

Preparation, characterization, and *in vivo* pharmacokinetics of thermosensitive *in situ* nasal gel of donepezil hydrochloride

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Donepezil hydrochloride thermosensitive *in situ* gel for nasal delivery was prepared by using Poloxamer 407 and Poloxamer 188 as thermoreversible polymers, hydroxypropyl- β -cyclodextrin and ethylparaben as permeation enhancer and preservative, respectively. The gelation temperature and time, pH value of the gel formulation were found to meet the requirements for nasal administration. The *in vitro* erosion and *in vitro* release tests exhibited obvious drug sustained release behavior. Meantime, main pharmacokinetic parameters such as t_{\max} , c_{\max} and AUC in plasma as well as in brain were significantly different between the nasal gel formulation and intragastric drug solution in rats ($p < 0.01$). The relative bioavailability and drug targeting efficiency of the gel formulation were calculated to be 385.6 and 151.2 %, respectively. Thus, the drug gel formulation might be a potential new delivery system for treatment of Alzheimer's disease due to its higher bioavailability and better distribution to brain when compared to oral route.

Keywords: donepezil hydrochloride, thermosensitive *in situ* gel, nasal delivery, bioavailability, brain targeting

Accepted November 18, 2019

Published online December 16, 2019

Alzheimer's disease (AD) is a central nervous system (CNS) disease characterized by loss of cognitive behavior, memory impairment, and disturbance in daily activities (1). The cholinergic deficit is thought to be one of the major pathological causes of this disease, thus some cholinesterase inhibitors such as risvastigmine, tacrine and donepezil hydrochloride (DPZ) have been clinically applied in order to improve the available levels of acetylcholine in brain tissues (2). DPZ is a specific and reversible AchE inhibitor and it also could produce neuroprotective effect (3). More importantly, the drug has fewer side effects than non-specific AchE inhibitors, so presently it has become a clinical first-line drug for treatment of mild to moderate AD in many countries (4).

As a CNS disease, treatment of AD by oral administration route is particularly challenging due to the existence of the blood-brain barrier (BBB) (5). So far, intranasal delivery has been thought as an important dosing route for some drugs with many advantages such as large mucosa surface area, rapid onset of action, avoidance of hepatic first pass effect, high

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bioavailability and especially circumvention of BBB (6). Furthermore, this administration route is non-invasive, of convenient application and good compliance. Therefore, intranasal dosing of drugs has been thought to be an ideal alternative to the control of AD (7).

Currently, the marketed dosage forms of DPZ are as oral tablets or capsules. Though this administration route is convenient for some of patients, the drug showed obvious gastrointestinal side effects and muscle convulsions (8). Moreover, the therapeutic efficacy of the DPZ oral preparations was not satisfactory due to the existence of BBB. Thus, in order to overcome the above shortcomings of oral route, some new dosage forms such as nano-suspension, microemulsion, liposomes and liposomal hydrogel of DPZ for nasal administration have been studied and furthermore the enhanced bioavailability as well as brain targeting characteristics of the above new nasal delivery systems also confirmed (9–12). However, the above novel delivery systems of DPZ for nasal route have many limitations such as complicated formulation and preparation process, inconvenient dosing, poor physicochemical stability, or short residence time in nasal cavity. Presently, drops and sprays are two main nasal preparations widely applied in clinical treatment, these traditional liquid formulations, however, are unfavorable because of short residence time and thus poor bioavailability. In recent years, *in situ* nasal gel has gained more attention for systemic drug delivery because of its advantages such as simple formulation, easy preparation process, better physicochemical stability, convenient use, accurate dose, long nasal residence time, high bioavailability, sustained drug release and potentially improved therapeutic efficacy (13). Thus, the aim of present research was to prepare a nasally administered thermosensitive *in situ* gel of DPZ and to further characterize its *in vitro* properties. *In vivo* pharmacokinetics, relative bioavailability, and especially brain targeting characteristics of the *in situ* gel formulation was also investigated in rats.

EXPERIMENTAL

Materials

Donepezil hydrochloride was bought from Hubei Hongxin Ruiyu Fine Chemical Co., Ltd. (China). Poloxamer 407 (P407) and poloxamer 188 (P188) were bought from BASF Co., Ltd. (China). Hydroxypropyl- β -cyclodextrin (HP- β -CD) was purchased from Shandong Fine Chemical Co., Ltd. (China). Ethylparaben was given by Aladdin Biochemical Technological Co., Ltd. (China). All other reagents were of analytical or chromatographic grades.

Preparation of thermosensitive *in situ* gel

Optimal formulation of DPZ thermosensitive *in situ* gel for nasal delivery was obtained from the results of our previous research and it was prepared by the cold method. Briefly, HP- β -CD (3.5 %, *m/V*) was first dissolved in suitable volume of distilled water by agitation at room temperature. The required amount of DPZ (5.0 %, *m/V*) and ethylparaben (0.05 %, *m/V*) were then dissolved successively in the above solution under ultrasonic condition. After cooling to room temperature, P407 (19.6 %, *m/V*) and P188 (8.2 %, *m/V*) was slowly added to the mixed solution and then fully dispersed by stirring. The dispersion was stored in cool condition overnight so that the poloxamer get completely dissolved and clear solution will be obtained. Finally, volume was adjusted to the requisite amount by using distilled water.

Gelation temperature and gelation time

The gelation temperature of the *in situ* gel formulation was measured by the inverted-tube method as reported previously (14). 2 mL of the DPZ gel formulation was transferred to a 10 mL test tube which was sealed and immersed into a thermostat water bath. The temperature of the bath was increased from 20 °C to 40 °C and a thermometer with a minimum scale of 0.1 °C applied to accurately determine the temperature change. When the upper surface of the formulation solution wouldn't move if the test tube was turned 90 degrees, the temperature was thought as the gelation temperature. Similarly, 2 mL of the DPZ gel solution was transferred to a 10 mL volumetric flask and then heated to 20 °C in a thermostat water bath. After which, the flask was placed into 32 °C thermostat water bath and the duration for all the liquid to change completely into immovable gel was immediately recorded with a stopwatch. The above time was identified as gelation time. The gelation temperature and gelation time were both determined in triplicate.

pH of gel formulation

The pH value of the gel solution was determined by using the pH meter, which was firstly calibrated prior to the test. Briefly, 1 mL of the prepared DPZ gel formulation was added into a 100 mL volumetric flask, then dilute with freshly distilled water to volume and mix. The pH value of the above solution was determined in triplicate.

In vitro erosion and release studies

In vitro corrosion and release of DPZ from the gel formulation were investigated simultaneously by the membraneless method (15). Briefly, 10 g of DPZ gel solution was transferred into a pre-weighted test tube, which was then placed in water bath (34±0.5 °C) and maintained for 5 min. After completion of sol-gel transition, 1.0 mL of pH 6.4 phosphate buffer pre-warmed at 34±0.5 °C, was gently added over the surface of the gel and the test tube was immediately shaken at 50 r min⁻¹ in a thermostat oscillator maintained at 34 °C. After dumping the medium at specified time interval, the test tube was cleared, weighted and replaced with the same volume and temperature of fresh phosphate buffer. The DPZ in the samples was determined by reported HPLC method with minor modification (16). The analysis was performed on a C₁₈ column (250×4.6 mm, 5.0 µm) at the column temperature of 30 °C. The mobile phase consisted of methanol and 20 mmol L⁻¹ ammonium acetate solution (containing 0.05 % formic acid) (55:45, V/V) and the flow rate was 1.0 mL min⁻¹. The injection volume was 20 µL and the detection wavelength was set at 271 nm.

Moreover, *in vitro* release studies were also performed to observe the DPZ sustained release from the *in situ* gel as compared with the drug solution by dialysis diffusion technique. Briefly, 2 mL of the *in situ* gel as well as drug solution equivalent to 100 mg of DPZ was placed in a dialysis bag (MWCO, 3500 Da) and then sealed at both ends. After which, the dialysis bags were placed in 500 mL of pH 6.4 phosphate buffer thermostatically maintained at 37±0.5 °C, and immediately stirred at speed of 100 rpm. At specific time intervals (5, 15, 30, 60, 120, 180, 240, 360 and 480 min), 2 mL of samples was withdrawn and filtered through a 0.45 µm syringe filter. The resultant filtrates were analyzed by above HPLC method.

Pharmacokinetics and brain targeting studies

Sprague-Dawley male rats weighing 250–320 g were selected for the study and an experimental protocol was approved by our Institutional Ethical Committee for animal and human tests (Grant No.YKD2018068). The rats were randomly divided into control and test groups. The control group received DPZ solution at a dose of 10 mg kg⁻¹ through intragastric (*i.g.*) route. The test group rats were firstly anesthetized with ether, after which, the trachea was cannulated with a polyethylene tube to stop nasal respiration. Meanwhile, an incision was made in the esophagus, after which, a length of polyvinyl chloride (PVC) pipe was inserted into it, the DPZ gel solution was then given at the same dose with a microliter syringe stretching into the back of the nasal cavity *via* the PVC pipe. The PVC pipe was subsequently pulled out and the esophagus ligated to prevent peroral absorption. For the control group, five rats per time point were sacrificed at 5, 15, 30, 45, 60, 90, 120, 180, 240, 360, 480 min following *i.g.* administration of the drug solution. Blood was collected and placed into heparinized PE tubes, immediately centrifuged at 10,000 rpm for 10 min and plasma separated. Meantime, the brain was isolated, rinsed with normal saline and weighed followed by homogenization using glass homogenizer with suitable amount of saline in ice-water bath. Homogenized brain tissue was centrifuged and the brain supernatant separated. For the test group, the same number of animals each time point were also sacrificed at 3, 5, 10, 20, 30, 45, 60, 90, 120, 180, 240 min after nasal administration of the DPZ *in situ* gel. The blood and brain tissues were pretreated as described above for *i.g.* dosing. Finally, the separated plasma and brain supernatant were stored at –20 °C until being assayed.

Plasma and brain sample analysis

In order to further improve the analytical sensitivity, the pretreatment process of blood and brain samples involving concentrating the sample and extracting the drug, was completed prior to analysis. For analysis of drug in plasma, 20 µL of 0.1 mol L⁻¹ NaOH solution and 300 µL of *n*-hexane/dichloromethane/ethyl acetate mixture (45:40:15, V/V/V) were successively added to 100 µL plasma sample and vortex-mixed for 3 min. After which, the mixture was centrifuged at 10,000 rpm for 10 min, and then the supernatant was transferred to a clean PE tube and evaporated to dryness under a nitrogen gas stream in a 40 °C water bath. The residue was then reconstituted in 400 µL methanol, and after centrifugation, 20 µL of the supernatant was injected for HPLC analysis. For drug analysis in brain homogenates, 20 µL of 0.1 mol L⁻¹ NaOH solution and 300 µL of methyl *t*-butyl ether were successively added to 100 µL of brain supernatant sample and vortex-mixed for 3 min. The mixture was also centrifuged at 10,000 rpm for 5 min, and then 200 µL of supernatant was transferred to a PE tube and evaporated to dryness. The residue were treated as described above for plasma samples and assayed for DPZ content. Finally, DPZ content in plasma and brain samples was determined according to a modified HPLC method (17). Chromatographic separation was achieved with a Inert Sustain C₁₈ column (150 mm×4.6 mm, 5 µm). The mobile phase consisted of 0.01 mol L⁻¹ sodium dihydrogen phosphate solution and acetonitrile in a ratio of 73:27 (V/V). The mobile phase was set at a flow rate of 1.2 mL min⁻¹. The eluate was monitored at 390 nm with an excitation wavelength of 325 nm. The concentration of DPZ in plasma or brain samples was estimated according to each calibration curve over the concentration range from 10.0 to 800.0 ng mL⁻¹.

Pharmacokinetic analysis was performed by a model-independent method and the mean concentrations of DPZ in plasma and brain samples were plotted against time. The peak plasma or brain concentrations (c_{\max}) as well as the time to reach these peaks (t_{\max}) were read as raw data. The area under the plasma or brain concentration-time curve (AUC_{0-t}) from time zero to the time of the last measured concentration (c_t) was calculated by means of the linear trapezoidal rule. The bioavailability (F_r) of the DPZ gel formulation relative to *i.g.* drug solution was calculated. Drug targeting efficiency (DTE %) of the *in situ* gel after nasal administration versus *i.g.* drug solution was estimated using the following equation(18):

$$\text{DTE (\%)} = (AUC_{\text{brain } 0-t} / AUC_{\text{plasma } 0-t})_{i.n.} / (AUC_{\text{brain } 0-t} / AUC_{\text{plasma } 0-t})_{i.g.} \times 100 \%$$

where AUC_{brain} and AUC_{plasma} is the area under the brain tissue concentration-time and plasma concentration-time curves after *i.n.* and *i.g.* administration, respectively.

RESULTS AND DISCUSSION

Preparation of in situ gel

Comprehensively taking into consideration the limitation of volume of human nasal cavity and clinical doses of oral preparations, the drug concentration of the DPZ gel formulation was established to be 5.0 % (*m/V*). As main excipients in the gel formulation, the ideal concentration of P407 as well as P188 was found to be 19.6 % (*m/V*) and 8.2 % (*m/V*), respectively, which was confirmed by our previous research (19). Owing to their higher aqueous solubility and better safety, several β -CD derivatives such as HP- β -CD, SBE- β -CD and DM- β -CD have been widely applied as transmucous absorption enhancers in nasal formulations. The *ex vivo* sheep nasal mucosa penetration test performed by our research group has confirmed that 3.5 % (*m/V*) HP- β -CD showed the best permeability enhancing effect among the above cyclodextrin enhancers (19). In addition, based on the concentration of ethylparaben often applied in oral and ophthalmic preparations, 0.05 % (*m/V*) of the preservative was used in the drug gel formulation. Finally, it needs to be emphasized that in the preparation process of the DPZ *in situ* gel solution, the above HP- β -CD used in the formulation, could also increase the solubility of the poor water-soluble ethylparaben by complexation reactions, which made it easier to dissolve the preservative in the HP- β -CD solution.

The polymers P407 and P188 are the most popular thermosensitive gelling agents used for the topical applications through the nasal, ocular and cutaneous routes and also have lower adverse effect on tissues compared with other stimuli-responsive gelling systems (20). Furthermore, 0.05 % ethylparaben is often used in oral and ophthalmic preparations and HP- β -CD used as additive in the intravenous injections. Based on the above analysis, it could be anticipated that the DPZ gel formulation may have lower toxicity and higher safety *in vivo*. Certainly, a study on the toxicity and human safety of the formulation still needs to be further investigated.

Gelation temperature and gelation time

Gelation temperature and gelation time are two important quality indexes for the DPZ thermoreversible *in situ* gel since they have been proved to be closely correlated with

the storage condition, convenience of administration, nasal residence time and *in vivo* drug bioavailability of *in situ* nasal gel. Gelation temperature value of the DPZ gel formulation was found to be approximately 32.5 °C within the nasal physiological temperature range 29–34 °C, thus suitable for thermosensitive nasal gel formulations. Furthermore, gelation time of the gel formulation was determined to be only about 40 s, suggesting rapid formation of immovable gel after its contact with nasal mucosa. In short, the above gelation temperature and gelation time are generally considered as optimal, which might reduce or even avoid loss of drug due to transportation of the formulation to the nasopharynx and then to the gastrointestinal tract.

pH of formulation

The pH of the gel formulation was determined after suitable dilution because of the following reasons. Firstly, the gel formulation has relatively higher viscosity, so it was not easy to obtain a stable readings of pH meter in testing process. Secondly, considering the high rate of nasal mucus turnover, for example, resting-state daily production, 75–150 mL, thus the dose volume of the gel solution may be diluted in nasal cavity by at least 100 times following nasal dosing. Finally, according to the chemical structures of the drug and additives in the gel formulation, pH of the gel formulation theoretically seems to be nearly neutral. Therefore, diluting the gel sample by distilled water may have insignificant effect on the pH of the gel solution.

Previous research performed by our group has confirmed that DPZ had high oil-water partition coefficient (lipophilicity) and better chemical stability in an approximately pH 7.0 aqueous media, and furthermore ethylparaben also possesses relatively good bacteriostatic effect in the pH range 4.0–8.0. Additionally, human nasal physiological pH range was reported to be within 5.5–6.5 and it has been proved that the nasal mucosa can tolerate pH range of 3–10 (21) and greater deviation from nasal physiological pH could lead to severe irritancy or damage to mucosa. Thus, taking the above factors into consideration, neutral pH value of the gel formulation seemed to be appropriate. Since the pH value of the prepared gel formulation was determined to be about 7.0, it is unnecessary to adjust the pH value of the gel solution in the final preparation process of the DPZ *in situ* gel.

In vitro erosion and release studies

The membraneless model was applied to investigate the gel corrosion and DPZ release from the thermosensitive gel formulation in our research as this model is quite similar to the *in vivo* condition. The concentrations of DPZ in the release media were calculated from the standard calibration curve ($Y = 69970c - 127740$, $R^2 = 0.999$, concentration range of 0.8–200.0 $\mu\text{g mL}^{-1}$) of DPZ and the cumulative release of drug as well as gel corrosion was plotted against time. As shown in Fig. 1, the *in vitro* cumulative erosion and cumulative drug release were both approximately 90 % in 45 min. The drug release from the gel formulation seemed rapid, which might be mainly attributable to the use of big volume of release media in the corrosion and drug release study, for example, 1.0 mL of phosphate buffer each time, which far larger than the actual volume of nasal mucus, was added over the surface of the gel. Furthermore, the sample-containing test tube was shaken at higher speed, resulting in the faster gel corrosion. The *in vitro* erosion and *in vitro* drug release kinetics of the DPZ *in situ* gel were both fitted to the zero order model, and moreover, there

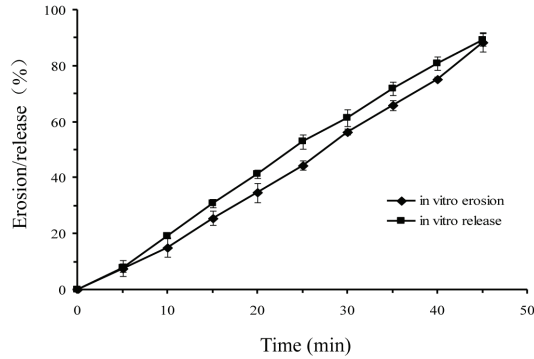


Fig. 1. The *in vitro* erosion and *in vitro* drug release kinetics of DPZ *in situ* gel with membraneless method. Data are represented as mean \pm SD ($n = 3$).

was a good correlation relationship between them ($R^2 = 0.9918$), which indicated that release of DPZ from the gel formulation was mainly controlled by gel corrosion.

Meanwhile, the DPZ sustained release from the gel formulation was also investigated by dialysis membrane diffusion method. An obvious drug sustained-release characteristics was observed in Fig. 2, for example, the cumulative release of DPZ from the gel formulation was only less than 80 % till 480 min, and 85 % of the drug, however, was released from the drug solution within 120 min. To explain the mechanism of DPZ release from the gel formulation, the *in vitro* release data of the drug were kinetically evaluated by using mathematical models. The order of the correlation coefficient (R^2) values of model fitting was found to be Weibull model > Riger-Pappas model > Higuchi model > First-order model > Zero-order model. The best fitted model was the Weibull model ($R^2 = 0.9947$), suggesting that the drug release from the gel formulation was controlled by drug diffusion as well as gel corrosion, probably due to the use of semi-permeable membrane in the *in vitro* release test.

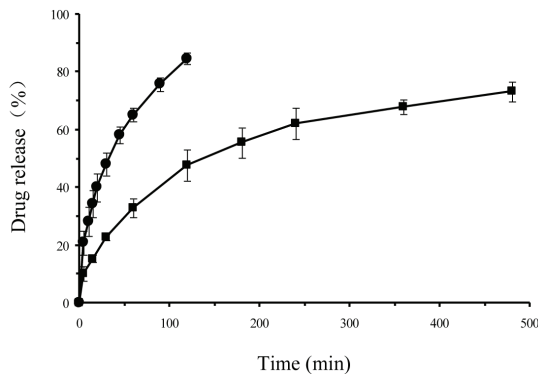


Fig. 2. The *in vitro* release profiles of DPZ *in situ* gel (squares) as well as the drug solution (circles) with dialysis diffusion method. Data are represented as mean \pm SD ($n = 3$).

In vivo pharmacokinetics and brain targeting characteristics

For the test group rats, after nasal dosing of the DPZ gel solution via a microliter syringe, the esophagus was ligated in order to prevent the gel formulation from entering GI tract and further avoid peroral absorption. As a result, the administered dose of the drug for nasal absorption might be accurate. Moreover, since nasal solution has shorter residence time relative to nasal *in situ* gel, a comparison with nasal DPZ solution has not been implemented in our research.

DPZ concentrations in plasma and brain after nasal administration of the *in situ* gel and *i.g.* drug solution to rats were calculated from each calibration curve equation. For the quantitative analysis of DPZ in plasma samples, the obtained calibration equation ($A = 111.14c + 10000$, $R^2 = 0.9982$) showed good linearity over the concentration range of 10.0–800.0 ng mL⁻¹. Accuracy of DPZ determination was found to be in range of 96.94–103.50 %. Intra-day and inter-day precision values were both lower than 5.0 %. Mean extraction recovery of DPZ in rat plasma at 50.0, 200.0, 600.0 ng mL⁻¹ were 97.92, 104.46 and 98.40 %, respectively. Additionally, as for the analysis of drug in brain samples, the linearity range was also from 10.0 to 800.0 ng mL⁻¹ with correlation coefficient of 0.9994 and the standard curve equation was $A = 105c + 4888.2$. Intra-day and inter-day precision values were both less than 2.0 %. Accuracy of DPZ determination was found to be in range of 96.29–104.28 %. Mean extraction recovery of DPZ in rat brain at 50.0, 200.0, 600.0 ng mL⁻¹ was between 97.12 and 104.43 %. In short, the above established methods were suitable for the determination of DPZ in plasma as well as brain tissues.

The mean plasma concentration-time curves of DPZ after nasal administration of the gel formulation and *i.g.* drug solution at dose of 10 mg kg⁻¹ and obtained pharmacokinetic parameters are shown in Fig. 3 and Table I, respectively. It could be found that the drug following nasal dosing was more rapidly absorbed from the *in situ* gel than from the drug solution after *i.g.* administration. The obtained c_{\max} of the drug *in situ* gel was 4113.41 ng mL⁻¹, which was approximately 15 times higher than that for the *i.g.* drug solution. Mean-time, t_{\max} for the nasal route was found to be only 10 min, which was much less than 55

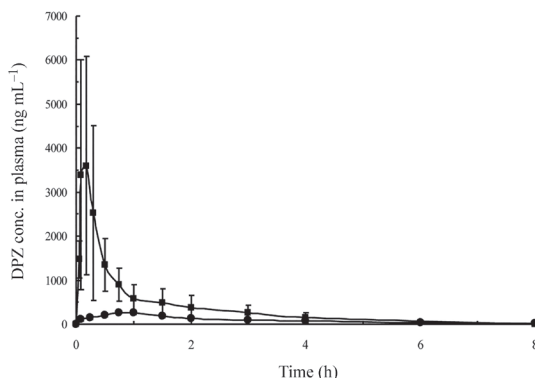


Fig. 3. Mean plasma concentration-time curves of DPZ after nasal administration of *in situ* gel (squares) as well as intragastric drug solution (circles) at the dose of 10 mg kg⁻¹ to rats. Data are represented as mean \pm SD ($n = 5$).

min for *i.g.* route. Furthermore, when the *AUCs* of both the dosing route were compared, the DPZ gel formulation had higher *AUC* value, for instance, $AUC_{0-\infty}$ for nasal route was up to 3010.10 ng h mL⁻¹, $AUC_{0-\infty}$ for the latter route, however, was only 780.66 ng h mL⁻¹. Therefore, there was significant difference in t_{max} , c_{max} and *AUC* values between the nasal gel formulation and the *i.g.* drug solution ($p < 0.01$). The relative bioavailability of the DPZ gel formulation was calculated to be up to 385.6 %, suggesting significantly enhanced bioavailability as compared to *i.g.* route. Compared to *i.g.* drug solution, the rapid absorption rate and significantly improved bioavailability of the DPZ *in situ* gel for nasal delivery in rats may be attributed to avoidance of hepatic first pass metabolism and enhanced drug permeability through the nasal mucosa by HP- β -CD. Moreover, the prolonged nasal residence time of the gel formulation also could result in better bioavailability. It also needs to be pointed out that the above results were in agreement with previous report by Al Asmari *et al.* (11), which proved that nasally administered liposomal formulation of DPZ showed significantly increased bioavailability in rats. The gel formulation in our study, however, have more advantages such as simple formulation, easy preparation process, better physicochemical stability, long nasal residence time over the liposomal formulation. Furthermore, the DPZ *in situ* gel could tightly adhere to the nasal mucosa and thereby prolonged the release and contact time. Thus, the use of the gel formulation resulted in higher systemic bioavailability as compared with the liposomal formulation.

As depicted in Fig. 4 and Table II, the peak concentration of DPZ in brain (c_{max}) and the time to reach it (t_{max}) following nasal administration of the *in situ* gel to rats were found to be 11218.34 ng mL⁻¹ and 14 min, respectively, for the *i.g.* drug solution, however, the values of c_{max} and t_{max} were 757.85 ng mL⁻¹ and 90 min, respectively, suggesting rapid and greater transport of DPZ into the brain ($p < 0.01$). Moreover, it was observed that DPZ concentration in the brain after intranasal *in situ* gel was far higher at almost all time points than after the *i.g.* drug solution. The AUC_{0-t} of DPZ in brain for the gel formulation was found to be more than five times greater compared with *i.g.* drug solution ($p < 0.01$). The above results demonstrated better bioavailability of DPZ in brain after nasal administration of the *in situ* gel compared to *i.g.* drug solution. Meanwhile, the DTE value was calculated to be 151.16 %, which was greater than 100, also suggesting superior drug delivery to the brain with the *in situ* gel. Our findings were similar to the results of the previous studies showing increased brain bioavailability of liposomal DPZ and the drug nanosuspension following intranasal dosing in rats (10, 11). Moreover, the brain targeting of nasally administered

Table I. Main pharmacokinetic parameters of donepezil hydrochloride in plasma after nasal administration of the *in situ* gel to rats using intragastric drug solution as control. Data are represented as mean \pm SD (n = 5).

Parameter	Drug solution (<i>i.g.</i>)	<i>In situ</i> gel (<i>i.n.</i>)
t_{max} (min)	55 \pm 7.75	10 \pm 5.48 ^a
c_{max} (ng mL ⁻¹)	269.35 \pm 27.06	4113.41 \pm 2729.68 ^a
AUC_{0-t} (ng h mL ⁻¹)	735.68 \pm 108.41	2718.25 \pm 1373.59 ^a
$AUC_{0-\infty}$ (ng h mL ⁻¹)	780.66 \pm 127.71	3010.10 \pm 1705.56 ^a
F_r		385.58 %

^a $p < 0.01$, compared to the control; F_r – bioavailability relative to intragastric drug solution.

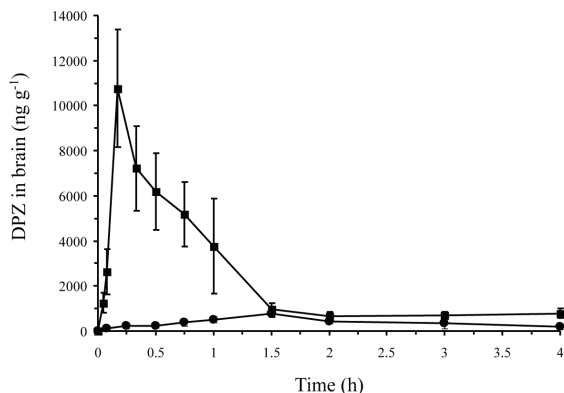


Fig. 4. Mean brain concentration-time curves of DPZ after nasal administration of the *in situ* gel (squares) and intragastric drug solution (circles) at the dose of 10 mg kg^{-1} to rats. Data are represented as mean \pm SD ($n = 5$).

DPZ was also confirmed by the pharmacokinetic study accomplished in rabbits, which indicated that intranasal delivery of the drug through liposomal hydrogel increase the brain intake of the drug compared to the oral route (12). There can be several reasons for the increased brain delivery for the DPZ gel formulation. Firstly, one possible reason may be that the better bioavailability and the greater plasma concentrations of DPZ with the gel formulation could result in more easily diffusing of the drug across the BBB due to the higher concentration gradient from the systemic circulation to the brain. In this respect, HP- β -CD might play an indirect role in improving brain bioavailability of DPZ gel formulation since it could promote the absorption of the drug across nasal mucosa and therefore significantly improved the plasma concentration and the bioavailability of the drug in plasma, which eventually led to enhanced transport of the drug from the blood system into the brain. Another possible reason may be that P188 in the gel formulation probably acted as absorption enhancers and furthermore inhibited the efflux system, especially *P*-glycoprotein, also increasing the transport across BBB. Finally, the preferential delivery into brain of the *in situ* gel may result from the direct transport of DPZ into brain *via* the olfactory or trigeminal nerves.

Table II. Main pharmacokinetic parameters of donepezil hydrochloride in brain after nasal administration of the *in situ* gel to rats using intragastric drug solution as control^a

Parameters	Drug solution (<i>i.g.</i>)	<i>In situ</i> gel (<i>i.n.</i>)
t_{max} (min)	90 ± 0.00	14 ± 5.48^b
c_{max} (ng g^{-1})	757.85 ± 137.82	11218.34 ± 1953.25^b
AUC_{0-t} (ng h g^{-1})	1581.32 ± 353.56	8832.22 ± 1397.13^b
DTE (%)		151.16

^a Data are represented as mean \pm SD ($n = 5$). ^b $p < 0.01$, compared to the control; DTE – drug targeting efficiency.

Due to the lack of nasal spray pumps and specific instruments, the aerosol performance of the gel formulation related to the several important parameters including droplet size, spray pattern, deposition pattern, *etc.* was not accomplished in this study, thus our research group wishes to complete it in near future.

CONCLUSIONS

DPZ thermosensitive *in situ* gel for nasal delivery was prepared by using P407 and P188 as main gel materials. The gel formulation exhibited ideal *in vitro* properties, such as suitable gelation temperature, short gelation time, approximately neutral pH, obvious sustained drug release. Moreover, pharmacokinetic studies confirmed that the bioavailability of the drug *in situ* gel formulation after nasal administration to rats was significantly increased compared to the drug solution after *i.g.* dosing. Brain targeting investigation also proved the remarkably enhanced transport of DPZ into the brain by intranasal *in situ* gel formulation. Thus, it is anticipated that the DPZ thermosensitive nasal *in situ* gel will be an ideal novel delivery system for the treatment of AD's disease due to lots of advantages over the other new delivery systems of the drug such as simple formulation, easy preparation process, convenient dosing, rapid onset of action, decreased dosing frequency, high bioavailability, significant brain targeting and potentially improved therapeutic efficacy. Finally, the pharmacokinetics and brain targeting of the DPZ *in situ* gel formulation in humans as well as its toxicity or irritation to animal or human nasal mucosa are unknown, so a further research is also necessary and furthermore very important.

Acknowledgements. – This study was supported by Grants from the Natural Scientific Foundation of Inner Mongolia (Grant No. 2018MS 08107).

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