

## Research Article

# Formulation and evaluation of matrix microspheres for simultaneous delivery of salbutamol sulphate and theophylline

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## Abstract

**Purpose:** The objective of this study was to formulate and evaluate a matrix microsphere system for simultaneous delivery of two anti-asthmatic drugs Salbutamol sulphate and Theophylline which are often indicated for the management of asthma, their frequent dosing may reduce compliance, thus making a prolonged release formulation necessary. Ethylcellulose was used as a retardant polymer and its compatibility with the drugs and the drug-drug compatibility were established through IR, XRD and DSC studies.

**Method:** Microspheres were prepared by emulsion solvent evaporation using acetone/light liquid paraffin system. Tween 80 was used as the dispersing agent and cyclohexane was added to harden the microspheres. The prepared microspheres were characterized for their micromeritic properties and drug loading, as well as by infrared spectroscopy (IR), differential scanning calorimetry (DSC), x-ray powder diffractometry (XRD) and scanning electron microscopy (SEM). The *in vitro* release studies were performed in pH 7.4, phosphate buffer.

**Result:** The prepared microspheres were white, free flowing and spherical in shape. The drug-loaded microspheres showed 67-91% of entrapment and release was extended upto 6 to 8 h. The infrared spectra, differential scanning calorimetry thermographs and XRD spectra all showed the stable character of both the drugs in the drug-loaded microspheres and revealed the absence of drug-polymer interactions. Scanning electron microscopy study revealed that the microspheres were spherical and porous in nature.

**Conclusion:** The matrix microspheres have a potential for the prolongation and simultaneous delivery of the anti asthmatic drugs (salbutamol and theophylline).

**Keywords:** Matrix microspheres, Salbutamol sulphate, Theophylline, Ethylcellulose, Emulsion solvent evaporation technique.

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## INTRODUCTION

Patients suffering from chronic diseases like asthma, diabetes and epilepsy may have to take drugs everyday for the rest of their life<sup>3</sup>. WHO estimates the number of asthmatic patients to be around 100 to 150 millions around the world and India contribute 10 % of the total and its incidence is escalating every decade at an alarming rate. In management of chronic diseases like asthma, compliance to the dosage regimen is the key to a successful therapy. Patient may be treated with more than one drug and compliance is found to be low in such cases.

The short half life (4 to 6 hours) with extensive first pass metabolism of salbutamol and the propensity for interaction and narrow therapeutic index (10 to 20 µg/ml) of theophylline are well known. Although salbutamol and theophylline are often indicated for the management of asthma, their frequent dosing may reduce compliance, thus making a prolonged release formulation necessary. Theophylline produced an additive effect when used in combination with salbutamol sulphate<sup>4</sup>. Different controlled release dosage forms for simultaneous delivery of salbutamol and theophylline have been proposed by different authors. These include transdermal patches<sup>5</sup>, Matrix tablets<sup>6</sup> and Osmotic pump tablets<sup>7</sup>. Both salbutamol and theophylline are readily and well absorbed along the gastrointestinal tract. Even when salbutamol is given as an inhalation, it has been suggested that majority of the dose is swallowed and absorbed from the gut<sup>8</sup>. Microencapsulation is used to modify and retard drug release. Microspheres, due to its small particle size, are widely distributed throughout the gastrointestinal tract which improves drug absorption and reduces side-effects due to localized build-up of irritating drugs against the gastrointestinal mucosa<sup>9</sup>. The objective of the present study includes: (a) formulation of a sustained release matrix microsphere system containing salbutamol

sulphate and theophylline using ethylcellulose as the retardant polymer which will release the drug at the gastrointestinal tract for a prolonged duration to promote patient compliance; (b) to study the effects of drug to polymer ratio on in-vitro drug release and (c) to fit the drug release data to various drug release models.

## MATERIALS AND METHODS

### Materials

The test drugs, salbutamol sulphate IP and anhydrous theophylline IP (Ducbill drugs, Kolkatta), the polymeric matrix former ethylcellulose (22 cps grade determined at 80:20 toluene: ethanol, Wilson Brothers, Mumbai), and the dispersing agent Tween 80 (Rankem, New Delhi) were obtained and all other chemicals and reagents used were of analytical grade.

### Determination of Saturation solubility

The saturation solubility of both Anhydrous theophylline and salbutamol sulphate were separately determined at distilled water and phosphate buffer pH 7.4 to observe the suitability of the solution as a dissolution medium (Table 2). Excess amount of the drugs were taken and dissolved in a 1 ml of phosphate buffer (pH 7.4) and distilled water and extracted for 24 hours. After suitable dilutions, the concentrations were determined spectrophotometrically at 276 nm for salbutamol sulphate and 270.5 nm for Theophylline<sup>10</sup>.

### Preparation of microspheres

The microspheres were prepared by emulsion solvent evaporation technique using the formulation as shown in Table 1. In this method 1219.2 mg of ethylcellulose was dissolved in 15 ml of acetone and a given amount of the drugs were dispersed in it to make different drug to polymer ratio of 1:1, 1:2 and 1:3 and stirred for about 10 minutes. Then the polymer drug dispersion was poured into 50 ml of liquid paraffin (light) containing varying concentrations of Tween 80. The whole system was then stirred for about 4 hours at 900 RPM. After stirring process is over the liquid paraffin (light) was decanted off and the microspheres formed were collected

by filtration in a Whatman filter paper and washed with Cyclohexane to completely

$$\text{Drug entrapment efficiency} = \frac{\text{Experimental drug content}}{\text{Initial drug content in the formulation}} \times 100 \quad \dots 1$$

remove the remaining oil and dried at 50 °C in Vacuum drier (NSW, India) for 6 hours and collected for further studies.

#### **Particle size determination**

The particle size of the microspheres was determined by microscopic method<sup>11</sup>. For each batch of the microspheres, 100 particles were counted and done in triplicate.

#### **Drug entrapment efficiency**

The amounts of salbutamol sulphate and theophylline present in the microsphere were determined by extraction in distilled water<sup>12</sup>. 50 mg of the crushed and powdered microsphere was taken and extracted in 50 ml of distilled water and stirred for 15 minutes at 1500 rpm. The solution was filtered and diluted with 0.05 M NaOH and absorbance

$$\text{Drug entrapment efficiency} = \frac{\text{Experimental drug content}}{\text{Initial drug content in the formulation}} \times 100 \quad \dots 2$$

measured spectrophotometrically (U-2001, Hitachi) at 244.6 nm and 276.5 nm and concentrations were determined by employing simultaneous equations method<sup>10,5</sup>.

#### **In vitro drug release study**

The in-vitro release study of the microsphere was carried out using USP rotating basket method at 50 rpm at 37 °C. Dissolution study was performed in Phosphate buffer pH 7.4 taking 900 ml for each study. 100 mg of the microsphere was placed in the dissolution medium and test samples were taken from the medium at predetermined time intervals over a period of 12 hours and the samples were analyzed for salbutamol sulphate and theophylline content UV spectrophotometer.

#### **Infrared Spectroscopy**

IR spectra of the pure drugs and microspheres were recorded using Perkin-

Elmer model 883 IR-spectrophotometer between the ranges of 500 to 4000 cm<sup>-1</sup> by making a pellet of the samples with KBr. The resultant spectra were then compared with standard reference (IP 1996) and observe for any type of deviation from the standard.

#### **Differential Scanning Calorimeter Analysis (DSC)**

DSC thermogram of the pure drugs and the microspheres were recorded with a differential scanning calorimeter (Universal V2.5H TA Instrument) from 20 to 550 °C at a heating rate of 20 °C/minute.

#### **X-Ray Diffraction Spectroscopy (XRD)**

X-ray diffraction spectrum of the pure drug and microspheres were recorded with Phillips PW 1830 X-ray generator fixed with PW 1710 diffractometer (Phillips Industrial & Electro-acoustic Systems Division, Almelo, The Netherlands). The XRD was performed at the angle between 5-60 ° 2θ.

#### **Scanning Electron Microscopy (SEM)**

Scanning electron microscopy (Hitachi S-3600N, Japan) was done to characterize surface topography of the microspheres. Photomicrograph of the microspheres before and after the release of drugs was taken. The quality of the microspheres (with respect to surface properties) and the nature and size of pores developed on the surface can be studied. The changes that occur during in-vitro dissolution studies may have implications to the performance of the microspheres.

#### **Release Kinetics**

Data obtained from in vitro release studies were fitted to various kinetic equations to find out the mechanism of drug release from the ethylcellulose microsphere. The kinetic models used were:

$$Q_t = k_0 t \quad (\text{zero-order equation}) \quad \dots 3$$

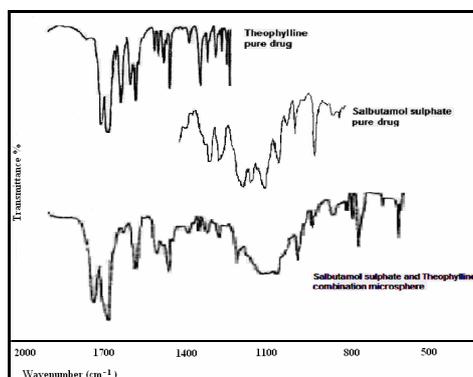
$$\ln Q_t = \ln Q_0 - k_1 t \quad (\text{first-order equation}) \quad \dots 4$$

**Table 1:** Formulation, drug entrapments and mean particle size of microspheres

Formulation code	Salbutamol sulphate (%)	Theophylline (%)	Mean particle size (µm)	Amount of polymer (mg)	Amount of theophylline (mg)	Amount of salbutamol sulphate (mg)
T1	78.667±2.45	82.862±1.75	730.55±1.582	1219.2	1200	19.2
T2	76.534±3.67	83.468±1.32	689.71±1.281	1219.2	600	9.6
T3	83.45±1.25	86.789±0.986	695.64±2.565	1219.2	400	6.4

**Table 2:** Correlation coefficient ( $r^2$ ), Constant (K) and Diffusion exponent (n) after fitting of dissolution data into various release kinetic models Theophylline

Formulation (Drug:Polymer)	Zero-Order		First-Order		Higuchi Model		Korsenmeyer-Peppas Model	
	$r^2$	$K_0$	$r^2$	$K_1$	$r^2$	$K_h$	$r^2$	n
Theophylline								
1:1	0.7406	6.9769	0.9707	0.104	0.9842	26.287	0.9907	0.2506
1:2	0.7908	5.5131	0.9687	0.068	0.9523	23.334	0.9926	0.2798
1:3	0.8461	4.4599	0.955	0.037	0.9782	18.44	0.9902	0.2334
Salbutamol Sulphate								
1:1	0.767	9.5228	0.9579	0.213	0.9525	33.335	0.9811	0.4225
1:2	0.7951	6.8991	0.9873	0.131	0.9879	27.889	0.9919	0.4447
1:3	0.8671	4.8203	0.9636	0.043	0.9763	19.903	0.9973	0.3037



**Figure 1:** IR spectra salbutamol sulphate (pure drug), theophylline (pure drug) and microsphere containing the two drugs (1:3 drug:polymer).

$$Q_t = K.S.\sqrt{t} = k_H.\sqrt{t} \dots\dots\dots 5$$

(Higuchi eqn based on Fickian diffusion)

Where,  $Q$  is the amount of drug release in time  $t$ ,  $Q_0$  is the initial amount of drug in the microsphere,  $S$  is the surface area of the microcapsule and  $k_0$ ,  $k_1$ , and  $k_H$  are rate constant of zero order, first order and Higuchi rate equations respectively. In addition to these basic release models, there are several other models as well. One of them is Peppas and Korsenmeyer equation<sup>13,14</sup> (power law).

$$M_t / M_\infty = k \cdot t^n \dots\dots\dots 6$$

Where  $M_t$  is the amount of drug release at time  $t$  and  $M_\infty$  is the amount release at time  $t = \infty$ , thus  $M_t / M_\infty$  is the fraction of drug released at time  $t$ ,  $k$  is the kinetic constant, and  $n$  is the diffusion exponent which can be used to characterize both mechanism for both solvent penetration and drug release. Determining the correlation coefficient assessed fitness of the data into various kinetic models. The rate constants, for respective models were also calculated from slope.

## RESULTS

The physicochemical compatibility of the drugs and the polymer was established through infrared spectroscopy (Figure 1), differential scanning calorimetry and x-ray diffraction spectroscopy. Results from IR, DSC and XRD studies do not show any interaction or degradation.

### Scanning Electron Microscopy (SEM):

Scanning electron microscopy of drugs-loaded ethylcellulose microspheres (Figure 2) reveals that the microspheres possess a rough and rugged surface. The micrograph taken after 12 hours release studies also reveals porosity developed at the surface. The micrographs show that porosity developed but structure is retained as 12 hours drug release study was carried out.

**Entrapment Efficiency:** The entrapment efficiency determined at phosphate buffer of pH 7.4 shows the entrapment efficiency of both drugs increase as the polymer concentration is increased (Table 1). The entrapment efficiency for salbutamol sulphate was in the range of 78 (for 1:1 drug:polymer)

to 83 (for 1:3 drug:polymer) and for theophylline it was in the range of 82 (for 1:1 Drug:Polymer) to 86 (for 1:3 Drug:Polymer).

**Mean Particle Size:** The particle size of the microspheres (Table 1) was in the range of 690  $\mu\text{m}$  to 730  $\mu\text{m}$ .

**Drug Release:** The effect of variation in drug to polymer ratio on drug release was studied on salbutamol sulphate and theophylline combination microspheres in a dissolution medium of phosphate buffer of pH 7.4. Different drug to polymer ratio taken were 1:1, 1:2 and 1:3 and increase in polymer concentration resulted in a decrease of drugs release rate. The release of salbutamol sulphate was found to be higher as compared to theophylline release. Sustained release up to 12 hours was achieved when drug: polymer was taken up to 1:3.

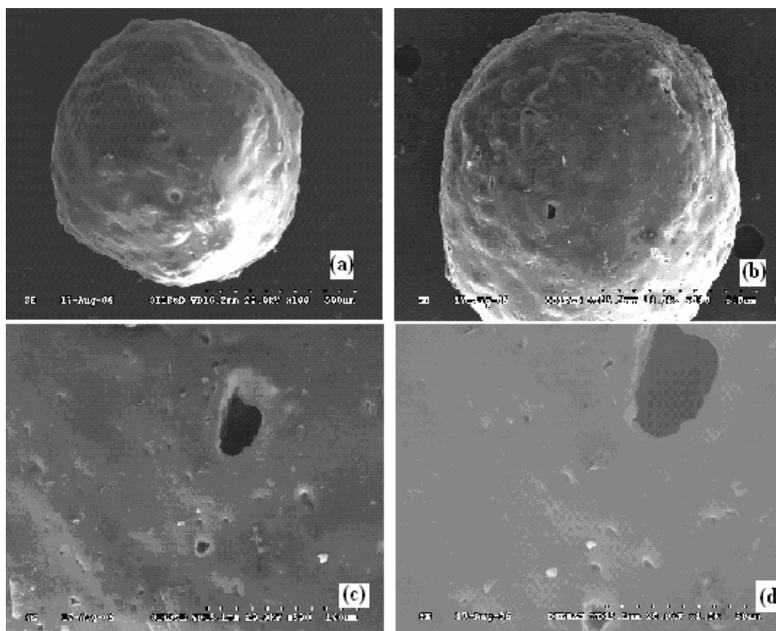
The release data obtained were fitted into various release kinetic models and release kinetics table (Table 2) was prepared.

## DISCUSSION

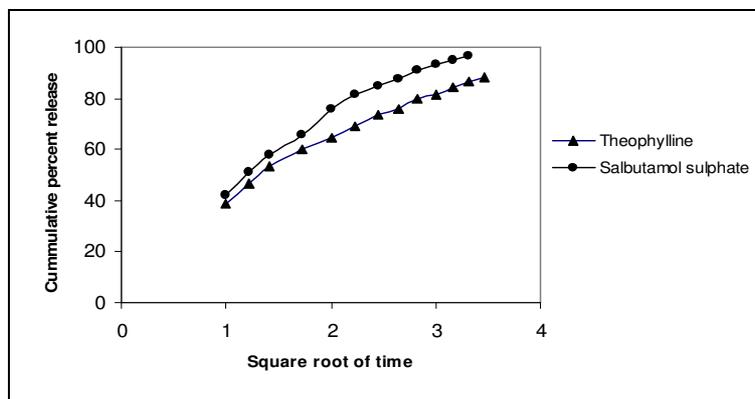
A matrix microsphere system for sustain release of two drugs has been formulated. For this type of system it is important that the presence of one drug species do not alter the physicochemical properties of the other<sup>13</sup>.

Scanning electron microscopy of drugs-loaded ethylcellulose microspheres reveals that the microspheres possess a rough and rugged surface. It was also observed that after dissolution was performed, bigger and more pores are developed. The surface porosity is crucial for drug release in microspheres prepared with ethylcellulose. Since the polymer is not biodegradable, the release of the drugs from microspheres takes place by dissolution and diffusion through the pores. Ethylcellulose allows water to permeate through its surface without itself dissolving in it. The micrographs show that porosity developed but structure is retained as 12 hours drug release study was carried out.

The entrapment efficiency of both drugs was found to increase as the polymer concentration is increased, as there were more polymeric binding sites for the drugs. And as the amount of drug taken is decreased



**Figure 2:** Scanning electron micrographs of salbutamol sulphate-theophylline combination microsphere (1:3 drug:polymer) (a) before drug release; (b) after 12 hours of drug release; (c) and (d) show magnifications of (b).



**Figure 3:** Higuchi plots for drug release from microsphere (1:2, drug:polymer, 1% Tween 80)

while taking constant amount of the polymer, the particle size is decreasing. This may be due to the decrease in viscosity of the drug-polymer dispersion since constant amount of polymer was dissolved in acetone and the amount of drug taken was reduced to obtain a higher polymer concentration<sup>15</sup>.

The effect of variation in drug to polymer ratio on drug release was due to the increase in wall thickness that results in longer diffusion path<sup>16</sup>. The release of salbutamol sulphate was found to be higher as compared to theophylline release, this may be due to the higher solubility of salbutamol sulphate than theophylline in the dissolution medium.

Dissolution studies show a sustained release upto 12 hours when drug:polymer was taken up to 1:3 this may be due to an increase in the tortuosity of the diffusion pathlength as the amount of the polymer is increased. From the release data obtained and the release kinetics table (Table 2) it can be observed that the best-fit release kinetics was achieved with Higuchi plot followed by zero order and First order. The release of both the drugs was influenced by the drug to polymer ratio and was found to be diffusion controlled. So, the release of salbutamol sulphate and theophylline from the ethylcellulose microspheres was found to exhibit diffusional characteristics and closely follows Higuchi Model and also highly correlated with first-order release model (Figure 3).

## CONCLUSION

Salbutamol sulphate-theophylline combination microspheres was prepared successfully using the emulsion solvent evaporation method. Taking of different polymer:drug ratio is found to influence the size, entrapment efficiency and release characteristics of the microspheres. The assessment of the release kinetics revealed that drug release from the microspheres followed Higuchi Model. The mechanism of drug release from microspheres was diffusion-controlled. Since the polymer and the drugs were found to be compatible and the release mechanism is characterized, there is a great scope for the formulation of these antiasthmatic drugs as a microsphere system. Since microspheres are better distributed and facilitate the absorption of drugs from the intestines, this type of formulation would help in controlling asthmatic attacks to millions of patients suffering from it.

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