitrify, both the isothermal and non-isothermal kinetics of crystallization have been investigated.

The isothermal cold crystallization kinetics of NMS were studied by using BDS technique. In this experiment, we performed time-depending dielectric measurements at four different temperatures (318 K, 323 K, 328 K and 330.5 K). For each measurement a new sample was prepared by employing the vitrification method. Using dielectric spectroscopy the crystallization process can be followed by measuring both the real (ϵ') and imaginary (ϵ'') parts of the complex dielectric permittivity. In the ϵ' , devitrification is reflected by a decrease

in the static permittivity (Figure 3a), whereas in the ϵ'' by a decrease in the intensity of the α -loss peak (Figure 3b).

Figure 3. Dielectric spectra of the real (a) and imaginary (b) parts of the complex dielectric permittivity during an isothermal cold crystallization of NMS at fixed temperature equal to 238 K.

Usually, the progress of cold crystallization is analyzed by normalized dielectric constant (ϵ'_N), defined as follows:

$$\epsilon'_N(t) = \frac{\epsilon'(0) - \epsilon'(t)}{\epsilon'(0) - \epsilon'(\infty)} \quad (5)$$

where $\epsilon'(0)$ is the dielectric constant at the beginning of the crystallization process, $\epsilon'(t)$ is the value of dielectric constant at time t , and $\epsilon'(\infty)$ is the long-time limiting value. The normalized kinetics curves of NMS are presented together in Figure 4.

Figure 4. Normalized dielectric constant ϵ'_N as a function of time from crystallization processes occurring at: 318.15, 323.15, 328.15 and 330.65 K. The lower inset presents the Avrami plots in terms of eq. 6 for each crystallization temperature in the time ranges where the dependences are linear. The upper inset shows the temperature dependence of $\log k$ – here, the solid line denotes the linear fit of eq. 7.

To analyze the time dependence of ε'_N we used the Avrami model.³² It should be mentioned that this model is the most commonly applied method of analyzing the crystallization process at a fixed temperature, and is described by the following formula:

$$\varepsilon'_N(t) = 1 - \exp(-K(t)^n) \quad (6)$$

where $K=k^n$ is the rate constant, and n is the Avrami parameter that is directly related to the nucleation dimensionality. Usually parameter n falls in the range of 2-3.³³ To determine values of both K and n parameters we prepared a so-called Avrami plot (see lower inset of Figure 4), which is based on the equation:³⁴

$$\log(-\ln(1 - \varepsilon'_N(t))) = \log K(t) + n \log t \quad (7)$$

As can be clearly seen, if the $\log(-\ln(1 - \varepsilon'_N(t)))$ vs. $\log t$ is linear, the n and $\log K$ can be determined as the slope and the intercept of this dependence, respectively. Avrami parameters obtained in this way are shown in Table 1. The value of Avrami exponent n changes in the standard way, i.e. decreases with temperature (from 2.9 to 1.8), indicating that the crystallization process of NMS at isothermal conditions is three- and two-dimensional at high and low temperatures, respectively. Additionally, since the parameter K increases with increasing crystallization temperature, we can conclude that the dominant mechanism of isothermal devitrification in the case of NMS is the diffusion of molecules.³⁵

T [K]	n	logK
318.15	2.86 ± 0.01	- 11.33 ± 0.04
323.15	2.30 ± 0.02	- 8.49 ± 0.07
328.15	2.18 ± 0.03	- 7.36 ± 0.10
330.65	1.89 ± 0.01	- 5.98 ± 0.03

Table 1. Isothermal cold crystallization kinetics parameters of NMS obtained from the Avrami model according to eq. 7.

In order to estimate the activation energy of isothermal cold crystallization (E_{al}) for NMS we plotted the logarithm of k ($k=K^{1/n}$) versus reciprocal temperature (see upper inset of Figure 4). Since the $\log k$ vs. $1000/T$ dependence, in measured temperature range, is linear, the value of E_{al} may be found by fitting Arrhenius law to the experimental points:

$$\log k = \log k_0 - \frac{E_{al}}{RT} \log e \quad (8)$$

where k_0 and E_{al} are fitting parameters, and R is the universal gas constant. It should be pointed out the temperature dependence of crystallization might not be linear if the measured temperature range would be wider. Crystallization activation energy of pure NMS obtained in this way is equal to 126 ± 8 kJ/mol.

The cold crystallization kinetics of NMS was also studied at non-isothermal conditions. For these measurements the DSC technique was employed. During the experiment the sample was heated from 270 K to 430 K at different heating rates $\phi = 5, 10, 20$ and 30 K/min. As can be seen in Figure 5 and Table 2, with increasing heating rate the values of T_g , T_o (temperature of crystallization onset) and T_p (temperature of crystallization peak) shift toward higher temperatures.

Heating Rate [K/min]	T_o [K]	T_p [K]	T_g [K]
5	342	348	293
10	348	354	294
20	352	360	296
30	359	365	298

Table 2. Temperatures of crystallization onsets, peaks and glass transitions determined from DSC thermograms measured at different heating rates.

Figure 5. DSC thermograms of NMS measured at different heating rates (5 K/min, 10 K/min, 20 K/min, 30 K/min). The upper inset shows an enlarged temperature region of the glass transitions. The lower inset presents an Augiss and Bennett plot for the crystallization peak of NMS. The orange line indicates the linear fit to eq. 9.

Based on the parameters collected in Table 2, we determined the activation energy of non-isothermal cold crystallization (E_{aNI}) of NMS by using the Augiss and Bennett approach (see lower inset of Figure 5):³⁶

$$\ln\left(\frac{\phi}{T_p - T_o}\right) = C_{AB} - \frac{E_{aNI}}{RT_p} \quad (9)$$

where C_{AB} and E_{aNI} are fitting parameters and R is the universal gas constant. The activation energy obtained using this method is significantly greater (over 35%) than the crystallization activation energy of NMS at isothermal conditions, and is equal to 210 ± 14 kJ/mol.

Further analysis of non-isothermal cold crystallization kinetics of NMS was carried out based on the temperature dependence of relative degree of crystallinity X .³⁷ The $X(T)$

dependence, for each heating rate, is presented in Figure 6a, and was obtained by using the following equation:

$$X(T) = \frac{\int_{T_0}^T (dH_c/dT)dT}{\int_{T_0}^{T_\infty} (dH_c/dT)dT} \quad (10)$$

where T_0 and T_∞ represent the temperatures at which the crystallization process begins and ends, respectively, while dH_c denotes the enthalpy of crystallization that is released during an infinitesimal temperature range dT .

Figure 6. (a) The temperature dependence of crystallization degree determined from eq. 10 at different heating rates. (b) The time dependence of crystallization degree obtained by transforming the temperature scale into the time scale using eq. 11. The inset presents the Avrami plots, which have been constructed from the non-isothermal crystallization data for NMS.

After determination of the $X(T)$ dependence it was possible to transform the temperature scale into the time scale and consequently obtain $X(t)$ function. For this transformation the equation listed below has been used:

$$t = \frac{T - T_0}{\phi} \quad (11)$$

where t denotes the time of crystallization and ϕ is the heating rate. The plots of the relative degree of crystallinity as a function of time at different heating rates are shown in Figure 6b. $X(t)$ dependence was next analyzed by the Avrami model – inset of Figure 6b. Obtained values of n' and $\log K'$ are shown in Table 3.

ϕ [K/min]	n'	$\log K'$
5	5.86 ± 0.05	-13.20 ± 0.11
10	6.78 ± 0.05	-14.07 ± 0.11
20	6.52 ± 0.08	-11.89 ± 0.14
30	5.57 ± 0.07	-9.04 ± 0.11

Table 3. Non-isothermal cold crystallization kinetics parameters of NMS obtained from the Ozawa model according to eq. 11.

As can be seen, the n' parameter changes from 5.6-6.8, indicating that the non-isothermal cold crystallization of NMS is much more complicated than crystallization at fixed temperature, where parameter n oscillated around 2.5. This is probably due to the formation of more than one crystallographic structure at non-isothermal conditions. The formation of various polymorphs also might explain higher E_a of crystallization at non-isothermal conditions than at fixed temperature.

THE EFFECTS OF POLYMERIC ADDITIVES ON PHYSICAL STABILITY OF AMORPHOUS NMS

In this part of the paper we shall investigate how three polymers that are frequently used in the pharmaceutical industry, Soluplus (SOP), inulin (INL) and poly(vinylpyrrolidone) (PVP), affect the physical stability of the amorphous form of NMS. Since one of the main factors affecting the re-crystallization of disordered pharmaceuticals is molecular mobility, the NMS + polymer systems have been investigated by means of BDS technique. The dielectric loss spectra of systems containing NMS drug and the polymeric additive were measured at the temperature range from 173.15 K to 338.15 K with step of 2.5 K. The representative spectra of the binary amorphous mixtures with 5% of the polymers are presented in Figure 7. As can be seen, all three mixtures exhibit one well resolved secondary relaxation process below the glass transition temperature (lower panels of Figure 7 a, b and c). All these processes are similar to that found in pure NMS, and therefore it is concluded that they originate from dipole reorientation of the drug's molecules. Above the T_g a well pronounced relaxation peak, corresponding to the structural α -relaxation process, is observed in the dielectric spectra (see upper panels of Figure 7 a, b and c). It moves toward higher frequencies with increasing temperature, and its intensity begins to rapidly decrease at $T = 326$ K for NMS + 5% INL, and $T = 343$ K – for NMS + 5% SOP and NMS + 5% PVP. This drop in intensity of the α -peak is due to the crystallization onset.

Figure 7. Dielectric loss spectra of binary amorphous mixtures of NMS drug and 5% weight of INL (black panels), SOP (cyan panels) and PVP (pink panels) obtained on heating. The upper panels present spectra collected above the glass transition temperature, whereas lower panels show spectra below the T_g .

As can be seen, the devitrification of the mixture containing 5% of INL begins at lower temperature than for the other mixtures. Therefore we can conclude that INL stabilizes NMS the least. Similar result has been obtained by using DSC experiment (see Figure 8). The heating rate applied in these DSC experiments is equal to 10 K/min.

Figure 8. The DSC thermograms of pure NMS, NMS + 5% of INL, NMS + 5% of SOP, and 5% of PVP measured at heating rates equal to 10 K/min.

In order to check how higher concentration of PVP and SOP will affect on NMS's physical stability, we additionally prepared for further studies the mixtures of NMS + SOP and, NMS + PVP with a higher (20% weight) concentration of the polymers. From the analysis of dielectric loss spectra collected both above and below the T_g of all examined systems (i.e. NMS + 5% INL, NMS + 5% SOP, NMS + 5% PVP, NMS + 20% SOP and NMS + 20% PVP), the temperature dependences of α - and β - relaxation times for all of these mixtures (see Figure 9) were obtained. To determine the τ_α and τ_β we employed the same method used previously for pure NMS.

Figure 9. Relaxation maps of NMS and mixtures containing NMS and 5% or 20% polymer (INL,SOP or PVP). Temperature dependence of τ_α in the supercooled liquid has been described by VFT equation. The temperature dependence of τ_β was fitted by using the Arrhenius equation.

The Vogel-Fulcher-Tamman equation was used to describe the $\tau_\alpha(T)$ dependences, while the $\tau_\beta(T)$ dependences were fitted to the Arrhenius law. The fitting parameters for the VFT and the Arrhenius equations are collected in Table 4. As can be seen in Figure 9 the greatest changes in the temperature dependence of structural relaxation times can be observed in the case of the mixtures containing PVP, suggesting that this polymer may be the most successful at stabilizing the amorphous form of NMS drug.

Sample	<u>α-process</u>			<u>β-process</u>	
	T_0 [K]	$\log \tau_{\infty-\alpha}$	B	$\log \tau_{\infty-\beta}$	E_β [kJ/mol]
Pure NMS	216.1 ± 3.8	-19.4 ± 0.7	3711 ± 310	-14.2 ± 0.2	41.9 ± 0.9
NMS + 5% INL	182.4 ± 5.4	-24.8 ± 1.0	6609 ± 568	-14.4 ± 0.1	42.4 ± 0.5
NMS + 5% SOP	210.9 ± 3.9	-19.2 ± 0.6	3881 ± 299	-11.8 ± 0.2	30.4 ± 0.7
NMS + 5% PVP	213.0 ± 2.2	-19.1 ± 0.4	3841 ± 171	-12.1 ± 0.1	31.5 ± 0.5
NMS + 20% SOP	198.1 ± 6.6	-19.3 ± 0.8	4499 ± 485	-13.4 ± 0.1	33.9 ± 0.4
NMS + 20% PVP	237.9 ± 4.8	-15.7 ± 0.6	2588 ± 259	-12.6 ± 0.1	30.8 ± 0.3

Table 4. The fitting parameters from the VFT and Arrhenius equations.

In order to check how effectively the stability of disordered NMS was improved by using the examined polymers, the isothermal kinetics of crystallization, at four various temperatures for NMS's mixtures containing 5% INL, 5% SOP, 5% PVP, 20% SOP and 20% PVP, have been measured. Since the tested mixtures exhibit different tendencies to crystallize, it was not always possible to measure them at the same temperatures. The crystallization kinetics studies were carried out at 318 K, 323 K, 328 K and 333 K for NMS + 5% INL; at 328 K, 330.5 K, 333 K and 338 K for NMS + 5% SOP; at 328 K, 330.5 K, 333 K and 338 K for NMS + 5% PVP, at 343 K, 353 K, 358 K and 363 K for NMS + 20% SOP; and at 360.5 K, 363 K, 365.5 K and 368 K for NMS + 20% PVP. For each measurement a new sample was prepared by the vitrification method.

The method used to analyze the crystallization kinetics of all samples was analogous to that used for pure NMS, i.e. using the Avrami model. The temperature dependences of the logarithm of the crystallization rate k are presented in Figure 10. This plot has the fundamental purpose of answering the following two questions: i) how do the polymeric additives change the crystallization activation energy?; and ii) what is the difference in the $\log k$ values of pure NMS and various binary amorphous mixtures at fixed temperature?

Figure 10. The temperature dependence of $\log(k)$ of pure NMS and binary amorphous mixtures containing the drug and 5% or 20% polymer (INL, SOP or PVP). Solid lines in this figure denote the linear fit to the Arrhenius equation. The inset a presents the crystallization half-life time of NMS and all prepared mixtures checked at a temperature of 328 K. The inset b present the $\log_{10}k$ as a function of $\log_{10}\tau_a$.

The crystallization activation energy of the mixture containing 5% INL is comparable to the value of 126 kJ/mol estimated for the pure NMS. In contrast, both 5% additive of SOP and 5% additive of PVP result in an increase in this value. The E_a of NMS + 5% SOP and NMS + 5% PVP are equal to 146 ± 6 kJ/mol and 205 ± 16 kJ/mol, respectively. A greater amount of SOP polymer (20%) brings about an increase in the activation energy of crystallization to 156 ± 4 kJ/mol. However, the most significant effect has been observed after adding 20% PVP to NMS. In this case, the activation energy reached the highest value of 324 ± 28 kJ/mol.

A useful measure of the capability of polymers to stabilize the NMS drug is the crystallization half-life time ($t_{1/2}$).³⁸ It defines the time required to reach half of the final crystallinity and it has been calculated for each measured sample at $T = 328$ K by using the following relation:

$$t_{1/2} = \left(\frac{\ln 2}{K} \right)^{1/n} \quad (12)$$

The values of $t_{1/2}$ are plotted versus concentration of polymers in the inset of Figure 10. As can be seen, the half-life time at $T = 328$ K for pure NMS is equal to 33 min, while for samples

containing 5% INL, SOP and PVP $t_{1/2} = 1.8$ h, $t_{1/2} = 6.7$ h and $t_{1/2} = 8.8$ h, respectively. If the concentration of SOP and PVP is raised to 20% the values of $t_{1/2}$ are equal to 7.3 days for SOP and 40 years for PVP.

The results presented above indicate that INL has indeed the least stabilizing effect on the examined drug from all tested polymers, while the highest inhibition of crystallization was observed with mixtures containing PVP – the polymer which has the greatest impact on NMS's molecular mobility.

In order to find out if the suppression of the drug's re-crystallization in NMS-PVP mixtures is caused only by the antiplastization effect, we examined how the glass transition temperatures of NMS-PVP mixtures change with the polymer concentration. It has been demonstrated many times that if additional interactions between the drug and polymer exist, the experimentally determined T_g values for different concentrations of the additive should deviate from the theoretical dependence that was proposed by Gordon and Taylor:^{39,40}

$$T_g = \frac{w_1 T_{g1} + K w_2 T_{g2}}{w_1 + K w_2} \quad (13)$$

where w_1 and w_2 are the weight fractions of each component, and T_{g1} and T_{g2} correspond to the glass transition temperature of each component. T_g is the glass transition temperature of the mixture, while K is a measure of the interaction between the components, and can be defined as follows:⁴¹

$$K \approx \frac{\Delta C_{p2}}{\Delta C_{p1}} \quad (14)$$

where ΔC_p denotes the change in heat capacity at T_g .

It can be clearly seen in Figure 11 that the T_g values of the NMS-PVP mixture grow continuously with increasing polymer content. A noticeable discrepancy between the predicted values of glass transition temperature and the experimentally determined data was observed. Therefore one can expect that, apart from an antiplasticizing effect also the specific interactions between the drug and excipient, and/or polymeric steric hindrances, may also play an important role in improving NMS's physical stability in binary amorphous NMS-PVP systems.

Figure 11. Glass transition temperatures of the NMS–PVP mixtures. The pink circles correspond to the experimentally determined T_g values, whereas the dashed line represents the prediction of the Gordon–Taylor/Kelley–Bueche equation, calculated from the T_g values of the pure NMS and pure PVP K15. In the inset the chemical structures of both compounds are presented.

To check if specific chemical interactions indeed exist in NMS-PVP mixtures, we performed a series of FTIR experiments. Table 5 presents the position of the main absorption peaks (band assignment was carried out based on work of Galico et al.⁴² and Tajber et al.⁴³) and Figure 11 shows selected FTIR spectra of the amorphous samples. It can be seen that, with exception of the CO stretching vibration of PVP, the bands remained practically at the same position, regardless of the sample composition. The ν_{CO} band showed, however, a blue shift as the NMS content in the sample increased, which indicates that this chemical group becomes less H-bonded in the presence of the drug, most likely due to the more hydrophobic environment limiting the moisture content. Overall, FTIR analysis indicates that NMS and PVP did not interact in the amorphous phase and the overall net of H-bond interactions decreased, consistent with large negative deviations of the T_g values from the prediction, as presented in Figure 11. Consequently, the suppression of NMS's crystallization in NMS-PVP mixtures should be caused by two mechanisms: the polymeric steric hindrances as well as the antiplastization effect exerted by the additive.

NMS (% w/w)	NMS						PVP	
	ν_{NH}	$\nu_{\text{CC}_{\text{ring}}}$	ν_{aNO_2}	$\rho_{\text{rNH}} + \rho_{\text{rCH}_{\text{ring}}} + \nu_{\text{aSO}_2}$	$\nu_{\text{CN}_{\text{nitro}}} + \nu_{\text{sNO}_2}$	ν_{sSO_2}	ν_{CO}	ν_{CN}
100	3284	1599,1589	1522	1343	1318	1154	N/A	N/A
95	3284	1599,1589	1522	1343	1318	1155	1678	N/A
90	3285	1599,1588	1521	1343	1317	1154	1680	N/A

80	N/A	1598,1588	1524	1341	N/A	1156	1676	N/A
60	N/A	1597,1587	1523	1341	N/A	1157	1676	N/A
40	N/A	1596,1587	1523	1341	N/A	1157	1673	1289
20	N/A	1597,1587	1523	1340	N/A	1156	1670	1289
10	N/A	N/A	1524	1339	N/A	1157	1665	1290
0	N/A	N/A	N/A	N/A	N/A	N/A	1664	1290

Table 5. Position of principal FTIR bands of amorphous NMS, PVP and NMS-PVP mixtures (N/A – not applicable; ν – stretching, ν_a – asymmetric stretching, ν_s – symmetric stretching, and ρ_r – rocking vibration).

Figure 12. FTIR spectra of amorphous NMS-PVP systems

CONCLUSIONS

In this paper we investigated how three well known polymers, inulin, Soluplus and PVP, affect the physical stability and molecular mobility of amorphous nimesulide drug. Using broadband dielectric spectroscopy the molecular mobility of pure drug and mixtures containing 5% or 20% wt. of the polymers has been completely characterized. Besides the α -relaxation, NMS and binary amorphous mixtures exhibit one secondary mode. Both the BDS and DSC studies have shown that pure drug easily re-crystallized. To get a full picture of the tendency of amorphous NMS to devitrify, we measured isothermal and non-isothermal kinetics of crystallization. Based on the data we calculated the isothermal and non-isothermal crystallization activation energies. Their values are equal to 126 ± 8 kJ/mol and 210 ± 14 kJ/mol, respectively. The significantly greater (over 35%) value of E_{aNI} than E_{aI} can result from the formation of polymorphs during NMS's crystallization. We also found that the dominant mechanism of devitrification in the case of NMS drug is the diffusion of molecules.

In order to experimentally investigate the effectiveness of crystallization inhibition of the drug by polymeric additives we measured the isothermal crystallization kinetics of all tested substances. Based on these studies we calculated the crystallization activation energies as well as the crystallization half-life times at 328 K of all measured substances. The E_a and $t_{1/2}(\text{at } 328 \text{ K})$ of pure NMS, NMS + 5% INL, NMS + 5% SOP, NMS + 5% PVP, NMS + 20% SOP, and NMS + 20% PVP are equal to 126 ± 8 kJ/mol and 33 min, 126 ± 14 kJ/mol and 1.8 h, 146 ± 6 kJ/mol and 6.7 h, 205 ± 16 kJ/mol and 8.8 h, 156 ± 4 kJ/mol and 7.3 days, and 324 ± 28 kJ/mol and 40 years, respectively.

Our studies indicate that INL is the worst stabilizer for NMS, whereas the most significant stabilization effect can be achieved by adding the polymer PVP to NMS. Probably, the two mechanisms are responsible for suppression of NMS's re-crystallization in NMS-PVP system: the polymeric steric hindrances, and the antiplastization effect exerted by the excipient.

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