DEVELOPMENT AND EVALUATION OF LAMOTRIGINE SOYA LECITHIN SOLID DISPERSION: IN VITRO AND PHARMACODYNAMIC INVESTIGATION

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ABSTRACT

Objective: Epilepsy is a common neurodegenerative disorder characterized by spontaneous and repeated attacks of convulsions. It requires immediate pharmacotherapy to prevent its progression to status epilepticus. However, most of the anticonvulsant drugs are poorly water-soluble and demonstrate the delayed onset of action. Thus there is a need to improve its solubility for the better pharmaceutical profile. The objective of the present investigation was to enhance the solubility of lamotrigine incorporating soya lecithin as a phospholipid carrier by solid dispersion technique.

Methods: Solid dispersions of lamotrigine were prepared with soya lecithin by the solvent method. The effect of concentration of phospholipid and solvents on aqueous solubility and dissolution profile of lamotrigine was analyzed.

Results: Ethanol increased lamotrigine solubility with soya lecithin in ratio 5:1. X-ray diffraction and scanning electron micrograph indicated a smaller crystallite size of lamotrigine with fairly uniform size distribution in the lamotrigine-soya lecithin solid dispersion. The resultant solid dispersion also significantly delayed the onset of clonic convulsion (875.8 s) as compared to control (85.5 s) and offered complete protection (100%) against the pentylenetetrazole induced seizures in the rat as compared to control (33.33%). Also, solid dispersion with maximum drug content (77.68%) and dissolution rate (91.40%) was formulated as an orodispersible tablet and characterized for its pharmaceutical properties.

Conclusion: It can be concluded that the solid dispersion of lamotrigine incorporated with soya lecithin demonstrated enhanced solubility and dissolution rate may have potential clinical application.

Keywords: Lamotrigine, Phospholipids, Solid dispersion, Anticonvulsant activity, Pentylenetetrazole

INTRODUCTION

Epilepsy is a very common neurodegenerative disorder characterized by repeated episodes of epileptic attacks. It required immediate medical help to prevent its progression to status epilepticus [1]. Although pharmacological treatment has been more pronounced due to the availability of new generation anticonvulsant drugs, some of them are poorly water-soluble and show delayed onset of action following their oral administration [2]. Low aqueous solubility is a major problem encountered with the formulation development of such drugs [3]. If the rate of dissolution of the drug is significantly slower than the rate of absorption, the dissolution of the drug becomes the rate-limiting step in the absorption process. Different formulation approaches have been used to improve the oral absorption of drugs with low water solubility by increasing the dissolution rate and solubility [4, 5].

Lamotrigine is a second-generation antiepileptic drug that shows a broad spectrum of actions against partial, generalized tonic-clonic seizures and Lennox-Gastaut syndrome either alone or as adjunctive therapy [6, 7]. Comparatively, lamotrigine shows relatively few side effects and does not require blood monitoring in monotherapy [8, 9]. However, peak plasma concentration reaches after 1.4 h its oral administration. This delay in the onset of action in spite of good bioavailability is due to its low aqueous solubility (0.17 g/l) [2, 8]. Hence it is required to improve the solubility and dissolution of lamotrigine.

Enhancement of the solubility of poorly water drugs is a difficult task. Several methods have been used for solubility enhancement such as salt formation [10], complexation [11], solid dispersion [12], microcrystals [13], microemulsion [14] etc. One successful approach to enhance the solubility is solid dispersions using an appropriate carrier. The dissolution rates may be improved by dispersing drugs in polymeric carriers by fusion and solvent evaporation methods [3]. Though various solid dispersion techniques are available, the solvent method is advantageous as it requires low temperature for the evaporation of organic solvents that prevents the decomposition of drugs and carriers [15]. The preparation of solid dispersion using soluble carriers increases the solubility and dissolution rate of the drug, as it is exposed to the dissolution medium in the very fine particulate form [8, 16]. Hydrophilic polymers have been used to enhance the solubility and dissolution rate of lamotrigine. Some of the investigators have prepared and evaluated the formulations of solid dispersion of lamotrigine using hydrophilic polymers [8, 17]. However, it requires a great amount of the carrier for the preparation of solid dispersion. Phospholipids are required in much lower concentration for increasing the dissolution rate of poorly water-soluble drugs. These forms spherical bilayer structures (liposomes) when it is dispersed in aqueous media into which the drug is entrapped or sequestered [18-20] and released rapidly into the dissolution medium. Besides the assimilation of drugs with lipids or lipid-like compounds promotes the oral absorption of the drug by intrinsic lipid pathway [21].

Soya lecithin, the phospholipid is a multi-functional surface-active agent having various applications. Phospholipids due to their amphiphilic property have the solubilizing potential for poorly water-soluble drugs [22, 23] can be used as a carrier for the preparation of solid dispersion. With this background, the present investigation was carried out to enhance the solubility and dissolution of lamotrigine incorporating soya lecithin as a phospholipid carrier by solid dispersion technique. Also to determine the anticonvulsant activity of prepared formulation in rats and to develop an orally disintegrating tablet containing solid dispersion to achieve fast release of drug from the dosage form.

MATERIALS AND METHODS

Materials

Lamotrigine was obtained as a gift sample from Glenmark, Mumbai, soya lecithin was purchased from Himedia Lab. Mumbai, DMSO, and methanol were procured from Merck chemicals, Mumbai. Ethanol
was procured from Changshu Hongsheng Fine Chemical Co. Ltd Jiangsu Province. Chloroform was purchased from Loba Chemie Mumbai, croscarmellose sodium from Akhil Healthcare Mumbai and cross povidone was obtained from Sigma Aldrich, Mumbai. All the chemicals used in experiments were of analytical grade.

Preparation of phospholipid solid dispersions and physical mixture
Solid dispersions of lamotrigine and phospholipids were prepared by the solvent method [24, 25] using chloroform, DMSO, and ethanol as solvent. Weighed amounts of phospholipid and lamotrigine (ratio of 1:1, 1:3, 1:5 and 1:10) were added to the solvent and dissolved with gentle stirring further sonicated for 30 min after complete dissolution solvent was removed at room temperature or up to 60 °C temperature if required. Further drying was accomplished in a vacuum desicator overnight. Physical mixtures were prepared by gently triturating appropriate quantities of lamotrigine and phospholipid using a mortar and pestle. Solid dispersions and physical mixtures were passed through an 80-mesh sieve to obtained uniform size powder for further studies. The formulation composition for solid dispersion is depicted in table 1.

Table 1: Composition of lamotrigine solid dispersion

| Ingredients                       | S1   | S2   | S3   | S4   | S5   | S6   | S7   | S8   | S9   | S10  | S11  | S12  |
|-----------------------------------|------|------|------|------|------|------|------|------|------|------|------|------|
| PLP:LTG                           | 1.1  | 1.5  | 1.5  | 1:10 | 1:1  | 1:5  | 1:10 | 1:1  | 1:5  | 1:1  | 1:5  | 1:10 |
| DMSO+Ethanol (ml) (2:1)           | 10   | 10   | 10   | 10   | -    | -    | -    | -    | -    | -    | -    | -    |
| Ethanol ml                        | -    | -    | -    | -    | 10   | 10   | 10   | 10   | -    | -    | -    | -    |
| Chloroform+ Ethanol (ml) (2:1)    | -    | -    | -    | -    | -    | -    | -    | -    | 10   | 10   | 10   | 10   |

Saturation solubility
Saturation solubility was performed, adding an excess of the amount of sample in 20 ml distilled water in a screw-capped flask and shaken in a rotary flask shaker (20 rpm) at room temperature (37±0.5 °C) for 48 h. Once equilibrium was achieved appropriate aliquots were withdrawn and filtered through Whatman filter paper no. 45. The filtrate was analyzed for drug spectrophotometrically at 307 nm [26].

Drug content
Drug content was determined by dissolving solid dispersion or physical mixture equivalent to 10 mg of drug in 10 ml methanol following ultra-sonication for 20 min. The volume was adjusted to 100 ml with a 6.8 pH buffer. The solution was filtered through Whatman filter paper no. 45, further suitably diluted with 6.8 pH buffer and the absorbance was measured at 307 nm using a double-beam UV spectrophotometer [27].

Dissolution studies
The dissolution studies were carried out using the TAB dissolution test apparatus (Cambell DRS Company). The dissolution flasks were immersed in the water bath equipped with an external temperature control unit. 10 mg of the sample was dispersed on the surface of the dissolution medium at the beginning of the study. A USP standard paddle continuously stirred the dissolution medium (900 ml of 6.8 pH buffer) at 100 rpm at 37±0.5 °C. A sink condition was maintained in the dissolution medium. Samples were taken at pre-determined intervals and the concentration of lamotrigine present in the dissolution medium was determined using a double-beam UV spectrophotometer at 307 nm [25].

Characterization of lamotrigine phospholipid solid dispersions

Differential scanning calorimeter (DSC)
The DSC analysis was performed using Mettler Toledo DSC82e using aluminum crucibles with about 6.5-10 mg of samples, under dynamic N₂ atmosphere (10 ml min⁻¹) and heating rate of 20 °C min⁻¹ under a temperature range from 25 °C to 350 °C.

Powder X-ray diffraction studies (PXRD)
X-ray diffraction study was performed to determine the crystallinity of the prepared solid dispersion using Bruker X-ray diffractometer AXS D8 Advance using a Cu radiation (1.5406 Å) at a voltage 2.2 kV, and Detector-Si(Li)PSD. Diffractogram was scanned in the range from 3 °C to 80 °C (2θ) with a resolution of 0.02 °C and scanning speed of 2.0 °C min⁻¹.

Fourier-transform infrared spectroscopy
Fourier-transform infrared spectroscopy (FTIR) spectrum was recorded on Thermo Nicolet, Avatar 370 using the potassium bromide pellet method. About 2 mg of the sample was mixed with potassium bromide, and the mixture was compressed at a pressure of 5 tons for 5 min in a hydraulic press to form the pellets. The sample was scanned at 4 cm⁻¹ resolution.

Scanning electron microscopy (SEM)
A JSM-6390LV field emission scanning electron microscope (JEOL, Peabody, MA) was used to determine the particle shape and size of lamotrigine and solid dispersion. Photomicrograph was taken at 150x0 to 350x0 magnification.

Pharmacodynamic study
The study was carried out to investigate the anticonvulsant effect of an optimized batch of solid dispersion in PTZ induced seizures in rats through oral administration. It is commonly used in preclinical paradigms for predicting the anti-convulsant activity of lamotrigine after its acute administration. Adult male Sprague Dawley rats (National Institute of Nutrition; Hyderabad, India) weighing 200–280 g were housed in standard laboratory conditions of temperature (23±1 °C) and relative humidity (55±5%), with free access to food (NIN, Hyderabad, India) and water. Animals were divided into different groups (n=4), fasted overnight before the experiments and transferred to the laboratory at least 1 h before the beginning of the experiment. The experiments were performed during the light cycle in between 9.00 to 13.00 h. All the experimental protocols were approved by the Institutional Animal Ethics Committee (853/IAEC/2018-19/25). Oral doses of solid dispersion (equivalent to lamotrigine: 5 mg/kg, 10 mg/kg) and pure drug in saline were administered using a polyethylene tube. The tube was inserted about 5-6 mm deep for the proper delivery of drugs orally. 1 h following drug administration, animals were subcutaneously injected with PTZ (60 mg/kg) and the onset of clonic-tonic convulsions and percentage protection in each group were recorded.

Development of orodispersible tablet
For the preparation of oral dispersible tablet using solid dispersion formulation with maximum drug content and better dissolution profile was selected. The formulation composition of lamotrigine orodispersible tablets is depicted in table 2. Tablets were prepared by the direct compression method by taking solid dispersion equivalent to 25 mg of lamotrigine. Microcrystalline cellulose (MCC) and mannitol were used as directly compressible agents. Sodium starch glycolate (SSG), crospovidone (CPD), croscarmellose sodium (CCS) were used as super disintegrant agents in different batches. Aspartame was added as the sweetening agent. Talc and magnesium stearate were added as lubricant and glidant and compressed as a tablet using flat-faced punches. The hardness of the tablets was kept constant and was measured with a hardness tester. Tablet blend and tablets were evaluated for various pre-compression and post-compression parameters [17].
showed a significant increase of 96.68±0.41 % in the extent of lamotrigine to soya lecithin on the dissolution of lamotrigine. Preformulation studies were performed for multiple comparison test. Differences between formulations were performed by one-way analysis of variance followed by Bonferroni's test. All the results are reported as mean ± standard deviation (SD) or ±standard error mean (SEM). Statistical comparisons were performed by one-way analysis of variance followed by Bonferroni's multiple comparison test. Differences between formulations were considered to be statistically significant at P<0.05 in all cases.

RESULTS AND DISCUSSION

Preparation and characterization of lamotrigine solid dispersion

Solid dispersion of lamotrigine-soya lecithin was prepared by the solvent method. Preformulation studies were performed for lamotrigine solubility in ethanol, chloroform, DMSO, ethanol-chloroform and DMSO-ethanol to select the solvent system for the preparation of solid dispersion. Based on the results solid dispersions were prepared by using DMSO-ethanol (2:1), chloroform-ethanol (2:1) and ethanol as solvents. The aqueous solubility of pure lamotrigine was found to be 0.1515 mg/ml. Solid dispersions were prepared by using the various ratio of lamotrigine and soya lecithin using different solvent were analyzed for aqueous solubility. Results revealed an increase in solubility was for the batch S7 prepared by using ethanol as a solvent in comparison to other solvents. Solid dispersion prepared with DMSO and ethanol showed an increase in solubility of lamotrigine as compared to that of the pure drug, but the preparation was more viscous and difficult to dry. Batches prepared with ethanol and chloroform as solvent showed an increase in aqueous solubility, but results were more enhanced for the solid dispersions prepared with ethanol as solvent. Batch S7 consisting of soya lecithin and lamotrigine in ratio 1:5 showed a significant increase in aqueous solubility of lamotrigine i.e. 3.1814 mg/ml as compared to its lower ratios (1:1 and 1:3). However, a further increase in ratio showed a decreased solubility. A physical mixture of lamotrigine with soya lecithin showed an increase in aqueous solubility than pure drug, but it was very less as compared to aqueous solubility of various solid dispersions. Table 3 depicts the results for the effect of solvents and the ratio of phospholipid and lamotrigine on the saturation solubility of lamotrigine.

The drug content of the formulation was ranging from 41.39 % to 77.68 %, indicating maximum entrapment of drug in the formulation. The maximum amount of drug content 77.68±1.35% was obtained for solid dispersion with a 1:5 ratio of soya lecithin and lamotrigine using ethanol as solvent. The percentage of drug content proportionally increased with the drug concentration.

Table 2: Composition of lamotrigine orodispersible tablets

| Ingredients (mg) | F1  | F2  | F3  | F4  | F5  | F6  |
|-----------------|-----|-----|-----|-----|-----|-----|
| SD LTG          | 26.05 | 26.05 | 26.05 | 26.05 | 26.05 | 26.05 |
| SSG             | 6   | 6   | 6   | 6   | 6   | 6   |
| CCS             | -   | -   | 8   | 8   | 8   | 8   |
| CPD             | -   | -   | -   | 6   | 8   | -   |
| Mannitol        | 49.15 | 47.45 | 49.15 | 47.45 | 49.15 | 47.45 |
| MCC             | 8.5 | 8.5 | 8.5 | 8.5 | 8.5 | 8.5 |
| Acpartame       | 3   | 3   | 3   | 3   | 3   | 3   |
| Tak             | 3   | 3   | 3   | 3   | 3   | 3   |
| Aerosil         | 2   | 2   | 2   | 2   | 2   | 2   |
| Mg stearate     | 2   | 2   | 2   | 2   | 2   | 2   |
| Total Weight    | 100 | 100 | 100 | 100 | 100 | 100 |

Table 3: Saturation solubility of lamotrigine, solid dispersions and physical mixture

| Formulation code | Media          | Solubility (mg/ml) | Drug content*(% w/w) |
|------------------|----------------|-------------------|----------------------|
| LTG              | 6.8 Buffer     | 0.4882            | -                    |
| LTG              | Ethanol        | 1.8232            | -                    |
| LTG              | water          | 0.1515            | -                    |
| S1               | water          | 1.6501            | 41.39±0.77           |
| S2               | water          | 1.7526            | 49.84±1.68           |
| S3               | water          | 1.8804            | 61.69±1.75           |
| S4               | water          | 2.2552            | 69.37±0.57           |
| S5               | water          | 0.1942            | 39.83±1.95           |
| S6               | water          | 2.9986            | 57.46±5.90           |
| S7               | water          | 3.1814            | 77.68±1.35           |
| S8               | water          | 3.0400            | 72.55±1.27           |
| S9               | water          | 1.9821            | 40.4±1.94            |
| S10              | water          | 2.0447            | 50.84±0.86           |
| S11              | water          | 2.2356            | 70.32±0.64           |
| S12              | water          | 2.3252            | 68.56±1.16           |
| PM               | water          | 0.1787            | 62.7±1.06            |

Results are shown as mean±SD (* n=3)

Dissolution studies

The lamotrigine-soya lecithin solid dispersions prepared from ethanol exhibited greater dissolution as particles formed were fine, non-waxy and discrete. Fig. 1 shows the effect of different ratios of lamotrigine to soya lecithin on the dissolution of lamotrigine. Increasing the soya lecithin content in this system from 5:1 to 1:1 showed a significant increase of 96.68±0.41% in the extent of dissolution at 75 min; also the initial rate of dissolution at 5 min was increased. However, the results were slightly decreased for the system containing lamotrigine and soya lecithin ratio 10:1. There were no significant differences in dissolution between 5:1 and 10:1 ratios of lamotrigine to soya lecithin. This illustrates that a small amount of the carrier phospholipid was adequate for a significant increase in the rate and extent of dissolution. An increase in dissolution rate was observed for the solid dispersions as compared to physical mixture and lamotrigine. This might be because of the decrease in particle size of the crystallites, resulting in greater wetting and increase in surface area of particles or because the crystallites are coated with the phospholipid or both. When placed into an aqueous medium, phospholipids rapidly form liposomal structures on dispersion resulting in an effective increase in the
saturation concentration of drugs in the diffusion layer during the dissolution process through the release of substantially entrapped finely dispersed drug from the lipid bilayer or the aqueous compartments. This may be another reason for the increased dissolution of lamotrigine from the lamotrigine-soya lecithin solid dispersion.

Characterization of lamotrigine phospholipid solid dispersions

Differential scanning calorimeter (DSC)

DSC thermogram of lamotrigine, lamotrigine-soya lecithin physical mixture and solid dispersion of lamotrigine-soya lecithin prepared by using ethanol are depicted in fig. 2. The studies revealed the formation of lamotrigine solid dispersion with and broaden endothermic peak at 206.69 °C by the solvent method. DSC thermograph of pure lamotrigine exhibited a sharp endothermic peak at 217.48 °C suggests the melting temperature of lamotrigine which has shifted backward and becoming much broader, suggests the formation of crystals of solid dispersion with soya lecithin.

X-ray diffraction studies (XRD)

XRD analysis of lamotrigine, lamotrigine-soya lecithin physical mixture and solid dispersion of lamotrigine-soya lecithin prepared by using ethanol are depicted in fig. 3. The X-ray diffractogram of lamotrigine shows a sharp and intense peak indicating the crystalline nature of the drug in pure form. The X-ray diffractogram of the physical mixture and solid dispersion were obtained to determine if there is a loss or modification of the pure drug's crystal structure after it is formed into a solid phospholipid dispersion, and to determine if any new crystalline phases may have formed. Fig. 3 indicates that there is virtually no difference in crystallinity between
prepared solid dispersion and pure lamotrigine proposed that enhanced dissolution might be due to the increased surface area of the drug crystallites after the formation of the solid dispersions and not due to a change from a crystalline to an amorphous state [25].

Fig. 3: XRD spectral analysis of a) lamotrigine, b) lamotrigine soya lecithin physical mixture and c) Solid dispersion of lamotrigine using ethanol

FTIR analysis

Results of FTIR spectral analysis of lamotrigine, soya lecithin, physical mixture of lamotrigine-soya lecithin and solid dispersion of lamotrigine-soya lecithin are depicted in fig. 4. FTIR spectra of lamotrigine showed characteristic absorbance at 3448 cm⁻¹ specifying NH stretching of an amino group (Aromatic), 3209 cm⁻¹ shows aromaticity (aromatic CH stretching), 1620 cm⁻¹, 1486 cm⁻¹ indicates C=C ring stretching and 962 cm⁻¹ shows C-Cl stretching of halides. In FTIR spectra of physical mixture characteristics peak of both lamotrigine and soya lecithin were observed. It suggests no significant interaction between drug and excipient. However, the characteristic absorbance peaks of lamotrigine were found to be shifted to lower values and decreased in intensity also, suggesting the formation of solid dispersion of lamotrigine with soya lecithin by using ethanol as solvent.

Thus, based on DSC, XRD, and FTIR analysis, the formation of crystallites of solid dispersion was affirmed.

Fig. 4: FTIR spectral analysis of a) lamotrigine, b) soya lecithin, c) lamotrigine soya lecithin physical mixture and d) Solid dispersion of lamotrigine using ethanol
Scanning electron microscopy (SEM)

Fig. 5 shows SEM images of the lamotrigine and solid dispersion of lamotrigine-soya lecithin. As observed from the fig. 5 particles of lamotrigine–soya lecithin solid dispersion showed the regular shape and smooth surface and significant reduction of the lamotrigine crystallite size as compared to pure lamotrigine. This allows faster solvating of the drug and the faster breakup of the solid dispersions. This could be because the crystallites are less than a micron in size. This may be the reason for the increased dissolution of lamotrigine from the lamotrigine-soya lecithin solid dispersion.

Fig. 5: SEM image of a) Lamotrigine and b) Solid dispersion of lamotrigine using ethanol

Pharmacodynamic study

As shown in fig. 6, administration of PTZ to vehicle-treated rat produced clonic convulsions in all animals, and the onset of such convulsions was 85.5±14.36 s. Oral administration of solid dispersion of lamotrigine (10 mg/kg) significantly delayed the onset of clonic-tonic seizures in PTZ injected animals up to 691.25 s and provided 100% protection against mortality induced by PTZ. The results indicate the greater anticonvulsant effect of lamotrigine-soya lecithin solid dispersion in PTZ-induced seizures as compared to pure lamotrigine. The results of the study are also supported by our earlier observations demonstrating the anticonvulsant effect of lamotrigine in PTZ induced clonic tonic seizures [28, 29].

Fig. 6: Effect of lamotrigine solid dispersion on the mean onset time of convulsions against pentylenetetrazole-induced seizures. 
***P<0.001 as compared to control group; #P<0.001 as compared to the control group. Data are represented as the mean onset time of convulsions±SEM. Statistical analysis by ANOVA followed by Bonferroni’s multiple comparison test (n = 4)

Characterization of lamotrigine orodispersible tablets

Precompression studies

Solid dispersion of batch S7 consisting of 1:5 ratio of soya lecithin-lamotrigine was obtained as discrete, uniform and non-greasy particles showed maximum drug content, better dissolution profile as compared to other batches and significant anticonvulsant activity so it was selected for the formulation of orodispersible tablets. The tablet blend was prepared by using various super disintegrants such as SSG, CCS, and CPD. The powder blends were evaluated for various
pre-compression parameters of powder blend and results are depicted in table 4. The bulk density found to be in range 0.523±0.003 to 0.558±0.002 g/cm², which was within acceptable limits. The angle of repose of all the formulations was found to be less than 25° indicating excellent flowability of the powder blend. Another important parameter to determine powder compressibility and flowability is the Hausner's ratio. A ratio greater than 1.5 indicates cohesive and poor flow property of powder blend. Results revealed the values less than 1.5 indicates good flow and compression property of the powder blends. Results of Carr’s index also indicate acceptable flow property and compression behavior powder blend to be compacted as tablets.

| Formulations | Bulk density (g/ml) | Tapped density (g/ml) | Carr’s index (%) | Hausner's ratio | Angle of repose (0) |
|--------------|---------------------|----------------------|------------------|----------------|--------------------|
| F1           | 0.54±0.001          | 0.76±0.003           | 28.38±1.23       | 1.396±0.025    | 19.28±0.65         |
| F2           | 0.55±0.003          | 0.727±0.004          | 26.77±0.98       | 1.302±0.031    | 19.89±0.88         |
| F3           | 0.55±0.004          | 0.738±0.002          | 25.33±0.56       | 1.339±0.011    | 20.12±0.76         |
| F4           | 0.523±0.002         | 0.691±0.003          | 21.10±0.57       | 1.321±0.014    | 20.33±0.65         |
| F5           | 0.56±0.01           | 0.716±0.006          | 23.22±0.87       | 1.269±0.023    | 19.66±0.63         |
| F6           | 0.534±0.005         | 0.696±0.004          | 22.27±1.05       | 1.303±0.046    | 19.35±1.03         |

Results are shown as mean±SD (n=3)

**Post dispersible studies**

Orodispersible tablets were evaluated for post-compression properties and results are depicted in table 5. For all formulations, results of weight variation were found to be within limit 100±1.44%. The hardness test indicated a good mechanical strength of 3.33±0.87 kg/cm². The friability was obtained to be less than 1% which indicates that all the batches have good mechanical resistance. The tablets were evaluated for the in vitro disintegration time, results revealed less disintegration time 22.59 s for the tablets consisting of CCS. The disintegration time was more with SSG which might be due to further penetration of the disintegration medium and hindered the disintegration of tablet content. Thus the tablet disintegration is retarded to some extent with the tablet containing SSG when compared with the disintegration time of the tablet containing CCS. The release profile for a formulation containing various super disintegrant and magnesium stearate as a lubricant is depicted in fig. 7. The drug content of the formulation was ranging from 92.31% to 95.27 % indicating maximum entrapment of drug in the formulation. F1 formulation gives 89.08 % at 15 min and F3, F4 also shows significant release 96.34 %, 100.06 % respectively. These results revealed that maximum drug release was observed when lamotrigine was combined with soya lecithin in a 1:5 ratio for solid dispersion prepared by the solvent method using ethanol as solvent and CCS as a super disintegrating agent.

| Code | Weight variation (mg) | Hardness* (kg/cm²) | Friability (%) | Thickness* (mm) | Disintegration* (s) | Drug content* (%) |
|------|-----------------------|-------------------|----------------|-----------------|--------------------|------------------|
| F1   | 100±1.0               | 3.50±0.47         | 0.60±0.05      | 3.11±0.14       | 26.11±0.41         | 82.31±0.77       |
| F2   | 100±1.12              | 3.40±0.18         | 0.70±0.04      | 3.09±0.23       | 24.45±0.52         | 91.04±0.68       |
| F3   | 100±1.43              | 3.30±0.43         | 0.60±0.03      | 3.18±0.19       | 23.33±0.63         | 89.39±1.75       |
| F4   | 100±1.44              | 3.33±0.87         | 0.50±0.02      | 3.13±0.17       | 22.59±0.55         | 95.27±0.57       |
| F5   | 100±1.83              | 3.29±0.50         | 0.90±0.01      | 3.12±0.05       | 25.15±0.47         | 86.02±1.95       |
| F6   | 100±1.13              | 3.5±0.83          | 0.80±0.03      | 3.16±0.11       | 27.32±0.12         | 90.97±0.90       |

Results are shown as mean±SD (* n=3, $ n= 20, # n=6)

**Fig. 7: Dissolution profile of various formulation of lamotrigine solid dispersion prepared by using ethanol**

**CONCLUSION**

In this study, we have prepared the orally disintegrating tablet consisting of solid dispersion of lamotrigine with soya lecithin as phospholipid. Solid dispersion of soya lecithin-lamotrigine using ethanol as a solvent showed a significant increase in solubility over and above the increase in rate and extent of dissolution, which may increase the bioavailability of lamotrigine. The improved dissolution behavior of drug-phospholipid solid dispersions is expected due to the significant increase in the drug’s total surface area during the dissolution process. Solid dispersion of lamotrigine with soya lecithin also showed significant antiepileptic activity as compared to pure lamotrigine. Also, optimized solid dispersion on oral administration delayed the onset of convulsions and offered complete protection against the PTZ induced seizures compared to pure lamotrigine. ODT consisting of solid dispersion exhibited faster and greater release of lamotrigine. Thus the formulation of lamotrigine solid dispersion as ODT offers promising advantages over conventional dosage with its immediate onset of action.

**AUTHORS CONTRIBUTIONS**

All the authors have contributed equally.

**CONFLICT OF INTERESTS**

The authors report no conflict of interest.

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