

# SEBDENIA POLYDACTYLA—A NOVEL SOURCE OF CARRAGEENAN

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**Key Word Index**—*Sebdenia polydactyla*; Gigartinales; polysaccharide; carrageenan; phycocolloid.

**Abstract**—The polysaccharide extracted from *S. Polydactyla* with sodium hydrogen carbonate at 90° accounted for 32% of the dry weight of the alga. It was characterized as  $\lambda$ -carrageenan by IR and physico-chemical analysis.

## INTRODUCTION

Some of the red seaweeds of the order Gigartinales contain considerable amounts of polysaccharides [1]. However, *Sebdenia polydactyla* (Boergs) Balak, [2] a red alga from India, has not been chemically investigated. In this paper we show that it contains the phycocolloid,  $\lambda$ -carrageenan.

## RESULTS AND DISCUSSION

A water-soluble polysaccharide, which constituted ca 32% of the dry weight of the alga, was isolated as a white powder which on hydrolysis gave the monosaccharides, D-galactose and 3,6-anhydrogalactose. All its properties (Table 1) support the identity of the polysaccharide from *Sebdenia* as a  $\lambda$ -carrageenan.

Table 1. Content and properties of polysaccharide from *S. polydactyla*

Appearance	White powder
Yield (%)	32 $\pm$ 0.5
Total ash	25.97
Total sulphate (%)	17.2
Galactose (%)	32.8
3,6-Anhydrogalactose (%)	2.4
Aqueous gel strength (1 g in 1% KCl soln.)	No gel formation
Optical rotation (0.25% in H <sub>2</sub> O)	Positive, $[\alpha]_D^{25}$ 55°
Solubility in KCl soln. (3 M KCl)	Completely soluble, polysaccharide precipitated with <i>iso</i> -PrOH from the filtrate
Methylene Blue test	Positive, the phycocolloid precipitates Methylene Blue.
Solubility in H <sub>2</sub> O	Dissolves in cold H <sub>2</sub> O, but completely above 70°.
Solubility in milk	Insoluble in cold but soluble in hot milk
Milk reactivity at 25° (0.154% polysaccharide in homogenized milk)	< 50 g cm <sup>-2</sup>

IR spectra [3–6]	Intensity of absorption bands (cm <sup>-1</sup> )*			
	1240	930	840–850	805
Difco-Bacto agar (0140–01)	w	s	—	—
$\kappa$ -Carrageenan (C-1263)	vs	s	s	—
$\lambda$ -Carrageenan (C-3889)	vs	w	w	—
i-Carrageenan (C-4014)	vs	s	s	vs
<i>Sebdenia</i> polysaccharide	vs	w	w	—
<i>Sebdenia</i> polysaccharide (alkali modified)	vs	m	w	—

\*w, weak; m, medium; s, strong; vs, very strong.

## EXPERIMENTAL

*Sebdenia polydactyla* (Sebdeniaceae) was collected during Feb. from the low-tide water mark at Okha (22° 18'N, 69° 06'E) on the west coast of India. A voucher specimen identified by one of the authors (VDC) an algologist is deposited in CSMCRI Herbarium, Bhavnagar, India. The plants after collection were washed thoroughly in seawater, rinsed with H<sub>2</sub>O and dried in a hot air circulating oven at 60° followed by grinding. Carrageenan was extracted by the method of ref. [7].

Dry depigmented algal powder was initially extracted with 0.5 M NaHCO<sub>3</sub> soln at 90° for 3 hr. The hot viscous soln was pressure filtered and the residues re-extracted as above for another 30 min. Both filtrates were combined and filtered through glass fibre filter paper (Whatman GF/C). The phycocolloid was pptd with CTAB. The pptd material was washed sequentially with H<sub>2</sub>O, a saturated soln of NaOAc in EtOH, EtOH and Et<sub>2</sub>O and then dried *in vacuo* over P<sub>2</sub>O<sub>5</sub>.

Fractionation of the polysaccharide was carried out with KCl as described in ref. [5]. The samples were alkali modified following the procedure of ref. [8] with NaBH<sub>4</sub>. Aqueous gel strength was determined with 1% polysaccharide in 1% KCl soln. Sulphate was estimated gravimetrically after hydrolysis of the polysaccharide with MHCl for 2 hr at 105° in sealed tubes. The 3,6-AG content was obtained as per the modified resorcinol method of ref. [9] and galactose by PC [10]. Films for IR studies were prepared by evaporating a 0.2% soln over Hg [11]. Difco-Bacto agar (0140-01, Difco Labs USA),  $\kappa$ -carrageenan (C-1263),  $\lambda$ -carrageenan (C-3889) and  $\iota$ -carrageenan (C-4014) were procured from Sigma. The milk reactivity was determined by the standard procedure [12] and the Methylene Blue test was performed using a 1% soln of polysaccharide in H<sub>2</sub>O. KCl solubility was ascertained as follows:

The powdered polysaccharide was dissolved at 0.1% concn by warming to 70° in 3 M KCl, cooled to 25° and filtered. Obser-

vations for the dissolution of polysaccharide in KCl soln were made. To the filtrate, 2.5 vol. *iso*-PrOH was added to detect the reprecipitation of the algal polysaccharide.

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