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## **BIO POLYMERS: A POTENTIAL CARRIER FOR PHARMACEUTICAL INDUSTRY**

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#### **Keywords:**

Bio Polymer, Excipient, Biocompatible, Gum & Mucilage, Sustain Release & Control Release Dosage Form

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ABSTRACT: Apart from the API, the excipient plays a crucial role in a formulation because of their versatile application, which enhances different physical and chemical characteristics in a dosage form. At present scenario natural polymers are Considered a leading constituent as an excipient to formulate several innovative approaches for designing the drug delivery system. The components which are obtained from nature is always preferred for our health due to their fewer side effects. Hence, biopolymers have the additional privilege of being incorporated as an excipient in pharmaceutical formulations. Demand for natural polymers is enhancing day by day (increased 7.1% in every year as per scientific data) because of their versatile application in drug delivery systems and multiple utilization of several domains in the biomedical section. Low toxicity, compatibility, sustainability, biodegradability, swelling index, release pattern are the major Parameters of the biopolymers which alter the different pharmacokinetic and pharmacodynamic characteristics of a pharmaceutical dosage form. The major aim of this current review is to impart some knowledge related to the widespread biopolymers. And emphasize the various fundamental applications of biopolymers in the pharmaceutical industry as an excipient and implementation of the biopolymers to design the innovative approaches for drug delivery systems. From this study, it was concluded that biopolymers have been effectively used as an excipient in the manufacturing of dosage forms and play a crucial role as a carrier mediated drug delivery system.

**INTRODUCTION:** Polymers are those substances in which the monomers (repeating molecular units) are attached with the support of a chemical or covalent bond and form a macromolecule, or a large molecule. Usually, polymers are categorized into three different groups like synthetic, semisynthetic, and natural polymers <sup>1</sup>. Synthetic polymers are those substances which are integrated by using the chemical processes in research laboratories utilizing different techniques. Polyethene, polystyrene, nylon, synthetic rubber,



polyvinyl chloride, Teflon, *etc.* are examples of synthetic polymers. Semi synthetic polymers are the substances which are extracted from natural sources, but they go through some chemical treatment to expand their physical parameters. Rubber, rayon, gun cotton, and different cellulose derivatives like cellulose acetate, cellulose nitrate, *etc.* are examples of semi synthetic polymers. Natural polymers are those components which originate from any type of natural derivative, like animal or plant sources<sup>2</sup>.

In recent years, natural polymers have drawn the main attention of the research scientist due to the enormous implementation of natural polymers in drug delivery systems. Nowadays, natural polymers are considered a key element as an excipient to develop innovative approaches for delivering drugs <sup>3</sup>. The importance of natural polymers is increasing

day by day because of their versatile application in drug delivery systems and the multiple utilization of several domains in biomedical sections <sup>4</sup>. Also, the scientists are immensely interested in the natural polymers on account of their remarkable utility beyond the synthetic or semisynthetic polymers. The benefits of natural polymers include that they are stable, nontoxic, compatible, and widely available at an affordable price (cheap)  $^{5}$ . Natural polymers also have the advantage that they are generally biodegradable in nature; these are known as biopolymers <sup>6</sup>. Now polymer serve as an important role for pharmaceutical industries, and it is treated as a backbone for drug delivery system. It is very essential for controlling the drug discharge rate from a formulation<sup>7</sup>.

So, in the modern era, the natural polymer acts as an inert transport system which delivers the API to a specific region or site in our biological system, which is familiar as a targeted drug delivery system. The natural polymer modified the pharmacokinetic and pharmacodynamic parameters, as well as altered the stability and solubility properties, and enhanced the plasma  $\frac{1}{2}$ life of a drug molecules <sup>8</sup>.

**Benefits of Natural Polymers Over Synthetic and Semi-Synthetic Polymers:** There are several benefits to using a natural polymer in a polymerbased drug delivery system rather than a synthetic or semisynthetic polymer. Natural polymers enhance the pharmacokinetic behaviours of drug molecules and also improve the distribution properties of drugs in biological systems <sup>9</sup>.

## **Advantages of Natural Polymers:**

**Biodegradable:** Biodegradation takes place in living organisms by enzyme degradation or chemical deterioration processes. Natural polymers do not exhibit any injurious responses to humans or the environment because they are degraded by living organisms, so they are eco-friendly in nature. But synthetic or semi synthetic polymers are composed of various types of chemical components which do not degrade within living organisms, so they are harmful to the surroundings and also to human beings<sup>10</sup>.

Non-toxic and Biocompatible: Natural polymers consist of replicate units of monomers, which are

generally carbohydrate-based in nature, so they are non-toxic in comparison with synthetic or semi synthetic polymers<sup>11</sup>.

Less side Effects and Safety: Natural polymers are extracted from natural origin and do not contain any type of chemical, so they are safe for humans. In the case of synthetic or semi-synthetic polymers, which are formulated by using different types of chemicals and undergo various chemical processes, they are harmful to the environment as well as human beings <sup>3</sup>.

**Commercial Benefits:** Natural polymers are more economical because their manufacturing costs are lower compared to synthetic or semi-synthetic polymers.

**Easily Available:** Biopolymers are originated from plant or animal origin, which are effortlessly obtainable in nature, and the sources are renewable. The cultivation of different plants is promoted by the government in developed countries, from which natural polymers are obtained (like tragacanth, guar gum, *etc*  $^{12}$ .

**Eco-friendly Processing:** Natural polymers are easily collected from various natural sources throughout the year in different seasons in sufficient quantities. The formulation process is also eco-friendly and simple compared to the synthetic polymer formulation.

**Better Acceptance and Patient Tolerance:** Natural polymers have better acceptability in comparison with synthetic and semisynthetic polymers. Some polymers are originated from natural, edible sources. So, this are more acceptable to the patient <sup>9</sup>.

**Characteristics of Natural Polymers:** Natural polymers exhibit different properties depending on their chemical, physiochemical, and mechanical characteristics.

**Chemical Property:** Polymers contain repeated molecular units, which are known as monomers. If one monomer is repeatedly present in a polymer, this is known as a homopolymer, and the arrangement of the different monomer units within a polymeric chain is known as a copolymer  $^{13}$ .

Basically, a polymer is the repetition unit of a monomer in a single or branch chain. They are interacting with each other through different forces of attraction. The intermolecular force between the polymeric chain is insistent by means of the polymer's monomer units and the attractive force present in the polymer chain. Generally, polymers consist of an extended chain, so the attraction, or intermolecular force, is enhanced between the molecules. The branch chain also creates hydrogen bonding, or ionic bonding, with its own chain. If any amide group or carbonyl group is present in a polymer, hydrogen bonding takes place <sup>14</sup>.

**Mechanical Property:** The mechanical property describes a polymer's ability to withstand different tensile strengths. The tensile strength and melting point are increased if a well-built intermolecular force or a strong hydrogen bond are present within the polymeric chain. Bond distortion in nature describes the elasticity property of a natural polymer. The rate of movement of drug particles throughout the polymeric bed is related to the transportation property, which is recognized as per the diffusion behaviour of a natural polymer.

**Transport Property:** Transport property is very important in the case of polymers because it is associated with diffusivity (which means the speed of the molecules throughout the polymer matrix). The transport property of a polymer is directly connected to the various applications of polymers like films and membranes<sup>15</sup>.

**Classification of Natural Polymers:** Natural polymers perhaps classified into three major groups according to their source of origin. Such as,

**Plants' Sources of Origin:** They are extracted from any type of plant source. e.g., gum acacia, gum tragacanth, pectin, agar, starch, *etc*.

Animals' Sources of Origin: They are isolated or extracted from any animal source. E.g., chitosan, albumin, gelatine, collagen, etc.

**Microbes' Sources of Origin:** They are derived from any type of microbe, like bacteria, fungi, molds, algae, etc., e.g., dextran, xanthan, pullulan, carrageenan, *etc*<sup>3</sup>.

According to its chemical composition, the biological polymer could be categorized into two main groups. Such as

**Natural Polymers Consist of Proteins:** Under this classification, the main composition of the natural polymer is protein.

The amount of protein is higher in comparison with other chemical group, which is also present in natural polymers. Other than protein, there are many groups present, like gliadin, glutenin, carbohydrates, *etc.*, e.g., soy protein, white gluten, zein, collagen, *etc*<sup>9</sup>.

**Natural Polymers Consist of Polysaccharides:** This classification is established on the presence of polysaccharides in the natural polymer.

The major chemical component is polysaccharides, along with different chemical groups. e.g., starch, pectin, cellulose, agar, chitosan, etc<sup>10</sup>.



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# Some Common Natural Polymers in the Pharmaceutical Industry:

Locust Bean gum: This polymer originates from Ceratonia siliqua Linn, associated with the family Leguminosae. It is also referred to as carob bean gum because it is excreted from the brown pod of the locust bean plant. This gum is pulverized, and it is prepared by grinding the seeds (endosperm) of the brown pod collected from the locust tree <sup>16</sup>. This polymer is a natural carbohydrate that contains galactose and mannose in a 1:4 ratio. The major component of locust gum is a neutral galactomannan polymer composed of 1,4-linked Dmannopyranosyl groups with a D-galactopyranosyl unit substituted on C6 of every fourth or fifth chain unit. It is a neutral polymer, and the solubility and consistency of locust gum are a little bit changed in the pH vallue  $3-11^{17}$ .

## Uses:

- It is extensively exploited in pharmaceutical formulations as a stabilizing agent, binder, flocculating agent, and thickening agent.
- It is also applied to formulated the propranolol hydrochloride-controlled delivery system to prevent first-pass metabolism.
- Locust gum is also employed to prepare matrix tablets with and without a cross-linking agent.
- Diclofenac sodium mini-matrix tablets for sustained release can be formulate by locust gum polymer.
- TIMERx® is an economically accessible tablet formulated by using locust and xanthan gum, which exhibited *in-vitro* & *in-vivo* extended delivery <sup>18</sup>.

**Karaya gum:** It is the dehydrated mucilaginous discharge derived from the tree *Sterculia urens roxb* under the family Sterculiaceae. It's also recognized by Indian Tragacanth, Sterculia, Karaya, or Bassora tragacanth gum and is available in Pakistan, India, and some places in Africa. Karaya gum is a fractionally acetylated polymer of rhamnose, glucuronic acid, and galactose. It consists of a major amount of D-glucuronic acid with D-galacturonic acid residue with branched heteropolysaccharide. No methoxy groups are

present in this polymer <sup>19</sup>. The gum is producing a viscous consistency when absorbing water, though it is commercially less soluble. It absorbed water very quickly and expanded many times of its initial volume. Karaya varies from tragacanth in such it has no starch & stains pink when used with ruthenium red solution <sup>20, 21</sup>.

#### Uses:

- In the pharmaceutical field, karaya gum is broadly used as a suspending, stabilizing, thickening, and mucoadhesive agent.
- It is applied as a dental adhesive, bulk laxative, and matrix-forming agent.
- Other than the pharmaceutical industry, it is also applied in the paper, food, and textile sectors. using karaya gum and hydroxypropyl methylcellulose (HPMC) as polymers <sup>1, 21</sup>.
- Karaya gum is used to prepare gastric drifting medication for a sustained-release drug delivery system by preparing a matrix system to deliver the drug for eight hours.

**Rosin:** It is also recognized by the Greek name pitch, or colophony, derived from the plants Pinus palustris (pine) and conifers. The major sources of this polymer are *Pinus soxburghii*, *Pinus longifolium, and Pinus toeda*. It is a non-evaporated component prepared by vaporizing volatile components with the help of heat, which is provided to the fresh liquid resin. Rosin is collected from oleoresin, and it is a biopolymer with a molecular weight of 400 DA. The major components are abietic and pimaric acids, which are present in resin, and a certain number of Non acidic components and esters are also present in this polymer <sup>1,4,5</sup>.

- Rosin considered as a polymer in various drug delivery techniques because of its physico-chemical properties and ease of accessibility.
- Rosin is a good film-forming biopolymer (due to the existence of esters), which is extensively utilized to formulate different kinds of controlled and sustained-release drug delivery techniques. This polymer was also applied to produce the film of the enteric coating tablet.

- It is also used to prepare transdermal dosage forms.
- It was recorded that rosin had been utilised to formulate the micro encapsules.
- When it was combined with dibutyl phthalate and polyvinyl pyrrolidone, it fabricated a smooth film that enhanced the tensile strength and elongation properties.
- Rosin has been utilised to develop a sustainedrelease microsphere of diclofenac sodium <sup>4, 5, 23</sup>.

**Honey Locust gum:** This polymer is obtained from the plant *Gleditsia triacanthos*, which belongs to the family Leguminosae. Honey locust gum is collected from the seeds, which contain fat, proteins, fibers, and carbohydrates. 88% D-galacto-D-mannoglycan, 1% cellulose, 4% pentan, 1% ash, and 6% proteins are present in this polymer. Plant seed galactomannan is created of a 1-4 linked D-mannan backbone and a 1-6 linked galactomannan. Honey locust gum is not freely soluble in cold water; for achieving maximum viscosity, solubilization and hydration are mandatory <sup>5, 17, 24, 25</sup>.

## Uses:

- Pharmaceutically, it is applied to form different types of sustained-release devices.
- Loctus gum is used to prepare a matrix tablet of theophylline, which sustains the drug delivery of theophylline in a controlled manner <sup>17, 24, 25</sup>.

*Tamarind gum: Tamarindus indica*, a member of the Leguminoseae family, is a biopolymer also known as Tamarind Kernel Powder (TKP). is present in the albumen of the kernel of the tamarind tree. It is polysaccharide in nature and consists of glucosyl, xylosyl, and galactosyl in a 3:2:1 ratio. Xyloglucan is present in this polymer, which is an important polysaccharide in the primary cell walls of higher plants.

Xyloglucan has a (1,4)-D-glucan backbone that is partially substituted at the O-6 position of its glucopyranosyl residues with "-D-xylopyranose." Tamarind gum is not soluble in organic solvents, but it has an extremely viscous consistency with a wide pH tolerance and gelling capacity when placed in hot water <sup>5, 26, 27, 28</sup>.

#### Uses:

- In the pharmaceutical industry, it is used as a thickening, stabilizing, binding, and gelling agent.
- This polymer has a large drug-retention capacity and tremendous thermal strength. For this reason, it was utilized as an excipient for a hydrophilic drug delivery system.
- This polymer is mucoadhesive, biocompatible, and non-carcinogenic in nature.
- Furthermore, it is employed for the preparation of hydrogels and spheroids.
- By using this polymer, different types of nasal preparation and mucoadhesive dosage forms for ocular purposes are prepared <sup>5, 26, 27, 28</sup>.

Carrageenans: This natural polymer is obtained from seaweed, Irish moss, or carrageen. The source of this polymer is *Chondrus crispus*, belonging to the family Rhodophyceae. Water or aqueous alkali is used to extract the carrageenans, and they are retrieved by alcoholic precipitation followed by the freeze drving or drum-drying method. sulfated polysaccharides Carrageenans are consisting of anhydrogalactose and galactose. This polymer produces a gel-like or viscous consistency in the occurrence of proteins because of the formation of a complex among the amino acids and the carrageenans. Generally, carrageenan is classified into three different types <sup>1, 4, 29</sup>.

- Carrageenans are used as additives for preparing different types of liquid dosage forms, like antacids and emulsions.
- It acts as a suspending, emulsifying, and thickening agent in the pharmaceutical field.
- Carrageenans increase homogeneity in colloidal suspensions and give a stable form of an insoluble drug through emulsion preparation.
- It is a very good alternative to gelatine for preparing capsules.
- Different types of cosmetics, like creams, toothpaste, and lotions, are also prepared by using carrageenans.

- Some studies indicate that carrageenans are appropriate for the manufacturing of controlled-release tablets.
- Carragenans were also used to formulate hydrogel beads for controlling drug delivery, which increased the thermostability of the hydrogel matrix <sup>1, 4, 30, 31, 32</sup>.

**Okra gum:** Okra gum is also recognised as Abelmoschus gum, collected from the plant *Abelmoschus esculentus*, which associate with family Malvaceae. This polymer is excavated from the fresh fruit, which is commonly known as laddish finger, a popular and easily available vegetable in India. Okra gum is polysaccharide in nature, composing L-galacturonic acid, D-galactose, and L-rhamnose acid, with some parts of glucose, mannose, xylose, and arabinose <sup>5, 33</sup>.

#### Uses:

- ✓ In the pharmaceutical domain, it is utilised as a suspending agent and binding agent for tablet preparation. Okra gum is a bioadhesive, filmforming agent, and controlled-release polymer.
- ✓ Okra gum is also used as a binding agent for the preparation of paracetamol tablets.
- ✓ Okra polymer is implemented to prepare a floating delivery system which prolongs the gastric retention time <sup>5, 33, 34, 35</sup>.

*Dillenia indica*: This natural polysaccharide originated from the fruit of *Dillenia indica* Linnaeus, which comes under the family Dilleniaceae. It is also recognized by the elephant apple. It is widely available in India, particularly in Assam and the northeast.

In Assam, this fruit is used to prepare different types of pickles, sauces, jams, and jellies <sup>36</sup>. Water was utilized as a major solvent for the extraction process of the mucilage from the Dillenia fruit, and acetone was used to precipitate the mucilage <sup>37</sup>.

*Dillenia indica* is a polysaccharide in nature which contains glycosides, steroids, flavonoids, and reduces sugar. Different chemical components like 3, 5, and 7-trihydroxy-3', 4'-dimethoxy flavone, betulinic acid,  $\beta$ -sitosterol, and stigmasterol are also present in this mucilage. *Dillenia indica* is also a very good source for pectin<sup>38</sup>.

#### Uses:

- Diallinia mucilage is a promising excipient in the pharmaceutical domain. The dillenia mucilage has a very good mucoadhesive property, which was utilized to formulate a nasal gel containing domperidone and oxytocin <sup>39, 40, 41</sup>.
- Dillenia indica is also applied as a polymer in prepared controlled-release formulations. Sharma et al. (2014 prepared a microsphere by using Dillenia indica. The prepared microsphere had excellent mucoadhesive properties, adhering for 3.5 hours <sup>42</sup>.
- This polymer is also implemented as a sustainrelease polymer for microsphere formulation<sup>43</sup>.

Fenugreek Mucilage: This mucilage is acquired from the seed of Trigonella foenum-graceum, generally recognized as fenugreek, an herbaceous plant belonging to the family Leguminous. Fenugreek seeds consist of a high amount of mucilage, but this mucilage fails to dissolve in water. It has a viscous, tacky consistency with fluids. The fenugreek seed also swells up and becomes slippery when interact to fluids <sup>44, 45</sup>. Different types of sugar, like fructose, galactose, lactose, and mannose, are present in fenugreek mucilage. Other than sugar, proteins and amino acids like alanine, arginine, aspartate, cysteine, etc. are also present in this mucilage. The presence of different chemical groups like steroids, terpenoids, flavonoids, tannins, coumarins, alkaloids, and saponins is identified from the aqueous extraction of fenugreek mucilage 46, 47.

- Fenugreek mucilage is utilized as an adjuvant in the pharmaceutical domain.
- It's exploited as a gelling agent, tablet binder, sustaining agent, emollient, and demulcent for different pharmaceutical preparations.
- It also has mucoadhesive properties  $^{48}$ .
- Fenugreek mucilage has the potential to be utilized as a suspending agent <sup>49</sup>.
- Fenugreek mucilage acts as a promising super disintegrating agent which showed very good anti-inflammatory action along with diclofenac sodium<sup>50</sup>.

Fenugreek mucilage is also incorporated into the prepared beads. The optimized beads showed better mucoadhesive properties, which controlled the release of metformin over a period of time after oral administration<sup>51</sup>.

*Hibiscus mucilage*: It is collected from the leaves of *Hibiscus rosa-sinensis*, which belongs to the family Malvaceae. This plant is also known as China rose or Chinese hibiscus, which is a popular flowering shrub. Some chemical components, like methyl sterculate, methyl-2-hydroxysterculate, 2hydroxysterculate malvate, cyclopropanoids, and rosasterol, are present in this shrub. The mucilage that is acquired from the fresh leafage of *Hibiscus rosa sinensis* consists of L-rhamnose, D-galactose, D-galactouronic acid, and D-glucuronic acid <sup>5, 44</sup>.

#### Uses:

- In the pharmaceutical domain, it is applied as an emulsifying agent, disintegrating agent, sustained release agent, suspending agent, etc.
- Commonly, hibiscus leaves are served as traditional medicine to treat constipation, different types of skin diseases, and to reduce the burning sensation <sup>5, 52</sup>.
- A dispersible tablet of aceclofenac is also formulated by using hibiscus mucilage, and it is applied as a super disintegrate agent, which shows better results than synthetic super disintegrants like Ac-di-sol <sup>53</sup>.
- Hibiscus mucilage can be utilized as a sustained-release polymer <sup>54</sup>.

Aloe Mucilage: Aloe mucilage originates from the leaf portions of *Aloe barbadensis Miller*, which associate with the family Liliaceae. The exudate was collected from the cells adjoining the vascular bundles and the central parenchyma tissue of this plant.

This exudate is a yellowish gel in nature and contains 1,8-dihydroxyanthraquinone derivatives together with their glycosides. Some scientists reported that partially acetylated mannan, or else acemannan, is the major polysaccharide, while others confirmed the pectic component as the prime polysaccharide. Arabinan, arabinorhamnogalactan, galactan, galactogalacturan, glucogalactomannan, galactoglucoarabinomannan, and glucuronic acidcontaining polysaccharides are also present in the aloe gel. The major active component is aloin, which is a blend of barbaloin, isobarbaloin, aloe emodin, and resins <sup>5, 27, 44</sup>.

## Uses:

- It has been utilized for the formulation of different topical products like gel, ointment, etc.
- Aloe vera mucilage is very popular in the cosmetic industry.
- It is not only an excipient; aloe mucilage also serves as an anti-inflammatory and anti-diabetic agent <sup>55, 56</sup>.
- It is also beneficial for the composition of tablets and capsules.
- It was established that aloe mucilage has the capability to retard drug release and has very good swelling properties <sup>57</sup>.

Albizia gum: The major origin of albizia gum is the plant *Albizia zygia*, belonging to the family Leguminosae. The gum is collected from the trunk portion of this tree, and it is round, elongated, and of variable color, ranging from yellow to dark brown. It is composed of  $\beta$ -1-3-linked D-galactose units with a few  $\beta$ 1-6-linked D-galactose units. Albizia gum has the possibility of being a substitute for Arabic gum as a natural emulsifying agent considered in the food and pharmaceutical unit <sup>27</sup>, <sup>58</sup>.

- 1. In the pharmaceutical domain, it is applied as a binding and suspending agent. This gum is also applied as a coating substance for drug delivery design that target the colon.
- 2. Albizia and khaya gums were studied as coating substances for direct compression colon-targeted tablets, where paracetamol and indomethacin were used as model drugs.
- 3. These gums were exploited as a coating material, which was broken down within the colon by microflora and Delivered the drug 59, 60.

**Khaya gum:** *Khaya senegalensis* & *Khaya grandifoliola*, both associated with the family Meliaceae, are the main sources for Khaya gum.

It is a polysaccharide where galacten is present at the 1, -3 linked  $\beta$ -D galactopyranosyl residues, which are concentrated in the inner chain. Khaya gum is also composed of both D-glucoronic and D-galatoronic acids. Different studies about Khaya gum indicate that it has many similarities to acacia gum. The methylation of the gum also proved that it was similar to acacia gum. Khaya gum is semi-transparent in nature <sup>27, 61</sup>.

Uses:

- 1. In the pharmaceutical domain, it is exploited as a binding agent, suspending agent, and disintegrating agent, and in addition to other polymers, it shows good results for control or sustained release formulations.
- **2.** Khaya gum was able to control the drug delivery for up to 5 hours  $^{62}$ .
- **3.** Khaya gum has a potential binding property to formulate tablets <sup>63</sup>.
- 4. Khaya gum exploited as a suspending agent at 0.2% w/v concentration<sup>64</sup>.

**Gum damar:** It is derived from the plant *Shoreawiesneri*, which belongs to the family Dipterocarpaceae. Generally, it is whitish to yellowish in color. The major chemical component of gum damar is resin. It is composed of 40% alpha resin, which dissolves in alcohol, and 22% beta resin. Dammarolic acid is also exhibit in gum damar <sup>27</sup>.

Uses:

- In the pharmaceutical domain, it is mainly used as an emulsifier, stabilizer, and binding agent for the manufacturing of dosages.
- It produces a water-resistant coating, which is very popular for forming dental accessories.
- Gum damar has the potential property to formulate a sustain-release matrix  $^{65}$ .
- Damar gum has a very good film-forming capacity for coating purposes <sup>66</sup>.
- Gum damar can be utilised to formulate microparticles which use a microencapsulating agent for sustained delivery <sup>67</sup>.

**Hakea gum:** It is the dried exudate collected from the plant *Hakea gibbose*, which belongs to the family Proteaceae. Hakea gum is composed of Larabinose and D-galactose linked, as in gums that are acidic arabinogalactans. Glucuronic acid, arabinose, mannose, galactose, and xylose are the different sugars that are also present in the Hakea gum in a 12:43:32:5:8 ratio. Hakea gum is partially soluble in water  $^{27, 68, 58}$ .

## Uses:

- Hakea gum is applied as a sustained-release & bioadhesive material in the development of buccal tablets<sup>68</sup>.
- Hakea gum was implemented to formulate a buccal tablet of chlorpheniramine maleate <sup>69</sup>.
- It was also perceived that hake a gum has potential properties for sustaining the delivery of drug molecules and also has bio adhesive and mucoadhesive properties <sup>70</sup>.

Hupu gum: Hupu gum is also recognise as Kondagugu gum, is a natural polysaccharide obtained from the plant Cochlospermum gossypium, which associated with the family Bixaceae. Hupu gum is fabricated with sugars like galactose, arabinose, rhamnose, mannose-D glucose, D-glucouronic acid, and D-galactouronic acid. Tannin, protein, uronic acid, and some soluble fibers are also present on Hupu gum<sup>27, 58, 71</sup>.

## Uses:

- **1.** Hupu gam is utilize for the medication of diarrhoea, dysentery, cough, pharyngitis, etc.
- 2. In the pharmaceutical industry, it is utilized as an excipient, which is applied as a substitute for gum tragacanth.
- **3.** Hupu gum is utilized as a sustained-release polymer which can formulate a gastric floating delivery system.
- **4.** It has very good emulsifying properties, even at very low concentrations <sup>72</sup>.
- **5.** Kondagogu gum used as a mucoadhesive polymer to formulate microspheres has potential mucoadhesive properties <sup>73</sup>.

**Tara gum:** The source of tara gum is the endosperm part of the seed of *Caesalpinia spinosa*,

which belongs to the family Fabaceaeor Leguminosae.

It is a natural polysaccharide that mainly consists of galactomannan, which is a dietary fibre. The major chemical composition is comparable to the main constituents of loctus bean gum and guar gum, which consist of a linear main chain of (1-4)- $\beta$ -D-mannopyranose units with  $\alpha$ -D-galactopyranose units attached by (1-6) linkages. Galactose and mannose are present in tara gum in a 1:3 ratio. Generally, tara gum is extremely viscous in nature, and it produces a very viscous solution even at very low (1%) concentrations <sup>5, 27, 58</sup>.

Uses:

- In the pharmaceutical domain, it is utilized as an emulsifying component, a thickening agent, and a stabilizer in various formulations.
- Tara gum is also applied to formulate controlled-release emulsions and tablets.
- Tara gum is implemented as a controlledreleased polymer which formulates a matrix tablet of ambroxol hydrochloride<sup>74</sup>.
- Fernandes et al. (2021 demonstrated the application of tara gumin toothpaste for delivery of fluoride <sup>75</sup>.

**Moi gum:** *Lannea coromandelica*, which is also recognized as an Indian ash tree belonging to the family Ancardiaceae, is the primary source of Moi gum. The aforementioned gum is acquired from various parts of these plants, including the leaves, fruits, stems, and bark. Generally, the color of fresh gum becomes yellowish white, but after drying, it changes into a dark color. Moi gum is a type of arabino-galactan, which is similar to gum Arabic. The major chemical constituent is D-galactose, which is present in 69.5%; 2.5% L-rhamnose; 11% L-arabinose; 17% 4-o-methyle uronic acid; and 1.38% protein, which is furthermore present in moi gum. Arabino-3,6-galactan is produced after the hydrolysis of the moi gum and mucilage <sup>58, 76</sup>.

Uses:

✓ Although moi gum is not extensively used in the pharmaceutical field, researchers have conducted a number of studies to evaluate its properties. ✓ Moi gum is utilized as a microencapsulating agent to formulate polymers which sustain the drug released <sup>77</sup>.

Leucaena leucocephata gum: Leucaena *leucocephata* is the main source of this gum, which belongs to the family Fabaceae. The gum is present on the seeds and leaves of these plants. Mimosine is a toxic and non-protein component which is present in the seeds, bark, and leaves of Leucaena plants. Tanin and oxalic acid are exhibit in the seed. Generally, 25% gum is present on the seed, which is colourless to reddish brown in color. The pH of this gum is 4.2, which is acidic in nature. Leucaena leucocephata gum is moderately soluble in water but practically insoluble in semi-polar solvents like chloroform, acetone, ethanol, etc., but in the presence of water, this gum swells up to five times its original size. The viscosity is dependent on the concentration; it will increase with the concentration of gum<sup>27, 78</sup>.

#### Uses:

- The seed gum of *Leucaena leucocephata* is applied as a suspending, emulsifying, disintegrating, and binding agent in the pharmaceutical domain.
- It's used as an emulsifying agent, which showed better emulsifying properties compared to gum acacia <sup>79</sup>.
- Verma et al. 2007 formulated a tablet applying ibuprofen as a standard drug for evaluating the disintegrating properties of Leucaena leucocephata gum<sup>80</sup>.
- Other than the excipient, Leucaena leucocephata gum also has medicinal properties. This gum is very effective for treating stomach upset and as a contraceptive agent.
- The extract substance of the seeds has antidiabetic, anti-microbial, and antioxidant properties<sup>81</sup>.

**Bhara gum:** The bark portion of *Terminalia bellerica*, associated with the family Combretaceae, is the primary source of the bhara gum. Generally, it is yellowish in color.

Uses:

- ✓ Bhara gum is utilized in the cosmetic industry as an emulsifier, demulcent, and purgative agent.
- ✓ Bhara gum is implemented as a sustainedrelease polymer which sustains drug delivery for 10 hours<sup>83</sup>.

**Delonix regia gum:** Delonix regia gum is procured from the seed part of the plant Deloxia regia in the family Leguminosae. It is also recognized by the flamboyant flame tree, flame tree, flame of the forest etc., but in India it is popular by the name of Gulmohor. The color of this gum is yellowish or reddish brown. Generally, this gum is thick and viscous in nature. Delonix regia gum is an inherent polysaccharide composed of protein, fiber, and crude fat. This gum is water-soluble in nature<sup>84, 85</sup>.

Uses:

- **1.** In the pharmaceutical sector, it is applied as a binding and disintegration agent for the manufacturing of tablets.
- **2.** In higher concentrations, it is exploited as a polymer in order to develop sustained-release formulations <sup>86</sup>.
- **3.** *Delonix regia* gum also has the binding capacity to formulate tablets, and in lower concentrations, it is balanced between binding and disintegration properties when the fastest disintegration is required <sup>87</sup>.
- **4.** Delonix regia gum has the potential to be applied a suspending agent  $^{88}$ .
- 5. Adetogun *et al.* 2007 established that *Delonix regia* gum may be applied as a binding agent as a substitute for tragacanth and gum acacia  $^{89}$ .

**Almond gum:** It is procured from the wood *Prunus amygdalus* or *Amygdalus communis*, which belongs to the family Rosaceae. This gum is a clear extrusion from the injured parts of the trunk and branches of the sweet almond tree. This gum is

typically white, red, or yellow in color and is water-soluble. Almond gums contain aldobionic acid, L-galactose, L-arabinose, D-mannose, and other sugars, but galactose and arabinose are the primary sugars found in almond gum. Protein, fat, and different minerals like sodium, potassium, magnesium, calcium, and iron are present in this gum. Furthermore, it is water-soluble in nature <sup>58</sup>, <sup>90, 91</sup>.

## Uses:

- **1.** In the pharmaceutical sector, almond gum is applied as a suspending, emulsifying, thickening, adhesive, and stabilizing agent.
- **2.** This gum is also applied as a controlled and sustained-release polymer
- 3. Almond gum was used to formulate the indomethacin sustained-release matrix tablet, which had potential for increasing therapeutic activity as well as reducing the dosing frequency  $9^{2}$ .
- 4. As per the study, 2% w/v almond gum binder showed the optimal result as a binding component in tablets <sup>93</sup>.

*Cassia tora* **mucilage:** The source of this gum is the seeds of *Cassia tora*, belonging to the family Caesalpiniaceae. *Cassia tora* mucilage is popularly known by the name of charota. It is translucent, amorphous, and soluble in water. Generally, Cassia toramucilage is soluble in cold water, but it forms a colloidal dispersion in warm water and is insoluble in acetone, ethanol, methanol, etc. The major chemicals are polysaccharides like glucomannon and galactomannon, and different chemicals like cinnamaldehyde, tannins, coumarins, mannitol, and resins are also present in *Cassia tora* gum. Some essential oils like eugenol, pinene, etc. and volatile oils like cassia oil are present in cassia tora gum <sup>58</sup>, <sup>94, 95</sup>.

## Uses:

This mucilage has the potential properties of a suspending and binding agent. Several studies were done to evaluate this property and reported that cassia tora gum works as a binding and suspending agent in pharmaceutical preparations <sup>58</sup>.

- Cassia tora acts as an economic binding agent with an 8% w/v concentration, and in this specific concentration, it was almost equivalent to the 8% w/v guar gum mucilage <sup>96</sup>.
- Cassia tora also used as super disintegrating agent to formulate orodispersible tablet of Rosuvastatin <sup>97</sup>.
- Tora mucilage's suspending ability is evaluated by comparing it to that of tragacanth, gum acacia, and gelatine. After evaluation, the degree of susceptibility can be arranged in the following order: Cassia tora> Tragacanth gum> Acacia gum> Gelatine <sup>58</sup>.

TABLE 1: RECENT ADVANCEMENT OF BIOPOLYMER IN PHARMACEUTICAL DOSAGE FORM DESIGN
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Name of biopolymer	Application in dosage form design
Locust Bean gum	Antibacterial Wound Dressing Film <sup>98</sup> .Super disintegrating agent in Oro dispersible tablet <sup>99</sup> .
	Fast disintegrating tablet <sup>100</sup> . Matrix Tablet <sup>101</sup> , Hydrogel <sup>102</sup> , Mucoadhesive Buccal Tablets <sup>103</sup> ,
	Niosome loaded Hydrogel for topical apllication <sup>104,</sup> Nanoparticls <sup>105</sup> ,
Karaya gum	Floating drug delivery system <sup>22</sup> , Sustain release matrix tablet <sup>101</sup> , Vaginal Film <sup>106</sup> ,
	Hydrogel <sup>107</sup> , Mucoadhesive tablet <sup>108</sup> , Microparticles <sup>109</sup> , Effervescent Floating Matrix
	Tablets <sup>110</sup> ,
Honey locust gum	Microspheres <sup>111</sup> , Sustain release matrix tablet <sup>25</sup> , Microparticle <sup>112</sup> .
Tamarind gum	Nanoparticles <sup>113,114</sup> , Hydrogel <sup>115,116</sup> , Coating materials for gauze <sup>117</sup> , Ocular delivery <sup>118,119</sup> ,
	Microcapsules <sup>120</sup> , Mucoadhesive Tablet <sup>121</sup> , Emulsifier in nanoemulsion <sup>122</sup> , Disintegrating
Moringa gum	Nanovel <sup>124</sup> Encapsulating Agent <sup>125,126</sup> Nanoparticles <sup>127</sup> Hydrogel <sup>128</sup> Nanometric carrier <sup>129</sup>
Worliga gain	colon specific drug delivery system <sup>130</sup> . Binding agent for tablet formulation <sup>131</sup> .
Okra gum	Nanoliposom <sup>132</sup> , Nanoparticles <sup>133, 134, 135</sup> , Mucoadhesive Tablet <sup>136</sup> , Binding agent <sup>137</sup> , Film
	formingagent <sup>138</sup> .
Dillienia indica gum	Nasal Gel <sup>39,40,41</sup> , Microparticles <sup>42,43</sup> , Microbeads <sup>139</sup> , Nanoparticles <sup>140,141</sup> , Buccal patches <sup>142</sup> .
Fenugreek mucilage	Matrix tablet <sup>48</sup> , Suspending agent <sup>49</sup> , Super disintegrating agent <sup>50</sup> , Mucoadhesive beads <sup>51</sup> ,
	Hydrogel <sup>143</sup> , Buccal Patches <sup>144</sup> , Bio adhesive tablet for sustain release <sup>145</sup> , Nanoparticulate
	system for ocular drug delivery system <sup>146</sup> .
Hibiscus mucilage	Disintegrating agent <sup>53, 147</sup> , Wound healing activity <sup>148</sup> , Mucoadhesive Beads <sup>149</sup> , Sustain
	release matrix tablet <sup>150</sup> ,
Aloe Mucilage	Matrix tablet <sup>57</sup> , Hydrogel <sup>151,152</sup> , Microencapsulating agent <sup>153</sup> , Binding agent <sup>154</sup> , Bioactive Film <sup>155</sup> ,
Albizia gum	Coating Materials for tablet <sup>59, 60</sup> , Oral Dissolvable Films <sup>156</sup> , Microbeads <sup>157</sup> , Binding agent <sup>158</sup> ,
Khaya gum	Binding agent <sup>63</sup> , Suspending agent <sup>64</sup> , Oral Dissolvable Films <sup>156</sup> , Colon targeted drug delivery
	system (Tablet) <sup>158</sup> , Matrix tablet <sup>159</sup>
Gum damar	Film forming agen <sup>66</sup> , Microencapsulating agent <sup>67</sup> , Sustain release tablet <sup>160</sup> , Nanocapsules <sup>161</sup> ,
	Sustained release matrix <sup>162</sup> ,
Hakea gum	Sustain release and mucoadhesive property <sup><math>10</math></sup> ,
Hupugum	Mucoadhesive Microcapsules <sup>7,5</sup> , Microcrystals <sup>105</sup> , Solid mixture of poorly water soluble
-	$drug^{104}$ , Colon targeted drug delivery system <sup>103</sup> ,
Tara gum	Controlled delivery matrix <sup>1</sup> , Toothpaste <sup>1</sup> , Thickening agent in emulsion preparation <sup>10</sup> ,
	Microencapsulating agent <sup>-1</sup> , Nanocrystal film <sup>-1</sup> ,
Moi gum	Microsphere , $\frac{79}{12}$ is interaction $\frac{80}{10}$ for the large $\frac{169}{10}$
Leucaena leucocepnatagum	Emuisitying agent, disintegrating agent, Sustain release carrier,
Dilaraguili Delonix regia sum	Microsphere Binding agant for formulation of tablat <sup>86,88</sup> Suspanding agant <sup>87</sup>
Detonix regia guin	Mucoodhesiye papostructure lipid carrier <sup>170</sup> sustained release tablet <sup>171</sup> pH and time
	dependent colon targeted drug delivery system <sup>172</sup>
Almond Gum	Sustain release matrix tablet <sup>92</sup> . Binding agent for tablet <sup>93</sup> . Bio composite Film <sup>173</sup> Sustained
	release pellets <sup>174</sup> . Nanoparticles <sup>175</sup> .
Cassia tora mucilage	Binding agent <sup>96</sup> , encapsulating agent <sup>176</sup> .

**CONCLUSION:** In the last few decades, natural polymers have drawn huge attention from research scientist due to their tremendous implementation in drug delivery approach. In the present scenario, natural polymers are considered essential

ingredients which play a critical role in developing different innovative approaches for delivering the drug into the targeted area. Biopolymers are more acceptable because of their numerous advantages over synthetic and semi-synthetic polymers. But

the selection of a natural polymer is an important criterion for incorporating it into the drug delivery approach. In the polymer and drug interaction study, the degradation pattern of the polymer is also playing a crucial role in selecting a polymer as the carrier of a drug delivery technique. From the above discussion, it can be concluded that several natural polymers have a diverse role in the pharma industry in different prospects as well as drug delivery systems. It was established that natural polymers have been efficiently utilized as an excipient in the manufacturing of dosage forms and play a crucial role as a carrier-mediated drug system. Numerous delivery polysaccharides obtained from plant origins have been prospective criteria as carriers for sustained or controlledrelease drug delivery systems. So, there will be a huge scope for research scientists to develop innovative approaches by using Biopolymers as carriers to deliver the drug in a controlled or sustained manner.

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#### **REFERENCES:**

- 1. Gupta M, Thakur A and Gupta J: Recent Advances of Polymers in Pharmaceutical Area: A Review. International Journal of Advance Research and Innovative Ideas in Education 2023; 9(3): 2395-4396.
- Alkadi H: A Review on the Role of Polymers in Pharmaceutical Applications. Venoms and Toxins 2021; 1: 41-55.
- 3. Opris O, Mormile C, Lung I, Stegarescu A, Soran M and Soran A: An Overview of Biopolymers for Drug Delivery Applications. Applied sciences 2024; 14(1383): 1-33. https://doi.org/10.3390/app14041383
- Deb PK, Kokaz SF, Abed SN, Paradkar A and Tekade RK: Pharmaceutical and Biomedical Applications of Polymers. Basic Fundamentals of drug delivery 2019; 203-267. doi.org/10.1016/B978-0-12-817909-3.00006-6
- Fazal T, Murtaza B T, Shah M, Iqbal S, Rehman M, Jaber F, Dera A, Awwad N S and Ibrahium H A: Recent developments in natural biopolymer based drug delivery systems. Royal Society of Chemistry 2023; 13(33): 23087–23121. doi: 10.1039/d3ra03369d
- Jain S, Jain A, Jain R and Chauhan N: Potential of natural polymeric materials in pharmaceutics. Pharmacological Research - Natural Products 2024; 2(2024): 100014. doi.org/10.1016/j.prenap.2024.100014
- 7. Esther k, and Ayodeji: Review of the Impact of Polymers on the Pharmaceutical Industry. International journal of novel research and development 2024; 9(1).
- 8. Pal R, Pandey P, Saxena A and Thakur S: The Pharmaceutical Polymer's; A Current Status in Drug Delivery: A Comprehensive Review; Journal of Survey in Fisheries Sciences 2023; 10(1): 3682-3692.

- Alaswad S, Mahmoud A and Arunachalam P: Recent Advances in Biodegradable Polymers and Their Biological Applications: A Brief Review. Polymer 2022; 14(22): 1-15. https://doi.org/10.3390/polym14224924
- Samir A, Ashour F, Hakim A and Bassyouni M: Recent advances in biodegradable polymers for sustainable applications. Materials degradation 2022; 6(68): 1-28.doi.org/10.1038/s41529-022-00277-7
- Gupta M K, Suryawanshi M and Shrivastava B: A Bird Eye View on Natural Gums and Mucilage used in Drug Delivery System. International journal of pharmaceutical science & nanotechnology 2023; 16(1). doi.org/10.37285/ijpsn.2023.16.1.10
- 12. Dhanal S, Tamboli F, More H, Kamble S and Shere S: Use of natural polymers as excipients in the pharmaceutical industry. International journal of advanced scientific research 2021; 6(2): 55-58.
- Reddy M B, Ponnamma D, Choudhar R, and Sadasivun K: A Comparative Review of Natural and Synthetic Biopolymer Composite Scaffolds. Polymers 2021; 13 (1105). doi.org/10.3390/polym13071105
- Sivakumar P: Overview on pharmaceutical polymers. Journal of Pharmacy and Pharmaceutical Sciences 2021; 10(5):953-975. doi:10.20959/wjpps20215-18964
- 15. Senv D, Mohite B and Kayande N: Review on polymer. International Journal of Pharmaceutical Sciences and Medicine 2019; 4(10): 1-15.
- Soumya RS, Raghu KG and Abraham A: Locust bean gum-based micro- and nanomaterials for biomedical applications. Micro and Nano Technologies 2022; 241-253. doi.org/10.1016/B978-0-323-90986-0.00012-1
- Petitjean M and Isasi JR: Locust Bean Gum, a Vegetable Hydrocolloid with Industrial and Biopharmaceutical Applications. Molecules 2022; 27(23): 8265. doi.org/10.3390/molecules27238265
- Duhan N, Barak S and Mudgil D: Chemistry, Biological activities, and uses of locust been gum. Gums, resins and latexes of plant origin 2022; 225-239. doi.org/10.1007/978-3-030-91378-6\_9
- Prasad N, Thombare N, Sharma SC and Kumar S: Recent development in the medical and industrial applications of gum karaya: a review. Polymer Bulletin 2022; 80(150): 3425-3447. doi:10.1007/s00289-022-04227-w
- 20. Sahu P, Pisalkar PS, Patel S and Katiyar P: Physicochemical and Rheological Properties of Karaya Gum (*Sterculia urens* Roxb.). International Journal of Current Microbiology and Applied Sciences 2019; 8(4): 672-681.
- 21. Raj V, Lee JH, shim JJ and Lee J: Recent findings and future directions of grafted gum karaya polysaccharides and their various applications: A review. Carbohydrate Polymer 2021; 258. doi.org/10.1016/j.carbpol.2021.117687
- 22. Gangadharappa HV, Rahamath-Ulla M, Kumar PTM and Shakeel F: Floating drug delivery system of verapamil hydrochloride using karaya gum and HPMC. Clinical Research and Regulatory Affairs, 2010; 2(1): 13–20. doi:10.3109/10601331003604762
- Nande V, Barabde U, Morkhade D, Patil A and Joshi S: Sustained Release Microspheres of Diclofenac Sodium Using PEGylated Rosin Derivatives. Drug Development and Industrial Pharmacy 2007; 33(10): 1090-1100. doi.org/10.1080/03639040601180101
- 24. Gorde N and Pawar H: Locust bean gum: a comprehensive review. International Journal of Medical, Pharmaceutical and Biological Sciences 2023; 3(1): 1-8.

- 25. Linge V, Bhagwat D and Kore U: Natural gum and mucilage for biopharmaceutical applications. International journal of creative research thoughts 2022; 10(8): 444-458.
- 26. Amiri M, Mohammadzadeh V, Yazdi M, Barani M, Rahdar and Kyzas G: Plant-Based Gums and Mucilages Applications in Pharmacology and Nanomedicine: A Review. Molecules 2021; Mar; 26(6): 1770. doi: 10.3390/molecules26061770
- 27. Muruganantham, Krishnaswami V, Manikandan A, Aravindaraj, Suresh J, Murugesan M and Kandasamy R: Gum as pharmaceutical excipient: An overview. Gums, resins and latexes of plant origin 2022; 145-149.
- 28. Naeem A, Batool R, Aziz M, Tahseen S, Rohi M, Zafar A, Mubeen R and Saleem B: *Tamarindus indica* seed extraction, application, and physiochemical characterization: a review. Journal of population therapeutics and clinical pharmacology 2023; 330(18). doi.org/10.53555/jptcp.v30i18.3379
- Volodko A, Son E, Glazunov V, Davydova V, Sinkler E, Aleksandrova S, Blinova M and Yermak I: Carrageenan films as promising mucoadhesive ocular drug delivery systems. Colloids and Surfaces B: Biointerfaces 2024; 237. doi.org/10.1016/j.colsurfb.2024.113854
- Neamtu B, Barbu A, Mihai N, Neamtu C, Popescu D, Zahan M, and Miresan V: Carrageenan-Based Compounds as Wound Healing Materials. International journal of molercular science 2022; 23(16): 9117. doi: 10.3390/ijms23169117
- Qureshi D, Nayak SZ, Maji S, Kim D, Banerjee and Pal K: Carrageenan: A wonder polymer from marine algae for potential drug delivery application. Current pharmaceutical design 2019; 25(11): 1172-1186. 10.2174/1381612825666190425190754
- Khalid M, Patel T J, Chand G, Ahmad M, Kumar D, Aijaz M, Mishra N and Thakur A: Formulation and Evaluation of Ramosetron Hydrochloride Sustained Release Matrix Tablet Using Different Ratios of Chitosan and K-Carrageenan Polymer. European chemical bulletin 2023; 12(8): 6033-6055. doi:10.48047/ecb/2023.12.si8.5142023.25/07/2023
- 33. Sonika, Dhiman S, Singh TG, Arora G and Arora A: *Moringa gum*: a comprehensive review on its physicochemical and functional properties. Plant Archives 2020; 20(1); 3794-3805.
- Shah R, Adnaik R, Adnaik P and Patil S: Extraction, isolation and characterization of okra mucilage as potential source binder in tablet. Asian journal of pharmaceutical technologies 2023; 13(3): doi: 10.52711/2231-5713.2023.00032
- 35. Manyam S, Arumilli S, Pakalapati P and Sarella P: Formulation and Evaluation of Sustained Release Atorvastatin Tablets Using Natural Polymers, with a Focus on Okra Gum. International Journal of Pharmaceutical Sciences and Medicine 2023; 8(10): 11-22. doi:10.47760/ijpsm.2023.v08i10.003.
- 36. Rai H, Upadhyay S and Sajwan A: An overview of *Dillenia indica* and their properties. The Pharma Innovation Journal 2020; 9(6): 41-44.
- Gogoi B, Saharia V, Patowary P and Mawlieh B: *Dillenia indica* linn. -a multipurpose medicinal plant of assam. British journal of medical and health research 2020; 7(5): 51-63. doi:10.46624/bjmhr. 2020.v7.i5.006
- Saikumar A, Nickhil C and Badwaik L: Physicochemical characterization of elephant apple (*Dillenia indica* L.) fruit and its mass and volume modeling using computer vision. Scientia Horticulturae 2023; 314. doi.org/10.1016/j.scienta.2023.111947

- 39. Saikia D, Kesavan R, Inbaraj B, Dikkala P, Nayak P and Sridhar K: Bioactive Compounds and Health-Promoting Properties of Elephant Apple (*Dillenia indica* L.): A Comprehensive Review. Foods 2023; 12(16): 2993. doi: 10.3390/foods12162993
- 40. Fahaduddin and Bal T: Fabrication and evaluation of *Dillenia indica*-carrageenan blend hybrid superporous hydrogel reinforced with green synthesized MgO. International Journal of Biological Macromolecules nanoparticles as an effective wound dressing material 2024; 265(2). doi.org/10.1016/j.ijbiomac.2024.130835
- 41. Bora A, Kalita P, Adhikari R, Das A, Zaheer R, Laskar M and Patha K: Harnessing the Therapeutic Potential of Dillenia indica: An Overview of Recent Dosage Form Developments. Current drug discovery technologies 2024; doi: 10.2174/0115701638292980240407135246
- 42. Ghosh D, De S, Deka D and Das G: Amphiphilic polyphenol incorporated hydrogel derived from mucoadhesive of *Dillenia indica*. Potential antioxidant and adsorbent. International Journal of Biological Macromolecules 2024; 254(1). doi.org/10.1016/j.ijbiomac.2023.127759
- Chakraborty S, Khandai M, Dhibar M, Yadav S, Kumari H: Isolation and Characterization of Dillenia Fruit Mucilage: A Novel polymer for microsphere Delivery of Losartan potassium. Drug delivery letters 2020; 10(4): 314-325. doi: 10.2174/2210303110999200802030342
- 44. Pawan P, Mayur P and Ashwin S: Role of natural polymers in sustained release drug delivery system: applications and recent approaches. International Research Journal of Pharmacy 2011; 2(9): 6-11.
- 45. Prajapati V, Desai S, Gandhi S and Roy S: Pharmaceutical applications of various natural gums and mucilages. Gums, Resins, and latexes of plant origin 2022; 25-57. doi.org/10.1007/978-3-030-91378-6\_2
- 46. Zemzmi J, Rodenas L, Blas E, Najar T and Pascual JJ: Characterization and *in-vitro* evaluation of fenugreek (*Trigonella foenum-graceu*) seed gum as a potential prebiotic in growing rabbit nutrition. Animals 2020; 10(6): 1-15.doi: 10.3390/ani10061041
- 47. Ayoub A and Rahman S: Study of the Physical and Functional Properties of Fenugreek Gum. IOP Conference Series Earth and Environmental Science 2023; 1252(1):012170. doi:10.1088/1755-1315/1252/1/012170
- 48. Dhull S, Bamal P, Kumar M, Bangar S, Chawla P, Singh S, Mushtaq W, Ahamed M, and Sihag S: Characterization of Fenugreek Seeds Mucilage and its Evaluation as Suspending Agent. Legume Science 2022; 1-14. doi.org/10.1002/leg3.176
- 49. Gorakhnath GS and Hingane LD: Characterization of Fenugreek Seeds Mucilage and its Evaluation as Suspending Agent. International Journal for Research in Applied Science & Engineering Technology 2022; 10(6):4382-4386 doi.org/10.22214/ijraset.2022.44934
- 50. Kumari N and Sharma R: An Immediate Release tablet of Carvedilol with Natural Superdisintegrants Fenugreek Seed Mucilage and synthetic Superdisintegrants. Asian journal of pharmacy and technology 2020; 10(3):156-164. doi: 10.5958/2231-5713.2020.00027.6
- Nayaka AK, Pal D and Das S: Calcium pectinatefenugreek seed mucilage mucoadhesive beads for controlled delivery of metformin HCL. Carbohydrate Polymers 2013; 96(1): 349-357. doi.org/10.1016/j.carbpol.2013.03.088
- 52. Yahaya N, Anuar A and Saidin N: Hibiscus Rosa-Sinensis mucilage as a functional polymer in pharmaceutical applications: a review. International Journal of Applied

Pharmaceutics 2023; 15(1): 44-49. doi:10.22159/ijap.2023v15i1.46159

- 53. Bhadoria J and LikhariyaM:A Review on Orodispersible Tablet by Using Hibiscus rosa sinesis as Natural Superdisintegrant. International Journal of Pharmaceutical Sciences & Medicine 2022; 7(6):59-69.
- 54. Sahu A, Chouksey K and Ganju K: Formulation and evaluation of ciprofloxacin hydrochloride sustained release tablets using *Hibiscus rosa-sinensis* mucilage. Journal of Advanced Scientific Research 2022; 13(8): 71-78.
- 55. Sanchez M, Burgos E, Iglesias I and Serranillos M: Pharmacological Update Properties of Aloe Vera and its Major Active Constituents. Molecules 2020; 25(6): 1324.doi: 10.3390/molecules25061324
- 56. Prakash K and Tiwari S: Pharmaceutical assessment of Aloe Vera Skin Gel: A herbal formulation and its potential benefits. World Journal of Biology Pharmacy and Health Sciences 2023; 15(3):43-50. doi:10.30574/wjbphs.2023.15.3.0375
- 57. Mahmood A, Erum A, Tulain U, Shafiq S, Malik S, Khan M and Alqahtani M: Aloe vera-based polymeric network: a promising approach for sustained drug delivery, development, characterization, and *in-vitro* evaluation. Recent Advances in Gels Engineering for Drug Delivery 2023; 9(6), 474; https://doi.org/10.3390/gels9060474
- Mundhar D, Wankhed D, Deshmuk L and Savarka S: A review on gums and mucilages used in pharmaceutical industry. International journal of creative research thoughts 2023; 11(10):2320-2882
- 59. Kwakye KO, Adom ENN, and Kipo SL: Preparation and in vitro characteristics of tablet cores coated with albizia, albizia/khaya and albizia/hpmc films. International Journal of Applied Pharmaceutics 2009; 1(1):22-29.
- Odeku OA and Fell JT: *In-vitro* evaluation of khaya and albizia gums as compression coatings for drug targeting to the colon. Journal of Pharmacy and Pharmacology | Oxford Academic 2005; 57(2): 163-168. doi: 10.1211/0022357055362.
- Ozoude C, Azubuike C, Ologunagba M, Tonuewa and Igwilo C: Formulation and development of metforminloaded microspheres using *Khaya senegalensis* (Meliaceae) gum as co-polymer. Future Journal of Pharmaceutical Sciences 2020; 6:120.
- Saadiya M, Oyi R and Ibrahim Y: Characterization of Khaya Senegalensis Gum: Effects of Drying Methods on the Physicochemical Properties. Journal of pharmaceutical research 2023; 22(3): 142-151. doi: 10.18579/jopcr/v22.3.23.6
- 63. Adenuga YA, Odeku OA, Adegboye TA and Iteola OA: Comparative Evaluation of the Binding Properties of Two Species of Khaya Gum Polymer in a Paracetamol Tablet Formulation. Pharmaceutical Development & Technology 2008; 13(6): 473-480. doi: 10.1080/10837450802179338
- 64. Rajeswari S, Patil R, Tatiya A and Dipashri D: A review on natural gums and mucilage used as suspending agents in various suspension. European journal of pharmaceutical and medical research 2021; 8(3).
- 65. Morkhade DM, Fulzele SV, Satturwar PM and Joshi SB: Gum copal and gum damar: Novel matrix forming materials for sustained drug delivery. Indian journal of pharmaceutical sciences 2006; 68(1):53- 58. doi: 10.4103/0250-474X.22964
- 66. Mundada A, Satturwar P, Fulzele S, Joshi S and Dorle A: Characterization and Evaluation of Novel Film Forming Polymer for Drug Delivery. Iranian Journal of Pharmaceutical Research 2011; 10(1): 35-42.

- 67. MorkhadeAtmaram R and Dipali P: A Review on Herbal Excipient. International Journal of Research Publication and Reviews 2022; 3(12): 1090-1093.
- Tall A, Diallo A, Erouel M, Seck M, Chouiref L, Saddi M, Wederni M, Ly E, Diallo A, Noureddine B, Kobor D and Khirouni K: Electrical and Dielectrical Properties of Khaya Gum Biopolymer Thin Filmcoated by Spray Pyrolysis Technique. Journal of Sol-Gel Science and Technology 2022; 104. doi:10.1007/s10971-022-05952-4
- Lankalapalli S and Sandhala D: A review on natural gums and their use as pharmaceutical excipients. International journal of pharmaceutical science & research 2019; 10(12): 5274-5283.
- Singh R and Barreca D: Analysis of gums and mucilages. In book: Recent Advances in Natural Products Analysis 2020; 663-676. doi:10.1016/B978-0-12-816455-6.00021-4.
- 71. Ramakrishnan R, Palanisamy S, Sumitha N, Padinjareveetil A, Sabarinath S, Waclawek S, Uyar T, Cernik M, Varma R, Cheong J Vellora V and Padil T: Regenerable and Ultraflexible Sustainable Film Derived from Tree Gum Kondagogu for High-Performance Electromagnetic Interference Shielding. ACS Sustainable Chemistry. Engineering 2023; 11(19): 7344–7356. https://doi.org/10.1021/acssuschemeng.2c07743
- Mahajan Y, Sharma D, Kumar P and Ashawat M: Natural Polymers as Excipient in Formulation of Novel Drug Delivery System. International journal in pharmaceutical sciences 2023;1(8);50-71.
- 73. Putta KS, kumar AA and Kumar AA: Formulation and invitro evaluation of mucoadhesive microcapsules of Glipizide with gum Kondagogu. Journal of Chemical and Pharmaceutical Research 2010; 2(5): 356-364.
- 74. Ramana G, Reddy KD and Sravanthi O: Design and Evaluation of Natural Gum Based Oral Controlled Release Matrix Tablets of Ambroxol Hydrochloride. Der Pharmacia Lettre 2012; 4(4): 1105-1114.
- 75. Fernandes N, Meira I, Alves V, Sampaio f and Oliveira A: Tara Gum as a controlled delivery system of fluoride in toothpaste: *in-vitro* enamel remineralization study. *Pesquisa Brasileira* em Odontopediatria e Clínica Integrada 2020; 1-7. (doi.org/10.1590/pboci.2021.003)
- 76. Mate J and Mishra S: Exploring the Potential of Moi Gum for Diverse application: A Review. Journal of Polymers and The Environment 2020; 28(5):1579-1591. doi:10.1007/s10924-020-01709-8
- 77. Mate j and Mishra S: Exploring the Potential of Moi Gum for Diverse Applications: A Review. Journal of Polymers and the Environment 2020; 28(5). doi:10.1007/s10924-020-01709-8
- MuraniV: A Review on: Extraction of Mimosine from Leucaena leucocephala tree & their Phytochemical study and Pharmacologycal Activity. International journal of scientific research 2023; 12(4): 21-23.
- Mittal N and Kaur G: Leucaena leucocephala (Lam.) galactomannan nanoparticles: Optimization and characterization for ocular delivery in glaucoma treatment. International journal of biological macromolecules 2019; 139: 1254-1262.
  https://doi.org/10.1016/j.jjiiamaa.2010.08.107

https://doi.org/10.1016/j.ijbiomac.2019.08.107

- 80. Verma PRP and Razdan B: Studies on disintegrant action of *Leucaena leucocephala* seed gum in ibuprofen tablet and its mechanism .Journal of Scientific and Industrial Research 2007; 66(7).
- 81. Zayed Z M, Sallam S and Shetta N: Review article on leucanealeucocephala as one of the miracle timber trees.

International Journal of Pharmacy and Pharmaceutical Sciences 2018; 10(1): 1-7.

- Lodhi D, Verma M and Golani P: Review of natural gum as a sustained release excipient in tablet form. World journal of pharmaceutical and medical research 2021; 7(10):85 – 91.
- Nayak BS, Nayak UK, Patro KB and Rout PK: Design and Evaluation of Controlled Release Bhara Gum Microcapsules of Famotidine for Oral Use. Research Journal of Pharmacy and Technology 2008; 1(4): 433-436.
- 84. Tiwari S and Talreja S: *Delonix regia*'s Potential Health Benefits: Investigating the Medicinal Wonder. International journal in pharmaceutical science 2023; 1(8):223-231. doi:10.5281/zenodo.8280328
- 85. Shirsath V and Sonawane S: Ethanomedicinal, phytochemical constituents and pharmacological activities of *Delonix regia*: a review. World Journal of Pharmaceutical Research 2022; 11(16): 296-311.
- Rodriguez W, Cerqueira M, Guerrero L, Pastrana L and Vega M: *Delonix regia* galactomannan-based edible films: Effect of molecular weight and k-carrageenan on physicochemical properties. Food hydrocollids 2020; 103. doi.org/10.1016/j.foodhyd.2019.105632t
- 87. Adetogun EG and Alebiowu G: Properties of *Delonix regia* seed gum as a novel tablet binder. Acta Poloniae Pharmaceutica-Drug Research 2009; 66(4):433-438.
- Barhate AN, Virkar AR, Beldar GB and Shinde S: Evaluation of *Delonix regia* Seed Gum as Suspending Agent in Paracetamol Suspensions. International journal of pharmtech research 2014; 6(2):530-536.
- Adetogun EG and Alebiowu G: Influence of *Delonix regia* seed gum on the compressional characteristics of paracetamol tablet formulation 2007; 17(6): 443-445. (DOI:10.1016/S1773-2247(07)50086-9
- Hedayati S, Ansarifar E, Tarahi M, Tahsiri Z, Baeghbali V and Niakousari m: Influence of Persian Gum and Almond Gum on the Physicochemical Properties of Wheat Starch. Gels 2023; 9(6): 460. https://doi.org/10.3390/gels9060460
- 91. Chaudhari SR and Dhuppad UR Extraction and Characterization of Okra and Almond Gum as a Pharmaceutical Aid. International Journal of Drug Delivery Technology 2023: 13(04):1503-1508. Doi: 10.25258/ijddt.13.4.58
- 92. Gayathri M, Chandrasekaran M, Radhakrishnan, Kuppusamy G and Singn S: Optimizing Badam gun towards tableting excipients. Research Journal of Pharmacy and Technology. 2020;13(12): 6176-6181. doi: 10.5958/0974-360X.2020.01077.X
- 93. Sunayana S, Gowda DV, Gupta N B, Sivadasu P and Manjunath M: Formulation development and evaluation of almond gum based sustained release. Asian Journal of Pharmaceutical and Clinical Research2018; 11(12):166-169.
- Sahu P, Patel S, Pisalkar PS, Katiyar P and Khokhar D: Nutritional, physical and thermal properties of Cassia tora (Charota) seeds. The Pharma Innovation Journal 2023; 12(9): 2291-2299.
- 95. Zibaee E, Javedi B, Sobhani Z, Akaberi M, Farhadi F, Amiri M, Baharara H, Sahebkar A and Emami S: Cassia species: A review of traditional uses, phytochemistry and pharmacology. Pharmacological Research-Modern Chinese medicine 2023; 9: 65. doi.org/10.1016/j.prmcm.2023.100325
- 96. Singh S, Bothara S, Singh S, Patel DR, and Mahobia NK: Pharmaceutical Characterization of Cassia tora of Seed Mucilage in Tablet Formulations. Der Pharmacia Lettre 2010; 2(5): 54-61.

- 97. Chouhan S, Darwhekar G and Gupta A: Formulation and evaluation of orodispersible tablet of rosuvastatin using cassia tora seeds mucilage as natural superdisintegrant. International Journal of Pharmaceutical Sciences and Medicine 2020; 5(7): 16-38.
- 98. Akkaya NE, Ergun C, Saygun A, Yesilcubuk N, Akel-Sadoglu N, Kavakli HI, Turkmen HS and Giz CH: New biocompatible antibacterial wound dressing candidates; agar-locust bean gum and agar-salepfilms. International journal of biological macromolecules 2020; 155: 430-438. doi: 10.1016/j.ijbiomac.2020.03.214
- 99. Malik K, Arora G and Singh I: Locust bean gum as super disintegrant - Formulation and evaluation of nimesulideorodispersible tablets. Polymers in Medicine 2011; 41(1):17
- 100. Singh H, Majumdar A, Malviya N and Saxen S: Design, Development and Characterization of fast disintegrating tablet of amlodipine besylate by using superdisintegrantsplantago ovata and locust bean gum. International journal of pharmaceutical science and research 2020;11(12): 6166-6172.
- 101. Moin A and Shivakumar H: Formulation and in vitro evaluation of sustained-release tablet of diltiazem: Influence of hydrophilic gums blends. Journal of pharmaceutical research 2010; 9(3): 283-291.
- 102. Sandolo C, Coviello T, Matricardi P and Alhaique F: Characterization of polysaccharide hydrogels for modified drug delivery. European Biophysics Journal 2007; 36(7): 693-700. doi: 10.1007/s00249-007-0158-y
- 103. Vijayaraghavan C, Vasanthakumar S and Ramakrishnan A: In vitro and in vivo evaluation of locust bean gum and chitosan combination as a carrier for buccal drug delivery. Archiv der Pharmazie 2008; 63(5): 342-347
- 104. Marianecci C, Carafa M, Di Marzio L, Rinaldi F, Di Meo C, Alhaique F, Matricardi P and Alhaique F: New vesicle-loaded hydrogel system suitable for topical applications: Preparation and characterization. The Journal of Pharmacy & Pharmaceutical Sciences 2011; 14(3): 336-346 doi: 10.18433/j3160b
- 105. Braz L, Grenha A, Corvo M, Lourenco J, Ferreira D, Sarmento B and Costa AM: Synthesis and characterization of Locust Bean Gum derivatives and their application in the production of nanoparticles. Carbohydrate Polymer 2018; 1(181): 974-985 doi 10.1016/j.carbpol.2017.11.052
- 106. Martin A, Chinarro E, Cazorla R, Notario P F, Veiga MD, Rubio J and Tamayo A: Optimized hydration dynamics in mucoadhesive xanthan- based trilayer vaginal films for the controlled release of tenofovir. Carbohydrate Polymer 2022; 278: 1-14. doi: 10.1016/j.carbpol.2021.118958
- 107. Pandey S, Son N and Kang M: Synergistic sorption performance of karaya gum crosslink poly(acrylamide-coacrylonitile) @ metal nanoparticle for organic pollutants, International Journal of biological macromolecules 2022; 210: 300-314.
- 108. Munot N, Kandekar U, Rikame C, Patil A, Sengupta P, Urooj S and Bilal A: Improved Mucoadhesion, permeation and In Vitro Anticancer Potential of synthesized Thiolated Acacia and Karaya Gum Combination: A Systemic Study. Molecules 2022; 27(20) 1-15. doi: 10.3390/molecules27206829
- 109. Bin MN, Nadeem MS, Gilani SJ, Imam SS, Alshehri S and Kazmi I: Novel karaya gum micro-particles loaded *Ganoderma lucidum* polysachharide regulate sex hormone, oxidative stress and inflammatory cytokine levels in cadmium induced testicular toxicity in experimental animals. International journal of biological

macromolecules. 2022; 138: 338-346. doi: 10.1016/j.ijbiomac.2021.11.072

- 110. Rahamathulla M, Saisivam S, Alshetaili A, Hani U, Gangadharappa HV, Alshehri S, and Ghoneim MM: Design and Evaluation Of losartan Potassium Effervescent Floating Matrix Tablets: *in-vivo* X-ray Imaging and Pharmacokynetic studies in Albino Rabbits. Polymers (basel) 2021; 2-16: 13(20). doi.org/10.3390/polym13203476
- 111. Hasnain S, Nayak A, Ansari M and Pal D: Pharmaceutical Applications of Locust Bean Gum. Natural Polymers for Pharmaceutical Applications 2019; 139-162. doi:10.1201/9780429328251-6.
- 112. Sharma N, Deshpande R, Sharma D and Sharma R: Development of locust bean gum and xanthan gum based biodegradable microparticles of celecoxib using a central composite design and its evaluation. Industrial Crops and Products 2016; 82: 161-170. doi:10.1016/j.indcrop.2015.11.046
- 113. Amaldos MJN, Najar IA, Kumar J and Sharma A: Therapeutic efficacy of rifaximin loaded tamarind gum polysaccharide nanoparticles in TNBS induced IBD model wistarrats. Reports on Practical Oncology and Radiotherapy. 2021; 26(5):712-729. doi: 10.5603/RPOR.a2021.0100
- 114. Raj S, Fuloria S, Subramaniyan V, Sathasivam K, Kumari U, Unnikrishnan Meenakshi D, Porwal O, Hari Kumar D, Singh A, Chakravarthi S and Kumar FN: Evaluation of antitumor efficacy of Chitosan-Tamarind Gum Polysaccharide Polyelectrolyte Complex Stabilized Nanoparticles of Simvastatin. International Journal of Nanomedicine 2021; 16: 2533-2553. doi: 10.2147/IJN.S300991
- 115. Bagal-Kestwal DR and Chiang BH: Tamarindus indica seed-shell nanoparticles-silver nanoparticles-Ceratonia silique bean gum composite for copper-micro mesh grid electrode fabrication and its application for glucose detection in artificial salivary samples. International Journal of Biological Macromolecules 2021; 189:993-1007. doi: 10.1016/j.ijbiomac.2021.08.148
- 116. Balakrishnan B, Sarojini B, Kodoth A, Dayananda S and Venkatesha R: Fabrication and characterization of tamarind seed gum based novel hydrogel for the targeted delivery of omeprazole magnesium. International journal of biological macromolecules 2024; 258(1). doi.org/10.1016/j.ijbiomac.2023.128758
- 117. Sittikijyothin W, Phonyotin BSangnim T, and Huanbutta K, Using carboxymethyl gum from Tamarindus indica and Cassia fistula seeds with Chromolaena odorata leaf extract to develop antibacterial gauze dressing with hemostatic activity. Research in Pharmaceutical Science. 2021; 16(2):118-128. doi: 10.4103/1735-5362.310519
- 118. Chun T, Mac Calman T, Dinu V, Phillips-Jones MK and Harding SE: Hydrodynamic Compatibility of Hyaluronic Acid and Tamarind Seed Polysaccharide as Ocular Mucin Supplements. Polymers (Basel) 2020;12(10): 2-11.
- 119. Rawooth M, Qureshi D, Hoque M, Prasad MPJG, Mohanty B, Alam MA, Anis A, Sarkar P and Pal K: Synthesis and characterization of novel tamarind gum and rice bran oil-based emulgels for the ocular delivery of antibiotics. International Journal of Biological Macromolecules 2020; 1(164): 1608-1620.
- 120. Oliveira WQ, Wurlitzer NJ, Araújo AWO, Comunian TA, Bastos MDSR, Oliveira AL, Magalhaes HCR, Ribeiro HL, Figueiredo RW and Sousa PHM: Complex coacervates of cashew gum and gelatin as carriers of green coffee oil: The effect of microcapsules application on the rheological and

sensorial quality of a fruit juice. Food Research International 2020; 131: 109047. doi: 10.1016/j.foodres.2020.109047

- 121. Kiniwa R, Miyake M, Kimura SI, Itai S, Kondo H and Iwao Y: Development of muco-adhesive orally disintegrating tablets containing tamarind gum-coated tea powders for oral care. International Journal of Pharmaceutics: X 2019; 1-10. doi: 10.1016/j.ijpx.2019.100012
- 122. Sinha R, Singh R, Chandrakar M and Parihar A: Designing of Curcumin nanoemulsion by using Tamarind gum as natural emulsifying agent. Research journal of pharmacy and technology 2024; 17(1): 172-178. doi: 10.52711/0974-360X.2024.00028
- 123. Huanbutta K, Yunsir A, Sriamornsak P and Sangnim T: Development and in vitro/in vivo evaluation of tamarind seed gum-based oral disintegrating tablets after fabrication by freeze drying. Journal of drug delivery science and technology 2019; 54. doi.org/10.1016/j.jddst.2019.101298
- 124. Ranote S, Musioł M, Kowalczuk M, Joshi V, Chauhan GS, Kumar R, Chauhan S and Kumar K: Functionalized Moringa oleifera Gum as pH-Responsive Nanogel for Doxorubicin Delivery: Synthesis, Kinetic Modelling and In Vitro Cytotoxicity Study. Polymers (Basel). 2022; 14(21): 2-19. doi.org/10.3390/polym14214697
- 125. George TT, Oyenihi AB, Rautenbach F and Obilana AO: Characterization of Moringa oleifera Leaf Powder Extract Encapsulated in Maltodextrin and/or Gum Arabic Coatings Foods. 2020; 10(12). 2-19. doi.org/10.3390/foods10123044
- 126. Castro C, Espinoza C, Ramos R, Boone-Villa VD, Aguilar MA, Martínez GCG, Aguilar CN and Ventura JM: Spraydrying encapsulation of microwave-assisted extracted polyphenols from Moringa oleifera: Influence of tragacanth, locust bean, and carboxymethyl-cellulose formulations. Food Research International. 2022; 144: 1-14. doi: 10.1016/j.foodres.2021.110291
- 127. Irfan M, Munir H and Ismail H: Moringa oleifera gumbased silver and zinc oxide nanoparticles: green synthesis, characterization and their antibacterial potential against MRSA. Biomaterial Research 2021; 25(17): 2-8. doi.org/10.1186/s40824-021-00219-5
- 128. Ahmad S, Manzoor K, Purwar R and Ikram S: Morphological and Swelling Potential Evaluation of Moringa oleifera Gum/Poly(vinyl alcohol) Hydrogels as a Superabsorbent. ACS Omega 2020; 5(29): 17955-17961. doi: 10.1021/acsomega.0c01023
- 129. Rimpy, Abhishek, Ahuja M: Evaluation of carboxymethyl moringa gum as nanometric carrier. Carbohydrate Polymer 2017; 174: 896-903. doi: 10.1016/j.carbpol.2017.07.022
- 130. Kotadiyal R, Savant N P and Upadhyay U: Colon targeted moringa gum compression coated tablets of capecitabin: a factorial approach. Pharmacophore 2019; 10(1): 21-29.
- 131. Badwaik H, Hoque A, Kumari L, Sakure K, Baghel M and Giri T: Moringa gum and its modified form as a potential green polymer used in biomedical field. Carbohydrate Polymers 2020; 249
- 132. Hodaei M and Varshosaz J: Cationic Okra gum coated nanoliposomes as a pH-sensitive carrier for co-delivery of hesperetin and oxaliplatin in colorectal cancers. Pharmaceutical Development & Technology 2022; 27(7): 773-784. doi: 10.1080/10837450.2022.2119249)
- 133. Upadhyay P, Agarwal S and Upadhyay S: Hydrophobically Modified Abelmoschus esculentus Polysaccharide Based Nanoparticles and Applications: A Review. Current Drug Discovery Technologies 2022; 19(6). doi: 10.2174/1570163819666220801121857

- 134. Brar V and Kaur G: Thiolated okra chitosan nanoparticles: preparation and optimisation as intranasal drug delivery agents. Journal of Microencapsulation 2020; 37(8): 624-639. doi: 10.1080/02652048.2020.1836057
- 135. Naderi M, Sabouri Z, Jalili A, Zarrinfar H, Samarghandian S, and Darroudi M: Green synthesis of copper oxide nanoparticles using okra (Abelmoschus esculentus) fruit extract and assessment of their cytotoxicity and photocatalytic applications. Environmental technology & innovation 2023; 32: 1-14. doi.org/10.1016/j.eti.2023.103300
- 136. Kurra P, Narra K, Orfali R, Puttugunta SB, Khan SA, Meenakshi DU, Francis AP, Asdaq SMB and Imran M: Studies on Jackfruit-Okra Mucilage-Based Curcumin Mucoadhesive Tablet for Colon Targeted Delivery. Frontiers in pharmacology 2022; 13: 1-13. doi: 10.3389/fphar.2022.902207
- 137. Chatterjee S and Majumdar R: Novel approach of extraction and characterization of okra gum as a binder for tablet formulation. Asian Journal of Pharmaceutical and Clinical Research 2019;12(1):189 doi:10.22159/ajpcr.2018.v12i1.29053
- 138. Rajalakshmi M and Sangeetha S: Okra mucilage method of extraction and a novel strategy for pharmaceutical drug delivery system. Journal of Pharmaceutical Negative Results 2023;14(3): 542-550)
- 139. Ajala TO and Silva BO: The design of ibuprofen-loaded microbeads using polymers obtained from Xanthosoma sagittifolium and Dillenia indica. Polymer Medicine 2020; 50(1): 21-31. doi: 10.17219/pim/122015
- 140. Huang Q, Luo A, Jiang L, Zhou Y, Liu Qand Zhang C: Disinfection efficacy of green synthesized gold nanoparticles for medical disinfection applications. African Health Science 2019; 19(1):1441-1448.
- 141. Nayaka S, Bhat M, Udayshankar A C, Lakshmeesha T R, Geetha N and JogaiahS: Biosynthesis and characterization of Dillenia indica -mediated silver nanoparticles and their biological activity. Applied Organometallic Chemistry 2020; 34(4): 1-9. doi:10.1002/aoc.5567
- 142. S Hasnain, Guru P, Rishishwar P, Ali S, Tahir Ansari M and Nayak A: Atenolol releasing buccal patches made of Dillenia indica L. fruit gum: preparation and ex vivo evaluations. Discover Applied Sciences 2020; 2(57). https://doi.org/10.1007/s42452-019-1756-x
- 143. Hussain HR, Bashir S, Mahmood A, Sarfraz RM, Kanwal M, Ahmad N, Shah HS, and Nazir I: Fenugreek seed mucilage grafted poly methacrylate pH-responsive hydrogel: A promising tool to enhance the oral bioavailability of methotrexate. International journal of biological macromolecules 2022; 202:332-344.doi: 10.1016/j.ijbiomac.2022.01.064
- 144. Adhikari S and Panda S: Development and Evaluation of Atenolol Containing Mucoadhesive Buccal Patches. International journal of pharmaceutical review and research 2019; 57(2): 29-36.
- 145. Momin MM, Kane S and Abhang P: Formulation and evaluation of bilayer tablet for bimodal release of venlafaxine hydrochloride. Frontiers Pharmacology 2015; 6:144.
- 146. Pathak D, Kumar P, Kuppusamy G, Gupta A, Kamble B and Wadhwani A: Physicochemical characterization and toxicological evaluation of plant-based anionic polymers and their nanoparticulated system for ocular delivery. Nanotoxicology. 2014; 8(8):843-855.
- 147. Kumari S, Panda S, Durgarao S, Chinnababu K and Rao K: Formulation and Evaluation Fast Dissolving Tablet of Hibiscus rosa-sinensis Leaf Mucilage as Superdisintegrant.

International Journal of Pharma Research and Health Sciences 2019; 7 (5): 3056-3061

- 148. Bakr RO, Amer RI, Attia D, Abdelhafez MM, Al-Mokaddem AK, El-Gendy AEG, El-Fishawy AM, Fayed MAA and Gad SS: In-vivo wound healing activity of a novel composite sponge loaded with mucilage and lipoidal matter of Hibiscus species. Biomedicine and Pharmacotherapy 2021; 135: 1-14.doi.org/10.1016/j.biopha.2021.111225
- 149. Kurra P, Narra K, Puttugunta SB, Kilaru NB and Mandava BR: Development and optimization of sustained release mucoadhesive composite beads for colon targeting. International journal of biological macromolecules 2019; 139:320-331. doi: 10.1016/j.ijbiomac.2019.07.190
- 150. Ahad H, Chinthaginjala H, Karar A, Seed M and Alward A: Effective management of rare lymphangioleiomyomatosisusing sirolimus: tablet matrix with hibiscus rosa sinensis leave mucilage.Research journal of pharmaceutical dosage form and technology 2021; 13(4). doi: 10.52711/0975-4377.2021.00045
- 151. Frasat T, Tulain UR, Erum A, Saleem U, Sohail MF and Kausar R: Aloe vera and artemisia vulgaris hydrogels: exploring the toxic effects of structural transformation of the biocompatible materials. Drug Development and Industrial Pharmacy 2021; 47(11):1753-1763. doi: 10.1080/03639045.2022.2050751
- 152. Bhar B, Chakraborty B, Nandi SK and Mandal B: Silkbased phyto-hydrogel formulation expedites key events of wound healing in full-thickness skin defect model. International journal of biological macromolecules 2022; 1(203):623-637. doi: 10.1016/j.ijbiomac.2022.01.142
- 153. Otalora MC, Wilches-Torres A and Gomez Castano JA: Spray-Drying Microencapsulation of Pink Guava (Psidium guajava) Carotenoids Using Mucilage from Opuntia ficusindica Cladodes and Aloe Vera Leaves as Encapsulating Materials. Polymers (Basel). 2022; 14(2): 1-16. doi.org/10.3390/polym14020310
- 154. Jadav A, Shewale A and Bhutkar M: Evaluation of aloe vera and hibiscus rosa sinensis mucilage as a binder in different tablet formulation. Asian journal of pharmacy and technology 2020; 10(1).
- 155. Aguirre-Joya JA, Pastrana-Castro L, Nieto-Oropeza D, Ventura-Sobrevilla J, Rojas-Molina R and Aguilar CN: The physicochemical, antifungal and antioxidant properties of a mixed polyphenol based bioactive film. Heliyon 2018; 4(12):1-14. doi: 10.1016/j.heliyon.2018.e00942
- 156. Bonsu MA, Ofori-Kwakye K, Kipo SL, Boakye-Gyasi ME and Fosu MA: Development of Oral Dissolvable Films of Diclofenac Sodium for Osteoarthritis Using Albizia and Khaya Gums as Hydrophilic Film Formers. Journal of Drug Delivery 2016; 1-11. doi: 10.1155/2016/6459280
- 157. Odeku OA, Okunlola A and Lamprecht: A Microbead design for sustained drug release using four natural gums. International Journal of Biological Macromolecules 2013; 58:113-120. doi: 10.1016/j.ijbiomac.2013.03.049
- 158. Prabhu P, Ahamed N, Matapady HN, Ahmed MG, Narayanacharyulu R, Satyanarayana D and Subrahmanayam E: Investigation and comparison of colon specificity of novel polymer khaya gum with guar gum. Pakistan Journal of Pharmaceutical Science 2010; 23(3): 259-265.
- 159. Odeku OA and Fell JT: Effects of the method of preparation on the compression, mechanical, and release properties of khaya gum matrices. Pharmaceutical development & technology 2006; 11(4): 435-441.doi: 10.1080/10837450600770544

- 160. Wadher KJ, Kakde RB and Umekar MJ: Formulation and evaluation of a sustained-release tablets of metformin hydrochloride using hydrophilic synthetic and hydrophobic natural polymers. Indian Journal of pharmaceutical sciences 2011; 73(2): 208-215. doi: 10.4103/0250-474x.91579
- 161. Rani R, Narsiman B, Varma RS and Kumar R: Gumbased nanocapsules comprising naphthoquinones enhance the apoptotic and trypanocidal activity against Trypanosoma evansi. European journal of pharmaceutical science 2022; 171: 2-12. (doi.org/10.1016/j.ejps.2022.106118)
- 162. Fulbandhe VM, Jobanputra CR, Wadher KJ, Umekar MJ and Bhoyar GS: Evaluation of release retarding property of gum damar and gum copal in combination with hydroxypropyl methylcellulose. Indian Journal of Pharmaceutical Science 2012; 74(3): 189-194. doi: 10.4103/0250-474X.106059
- 163. Kr V, Yalavarthi PR, Vadlamudi HC, Kalluri JKY and Rasheed A: Process, Physicochemical Characterization and In-Vitro Assessment of Albendazole Microcrystals. Advance Pharmaceutical Bulletin 2017; 7(3): 419-425.
- 164. Siraj S, Athar S, Khan G, Raza S and Ansari M: Review on solid dispersion of poor water-soluble drug by using natural polymers. The Pharma Innovation Journal 2019; 8(1): 631-636
- 165. Yalavarthi PR, Vulava J, Vadlamudi HC, Balambhaigari RY and Nair R: Modified pulsincap of ibuprofen--a novel approach for chronotherapy. Current Drug Delivery. 2013; 10(3): 299-308. doi: 10.2174/1567201811310030006
- 166. Barbosa BST and Garcia-Rojas EE: Double emulsions as delivery systems for iron: Stability kinetics and improved bioaccessibility in infants and adults. Current Research in Food Science 2022; 5:718-725.
- 167. Santos MB, de Carvalho CWP and Garcia-Rojas EE: Microencapsulation of vitamin D3 by complex coacervation using carboxymethyl tara gum (*Caesalpinia spinosa*) and gelatin A. Food Chemistry 2021; 343: 2-10. doi: 10.1016/j.foodchem
- 168. Ma Q, Hu D and Wang L: Preparation and physical properties of tara gum film reinforced with cellulose nanocrystals. International Journal of Biological Macromolecules 2016; 86: 606-612. doi: 10.1016/j.ijbiomac.2016.01.104

- 169. Jeevanandham S, Sekar M, Dhachinamoorthi D, Muthukumaran M, Sriram N, and Joysaruby J: Sustainrelease of various drugs from *leucaena leucocephala* polysaccharide. Journal of Young Pharmacist. 2020 2(1):15-20. doi: 10.4103/0975-1483.62207
- 170. Devkar TB, Tekade AR and Khandelwal KR: Surface engineered nanostructured lipid carriers for efficient nose to brain delivery of ondansetron HCl using *Delonix regia* gum as a natural mucoadhesive polymer. Colloids Surf B Biointerfaces. 2014; 122:143-150. doi: 10.1016/j.colsurfb.2014.06.037
- 171. Krishnaraj K, Chandrasekar MJ, Nanjan MJ, Muralidharan S and Manikandan D: Development of sustained release antipsychotic tablets using novel polysaccharide isolated from *Delonix regia* seeds and its pharmacokinetic studies. Saudi Pharmaceutical Journal 2012; 20(3): 239-48. doi: 10.1016/j.jsps.2011.12.003
- 172. Dabhi C, Randale S, Belgamwar V, Gattani S and Tekade A: Predictable pulsatile release of tramadol hydrochloride for chronotherapeutics of arthritis. Drug Delivery 2012; 17(5): 273-81. doi: 10.3109/10717541003706240
- 173. Venkatesan R, Sekar S, Raorane CJ and Kim SC: Hydrophilic Composites of Chitosan with Almond Gum: Characterization and Mechanical, and Antimicrobial Activity for Compostable Food Packaging. Antibiotics (Basel).2022; 11(11) doi.org/10.3390/antibiotics11111502
- 174. Kanteti RV, Sarheed O, Yadav H, Islam Q and Boateng J: Studies on Almond Gum and Gelucire-Based Pellets Prepared by Extrusion and Spheronization for Sustained Release. Turkish Journal of Pharmaceutical Sciences 2022; 19(5):521-529. doi: 10.4274/tjps.galenos.2021.05252
- 175. Nithiyavathi R, John Sundaram S, Theophil Anand G, Raj Kumar D, Dhayal Raj A, Al Farraj DA, Aljowaie RM, AbdelGawwad MR, Samson Y and Kaviyarasu K: Gum mediated synthesis and characterization of CuO nanoparticles towards infectious disease-causing antimicrobial resistance microbial pathogens. Journal of Infection and Public Health 2021; 14(12):1893-1902.doi: 10.1016/j.jiph.2021.10.022
- 176. Rodrigues FJ, Cedran MF, Pereira GA, Bicas JL and Sato HH: Effective encapsulation of reuterin-producing *Limosilactobacillus reuteri* in alginate beads prepared with different mucilages/gums. Biotechnology reports 2022; 1-11. doi: 10.1016/j.btre. 2022.e00737

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