SYNTHESIS AND CHARACTERIZATION OF A NOVEL MUCOADHESIVE DERIVATIVE OF PSYLLIUM SEED POLYSACCHARIDE

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INTRODUCTION

Natural polysaccharides have widespread applications as biopolymers as they are economical, easily available in a variety of structures. They can be modified easily by the chemical and biochemical process. Along with other properties like stability, safety and biodegradability, they suggest use in targeted drug delivery systems. Also, they find extensive applications in pharmaceutical and food industry because of their diversity in structure and properties [1, 2].

Psyllium, also known as ispaghula comprises of seed husks of Plantago ovata forsk (family Plantaginaceae). The laxative and prolonged action of Plantago husk have been attributed to the gel forming fraction of the husk. Psyllium mucilage obtained from psyllium husk is white fibrous hydrophilic material that forms a clear colourless mucilaginous gel by absorbing water. The polysaccharides extracted from the husk of Plantago ovata have been chemically characterised to contain a high proportion of hemicellulose which is the alkali soluble fraction of the husk. It consists of highly branched acidic arabinose and xylose forming the side chains [3].

Psyllium husk has been reported for the treatment of constipation, diarrhea, irritable bowel syndrome, inflammatory bowel disease ulcerative colitis, colon cancer, diabetes and hypercholesterolemia [4].

Phosphorylated PSY has been investigated as a release retardant and binder [5, 6]. Acrylamide-crosslinked psyllium husk is reported as having increased mucoadhesive properties [11]. Carboxymethyl dextran was modified by the covalent attachment of cysteine which improved the mucoadhesive properties [12].

In the present study, an attempt was made to improve the mucoadhesive properties of PSY by thiol-functionallization [13]. Thiolation was observed in the surface morphology of psyllium from fibrous to granular and resulted in 82% swelling in deionized water. The percent increase in mucoadhesive retention time of TPSY was found to be 50.31% as compared to PSY and 128.30% as compared to carbopol. The percent increase in mucoadhesive retention time of TPSY was found to be 110% as compared to PSY and 50% as compared to carbopol.

Conclusion: Mucoadhesive retention time were greater of tablets containing a higher amount of TPSY. Further, the acute oral toxicity studies and repeated oral toxicity for TPSY proved it as non-toxic and hence safe for human use.

Keywords: Psyllium seed polysaccharide, Crosslinking, Microwave assisted, Thiolated psyllium seed polysaccharide

MATERIALS AND METHODS

Materials

Psyllium seeds were obtained from Manakarnika Aushadalay, Pune. Thiglyclic acid was a gift sample from Molychem, Mumbai. All the other chemicals, solvents and reagents used were of analytical grade and were procured locally. Dialysis membrane (HiMedia LA 401) was used for purification of crosslinked conjugate. Ciprofloxacin HCL (CIP) was supplied by Aarti Drugs Ltd. Mumbai.

Acute oral toxicity and repeated oral toxicity studies for TPSY were also conducted. A simplex centroid design containing 7 experimental runs to evaluate the effect of polymers i.e. PSY, TPSY and carbopol on the release of drug. The thiolation was confirmed by -SH stretch in Fourier Transform infrared spectra at 2353 cm⁻¹. Thiolation was observed in the thiolated PSY (TPSY) by a change in the surface morphology of psyllium from fibrous to granular and resulted in 82% swelling in deionized water. TPSY was found to contain 102.35 mmol of thiol groups/g as determined by the Ellman’s method. The percent increase in mucoadhesive retention time of TPSY was found to be 50.31 % as compared to PSY and 128.30 % as compared to carbopol.

Concentration: Mucoadhesive strength and mucoadhesive retention time were greater of tablets containing a higher amount of TPSY. Further, the acute oral toxicity studies and repeated oral toxicity for TPSY proved it as non-toxic and hence safe for human use.

Keywords: Psyllium seed polysaccharide, Crosslinking, Microwave assisted, Thiolated psyllium seed polysaccharide

AbSTRACT

Objective: In the present study, thiol-functionalization of psyllium seed polysaccharide (PSY) was cross-linked with thiglyclic acid by esterification in an attempt to reveal the mucoadhesive properties of thiolated psyllium seed polysaccharide (TPSY).

Methods: The crosslinking was carried out by the microwave-assisted method. A simplex centroid design was employed to study the mucoadhesive strength, mucoadhesive retention time and drug release profile. Comparative evaluation of carbopol-based ciprofloxacin hydrochloride (HCl) tablets containing PSY and TPSY was carried out. Acute oral toxicity studies and repeated oral toxicity for TPSY were also conducted.

Results: Thiolation was confirmed by -SH stretch in Fourier Transform infrared spectra at 2353 cm⁻¹. Thiolation was observed in thiolated PSY (TPSY) by a change in the surface morphology of psyllium from fibrous to granular and resulted in 82% swelling in deionized water. TPSY was found to contain 102.35 mmol of thiols/g as determined by the Ellman’s method. The percent increase in mucoadhesive retention time of TPSY was found to be 50.31 % as compared to PSY and 128.30 % as compared to carbopol. The percent increase in mucoadhesive retention time of TPSY was found to be 110% as compared to PSY and 50% as compared to carbopol.

Conclusion: Mucoadhesive strength and mucoadhesive retention time were greater of tablets containing a higher amount of TPSY. Further, the acute oral toxicity studies and repeated oral toxicity for TPSY proved it as non-toxic and hence safe for human use.

Keywords: Psyllium seed polysaccharide, Crosslinking, Microwave assisted, Thiolated psyllium seed polysaccharide

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Methods

Thiolation by microwave assisted method

TPSY was synthesised by microwave assisted esterification of PSY with thioglycolic acid in the presence of hydrochloric acid. The reaction was carried out with 2 moles of thioglycolic acid for every mole of the hydroxyl group in PSY. Psyllium seed powder was soaked in water (1:50) overnight and the pH of the mixture was adjusted to 12 with sodium hydroxide (2.5% w/v) solution [16]. The mixture was transferred to a two-neck round bottom flask and placed on the turntable of a microwave oven (Microwave Catalyst system CATA 2R). An electronic thermometer with an alarm set at 70 °C was incorporated into the reaction vessel. This was to ensure that the reaction temperature did not cross 70 °C. Microwave irradiation at 340 W was performed for 10 min followed by 425 W for 10 min [17]. The reaction mixture was cooled and precipitated by adding methanol followed by washing of precipitate with methanol to remove the unreacted thioglycolic acid. TPSY so obtained was frozen at -40 °C for 4h followed by lyophilization in a laboratory model freeze drier (2.5 Freezone Labconco equipment limited) at -90 °C and 0.0010 mbar pressure for 24 h.

Purification by dialysis

Dialysis membrane (HiMedia LA 401) with a cut-off molecular weight between 5000-10000 daltons was soaked overnight in distilled water. A section of required dimensions was cut and one end was tied and filled with the synthesized product. The other end was then tied securely to prevent any leakage and it was immersed in a beaker containing 50 ml dialysing solvent i.e. methanol. The temperature was maintained at 25 °C to prevent any leakage and it was immersed in a beaker containing 50 ml dialysing solvent i.e. methanol. The temperature was maintained at 25 °C and the system stirred for 24 h with a magnetic stirrer. The sample was then removed and air dried for 3 h. The dialyzed product was then subjected to further characterization [18].

Determination of thiol substitution

The degree of thiol group substitution was determined by Ellman’s method. Aqueous dispersions (0.5%, w/v) of TPSY and PSY (as control) were prepared and diluted with phosphate buffer (5M, pH 8.0) to a concentration of 0.15% w/v. An aliquot of 5 ml of the polymer solution was allowed to react with 5 ml of Ellman’s reagent (5.5-dithio bis (2-nitrobenzocid acid) (0.3%) w/v) for 2h at room temperature, followed by measurement of absorbance at 450 nm. The numbers of thiol group substituents per gram of TPSY were determined using a calibration curve prepared by reacting standard solutions of thioglycolic acid with Ellman’s reagent [19].

Characterization of psyllium seed polysaccharide (PSY) and thiolated psyllium seed polysaccharide (TPSY)

The PSY and TPSY were characterized for physicochemical properties (solubility studies, swelling index and micromeritic properties), spectral analysis and thermal analysis. Acute oral toxicity studies and repeated oral toxicity were also conducted as per OECD Guidelines 423.

Swelling Index

Swelling index is indicative of the ability of polysaccharides to swell after absorbing fluids available at the site of absorption, which is mandatory for mucoadhesion. Swelling behavior of PSY and TPSY was studied at different pH such as 0.1 N HCl (pH 1.1), phosphate buffer pH 6.8, deionized water and 0.5M NaOH (pH 13.7). For this, 1 g of PSY and TPSY were added separately in 25 ml of the respective media in stoppered volumetric flask at 37 °C and kept overnight with intermittent shaking. The volume occupied by the PSY was measured using a measuring cylinder and swelling index was calculated from the initial and final volume of PSY and TPSY respectively [20].

Solubility profile

The solubility of PSY and TPSY was evaluated by modified gravimetric method at 37 °C in various solvents such as acetone, alcohol, ether, chloroform, dichloromethane, dimethyl amine, tri methylamine, diethyl ether, ethyl acetate, dimethyl sulfoxide, isopropyl alcohol, 0.5N sodium hydroxide, N-methyl-2-pyrrolidone, N,N-dimethyl formamide, hydrochloric acid, ammonia, and conc. nitric acid by adding PSY and TPSY in increments of 0.1 g followed by sonication for 5 min [20].

Micrometric properties

The bulk density, tapped density, angle of repose, Hausner’s ratio and Carr’s index of PSY and TPSY was evaluated [21].

Rheological properties

The muclages of PSY and TPSY were prepared in distilled water in strengths of 0.4, 0.5, 0.6, 1 and 2 % w/v. The muclages were homogenized using tissue homogenizer to break any lumps and subjected to rheometry studies at different shear rates using Brookfield viscometer (Brookfield DV-E Viscometer) using Helpath spindle-E (6) at 25°C [20].

Mucoadhesive strength

The mucoadhesive strength of PSY and TPSY pellets were evaluated using Texture Analyzer (CEB Texture Analyzer, Brookfield Engineering Labs, Inc, Model Texture Pro CT V1.4 Build 17). The pellets (100 mg) were prepared using a hydraulic press (Mini Press II MT, Make: Rimek). Fresh sheep gastric mucosa was procured from the local slaughter house and was washed with distilled water followed by 0.1N HCl and was carefully attached using a double-sided adhesive tape to the cylindrical probe (TA 3/100 Probe). The probe was then removed at a speed of 1 mm/s and to a distance of 15 mm, and maximum detachment force (N) was determined with data rate 15 points/sec. A different mucosa sample was used for every new sample. The mucoadhesive strength, i.e. the maximum force required for separating the tablet from the mucosal surface was obtained [22].

Mucoadhesion time

The mucoadhesion time was evaluated by in vitro adhesion testing method known as the wash-off method using disintegration apparatus (Make: VEEGO VTD-2). Gastric mucosa was fixed on a glass slide using a double-sided adhesive tape and the pellet was wetted with a drop of 0.1N HCl and affixed to the mucosal tissue. The glass slide was then tied to the paddle with cotton thread of a USP disintegration apparatus containing 900 ml of 0.1N HCl and rotated at 37±0.5 °C. After 2 min, a stirring rate of 25 rpm was applied to simulate the gastric environment, and time taken for the tablet to detach from sheep gastric mucosal tissue was recorded as mucoadhesion time [23].

Spectral analysis

Fourier transform infrared spectroscopy (FTIR)

Infra-red spectrum of PSYand TPSY was recorded out on Jasco FTIR-401, Japan at transmittance mode. Baseline correction was done using dried potassium bromide (KBr). KBr was initially dried and mixed in a ratio of 1:4 with PSYand TPSY. A small amount of triturated sample was kept into the sample holder and scanned from 400 cm⁻¹ to 400 cm⁻¹ in FTIR spectrophotometer (FTIR 8400S, Shimadzu, Japan).

Powder X-ray diffraction analysis (PXRD)

PXRD patterns were recorded in the reflection mode on a Siemens D-500 X-ray diffractometer. Diffractograms were registered at Bragg angle (2θ) range of 5-40° at a scan rate of 2.5° per minute and step size of 0.02°. X-ray diffractogram on powdered samples were obtained using a Bruker AXS Advanced X-ray diffractometer under the following operating conditions: 40 kV and 40 mA with Cu-Kα radiation at k 1.5418 Å and acceptance slot at 0.1 mm. About 15-20 mg of the sample was spread on a sample stage, and the relative intensity was recorded in the scattering range (2θ) of 0-60°C in steps of 0.1°C.

Field emission scanning electron microscopy (FESEM)

The surface morphology of PSY and TPSY were determined using FESEM (Instrument: FEL, model: Nova nano SEM 450). A field emission gun produces an electron beam with an extremely high...
current density, obtained by applying an intense electric field to a tungsten single crystal with a needle-shaped tip. This requires ultra-high vacuum (10\textsuperscript{-10} Torr). The samples were sprinkled on a double adhesive tape stuck to an aluminum stub. The stubs were then coated with platinum to a thickness of about 10Å under an argon atmosphere using a gold sputter module in a high-vacuum. The stub containing the samples was placed in the FESEM chamber.

**Differential scanning calorimetry (DSC)**

The thermograms of PSY and TPSY were obtained by a DSC (DSC 821, Make: METTLER) at a heating rate of 10 °C/min from 30 to 300 °C in a nitrogen atmosphere (30 ml/min).

**Acute oral toxicity studies for TPSY**

An acute oral toxicity studies was obtained from Institutional Animal Ethics Committee (Approval No.: CPCSEA/IAEC/PT–05/01–2K16). An acute oral toxicity study was conducted according to OECD guidelines 423. The study was performed in healthy male Wistar rats (200-300 g). Pilot study was carried out using six male Wistar rats in a group with the starting dose of 300 mg/kg body weight. Animals were monitored continuously immediately after dosing up to first 4 h. The animals were observed for any sign of toxicity or mortality after 24 h and 14 d. When 300 mg/kg dose was proved to be non-toxic, in the main study 2000 mg/kg dose was screened for checking toxicity using the above-mentioned procedure [24].

**Repeat dose (28 d) oral toxicity study**

Approval for animal studies was granted by CPCSEA with Approval number: CPCSEA/AEAC/PT-05/01-2K16. A 28 d repeat dose toxicity study was performed as per OECD Guidelines 407 (Adopted: 3 October 2008) for the optimised TPSY. Male Wistar rats, divided into 4 groups of five rats, were used for the study. The animals in group 1 served as vehicle controls and were administered normal saline daily by oral gavage. The animals in groups 2, 3 and 4 were administered a daily dose of 70, 175 and 350 mg/kg body weight of TPSY as aqueous suspension by oral gavage for 28 d respectively.

All the animals were observed daily for clinical signs of toxicity and morbidity. Individual animal body weights were recorded at receipt (on day 1) and weekly thereafter and at the time of sacrifice (fasting body weights). Histological examinations of various organs (brain, kidney, liver, small intestine and stomach) were carried out after 28 d using haematoxylin and eosin as staining agents. The organs were preserved in 10% formalin solution.

**Preparation of granules**

Granules were prepared by wet granulation technique using varying concentrations of PSY, TPSY and PVPK30 for comparative study. CIP was mixed with lactose and granulated using PSY, TPSY and PVP K30 (1% w/v solution in IPA) as granulating agents separately. The wet mass was passed through sieve no.12 and the granules were dried at 45 °C for 30 min. The dried granules were further passed through mesh no.16 and then lubricated with the mixture of talc and magnesium stearate. The granules were evaluated and further compressed into tablets using 10.5 mm flat punch in rotary tablet press (Mini 2 Remake).

**Experimental design**

A simplex centroid design containing 7 experimental runs to evaluate variable drug to thiomer concentration was employed to determine their effect on three responses i.e. mucoadhesive strength, retention time and percent drug release and their interaction therein. The simplex centroid design for a 3-component system (AA, BB and CC) is represented by an equilateral triangle in a 2-dimensional space shown in fig. 1. The amounts of thiolated psyllium (A+TPSY), psyllium (B=PSY) and carbopol (C=carbopol) were selected as independent variables (table 1) [25].

![Fig. 1: Equilateral triangle representing simplex centroid design for 3 components (A,B,C).](image)

**Table 1: Composition of formulation of ciprofloxacin HCl tablets prepared according to simplex mixture design with translation of coded levels in actual units (F1-F7)**

| Ingredients        | Formulations | F1     | F2     | F3     | F4     | F5     | F6     | F7     |
|--------------------|--------------|--------|--------|--------|--------|--------|--------|--------|
| Ciprofloxacin HCl  | 125          | 125    | 125    | 125    | 125    | 125    | 125    | 125    |
| TPSY (A)           | 70 (1)       | 40 (0) | 40 (0) | 55 (0.5)| 55 (0.5)| 40 (0) | 55 (0.5)| 50 (0.33)|
| PSY (B)            | 40 (0)       | 70 (1) | 40 (0) | 55 (0.5)| 40 (0) | 55 (0.5)| 50 (0.5)| 50 (0.33)|
| Carbopol1(C)       | 40 (0)       | 40 (0) | 70 (1) | 55 (0.5)| 55 (0.5)| 50 (0.5)| 50 (0.33)|
| PVP K30            | 35           | 35     | 35     | 35     | 35     | 35     | 35     | 35     |
| Talc               | 7            | 7      | 7      | 7      | 7      | 7      | 7      | 7      |
| Magnesium stearate | 7            | 7      | 7      | 7      | 7      | 7      | 7      | 7      |
| Lactose            | 26           | 26     | 26     | 26     | 26     | 26     | 26     | 26     |
| Total              | 350          | 350    | 350    | 350    | 350    | 350    | 350    | 350    |

All values are in mg. *A is TPSY, B is PSY and C is Carbopol

**Evaluation of tablets**

**Physical properties of tablets**

The hardness of the tablets was evaluated using Monsanto Hardness Tester and was also evaluated for weight variation. The tablets were also evaluated for friability using Roche friabilator [20]. Ten tablets were placed in the Roche friabilator, which was then operated for 100 revolutions for 4 min at 25 rpm. After 100 revolutions the tablets were dedusted and reweighed. Percentage friability was calculated by the following formula:

\[
\text{Percentage friability} = \frac{[\text{Initial weight of tablets} - \text{Final weight of tablets}]}{\text{Initial weight of tablets}} \times 100
\]

**Mucoadhesive strength and mucoadhesion time**

The mucoadhesive strength of the tablets was evaluated using Texture Analyzer (CEB Texture Analyzer, Brookfield Engineering Labs, Inc., Model Texture Pro CT V1.4 Build 17) and mucoadhesion time was evaluated using disintegration apparatus on fresh sheep gastric mucosa by the procedure described previously.

**In vitro dissolution studies and release kinetics**

The in vitro dissolution studies were carried out in 900 ml 0.1N HCl using USP Dissolution Apparatus type I at 37 °C and 100 rpm. Dissolution fluid was withdrawn at regular time intervals during the studies and sink condition was maintained by replacing with an
equal volume of drug-free dissolution fluid. The samples were suitably diluted with fresh dissolution fluid. The samples were then analysed on a Jasco UV/VIS spectrophotometer (V-530) at λ max of 276 nm. Absorbance was measured and percent drug release was determined. The data obtained from in vitro release studies were fitted into various kinetic equations viz., zero order, first order, Higuchi matrix and Korsmeyer-Peppas to know the mechanism of drug release. The equation with high regression coefficient (r²) and n values for formulation will be the best fit of release data. Goodness-of-fit was evaluated using DD Solver 1.0 software.

RESULTS AND DISCUSSION

Crosslinking of PSY

Microwave irradiation enhances the reaction rate by providing energy efficient internal heating by direct coupling of microwave energy. In the conventional method, reactants are slowly activated by a conventional external heat source [14]. Microwave method was best suited for the reaction compared to the conventional method as the percent yield was greater in microwave i.e. 74% as compared to conventional method i.e. 60%. The conventional method required 2 h for completion of reaction whereas with microwave method the reaction was completed in 20 min.

The covalent attachment of thiglycolic acid to psyllium was achieved by ester bond formation between the hydroxyl group of arabinoxylan moieties of psyllium and carboxyl group of thiglycolic acid as depicted in Fig. 2. The finely ground product appeared as a brownish yellow odorless powder, which formed a gel in water.

The average yield amounted to 74% of the utilized amount of psyllium. The unreacted thiglycolic acid was removed by repeated washings with methanol. TPSY was found to contain 102.35 mmol of thiol groups/g, as determined by the Ellman’s method. Dialysis with excess methanol was carried out for removal of unreacted PSY from the freeze-dried sample [26, 27].

Characterization of psyllium seed polysaccharide (PSY) and thiolated psyllium seed polysaccharide (TPSY)

Swelling profile of PSY and TPSY

The percent swelling of PSY and TPSY was studied in phosphate buffer pH 6.8, 0.1 N HCl, 0.5 N NaOH and deionised water. For PSY, maximum swelling of 70% was observed in phosphate buffer pH 6.8 whereas, in deionized water, 55% swelling was observed. No swelling was observed in HCl For TPSY, 82% swelling was observed in deionized water, 75% in phosphate buffer pH 6.8, 35% in 0.5 N NaOH and 52% in 0.1 N HCl. Swelling is an important criteria for mucoadhesion along with molecular weight and concentration of the polymer and flexibility of polymer chains [28]. Swelling of the polymer was found to increase with respect to time and rate of hydration. The initial contact time affected the percent swelling with higher swelling observed with greater contact time. Inter-penetration of polymer chains was enhanced due to increased swelling which facilitated stronger anchoring onto the mucosal surface leading to greater mucoadhesion [29]. Thiol groups on TPSY further strengthen the interaction between the polymer and mucosa thereby enabling maximal mucoadhesion [30].

Solubility profile

Solubility can be defined as the spontaneous interaction of two or more substances to form a homogenous molecular dispersion. PSY was found to form gel in DMSO, ammonia, 0.5N NaOH, water and formamide whereas TPSY was found to be slightly soluble only in formamide and formed a gel in 0.5N NaOH, dimethyl sulfoxide, ammonia and water. The formation of gel of TPSY is indicative of its hydrogel nature which can be exploited to design sustained release dosage forms. Hydrogels are three-dimensional, hydrophilic, polymeric networks, with chemical cross-links, capable of imbibing large amounts of water [31]. Hydrogels have been reported as drug delivery systems for modified release dosage forms. Chitosan hydrogels have been used for insulin delivery through oral route and hyaluronic acid hydrogels are reported for drug delivery and tissue engineering [32].

Rheological data

Rheological investigations of 1% w/v dispersions of PSY and TPSY revealed a continued decrease in viscosity with an increase in shear rate indicating pseudoplastic behavior (fig. 3). Similar results were obtained for tamarind seed polysaccharide [29]. Pseudoplastic systems are shear thinning in nature and are characterised by structural rigidity at rest. Application of shear results in the breakdown of the structure and an apparent fall in viscosity due to the release of entrapped water molecules [33]. TPSY exhibited a sudden breakdown in structure with increase in shear rate. At higher strength, the mucilage has a very rigid structure which can be attributed to the formation of hydrogen bonds between the polysaccharide and water.

The viscosity of TPSY at any shear rate was found to be significantly higher than that of PSY. The viscosity of 1% dispersion of TPSY at a shear rate of 10 rpm was found to be 1.75 folds higher than that of PSY. The higher viscosity of TPSY at low shear rates will facilitate its mucoadhesive properties as it will contribute to increased residence time of the drug delivery system at the site of use. At the same time, shear thinning properties of TPSY will allow greater flexibility of the polymer chains which is a prerequisite for mucoadhesion.
Mucoadhesive strength

The mucoadhesive strength of a polymer is dependent on molecular weight of the polymer, contact time with the membrane, the degree of swelling of the polymer and concentration of polymer [29]. The mucoadhesive strength of PSY and TPSY was studied using sheep intestinal mucosa (table 2). TPSY was found to have higher mucoadhesive strength as compared to PSY and carbopol. Carbopol is a hydrogel which swells on contact with water. The extent of swelling depends upon the degree of crosslinking. Hydrogen bonding with mucin is responsible for mucoadhesive properties of hydrogels [35]. Thiolated polymers act by forming covalent disulphide bonds with cysteine-rich subdomains of mucin glycoproteins which build the mucus gel layer [36]. The bond strength of covalent bonds ranges from 50-200 kcal/mole while a hydrogen bond has the strength of 2-8 kcal/mole [37]. The higher mucoadhesive strength of TPSY can thus be attributed to the stronger bonds formed between the polymer and mucin. The percent increase in mucoadhesive strength of TPSY was found to be 50.31 % as compared to PSY and 128.30 % as compared to carbopol.

| Polymer | Mucoadhesive strength (g) | Mucoadhesive retention time (min) |
|---------|---------------------------|----------------------------------|
| PSY     | 16.1±1.35                 | 62±0.48                          |
| Carbopol| 10.6±0.95                 | 70±0.28                          |
| TPSY    | 24.2±1.17                 | 195±0.72                         |

Table 2: Evaluation of mucoadhesive strength and mucoadhesive retention time of PSY and TPSY pellets

Mucoadhesive retention time

In vitro retention time is an important physical parameter of mucoadhesive polymers. PSY and carbopol were found to have lower retention time as compared to TPSY (table 2). The mucoadhesive retention time of TPSY was found to be 110 % higher than PSY and 50 % higher than carbopol. Increased residence time was observed with increase in the concentration of mucoadhesive polymers in the formulations. This test reflects the mucoadhesive capacity of polymers used in formulations. The results revealed that TPSY showed better bio-adhesion than PSY and carbopol polymer. These results co-related well with results of mucoadhesive force and swelling index.

Spectral analysis

Fourier transform infrared spectroscopy (FTIR)

The fig. 4 exhibits the FTIR spectrum of PSY and TPSY in the frequency region from4000 to 400 cm⁻¹. The spectra of PSY shows a broad absorption band at 3401 cm⁻¹ which can be attributed to OH stretching of alcohols. The peaks appearing at 1000, 740, and 590 cm⁻¹ may be due to polymer backbone bendings. The IR spectra of TPSY show the absorption band due to OH stretching at 3425 cm⁻¹, while the–SH stretch appears at 2353 cm⁻¹ which confirms the thiolation. The C–O–C stretch appears at 1000 cm⁻¹ in TPSY, while the peaks at 770 and 447.50 cm⁻¹ are for polymer backbone bendings.

Powder x-ray diffraction analysis (PXRD)

XRD patterns are recorded using X-ray diffractometer. Samples were analyzed between 0-80° (2θ). The halo pattern evident in the diffractogram was indicative of the amorphous nature of both PSY and TPSY depicted in fig. 5. During crystallization process, short chain polymers organize themselves into highly ordered crystalline structures more readily than long chain polymers. Polymer morphology is also dependent on the size and shape of the substituent groups [33]. The diffractograms support the DSC findings and we may infer that the thiolated polymer is amorphous in nature.

Differential scanning calorimetry (DSC)

Fig. 6 compares the DSC thermograms of PSY and TPSY. The thermal curve of PSY shows broad endotherm at 88.79 °C while the thermogram of TPSY displayed a broad endotherm at 100 °C as shown in fig. 6. Thus the shift in the endothermic peaks in the thermal curve of TPSY indicates the modification of PSY [38]. The enthalpy values for PSY were found to be-0.800 mW and that of TPSY were found to be-0.780 mW. The enthalpy of TPSY was higher which indicates that TPSY is more thermally stable as compared to PSY. Enthalpy is a state function that depends only on the prevailing equilibrium state identified by the variables internal energy, pressure, and volume [39, 40]. Previously reported DSC studies on xyloglucan and thiomer (modified L-cysteine HCl) showed a broad endotherm at 54 °C and 74.59 °C respectively. The thermoderm of thiomer exhibited a sharp endothermic peak 212.57 °C with heat flow 113.96 mJ/g which might be due to combustion of thiomer [41]. Previously reported DSC studies on xyloglucan and thiomer (modified thioglycolic acid) showed broad endotherm at 85.29 °C while the thermal curve of thiolated TSP showed a broad endotherm at 81.78 °C [42].
Field emission scanning electron microscopy (FESEM)

In FESEM, the gun emits the electrons from the much smaller area and the coherency is higher and thus provides better resolution. Fig. 7 shows the shape and surface morphology of PSY and TPSY, examined under an FESEM. The shape of PSY and TPSY particles was found to be polyhedral. A close examination of surface morphology revealed the fibrous appearance of PSY and granular appearance of TPSY.

Acute oral toxicity studies for TPSY

Acute toxicity refers to the adverse effects that occur on first exposure to a single dose of polymer. Acute oral toxicity studies were conducted according to OECD guidelines. A pilot study was carried out using six male Wistar rats in a group with the starting dose of 300 mg/kg body weight. Animals were monitored continuously immediately after dosing up to first 4 h and thereafter for any sign of toxicity or mortality after 24 h and 14 d. When 300 mg/kg dose was found to be non-toxic, a higher dose of 2000 mg/kg was screened.

At a dose of 2000 mg/kg, the mice showed lethargy during first 4 h of observation. A slight increase in body weight was observed after 14 d but no mortality was seen (table 3). There were no clinical signs of toxicity like skin rashes, hyperactivity, excessive grooming and kicking of rear legs. Administration of TPSY up to 2000 mg/kg was well tolerated by the animals. Mean body weight changes and feed consumption were not affected by administration TPSY by oral gavage. Animals in the test groups continued to gain body weights during the study.
Repeat dose oral toxicity studies

The objective of the repeat dose oral toxicity studies is to identify toxicity, to identify the organs most affected and to determine the doses at which each effect occurs. Male Wistar rats, divided into 4 groups of five rats each, were used for the study. The animals in group 1 served as vehicle control and were administered normal saline daily by oral gavage. The animals in groups 2, 3 and 4 were administered a daily dose of 70, 175 and 350 mg/kg body weight of TPSY as aqueous suspension by oral gavage for 28 d respectively. All the animals were observed daily for clinical signs of toxicity and morbidity. Individual animal body weights were recorded at receipt (on day 1) and weekly thereafter and at the time of sacrifice (fasting body weights). The animals were harvested after 28 d and histological examinations of various organs (brain, kidney, liver, small intestine and stomach) were carried out. The organs were preserved in 10 % formalin solution.

On administration of TPSY, the animals in all groups were found to survive at the end of 28 d of study. The histological results are shown in table 4. Liver tissue showed minimal degeneration of hepatocytes compared to control at the highest dose of 350 mg which was however reported to be non-toxic.

Mild tubular degeneration was observed in kidneys. Stomach tissue displayed moderate changes compared to control, with minimal degeneration of mucosal cells at highest dose of 350 mg. Large intestine showed marked submucosal edema but the mucosal tissue appeared to be normal at the highest dose of 350 mg. No abnormality was observed in brain tissues at a low and medium dose. Mild submucosal edema was observed at 350 mg/kg. These results point to the negligible toxicity associated with the thiolated polymer and hence it can be considered to be safe for use as a pharmaceutical excipient.

Evaluation of tablets

The diameter and thickness of all the batches were found to be 10.51-10.58 mm and 4.30-4.38 mm respectively. The weight variation was found to be 0.349-0.354. The hardness and friability were found to be 5.2-5.8 kg/cm² and 0.14-0.21 % respectively. All the parameters were found to be within official limits.

Simplex-centroid design

A simplex centroid design containing 7 experimental runs to evaluate variable drug to thiomer concentration was employed to determine their effect on three responses i.e. mucoadhesive strength, retention time and percent drug release (table 5).

Table 4: Histological results of repeated dose toxicity studies of TPSY (n=5)

| Organs       | Normal saline | TPSY dose strength |
|--------------|---------------|--------------------|
|              | Control       | 70 mg/kg           | 175 mg/kg          | 350 mg/kg          |
| Liver        | Normal architecture of liver, nothing abnormal detected. | Nothing abnormal detected. | Minimal degeneration of hepatocytes and infiltration of inflammatory cells. | Minimal degeneration of hepatocytes and infiltration of inflammatory cells. |
| Kidney       | Normal architecture of kidney, nothing abnormal detected. | Minimal tubular degeneration. | Mild tubular degeneration and congestion of blood vessels. | Mild tubular degeneration. |
| Stomach      | Normal architecture of Stomach, Nothing abnormal detected. | Minimal degeneration of mucosal lining cells. | Mucosal cells were enlarged and were darkly stained. | Minimal degeneration of mucosal cells, accumulation of dark staining material in the gland. |
| Brain        | Normal architecture of the brain, Nothing abnormal detected. | Nothing abnormal detected. | Nothing abnormal detected. | Minimal neuronal degeneration. |
| Large intestine | Papillary projections with goblet cells. | Mucosal lining no infiltration of inflammatory cells. | Minimal infiltration of inflammatory cells and congestion. | Marked submucosal edema, mucosal tissue appears normal. |
Table 5: Responses for tablets prepared using simplex centroid design

| Formulation | Response 1 mucoadhesive strength (g) | Response 2 mucoadhesive time (min) | Response 3 % drug release (%) |
|-------------|--------------------------------------|-----------------------------------|------------------------------|
| F1          | 24.78±0.21                           | 441±0.61                          | 90±0.33                      |
| F2          | 15.24±0.43                           | 68±0.46                           | 88±0.27                      |
| F3          | 14.57±0.56                           | 62±0.22                           | 86.5±0.58                    |
| F4          | 20.96±0.61                           | 334±0.45                          | 91.5±0.68                    |
| F5          | 19.24±0.24                           | 331±0.41                          | 91.8±0.48                    |
| F6          | 13.4±0.51                            | 122±0.54                          | 95.8±0.71                    |
| F7          | 17.6±0.46                            | 247±0.82                          | 94.77±0.821                  |

*mean±SD; n=3
Mucoadhesive strength (g)

TPSY shows enhanced mucoadhesive strength as compared to PSY and carbopol. This can be attributed to cross linking of PSY with thioglycolic acid (Equation 1). The low coefficients for terms AB, AC and BC point to the absence of synergistic effect of polymers. The higher coefficient for term A (TPSY concentration) is indicative of its greater mucoadhesive strength than terms B (PSY concentration) and C (carbopol). This supports our earlier findings regarding the strong covalent bond that thiol functionalization forms with the mucin than the hydrogen bonds which are responsible for the mucoadhesive strength of PSY and carbopol. The Model F-value of 1287.38 implied that the quadratic model is significant which is further substantiated by values of "Prob>F" less than 0.0500 for A, B and C.

\[ \text{Mucoadhesive Strength} = 24.79(A) + 15.25(B) + 14.58(C) + 3.59(AB) - 1.95(AC) - 6.23(BC) \] Eq. 1

Fig. 9: Response surface graph for mucoadhesive strength where X1=A=TPSY, X2=B=PSYand X3=C=Carbopol

Mucoadhesive time (min)

The mucoadhesive retention time for TPSY was higher as compared to PSY and carbopol which reinforces our findings of mucoadhesive strength. The increased retention time of TPSY can be due to enhanced mucoadhesive strength of TPSY (Eq.2). This may be attributed to the cross linking by thiol moiety. The Model F-value of 269.41 implied the significance of the quadratic model. Values of "Prob>F" less than 0.0500 for A supported the above inference.

\[ \text{Mucoadhesion Time} = 7.20(A) + 1.79(B) + 1.19(C) + 3.84(AB) + 3.84(AC) + 3.06(BC) \] Eq.2

Fig. 10: Response surface graph for mucoadhesion retention time where X1=A=TPSY, X2=B=PSYand X3=C=Carbopol

Percent drug release

TPSY showed greater drug retardation as compared to PSY and carbopol (Eq.3). The Model F-value of 1997.99 implied that the quadratic model was significant. Values of "Prob>F" less than 0.0500 for A, B, C, AB, AC and BC indicated significant quadratic model. The release of the drug from the formulation was found to be comparable with a formulation containing carbopol. The percent increase in mucoadhesive retention time of TPSY was 110% as compared to PSY and 50% as compared to carbopol. The results of acute oral toxicity studies and repeat dose (28 d) oral toxicity study for thiolated conjugate pointed to the negligible toxicity associated with the thiolated polymer and hence can be considered to be safe for use as a pharmaceutical excipient. We may thus conclude that thiol modification of psyllium resulted in improved its mucoadhesive properties.

\[ \text{Drug Release} = 89.99(A) + 87.99(B) + 86.49(C) + 10.14(AB) + 14.34(AC) + 34.42(BC) \] Eq. 3

Fig. 11: Response surface graph for drug release where X1=A=TPSY, X2=B=PSYand X3=C=carbopol

Fig. 12: In vitro drug release from formulations in 0.1N HCl (mean±SD n=3)

CONCLUSION

Modification of psyllium was carried out by esterification with thioglycolic acid. Thiolated conjugate was characterized by FT-IR, XRD, DSC and FESEM study. Modified psyllium was employed for formulating mucoadhesive tablets using ciprofloxacin HCl as the model drug. The percent increase in mucoadhesive strength of TPSY was found to be 50.31 % as compared to PSY and 128.30 % as compared to carbopol. The percent increase in mucoadhesive retention time of TPSY was 110% as compared to PSY and 50% as compared to carbopol. The results of acute oral toxicity studies and repeat dose (28 d) oral toxicity study for thiolated conjugate pointed to the negligible toxicity associated with the thiolated polymer and hence can be considered to be safe for use as a pharmaceutical excipient. We may thus conclude that thiol modification of psyllium resulted in improved its mucoadhesive properties.
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CONTRIBUTION

The corresponding author, Monica RP Rao has supervised the project. Ms. Snehal Galkowd has undertaken the project and preparing the draft of the manuscript. Ms Prachi Shevate has helped in drafting and editing the manuscript.

CONFLICT OF INTERESTS

There is no conflict of interest, financial or otherwise associated with this project.

REFERENCES

1. Chauhan P, Mehta N. Probiotic assisted colon targeted drug delivery system: research scope. Asian J Pharm Clin Res 2011;4:12-5.
2. Khan M, Ansari V, Kushwaha P, Kumar A, Akhtar J. Mucoadhesive microspheres for controlled delivery of drugs. Asian J Pharm Clin Res 2015;8:1-20.
3. Kumar R, Setia A, Mahadevan N. Grafting modification of the polysaccharide by the use of microwave irradiation—a review. Int J Recent Adv Pharm Res 2012;2:45-53.
4. Aminabhavi T, Chaturvedi K, Ganguly K, Nadagouda M. Polymeric hydrogels for oral insulin delivery. J Controlled Release 2013;165:129–38.
5. Singh B. Psyllium as therapeutic and drug delivery agent. Int J Pharm 2007;34:1-14.
6. Rao MRP, Warrier D, Rao S. Evaluation of phosphorylated psyllium seed polysaccharides a release retardant. Indian J Pharm Sci 2015;77:605-12.
7. Singh B, Sharma V. Design of psyllium–PVA-acrylic acid based novel hydrogels for use in antibiotic drug delivery. Int J Pharm 2012;389:94–106.
8. Singh B S, Kumar. Synthesis and characterization of psyllium-NVP based drug delivery system through radiation crosslinking polymerization. Nuclear Instrum Methods Phys Res 2008;266:3417–30.
9. Lalge M, Sharma P, Bhandari A, Garud A, Garud N. Mucoadhesive drug delivery system: a review. Crit Rev Pharm Sci 2014;3:17-29.
10. Sarti F, Staff A. Thiolated hydroxyethylcellulose: synthesis and in vitro evaluation. Eur J Pharm Biopharm 2010;76:421-7.
11. Roldoa M, Kwon M. Mucoadhesive thiolated chitosans as platforms for oral controlled drug delivery. synthesis and in vitro evaluation. Eur J Pharm Biopharm 2000;54:7:115-21.
12. Shahnaz G, Perera G. Synthesis, characterization, mucoadhesion and biocompatibility of thiolated carboxymethyl dextran-cysteine conjugate. J Controlled Release 2010;144:32-9.
13. Kaur H, Yadav S. Synthesis, characterization and evaluation of thiolated tamarind seed polysaccharide as a mucoadhesive polymer. Carb Pol 2012;90:1543-9.
14. Bhosale R, Gangadanapppa H, Moin A, Gowda D, Osmani R. A review on grafting modification of polysaccharides by microwave irradiation-distinctive practice for application in drug delivery. Int J Curr Pharm Res Rev 2015;6:8-17.
15. Divate V, Dongade Desed S. An efficient microwave assisted multicomponent synthesis of some 7-amino-3-(substituted phenyl)-5-(substituted phenyl)-[1,2,4]triazolo-[4,3-a]pyrimidine-6-carbonitrile derivatives. Int J Curr Pharm Res 2014;6:20-4.
16. Bhattia M, Ahuja M. Thiol modification of psyllium husk mucilage and evaluation of its mucoadhesive applications. Hindawi Publishing Corporation Sci World J 2015;3:1-7. http://dx.doi.org/10.1155/2015/284182
17. Sena G, Mishra S. Microarray initiated synthesis of polyacrylamide grafted psyllium and its application as a flocculant. Int J Biol Macromol 2012;50:369-75.
18. Berg J, Tymoczko J, Stryer L. Biochemistry 5th ed. WH Freeman. New York; 2002.
19. Khandelwal KR, Practical pharmacognosy, Techniques and experiments. 19th ed. Niral Prakashan, Pune; 2008. p. 159.
20. Rao MRP, Kambete M, Lunavat H. Study of rheological properties of psyllium polysaccharide and its evaluation as a suspending agent. Int J PharmTech Res 2013;5:1197-1.
21. Patrick S, Martin’s physical pharmacy and pharmaceutical sciences. 5th ed. Lippincott Williams and Wilkins, Philadelphia; 2000. p.587-8.
22. Krishnarajan D. Effect of cellulose and non-cellulose polymers on ciprofloxacin extended-release tablets. J Chem Pharm Res 2012;4:3617-23.
23. Surawase R, Maru A. Formulation and evaluation of metoprolol succinate buccal tablet containing tamarind seed polysaccharides. Int J Pharm Pharm Sci 2011;3:1491.
24. OECD guideline for testing of chemicals—423 Acute Oral Toxicity—Acute Toxic Class Method; 2001.
25. Madgulkar A, Bhailekar M. Optimization of carboxymethyl-xylloglucan-based tramacol matrix tablets using simplex centroid mixture design. Hindawi Publishing Corporation J of Pharma 2013;1-11. http://dx.doi.org/10.1155/2013/396468
26. Indian Pharmacopoeia. The government of India, Ministry of Health and Family Welfare. Vol. 3. Published by the Indian Pharmacopoeia Commission; 2010. p. 1247.
27. Singh V, Kumar P, Sanghi R. Use of microwave irradiation in the grafting modification of the polysaccharides—a review. Prog Polym Sci 2012;37:604-64.
28. Bhattacharai N, Ramay H. PEG-grafted chitosan as an injectable thermosensitive hydrogel for sustained protein release. J Controlled Release 2005;103:609–24.
29. Sravani B, Deveswaran R, Bhardwaj S, Basavanaj B, Madhavan V. Development of sustained release metformin hydrochloride tablets using a natural polysaccharide. Int J App Pharm 2012;4:23-9.
30. Fang P, Khurana S, Madhav S. Mucoadhesive drug delivery: mechanism and methods of evaluation. Int J Pharmacol Biol Sci 2011;3:458.
31. Muraleedhara KK. Mucoadhesive vaginal drug delivery system: a review on advanced status. Int J Pharm Res Anal 2013;3:33-46.
32. Covello T, Matricardi P, Mianacci C, Aliaquie F. Polysaccharide hydrogels for modified release formulations. J Controlled Release 2007;119:5-24.
33. Chaturvedi K, Ganguly K. Polymorphic hydrogels for oral insulin delivery. J Controlled Release 2013;165:129–38.
34. Patrick S. Martin’s physical pharmacy and pharmaceutical sciences. 6th ed. Lippincott Williams and Wilkins, Philadelphia; 2000. p.581.
35. Huang Y, Leobandung W. Molecular aspects of muco-bioadhesion: tethered structures and site-specific surfaces. J Controlled Release 2006;65:63-71.
36. Qaqish R, Amiji M. Synthesis of a fluorescent chitosan derivative and its application for the synthesis of chitosan-mucin interactions. Carbohydr Polym 1999;38:99-107.
37. Bembik Schurch A. Thiomers: a new generation of mucoadhesive polymers. Adv Drug Delivery Rev 2005;58:1569-82.
38. Patrick S. Martin’s physical pharmacy and pharmaceutical sciences. 5th ed. Lippincott Williams and Wilkins, Philadelphia; 2000. p. 26, 67.
39. http://pclcvruedu/tutorial/enhanced/files/polymers/orient/orient.htm [Last accessed on 20 Apr 2017]
40. https://en.wikipedia.org/wiki/Enthalpy. [Last accessed on 20 Apr 2017]
41. Madgulkar A, Bhailekar M. Synthesis and characterization of a novel mucoadhesive derivative of xylloglucan. Carbohydr Polym 2016;135:356–62.
42. Kaur H, Yadav S. Synthesis, characterization and evaluation of thiolated tamarind seed polysaccharide as a mucoadhesive polymer. Carbohydr Polym 2012;90:1543-9.
43. Kottke K, Edward M. Tablet dosage forms. In: Banker GS, Lachman L, Lieberman HA, eds. Pharmaceutical Dosage Forms: Tablets. New York: Marcel Dekker, Inc, 2002;1:287-333.

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