ABSTRACT

Objective: The objective of this current study is to fabricate ocuserts to control the drug release from chosen bioadhesive polymeric matrices to enhance patient compliance. Ciprofloxacin HCl (CFX HCl) was selected as a model drug.

Methods: Different bioadhesive polymers with different film forming capabilities namely Hydroxy Propyl Methyl Cellulose (HPMC K4M), Poly Vinyl Alcohol (PVA), Sodium Carboxy Methyl Cellulose (Na CMC), Hydroxy Propyl Cellulose (HPC), Sodium Alginate (Na Alg.), pullulan and Xanthan Gum (XG) in different ratios were used in fabricating ocuserts using solvent-casting technique. Propylene Glycol (PG) was used as a plasticizer to facilitate the fabrication process. Characterization tests of the developed ocuserts were performed as well as bioadhesive tests and in vivo release studies of the incorporated drug. The obtained results were analysed using different release kinetic models. Stability of the selected ocuserts was investigated at 40±1.5 °C and 75±5% Relative Humidity (RH) for three months' storage period. In vivo ocular irritation test was performed to investigate the safety of the formula in rabbits' eyes as well as to test the release profile and thus to estimate In vitro In vivo correlation.

Results: All the prepared ocuserts showed the uniformity of film characterization and bioadhesion strength ranged from 240 ±66 and 158±52dyne/cm². Selected formula from the in vitro release study tested for in vivo study showed the slow release of ciprofloxacin drug up to 24 h with no signs of eye irritancy. Results for In vitro In vivo correlation showed an excellent correlation with R² value of 0.9982.

Conclusion: PVA based ocuserts proven to be a promising once-daily, effective and safe ocular delivery system of the drug.

Keywords: Polymeric matrix, Ocuserts, In vivo tests, Once-daily

INTRODUCTION

Eye is an essential organ with unique qualities and enable us to see the world. Drug delivery through ocular route is local more than systemic to bypass the high blood concentration of the drug, which may cause eye damage [1].

The solitary anatomy, biochemistry, and physiology of the eye challenge the formulator to circumvent the protective barriers of the eye without causing any tissue damage because of the impervious nature of the eye [2]. About 70% of the total ophthalmic preparations available in market are conventional ophthalmic dosage forms (eye drops and ointments). For these dosage forms, frequent dosing is the main disadvantage which from an industry point of view, could increase the bioavailability of the drug in a controlled and sustained manner [3–6].

Ocuserts (synonyms of Ocular inserts) are drug delivery devices constructed from polymeric materials with solid or semi-solid consistency, delivering the incorporated drug to the ocular surface by placing it in the conjunctival sac. Ocuserts demonstrate many advantages such as delivering an accurate dose, minimizing systemic drawbacks of ocular remedies, prolong the ocular residence time thus reducing the frequency of administration leading to improving patient compliance, possibility of releasing drugs at a slow and constant rate which from an industry point of view, could increase shelf life stability [7]. Ocular route provides large absorption surface area and high vascularization offering good penetration for hydrophilic, low molecular weight drugs achieving fast onset of action [8]. Compared to oral delivery, the ocular administration provides a potential of dose reduction because it avoids the hepatic first pass metabolism. Therefore, in emergency therapy, ocular administration of conventional drugs would be considered as an alternative to other administration routes [9].

Ocuserts applied behind the eyelid were found to prolong the retention time and precision of dosing [10]. However, films were found to tend to move across the surface of the eye, thus resulting in irritation. It has been shown that the addition of mucoadhesive polymers to ocular films, which can adhere to the epithelial surface [11] reduced film movement across the eye, minimizing ocular irritation and burning sensations [12]. The main disadvantage reported for ocuserts is the annoying sensation accompanying its insertion in the eye, but the numerous advantages of ocuserts supersede this single disadvantage [13] as seen by the implementation of this technology in several successfully marketed ocuserts (Ocusert®, Ocufit®SR, and Minidisco9) [14].

Many ocular regions have poor accessibility to systemic circulation due to the presence of protective barriers which makes the local delivery via topical administration is the favoured route for the treatment of ocular diseases. Typical conditions that require ocular administration include ocular infections and disorders (like conjunctivitis and glaucoma) [2].

CFX HCl is a broad-spectrum fluoroquinolone antibiotic that can be taken orally or intravenously [10].

CFX HCl is used systematically in numerous microbial contaminations such as dermal, pulmonary, urinary tract infections and topicaly in conjunctivitis and anterior ocular infections [15].

CFX HCl was the most used antibacterial agent worldwide, and the fifth regularly used generic antibacterial in the USA during the last decade in the 20th century [16].

The aim of the current study was to develop bioadhesive ocuserts for the topical delivery of CFX HCl suitable for a once a day application employing different bioadhesive polymers.

MATERIALS AND METHODS

Materials

CFX HCl was a gift sample from Eipico, Egypt. Pullulan was purchased from VWR International LLC, West Chester, PA, USA. HPC, HPMC K4M,
PG, and Glycerin were obtained from ADWIC, El-Nasr Chemical Co., Cairo, Egypt. Na Alg., Di-Sodium Hydrogen Phosphate (Na2HPO4) and Potassium Di-Hydrogen Phosphate (KH2PO4) were supplied from El-Nasr pharmaceuticals company, Cairo, Egypt. Na CMC and PVA were supplied from Sigma Company for pharmaceuticals, Cairo, Egypt and were used without further purification.

**Methods**

**UV scanning solution preparation**

A stock solution of CFX HCl was prepared by placing accurately weighed 100 mg in a 100 ml measuring flask and freshly prepared phosphate buffer pH 7.4 was added till reaching the mark. From the resultant solution, 1 ml was taken and completed to 100 ml with phosphate buffer pH 7.4 and then scanned in the 200-400 nm using a UV-spectrophotometer (Shimadzu UV spectrophotometer, 1601-PC double-beam spectrometer, Kyoto, Japan).

**FTIR spectroscopy**

Drug identification and detection of any possible incompatibility existed between the excipients is done through Fourier transform infrared (FTIR) spectroscopy. CFX HCl powder and the formula (F-I) blend were examined using FTIR spectrophotometer (model Impact 410, Milwaukee, WI, USA) with a scanning range 500 to 4000 cm⁻¹ and the resolution was 1 cm⁻¹. KBr disk method was used to obtain the FTIR spectra of CFX HCl and the formulation (F-I) blend. The infrared peaks of pure CFX HCl were analysed and compared with the peaks obtained for the formulation (F-I) blend to identify if there is the deletion of or additional formation of peaks.

**Fabrication of ocuserts**

CFX HCl ocuserts were prepared by solvent casting technique using HPMC K4M, pullulan, CMC, HPC, Na Alg., PVA and XG as biodegradable bioadhesive polymers [17, 18]. All polymers were completely dissolved in distilled water (left overnight for uniform dispersion) except PVA which was dissolved in hot water at 70 °C. After dissolving the polymers in water, PG as a plasticizer was added (2.5% w/w of polymer). Bath sonicator (Model SI101H 230, Sonix IV, CA, USA) was used to remove air bubble appeared after addition of CFX HCl into the polymeric solution. After complete mixing of drug and polymer, 10 ml of the clear solution was poured into the clean petridish moistened with glycerin. The petri dish was covered with an inverted glass funnel of stem orifice 0.6 cm in diameter with a cotton plug closing the stem of the funnel. Clearance was provided for the escape of the solvent vapors by raising the base of the funnel (2 cm) just above the resting surface. The funnel was an aid to control the rate of evaporation of the solvent and reducing the blistering of the surface of the deposited film [19]. After complete evaporation of the solvent, cast films were obtained, cut into definite circular pieces by cork borer (8 mm), wrapped in an aluminum foil and stored in a CaCl2 desiccator at room temperature in a dark place for further evaluation studies.

**Characterization of film fabrication**

The thickness of the ocusert was measured using a film thickness tester (Vernier caliper, Shanghai, China) at various regions of the film (n = 3) [20, 21].

Three films of the same size were weighed on an electronic digital balance (Mettler AJ 100, Switzerland). The average weight, as well as the weight variation, were calculated [20-22].

Petri dish containing 0.5 ml distilled water was used to moisten the ocusert for 30 s. Then the pH meter electrodes (Digital pH meter, Toshniwal Pvt. Ltd., India) were brought in contact with the surface of the ocusert and left for 1 min for equilibration. The pH value was noted and recorded [20, 21].

To determine moisture loss, weighted ocusert was placed in a desiccator containing anhydrous calcium chloride. After three days the film was reweighed, and moisture loss was calculated using the following equation [23, 24].

\[
\text{Moisture loss (\%) } = \left( \frac{W_i - W_f}{W_i} \right) \times 100 \quad \ldots \quad (1)
\]

Where \( W_i \) is the initial weight, and \( W_f \) is the final weight of the film. The study was done in triplicate for each ocusert formulation.

For Moisture gain study, weighted ocusert was placed in a desiccator containing 100 ml of saturated solution of aluminum chloride to maintain 80% humidity. After three days film was reweighed, and moisture gain was calculated using the following equation [23, 24].

\[
\text{Moisture gain (\%) } = \left( \frac{W_f - W_i}{W_i} \right) \times 100 \quad \ldots \quad (2)
\]

Where \( W_i \) is the initial weight, and \( W_f \) is the final weight of the film. The study was done in triplicate for each ocusert formulation.

**Table 1: Formulation of ocuserts with the different film-forming agents**

<table>
<thead>
<tr>
<th>Formula code</th>
<th>HPMC K4M</th>
<th>PVA</th>
<th>CMC</th>
<th>HPC</th>
<th>Na Alg.</th>
<th>Pullulan</th>
<th>XG</th>
</tr>
</thead>
<tbody>
<tr>
<td>F-I</td>
<td>1.5%</td>
<td></td>
<td>1.5%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F-II</td>
<td></td>
<td></td>
<td></td>
<td>1.5%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F-III</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.5%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>F-IV</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2%</td>
<td></td>
</tr>
<tr>
<td>F-V</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.5%</td>
<td></td>
</tr>
<tr>
<td>F-VI</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.5%</td>
</tr>
<tr>
<td>F-VII</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*All formulae were prepared with 2.5% PG as a plasticizer and 17 mg of CFX HCl.

**Fig. 1: Schematic diagram of the device used for bioadhesion test**

(A) modified-balance (B) weight (C) rubber stopper (D) cornea with ocusert (E) glass vial (F) support to the vial to adjust weight.

Bioadhesion characterization

Ex vivo bioadhesive strength

The freshly excised cornea of a cow was used as a model mucous membrane for the measurement of bioadhesive strength. Fresh cow cornea was obtained from a local slaughterhouse and used within 2 h of slaughter. The mucosal membrane was separated by removing the underlying fat and loose tissues. The membrane was washed with distilled water and then with isotonic phosphate buffer pH 7.4 at 37 °C.

The bioadhesive strength of ocusert (n = 3) was measured on a modified two-arm physical balance as illustrated in fig. 1 [25, 26].

The pan at the left arm of the balance was detached, and a vertical thread was tied to its end. A rubber stopper, hanging downward, was hung to the lever of the left arm. The ocusert to be tested adhered to the downward facing side of the rubber stopper. Cow cornea was tied onto the open mouth of a glass vial filled with isotonic phosphate buffer. The vial was fitted in the centre of a glass beaker filled illustrated ed with simulated tear fluid (STF) with pH 7.4, 37 °C±1 °C. The apparatus was set such that the vial (mucosal membrane tied on it, facing upward) lies exactly below the rubber stopper (patch adhered onto it, facing downward). The rubber stopper was lowered so as to make the ocusert come in contact with the membrane. After facilitating the contact between the two, weight was put on the right limb of balance and increased gradually until the ocusert got detached from the cornea. The weight (gram force) required to detach the ocusert from the mucosal surface gave the measure of detachment strength, calculated by:

\[
\text{Detachment stress } (\text{dyn/cm}^2) = \frac{W_g}{A} \quad (3)
\]

where w is the weight required for the detachment of ocusert, g is the acceleration due to gravity considered as 980 cm/s, and A is the area of the mucosal surface exposed (cm²) [25–27].

Ex vivo bioadhesion time

The ex vivo bioadhesion time was ascertained (n = 3) after application of the ocusert onto freshly cut cow cornea. The fresh cornea was fixed in the inner side of the beaker, above 2.5 cm from the bottom, with cyanoacrylate glue. One side of each ocusert was wetted with one drop of isotonic phosphate buffer pH 7.4 and pasted to the cornea by applying a light force with a fingertip for 30 s. The beaker was filled with 500 ml of STF pH 7.4 and was kept at 37 °C±1 °C. Ocusert adhesion was monitored up to 6 h and the time required for the ocusert to detach from the cornea was recorded as the mucoadhesion time [25–27].

In vitro drug release study

The in vitro drug diffusion from the ocuserts was studied using the cylindrical glass tube (Internal diameter 15 mm and length 100 mm). The diffusion cell membrane (Prehydrated cellophane membrane) was tied to one end of the cylindrical tube, which acted as a donor compartment and the ocusert was placed inside this compartment. The entire surface of the membrane was in contact with 25 ml of distilled water and then with isotonic phosphate buffer pH 7.4. The diffusion cell membrane remained in pH 7.4 and was kept at 37 °C±1 °C. Ocusert adhesion was monitored up to 6 h and the time required for the ocusert to detach from the cornea was recorded as the mucoadhesion time [25–27].

The accelerated stability studies were carried out in accordance with the ICH guidelines [31]. Enough ocuserts (packed in aluminium foil) were stored, with RH of 75 % and at a temperature of 4±0.5 °C for three months. The samples were tested for drug content after 0, 7, 15, 30 and 90 days respectively.

In vivo ocular irritation test

Approval for the use of animals in the study was obtained from the Al-Azhar University, Faculty of Pharmacy, Animal Ethics Committee (Ref. No.166/2018) which comply with the 3ARRIVE guidelines. The current experiment was carried out in accordance with the U. K. Animals (Scientific Procedures) Act, 1986 and associated guidelines, EU Directive 2010/63/EU for animal experiments. In addition, all institutional and national guidelines for the care and use of laboratory animals were followed.

Our in vivo study aims to determine the objective irritation potential of the prepared ocusert and the in vivo drug profile of the selected ocusert. The rabbit was chosen as a model for this study because its eye simulates an adult human eye with respect to size, shape, physiology, and composition of tears [32].

New Zealand rabbits of either sex weighing 2.5 to 3.5 kg were used to measure the in vivo ocular irritation test in the eye. The rabbits were purchased from the Nile Company for pharmaceuticals and chemical industries (Al Sawah St, Cairo, Egypt).

The animals were housed in cages in the animal house (located in the faculty of pharmacy (boys branch), Al-Azhar university, Cairo, Egypt) under controlled conditions of temperature (27±2°C) and light. They were fed with standard laboratory diet, and water was provided ad libitum. Ethical clearance for the handling of experimental animals was obtained. The rabbits were fed balanced diet pellets and maintained in a temperature-controlled room, at 20 °C to 24 °C before the experiment. 6 animals were used in the experiment where free leg and eye movement was allowed.

The investigated ocusert was placed in the left eye while the right eye of each rabbit was considered the control. The ocusert was sterlized by using UV radiation before in vivo study. The ocuserts and other materials were exposed to UV radiation for 1 h [33]. After sterilization, ocusert was transferred into polyethylene bag with the help of forceps inside the sterilization chamber itself. The observations based on scoring approach (0 = normal; 3 = worst) according to Peyman scale. [34] Peyman et al. [34] established the safety of the developed ocuserts in the rabbit eye. Ocusert was removed each hour for the first eight hours to determine the amount of drug released and hence evaluate in vitro in vivo correlation. The amount of drug remaining in each ocusert was determined as per the assay method of the drug in ocuserts given in drug content. Cumulative percent drug released [CDR] in vivo was calculated [35, 36].

Sterility test

Any preparation intended to be placed in the eye must be sterile, therefore, testing the sterility is a very important evaluation parameter. The tests for sterility were done by detecting the presence of viable forms of bacteria, fungi, and yeast in or on preparations. The tests were carried out under strict aseptic techniques to avoid accidental contamination of the preparation [37].

The sterility test was performed according to the guidelines of Indian Pharmacopoeia applying direct inoculation method. 2 ml of prepared CFX HCl ocusert solution was removed with a sterile needle and aseptically transferred to Medium A (fluid thioglycolate medium) and Medium B (soyabean-casein digest medium) separately. After mixing with each medium, incubation of each medium for 7 d was done (The incubation temperature was 30 °C to 35 °C in the case of fluid thioglycolate medium and 20 °C to 25 °C in the case of soyabean-casein digest medium) [37].

Microbiological studies

Staphylococcus aureus (S. aureus; ATCC® 25923) test microorganism purchased from American Type Culture Collection (PO Box 1549, Manassas, VA 20108 USA) was used to assess the biological activity of the selected ocusert formulation. The test organism was seeded in
nutrient agar and allowed to solidify in the petri dish. An ocusert was carefully placed over the agar layer at a suitable distance [38]. The plates were then incubated at 37±0.5 °C for 24 h. After incubation, the obtained zone of inhibition was compared with the control ocusert (i.e. ocusert with no drug).

RESULTS AND DISCUSSION

UV scanning

The stock solution was scanned in the 200-400 nm UV regions. The wavelength adopted for absorbance measurement was the observed wavelength maximum ($\lambda_{\text{max}}$) at 278 nm.

FTIR spectroscopy

The characteristic absorption bands of CFX HCl at 1281.59 and 1612 cm$^{-1}$ (fig. 2b) was due to the stretching vibration of C–F bond and the vibration of phenyl framework conjugated to – COOH, respectively. The stretching vibration at 1705.07 cm$^{-1}$ was due to – COOH and the stretching vibrations of C–H from the phenyl framework at 2962.66 and 2908.65 cm$^{-1}$ were also observed. All the peaks were also found in drug-loaded ocusert that confirms the presence of the drug in the polymers without any interaction.

Fabrication of ocuserts

Ocuserts were prepared using polymers pullulan, HPC, HPMC K4M, CMC, Na Alg., PVA and XG. The selection of polymers depended on their film-forming properties, their biodegradability and retardant to biodegradability to provide sustained release pattern. As shown in fig. 3, the prepared ocuserts were found to be satisfactory uniform, transparent and flexible [39].

Characterization of the prepared ocuserts

The prepared ocuserts were evaluated for their thickness at three random points. All ocuserts showed uniformity in thickness as indicated by low SD of the measured thickness. Variations in the ocusert thickness is reflected by the different amount of polymer needed for ocusert fabrication (table 2).

Uniformity of the weight of the ocuserts is illustrated by low SD obtained after measurement. The mean weight value varied between 2.44±0.15 mg to 8.4±1.3 mg as it’s affected proportionally by the ocusert thickness.

The mean values of three replicates of both % moisture loss and % moisture gain were recorded in table 2. Formulae F–V and F–VI become brittle upon exposure to moisture loss while formulae III and IV showed gelation of the film and in formula V disintegration of the film into fragments took place upon moisture gain. These observations are the net result of different hydration properties of different polymers.

All prepared formulae showed uniform drug content in the range of $94.14±8.22\%$ to $98.22±5.02\%$ with low SD values.

Folding endurance of all batches was found between $19±6$ to $72±11$, the significant difference in the folding endurance of the prepared films is due to the use of different polymer types and concentrations.

The values of surface pH vary between 6.2 to 7.2 which give rise to the assumption that the ocuserts will cause no irritation upon insertion in the eye.
Bioadhesion may be defined as the adhesion between a polymer and a biological membrane, e.g., mucus. The strength of bio-adhesion is affected by numerous factors such as the molecular weight of polymers, contact time with mucus, swelling rate of the polymer, and biological membrane used in the study. All ocuserts showed appreciable bioadhesive detachment stress that ranged between 240±66 and 158±52 dyne/cm² (Fig. 4) indicating a potential of sustaining the stay and enhancing contact with cornea. Various mechanisms have been proposed to explain the in vitro bioadhesion or mucoadhesion phenomena such as electrical double layers, electrostatic attractions, hydrogen bonding, Van der Waals force, hydrophobic bonding, wetting, diffusion-interpenetration, physical entanglements, and surface-free energy [40].

Most of the hydrophilic polymers can absorb water and swell. This can increase the potential to adhere onto mucosal surfaces. This is the simplest mechanism of adhesion and has been defined as “adhesion by hydration”.

Na CMC can increase surface charge density, and the carboxylic group can form hydrogen bonds with tissue [40]. HPMC is the long chained, non-ionic polymer and the mucoadhesive property could be due to the formation of physical or hydrogen bonding with the mucus components. HPMC can relieve the dryness and irritation even in the case of reduced mucus secretions [40, 41]. Na CMC and HPMC show faster hydration rate and thereby swelling which helps in the interpenetration of mucus and polymer resulting in bio-adhesion.

HPMC is a non-ionic polymer containing only hydroxyl groups, which can form weak hydrogen bonds with mucous layers. Furthermore, owing to its slow rate of hydration it can form a strong surface gel that efficiently adheres onto the mucosal surface and remains in contact for a longer time. For this reason, it can be characterized as one of the most effective mucoadhesive polymers [25, 42].

The highest bioadhesive force showed by F-IV containing HPC while the lowest value showed by F-VI containing pullulan.

### Table 2: Physicochemical evaluation of different formulations*

<table>
<thead>
<tr>
<th>Formulae</th>
<th>Thickness (mm)</th>
<th>Weight (mg)</th>
<th>Moisture loss %</th>
<th>Moisture gain %</th>
<th>Drug content %</th>
<th>Folding endurance</th>
<th>Surface pH</th>
</tr>
</thead>
<tbody>
<tr>
<td>F-I</td>
<td>0.26±0.05</td>
<td>4.6±0.24</td>
<td>21.6±7.4</td>
<td>36.4±7.8</td>
<td>94.6±7.4</td>
<td>56±7</td>
<td>6.23</td>
</tr>
<tr>
<td>F-II</td>
<td>0.34±0.02</td>
<td>8.4±1.3</td>
<td>24.9±6.6</td>
<td>33.2±11.4</td>
<td>97.0±6.6</td>
<td>72±11</td>
<td>6.2</td>
</tr>
<tr>
<td>F-III</td>
<td>0.24±0.07</td>
<td>4.6±0.07</td>
<td>6.8±2.7</td>
<td>Gelation</td>
<td>98.2±5.02</td>
<td>45±6</td>
<td>6.22</td>
</tr>
<tr>
<td>F-IV</td>
<td>0.22±0.00</td>
<td>4.9±0.63</td>
<td>12.8±5.3</td>
<td>Gelation</td>
<td>95.7±12.7</td>
<td>24±4</td>
<td>7.1</td>
</tr>
<tr>
<td>F-V</td>
<td>0.31±0.04</td>
<td>5.7±0.05</td>
<td>Brittle</td>
<td>Disintegrated</td>
<td>98.0±9.7</td>
<td>22±6</td>
<td>6.66</td>
</tr>
<tr>
<td>F-VI</td>
<td>0.32±0.02</td>
<td>6.42±0.26</td>
<td>30.5±8.5</td>
<td>34.7±13.8</td>
<td>96.37±10.2</td>
<td>62±14</td>
<td>6.8</td>
</tr>
<tr>
<td>F-VII</td>
<td>0.31±0.04</td>
<td>2.4±0.15</td>
<td>Brittle</td>
<td>35.4±11.2</td>
<td>94.14±8.22</td>
<td>19±8</td>
<td>7.2</td>
</tr>
</tbody>
</table>

* (Results are expressed as mean±SD, n=3)

### Ex vivo bioadhesion strength

Ex vivo bioadhesion characterizations

The ex vivo bioadhesion time (residence time) of ocuserts varied from 2360±140 to 260±32 s. (Fig. 5). It was observed that a gradual increase in the residence time occurred with a concomitant increase in the polymer viscosity. The observation can be assigned to the inherent property of the polymer HPMC that although showing significantly higher swelling is less water affinity and hence tends to retain its structure better. In addition, increased viscosity led to the formation of a surface gel that maintained its structural integrity for a longer period of time, thereby resulting in increased residence time.

### In vitro drug release study

The cumulative percent of CFX HCl released from ocuserts as a function of time is shown in Fig. 6 which revealed that 99% of the drug was released from HPMC based ocuserts in about 5 h.

Na Alg and pullulan based ocuserts showed very close results (i.e., Na Alg based ocuserts released about 94% and pullulan based ocuserts released about 92% in 8 h).

PVA, HPC and XG based ocuserts released 98%, 92% and 90% respectively in about 24 h. The most sustaining effect is attained by Na CMC based ocuserts where it shows only the release of 34% in about 24 h.

The enormous difference in drug release may be explained by the change in the polymer forming the ocuserts. Generally, drug release from the polymeric matrices is elicited by the ease of water accessibility into the matrix, which breaks the polymer-polymer bonds and thus simultaneously leads to binding of water and polymer molecules, separation of polymer chains, swelling to form a gel, and finally dispersion of polymer chains in the medium. "The drug dissolves in the gel and diffuses to the exterior with a rate depending on its concentration gradients and its diffusion ability through the gel. Concurrently, the latter is eroded with a rate depending on polymer molecular weight and hydrodynamics of release medium. The drug release pattern depends on the relative rates of these processes” [43].
As, HPMC is a hydrophilic polymer which facilitates ease of water penetration into the polymer matrix and hence the ease of drug diffusion. Although pullulan is also hydrophilic polymer composed of polysaccharide, it shows more sustaining in drug release than HPMC where a lesser amount is sufficient in fabricating film with satisfactory mechanical strength.

Na Alg. showed more retardation in drug release reached to 8 h to complete release of the incorporated drug. This may be explained by features of Na Alg. in which hydrophilic functional groups of the polymer and water molecules tend to bind together by hydrogen bonding leading to excessive swelling results in retarding of drug release [44].

In case of PVA (a hydrophilic polymer); but in its processing in formulating a film, it was heated to 70 °C and plasticized by PG. Both heat and PG addition caused crosslinking of PVA resulting in retardation in drug release [45].

The release pattern of HPC can be explained by high swelling behavior which results into inclusion of water inside the polymer matrix resulting into retardation in drug release [46].

XG is considered a good matrix-forming material for sustained-release tablets as it tends to form viscous gels in the presence of water, whether they are used alone or in combination with other gums or polymers. This property also affects the drug release from ocuserts as seen in the dissolution behaviour [46].

In case of Na CMC there was great retardation in drug release although its hydrophilic character it had great swelling behaviour and inclusion of water molecules into the polymer matrix results in the formation of gel into the polymer matrix causes retardation in drug release as seen the remaining of gel in the diffusion cell at the end of 24 h but it may be promising in preparing formulations for several days but in ocular route it may be annoying to the patient so its excluded from further study.

By overviewing the aforementioned release pattern, FIV was selected as a promising ocuserts providing the overall incorporated drug in 24 h comply with the hypothesis of the study by formulating once a day dose.

Considering the drug release kinetics when the data was plotted as cumulative % of the drug released vs. time according to zero order equation, F. I, VI and VII showed a fair linearity indicating higher correlation than first order and Higuchi equation, with the highest R2 values compared to other plots, while, F. II, III, and IV showed a fair linearity with first order indicating higher correlation than zero order and Higuchi equation, with the highest R2 values compared to other plots except for Alginate showing diffusion release. The zero-order kinetics [47] reveals delivery of the drug in a sustained manner whereby the drug is held in a reservoir representing the ocusert and is released at a constant rate to provide a constant concentration in the cornea which improves patient compliance [48, 49].

**Table 3: R² correlation values of different kinetic models of CFX HCl ocuserts**

<table>
<thead>
<tr>
<th>Formula</th>
<th>R² of zero</th>
<th>R² of first</th>
<th>R² of higuchi</th>
</tr>
</thead>
<tbody>
<tr>
<td>F-I</td>
<td>0.998568</td>
<td>0.82807</td>
<td>0.996927</td>
</tr>
<tr>
<td>F-II</td>
<td>0.94915</td>
<td>0.99503</td>
<td>0.978563</td>
</tr>
<tr>
<td>F-III</td>
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<td>0.99716</td>
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<td>0.943501</td>
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<td>0.96156</td>
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<td>F-VI</td>
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<td>0.96004</td>
<td>0.995587</td>
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<tr>
<td>F-VII</td>
<td>0.99752</td>
<td>0.98684</td>
<td>0.97749</td>
</tr>
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</table>

**Stability studies**

Formula F-II was subjected to accelerated stability study to determine the physical stability of the formulation. There were no significant changes regarding the physical properties at the end of the three months and drug content during the study period. The overall degradation is less than 1.25%. A tentative shelf-life of one year may be assigned to formulation as per ICH guidelines. No change in physical appearance of ocusert was reported during the period of study thus revealing that F II passed the stability test indicating that it was chemically, physically and microbiologically stable at the examined temperature for 3 mo. However, its shelf life needs to be established by further studies at different temperatures and humidity conditions.

**UV irradiation**

The efficiency of the sterilization process is indicated mainly by sterility test where the ocuserts is examined for sterility by two media where it showed no sign of turbidity after 7 d for Medium A (fluid thioglycollate medium) and Medium B (soyabean-casein digest medium) indicating excellent sterility of the tested ocuserts.

**In vivo drug release and irritation study**

The results of *in vivo* drug release study are presented in fig. 7. The ocusert for the first 5 h can be removed from the rabbit eye and tested for drug remaining. After the first 5 h the ocusert adhesive to the ocular surface and cannot be removed. The rabbits also examined for eye irritancy for 7 d after the study and no rabbit show any sign of irritancy.
(score 0) for all formulations. The in vivo drug release study from F. II was found to be in accordance with that of the in vitro drug release study. Hence, we tried to correlate in vivo results with the in vitro percentage drug release. The correlation value was found to be 0.9982 [50]. Therefore, the formula (F-II) exhibited strong in vitro-in vivo correlation revealing the efficacy of the formulation (fig. 8). No drag out of circular inserts at the time of experiment was happened which suggest that the dimension (8 mm) was suitable as ocuserts [50].

Fig. 7: In vivo cumulative percent release of formula II, (results are expressed as mean±SD, n=3)

Fig. 8: In vitro In vivo correlation of formula II, (results are expressed as mean±SD, n=3)

Fig. 9: Sequence of photos of in vivo experiment showing (a) before insertion of ocusert (b) during removal of ocusert (c) and after the complete dissolving of ocusert after 24 h
Microbiological studies

Testing the selected ocuserts (Formulae F-II and F-III) against staphylococcus showed good antimicrobial activity indicated by clear zones of inhibition (3 mm diameter for each formulae) when tested microbiologically on solidified agar. While control ocusert (occusert with no drug) shows no zone of inhibition as represented in fig. 10 [51].

CONCLUSION

Various bioadhesive formulations of C.F.X HCl ocuserts were prepared using a solvent casting method and evaluated. Ocuserts (formula F-II) consisting of a 1.5% PVA with 2.5% PG satisfied all the pharmaceutical parameters of bioadhesive ocuserts and demonstrated controlled release of the drug in the eye over the period of 24 h. Formula F-II is considered a promising formulation suitable for a once a day dose and thereby improving the patient compliance by providing benefits of reducing in the frequency of administration by controlled drug release. Pharmacokinetics and pharmacodynamics studies in human beings are needed to be carried out to establish the therapeutic utility of this system.

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AUTHORS CONTRIBUTIONS

All the author have contributed equally

CONFLICT OF INTERESTS

No conflicts of interest were reported by the authors. Only the authors are responsible for the scientific hypothesis, performance of the work and writing of this paper.

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Fig. 10: Photo represents the microbiological study of formula II and III ocusert against S. Aureus


