INTRODUCTION

Oral delivery of drugs is by far the most preferable route of drug delivery due to the ease of administration with more patient compliance. These systems have progressed from immediate release to site specific delivery over a period.[1,2] An ideal drug delivery system should have two main properties that are, containing a single dose or less frequent dosing for the whole duration of treatment and the dosage form must release active drug directly at the site of action.[3-5] The recent developments of floating and bioadhesive drug delivery systems (DDS) considering the role physiological environment and formulation variables affecting gastric retention, approaches to design single-unit and multiple-unit floating systems. Among the methods described, floating drug delivery and bioadhesive drug delivery systems (DDS) are promising systems in gastro retention with few limitations, which has a great impact on the drug delivery to its intended site of administration.[6-11]

The major disadvantage of the floating system is a requirement of a sufficiently high level of fluids in the stomach for the system to float. The floating DDS are effective only when the fluid level in the stomach is sufficiently high. Nonetheless, as the stomach empties dosage form reaches to the pylorus, the buoyancy of...
the dosage form may be impeded. Thus, bioadhesive DDS are suffering from the effect of mucous turnover. The mucous secreted by the mucosa lining of stomach wall may detach the dosage form from the wall of the stomach which get emptied from the stomach along with its contents. This limitation can be overcome by making the floating system eventually adhere to the mucous lining of the stomach wall.\textsuperscript{[22]}

Thus, FBDDS offers the advantage of increased gastric residence time of drugs over normal floating DDS. The FBDDS can be formulated by incorporating bioadhesive polymers to normal floating DDS.

Floating bioadhesive systems can persist in the stomach for several hours and hence considerably extend the gastric residence time of therapeutics. Due to the extended gastric retention the delivery system enhances bioavailability. It has applications also for delivery of drug to the upper gastric tract. Floating and bioadhesive delivery scaffold system helps to provide better availability of new products with new therapeutic possibilities and substantial benefits for patients.\textsuperscript{[13-15]}

Several classes of medications collectively referred to as antihypertensive drugs for treating hypertension. Therapeutic agents within a particular class generally share similar pharmacologic mechanisms of action and in many cases have an affinity for similar cellular receptors. Atenolol is a beta-adrenergic blocking agent that blocks the effects of adrenergic drugs.\textsuperscript{[16]}

Moreover, the site of absorption of atenolol is in the stomach. Dosage forms that are retained in the stomach would increase the absorption, improve drug efficiency, and decrease dose requirements. Due to its high permeability in nature controlled drug delivery is required for prolonged gastric retention may offer numerous advantages including an increase in the extent of absorption, improved bioavailability, and therapeutic efficacy.\textsuperscript{[9]} The objective of the present study is to develop a floating-bioadhesive drug delivery system exhibiting a unique combination of floatation and bioadhesion to prolong residence in the stomach using atenolol as a model drug.

### MATERIALS AND METHODS

Atenolol as a gift sample from Vapi Care Pharma Limited, Vapi, Gujarat; Hydroxypropyl methylcellulose K100M, hydroxypropyl methyl cellulose K15M, hydroxypropyl methyl cellulose K4M from Burgeon Pharmaceuticals, Chennai; Carbopol 934P, PVP K-30 from Triveni Chemicals, Vapi, Gujarat; Sodium bicarbonate, Magnesium stearate, talc from SD Fine Chemicals, Mumbai; Citric acid, Spray dried Lactose from Kawarlal and Co., Mumbai.

#### Preparation of floating bioadhesive tablets containing atenolol

Floating bioadhesive tablets of atenolol were prepared by employing various polymers like Carbopol 934P, HPMC K4M, HPMC K15M, and HPMC K100M in combination by direct compression method using compression machine. For the preparation of floating bioadhesive tablets, all components were screened through sieve number 60 and mixed thoroughly in a mortar and pestle for 10 min. Magnesium stearate and talc were added to the above blend as flow promoters.

In all the formulations, the amount of atenolol was kept constant at 50 mg. The polymers like Carbopol 934p, HPMC K4M, HPMC K15M, and HPMC K100M were used in different concentrations in combination. Total weight of the tablet was kept constant at 350 mg. The formulae of different floating bioadhesive tablets of atenolol are given in Table 1.

#### Preformulation studies of the optimized blends

The aforementioned polymeric blends were tested for the angle of repose, Hausner’s ratio, Carr’s index (% compressibility).

#### Evaluation of physical parameters

**Hardness and friability**

The hardness of prepared tablets was determined by using Monsanto hardness tester and measured in terms of kg/cm\(^2\). The friability test was performed to assess the effect of friction and shocks, which may often cause the tablet to chip, cap or break. Roche friabilator was used for testing the friability of prepared floating bioadhesive tablets. Ten tablets were accurately weighed and placed in the friabilator and operated for 100 revolutions. The tablets were de-dusted and reweighed and friability (F) was calculated.

#### Weight variation

The weight variation test was done by weighing 20 tablets individually, calculating the average weight and comparing the individual tablet weights to the average. The percentage difference

### Table 1: Composition of floating bioadhesive tablets of atenolol

| Ingredients (mg) | F1  | F2  | F3  | F4  | F5  | F6  | F7  | F8  | F9  | F10 | F11 | F12 | F13 | F14 | F15 |
|------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| Atenolol         | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  | 50  |
| HPMC K4M        | 70  | 60  | 50  | 40  | 30  | 60  | 60  | 60  | 60  | 60  | 60  | 60  | 60  | 60  | 60  |
| HPMC K15M       | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- | ----- |
| HPMC K100M      | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  | 30  |
| Carbopol 934P   | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 |
| Sodium bicarbonate | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  | 35  |
| PVP K 30        | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 | 17.5 |
| Magnesium stearate | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    | 7    |
| Talc             | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    | 3    |
| Spray dried lactose | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  | 120  |
in the weight variation should be within the permissible limits (7.5%). The percent deviation was calculated.

**Drug content**

Six tablets from each batch were weighed and powdered. Powder equivalent to the average weight of the tablet was accurately weighed and transferred into a 100 ml volumetric flask and dissolved in a suitable quantity of phosphate buffered saline buffer (100 ml). A portion of the sample was filtered and analyzed by an ultraviolet (UV) spectrophotometer at 225 nm.

**Buoyancy/floating test**

The *in vitro* buoyancy was determined by floating lag time. The tablets were placed in a 100 ml beaker containing 0.1 N HCl. The time required for the tablet to rise to the surface and float was determined as floating lag time. The duration of time the dosage form remained on the surface of the medium was determined as the total floating time.

**In vitro bioadhesive strength**

The test methods for determining mucoadhesion can be classified into two major categories: *in vitro*/*ex vivo* methods and *in vivo* methods. The most common methods are based on the measurement of either tensile or shear stress. In this study, an instrument was designed to evaluate the tensile force. This instrument consists of a modified physical balance. This method was used for determination of the *in vitro* bioadhesive strength. The balance was modified by replacement of one pan with the metal shaft 5 g heavier in weight than the pan. Fresh sheep mucosa obtained from local slaughterhouse was cut into pieces, washed with distilled water followed by 0.1 N HCl. A piece of the mucosa was fixed in a petri dish with instant adhesive. The two sides of the balance were made equal before the study, by keeping 5 g weight on the right hand pan. A weight of 5 g was removed from the right hand pan, which lowered the shaft along with the tablet over the mucosa. The balance was kept in this position for 3 min contact time. The weight was added slowly to the right hand pan until the tablet detached from the mucosal surface. This detachment force gave the bioadhesion strength of the bioadhesive tablet in gram (total weight on right hand pan minus 5 g).

**Swelling studies**

The extent of swelling was measured in terms of percentage weight gained by the tablet. One tablet from each formulation was weighed and kept in petri dish containing 15 ml of 0.1 N HCl. At the end of specified time intervals tablets were withdrawn from petri dish and excess buffer blotted with tissue paper and weighed. The percentage of weight gained by the tablet was calculated.

**In vitro drug release studies**

The dissolution of the floating bioadhesive tablet was performed using USP type II XXIII dissolution apparatus (paddle method) using 900 ml of 0.1 N HCl with pH 1.2 as the dissolution medium to mimic stomach, which was maintained at 37°C and stirred at 50 rpm. Aliquots of 5 ml of samples were withdrawn with a bulb pipette at different time intervals and replaced with equal volume of 0.1 N HCl at each withdrawal, filtered through Whatman filter paper number 1. The samples were then analyzed using UV spectrophotometer at 225 nm and the cumulative amount of drug released at various time intervals was calculated.

**Kinetic studies**

To analyze the release mechanism, several release models were tested such as:

- **Zero order**: \[ Q_t = Q_o + K_o t \] (1)
  
  Where \( Q_t \) is the amount of drug released at time \( t \), \( K_o \) is the apparent dissolution rate constant or zero order release constant and \( Q_o \) is the initial concentration of the drug in the solution resulting from a burst effect; in this case the drug release runs as a constant rate.

- **First order**: \[ \ln Q_t = \ln Q_o + K_1 t \] (2)
  
  Where \( K_1 \) is the first order release constant; in this case, the drug released at each time is proportional to the residual drug inside the dosage form.

- **Higuchi**: \[ Q_t = K_{H} \sqrt{t} \] (3)
  
  Where \( Q_t \) is the amount of drug released at time \( t \) and \( K_{H} \) is the Higuchi release rate constant; this is, the most widely used model to describe drug release from pharmaceutical matrices.

- **Korsmeyer–Peppas**: \[ Q_t/Q_o = K_p t^n \] (4)
  
  Where \( K_p \) is a constant incorporating structural and geometric characteristic of the drug dosage form and \( n \) is the release exponent, indicative of the drug release mechanism.

The value of \( n \) for a tablet, \( n = 0.45 \) for Fickian (Case I) release, \( >0.45 \) but \( <0.89 \) for non-Fickian (anomalous) release and 0.89 for Case II (zero order) release and \( >0.89 \) for super case II type of release.

Case II transport generally refers to the dissolution of the polymeric matrix due to the relaxation of the polymer chain and anomalous transport (non-Fickian) refers to the summation of both diffusion and dissolution controlled release.

**RESULTS AND DISCUSSION**

**Precompressional parameters**

Precompression parameters of all formulations blend were conducted for angle of repose, bulk density, tapped density, compressibility index, and Hausner’s ratio. Table 2 shows all the precompressional parameters of the prepared blends.
Physical parameters of the prepared tablets

Table 3 shows postcompressional parameters of the prepared tablets. The hardness of the tablets was found to be 5.0 ± 0.36–5.9 ± 0.33 kg/cm² and friability was found to be below 1% indicating good mechanical resistance. The thickness of the tablets was found to be 3.10 ± 0.04–3.20 ± 0.05. All the tablets passed weight variation test, as percentage weight variation was within the pharmacopeial limits that is, ±7.5%, mean percentage of drug content was found to be 98.5%.

Buoyancy lag time studies

All tablet formulations exhibited satisfactory flotation ability and remained buoyant for 10-24 h in the dissolution medium. The buoyancy lag time of tablets depends on the amount of sodium bicarbonate and citric acid involved in CO₂ formation and the concentration of polymers used. It was clearly observed that the reduction in the concentration of HPMC in each batch the floating lag time increased as well as floating duration decreased and also increase in the viscosity of HPMC polymers delayed the floating lag time and prolonged the drug release.

Swelling studies

The percentage water uptake of the formulations ranged from 70% to 142% [Table 4]. The formulation F15 shows maximum swelling index. It was observed that as we increase the concentration of Carbopol 934P and the viscosity of the hydrophilic polymers the water uptake capacity increased which results in an increase of swelling index. This may be because of the mobility of polymer chains was very dependent on the water content of the system.

In vitro bioadhesive study

In vitro bioadhesion evaluation test was conducted for all formulations, the result showed in Figure 1. The mean bioadhesive strength values were found in a range of 16.2-52.1 g for the floating bioadhesive tablets F1 to F15. There was a gradual increase in the bioadhesion strength in each batch that is, from F1 to F5, F6 to F10, and F11 to F15. This was due to the increase in the concentration of bioadhesive polymer Carbopol 934P. Maximum bioadhesion strength was found for formulations F11-F15 and low bioadhesion strength was found for formulations F1-F5, this may be expected that as the viscosity of the hydrophilic polymer increases and concentration of Carbopol 934P increases the adhesive property also increases.

In vitro dissolution studies

The dissolution rates of all floating bioadhesive tablets were studied by using USP type II apparatus (paddle type) in 0.1 N HCl. The release of atenolol from floating bioadhesive tablets depends on the type and concentration of polymer. Batches F1, F2, F3, F4, and F5 are composed with HPMC K4M as a hydrophilic polymer and Carbopol.
934P as bioadhesive polymer in increasing ratios of Carbopol 934P and decreasing ratios of HPMC K4M. The release profile depicted in Figures 2-4 shows that Carbopol 934P was helpful in retarding drug release. Batches from F6 to F10 were composed with HPMC K15M as hydrophilic polymer and Carbopol 934P as bioadhesive polymer in increasing ratios of Carbopol 934P and decreasing ratios of HPMC K15M, respectively. This might be due to high viscosity of polymer HPMC K15M and HPMC K4M.

Batches from F11 to F15 were composed with HPMC K100M as hydrophilic polymer and Carbopol 934P as bioadhesive polymer in increasing ratios of Carbopol 934P and decreasing ratios of HPMC K100M, respectively. This might be due to high viscosity of polymer HPMC K100M than HPMC K4M and HPMC K15M.

**Kinetic studies**

To investigate the mechanism of drug release from floating bioadhesive tablets, various kinetics models such as zero order, first order, Higuchi’s and Korsmeyer–Peppas equations were applied to the in vitro release data obtained from different formulations. The values of correlation-coefficient ($r^2$) for all the formulations were high enough to evaluate the drug dissolution behavior. The value of release exponent ($n$) was found to be a function of polymer used and the physicochemical property of a drug molecule itself. It was evident that the formulation F11 followed first order process as the correlation coefficient ($r^2$)

Table 4: Physical properties of floating bioadhesive tablets of atenolol ($n = 3$)

| Batch number | Percentage swelling index | Floating lag time (min) | Total floating time (h) |
|--------------|---------------------------|-------------------------|-------------------------|
| F1           | 70±2.3                    | 2.0±0.2                 | 15±1.0                  |
| F2           | 79±1.8                    | 2.4±0.4                 | 14±1.5                  |
| F3           | 91±2.6                    | 3.0±0.5                 | 13±2.6                  |
| F4           | 101±2.5                   | 3.5±1.0                 | 12±1.5                  |
| F5           | 112±2.8                   | 4.0±1.2                 | 10±1.0                  |
| F6           | 76±3.5                    | 2.2±0.5                 | 24±3.0                  |
| F7           | 89±4.4                    | 3.0±0.2                 | 24±2.5                  |
| F8           | 96±3.0                    | 3.6±0.8                 | 23±1.5                  |
| F9           | 120±2.5                   | 4.2±1.2                 | 23±1.0                  |
| F10          | 137±1.5                   | 5.0±1.5                 | 21±2.0                  |
| F11          | 80±2.0                    | 2.9±0.4                 | 24±4.5                  |
| F12          | 87±2.5                    | 3.4±0.6                 | 24±4.0                  |
| F13          | 104±1.0                   | 4.3±0.4                 | >24                     |
| F14          | 134±3.5                   | 4.8±1.0                 | >24                     |
| F15          | 142±2.6                   | 5.2±0.5                 | >24                     |
value was 0.988, respectively. This indicated that the dissolution rate of the drug was dependent on the concentration of dissolving species. Further, when the drug release data was put into Higuchi equation, good correlation coefficient ($r^2$) values 0.963-0.992 were obtained, indicating that the drug release was diffusion controlled [Figures 5-8].

The release data obtained were also put in Korsmeyer–Peppas model to find out $n$ values, which describe the drug release mechanism; good correlation coefficient ($r^2$) values 0.97-0.997 were obtained. The $n$ values were in the range of 0.623-0.719 indicating non-Fickain anomalous type transport mechanism [Tables 5 and 6].

### CONCLUSION

In this current work, an attempt was made to design floating bioadhesive drug delivery system of an antihypertensive drug. Floating-bioadhesive tablets of atenolol that exhibit a unique combination of floatation and bioadhesion for prolonged residence in the stomach were prepared by direct compression technique using polymers such as HPMC K4M, HPMC K15M, HPMC K100M, and Carbopol 934P. From aforesaid results, it can be concluded that among all the formulations the formulations with HPMC K100M with Carbopol 934P showed controlled release. The F11 formulation showed a satisfactory dissolution profile, detachment stress and floating characteristics, which can increase the gastric residence time as well as bioavailability and better patient compliance.

### Financial support and sponsorship
Nil.

### Conflicts of interest
There are no conflicts of interest.

| Time (h) | $\sqrt{t}$ | Log t | Cumulative release* Percentage | Log percentage | Cumulative retained Percentage | Log percentage |
|----------|------------|-------|--------------------------------|----------------|-------------------------------|----------------|
| 0        | 0          | 0     | 0                              | 0              | 100                           | 2.000          |
| 0.5      | 0.707      | 0.301 | 10.62                          | 1.026          | 89.38                         | 1.951          |
| 1        | 1.000      | 0.000 | 14.65                          | 1.166          | 85.35                         | 1.931          |
| 2        | 1.414      | 0.301 | 19.98                          | 1.301          | 80.02                         | 1.903          |
| 3        | 1.732      | 0.477 | 23.04                          | 1.362          | 76.96                         | 1.886          |
| 4        | 2.000      | 0.602 | 29.7                           | 1.473          | 70.3                          | 1.847          |
| 5        | 2.236      | 0.699 | 36.54                          | 1.563          | 63.46                         | 1.803          |
| 6        | 2.449      | 0.778 | 42.46                          | 1.628          | 57.54                         | 1.780          |
| 7        | 2.646      | 0.845 | 49.37                          | 1.693          | 50.63                         | 1.704          |
| 8        | 2.828      | 0.903 | 54.21                          | 1.734          | 45.79                         | 1.661          |
| 9        | 3.000      | 0.954 | 60.72                          | 1.783          | 39.28                         | 1.594          |
| 10       | 3.162      | 1.000 | 68.4                           | 1.835          | 31.6                          | 1.500          |
| 11       | 3.317      | 1.041 | 71.3                           | 1.853          | 28.7                          | 1.458          |
| 12       | 3.464      | 1.079 | 76.3                           | 1.883          | 23.7                          | 1.375          |
| 13       | 3.742      | 1.146 | 80.3                           | 1.905          | 19.7                          | 1.294          |
| 14       | 4.000      | 1.204 | 84.6                           | 1.927          | 15.4                          | 1.188          |
| 15       | 4.243      | 1.255 | 88.9                           | 1.949          | 11.1                          | 1.045          |
| 20       | 4.472      | 1.301 | 90.3                           | 1.956          | 9.7                           | 0.987          |
| 24       | 4.899      | 1.380 | 92.3                           | 1.965          | 7.7                           | 0.886          |
REFERENCES

1. Rao YM, Vani G, Chary RB. Bioavailability studies of paracetamol bioadhesive tablets. Indian J Pharm Sci 1999;38:379-82.
2. Amidon GL, Lennernäs H, Shah VP, Crison JR. A theoretical basis for a biopharmaceutic drug classification: The correlation of in vitro drug product dissolution and in vivo bioavailability. Pharm Res 1995;12:413-20.
3. Antonios GM, Nicolas AP. Bioadhesive analysis of controlled release system: An experimental method for testing the adhesion of micro particulate with mucus. J Control Release 1990;12:31-7.
4. Cheueh HR, Zia H, Rhodes CT. Optimization of sotalol floating and bioadhesive extended release tablet formulations. Drug Deliv Indian Pharm 1996;21:1725-47.
5. Baugmgartner S, Kristl J, Vrecer F, Vodopivee P, Zorko B. Optimization of floating matrix tablets and evaluation of their gastric residence time. Int J Pharm 2004;195:125-135.
6. Ranga RK, Siva P, Sajeeth CI. Formulation and evaluation of gastroretentive floating bioadhesive tablets of glipizide. Int J Res Pharm 2011;2:252-60.
7. Rajput GC, Majumdar FD, Patel JK, Patel KN, Thakor RS, Patel BP, et al. Stomach specific mucoadhesive tablets as controlled drug delivery system – A review work. Int J Pharm Biol Res 2010;1:30-41.
8. Zawar LR, Savaliya, PJ, Bari SB, Gottani SG. Formulation & evaluation of floating-mucoadhesive tablets of clarithromycin. Int J Pharm Biosci 2010;1:1-10.
9. Löbenberg R, Kim JS, Amidon GL. Pharmacokinetics of an immediate release, a controlled release and a two pulse dosage form in dogs. Eur J Pharm Biopharm 2005;60:17-23.
10. Chavanpatil MD, Jain P, Chaudhari S, Shear R, Vavia PR. Novel sustained release, swellable and bioadhesive gastroretentive drug delivery system for ofloxacin. Int J Pharm 2006;316:86-92.
11. Patel FV, Patel MN, Yeole GP. Studies on formulation and evaluation of ranitidine floating tablets. Indian J Pharm Sci 2005;67:703-9.
12. Reza MS, Quadir MA, Haider SS. Comparative evaluation of plastic, hydrophobic and hydrophilic polymers as matrices for controlled-release drug delivery. J Pharm Pharmacol Sci 2003; 6:282-91.
13. Rahman Z, Ali M, Khar R. Design and evaluation of bilayer floating tablets of captopril. Acta Pharm 2006;56:49-57.
14. Jain SK, Agrawal GP, Jain NK. A novel calcium silicate based microspheres of repaglinide: In vivo investigations. J Control Release 2006;113:111-6.
15. Hong SI, Oh SY. Dissolution kinetics and physical characterization of three-layered tablet with poly(ethylene oxide) core matrix capped by Carbopol. Int J Pharm 2008;366:121-9.
16. Srivastava AK, Wadhwa S, Ridhurkar D, Mishra B. Oral sustained delivery of atenolol from floating matrix tablets: formulation and in vitro evaluation. Drug Dev Ind Pharm 2005;31:367-74.

Table 6: Kinetic study of floating bioadhesive tablets of atenolol

| Formulation code | Zero order $r^2$ | First order $r^2$ | Higuchi $r^2$ | Korsmeyer–Peppas $n$ |
|------------------|-----------------|-----------------|-------------|-------------------|
| F1               | 0.924           | 0.978           | 0.988       | 0.623             |
| F2               | 0.955           | 0.950           | 0.986       | 0.997             |
| F3               | 0.954           | 0.969           | 0.992       | 0.964             |
| F4               | 0.982           | 0.937           | 0.971       | 0.996             |
| F5               | 0.979           | 0.959           | 0.971       | 0.988             |
| F6               | 0.974           | 0.887           | 0.979       | 0.996             |
| F7               | 0.961           | 0.917           | 0.980       | 0.996             |
| F8               | 0.920           | 0.982           | 0.979       | 0.985             |
| F9               | 0.898           | 0.983           | 0.976       | 0.983             |
| F10              | 0.877           | 0.971           | 0.968       | 0.977             |
| F11              | 0.909           | 0.988           | 0.970       | 0.978             |
| F12              | 0.898           | 0.983           | 0.965       | 0.976             |
| F13              | 0.909           | 0.99            | 0.971       | 0.976             |
| F14              | 0.912           | 0.985           | 0.966       | 0.973             |
| F15              | 0.915           | 0.983           | 0.963       | 0.97              |

Figure 7: Higuchi plot of optimized floating bioadhesive tablets of atenolol

Figure 8: Korsmeyer–Peppas plot of optimized floating bioadhesive tablets of atenolol