

Development and Optimization of Amoxicillin Floating Raft System to effectively treat *Helicobacter pylori* infection

Formulación y Optimización del Sistema Flotante de Amoxicilina para el tratamiento efectivo de la infección por *Helicobacter pylori*

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RESUMEN

Introducción: El objetivo del presente estudio fue formular y caracterizar el Sistema Flotante (FRS, siglas en Inglés) de Amoxicilina para prolongar el tiempo de residencia gástrica y liberación del fármaco para el enfoque efectivo del *Helicobacter pylori*.

Método: Para el presente estudio se seleccionaron como factores goma guar, Monoestearato de glicerilo (GMS), carbonato de calcio. Como reacciones, se seleccionaron el período de congelación (h), el lapso de flotación (min), y el porcentaje acumulado de liberación del fármaco (CDR). Para la experimentación se seleccionaron el diseño factorial 2³ con réplicas.

Resultados: Se observó que la goma guar y el GMS fueron los factores principales que afectaron el período de congelación y mostraron un efecto sinérgico (positivo). Mientras que la goma guar y el carbonato de calcio mostraron un efecto positivo y el GMS mostró un efecto antagónico (negativo) en el lapso de flotación. El porcentaje CDR mostró un efecto antagónico en todos los factores. Se emplearon curvas de nivel para identificar el diseño del espacio, análisis numéricos posteriores produjeron 12 soluciones óptimas en base a la deseabilidad. El FRS mostró un mayor AUC_{0-t'}, C_{max}, t_{max}, y t_{1/2} cuando se comparó con la formulación comercial, aproximadamente 2.30 cambios múltiples y prolongación con liberación sostenida por más de 24 h que pudo deberse a una mejor congelación.

Conclusiones: Se puede concluir que el sistema flotante se desarrolló satisfactoriamente por la aplicación del Diseño de Experimentos (DoE) con menores ensayos y utilizando fácilmente los excipientes disponibles para una mejor flotación, congelación y suministro constante del fármaco.

Palabras clave: Sistema flotante; *Helicobacter pylori*; diseño factorial; estudios *in vivo*

ABSTRACT

Introduction: The aim of the present study was to develop and to characterize the floating raft system (FRS) of Amoxicillin to enhance gastric residence time and drug release to target *Helicobacter pylori* effectively.

Method: In the present study, guar gum, glyceryl monostearate (GMS), calcium carbonate were selected as factors. Gelation duration (h), floating lag time (min), and % Cumulative drug release (CDR) were selected as responses. 2³ factorial design with replicates was selected for experimentation.

Results: It was observed that guar gum and GMS were the major factors affecting gelation duration, increase in the quantity of both guar gum and GMS increased gelation duration i.e., sustained gelation period (24 h). Floating time increased with an increase in the amount of guar gum and calcium carbonate, whereas an increase in the quantity of GMS decreased floating time. Guar gum, calcium carbonate, and GMS exhibited an antagonistic effect on % CDR. Contour plots were used to identify design space; further numerical analysis yielded 12 best solutions based on desirability. FRS exhibited greater AUC_{0-t'}, C_{max}, t_{max}, and t_{1/2} when compared to marketed formulation approximately 2.30 folds enhancement and prolongation with a sustained release for greater than 24 h that might be due to better gelation.

Conclusions: It can be concluded that the floating raft system was successfully developed by the Design of experiment (DoE) application with fewer trails and by utilizing easily available excipients for better floating, gelation, and sustained delivery of the drug.

Keywords: Floating raft system; *Helicobacter pylori*; factorial Design; *In vivo* studies

INTRODUCTION

About half a million new cases per year have been recorded with gastric and duodenal cancers majorly linked to *H. Pylori* infection ⁽¹⁾. The development of the proper floating retentive system is essential to eradicate *H. Pylori* since its absorption in the mucous layer is more effective than the basolateral membrane ⁽²⁾. Hence Floating Raft System (FRS) was selected to improve gelation, duration, and better drug release for effectively treating this infection. As of literature search, this drug was not developed in the form of FRS. Hence we selected this system to improve its availability at the target site for the eradication of *H. pylori*. When FRS comes in contact with gastric fluids, a viscous cohesive gel is formed, wherein each portion of the liquid swells forming a continuous layer called Raft.

Several dosage forms were formulated among FRS showed benefits like ease of manufacturing processes, administration, and to float faster with better compliance to the patient ⁽³⁾. The problems of bacterial resistance and side effects associated with Amoxicillin could be reduced by formulating gastro retentive FRS, as they provide an appropriate

prolongation of drug release near the bacterial ecological niche ⁽⁴⁾.

Design of Experiment (DoE) is an essential statistical approach that can be utilized to optimize and to select compatible stable formulation by the screening of variables. Adopting this technique minimizes the time where the formulation can be optimized by a few trails, thereby reducing the cost of final Amoxicillin product with in-built quality. Hence in the present study, the formula was developed and optimized by 2³ factorial design with replicates.

MÉTHODES

Preparation of the floating raft system

Sixteen formulations (Table 1) were designed as per 2³ factorial design (8 formulations+8 replications, Design expert software version 11.0) and were developed by the method as described below.

Preparing a dispersion medium in which the guar gum was dispersed in 90 mL of deionized water along with 0.25% w/v-sodium citrate and heated at 90°C. The medium stirred until it appears as a homogenous viscous liquid. Later cooled to 40°C and dispersed Amoxicillin drug in the medium and then followed by the addition of 10 mL of calcium carbonate dispersions with stirring. GMS was melted to 90°C and dispersed it into the above drug incorporated gum formulation by homogenization using a high-speed homogenizer ⁽²⁾ (Figure 1).

Table 1. Formulation table of Amoxicillin floating raft system (Factors & responses)

Formulation	Factor 1 A: Guar gum %	Factor 2 B: Calcium carbonate %	Factor 3 C: GMS %	Response 1 Gelation duration (h)	Response 2 FT Lag time (min)	Response 3 % CDR at 24 h
1	3.00	1.00	1.00	20	4	90.2
2	3.00	1.00	1.00	20	4.1	88.4
3	4.00	1.00	1.00	24	2.5	82.4
4	4.00	1.00	1.00	24	2	80.2
5	3.00	1.50	1.00	15	5	84.4
6	3.00	1.50	1.00	15	5.5	80.7
7	4.00	1.50	1.00	24	2.5	85.1
8	4.00	1.50	1.00	24	1.5	90.7
9	3.00	1.00	1.50	24	1.5	95.4
10	3.00	1.00	1.50	24	2	94.7
11	4.00	1.00	1.50	24	1	97.4
12	4.00	1.00	1.50	24	1.2	94.2
13	3.00	1.50	1.50	24	4	87.6
14	3.00	1.50	1.50	24	4	80.7
15	4.00	1.50	1.50	20	3.5	80.2
16	4.00	1.50	1.50	20	3.5	85.1

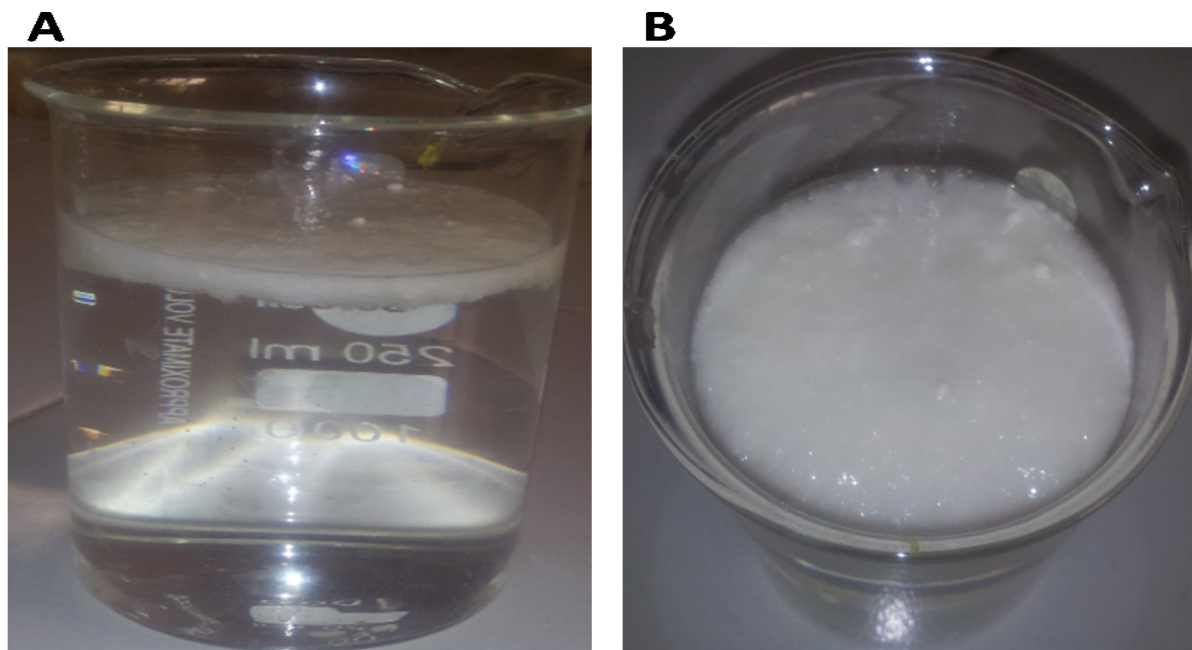


Figure 1. Amoxicillin Floating Raft System (FRS) A. Side view B. Top view

Invitro Gelation study

Prepared and collected pH 1.2 dissolution media in vials to maintain SGF (stimulated Gastric fluids) condition at $37\pm 0.5^{\circ}\text{C}$. Add a drop of formulation in each vial and make a visual observation on Gelation lag time (time taken to the formation of gel) and Gelation duration (time taken to dissolve)⁽⁵⁾.

Measurement of viscosity

FRS viscosity was measured by Brookfield viscometer by changing the angular velocities 10 -100 RPM at $23\pm 0.5^{\circ}\text{C}$ ⁽⁶⁾.

In-vitro floating study

USP dissolution apparatus II was used. 900 mL of pH 1.2 dissolution media was prepared to create SGF (Stimulated Gastric Fluid) condition. 10 mL of in-situ gel formulation was added slowly in each dissolution medium without disturbance at $37\pm 0.5^{\circ}\text{C}$, and visual observation was made on floating lag time (time for the formulation to appear on the medium surface) and floating duration (time taken to dissolve gel)⁽²⁾.

In vitro drug release

The study was performed using 0.1N HCl buffer (pH 1.2) and maintained at $37\pm 0.5^{\circ}\text{C}$ with 50 RPM. 10 ml of gel formulation was taken into the watch glass and kept in the dissolution medium. The sample was collected at each interval time and replaced the same amount of buffer medium, which was prewarmed and maintained at $37\pm 0.5^{\circ}\text{C}$. Spectrophotometrically measured at 210 nm. All dissolutions were performed in triplicate.⁽²⁾

In vivo Pharmacokinetic studies

A total of 4 healthy rabbits were randomly divided into two groups, and each group consisting of two animals. Group-I: Administration of marketed formulation (EDMOX). Group-II: Administration of optimized FRS. After a single oral administration, 0.25ml blood sample was collected from either left or right marginal ear vein using a 3ml needle at 0, 0.5, 1, 1.5, 3, 4.5, 6, 12 and 24h. Blood samples were analyzed further by HPLC technique for measurement of pharmacokinetic parameters like AUC_{0-t} , C_{max} , t_{max} and $t_{1/2}$. The measurements were assessed at $p < 0.05$ ^(7,8).

RESULTS AND DISCUSSION

In the present study, guar gum, calcium carbonate, and GMS were selected as factors and gelation time duration (h), floating lag time, and % cumulative drug release (%CDR) were selected as responses. These were selected based on the literature survey. After selection, preliminary studies were conducted, and the concentrations were fixed, followed by experimentation using the Design of experiments (Factorial design, Design expert software version 11). The study was conducted at 2 levels 3 factors with replications (8) (8+8 total of 16 experiments). It was observed that guar gum and GMS were the significant factors affecting gelation duration, increase in the quantity of both guar gum, and GMS increased gelation duration i.e., sustained gelation period (24 h). Floating time increased with an increase in the amount of guar gum and calcium carbonate, whereas an increase in the quantity of GMS decreased floating time.

An increase in the amount of guar gum, calcium carbonate, and GMS decreased % CDR i.e., sustained the release of the drug for 24 h (equation 1, 2, 3). All the desired factors exhibited probability at <0.05 , indicating significance at that level (Table 2). In all the cases, the predicted R^2 is closer to adjusted R^2 and adequate precision values were greater than 4 and indicated adequate signal. Hence it was used to navigate the design space. A significant factor that affects the formulation was identified by Pareto charts and by cooks distance. Pareto chart is a useful tool to identify statistically significant effects by t-value limits; the effects above the t-value indicated significant factors. The predicted vs. actual values plot revealed that the experimental values were in close agreement with the predicted values (Figure 2).

$$\text{Gelation duration} = -66.00000 + 23.50000 * \text{Guargum} + 64.00000 \text{ GMS} - 170000 \text{ Guargum} * \text{GMS} \quad \text{--- 1}$$

$$\text{Floating Lag time} = +1.22500 + 0.075000 * \text{Guar gum} + 7.35000 * \text{Calcium carbonate} - 1.60000 * \text{GMS} - 1.30000 * \text{Guar gum} * \text{Calcium carbonate} \quad \text{--- 2}$$

$$\begin{aligned} \% \text{CDR at 12h} = & +418.50000 - 114.60000 * \text{Guar gum} - 264.20000 \\ & * \text{Calcium carbonate} - 211.60000 * \text{GMS} + 89.10000 * \text{Guar gum} \\ & * \text{Calcium carbonate} + 79.90000 * \text{Guar gum} * \text{GMS} + 170.60000 \\ & \text{Calcium carbonate} * \text{GMS} - 62.40000 * \text{guar gum} * \text{calcium carbonate} * \text{GMS} \quad \text{--- 3} \end{aligned}$$

Table 2. ANOVA representing significance at $p < 0.05$

Source	Sum of squares	df	Mean square	F Value	p-value	
Model	112.75	3	37.58	11.00	0.0009	significant
A-Guar gum	20.25	1	20.25	5.93	0.0315	
C-GMS	20.25	1	20.25	5.93	0.0315	
AC	72.25	1	72.25	21.15	0.0006	
Residual	41.00	12	3.42			
Lack of fit	41.00	4	10.25			
Pure error	0.000	8	0.000			
Cor Total	153.75	15				

Contour plots (Figure 3) were used to identify design space; further numerical analysis yielded 12 best solutions based on desirability. To achieve desired criteria, i.e., gelation time closer to 24 h, floating lag time 1-2 min, and % CDR greater than 95%, the factors can be varied at 3.79-4.00 % of guar gum, calcium carbonate and GMS at 1 and 1.5% respectively with the desirability of 0.899-0.902. Calcium carbonate generates gas, which is obtained from calcium. When the formulation comes in contact with stomach acidic pH, carbon dioxide gas is evolved. It shows in situ release along with floating properties because of entrapping of gel network in the formulation and enhance gel floating on the surface of the dissolution medium (*In vitro*) or the stomach (*In vivo*). These Ca^{2+} ions are responsible for gelation through a cross-linking polymer chain. Lipid emulsification ameliorated the FRS characteristics. Incorporation of GMS enhanced the gelation efficiency with an increased Gelation period from a few hours to greater than 24h. The AUC_{0-t}

of the marketed formulation, was found to be 42.27 $\mu\text{g.h/ml}$. The increase in AUC_{0-t} was observed in the FRS (test) formulation, which was around 68.42 $\mu\text{g.h/ml}$. This indicated the drug availability for a long duration. The mean peak plasma concentration of test formulation C_{max} 6.97 $\mu\text{g/ml}$ gradually reached in 4 h. In the case of marketed formulation (R), the C_{max} was 3.02 $\mu\text{g/ml}$ gradually reached in 3 h. The C_{max} of the marketed formulation was less when compared with the test formulation. The increase in t_{max} indicated drug availability for a prolonged period. The plasma elimination half-life ($t_{1/2}$) of the marketed formulation was 4.72 h, and the FRS (T) formulation was 5.98 h, which were significantly different. The overall C_{max} , t_{max} , AUC_{0-t} and $t_{1/2}$ were completely different between both the test and marketed formulations (Fig 4). Therefore the prepared formulations released the drug for a prolonged period essential for the effective eradication of *H. pylori* infection.

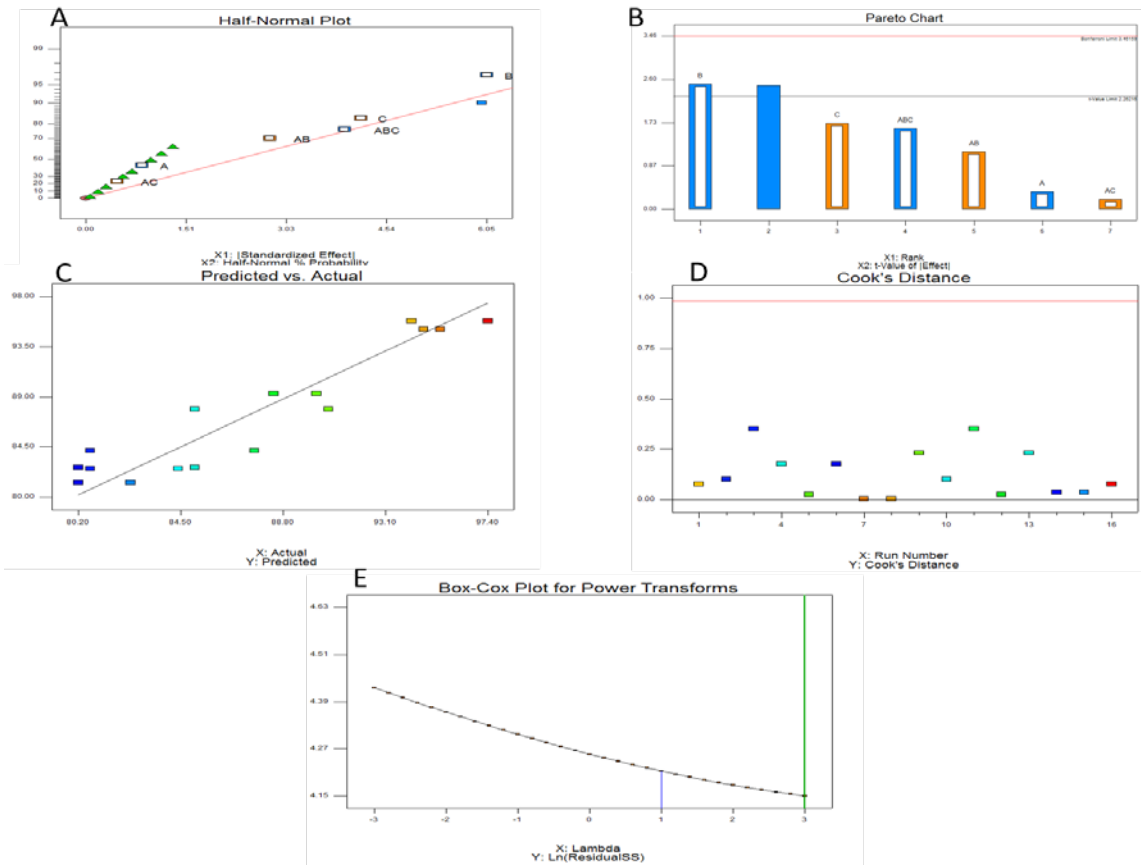


Figure 2. Diagnostic plots A. Half-Normal Plot B. Pareto chart C. Predicted Vs. Actual D. Cooks Distance E. Box-cox plot for Power transforms

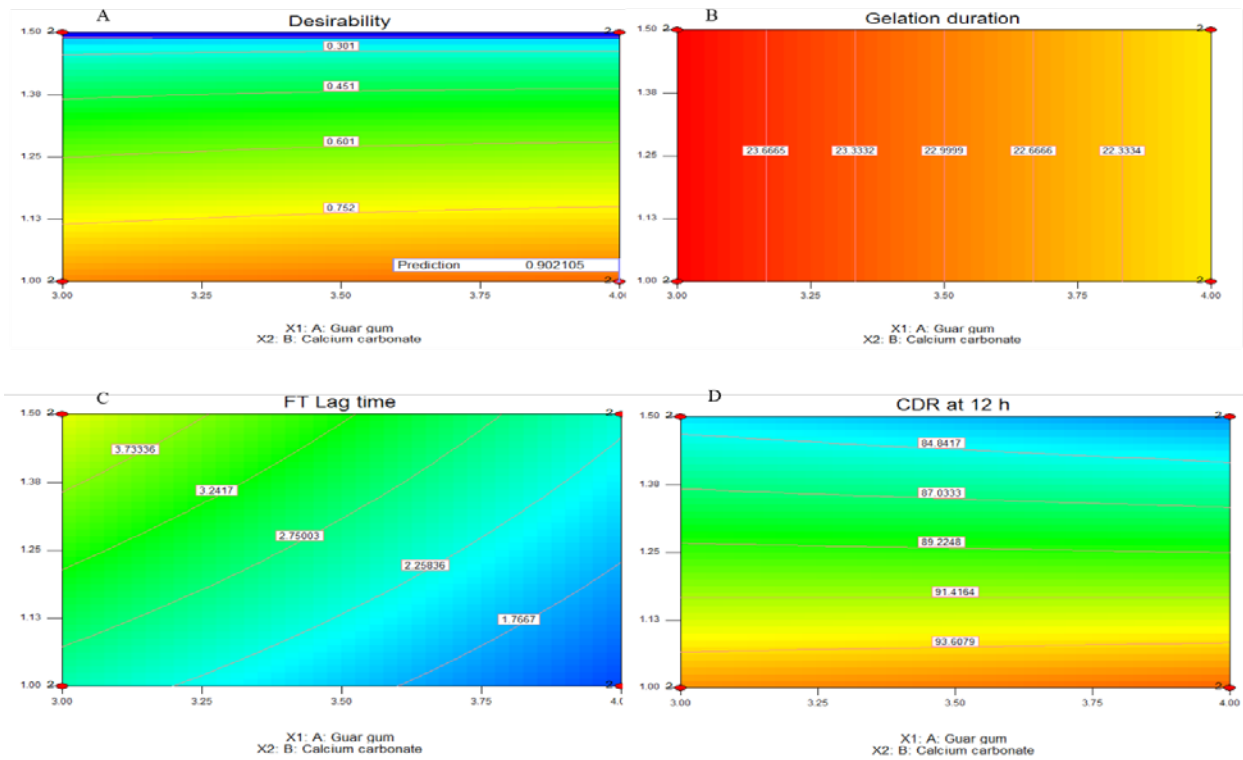


Figure 3. Contour plots under Design space

Figure 4. Mean plasma concentration-time profiles after oral administration of FRS (test) and marketed formulations; Each point represents mean \pm SE (n=4, at p<0.05).

CONCLUSION

Amoxicillin FRS was successfully developed and optimized by factorial Design with replicates to enhance gastric residence time and drug release to effectively target *Helicobacter pylori*. It can be concluded that the floating raft system was successfully developed by the Design of experiment (DoE) application with fewer trails & by utilizing easily available excipients for better floating, gelation, and sustained delivery of the drug. *In vivo* studies exhibited better pharmacokinetic parameters in FRS (test formulation) than marketed formulation that proved extended and prolonged release of drug at the required site for treating *Helicobacter pylori*.

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