Dual drug delivery system for targeting H. pylori in the stomach: preparation and in vitro characterization of amoxicillin-loaded Carbopol® nanospheres

Sree Harsha
Department of Pharmaceutical Sciences, College of Clinical Pharmacy, King Faisal University, Al-Asha, Saudi Arabia

Correspondence: Sree Harsha
College of Clinical Pharmacy, King Faisal University, Al-Asha 31982, Saudi Arabia
Tel +96635817175
Fax +96635817174
Mobile +966535485322
Email harsha1975@gmail.com

Background and methods: A dual (immediate/sustained-release) oral amoxicillin suspension was developed as a new dosage form to eradicate Helicobacter pylori. Carbopol®-loaded amoxicillin nanospheres could bind with the mucosa after delivery to the stomach and could increase the efficiency of the drug, providing both an immediate and a sustained action.

Results: The objective of this research was to develop amoxicillin nanospheres using a spray-drying technique and to investigate such features as their particle size, drug content, percentage yield, surface morphology, in vitro release, and stability. The nanospheres had a particle size range of 280–320 nm after optimizing the preparation method using a central composite design. The drug content and percentage yield was 85.3% ± 0.7% and 92.8% ± 0.9%, respectively. The in vitro release profile of the amoxicillin nanospheres was consistent with a Korsmeyer-Peppas pattern, and the release after one hour was 19%, while for the original drug, amoxicillin, under the same conditions, 90% was released in the first 30 minutes.

Conclusion: The nanospheres used in this study enabled controlled release of amoxicillin over an extended period of time for up to 12 hours and the formulation was stable for 12 months.

Keywords: nanospheres, nano spraydryer, optimization, amoxicillin, Helicobacter pylori

Introduction
Helicobacter pylori is a bacterium commonly found in the stomach. Infection with H. pylori is universally accepted as the main threat for chronic gastritis and gastric carcinogenesis at present, and was classified as a carcinogen by the World Health Organization in 1994. H. pylori resides primarily in the gastrointestinal mucosa or at the interface between the mucus layer and the epithelial cells of the pyloric antrum region of the stomach.

Amoxicillin, a semisynthetic penicillin, is an effective antibiotic for H. pylori infection. It is stable in acid, inhibits synthesis of the bacterial cell wall, and has good sensitivity to H. pylori in vitro. These triple attributes contribute to the efficacy of amoxicillin in medical applications. Nevertheless, there are some reports indicating that treatment with amoxicillin cannot completely eliminate H. pylori, suggesting that additional research is needed to improve its therapeutic effects.

One of the reasons for only partial elimination of H. pylori using amoxicillin may be degradation of the drug in gastric acid and also a short drug residence time in the stomach, such that an effective drug concentration cannot be achieved in the gastric mucosal layer or epithelial cell surface where H. pylori resides.
Hence, administration of large doses of amoxicillin on a daily basis is necessary for eradication of *H. pylori*, and poor patient compliance due to adverse effects such as diarrhea, nausea, and vomiting is not unusual. Consequently, it is expected that if local delivery of antimicrobial agents from the gastric lumen into the mucus layer can be achieved, the *H. pylori* eradication rate would be increased.\textsuperscript{13}

Several formulations have been suggested to help retain amoxicillin in the stomach, including gellan beads,\textsuperscript{12} nanocapsules,\textsuperscript{13} hydrogel nanoparticles,\textsuperscript{14} floating microballoons,\textsuperscript{15} and mucoadhesive microspheres,\textsuperscript{16} which are able to remain in the gastrointestinal tract for a longer time, and therefore be more effective in *H. pylori* eradication.

Various methods can be used to prepare polymer nanospheres, such as micro-emulsion, solvent evaporation, supercritical fluid technology, salting-out, dialysis, surfactant-free emulsion, mini-emulsion, and interfacial polymerization.\textsuperscript{17} Spraydrying is a well recognized, rapid, one-step process frequently used in pharmaceutical engineering to produce a dry powder from a liquid phase.\textsuperscript{18}

In light of the above issues, there is a pressing need to develop a new formulation that delivers amoxicillin into the stomach to bind with the gastric mucosa, increase the efficiency of the drug, and provide a sustained action.

To date, no research has been published on the preparation of a suspension containing a dual-release formulation of amoxicillin. The purpose of this study was to design Carbopol\textsuperscript{18}-loaded amoxicillin nanospheres as a new dual-release suspension containing pure amoxicillin as well as a granular nanosphere form of amoxicillin using a Buchi Nanosprayer B-90 (Buchi Labortechnik AG, Flawil, Switzerland) for *H. pylori* eradication therapy.

### Materials and methods

Amoxicillin trihydrate was kindly gifted for this research by Karnataka Antibiotics and Pharmaceuticals Ltd, Bangalore, India. Carbopol-934P was purchased from Sigma-Aldrich (St Louis, MO). All other reagents and chemicals used were of analytical grade.

### Formulation of nanospheres by spraydrying

Carbopol-loaded amoxicillin nanospheres were prepared using a spraydrying method.\textsuperscript{19–21} The experiments were designed to develop an efficient spraydrying technique. Response surface methodology (specifically, a randomized rotatable central composite design) was used because it allows clarification of the relationship between cause and effect of variables used in the procedure and output. Nanospheres were prepared by varying parameters such as the concentration of Carbopol-934P (0.5–2.0 g), inlet temperature (80°C–120°C), and flow rate (120–180 mL/hour).

A solution of Carbopol-934P (Carbopol-934P soaked for one hour in 100 mL of water) containing a predetermined quantity of amoxicillin was sprayed using a spray nozzle through a 5.5 µm diameter. All samples were filtered prior to the spraydrying process to avoid blockage of the nozzle. Flow of heated air aspirated by a pump rapidly removed the solvent from the solution, leading to formation of solid nanospheres. The dried powder was removed from the particle collector using a particle scraper and stored in a desiccator at 25°C for further characterization.

### Surface morphology

The surface morphology of the nanospheres was observed using scanning electron microscopy (FEI Quanta 200, Eindhoven, The Netherlands). The nanospheres were placed in graphite casing and coated with gold using an ion sputter machine (JEOL, Tokyo, Japan) and detected at 12 kV.\textsuperscript{22,23}

### Particle size

Analysis of amoxicillin nanosphere particle size was done using laser light diffraction and ultrasonic techniques. The nanospheres were diluted 10-fold with deionized water using a Malvern Zetasizer Nano Series Nano-ZS (Malvern Instruments Inc, Westborough, MA).\textsuperscript{24}

### Drug content

A known quantity of amoxicillin nanosphere particles was dissolved in 0.1 M HCl. The amoxicillin content was assayed using spectrometry at 272 nm by constructing a calibration curve (UV-1601 spectrophotometer, Shimadzu, Tokyo, Japan). The best fitting straight line equation on the calibration curve was found to be $A = 0.294x + 0.0071$.\textsuperscript{25,26} The experiments were performed in triplicate. The drug content was calculated as the ratio between the actual and theoretical amount of amoxicillin loaded for an optimized formulation.

### Product recovery

The efficiency of nanosphere recovery was determined as the percentage weight of the nanospheres obtained compared with the entire amount of drug and polymer initially used in the preparation process.\textsuperscript{26}
In vitro release and kinetics

Amoxicillin release studies were carried out for both the nanospheres and dry powder suspensions using the paddle method specified in United State Pharmacopeia (USP) XXVII. The in vitro release studies were carried out using the USP I basket method (50 rpm, 500 mL, and 37°C ± 1°C). Amoxicillin-loaded nanosphere suspensions equivalent to 250 mg of drug were placed in a dialysis cellulose membrane bag (D9402-100FT, Sigma-Aldrich) with a molecular weight cutoff of 12,400, tangled, placed into the baskets, and transferred to 500 mL of 0.05 M phosphate buffer. Aliquots were withdrawn at prespecified time intervals and an equal volume of dissolution medium was replaced into the flask to maintain a constant volume. The aliquots drawn were analyzed at 272 nm, and the data were used to determine a cumulative drug release profile for the nanospheres. Sink conditions were retained throughout. Analysis of the suspension containing only amoxicillin was carried out in a similar manner.27

Formulation of dry mixtures for reconstitution

Xanthan gum was used as the suspending agent in the suspensions. D-sorbitol was used to impart a sweet taste. Sodium citrate and citric acid were used to vary the pH of the study formulations. To produce a dry mixture for reconstitution, the powder components were shrunk to approximately the same particle size. Ingredients present in small quantities (buffer components, sodium benzoate 0.2% and sodium lauryl sulfate 0.05%) were mixed homogeneously. The same procedure was followed for xanthan gum (0.6%) and amoxicillin-loaded nanospheres (containing the equivalent to amoxicillin 250 mg). The constituents were mixed with a portion of D-sorbitol (20%) according to the geometric dilution principle. To formulate the reconstituted suspension, 10 mL of water was added to the dry suspension powder in two stages and stirred with a spoon until a uniform product was attained.28

Stability

The powdered nanospheres were put into a bottle and stored for 12 months at 3°C–5°C, 15°C–25°C, and 37°C. Surface morphology and amoxicillin content were examined periodically.26

Results and discussion

Appearance of nanospheres

The amoxicillin nanospheres formed under most of the experimental conditions selected were smooth, almost spherical, and free flowing (Figure 1). Other investigators have reported that spraydried particles become shriveled, which may be the case if the polymer concentration is not optimal, and the shriveling may be due to rapid drying in the spraying funnel.29–31 A logical assumption is that when the solution is sprayed at the most favorable temperature and concentration in a spraying funnel, the smaller particles would dry out more rapidly and the critical point of crust formation would be reached soon after atomization.

Table 1 Design summary for DOE runs of amoxicillin microsphere construction with variable and response constraints

| Study type | Response surface | Experiments | 20 |
|------------|------------------|-------------|----|
| Initial design | Central composite | Blocks | No blocks |
| Design model | Quadratic | Quadratic |        |
| Response  | Name | Units | Runs | Minimum | Maximum | Trans | Model |
| Y1        | % yield | % | 20 | 64 | 92 | None | Quadratic |
| Factor    | Name | Type | Low actual | High actual | Low actual | High actual | Low coded | High coded |
| A         | Carbopol concentration | Numeric | 0.5 | 1 | 1 |
| B         | Inlet temperature | Numeric | 80 | 120 | 1 |
| C         | Feed flow rate | % | 15 | 25 | 1 |

Abbreviation: DOE, Design of Experiments.
However, for larger particles, it takes longer to shape a shell, resulting in shriveled nanospheres, and spherical particles would release the drug in a controlled fashion at the target site.

Optimization of pharmaceutical preparations with respect to one or more features is important in formulation studies. In a typical pharmaceutical formulation experiment, the focus is on determining which process variables have the greatest effect on the results. In pharmaceutical manufacturing, there may be numerous potentially significant parameters due to the large number of ingredients typically used in a pharmaceutical formulation. If all these variables were included in the experimental design, an excessive number of experiments would need to be performed. The formulation scientist must first identify the parameters most likely to be important in the formulation process in order to limit the number of experiments needed to a reasonable extent. A delicate equilibrium is established between what is worth investigating and what is not worth investigating. The formulator’s skill, knowledge, and experience play a crucial role in selecting the appropriate variables.

Response surface methodology is the most appropriate method for modeling and analysis of trials in which the response of interest is nonlinear and influenced by numerous variables. In this experiment, a full central composite design was used to develop the particle size model. Analysis using this mathematical model were carried out with Design-expert 6.0.5© Statease Inc (Minneapolis, MN). The fit and summary experimental specifies that the quadratic central composite design model was more significant than the linear model and that the linear model had a significant lack of fit. Therefore, the quadratic model was preferred in order to develop the central composite design.

A central composite design was generated to study the key effects and interactions of three factors, ie, Carbopol (g), inlet temperature (°C), and feed flow rate (mL/hour) on particle size (Table 1). Twenty preparations of nanospheres were formulated as per Table 2 to interpret the effect of these three independent variables. Experiments were conducted in accordance with our earlier study in random sequence in a face-centered manner in order to evaluate the interaction. The particle size data are shown in Table 2. The experimental results show that the particle

### Table 2 Investigational design by central composite design of amoxicillin nanospheres

| Run | Factor 1 | Factor 2 | Factor 3 | Response 1 |
|-----|----------|----------|----------|------------|
| A   | Carbopol concentration (%) | B inlet temperature (°C) | C feed flow rate (%) | Particle size (nm) |
| 1   | 0.75     | 100      | 20       | 250        |
| 2   | 0.75     | 100      | 20       | 250        |
| 3   | 0.75     | 100      | 20       | 250        |
| 4   | 1        | 100      | 20       | 404        |
| 5   | 0.75     | 120      | 25       | 280        |
| 6   | 1        | 80       | 25       | 310        |
| 7   | 0.75     | 80       | 20       | 210        |
| 8   | 0.75     | 100      | 20       | 250        |
| 9   | 0.75     | 100      | 15       | 270        |
| 10  | 1        | 120      | 25       | 350        |
| 11  | 0.75     | 100      | 20       | 250        |
| 12  | 0.5      | 100      | 20       | 225        |
| 13  | 1        | 80       | 15       | 290        |
| 14  | 0.5      | 80       | 25       | 270        |
| 15  | 1        | 120      | 15       | 330        |
| 16  | 0.5      | 120      | 25       | 220        |
| 17  | 0.5      | 80       | 15       | 250        |
| 18  | 0.75     | 100      | 20       | 250        |
| 19  | 0.5      | 120      | 15       | 200        |
| 20  | 0.75     | 80       | 20       | 388        |

### Table 3 Numerical optimization solutions for amoxicillin nanospheres

| Number | Carbopol concentration | Inlet temperature | Feed flow rate | Particle size |
|--------|------------------------|-------------------|----------------|--------------|
| 1      | 1.86                   | 101.48            | 17.77          | 309          |
| 2      | 1.12                   | 80.24             | 21.49          | 287          |
| 3      | 1.78                   | 99.71             | 24.28          | 300          |
| 4      | 1.85                   | 106.42            | 19.68          | 314          |
| 5      | 1.24                   | 81.10             | 21.99          | 284          |
| 6      | 1.88                   | 104.05            | 19.58          | 318          |
| 7      | 1.71                   | 96.10             | 18.16          | 289          |
| 8      | 1.82                   | 93.80             | 20.66          | 312          |
| 9      | 1.79                   | 105.78            | 22.18          | 303          |
| 10     | 1.83                   | 106.59            | 16.93          | 299          |
size was between 200 nm and 404 nm, and was not affected by transformations, and this trailed a quadratic arrangement indicating that the response measured clutches good over a range of other factors, ie, Carbopol concentration, inlet temperature, and feed flow rate. This information would, in turn, help in setting the optimal particle size. The polynomial equation giving the final equation in terms of actual factors is:

\[
\text{Particle size} = +182.15 + 207.6 * \text{Carbopol concentration} -0.99 * \text{Inlet temperature} + 1.8 * \text{Feed flow rate}
\]

**Process optimization**

The objective of this study was to ensure a narrow particle size range of 280–320 nm, so results obtained between 200 nm and 404 nm were considered to be unacceptable. Another goal was to identify any constraints on design space and the vulnerability of the investigational model. This is important because it indicates the factors, responses, and the goal for each variable with respect to the measured response. In this study, 10 optimum solutions were created in quantitative terms (Table 3), ie, the probable behavior of the measured response in terms of analyzed factors in a random manner within the design space. The optimized methods were selected based on the numerical optimization data (Table 4) for which the predicted results were closest to the observed results.

From the numerical optimization solutions, three formulatons (ACN-2, ACN-5, ACN-8) were randomly selected and further evaluated for particle size (Table 4), and the observed result was very similar to the response predicted by the expert design software.

As shown in Table 5, \( P \leq 0.05 \) for any factor in analysis of variance indicates an important effect of a variable on particle size. From the F ratios given in analysis of variance, it can be concluded that inlet temperature and Carbopol concentration had a significant effect on particle size.

Analysis of variance for the response surface model revealed that the concentration of the polymer is the most important parameter for particle size (Figure 2). In general, a higher concentration of the polymer led to larger particle size, which is in agreement with earlier reports. The relationship between the dependent and independent variables was further elucidated using response surface plots. However, the three-dimensional plot (Figure 3) shows that increasing the inlet temperature led to a decreased particle size. The correlation coefficient observed in the current investigation was found to be low (15.11). From Figures 3 and 4 it can be seen that lower and higher levels of feed flow rate did not vary to a noticeable extent. The drug content and percentage yield of the optimized formulations were found to be 85.3% ± 0.7% and 92.8% ± 0.9%, respectively. It seems logical that loss of product occurred mainly because of deposition of residual nanoparticles in the spraying compartment or by loss during manual removal from the collecting electrode.

**Size distribution of nanoparticles**

The size distribution of nanoparticles prepared by the spraydrying method showed a narrow distribution pattern with a size range of 200–404 nm (Figure 5). Particle size uniformity is very important because the target area of the nanoparticle containing the drug depends on the size of the particle, and bioavailability and plasma drug levels will fluctuate if the size deviation is wide. It has been reported elsewhere that nanoparticles prepared by spraydrying have advantages over those prepared by mechanical stirring or ultrasonification procedures. Also, the particle size distribution range is broad when nanoparticles are prepared by methods other than spraydrying.

### Table 4 Optimized formulae and formulation codes for amoxicillin nanospheres

| Formulation code | Carbopol concentration | Inlet temperature | Feed flow rate | Predicted particle size nm | Experimental particle size nm |
|------------------|------------------------|-------------------|----------------|---------------------------|-------------------------------|
| ACN-2            | 1.12                   | 80                | 21             | 287                       | 292                           |
| ACN-5            | 1.24                   | 81                | 21             | 284                       | 280                           |
| ACN-8            | 1.82                   | 94                | 21             | 312                       | 318                           |

### Table 5 Summary of analysis of variance results for quadratic model of particle size for amoxicillin nanospheres

| Source | Sum of squares | DF | Mean square | F value | Prob > F |
|--------|----------------|----|-------------|---------|----------|
| Model  | 905.9          | 9  | 100.7       | 6.2     | 0.0044   |
| A      | 421.9          | 1  | 421.9       | 25.9    | 0.0005   |
| B      | 336.7          | 1  | 336.7       | 20.7    | 0.0011   |
| C      | 3.5            | 1  | 3.5         | 0.2     | 0.6523   |
| A2     | 0.4            | 1  | 0.4         | 0.0     | 0.8725   |
| B2     | 10.1           | 1  | 10.1        | 0.6     | 0.4485   |
| C2     | 71.5           | 1  | 71.5        | 4.4     | 0.0627   |
| AB     | 44.9           | 1  | 44.9        | 2.8     | 0.128    |
| AC     | 1.2            | 1  | 1.2         | 0.1     | 0.7948   |
| BC     | 14.9           | 1  | 14.9        | 0.9     | 0.3611   |
Design-expert plot

Particle size
X = A: Carbopol conc
Y = B: inlet temperature

Actual factor
C: feed flow rate = 20.0

![Design-expert plot showing the effect of feed flow rate and inlet temperature, and their mutual interaction on particle size.](image)

**Figure 2** Three-dimensional plot showing the effect of inlet temperature and Carbopol concentration, and their mutual interaction on particle size.

Design-expert plot

Particle size
X = B: inlet temperature
Y = C: feed flow rate

Actual factor
A: Carbopol conc = 1.00

![Design-expert plot showing the effect of feed flow rate and inlet temperature, and their mutual interaction on particle size.](image)

**Figure 3** Three-dimensional plot showing the effect of feed flow rate and inlet temperature, and their mutual interaction on particle size.
Spraydrying technology is very attractive due to the one-step procedure involved in preparation of drug-loaded nanoparticles and achieving a uniform particle size.\textsuperscript{34,35} A narrow particle size range could be useful for amoxicillin nanospheres, which would adhere when given orally to the mucosal membrane, providing a high concentration gradient of the drug towards the absorption membrane for \textit{H. pylori} eradication.

**In vitro drug release pattern**

Figure 6 shows in vitro drug release from the amoxicillin nanospheres and from amoxicillin suspension. Approximately 18% of the total amount of amoxicillin in the nanospheres was released in the first hour, reflecting the significant amount of amoxicillin adsorbed onto or incorporated near the surface of the nanospheres. In clinical practice, this would lead to a “burst effect”, enabling the preparation to have a rapid onset
of action in patients. However, complete drug release from the amoxicillin nanospheres in this study occurred in 12 hours. During this time, the amount released was 97.3%. In comparison with amoxicillin nanospheres, the amoxicillin suspension released amoxicillin very rapidly, with 90% of the amoxicillin released in 30 minutes. The results indicate that amoxicillin nanospheres had well controlled release efficacy.

The data obtained from our in vitro release studies were fitted to various kinetic equations (ie, first-order, Baker and Lonsdale, Hixson and Crowell, Korsmeyer-Peppas, and Higuchi) to determine the mechanism of drug release and the release rate, using a macro written for SigmaPlot version 9.01 (Figure 7). The correlation coefficient value, $R^2$, was taken into account to decide upon the relevance of the model/curve fit which best described the extent of fit. According to the $R^2$ values given by the different data fits for amoxicillin nanospheres, the Korsmeyer-Peppas model was identified as having the ideal fit ($R^2 = 0.9968$). According to the Korsmeyer-Peppas fit, drug release is determined by diffusion of the polymeric matrix and is governed by Fick’s law. The release pattern showed classical Fickian diffusion, aided by initial swelling of the nanospheres and release of drug particles adsorbed on the surface, leading to a “burst” release. Later on, the release pattern was more controlled and was sustained for a longer period of time due to diffusion.

Reconstitutable amoxicillin nanosphere suspension

Because of the risk of medication flooding to the suspending medium during storage, the suspension formulations were stored as dry suspensions for reconstitution before use. Xanthan gum was chosen as the suspending agent and stabilizer in the dispersing medium due to its safety and unobjectionable toxicological properties for food and pharmaceutical applications. It is water-soluble and imparts high viscosity at a small concentration with thixotropic flow characteristics. Amoxicillin showed no notable
changes in external morphology or drug content during storage at 3°C–5°C or at room temperature (15°C–25°C) for 12 months (Figure 8). However, characteristic clumping was observed at 37°C and a relative humidity of 75%.

**Conclusion**

The major conclusion of this work is that the dual-release suspension is a useful means of targeting drugs to the stomach and can be used for delivery of effective antibiotic therapy. Our findings can also be used to develop an effective blueprint for targeted stomach-specific drug delivery, and can be investigated further in vivo with other drugs, using a slight modification in the techniques used here for preparation of the nanospheres.

The amoxicillin nanosphere suspension developed for this study was found to have suitable physicochemical properties and to have a favorable particle size range. The nanospheres were found to release the drug to a maximum extent in vitro. This novel dual-release formulation adds to the already important domain of targeted drug delivery systems, and holds promise as an alternative to conventional drug delivery, enabling bacteria such as *H. pylori* which are notoriously difficult to treat to be managed effectively.

**Acknowledgment**

This work was supported by the Deanship of Scientific Research, King Faisal University, Saudi Arabia (research project number 130012, 2012).

**Disclosure**

The author reports no conflict of interest in this work.

**References**

1. Hejazi R, Amiji M. Stomach-specific anti-*H. pylori* therapy. I: preparation and characterization of tetracycline-loaded chitosan microspheres. *Int J Pharm*. 2002;235(1–2):87–94.
2. Krueger S, Roessner A, Kuester D. Murine models of *H. pylori*-induced gastritis and gastric adenocarcinoma. *Pathol Res Pract*. 2011;207(10):599–607.
3. Peterson WL. *Helicobacter pylori* and peptic ulcer disease. *N Engl J Med*. 1991;324(15):1043–1048.
4. Meurer LN, Bower DJ. Management of *Helicobacter pylori* infection. *Am Fam Physician*. 2002;65(7):1327–1336.
5. Romano M, Cuomo A. Eradication of *Helicobacter pylori*: a clinical update. *Med Gen Med*. 2004;17(6):19.
6. Lin CK, Hsu PI, Lai KH, et al. One-week quadruple therapy is an effective salvage regimen for *Helicobacter pylori* infection in patients after failure of standard triple therapy. *J Clin Gastroenterol*. 2002;34(5):547–551.
7. Liu Z, Lu W, Qian L, Zhang X, Zeng P, Pan J. In vitro and in vivo studies on mucoadhesive microspheres of amoxicillin. *J Control Release*. 2005;102(1):135–144.
8. Axon AT. The role of acid inhibition in the treatment of *Helicobacter pylori* infection. *Scand J Gastroenterol Suppl*. 2004;201:16–23.
9. Fontana G, Liciardi M, Mansueto S, Schillaci D, Giannona G. Amoxicillin-loaded polyethyleneoxyacrylate nanoparticles: influence of PEG coating on the particle size, drug release rate and phagocytic uptake. *Biomaterials*. 2001;22(21):2857–2865.
10. Patel JK, Chavda JR. Formulation and evaluation of stomach-specific amoxicillin-loaded Carbopol-934P mucoadhesive microspheres for anti-*Helicobacter pylori* therapy. *J Microencapsul*. 2009;26(4):365–376.
11. Naoki N, Yohko A, Masafumi N, Mayumi T, Megumi K, Yasuyuki O. Mucoadhesive microspheres containing amoxicillin for clearance of *Helicobacter pylori*. *Antimicrob Agents Chemother*. 1998;42(10):2492–2494.
12. Narkar M, Sher P, Pawar A. Stomach-specific controlled release gellan beads of acid-soluble drug prepared by ionotropic gelation method. AAPS Pharm Sci Tech. 2010;11(1):267–277.

13. Jain P, Jain S, Prasad KN, Jain SK, Vyas SP. Polyelectrolyte coated multilayered liposomes (nanocapsules) for the treatment of Helicobacter pylori infection. Mol Pharm. 2009;6(2):593–603.

14. Moogooee M, Rameanzadeh H, Jassar O, Omidi Y, Davaran S. Synthesis and in vitro studies of cross-linked hydrogel nanoparticles containing amoxicillin. J Pharm Sci. 2011;100(3):1057–1066.

15. Awasthi R, Kulkarni GT, Pawar VK, Garg G. Optimization studies on gastroretentive floating system using response surface methodology. AAPS Pharm Sci Tech. 2012;13(1):85–93.

16. Sambathkumar R, Venkateswaranurthy N, Vijayabaskaran M, Perumal P. Formulation of clarithromycin loaded mucoadhesive microspheres by emulsification-internal gelation technique for anti-Helicobacter pylori therapy. Int J Pharm Sci. 2011;3:173–177.

17. Rao J, Geckeler KE. Polymer nanoparticles: preparation techniques and size-control parameters. Prog Polym Sci. 2011;36(70):887–913.

18. Lee SH, Heng D, Kiong WNG, Chan HK, Tan RBH. Nano spray drying: A novel method for preparing protein nanoparticles for protein therapy. Int J Pharm. 2011;403(1–2):192–200.

19. Sham JOH, Zhang Y, Finlay WH, Roa WH. Löbenberg R. Formulation and characterization of spray-dried powders containing nanoparticles for aerosol delivery to the lung. Int J Pharm. 2004;269(2):457–467.

20. Pilcer G, Vanderbist F, Amighi K. Preparation and characterization of spray-dried tobramycin powders containing nanoparticles for pulmonary delivery. Int J Pharm. 2011;408(1–2):192–200.

21. Li X, Anton N, Arpagaus C, Belletteix F, Vandamme TF. Nanoparticles by spray drying using innovative new technology: The Buchi Nano Spray Dryer B-90. J Control Release. 2010;147(2):304–310.

22. Dhanaraju MD, RajKannan R, Selvaraj D, Jayakumar R, Vamsadhara C. Preparation and characterization of nanoparticles for anti-Helicobacter pylori infection. Int J Pharm. 2011;408(1–2):248–256.

23. Wang LY, Gu YH, Su ZG, Ma GH. Preparation and improvement of release behavior of chitosan microspheres containing insulin. Int J Pharm. 2006;311(1–2):187–195.

24. Joscelyne SM, Trägårdh G. Membrane emulsification a literature review. J Memb Sci. 2000;169(1):107–117.

25. Lu B, Zhang JQ, Yang H. Lung-targeting microspheres of carboplatin. Int J Pharm. 2003;265(1–2):1–11.

26. Harsha NS, Rani SH. Drug targeting to lungs by way of microspheres. Arch Pharm Res. 2006;29(7):598–604.

27. Singh SK, Chidrawar VR, Ushir YV, Vadalia KR, Sheth NR, Singh S. Pharmaceutical characterization of amoxicillin trihydrate as mucoadhesive microspheres in management of H. pylori. Pharm Tech. 2010;6(2):348–358.

28. Gomez-Estaca J, Baluguer MP, Gavara R, Hernandez-Munoz P. Formation of zein nanoparticles by electrohydrodynamic atomization: Effect of the main processing variables and suitability for encapsulating the food coloring and active ingredient curcumin. Food Hydrocolloids. 2011;28:82–91.

29. Devrim B, Bozkir A, Canef K. Formulation and evaluation of reconstitutable suspensions containing ibuprofen-loaded Eudragit microspheres. Acta Pol Pharm. 2011;68(4):593–599.

30. Raula J, Eerikinen H, Kauppinen EI. Influence of the solvent composition on the aerosol synthesis of pharmaceutical polymer nanoparticles. Int J Pharm. 2004;284(1–2):13–21.

31. Tonon RV, Grossi CRF, Hubinger MD. Influence of emulsion composition and inlet air temperature on the microencapsulation of flaxseed oil by spray drying. Food Res Int. 2011;44(1):282–289.

32. Burki K, Jeon I, Arpagaus C, Betz G. New insights into respirable protein powder preparation using a nano spray dryer. Int J Pharm. 2011;408(1–2):248–256.

33. Wang LY, Gu YH, Su ZG, Ma GH. Preparation and improvement of release behavior of chitosan microspheres containing insulin. Int J Pharm. 2006;311(1–2):187–195.

34. Mizoe T, Beppu S, Ozeki T, Okada H. One-step preparation of drug-loaded poly (dl-lactide-co-glycolide) injectable microspheres. Int J Pharm. 2011;36(70):887–913.

35. Burki K, Jeon I, Arpagaus C, Betz G. New insights into respirable protein powder preparation using a nano spray dryer. Int J Pharm. 2011;408(1–2):248–256.

36. Wang LY, Gu YH, Su ZG, Ma GH. Preparation and improvement of release behavior of chitosan microspheres containing insulin. Int J Pharm. 2006;311(1–2):187–195.