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Review

Carrageenan: A natural seaweed polysaccharide and its applications



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ABSTRACT

Polysaccharides have been gaining interesting aluab lication ne food and pharmaceutical fields. As they are derived from the natural so , they are e e, non-toxic, cheap, biodegradable and biocompatible. Carrageenan is or them, whic s the criteria of polysaccharide; it is a natural carbohydrate (polysacchari rom edible od seaweeds. The name Carrageenan is derived from the Chondrus crispus species of sea (Rhodophyceace) known as Carrageen Moss or on its applic gamed wide applicat Irish Moss, and Carraigin. A demahas been widely increasing in food and pharmaceutical sectors. Carrageena ns in experimental medicine, pharmaceutical industries. Through keen references of the reported literature on carformulations, cosmetics, and rageenan, in this review, we e described ut carrageenan, its properties, extraction and refining, and its food and pharmaceuti plications

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1. Introduction

Natural polymers and their application in the pharmaceutical industry are covered by the presence of synthetic polymers. Natural polymers are preferred above the synthetic as they are inert, safe, non-toxic, biocompatible, biodegradable, low cost, ecofriendly and abundantly available in nature (Guo, Skinner, Harcum, & Barnum, 1998). Probably one of the materials taking more benefit from these advantages is the polysaccharide family, which has been gaining interesting and valuable applications in the biomedical and specifically in biopharmaceutical field. Polysaccharides can be obtained from a number of sources including seaweeds, plants, bacteria, fungi, insects, crustacea, animals and even humans, and can be structurally tuned through genetic engineering (Colquhoun et al., 2001; Coviello, Matricardi, Marianecci, & Alhaique, 2007; D'Ayala, Malinconico, & Laurienzo, 2008; Kost & Goldbart, 1997; Laurienzo, 2010). The polysaccharide term gathers collectively quite diverse large carbohydrates that can be composed of only one kind of repeating monosaccharide (termed homopolysaccharides or homoglycans; e.g. starch, cellulose) or formed by two or more different monomeric units (heteropolysaccharides or heteroglycans; e.g. agar, alginate, carrageenan). The conformation of the polysaccharide chains is markedly dependent not only on the and ionic strength of the medium, particularly in the case of polyelectrolytes, but also on the temperature and the concentrat of certain molecules. Polysaccharides are divided into two su types: anionic and cationic polysaccharides. Seve nic and cationic polysaccharides are widely available i *c*ure have gained keen interest in food and pharmaceut deld.

Carrageenan, a naturally occurring nio polysaccharides extracted from certa ed (Kirk & ed s Othmer, 1992) of the Rhodophyceae ly (Chen, I Dustan, 2002; Rees, 1969; Rochas, Rinaudo . 1989; Sno , 1976; euchema, gigartina Stoloff, 1959), particularly from chandrus stellate, iridaea, hypnea, solig agardhiella carconema (Chen, n, 1973; Cheney, ston, & Brailey, cc2 s, West, & Guiry, 1982, 1983; McLachlan, Neish, & Shag McCa 1987; Chiovitti et al., 19 Mollion 1983; Murano, Toffanin, Mollet, Rahaoui, & Lem ; Pare Cecere, Rizzo, & Knutsen Garg, Mehta, & Mehta, 8; Parekh, Rao, & Chauhan, 1979; Parekh, 1), & C s to originate from the inha-1988). The ı "ca geenan f carraghes, on the south Irish coast where bitants of count extracts from and medicines were already used ars ago. The major constituent of such algae as early as 6 geenans, co-polysaccharides with the linear is the so-called backbone built up b b-galactose and 3,6-anhydro-α-p-galactose with variable density in the sulfated group (Campo, Kawano, da Silva, & Carvalho, 2009), Chondrus crispus, Gigartina stellata, Iridaea spp., Eucheuma spp. and Kappaphycus spp. (Fig. 1) are the chief raw materials used for carrageenan extraction.

Ideally, each disaccharide in the chain contains a β -deltagalactopyranose (G-unit) with either α -delta-galactopyranose (D-unit) or 3,6-anhydrogalactose (DA-unit). Other carbohydrate residues commonly exist in carrageenan, such as xylose, glucose, and uronic acids. The disaccharide units are variably sulfated, resulting in a sulfate content of 22–38% by weight in commercial carrageenan (Van de Velde & De Ruiter, 2002). Other cations, such as ammonium, calcium, magnesium, potassium, and sodium,

are also often present in the form of galactose esters (FAO, 2007; U.S. Pharmacopeia, 2010). Carrageenan is a good source of soluble fibre (Burtin, 2003). There are several different carrageenans with slightly varied chemical structures and properties (McHugh, 2003). The three most prevaler ghest commercial interest, polysaccharides of carra nan are kappa, and roperties (lambda carrageenan, serving differ 2007; U.S. Pharmacopeia, 2010; Van de Velde & L ter, 2002 S Food and Drug Administration have "G rally Re ized afe" (GRAS) list of products for consu on and topic cations of carrageenan. Carrageenan i extrer versatile gredient suitable for use in food and nonly tries (V de Velde, Lourenco, as no nutritional value, Pinheiro, & Bakke .002). igeena ely used in industry (Heertje, 1993; but have been Hood & Aller oeren, 197 aisgood, 1982; Van de Velde cently used in the pharmaceutical induset al., 2002) and mo. try as ex inient in pill tablets (Campo et al., 2009) and as a naterials of hygels (Hoffman, 2002). In pharmaceupot Applications (Takamatsu & Tosa, 1993; Van de Velde et al., tiq 20) and expe ental medicine, carrageenan is often used for th nflammatory agents (Zacharopoulos & Phillips, ting of an 199

The ad activity of carrageenan as a natural occurring insaccharide has been increasing widely for human applications as a strong position in the biomedical field. Due to their afferent chemical structure and physical properties this natural source can be used in the different applications, varying from tissue engineering to the preparation of drug vehicles for controlled release. This review article is prepared in order to focus on the present use and the diversified applications of carrageenan in pharmaceutical and food sectors.

2. Short summary on types, structure and properties of carrageenans

There are several different carrageenans with slightly varied chemical structures and properties (McHugh, 2003). Carrageenan formed by alternate units of p-galactose and 3,6-anhydro-galactose (3,6-AG) joined by α -1,3 and β -1,4-glycosidic linkage. According to the literature survey, carrageenan can be classified in 3 ways, based on the amount and position of sulphate groups, based on their family and based on its properties. Depending on the amount and position of the SO₃⁻ groups carrageenan are classified into $\lambda(lambda)$, $\kappa(kappa)$, $\iota(iota)$, $\upsilon(nu)$, $\mu(mu)$, $\theta(theta)$ and ξ(Ksi), all containing about 22–35% of sulphate groups (Stanley, 1987). Based on the family, Greer and Yaphe (1984) classified carrageenan mainly into three types as shown in the Table 1 (Anderson, Campbell, Harding, Rees, & Samuel, 1969; Glicksman, 1979; Hoefler, 2001; Hoffmann, Russell, & Gidley, 1996; Imeson, 2000; Moirano, 1977; Rees, 1969). The first family involves kappa (κ) family which contain a subclass like Kappa, iota, mu and nu carrageenan. The second class of family involves lambda carrageenan which, contain subclass like lambda, Xi and pi and the third class involves beta family containing subclass like beta and gamma carrageenan. Whereas Mollion, Moreau, and Christian (1986) introduced another family named omega family in which sulphate groups are on the C₆ of the 1,3-1, linked galactopyranosyl units. Alpha carrageenan was included as a subclass in beta family

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