

Preparation and Characterization of a Chemically Sulfated Cashew Gum Polysaccharide

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A goma do cajueiro (GC) foi sulfatada em piridina:formamida usando ácido clorossulfônico como reagente. A confirmação da sulfatação foi obtida por espectroscopia no infravermelho com transformada de Fourier (FTIR) pelo aparecimento da frequência de vibração em 1259 cm^{-1} devido ao estiramento assimétrico S=O. Os graus de sulfatação foram de 0,02, 0,24 e 0,88, determinados pela percentagem de enxofre. Os dados de ressonância magnética nuclear (RMN) 1D e 2D mostram que a sulfatação ocorreu nos carbonos primários do polissacarídeo. Um aumento de no mínimo 4% na viscosidade da solução foi observado após a sulfatação. Análise termogravimétrica (TGA) indica que o derivado é estável até cerca de $200\text{ }^{\circ}\text{C}$. A comparação entre os derivados sulfatados e carboximetilados foi feita no intuito de verificar a possibilidade de formação de complexos polieletrólitos, já realizada com os carboximetilados.

Cashew gum (CG) was sulfated in pyridine:formamide using chlorosulfonic acid as the reagent. Confirmation of sulfation was obtained by Fourier transform infrared (FTIR) spectroscopy through the presence of an asymmetrical S=O stretching vibration at 1259 cm^{-1} . The degrees of substitution were 0.02, 0.24 and 0.88 determined from the sulfur percentage. 1D and 2D nuclear magnetic resonance (NMR) data showed that the sulfation occurred at primary carbons. An increase of at least 4% of the solution viscosity was observed due to sulfation. The thermal gravimetric curves (TGA) indicate that the derivatives are stable up to *ca.* $200\text{ }^{\circ}\text{C}$. The sulfated CG is compared to carboxymethylated CG in order to verify the possibility of the use of the former in the preparation of polyelectrolyte complexes; the latter is already being used for this application.

Keywords: cashew gum, *Anacardium occidentale*, sulfation, NMR

Introduction

Sulfated carbohydrates are found in plants and animals. In plants, they occur in the intercellular and cell wall region of brown and red algae (fucoïdan, carrageenan).^{1,2} Much attention has been focused on the physiological role of sulfated polysaccharides and many attempts have been made to synthesize sulfated polysaccharides to obtain new pharmacological agents with possible therapeutic uses,³ which are strictly related to the presence of polyanionic charges.⁴ Polyanions can also be used to obtain polyelectrolyte complexes with polycations, such as chitosan. These polyelectrolyte complexes (PECs) have been proposed for many applications including scaffolds for pulp cell regeneration and matrixes for protein carriers

and antibiotics.^{5,6} The particle size, swelling and drug delivery behavior of PECs can be affected by the molecular parameters of oppositely charged polyelectrolytes. The formation of polyelectrolyte complexes (PECs) represents an easy way to produce various structures ranging from nanometer sizes (water-soluble aggregates and nanoparticles) to macroscopic states (physical gel and film).⁷

Brazil has approximately 680,000 ha planted with the cashew nut tree (*Anacardium occidentale*) and the average production of gum/tree/year is 700 g,⁸ with a potential annual cashew gum (CG) production of 50,000 tons.⁹ The gum from Ceará State has been characterized as a heteropolysaccharide containing (by wt.%) the following neutral monosaccharides: galactose (72-73%), glucose (11-14%), arabinose (4.6-5.0%) and rhamnose (3.2-4.0%).^{10,11} The ionic charge of the gum is due to glucuronic acid, however, this has a low content

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(4.7-6.3 wt.%). Different composition was found for gum from Pernambuco State: galactose (82%), glucose (6%), arabinose (4%), rhamnose (2%) and uronic acid (5%).¹² Independent of the origin, CG is mainly composed of three types of galactan units within the core, linked by C-1 and C-3, C-1 and C-6 and C-1, C-3 and C-6.^{10,11} Modifications have been carried out to cashew gum in order to increase its ionic charge in the carboxylate group, such as carboxymethylation¹³ and oxidation with TEMPO reagent.¹⁴ An overcharged new polysaccharide (carboxymethylated) has been used in the preparation and characterization of polyelectrolyte complex nanoparticles with chitosan to identify their application as a drug delivery system.¹⁵

In this study, sulfated derivatives of cashew gum were synthesized and characterized in order to obtain a more charged polyelectrolyte than the original gum through the introduction of a new chemical group, $-\text{OSO}_3\text{Na}$. The polyelectrolyte properties were compared with those of carboxymethylated CG,¹³ which were already used in the preparation of nanoparticles of PECs,¹⁵ to verify the possibility of obtaining polyelectrolyte complexes with sulfated CG.

Experimental

Origin and purification

Crude samples of CG were collected from native trees from Fortaleza (Ceará State, Brazil). They were purified as a sodium salt using a previously described method.^{11,16} Nodules free of bark were selected and dissolved in distilled water at room temperature to give a 5% (m/v) solution. The solution pH was adjusted to *ca.* 7.0 by addition of diluted aqueous NaOH. The clear solution was successively filtered through sintered glass and the polysaccharide precipitated with ethanol.

Sulfation

The sulfation procedure was based on the methodology described by O'Neill¹⁷ and Ono *et al.*¹⁸ with modifications. This method uses chlorosulfonic acid (CSA) as the sulfation agent. CSA is a versatile reagent widely¹⁹ and recently^{4,21-24} used in the sulfation of polysaccharide. The purified cashew gum sample (1 g) was swelled in pyridine:formamide (60:75 mL) with stirring at 25 °C (12 h). The solution was cooled to 4 °C and CSA was dropped into the mixture over a period of 30 min. The mixture was kept at this temperature overnight. The procedure was performed with the following acid volumes 2, 6 and 8 mL to obtain different CG gum/CSA molar ratios. The resulting solution

was neutralized with NaHCO_3 , dialyzed against water for 3 days and freeze dried. The solid product was dissolved in water and precipitated with 2 volumes of ethanol. The samples were denoted as CGS2, CGS6 and CGS8, with the number corresponding to the volume of CSA.

Measurements

The contents of nitrogen, carbon and sulfur were determined for the sulfated cashew gum (CGS) by elemental microanalysis using a Perkin-Elmer CHN 2400 instrument.

Fourier transform infrared (FTIR) spectra were recorded with KBr pellets on a FT-IR Shimadzu 8300 spectrophotometer.

1D (^1H and ^{13}C broad band, BB) and 2D (^1H - ^{13}C heteronuclear single-quantum correlation, HSQC) spectra of 3% (m/v) solutions in D_2O at 343 K were recorded on a Fourier transform Bruker Avance DRX 500 spectrometer. An inverse multinuclear gradient probe-head equipped with z-shielded gradient coils, and with Unix Silicon Graphics workstation was applied. Sodium 2,2-dimethylsilapentane-5-sulfonate (DSS) was used as the internal standard (0.00 ppm for ^1H). A distortionless enhancement through the polarization transfer (DEPT 135) spectrum was recorded in order to determine the hydrogenation of each carbon.

Thermogravimetric analysis (TGA) of the samples were carried out using a Shimadzu TGA-50 instrument at a heating rate of 10 °C min^{-1} over the temperature range of 25-700 °C. The air flow rate was maintained at 50 mL min^{-1} and the initial sample mass was 10 mg.

The elution volumes were determined by gel permeation chromatography using a Shimadzu LC-10AD chromatograph with a refractive index detector RID-6A at room temperature using an Ultrahydrogel linear column (7.8 × 300 mm), flow rate of 0.5 mL min^{-1} , polysaccharide solution concentration of 0.1% (m/v), water as the solvent and 0.1 mol L^{-1} NaNO_3 as the eluent. The sample volume was 50 μL .

Rheological studies of CG and CGS aqueous solutions (1%, m/v) were performed on a Rheometer AR550 (TA instruments) at 25 °C with 40 mm cone plate sensor. The effect of shear rate on the solution viscosity was evaluated.

Results and Discussion

Effect of chlorosulfonic acid concentration on yield and degree of sulfation

The degree of sulfation (DS) obtained in reactions with different CSA volumes was determined. The DS is defined

as the average number of $-\text{OSO}_3\text{Na}$ groups inserted in each monosaccharide unit. As the main constituents of cashew gum are hexoses,¹⁰ the DS was considered as the number of $-\text{OSO}_3\text{Na}$ groups *per* 6 carbons. From data on the S and C content, it is possible to calculate the DS, in a similar way to that used by Melo *et al.*²⁵ The CG contains 1% of protein with *ca.* 70% of amino acid without sulfur.²⁶ The S content from protein was very low and its contribution in the calculus insignificant. Hence, the DS of sulfated cashew gum can be determined using C and S% through equation 1.

$$\text{DS} = \frac{\left(\frac{\text{S}\%}{\text{atomic mass of S}} \right)}{\left(\frac{\text{C}\%}{\text{atomic mass of C}} \right)^6} = 2.25 \left(\frac{\text{S}\%}{\text{C}\%} \right) \quad (1)$$

Calculated DS are shown in Table 1. Different values for cashew gum derivatives were produced by varying the amount of sulfation reagent, SR, in this case chlorosulfonic acid. For cashew gum, the DS increases with the SR/sugar unit (SR/SU) ratio. A similar trend has been observed for the sulfation of polysaccharides from seeds of *Lupinus polyphyllus*²⁷ and cellulose,²³ but only up to SR/SU ratios of 2.6 and 4.5, respectively. At a higher sulfation reagent/polysaccharide ratio, a lower DS was observed for both polysaccharides.

With the comparison of DS of different polysaccharides at similar SR/SU ratios (Table 1), it can be concluded that the reaction conditions applied to the cashew gum

led to a less efficient sulfation than that reported by other authors.^{23,28} This result could be attributed to the polysaccharide concentration and temperature, being lower than that applied in the other studies (temperature above 30 °C and polysaccharide concentration greater than 5%), as well as the use of different sulfation agents. A temperature of 4 °C has been adopted in the case of cashew gum in order to avoid serious degradation of the polysaccharide.¹⁷ When the same temperature, polysaccharide concentration and solvents were utilized, DS values of 0.50 and 0.62 were obtained by Ono *et al.*,¹⁸ even with a greater sulfation reagent concentration (SR/SU = 31.6).

The reaction yields obtained for different CSA volumes are also listed in Table 1. An increase in the amount of CSA promoted a significant decrease in the sulfation yield. The comparison of the reaction yields of cashew gum and of other gum derivatives obtained using similar methodology was not possible, since this information was not available in either of the reports by O'Neill¹⁷ and Ono *et al.*¹⁸ The apparent yields of more than 100% were reported by Geresh *et al.*²⁹ in the sulfation of extracellular polysaccharides of red microalgae. In this case, although not specified, the yields were calculated as the mass of sulfated product in relation to the initial mass of polysaccharide. However, the more precise yields are those calculated taking into account that the introduction of each $-\text{SO}_3\text{Na}$ group causes an increase of 102 g mol⁻¹. In the case of cashew gum, galactose is the major constituent.¹⁰ Therefore, the final molar mass of sulfated product is expected to be 162 + DS × 102 g mol⁻¹, where 162 g mol⁻¹ is the molar

Table 1. Comparison between the sulfation reaction conditions and the characteristics of the obtained polysaccharide derivatives

| Sample | Polysaccharide conc. / (% m/v) | Sulfation reagent | SR/SU ^a | Temperature / °C | Yield ^b / (g %) | DS | Ref. |
|---------------------------------------|--------------------------------|----------------------|--------------------|------------------|----------------------------|------|-----------------------------------|
| Pullulan | 20 | SO ₃ -py | 4 | 60 | 91 | 1.50 | Mihai <i>et al.</i> ²⁸ |
| | 20 | SO ₃ -py | 4 | 80 | 90 | 1.22 | |
| | 20 | SO ₃ -DMF | 4 | 60 | 42 | 1.32 | |
| | 20 | SO ₃ -DMF | 4 | 80 | 6 | 1.22 | |
| Cellulose ^c | 5 | CSA-DMF | 1.5 | 30 | 67 | 0.52 | Wang <i>et al.</i> ²³ |
| | 5 | CSA-DMF | 3.0 | 30 | 36 | 1.93 | |
| | 5 | CSA-DMF | 4.5 | 30 | 32 | 2.01 | |
| | 5 | CSA-DMF | 6.0 | 30 | 35 | 1.53 | |
| Epimedium polysaccharide ^d | 2 | CSA-py | 0.86 | 60 | 20 | 1.12 | Lu <i>et al.</i> ²² |
| | 2 | CSA-py | 0.86 | 80 | 16 | 0.83 | |
| | 2 | CSA-py | 0.86 | 95 | 39 | 0.35 | |
| Cashew gum | CGS2 0.7 | CSA-py-FA | 4.9 | 4 | 53 | 0.02 | this study |
| | CGS4 0.7 | CSA-py-FA | 14.6 | 4 | 26 | 0.24 | |
| | CGS8 0.7 | CSA-py-FA | 19.4 | 4 | 15 | 0.88 | |

^aMolar ratio of sulfation reagent to sugar unit; ^bcalculated using equation 2; ^creaction time = 1 h; ^dhighest DS values were chosen, from different CSA/py ratios, CSA = chlorosulfonic acid, py = pyridine, DMF = dimethylformamide and FA = formamide.

mass of the linked galactose units. Equation 2 expresses the most appropriate way to calculate the yield (%):

$$\text{Yield} = 100 \frac{m_{\text{final}}}{\left[1 + \left(\frac{\text{DS} \cdot 102}{162} \right) \right] m_{\text{inicial}}} \quad (2)$$

where m_{final} and m_{inicial} are the mass of sulfated product and of polysaccharide, respectively.

There are many methods which can be used to sulfate polysaccharides, for instance, employing sulfuric acid, sulfur trioxide-pyridine, sulfur trioxide-dimethylformamide, chlorosulfonic-pyridine (py) or chlorosulfonic-pyridine-formamide. A comparison of the obtained yields from different methods applied to polysaccharides is shown in Table 1. The reaction conditions are very diverse in terms of polysaccharide concentration, molar ratio of sulfation reagent to sugar unit, time and temperature of reaction, sulfation agent and solvent. The cashew gum sulfation was carried out with the lowest gum concentration, highest SR/SU ratio and lowest temperature. The obtained yield for the sulfation of CG is in the range of those reported for cellulose and epidemium polysaccharide using CSA as the sulfation agent (16-67%).^{22,23} A higher yield was registered for pullulan sulfated with SO_3 -pyridine, but a much more concentrated polysaccharide solution was used.²⁸ A lower yield at higher concentrations of chlorosulfonic acid was observed for cashew gum and may be attributed to degradation of the polysaccharide chain.

DS is an important parameter in terms of PEC formation because the charge density is the main driving force of the reaction. In addition, the particle size of PECs was influenced by the molar ratio of cationic to anionic units.³⁰ Carboxymethylated cashew gum derivatives with DS values of 0.16 and 0.36 were used to obtain nanoparticles of PEC with chitosan.¹⁵ The authors observed that the particle size increased with DS. The sulfated cashew gum derivatives showed the same range of negative charge as that observed for carboxymethylated CG. The formation of nanoparticles of CGS with chitosan is expected, but the dependence on DS is different due to the diverse polysaccharide conformation, which will be the subject of further research.

Infrared spectroscopy

The presence of sulfate groups in CGS was identified by comparing the FTIR spectra of CG and CGS derivatives (Figure 1). The spectrum for the sulfated cashew gum shows a new band at 1259 cm^{-1} , attributed to the asymmetrical

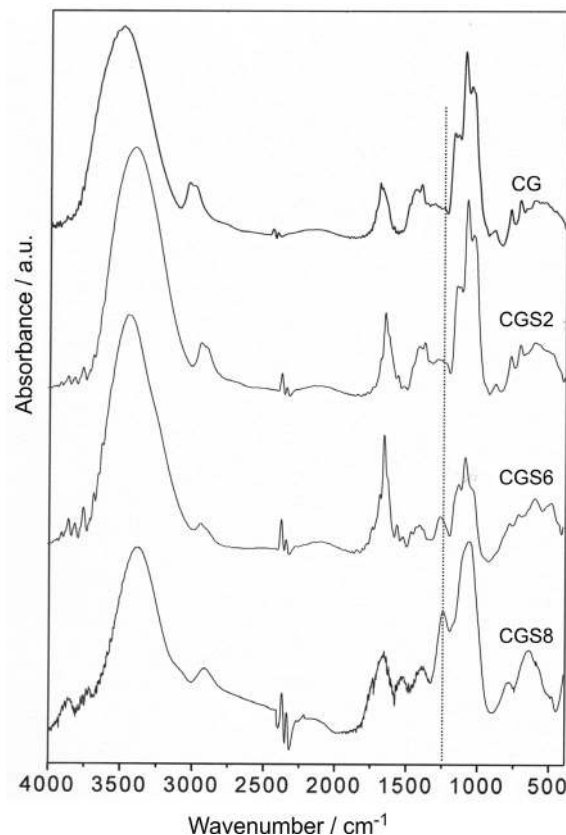


Figure 1. Infrared spectra obtained for unmodified (CG) and sulfated cashew gums (CGS2, CGS6 and CGS8).

stretching of the S=O vibration.³¹ This new band appears for all sulfated samples and becomes more intense as the DS increases, from CGS2 to CGS8. The profile of the $900\text{-}600 \text{ cm}^{-1}$ region for the CGS6 and, particularly, for the CGS8 gums, seems to differ from that of the CG. For cashew gum, the bands at 775 cm^{-1} can be assigned to the C–O–C bending vibration, and those at 711 and 603 cm^{-1} to deformation and scaffold vibrations, respectively.³¹ For CGS, the band at 711 cm^{-1} disappears and the other two are shifted to higher wavenumbers (781 and 638 cm^{-1}). A shift to higher values in the order of 10 to 40 cm^{-1} was also observed by Mahner *et al.*³¹ after sulfation of pullulan. This indicates that although not attributed directly to sulfate group vibration, the band shifts in the $800\text{-}600 \text{ cm}^{-1}$ region are correlated to the sulfation.

Nuclear magnetic resonance

The ^{13}C BB NMR spectrum (Figure 2) for CG is similar to that obtained by de Paula *et al.*¹⁰ The ^1H - ^{13}C HSQC spectrum for CG (Figure 3(a)) shows the correlation of previously assigned ^{13}C data¹¹ with anomeric protons: α -D-glucose (δ 4.95/100.7), α -L-rhamnose (δ 4.81/101.3), β -D-galactose ($1 \rightarrow 3$) (δ 4.69/104.1 and δ 4.43/104.2),

β -D-glucuronic acid (δ 4.55/103.8), and β -D-galactose (1 \rightarrow 6) (δ 4.39/105.0).

The ^{13}C BB NMR spectrum for CGS6 derivative (Figure 2) shows some differences in relation to unmodified gum. The anomeric signals decrease considerably due to glucose and rhamnose residue, probably because of the chain degradation. A new signal at δ 67.8 is observed for CGS6 in comparison with the CG spectrum. In ^{13}C DEPT NMR spectrum (not shown), the signal at δ 68.7 appeared with the opposite amplitude to those of CH_3 and CH , which can

be attributed to the sulfation of CH_2 primary carbons (C-6). The presence of a sulfate group causes an increase in the ^{13}C chemical shift of 6-8 ppm for the α carbons, and a shift of 0.4-0.7 ppm for protons.^{4,23,32} The ^1H - ^{13}C HSQC NMR spectrum for CGS6 (Figure 3(b)) shows a correlation with δ 4.2/67.8 (very small signal on the ^{13}C BB NMR spectrum), which confirms the substitution at CH_2 (C-6). Since protons H-6 from unsubstituted sugar residues occurred between 3.6 and 3.9 ppm (Figure 3(a)), the new signal at δ 4.2 indicates sulfation on the residues which have a primary group.³² The

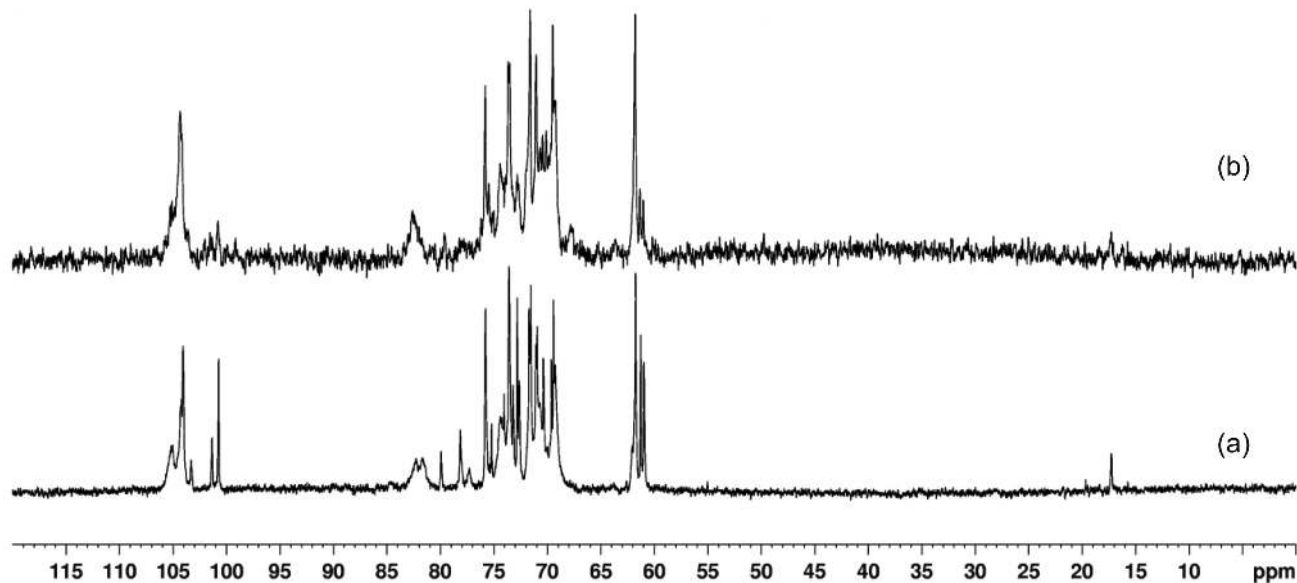


Figure 2. ^{13}C NMR spectra obtained for (a) unmodified cashew gum (CG) and (b) sulfated cashew gum (CGS6).

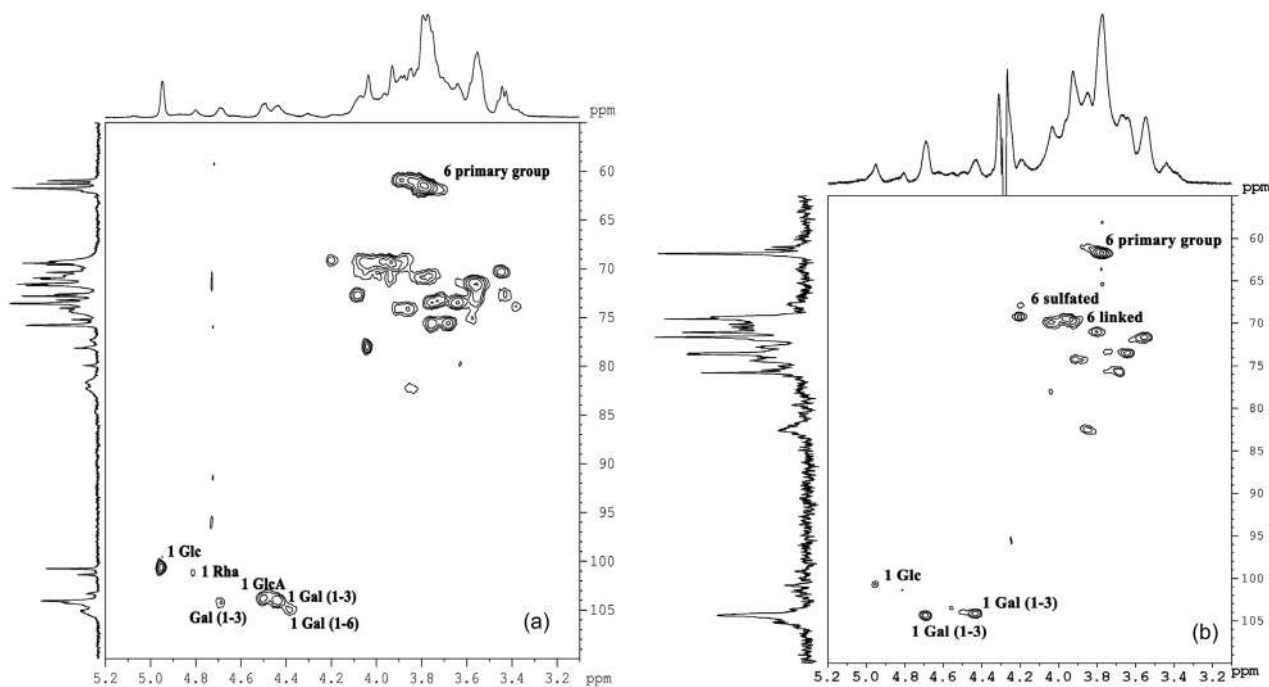


Figure 3. ^1H - ^{13}C HSQC spectra obtained for (a) unmodified cashew gum (CG) and (b) sulfated cashew gum (CGS6).

DEPT 135 NMR spectrum for CGS6 (see Supplementary Information) and the HSQC NMR experiment (Figure 3(b)) indicate that 6-O-substituted galactose units (δ 4.03/70.3) remain after the sulfation reaction.

Gel permeation chromatography

The molar mass and the degree of sulfation of the polysaccharide are important parameters influencing polyelectrolyte complexes.³⁰ The cashew gum has a peak molar mass of 2.3×10^4 g mol⁻¹. The absolute value of the average molar mass of sulfated products can only be determined by GPC if the chromatograph is coupled to a detector such as a multiangle laser light scattering (MALLS) instrument, in which no standards are required to calibrate the column. When other detectors are used, such as a refractive index detector, standards of different molar masses are required. Pullulan is frequently used for polysaccharides. However, this standard is not appropriate for charged polysaccharides due to the chain stiffening and length, as a consequence of electrostatic repulsion.

The GPC chromatograms for CG and sulfated derivatives can be seen in Figure 4. Taking into account that a MALLS detector was not available for this study, the discussion on the molar mass is based on the elution volumes and comparison. A shift to a lower elution volume was observed comparing the curves for CG ($V_e = 8.84$ mL) and CGS6 ($V_e = 8.74$ mL). If no chain degradation occurs, the sulfated samples will have a higher hydrodynamic volume and would be expected to elute at a lower volume than the starting polymer due to molar mass increase and repulsion between new charged groups. It is expected an increase in the molar mass due to the introduction of the $-\text{SO}_3\text{Na}$ group. In the CGS6 (DS = 0.24), if no degradation

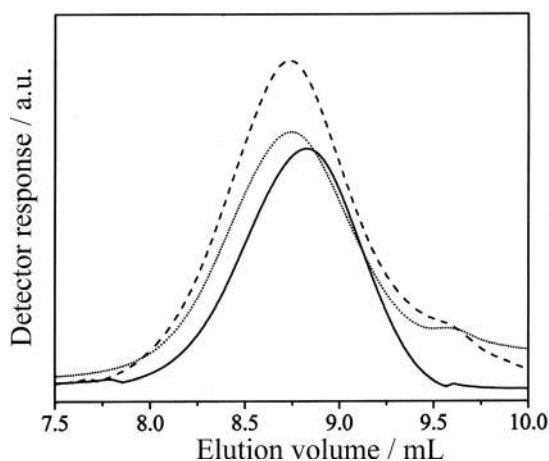


Figure 4. GPC chromatogram of aqueous solutions of unmodified CG (continuous line) and sulfated cashew gums CGS6 (dotted line) and CGS8 (dashed line).

occurs, the molar mass increases by around 15%. This higher molar mass is expected to change the elution volume by more than 0.10 mL. Thus, chain degradation probably occurs. The shoulder at a higher elution volume and the small yield are in agreement with this assumption.

CGS8 showed an elution volume similar to that of CGS6, even with a higher DS (0.88). Since the molar mass and also chain charge increase from CGS6 to CGS8, a higher elution volume was expected. Therefore, the chain degradation in CGS8 is higher than in CGS6. An increase in degradation as the DS increase was also observed in the sulfation of galactan²⁴ and lacquer polysaccharide.³³ Higher degradation with an increasing degree of derivatization was described for carboxymethylation¹³ and oxidation of cashew gum.¹⁴

Considering that no peak or shoulder at an elution volume comparable with that of the precursor gum was noted in CGS6 or CGS8 chromatogram, no unmodified gum remains. A different result was reported by Mihai *et al.*,²⁸ who found that 40% of the pullulan remains non sulfated in the $\text{SO}_3\cdot\text{py}$ complex at 80 °C over 4 h. These results were in agreement with the NMR data and the yield values.

Carboxymethylation of cashew gum leads to higher degree of degradation than that observed in the sulfation reaction.¹³ This is an important result when the objective is to prepare PECs, as concluded by Schatz *et al.*³⁰ in a study on the preparation of PEC using dextran sulfate and chitosan.

Thermal analysis

The thermogravimetric curves of the cashew gum and sulfated derivatives are shown in Figure 5 and the parameters obtained from the curves in Table 2. The comparison of the curves reveals that the thermal decomposition processes of the CG and CGS samples are different. The first event (T_{max} ca. 50 °C) is related to water loss and is present in all curves. Cashew gum shows an event at T_{max} 400 °C, which is not present in the case of sulfated samples. CGS gums show a new event at T_{max} 500-520 °C that is not present in the TGA of the original gum. Similar results were obtained for carboxymethylated cashew gum.³⁴ This event is probably related to thermal scission of carbohydrate segments containing sulfate groups, despite of the not proportional mass loss to the degree of sulfation.

The residues of the derivatives CGS6 and CGS8 at 700 °C increased by 15.8 and 23.9%, respectively, in comparison with the cashew gum residue (6.4%). The sulfated gum with higher DS showed the highest residue, as expected. The stability of the derivatives is around 200 °C.

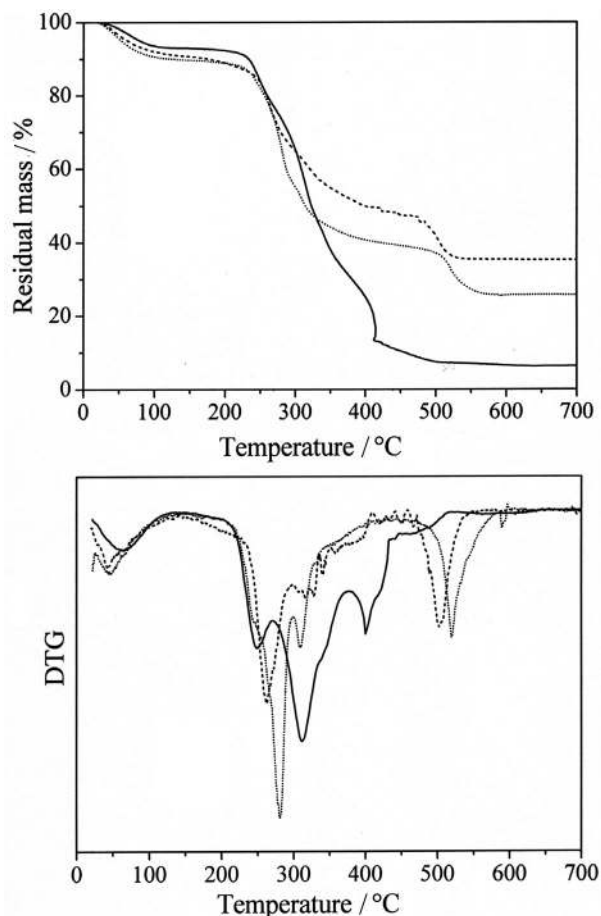


Figure 5. TGA and DTG curves obtained for unmodified CG (continuous line) and sulfated cashew gums CGS6 (dotted line) and CGS8 (dashed line) under synthetic air atmosphere.

Rheological studies

Aqueous solutions (1%) of CG and CGS samples have low viscosities (Figure 6). The CGS6 gum curve is very similar to the original gum across the whole shear rate range. However, the sample with higher DS (CGS8) shows a distinct behavior up to a shear rate of 80 s^{-1} . At 30 s^{-1} , for example, the viscosity increase of CGS8 is around 30% the value for the original gum. At a shear rate above 80 s^{-1} (Newtonian Plateau), the sulfation promotes an increase in viscosity of only 4% in comparison to the unmodified CG. The higher negative charge induces more repulsion

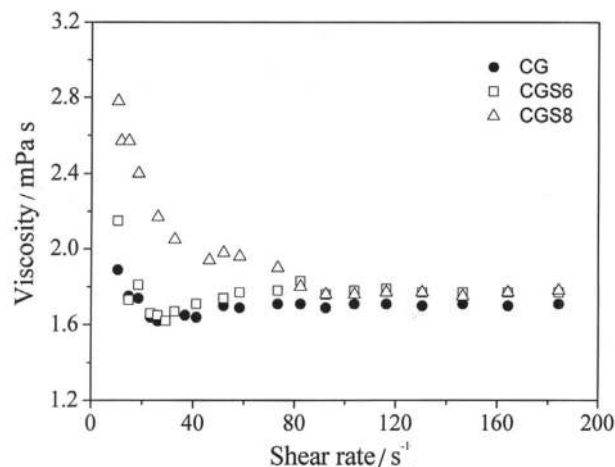


Figure 6. Effect of shear rate on the viscosity of aqueous solutions: unmodified CG, sulfated cashew gums CGS6 and CGS8. Gum concentration 1.0% (m/v) at $25 \text{ }^\circ\text{C}$.

between chains and a more expanded macromolecular conformation. However, Geresh *et al.*²⁹ found for sulfated extracellular polysaccharides that the viscosity of sulfated products was lower than that of the starting material, even with no evidence of degradation.

Conclusions

Three sulfated derivatives of cashew gum were synthesized and characterized. The new sulfate group was detected by FTIR and microanalysis. The sulfation occurs preferentially at the primary carbons of galactose units as observed in the 1D and 2D NMR experiments. The reaction also causes chain degradation, but to a smaller degree than carboxymethylation of CG. A slightly higher viscosity was observed for aqueous solutions of sulfated CG in comparison to the unmodified gum. The sulfation degree is low, but sufficient to increase the ionic charge of the polymer and to improve its polyelectrolytic properties. Sulfated cashew gum offers characteristics suitable for its use in the preparation of nanoparticles of polyelectrolyte complexes. It is more attractive because the cashew gum is a natural, abundant, non-toxic, low cost and regional raw material.

Table 2. Parameters from TGA and DTG curves for the CG and sulfated CGS samples in synthetic air atmosphere at $10 \text{ }^\circ\text{C min}^{-1}$

| Sample | Maximum temperature of event / $^\circ\text{C}$ | | | | | Residual mass at $700 \text{ }^\circ\text{C}$ / % |
|--------|---|-----|-----|-----|-----|---|
| | I | II | III | IV | V | |
| CG | 55 | 248 | 311 | 400 | – | 6.4 |
| CGS6 | 46 | 291 | 309 | – | 520 | 22.2 |
| CGS8 | 43 | 262 | 316 | – | 502 | 30.3 |

Supplementary Information

The DEPT data are provided as supplementary information and available free of charge at <http://jbcs.sbq.org.br> as PDF file.

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Preparation and Characterization of a Chemically Sulfated Cashew Gum Polysaccharide

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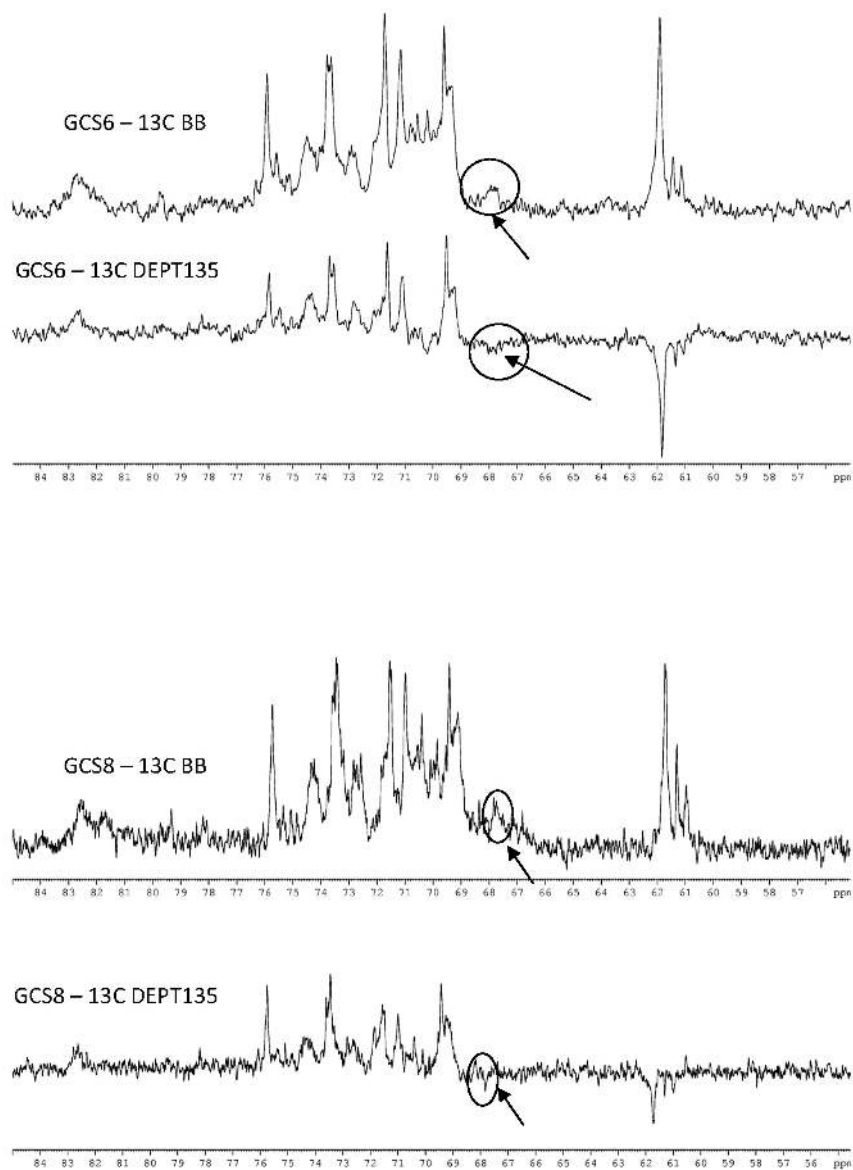


Figure S1. DEPT 135 and BB spectra of sulfated cashew gum (samples GCS6 and GCS8). In sulfated samples (GCS6 and GCS8) spectra a new signal is observed at δ 67.8 in both spectra. The same signal is visualized in opposite amplitude to those of CH_2 and CH signals in DEPT 135 experiment so based on this observation and that substitution of sulfate group on primary carbons causes an increase of chemical shift of 6 to 8 ppm^{4,23,32} we concluded that the sulfate substitution on sulfated cashew gum occur on primary carbons (C-6).

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