A Novel Topical Spray Formulation for Ginkgo Biloba for Antifungal Activity

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Abstract

The aim of this study is to prove the antifungal and antibacterial activity of Ginkgo biloba GKB from glycerin nanocapsules and nanoemulsions. GKB was formulated as spray formulation for topical application using ethyl alcohol 95%/glycerol/water based formulations. GKB spray was evaluated for their particle size distribution, pH, content uniformity, content per spray, in vitro release and antifungal activity. The antifungal activity was determined by Agar well-diffusion method. The formulae were tested for their skin permeation and the stability of the formulations. The results showed uniform formulae with nano-sized particles, pH compatible with the topical formulation and higher drug content. Diffusion studies of the optimized formulations through the abdominal rabbit skin showed higher penetrated drug over a period of 24 h. Stability studies indicated that formulations were stable. Skin irritation studies confirmed the compatibility of spray formulae with skin. Agar well-diffusion method showed that GKB was effective compared with fluconazole. The inhibition zones for all formulae were significant as compared with standard fluconazole. The results obtained showed that the topical spray formulation could be a promising and innovative therapeutic system as topical antifungal for the transdermal administration of GKB. The developed formulation can be used for delivery from spray formulation. This study proved the local activity of GKB as antifungal drugs.

Keywords: Topical spray; In vitro release; Ginkgo biloba

Introduction

Several different antifungal preparations were used to treat various fungal infections [1,2]. They are formulated as sprays, creams, solutions, ointment, shampoos, vaginal tablets, oral medicines, and injections [3]. The duration of treatment depends on kind of fungal infection, health problems of the infected patient, problems associated with immune system and types of formulations. The courses of treatment can be as short for few days or long for months [4].

Antifungal liquids or sprays (topical antifungals) can be used to treat skin fungal infections, scalp and nails [5]. These antifungal drugs can cause various type of adverse effect such as burning or redness at the site of application [6,7]. It was reported that 80 endophytic fungi were isolated from GKB, they were active against the chosen fungi using the agar diffusion method. The study indicated that GKB could be used as a fungicide to defend host plants against pathogenic organisms [8]. GKB has antifungal activity against some kind of endophytic. GKB extract holds flavonoid glycosides and unique terpene lactones as main active constituents [9].

Here GKB is presented for the 1st time as a new finding antifungal drug for human. It was formulated as novel spray formulations as safe natural product which have no side effects and are easy to use. GKB spray is a favorable antifungal medicine for curing all types of fungal infections, since it can be used to reduce itching, burning and scaling accompanied with all kinds of Tinea infections. The aim of this study was to formulate spray formula containing GKB and to verify the antifungal and antibacterial action of GKB. GKB was formulated as spray formulation for topical application using ethyl alcohol, glycerol and water established formulations. GKB spray was assessed for numerous parameters such as particle size distribution, pH, drug content per spray, content uniformity, in vitro release and antifungal activity. The established formulation was used for delivery from spray formulation. The antifungal activity was determined by Agar well-diffusion method.

Materials and Methods

Materials

Ginkgo biloba was kindly supplied from Lipits pharmaceuticals, Cairo, Egypt. Glycerol, ethyl alcohol (95%) and whatmann paper were purchased from, El-Gomhoria Chem. Co., Cairo (Egypt). Glass bottle and spray pumps were purchased from Measom Freer, Wigston, UK. All chemicals were of analytical grades. All glasswares were thoroughly washed with freshly prepared aqua regia (HCl: HNO₃, 3:1), extensively rinsed with Millipore water several times and oven-dried at 50°C for 2-3 hours before use.

Methods

Selection of excipients: Formulae were prepared using aqueous, non-aqueous diluents, co-solvents, and permeation enhancers as shown in Table 1. Ethyl alcohol 95% and water were selected as solvents for formulating of spray formulae due to the good solubility of GKB in these solvents [10]. Glycerol was used as a permeation enhancer and as coating material for GKB [11]. F₆ were found to be optimized formulation (Table 1).

Selection of containers and spray pumps: Glass bottles with a cubic figure carved at the base were employed for filling the transdermal spray formulae. The selected spray pumps can be used not only for body spray but also for hairspray, sunscreen, insect repellent, mist sprays, lens & glass cleaning solutions [12]. Spray pumps were chosen with

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over-cap (Diameter 18 mm, Height 39 mm, Neck Size 15 mmR4, Dose 120 µl, min 1,000 for Spray Pump) as it is more suitable for body spray.

**Formulation development of topical spray preparation:** GKB spray formulations were developed as topical solutions made up of ethyl alcohol 95%, and water dissolved in one phase [10]. Glycerol was used in this formula as enhancer and as coating material for GKB [11]. Briefly, GKB was dissolved in required amount of water and stirred for 1 hr as. The required amount of glycerol were added to the aqueous solution of GKB and stirred for another 1 hr. The dissolved GKB solution were dropped on the half amount of ethyl alcohol 95% and stirred with a magnetic stirrer. One mL of this solution was diluted with citrate buffer to 25 mL in volumetric flask. Then the solution was analyzed for GKB spectrophotometrically at λ max 380 nm against a blank solution prepared in the same manner using aqueous alcoholic spray [19-22].

Drug content%=[Concentration of entrapped drug]/[Total drug concentration] × 100 (1)

**In vitro skin permeation experiments:** In vitro penetration studies were carried out using abdominal male rat skin (150 ± 25 g). To obtain the skin of the rats, animals were first sacrificed. It was fixed on a modified Franz diffusion cell with a surface area of 7.0 cm². The dissolution medium was citrate buffer at pH 6 thermostatic at 37° and stirred with a magnetic stirrer. One mL of this solution was diluted with citrate buffer to 25 mL in volumetric flask. Then the solution was analyzed for GKB spectrophotometrically at λ max 380 nm against a blank solution prepared in the same manner using GKB spray [18,19,21,22]. Each penetration experiment was frequented 6 times. Percentage cumulative amounts of drug penetrated were plotted as a function of time. Approval of the Institutional Animal Ethical Committee was obtained prior to the commencement of skin irritation studies from the faculty of medicine, Asyut University, Asyut, Egypt. Moreover, the flux was calculated from the slope of the linear portion of the obtained curve. The permeability coefficient (Kp) of GKB crossways rat skin was calculated using Fick's first law of diffusion, which is expressed by the following equation:

\[ Kp = \frac{J}{C} \]  

Where; J is the flux (mg/cm²/hr) and C is the drug concentration in donor compartment.

**Stability studies:** The conventional formulations were set aside for stability testing. The chemical stability of the formulations was estimated by evaluation of the percentage drug residual in the formulations, drug release pattern, observing any change in pH, appearance, spray pattern, leakage rate and average weight per actuation assessed physical stability [14,18].

**Skin irritation studies**

The optimized formulations were sprayed on the scrap of shaved skin and observing the resulting reactions such as erythema and edema after 24 h. Briefly, five human volunteers were selected with age of 30–40 years, having no history of allergy against GKB, and the test formulation was applied to the back of hand on an area of 2 square inch. Then, the volunteers were observed for appearance of lesions or irritation [14,18].

**Antifungal activities:** The study of antifungal activity for GKB spray formula was carried out using different strains of fungi and bacteria. The antifungal activity of dry powder of GKB formulae was determined...
by agar well-diffusion method [23,24]. The media used was Mueller-
Hinton agar at only 4 mm deep, poured into 100 mm or 150 mm Petri
dishes. The pH level of the agar must be between 7.2 and 7.4. Briefly, the
work was done using aseptic technique. A sterile swab was placed into
the broth culture of a specific organism and then the excess liquid was
removed by gently pressing or rotating the swab against the inside of the
tube. To obtain uniform growth, the plate was streaked with the swab
in one direction, the plate was rotated at 90° and the plate was splashed
again in that direction. The plate was allowed to dry for approximately
5 minutes. Antibiotic discs containing specific antibiotics were
distributed onto the plate. Using a flame-sterilized forceps, each disc
was gently pressed to the agar to ensure that the disc is attached to
the agar. The plates were incubated overnight at an incubation temperature
of 37°C (98.6°F). Inoculation was made with a diluted broth culture to
match a 0.5 McFarland turbidity standard, which is roughly equivalent
to 150 million cells per mL. The inhibition zones for all GKB formulae
were measured as compared with fluconazole as a standard antifungal.

Statistical analysis: All statistical analysis was performed using
One-way analysis of variance ANOVA. Minitab16 Statistical Software
with the Tukey’s multiple comparison was employed for comparing
the formulations with each other. Statistically significant differences
were assumed when p<0.05. All values are expressed as their mean ±
standard deviation [18].

Result and Discussion

Spray formulae

Formula consisting of ethyl alcohol 95%: glycerol: water in the
ratio 7:2:1 exhibited preferred spray patterns with high and good
dispersibility of the GKB F3. The optimized formulae were selected
based on the transparency of the solution, dehydration rate, spray
display and thickness of the film formed. Formulation F1 was a simple
drug solution in solvent system and showed high clarity as it has a
thin film that adhered to the skin for periods up to 4 h. Comparing
the physicochemical characteristics of the films, spray patterns and
drug content, formulations F1, F2 and F3. F3 were found to be better
as compared to other formulations. The detailed results of qualitative
and quantitative tests for reference formulation and the optimized
formulations (F2 and F3) are summarized.

Size, zeta potential and PDI are factors that designate the stability of
the obtained nanoparticles [25-28]. The surface charge plays an essential
role in the stability of the obtained nanoparticles and the degree of
zeta potential is indicative of the colloidal stability of the system [29].
The results showed a satisfactory small and uniform particle size. The
formulated nanoparticles were of a uniform Z average diameter and
nano-size range, Figure 1. DLS recorded only one peak with intensity
of 100%. PDI (heterogeneity index or simply dispersity (Đ)) indicates
the stability of the product since it denotes the particles distribution in
colloidal solution. Smaller PDI suggest the homogeneity of the particle
size in suspension, ideally the value of PDI should be < 0.70 because
they indicated a particle size distribution falls within a narrow range of
sizes [27,30-33]. In the present study all PDI were of acceptable range
as recorded by DLS for all GKB spray, indicating the homogeneity of the
particle size in the obtained suspension as showed in Figure 1.

The zeta potential zeta potential of all particle types was positive,
Figure 2. GKB had a highly positively zeta potential, indicating that
GKB was coated with repulsive force enough to be stable. DLS recorded
also high-count rate (185 kpcs) indicates that the concentration of
nanoparticles was high enough for measurements.

The spray display was found to be affected by the size and shape of
actuator orifice as well as by the capacity of pump. Formula F3 showed
good spray displays in terms of uniform and spherical messe due to the
supple and cohesive film of the glycerol used.

F3 showed pH in the range of 6.0-6.5 which mimicking the skin pH.
The 100 μl pumps delivered average weight of 70 ± 5.0 mg per actuation

<table>
<thead>
<tr>
<th>Formulation</th>
<th>Flux (mg/cm².h)</th>
<th>Q 24 (mg/cm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>F1</td>
<td>0.0177</td>
<td>0.4374</td>
</tr>
<tr>
<td>F2</td>
<td>0.0266</td>
<td>0.4663</td>
</tr>
<tr>
<td>F3</td>
<td>0.0269</td>
<td>0.5678</td>
</tr>
</tbody>
</table>

*Amount of drug released per unit surface area after 24 h. Formulations F1 and F3 showed zero order kinetic model while formulation F2 showed root time kinetics
with R² value of 0.9608. [Formulations F2 and F3 (containing film formers and penetration enhancers) showed more flux and Q 24 value for the drug as compared to
Formulation F1].

Table 2: Permeation parameters of ginkgo biloba across rabbit abdominal skin.

Figure 1: Size distribution of ginko biloba solution for formula F3 as determined by zetasizer nano.

Figure 2: Size distribution of ginko biloba solution for formula F3 as determined by zetasizer nano.
(n=5). There was no statistically significant variation in the amount emitted per actuation indicating the effectiveness of the pump system in delivering reproducible amounts of the formulation per actuation (Table 1). Formulations F1, F2 and F3 showed average drug contents per spray of 100 to 104%. This test indicates the amount of the therapeutically active ingredient delivered per metered spray from the metered dose containers. Formulation F3 was found to show better uniformity in terms of content per spray. We inferred that the formulation F3 had more uniformity of the dosage of the active ingredient delivered through the metered pump compared to formulations F1 and F2.

**Ex vivo skin permeation**

Permeation properties of the GKB were illustrated initially by conducting diffusion studies on the simple organic drug solution (F3). The diffusion studies from the optimized formulations F1, F2 and F3 were carried out over a period of 24 h. Rabbit abdominal skin was used as animal model skin, because of its similarity with human skin ex vivo. As illustrated in Figure 3, a linear correlation was achieved between Q (the amount cumulative of penetrated GKB across the unit surface area of the skin) and time up to 24 h. The formulation F3 exhibited significantly higher penetration (P ≤ 0.05) of GKB as compared to formulation F1 and F2. The reason for this higher permeation is due to the presence of high concentration of glycerol. Formula F3 contained glycerol with high concentration as the permeation enhancers leading to more permeation of drug into the skin. The same results were confirmed the penetration enhancer of glycerol were presented previously [20,34,35].

**Statistical analysis**

The results illustrated in Table 2 confirmed the release kinetics fits the zero order kinetic models for all formulations. Analysis of variance showed that these correlations are statistically significant (P ≤ 0.05). Formula F3 holding higher concentration of glycerol as penetration enhancer showed more flux and Q24 value for the drug as compared to the other formulations F1 and F2.

**Stability studies**

Stability studies showed the formulations to be stable in terms of physicochemical parameters, drug content and drug release, under all the storage conditions for F3 as selected to be the best formulation as shown in Table 3.

**Skin irritation studies**

No erythema or edemas were observed in the primary skin
irritation studies of the optimized formulations on the rabbits. It found to be safe and non-irritant for transdermal application. The formulation developed was more efficient for a 24 h after application. However results need to be confirmed by pharmacokinetic studies.

**Antifungal activities**

This study proved the local activity of GKB as antifungal drugs. Vertical distribution of microfungi on decomposing GKB powder is shown in Table 4, in which the calculated occurrence frequencies are also indicated. Seven species of microfungi were recorded from powder extract of GKB. As listed previously, antifungal activities have been recorded for GKB as compared with fluconazole itself as reference antifungal. All the recorded results suggested the antifungal activity of GKB even fluconazole is better.

**Conclusions**

The evidence suggests that GKB is more effective as local antifungal action in comparing to fluconazole. GKB was successfully formulated as solution for spray pattern which can be used in the future trials as topical formulae on numbers of participants. From the results obtained in the present work, it can be concluded that GKB can be an innovative and promising approach for the transdermal administration of GKB. The diffusion studies indicated that the permeation of GKB formulations through the skin was much higher as compared to the diffusion of simple organic solution of the drug.

**References**


