ABSTRACT

Objective: The present investigation was undertaken with an objective of formulating oral disintegrating tablets of amlodipine besylate to enhance convenience and compliance of the elderly and pediatric patients for better therapeutic efficacy.

Methods: The tablets were prepared by using direct compression method and evaluated for weight variation, hardness, friability, wetting time, disintegration time and in vitro drug release study. Prepared tablets were evaluated for compatibility by Fourier transform infrared spectroscopy.

Results: Fourier transform infrared spectroscopy studies revealed that there was no physicochemical interaction between amlodipine besylate and other excipients. All the tablets hardness was found to be around 3.5 kg/cm² and friability of all the formulations was less than 1%. Drug content in all the formulations was found in the range of 97.05% to 99.13%.

Conclusion: The study clearly indicated that the type and concentration of superdisintegrants plays an important role in disintegration and dissolution of drug from oral disintegrating tablets. Among all the formulations, the maximum percentage of drug release and less disintegration time was found in F9 formulation containing 4% of crospovidone.

Keywords: Oral disintegrating tablets, Amlodipine besylate, ODTs, Crospovidone, Croscarmellose sodium

INTRODUCTION

Solid dosage forms are the most popular and preferred drug delivery systems. Most of the oral pharmaceutical dosage forms like conventional tablets and capsules are formulated to be swallowed or chewed. Many pediatric, geriatric and bedridden patients find difficulty in swallowing (i.e., dysphagia) these dosage forms [1]. It is estimated that 50% of the population is affected by this problem which results in a high incidence of non-compliance and ineffective therapy [2]. The concept of oral disintegrating drug delivery system emerged from the desire to improve patient compliance. An oral disintegrating tablet offers quick disintegration and dissolution of the tablet in the oral cavity within a minute without the need of water. According to European pharmacopoeia, oral disintegrating tablet should disperse or dissolve in less than three minutes. The medication can be absorbed partially or entirely into the systemic circulation from blood vessels in the oral mucosa; it can be swallowed as a solution to be absorbed from the gastrointestinal tract [3, 4].

Angina pectoris is chest pain due to ischemia of heart muscle due to obstruction or spasm of the coronary arteries. Amlodipine besylate is a long-acting calcium channel blocker mainly used in the treatment of chronic stable angina vasospastic angina and hypertension [5]. Amlodipine is listed in World health organization (WHO) under the list of essential medicines as antihypertensive medicine in 5 mg tablet. It inhibits the transmembrane influx of calcium ion into vascular smooth muscle and cardiac muscle. Its efficacy is not only limited to the treatment of cardiovascular diseases as it has shown to possess antioxidant activity and plays an important role in apoptosis. Peak plasma concentrations are reached 6-12 h. It has oral bioavailability of 64-90% and half-life of about 30-50 h [6]. Amlodipine besylate is slightly soluble drug the rate of absorption is controlled by the rate of dissolution. The rate of dissolution can be increased by the use of super disintegrants [7]. Oral disintegrating tablets are designed to be placed in mouth allowed to disperse or dissolve in the saliva and then swallowed without the aid of water. As the tablet disintegrates in the mouth, this could enhance the clinical effect of the drug through pre-gastric absorption [8]. The objective of the present study was to deliver the drug at a faster rate and to provide immediate onset of action in a shorter period of time with improved bioavailability. The basic approach in the development of oral disintegrating tablets is the use of superdisintegrants like croscarmellose sodium, sodium starch glycolate and crospovidone.

MATERIALS AND METHODS

Amlodipine besylate was obtained from Yarrow chemicals, Mumbai. Sodium starch glycolate, croscarmellose sodium and crospovidone, microcrystalline cellulose, and mannitol were purchased from Aman scientific industries, Vijayawada. Talc and magnesium stearate were purchase from Loba Chemie Pvt. Ltd. All other chemicals/solvents used were of analytical grade.

Estimation of ambroxol hydrochloride

Lab India double beam UV-Visible spectrophotometer was used to estimate amlodipine besylate in pure form and formulations developed using various concentrations of super disintegrants at 237.5 nm.

An analytical method for construction of calibration curve of amlodipine besylate

100 mg of amlodipine besylate was dissolved in 100 ml of 0.01N hydrochloric acid. 10 ml of this solution was further diluted to 100 ml of 0.01N hydrochloric acid to obtain 100 μg/ml From this solution (100 μg/ml) suitable working solutions of different concentrations of 3, 6, 9, 12, 15 and 18 μg/ml were prepared. The absorbance of these dilutions was measured at the determined lambda max of 237.5 nm. The standard graph of concentration versus absorbance was then plotted. Each point is an average of three determinations. Slope, intercept, and regression coefficients were calculated [9].

Drug-excipient compatibility study

Fourier transform infrared spectroscopic (FTIR) studies were conducted to know the compatibility between drug and excipients. In these studies pure amlodipine besylate and its mixture with sodium starch glycolate, croscarmellose sodium, and crospovidone
were grounded thoroughly with IR grade KBr and then compressed in a hydraulic press at a pressure of 10,000 psi. Each disc was scanned over a range of 400-4000 cm⁻¹ using FTIR instrument (FTIR-1600, Shimadzu, Japan). The characteristic peaks were observed and recorded.

Preparation of oral disintegrating tablets

Oral disintegrating tablets of amlodipine besylate were prepared by direct compression method. All the ingredients listed in table 1 were accurately weighed and passed through a # 60 sieve. Amlodipine besylate (10 mg) was mixed with various concentrations of super disintegrants, microcrystalline cellulose, and mannitol. Then magnesium stearate and talc were added to the above blend.

The flow property of the final blend was determined by measuring the angle of repose and Carr's index and they were found to be satisfactory to allow the mixture to be directly compressed into tablets on a 12-station rotary tablet punching machine (M/s. CEMACH Machinery, Co. Pvt. Ltd., India) using 8 mm punch [10].

Table 1: Composition of oral disintegrating tablets of amlodipine besylate

<table>
<thead>
<tr>
<th>Ingredients (mg/tablet)</th>
<th>F1</th>
<th>F2</th>
<th>F3</th>
<th>F4</th>
<th>F5</th>
<th>F6</th>
<th>F7</th>
<th>F8</th>
<th>F9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amlodipine besylate</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Sodium starch glycolate</td>
<td>2</td>
<td>4</td>
<td>6</td>
<td>2</td>
<td>4</td>
<td>6</td>
<td>2</td>
<td>4</td>
<td>6</td>
</tr>
<tr>
<td>Crosscarmelleose sodium</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
<td>66</td>
</tr>
<tr>
<td>Crospovidone</td>
<td>65</td>
<td>63</td>
<td>61</td>
<td>65</td>
<td>63</td>
<td>61</td>
<td>65</td>
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<tr>
<td>Mannitol</td>
<td>1</td>
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<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
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<tr>
<td>Microcrystalline cellulose</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>3</td>
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<tr>
<td>Sodium saccharin</td>
<td>3</td>
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<td>3</td>
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<td>3</td>
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<tr>
<td>Magnesium stearate</td>
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<td>3</td>
<td>3</td>
<td>3</td>
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</tr>
<tr>
<td>Talc</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
<td>150</td>
</tr>
</tbody>
</table>

Micromeritic properties of the blended powder

Angle of repose

The angle of repose of powder blend was determined by the fixed funnel method. The accurately weighed powder was taken in a funnel. The height of the funnel was adjusted in such a way that the tip of the funnel just touched the apex of the heap of the powder. The powder was allowed to flow through the funnel freely onto the surface. The diameter of the powder cone was measured and the angle of repose was calculated using the following equation [11].

\[ \tan \theta = \frac{h}{r} \]

Where, θ = angle of repose,

h = height of the pile,

r = radius of the pile base

Bulk Density and tapped density

A quantity of 2 g of powder from each formula, previously lightly shaken to break any agglomerates formed was introduced into a 10 ml measuring cylinder. After the initial volume was observed then individual tablet weight was compared with average weight [12].

Tablet hardness and friability

For each formulation, the hardness of 6 tablets was determined using the Monsanto hardness tester. Friability of tablets was measured by using Roche Friablator. Friability was evaluated from the percentage weight loss of 10 tablets tumbled in a friabulator at 25 rpm for 4 min. The tablets were deducted and the loss in weight caused by fracture or abrasion was recorded as the percentage weight loss. Friability below 1% was considered acceptable [12].

\[ \text{Friability} = \frac{\text{Initial weight} - \text{Final weight}}{\text{Initial weight}} \times 100 \]

Uniformity of drug content

10 Tablets were weighed from each formulation and triturated in a mortar to a fine powder. Powder equivalent to 10 mg of amlodipine besylate was extracted in 10 ml of methanol and liquid was filtered. The drug content was determined by measuring the absorbance at 237.5 nm (using a UV-Visible spectrophotometer, Lab India) after appropriate dilutions with methanol. The drug content was determined using a calibration curve. The mean percentage drug content was calculated as the average of three determinations.

Wetting time

The wetting time of the tablets can be measured using a simple procedure. Five circular tissue papers of 10 cm diameter are placed in a petridish with a 10 cm diameter. 10 ml of water containing amaranth, a water-soluble dye is added to petridish. A tablet is carefully placed on the surface of the tissue paper. The time required for water to reach the upper surface of the tablet is noted as a wetting time [13].

Water absorption ratio

A piece of tissue paper folded twice was placed in a small petridish containing 6 ml of water. A tablet was put on the tissue paper and allowed to completely wet. The wetted tablet was then weighed. Water absorption ratio, R was calculated using the following equation [14].

\[ R = \frac{W_a - W_b}{W_b} \times 100 \]

Where,

Wa = Weight of tablet after water absorption

Wb = Weight of tablet before water absorption
In vitro dispersion time

The amount of saliva available in the oral cavity is very limited (usually less than 6 ml). In a simplest method, 6 ml of phosphate buffer of pH 6.8 was taken in a 25 ml measuring cylinder. The temperature was maintained at 37 ± 2°C. An oral disintegrating tablet was placed into it and time required for complete disintegration of the tablet was noted [15].

In vitro drug release studies

In vitro drug release studies were carried out using Type II-USP dissolution test apparatus, Paddle (DBK dissolution testing apparatus, Mumbai, India) at 75 rpm. The dissolution medium consists of 500 ml of 0.01N hydrochloride acid. The medium was maintained at 37 ± 0.5°C. Aliquot (5 ml) of the solution was collected from the dissolution apparatus at 5, 10, 15, 20, 25 and 30 min. and replenishment with an equal volume of fresh dissolution medium. The withdrawn samples were analyzed by an UV-Visible spectrophotometer (Lab India) at 237.5 nm using 0.01N hydrochloride acid as a blank. Aliquots were withdrawn from a zone midway between the surface of dissolution medium and the top of rotating paddle [16].

RESULTS AND DISCUSSION

An analytical method for construction of a calibration curve

In order to conduct the in vitro drug dissolution studies calibration curve was plotted to determine R² and the equation of straight line is used to calculate drug release. Calibration curves of amlodipine besylate in 0.01N hydrochloride acid was constructed at lambda max of 237.5 nm and is shown in the fig. 1

![Calibration curve of amlodipine besylate](image)

**Table 2: Micromeritic properties of the blended powder**

<table>
<thead>
<tr>
<th>Formulation</th>
<th>Bulk density (gm/cc)</th>
<th>Tapped density (gm/cc)</th>
<th>Carr’s index (%)</th>
<th>Hausner ratio</th>
<th>Angle of repose (θ)</th>
</tr>
</thead>
<tbody>
<tr>
<td>F1</td>
<td>0.43±0.07</td>
<td>0.52±0.02</td>
<td>16.09±0.04</td>
<td>1.19±0.02</td>
<td>27.5±0.01</td>
</tr>
<tr>
<td>F2</td>
<td>0.44±0.05</td>
<td>0.51±0.03</td>
<td>14.06±0.03</td>
<td>1.16±0.01</td>
<td>29.0±0.04</td>
</tr>
<tr>
<td>F3</td>
<td>0.45±0.01</td>
<td>0.52±0.05</td>
<td>14.12±0.04</td>
<td>1.16±0.01</td>
<td>27.7±0.03</td>
</tr>
<tr>
<td>F4</td>
<td>0.43±0.09</td>
<td>0.52±0.08</td>
<td>16.39±0.03</td>
<td>1.19±0.02</td>
<td>29.7±0.04</td>
</tr>
<tr>
<td>F5</td>
<td>0.43±0.03</td>
<td>0.51±0.09</td>
<td>15.44±0.02</td>
<td>1.18±0.03</td>
<td>27.1±0.03</td>
</tr>
<tr>
<td>F6</td>
<td>0.44±0.03</td>
<td>0.51±0.09</td>
<td>12.74±0.07</td>
<td>1.14±0.04</td>
<td>28.3±0.06</td>
</tr>
<tr>
<td>F7</td>
<td>0.42±0.04</td>
<td>0.51±0.04</td>
<td>16.69±0.09</td>
<td>1.20±0.07</td>
<td>27.5±0.04</td>
</tr>
<tr>
<td>F8</td>
<td>0.42±0.05</td>
<td>0.50±0.04</td>
<td>15.71±0.03</td>
<td>1.18±0.03</td>
<td>28.0±0.01</td>
</tr>
<tr>
<td>F9</td>
<td>0.44±0.06</td>
<td>0.51±0.04</td>
<td>12.74±0.01</td>
<td>1.14±0.04</td>
<td>27.7±0.03</td>
</tr>
</tbody>
</table>

All the values are expressed as mean±SD, where n=3.

Oral disintegrating tablets of amlodipine besylate were prepared by direct compression method employing different concentrations of sodium starch glycolate, croscarmellose sodium and crospovidone. A total of nine formulations were designed. The formulation F1, F2, F3 having sodium starch glycolate as superdisintegrants. Formulation F4, F5, F6 having croscarmellose sodium as superdisintegrants. Formulations F7, F8, F9 having crospovidone as superdisintegrants. The flow properties of the powder blend were measured before compression. Results were shown in the table-2. The angle of repose values ranged from 27.5±0.03 to 29.7±0.04. Compressibility index and Hausner’s ratio values were ranged from 12.74±0.01 to 16.69±0.09 and 1.14±0.02 to 1.20±0.07 respectively. Low Hausner’s ratio (less than 1.25), compressibility index (less than 17 %) and angle of repose (less than 30°) indicated that the powder blend had good free-flowing property.

Oral disintegrating tablets of amlodipine besylate were prepared by direct compression technique. The prepared tablets were evaluated for different physicochemical properties and results were depicted in tables 3 and 4. All the tablets passed weight variation test as the % weight variation was within the pharmacopoeial limits of 7.5%. Hardness (3.6±0.11 to 3.9±0.12 kg/cm²) and friability was less than 1% in all the formulations indicated that tablets had good mechanical resistance. Uniformity of the amlodipine besylate was determined by double bean UV spectrophotometer (LABINDIA), the content of amlodipine besylate was found within official limits. Drug content of all the tablets was found in the range of 97.05% to 99.13%.
Wetting time evaluation was used to predict the penetration of medium into the structure of oral disintegrating tablets. The results of wetting time evaluation were shown in Table 4. The wetting time of all the formulations was found to be in the range of 24.48±1.1 to 91.0±1.2 sec. Water absorption ratio was important parameter for understanding the capacity of disintegrant to swell in the presence of a little amount of water. The difference in water absorption ratio among the formulations was due to water uptake and swelling behaviour of superdisintegrants. Water absorption ratio of all the formulation was found to be in the range of 56.2±0.2 to 92.85±0.9.

In vitro dispersion time is used to measure the time required by the tablets to produce complete dispersion. The most important parameter that needs to be optimized in the development of oral disintegrating tablets is the disintegration time of the tablets. As the concentration of super disintegrant increases the disintegration time decreases. In vitro dispersion time of all the formulated tablets was found to be within 40 sec. Fast disintegration was observed with crospovidone as it has very high capillary activity and hydration capacity. The water uptake stretches out the folded molecular chains lying between the crosslinks, causing an instant expansion of the polymer. The increase of volume creates an internal pressure exceeding that of tablet strength, which resulted in fast disintegration of tablet. The dissolution study of oral disintegrating tablets was performed in 0.01N hydrochloric acid using USP type II apparatus at 75 rpm. The comparative drug release profile was shown in Fig. 2.

The dissolution rate was found to be comparatively less for the formulations containing sodium starch glycolate. The maximum increase in the dissolution rate was observed with crospovidone. The order of dissolution rate with various superdisintegrants was found to be crospovidone>croscarmellose sodium>sodium starch glycolate. The maximum percentage of drug release was found in F9 formulation containing 4% of crospovidone, which shows 82.2% of drug release at the end of 10 min and 98.92% at the end of 30 min. Based on above results formulation F9 was selected as best formulation.
FTIR studies

The results of FTIR spectrum of pure amlodipine besylate and optimized formulation F9 was shown in fig. 3 and 4 respectively. The characteristic peaks of the drug were observed in the spectra of drug and optimized formulation F9 indicates that there is no interaction between the drug, super disintegrant and other excipients.

CONCLUSION

Oral disintegrating tablets of amlodipine besylate were successfully prepared by direct compression method using crospovidone, croscarmellose sodium and sodium starch glycolate. All the oral disintegrating tablets possessed good micromeritic, physicochemical and dissolution properties. Among the nine formulations prepared, the formulation F9 containing 4% of Crospovidone gave higher in vitro drug release of 98.92% at the end of 30 min. The developed oral disintegrating tablets may provide quick onset of action with improved oral bioavailability and enhanced patient compliance and therapeutic efficacy.

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AUTHORS CONTRIBUTIONS

All the author have contributed equally

CONFLICT OF INTERESTS

Declared none
REFERENCES