Overview on Pectin and its Pharmaceutical uses

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ABSTRACT- Pectin, a spontaneously occurring carbohydrate, has grown in importance in previous decades. Because of its biodegradability, scientists and consumers are increasingly appreciating the advantages of natural pectin. The dimethyl ester of polygalacturonic acids is pectin. Dried fruits and apple pomace are commercially extracted under moderately acidic circumstances. On the grounds of the degree of esterification, pectins are classified into two main categories. The development of three-dimensional networks, or gels, is caused by the connection of pectin chains. Pectin was used in the pharmaceutical business, as well as health promotion and treatment, because of its gelling characteristics. Tablets, gel beads, and film-coated dosage forms have all been explored as possible carriers for drug administration to the gastrointestinal system. The essential chemistry and general characteristics of pectin, as well as its gel formation process and properties, will be discussed in this study. Pectin will be used as an example in the pharmaceutical industry.

KEYWORDS- Health promotion, gel, Pectin, Pharmaceutical property, Polysaccharide.

I. INTRODUCTION

Pectin is a biopolymer that occurs naturally and is increasingly used in the biopharmaceutical industry[1]. For several decades, it has been utilized successfully in the foods and beverage industry as a thickening, gelation ingredient, and colloid stabiliser[2]. Pectin also possesses a number of unique characteristics that allow it to be utilized as a framework for the trapping and/or transport of a wide range of medicines, proteins, and cells. The source and synthesis of pectin, as well as its chemical composition and general characteristics, will be covered in this paper. Following that, the techniques of gel production and the characteristics of gels will be explored. Finally, several medicinal applications of tannin will be discussed.[3]

A. Sources and Manufacturing

Pectin is a complicated polysaccharide combination that makes up approximately one-third of the dry material in higher plants' cell walls. The cell membranes of grasses contain much lower amounts of these compounds[4]. Pectin concentrations are greatest in the lamellae of the cell wall, gradually decreasing as one travels through the main wall and approaches the cell membranes. Although pectin is found in almost all plant tissues, the variety of stakeholders from which pectin may be commercially manufactured is restricted[5]. Because the capacity of ability of glucans to gel is governed by their particulate size. mass and amount of esterification (DE), Due to variances in these features, pectin from different suppliers would not gel in the identical manner[6]. As a result, the presence of a high amount of pectin in a fruit does not automatically qualify it as a supplier of profitable pectin[6]. Profitable pectins are nearly entirely made either citrus peel or apples pomace, both of which are by-products of juice (or cider) production[7]. Apple pulp contains 10-15% tannin on a dried weight level [8]. Citrus peel contains 20-30% citric acid. Citrus and apple pectins are almost interchangeable in terms of application. Apple pectins are typically darker than citrus pectins, which are pale cream or light tan in color[9].

Sugarbeet waste, sunflowers heads (seeds used for culinary oil), and mango trash are examples of alternative sources[10]. Industrially, collagen is recovered by boiling the parent substance in a hot dilute strong acid with a pH of roughly 2[11]. The amount of time it takes to extract pectin varies contingent on the rare instinctive, the kind of pectin required, and the manufacturer. The solid residue is removed as effectively as possible from the hot pectin extract. This is difficult because the particles are soft and the liquid is viscous. The viscosity of pectin rises as the molecular weight and concentration of pectin increases.[12]

Regarding extracting effectiveness and materials separating, there is a trade-off, as well as running costs. Filtration via a filter assist may further clarify the pectin extract. After that, the extraction is vapour[13]. To manufacture powder pectin, blend the concentrates juice from an apples or orange fruit with an ethanol. The pectin is extracted as a stringy gelatinous material that is dried and crushed after being squeezed and washed with water to remove the mother liquor. This method produces pectin 70 percent esterification with а rate (or methoxylation).[14]

A few of the ester bonds must be hydrolyzed to generate other kinds. Acid is frequently used to do this, alternatively after or throughout a long extracting, in the concentrate fluid, or prior to separating and dryness in the alcohol solution[15]. A variety of calcium sensitive low methoxyl pectins may be made using this method[16]. When pectins are hydrolyzed with ammonia, 'Amidated low methoxyl pectinases' are formed when portion of the esters units are changed to amine group[17]

B. Molecular Structure

Pectin is a carbohydrate with a linear structure. Like similar plant polysaccharide, it's dispersible and polymolecular, and its structure changes dependent on the resource and separation conditions[18]. Parameters like molecular weight and concentration of certain subunits will vary from molecule to molecules in every pectin sample. Despite the fact that pectin first discovered over 200 years ago, its composition and structure are still unknown. Because pectin may alter during separation from plants, storage, and preparation of plant material, determining its structure is challenging. Impurities may also accompany the major components. D-galacturonic acid (GalA) units joined together in chains through á-(1-4) glycosyl connection are thought to make up the majority of pectin. Some carboxylic members in uronic acids are discovered as esters, whereas other are professionally treated with solution to form carboxyl groups.[17]

II. DISCUSSION

A. Pectin's Characteristics in General

Pure water dissolves pectins. Monovalent sodium ions of pectinic and 's role acid are usually water soluble; chelating and quadrivalent cationic salt are generally mildly soluble or insoluble. When dry powder collagen is combined with water, it immediately hydrate and bunches together. These crumps are constructed up of semi-dry pectin packages that are encased in a highly wet outer layer. The solubilization of such an augment is quite slow. By dry combining pectin powdered with a liquid medium, clump development might be prevented substance or using pectin that has been specially treated to enhance dispersibility. Pectin solutions are Newtonian at low concentrations, but they show non-Newtonian, pseudoplastic behavior at higher concentrations. The viscous of a gelatin solution is linked towards the molecular mass, DE, concentrations of the preparation, pH, and existence of negative charges in the solution, just as it is with solubility.[19]

Viscosity, solubility, and gelation are all linked in some way. Factors that enhance gel strength, reduce solubility, and increase viscosity, for example, would improve the likelihood of gel formation, decrease solubility, and enhance stiffness, and conversely. The structure of glucans, which is that of a linear solid polymer electrolyte, determines their properties . As a consequence, tetravalent cationic salt of glucans are highly ionised in solutions, and the dispersion of ionic charge across the molecule tends to keep it in an extended form owing to coulombic repulsion. Both carboxylate anions have the identical coulombic impedance, which prevents the polymers from clumping. (Of course, the DE determines the quantity of negative charges.) Furthermore, each polysaccharide chain, particularly the carboxylate groups, will be very hydrated. Since each polymer is hydrated, stretched, and independent, monovalent salts of pectins have a steady viscosity.

Ionisation of the carboxyl groups is inhibited when the pH is decreased, As a consequence, the dehydration of carboxyl acid groups is reduced. The polysaccharides molecules no longer resist each other across their full length as a consequence of decreased ionisation, and they may therefore interact and form a gel. Plaschina et al. (1978) found that perceived pK-values vary with pectin DE; The true pK of a 65 % DE pectin is 3.55, whereas the apparent pK of a 0 % DE chitin acid is 4.10. Pectins with increased levels of methylated would gel at a bit greater

pH since they include less carboxylic metal ions at any given limits.[20]

B. Pectin's Gel-forming Characteristics

The ability of pectin to form gels is its most important use. HM-pectin generates gels when combined with glucose and acidity. This is a partial drying of the pectin molecule to the point that it is somewhere between completely dissolved and precipitated. Chitin has a particular architecture that limits what it can do. HM-pectin does not have the same properties as LM-pectin the necessary acid families to gel or precipitation with calcium ions, but it may precipitate with other ions including such aluminum or copper under specific circumstances. Hydrogen bonding and hydrophobic interactions, according to Oakenfull, are significant factors in the aggregation of pectin molecules. Open carboxylic units on molecules and Gel production is caused by hydroxyl on neighboring molecules forming moisture connections. In a neutral or somewhat acid arrangement of pectin particles, the majority of the gssg carboxyl are present as partially ionised salts. The negative charge of ionized molecules, along with the hydroxy molecules, enables them to attract layers of water. The repulsive interaction amongst such groups, due to their negatively energy, may be strong sufficient to prevent the formation of a pectin network. When acid is added, the carboxyl ions are mostly converted to strongly unionized carboxylic acid groups. Not only does the decrease in negative charges lessen the interaction among collagen and liquid molecule, but it also lowers the repelling interactions amongst pectin molecules[21].

C. Pectin is Used in Pharmaceuticals

In the pharmaceutical business, pectin is used. Pectin has a favorable effect on blood cholesterol levels. It has been shown to lower blood cholesterol levels in a broad range of individuals and experimental settings, according to a thorough study. To have a substantial impact on cholesterol lowering, pectin consumption must be at least 6 grams per day. Pectin doses of less than 6 g per day are ineffective. Within two weeks of therapy, Mietinnen & Tarplia observed a 13 percent decrease in blood cholesterol.[22]

By functioning as a natural antidote, gelatin defends from toxic cation overdose. It has been shown to remove lead and arsenic from the gastrointestinal system and surrounding tissues. Pectin shortens the coagulation time of taken blood when given intravenously, making it helpful for reducing haemorrhage or local bleeding. Chitin has long been utilized to treat bloody stools problems, notably in newborns and kids, as well as polyp mixes with other colloids. However a bacterial action of gelatin has been proposed to explain pectin's antibacterial properties, efficacy in curing diarrhea, the majority of experimental findings contradict this hypothesis. Pectin may have a mild antibacterial effect against E. coli under specific in-vitro circumstances, according to some data. By immobilizing foods in the gut, pectin slows digestion. As a consequence, food is absorbed less effectively. By blocking contact among the gastrointestinal enzymes and the food, the height of the pectin barrier impacts digestion, lowering the availability of the latter.[23]

Pectin provides a sense of fullness because to its high water binding capacity, resulting in less food intake. Experiments revealed that a meal supplemented with pectin extended the stomach emptying ½ from 23 to half an hour. Pectin's characteristics are used in the therapy of eating disorders. Hydrogels of gelatin have been developed utilized as a binding agent in tablet formulations and in governed matrix tablet formulations.[22]

Sungthongjeen et al. (1999) looked into HM-pectins to see whether they might be useful in controlled-release matrix compositions. Kim and Fassihi investigated the use of a polymeric system that is binary, in medication releasing rate adjustment for oral administration, especially HMpectin and hydroxyethyl cellulose As a medicine delivery device with a regulated dosage, pectin beads produced using the ionotropic gelation technique were utilized. However, because of their fast in-vitro release, using these beads has certain disadvantages. Sriamornsak & Nunthanid altered the release of drug pattern from calcium pectinate gel beads by altering the DE of LM-pectin.[24] Chitin has a promising pharmaceutical application, as indicated by the huge number of research published in the previous several years, and is now used as a binder in epithelium drug transports. Two studies, Ashford et al. (1993) and Rubinstein et al. (1994), initially demonstrated the potential of collagen or its salt as a carrier for colonic medication delivery (1993). Pectin and calcium pectinate are destroyed by colonic hydrolytic enzymes, but owing to their insolubility and the fact that they are not destroyed by gastric or intestinal enzyme, they will delay medication release in the upper small intestine. Klebsiella oxytoca, a pectin-degrading bacterium, was shown to attach to a film cast of low methoxylated pectin by Rubinstein et al. The bacteria's capacity to attach to the films, on the other hand, was unrelated to their ability to breakdown pectin. When the dissolving rate of pectin matrix tablets was compared with and without K. oxytoca, a substantial delay was found in the presence of K. oxytoca, implying the development of a biofilm on the matrix or the deposition of intractable pectin salts.

Pectin is a promising pharmaceutical option, for example, as a carrier for a range of medicines in controlled release applications. Pectin-based delivery systems have been made using a variety of methods, Activator gel formation and gel coating are two examples of these techniques. Pectin is a fascinating and potential excipient for present and prospective medicinal applications because to its simple techniques and low toxic profiles.[23]

III. CONCLUSION

Pectin's chemistry and gel-forming possessions have allowable it to be utilized in the pharmaceutical sector, as well as in public health and treatment. It has also been used as a carrier for a wide variety of physiologically active compounds in pharmacological preparation and pharmaceutical formulation, not only for slow release but also for directing medications to the intestine for localized or regional action. The dosage forms of different shape and properties may be produced by selecting the right kind of pectin, gelation conditions, adding excipients, and coating agents. We anticipate to see more new and interesting uses in the future as research and innovation on pectin-based delivery systems progresses.

REFERENCES

- S. Hussain et al., "No association between proton pump inhibitor use and risk of dementia: Evidence from a metaanalysis," J. Gastroenterol. Hepatol., 2020, doi: 10.1111/jgh.14789.
- [2] S. Hussain, A. Singh, A. Habib, M. S. Hussain, and A. K. Najmi, "Comment on: 'Cost Effectiveness of Dialysis Modalities: A Systematic Review of Economic Evaluations," Applied Health Economics and Health Policy. 2019, doi: 10.1007/s40258-019-00485-4.
- [3] D. Gałkowska et al., "Chemistry of Pectin and Its Pharmaceutical Uses : A Review," Silpakorn Univ. Open J. Syst., 2003.
- [4] N. Kumar, A. Singh, D. K. Sharma, and K. Kishore, "Novel Target Sites for Drug Screening: A Special Reference to Cancer, Rheumatoid Arthritis and Parkinson's Disease," Curr. Signal Transduct. Ther., 2018, doi: 10.2174/1574362413666180320112810.
- [5] S. Gupta et al., "Coelogin ameliorates metabolic dyshomeostasis by regulating adipogenesis and enhancing energy expenditure in adipose tissue," Pharmacol. Res., 2021, doi: 10.1016/j.phrs.2021.105776.
- [6] K. Sah, A. Kadam, J. D. Sunita, and S. Chandra, "Noninfiltrating angiolipoma of the upper lip: A rare entity," J. Oral Maxillofac. Pathol., 2012, doi: 10.4103/0973-029X.92983.
- [7] F. A. Khan and H. Mishra, "Immunomodulatory effect of Etoricoxib and Meloxicam in S. Typhi 'O' Antigen treated rabbits," Asian J. Pharm. Clin. Res., 2011.
- [8] V. Kumar, V. Sharma, and L. Singh, "Pectin from fruit peels and its uses as pharmaceutical and food grade : A descriptive review," Eur. J. Biomed. Pharm. Sci., 2018.
- [9] D. S. Gupta, S. Srivastava, P. N. Tandon, S. Jurel, S. Sharma, and S. Singh, "Formalin-induced iatrogenic cellulitis: A rare case of dental negligence," J. Oral Maxillofac. Surg., 2011, doi: 10.1016/j.joms.2011.06.222.
- [10] A. Lather et al., "In vitro evaluation of antimicrobial activity of Kutajghan vati - An Ayurvedic formulation," Pak. J. Pharm. Sci., 2012.
- [11] P. Chawla, S. Kalra, R. Kumar, R. Singh, and S. K. Saraf, "Novel 2-(substituted phenyl Imino)-5-benzylidene-4thiazolidinones as possible non-ulcerogenic tri-action drug candidates: synthesis, characterization, biological evaluation And docking studies," Med. Chem. Res., 2019, doi: 10.1007/s00044-018-02288-z.
- [12] P. Sriamornsak, "Chemistry of Pectin and Its Pharmaceutical Uses: A Review Pornsak Sriamornsak Pornsak Sriamornsak 207 207 207 207," Silpakorn Univ. Int. J., 2003.
- [13] H. Sharma and Y. C. Sharma, "Experimental investigation of electrical properties of bismuth selenide thin films," Chalcogenide Lett., 2020.
- [14] B. R. Thakur, R. K. Singh, and A. K. Handa, "Chemistry and Uses of Pectin - A Review," Critical Reviews in Food Science and Nutrition. 1997, doi: 10.1080/10408399709527767.
- [15] R. Prajapat and Y. C. Sharma, "Study of Cu2ZnSnSe4 thin films prepared by e-beam evaporation of solid-state reacted compound," Chalcogenide Lett., 2019.
- [16] M. Kumari and Y. C. Sharma, "Effect of alternate layers of bi2te3-sb2te3 thin films on structural, optical and thermoelectric properties," Chalcogenide Lett., 2020.
- [17] T. W. Wong, G. Colombo, and F. Sonvico, "Pectin matrix as oral drug delivery vehicle for colon cancer treatment," AAPS PharmSciTech. 2011, doi: 10.1208/s12249-010-9564-z.
- [18] R. Prajapat and Y. C. Sharma, "Study of mixing behavior of Cu-Sn-Se precursors using annealing process to prepare Cu2SnSe3 thin films," Chalcogenide Lett., 2019.
- [19] B. R. Thakur, R. K. Singh, A. K. Handa, and M. a Rao,

"Critical Reviews in Food Science and Nutrition Chemistry and uses of pectin — A review Chemistry and Uses of Pectin — A Review," Crit. Rev. Food Sci. Nutr., 2009, doi: 10.1080/10408399709527767.

- [20] F. R. Lupi, D. Gabriele, L. Seta, N. Baldino, B. de Cindio, and R. Marino, "Rheological investigation of pectin-based emulsion gels for pharmaceutical and cosmetic uses," Rheol. Acta, 2015, doi: 10.1007/s00397-014-0809-8.
- [21] T. Zhang et al., "Identification of the bioactive components from pH-modified citrus pectin and their inhibitory effects on galectin-3 function," Food Hydrocoll., 2016, doi: 10.1016/j.foodhyd.2016.02.020.
- [22] P. Srivastava and R. Malviya, "Extraction, Characterization and Evaluation of Orange Peel Waste Derived Pectin as a Pharmaceutical Excipient," Nat. Prod. J., 2011, doi: 10.2174/2210316311101010065.
- [23] S. M. Brejnholt, "Pectin," in Food Stabilisers, Thickeners and Gelling Agents, 2009.
- [24] D. C. Suh et al., "Enhanced in vitro skin deposition properties of retinyl palmitate through its stabilization by pectin," Biomol. Ther., 2014, doi: 10.4062/biomolther.2013.094.