

Development and Evaluation of Isoniazid Loaded Silk Fibroin Microsphere

Desarrollo y evaluación de microesfera de seda Fibroin cargado de isoniácida

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RESUMEN

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Objetivo: La investigación experimental en curso está dedicada a la preparación de microesferas de pequeño tamaño y buena esfericidad mediante el método de separación de fases con isoniazida (INH) como fármaco modelo. La fibroína de seda tiene cualidades intrínsecas únicas como la biodegradabilidad, biocompatibilidad o propiedades de liberación y su capacidad de carga de fármacos ajustable. La aptitud de entrega de carga de las moléculas de fármaco en las esferas de seda estar supeditada a su carga, y la hidrofobicidad o subsiguiente alteración en los perfiles de liberación de fármacos.

Métodos: En el presente trabajo la microesfera de fibroína de seda cargada de isoniazida fue preparada utilizando el método de separación de fases. La microesfera fue evaluada por espectroscopia ultravioleta-visible, espectroscopia infrarroja con transformado de Fourier, se midió la eficiencia de atrapamiento y se estudió mediante microscopía electrónica de barrido.

Resultados: Estudios con el microscopio de escaneo de electrones revelaron que las microesferas de fibroína cargada de isoniazida eran esféricas. La eficacia de atrapamiento de las microesferas de formulación diferente de F1 a F5 estuvo en el rango de 53 a 68 %. F3 mostró un 68,47 % de eficiencia de atrapamiento y tras optimizar la formulación de liberación de fármacos fue de 93,56 %, a las 24 horas.

Conclusión: Esta investigación reveló una nueva formulación de base acuosa para las esferas de seda con forma controlable o la forma y el tamaño de la esfera. Las microesferas de seda cargadas de isoniazida pueden actuar como ideal formulación nano con estudios elaborados.

Palabras clave: fibroin de seda; nanoformulation; biodegradabilidad; isoniácida.

ABSTRACT

Aim: Current experimental investigation is dedicated to prepare microspheres with small size and good sphericity by Phase Separation method using Isoniazid (INH) as model drug. Silk fibroin has unique intrinsic qualities like biodegradability, biocompatibility or release properties and their tunable drug loading capacity. The delivery loading proficiency of the drug molecules in silk spheres be contingent on their charge, and hydrophobicity or subsequent in altered drug release profiles.

Methods: In the present work Isoniazid loaded silk fibroin microsphere was prepared by using phase separation method. Microsphere was evaluated for Ultraviolet-visible spectroscopy, Fourier Transform infrared spectroscopy, Entrapment efficiency, Scanning electron microscopy Studies.

Results: Scanning electron microscopy studies revealed that Isoniazid Loaded Silk Fibroin Microspheres were spherical. Entrapment Efficiency of Isoniazid loaded Microspheres of different Formulation from F1 to F5 was in range of 53 to 68 %. F3 showed 68.47 % entrapment Efficiency and the optimized formulation drug release was 93.56 % at 24 hours.

Conclusion: Experimental report disclosed a new aqueous based formulation method for silk spheres with controllable shape or size and sphere. Isoniazid loaded silk microspheres may act as ideal nano formulation with elaborated studies.

Keywords: silk fibroin; nanoformulation; biodegradability; isoniazid.

INTRODUCTION

Drug delivery research focused on micro and nano based delivery have made strong presence in healthcare industry. Microspheres based controlled delivery of antimycobacterial agents might be proficient through employing a variety of polymeric drug carriers. Even though knowledge among natural polymers is encouraging and extensive and may offer lots of benefits.¹⁻⁵ Tuberculosis is most health crisis during the world or contaminate more than eight million persons every year. Oral therapy with the presently working anti tubercular drugs is extremely effectual and still related amid a number of important problems.⁶⁻⁹ Further than 80% of TB cases were of pulmonary TB unaided and soaring drug dose has essential to be managed since merely little slice of the entire dose reaches the lungs. Anti Tubercular drugs delivery system which could be delivered using the respiratory route alongside with canister evade the daily dosing since they can help out in: (i) Express drug delivery to the unhealthy person; (ii) The targeting of alveolar macrophages which have uses via the mycobacteria while a secure place for their extended survival; (iii) reduced systemic toxicity of the drugs; and (iv) improved patient compliance. Furthermore in distinction the oral route of administration and inhaled drugs were not subjected to first pass metabolism. It is not gratified for the reason that could not steadily reside in the lung.¹⁰⁻¹² Silk fibroin show advanced mechanical properties and tunable degradation rates ranging from weeks to months *in vivo* due to control of excellent biocompatibility, crystallinity with low inflammatory and immunogenic response, and all aqueous material processing alternative to form films, gels, fibers, microspheres and sponges. For drug delivery, particularly protein drugs, silk materials exhibit controllable drug release kinetics and high encapsulation efficiency due to the crystalline beta-sheet formation.¹³⁻¹⁶ There are several techniques available for the preparation of drug loaded microspheres, such as evaporation or extraction methods, self-assembly, solvent displacement, emulsion-solvent phase separation, and spray drying. A new approach has preferred to fabricate silk microspheres through convenient sphere size, size and avoid with organic solvents along with a few extra insensitive circumstances during processing.¹⁷⁻²¹ Phase separation among polyvinyl alcohol and silk take place impulsively while the two polymer solutions are assorted and afterward casted into films. The influence of blending of PVA and silk on top of silk secondary arrangement changes also film mechanical and swelling properties was studied. The present study has to develop the phase separation of silk or PVA blends to produce silk spheres with controlled microspheres, size intended for efficacy in drug loading and release.²²⁻²⁶ Nearly numeral of recompenses to evolving sustained release formulation using silk fibroin for targeted drug delivery. PVA

microspheres are most widely studied drug delivery systems for the controlled release of drugs that is vaccines, antihypertensive agents, antibiotics, peptide drugs, anti-cancer agents and Proteins.²⁷⁻³⁰ The focus of the current study was to prepare Silk fibroin microspheres by phase separation technique using polyvinyl alcohol (PVA).³¹⁻³² The inspirations of many particulate parameters stood investigated in relation to Microsphere performance and in-vitro drug release characteristics. The aim was to prepare Silk fibroin/PVA microspheres loaded with Isoniazid as drug delivery system which might enhance efficiency of Isoniazid aimed to local anti-tubercular activity and reduced the toxicity.

MATERIALS AND METHODS

The following materials were used: Isoniazid obtained from Balaji Drugs, Polyvinyl alcohol obtained from Fisher Scientific, Ethanol (Changshuyangyuan chemicals, China), Calcium chloride obtained from Avarice and Silk Cocoons collected from Himachal Pradesh. All reagents used were of analytical grade.

Silk fibroin purification

Silk fibroin aqueous stock solutions have been prepared from cocoons of *Bombyxmoriouter*. Silk cocoons was boiled for 30 min in 0.02 M sodium carbonate and after that rinsed meticulously with distilled water. The Silk Fibroin extracted was dissolved in ternary solvent mixture comprises of CaCl₂ Ethanol: Water in the ratio of 1:2:8 respectively and boiled for 4 hour at 70 °C to form aqueous stock solution.³³ Synthesis of Silk fibroin purification shown in figure 1

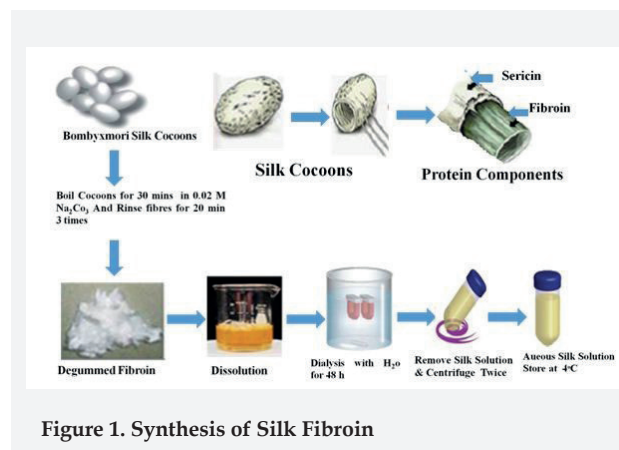


Figure 1. Synthesis of Silk Fibroin

Preparation of Poly Vinyl alcohol Solution:

A 5% (wt/vol) PVA solution by adding 0.25 g of PVA to 4 ml of deionized water and heated at 60 °C to dissolve the PVA.

Preparation of Isoniazid loaded Silk Fibroin Microspheres:

Microspheres were prepared using Phase separation method. The Microspheres were prepared by using different parts of Silk Aqueous Solution: Poly Vinyl alcohol Solution

and Isoniazid drug were selected as independent variables. Initially Parts of Silk aqueous solution was kept in beaker under stirring at room temperature. Consequently, Poly Vinyl alcohol Solution was dissolved in Silk aqueous solution to obtain a known concentration according to the formulation design magnetic stirring at room temperature for 45 to 60 mins and sonicate the solution for 30 s at 25% amplitude. Pour the solution into a Petri dish and allow it to cover the bottommost of the dish consistently. Dry the Liquid solution over-night in a fume hood. The film be able to kept for a insufficient weeks. Seal and cover the dish with Parafilm and store. Peel off the film from the dish and place one film in a 50-ml conical flask or tube. Add 20 ml of deionized water to the flask or tube and shake for 30 min at room temperature to dissolve the film. Centrifuge at 13,000 r.p.m. for 25 min at 4 °C and discard the supernatant. Resuspend the pellet in 5 ml of deionized water and used the suspension for Microspheres.³⁴

Determination of Partition Coefficient:

An estimated partition coefficient was determined by shake flask method from the ratio (1:1, 1:2, 2:1) of solubility of isoniazid in n-octanol/chloroform to that in water.

A stock solution of isoniazid was prepared in n-octanol/ chloroform and water such that the maximum concentration of the test substance, in each phase, always remains below 1.2 absorbance during the experiment. The organic and aqueous phase was separate and aqueous phase was analyzed by UV spectrophotometric method at 262 nm and the partition coefficient was calculated as follows:³⁵

Partition Coefficient; $\log_{10} P$ n-octanol or chloroform/ water =

$$\log_{10} \frac{\text{Total amount of drug} - \text{Saturation concentration of drug in water}}{\text{Saturation concentration of drug in water}} \times 100$$

Saturation concentration of drug in water

Saturation solubility study:

In this saturation solubility study, an excess quantity of Isoniazid was placed in the vial containing 10 ml and fill with deionized water. The vial was kept for 24 hours and the amount of the drug dissolved was analyzed after 24 hours using UV spectrophotometrically at 262 nm.³⁶

Formulation Design: Shown in Table 1

Table 1. Formulation design based on factorial design

Formulation Code	Silk Aqueous Solution used in Parts	Poly Vinyl alcohol Solution Used in Parts	Isoniazid Drug (mg)
F1	1	1	10
F2	1	2.5	10
F3	1	4	10
F4	1.5	5.5	10
F5	2	3	10

Evaluation of prepared Silk Fibroin of Microspheres.

Scanning electron microscopy

The samples were used for SEM solid as well liquid up to 50 mg or 0.5 ml liquid can be analyzed The morphologies of silk spheres were imaged using a SEM model No JEOL 5400 made by japan.Gold Ions Coating Is done 5-6 Mins. Images were taken with the installed software.

Entrapment efficiency (EE) of Microspheres

The entrapment efficiency of Microsphere was calculated using an indirect entrapment method. The prepared Microspheres suspension was centrifuged for 30 minutes at

40000 rpm. Then the supernatant is analyzed for the free drug content. The concentration of RIF in the supernatant was determined by UV-Visible spectrophotometry at 262 nm. The calculation was done to determine the amount of drug untrapped in the supernatant. The amount of drug entrapped was calculated by subtracting amount of free untrapped drug from the total amount of Isoniazid taken. The drug entrapment efficiency (EE) of microspheres was determined by using following formula:

$$\text{Entrapment efficiency (\%)} = \frac{\text{Total quantity of drug} - \text{free untrapped drug}}{\text{Total quantity of drug}} * 100$$

Total quantity of drug

In-vitro drug release study using dialysis membrane

Preparation of dialysis membrane

The dialysis membrane (Mol wt 12000 KD) was soaked overnight in a buffer in order to open the pores of the membrane. After period of 24 hours dialysis membrane was taken out and was used for the *in-vitro* drug release.

In-vitro drug release of Isoniazid Loaded Microspheres

A dialysis membrane was used to monitor Isoniazid release from the Microspheres. Dialysis membrane with molecular weight cut off 12,000 KD was used to determine the release of isoniazid from Microspheres. For determining the *in-vitro* drug release the 7.4 phosphate buffer solution was prepared. The dialysis membrane was soaked overnight in a buffer in order to open the pores of the membrane. The dialysis membrane was taken out from the buffer and was tied at one end. The Prepared Isoniazid Microspheres was introduced into the membrane and the other end was also tied with thread so that the sample should remain in the membrane. The membrane was introduced in the buffer medium. At frequent time intervals 1 ml aliquot was withdrawn and after sufficient dilution was analyzed spectrophotometrically at 262 nm. The sample Withdrawn stayed replaced via an equal quantity of fresh 7.4 phosphate Buffer saline. The drug release profile is presented graphically.

RESULT AND DISCUSSION

UV Spectroscopy (λ max determination):

The Standard Solution of Isoniazid deionized water and pH 7.4 Phosphate buffer (10 μ g/ml) was Scanned (200-400) nm by using Double Beam UV-Visible Spectrophotometer (UV-1800) manufacturer by Shimadzu Corporation .The λ max was found to be 262 nm.

Melting point of Drug

Melting point was determined by thiele tube method and was found to be in the range of 170° - 172° C which is in accordance with reference value

Saturation solubility study

Isoniazid, showed a solubility of 210mg/ml, in deionized water, falling in the category of very soluble drug as per I.P. 2007

Partition Coefficient

A negative log P (-0.420) confirms the hydrophilic nature of isoniazid. The experimental solubility and partition coefficient values substantiate the suitable candidature of isoniazid for the presently proposed study on hydrophilic drugs.

Evaluation of prepared Silk Fibroin drug loaded Microspheres.

Scanning electron microscopy (SEM)

The morphologies of silk spheres were imaged using a SEM model No JEOL 5400 shown in Figure: 2

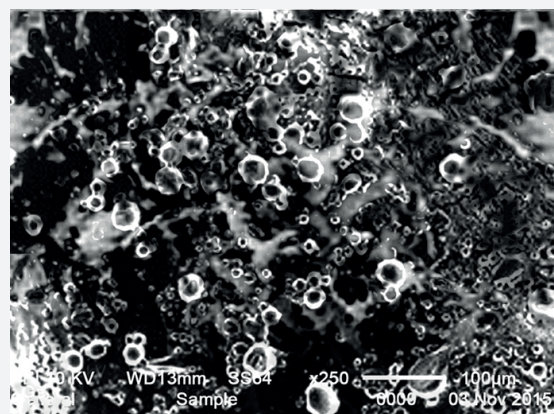


Figure 2. SEM Photomicrograph of silk microsphere

Entrapment efficiency (EE) of Isoniazid Loaded Microspheres.

The Entrapment Efficiency of Isoniazid loaded Microspheres were performed in triplicate different Formulation from F1 to F5 exposed in Table: 2 and Figure: 3

Table 2. Entrapment efficiency (EE) of Isoniazid Microspheres

Formulation Code	Silk Aq. Solution : PVA Parts	Drug Entrapment efficiency
F1	1:1	65.78 %
F2	1:2.5	59.44 %
F3	1:4	68.47 %
F4	1.5:5.5	63.93 %
F5	2:3	61.33 %

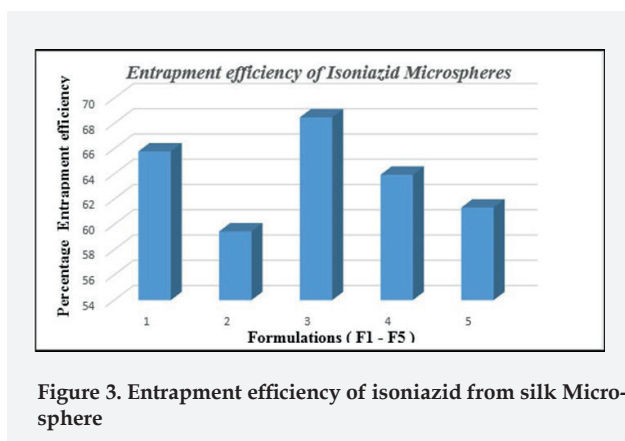


Figure 3. Entrapment efficiency of Isoniazid from silk Microsphere

In-vitro drug release from Isoniazid Loaded Microspheres in 7.4 phosphate buffer:

In-vitro release study of drug from Microsphere solution in 7.4 Phosphate buffer obtained between Percentage (%) cumulative release and Time (hours). Figure: 4

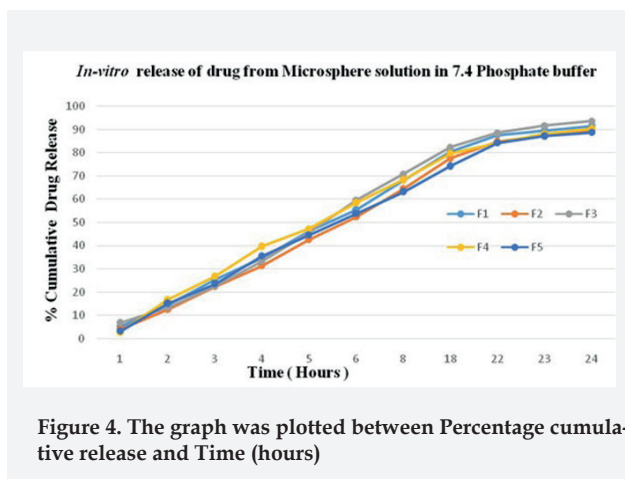


Figure 4. The graph was plotted between Percentage cumulative release and Time (hours)

CONCLUSION

Current experimental investigation was an attempt to develop a reconstituted microsphere of ant tubercular drug. Formulated silk microspheres were spherical, having narrow particle size distribution, glossy and dense surface, good entrapment efficiency and controlled drug release. The prepared microspheres could be used as a carrier for controlled release due to the biocompatibility and biodegradation of Silk Fibroin and long-term release of therapeutic proteins and peptides from Silk Fibroin matrix.

DISCUSSION

Isoniazid loaded silk fibroin microsphere was prepared by using phase separation method. All batches of Microsphere were preliminarily evaluated. Results disclosed the formation of Silk Microspheres and its drug delivery applications. Scanning electron microscopy studies revealed that

Isoniazid Loaded Silk Fibroin Microspheres were spherical. Entrapment Efficiency of Isoniazid loaded Microspheres of different Formulation from F1 to F5 was in range of 53 to 68 %. F3 showed 68.47 % Entrapment Efficiency and the optimized formulation drug release was 93.56 % at 24 hours. Stability Studies was Shows in 3 Months and no appreciable changes was noticed in drug release and Entrapment efficiency.

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