A Comparison of Saturated Solubility Enhancement via Spray Drying and Hot Melt Extrusion Processing

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Introduction

The permeability and solubility of a drug are key determinants of its oral bioavailability. Many new active pharmaceutical ingredients exhibit poor bioavailability due to limited aqueous solubility and thus has been recognized as a common and significant challenge for industry. Several techniques have been developed to enhance solubility including (a) salt formation (b) solutions in solvents and cosolvents (c) micelle systems and self-emulsifying drug delivery systems (d) particle size reduction and nanoparticles (e) complexation (f) pro-drugs and (g) amorphous solids and solid dispersions.

Solid solutions are similar to liquid solutions and consist of a single phase irrespective of the number of components. A solid solution of a poorly water soluble drug dissolved in a carrier is of particular interest as a means of improving oral bioavailability. In a solid solution, the drug’s particle size has been reduced to its molecular dimensions resulting in a more rapid dissolution rate. By judicious selection of a carrier, the dissolution rate of the drug can be increased by up to several orders of magnitude. Spray drying and hot melt extrusion are two common processing approaches for the formation of solid dispersions and solid solutions. We report a comparison of saturated solubility enhancement between spray drying and hot melt extrusion for a poorly soluble, new active pharmaceutical ingredient.

Materials & Methods

Study compounds were characterized and used as received from their suppliers. The polymers included Hypromellose Acetate Succinate (HPMCAS, Shin-Etsu), Eudragit RL (Evonik), Eudragit E (Evonik), Kollidon (PVP K90, BASF) Kollidon VA 64 (BASF) and SoluPlus (BASF).

Spray Drying (SD)

Spray dried samples of FSR-101 were prepared from a 10% solution of the drug and polymer in methanol using a Buchi B-290. The following conditions were prepared: Aspirator flow—100%, Nitrogen Gas flow rate: 550 – 600 L/Hr at 6 bar, solution flow rate 2-3 mL/min and a 0.7 mm nozzle tip diameter. The inlet temperature adjusted to maintain the outlet temp at 50-55 ºC. The resultant dispersions were then dried in a vacuum oven at 35 - 40 ºC for 24 hours.

Hot Melt Extrusion (HME)

Powders of the drug and polymer (25% w/w drug content) were blended in the Turbula mixer (Turbula Type T2 F) and introduced to the Haake Mixing Bowl (Haake PolyLab OS Rheo Drive 4 and Rheomix OS Mixing Bowl). The initial temperature was set according to the glass transition temperature of each polymer, generally about 25-50 ºC above the Tg. The mixing speed was set at 100 rpm.

Powder X-ray Diffraction Analysis (PXRD)

A Bruker D8 Advance diffraction meter equipped with a VANTEC-1 detector was utilized. Samples weighing approximately 100 mg were packed in 0.5-mm-deep cups and were spun at 15 rpm to minimize crystal orientation effects. The X-ray source was operated at 45 kV and 40 mA. Data for each sample were collected from 3° to 40° on the 2θ scale in continuous detector scan mode at a scan speed of 2 s/step and a step size of 0.04°/step.

Differential Scanning Calorimetry (DSC)

DSC analyses were performed using a Thermal Analysis Q2000 differential scanning calorimeter equipped with an autosampler. Samples were equilibrated at the desired RH overnight then crimped into hermetic aluminum pans with a pinhole lead. The samples were equilibrated at 25°C for 5 minutes followed by modulating the temperature at 0.32°C/min while increasing the temperature to 200°C using dry nitrogen at 40 mL/min.

Saturated Solubility Testing

Saturated solubility was measured by adding an excess of drug or dispersion to PBS solution at 37°C. The suspension was agitated at 37°C for at least 1 h and then centrifuged and filtered with a 0.45 µm membrane. The concentration of the supernatant was measured by HPLC.

Materials & Methods Continued

High-Performance Liquid Chromatography (HPLC)

Drug purity was determined using a Waters Alliance system with 2695X Separations Module, 2489 UV Vis Detector and Empower software. A gradient method was used, the column was XBridge phenyl with a 3.5 µm particle size, the flow rate was 1 mL/min, the column temperature was 30°C and the injection volume was 10 µL. The method was qualified for specificity (against the polymers), linearity, accuracy and precision.

Results & Discussion

Table 1. Purity & DSC Results of Hot Melt Extrudates

<table>
<thead>
<tr>
<th>Sample</th>
<th>Purity (%)</th>
<th>% Crystalline</th>
</tr>
</thead>
<tbody>
<tr>
<td>SoluPlus-HME</td>
<td>92.8</td>
<td>7.2</td>
</tr>
<tr>
<td>Eudragit E HME</td>
<td>88.4</td>
<td>11.6</td>
</tr>
<tr>
<td>Eudragit RL HME</td>
<td>89.4</td>
<td>10.6</td>
</tr>
<tr>
<td>HPMCAs 90/10</td>
<td>87.1</td>
<td>12.9</td>
</tr>
<tr>
<td>VA 90/40</td>
<td>93.5</td>
<td>6.5</td>
</tr>
</tbody>
</table>

The HME samples were analyzed by PXRD (Figure 1). All were found to be amorphous except for Kollidon 90. A single phase glass transition temperature was observed by DSC for the SoluPlus, Eudragit RL, Eudragit E and Kollidon VA-64 samples (Table 1). FSR-101 purity was determined to be acceptable for all samples except for the HPMCAs sample, which had ~20% degradation (Table 1). This observation is in contrast to the chemical stability of the spray dried dispersion.

The saturated solubility and morphology following a 10 day stability study of the HME samples was determined and is reported in Table 2. The saturated solubility of the SoluPlus, Eudragit RL, Eudragit E and Kollidon VA-64 samples was exceptional, and was greater than the spray dried dispersion. These samples were confirmed to be amorphous following a 10 day stability study.

Conclusions

Amorphous dispersions were successfully prepared by both spray drying and melt extrusion. For the spray drying process, HPMCAS was found to have the best saturated solubility, with no re-crystallization observed on stability. However, FSR-101 was found to degrade when processed by melt extrusion with this polymer. The saturated solubility of the amorphous dispersions prepared by melt extrusion exceeded those prepared by spray drying. This study demonstrates that amorphous dispersion can be prepared by spray drying and melt extrusion but the physicochemical properties of the dispersion are dependent upon the process.

References