Physicochemical characterization and evaluation of PGA bioflocculant in coagulation-flocculation and sedimentation processes


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This paper analyzes the effectiveness of three commercial products: aluminum sulfate, aluminum polychloride and polyglutamic acid-based bioflocculant, in water clarification and purification. Poly-γ-glutamic acid (PGA), a naturally occurring anionic polymer, is already produced on an industrial scale. A conventional jar test apparatus was used for flocculation experiments. The tests were performed using raw water from the Salto de Pirapora Water Treatment Plant (WTP) in the state of Sao Paulo, Brazil. The structure and elemental composition of PGA bioflocculant, which was characterized by Fourier transform infrared spectroscopy (FTIR), scanning electron microscopy (SEM), X-ray diffraction (XRD), X-ray fluorescence spectrometry (XRF) and thermogravimetric analysis (TGA), suggests that it is basically inorganic in nature, with high contents of calcium sulfate (87%) and γ-polyglutamic acid (5%), among other chemical compounds. The performance of PGA bioflocculant was superior to that of the other coagulants. In the dry season, 80 mg L⁻¹ of PGc21Ca removed 89.74% of apparent color, and residual turbidity was 5.78 NTU. In the rainy season, 90 mg L⁻¹ of PGc21Ca removed 96.89% of apparent color and residual turbidity was 1.93 NTU.

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1. Introduction

Conventional water treatment methods include coagulation, flocculation and sedimentation processes, usually followed by filtration and disinfection. Coagulation–flocculation is considered the most important process in surface water treatment, and it can be used as a pre-treatment, post-treatment, or even main treatment [1]. The main parameters used in the physical characterization of natural water are color, turbidity, the various fractions of solids levels, temperature, taste and odor. Although these are physical parameters, they provide valuable preliminary information to characterize the chemical quality of water, e.g., suspended solids levels (associated with turbidity) and concentrations of dissolved solids (associated with color), organic solids (volatile) and mineral solids (fixed), which are the compounds that produce odor, among other things.

The purpose of removing or reducing turbidity is to increase the efficiency of chlorination to disinfect water. In a water treatment plant (WTP) that uses the complete water cycle treatment, a decrease in process efficiency and in rapid-mix, flocculation and sedimentation operations causes clogging of the filters, which thus require more frequent washing, and may lead to increased turbidity and less effective disinfection of the water supply. Hence, the parameters of rapid-mix, flocculation and sedimentation markedly influence the overall removal efficiency of impurities contained in raw water. Therefore, these parameters should not be simply adopted, but established based on laboratory studies or a pilot plant. The coagulation–flocculation process also enables the identification of potential products that can be used as coagulants for water treatment, some of which are readily available, such as aluminum and iron-based coagulants, while others are experimental coagulants, such as PGA bioflocculant. According to the literature, the performance of each coagulant may also vary, depending on the type of water to be treated.

Coagulants are used extensively in the treatment of public water supply systems. The two most widely used coagulants are aluminum and iron salts [2]. Aluminum sulfate was commonly used in Brazil because of its low cost and high efficiency in the treatment of public water supply. However, aluminum is not biodegradable and high concentrations of this metal are harmful to health, possibly even aggravating disorders such as Alzheimer’s disease [3,4]. Currently, WTPs use prepolymerized aluminum salt, known as polyaluminum chloride (PAC). Due to its basicity, PAC generally releases less acid than other coagulants during hydrolysis, causing little variation in the pH level of treated water. For this
reason, media coagulated by PAC require little or no use of pH adjustment products. In addition to requiring only very slight pH adjustment, PAC is efficient in destabilizing particles, thus favoring the rapid formation of large uniform flakes without the need for flocculating aids. Although aluminum sulfate and PAC present favorable characteristics as inorganic coagulants, chemical formulations containing aluminum reportedly have harmful effects [5] and commonly produce a large amount of hazardous sludge containing residual metals [6]. Thus, there is a great demand for the development of a coagulant that poses no risk to the environment or to human health. To this end, numerous polymer matrices have been proposed for the removal of contaminants from water and for the treatment of wastewater. Biodegradable flocculants could represent an alternative to the conventional flocculants currently in use. Reports are available on the application of biofloculants such as chitosan, which may be used for both coagulation and flocculation [1,7]. Sellami et al. [8] evaluated cactus juice as a biofloculant in the coagulation-flocculation process for industrial wastewater treatment. Campos et al. [9,10] polymerized bentonites to obtain ionene derivatives, combining polysoap properties with organophilic properties for possible application in the removal of organic pollutants.

Poly-γ-glutamic acid (PGA) is commercially available as PGA21Ca, a sodium salt containing 14% poly-γ-glutamic acid, which is stabilized with minerals such as calcium sulfate [11]. The biopolymer γ-polyglutamic acid (γ-PGA) is anionic, water-soluble, biodegradable, biocompatible, edible and non-toxic to humans and the environment. Due to these special characteristics, it has been applied in the environmental sector as a coagulant agent. Several researchers have described the use of γ-PGA in water treatment [12–15]. Shih and Van [16] stated that poly-g-glutamic acid is a polypeptide flocculant and it is considered the best option because of its high yield, high flocculating activity and ability to flocculate a wide range of organic and inorganic compounds.

The purpose of this study was to evaluate the efficiency of commercial coagulants based on coagulation, flocculation and sedimentation tests. Experimental series were performed using traditional products such as aluminum sulfate, polyaluminum chloride and polyglutamic acid-based PGA biofloculant, a Japanese product described as effective in the decontamination of water for human consumption. Another aim of this study is to understand the behavior and physicochemical stability of PGA21Ca.

2. Material and methods

The biofloculant PGA21CaTM from Nippon Poly-Glu Co. Ltd. was supplied in powder form, with a recommendation for the use of 0.05–0.1 g L⁻¹. The product’s typical flocculation time is 15 min at an initial turbidity of 100 NTU, resulting in a final turbidity of <1 NTU. The aluminum sulfate and aluminum chloride were supplied by the Salto de Pirapora WTP.

Raw water was collected from the Salto de Pirapora WTP, because of its importance for the metropolitan area of Sorocaba. The sample was digested with concentrated nitric acid and heated to 100 °C in a block digestor to determine the cations in the raw water, using inductively coupled plasma optical emission spectroscopy (ICP-OES, Spectro ArcoS) in radial-view mode. The chromatographic method (HPLC/IC model 430) was used for the simultaneous determination of anions.

The PGA21Ca was characterized to determine its chemical composition, thermal stability, and morphology by means of X-ray fluorescence spectrometry (XRF), X-ray diffraction (XRD), elemental analysis, infrared spectroscopy, thermal analysis and scanning electron microscopy. Before subjecting the PGA21Ca to physicochemical characterization, it was sifted through a stainless steel sieve (Tyler, 200 mesh) with 0.075 mm sieve opening size, placed in stoppered flasks, and stored in a desiccator.

The PGA21Ca was analyzed by energy dispersive X-ray fluorescence (EDXRF) using a PANalytical Epsilon 3 spectrometer. Basal spacings were determined by X-ray diffraction (XRD) in a Rigaku MiniFlex diffractometer, using CuKα radiation (λ = 0.15416 nm) and a scintillation detector at 30 kV and 15 mA at a rotation of 5–80° at a scan of 2θ, applying 0.02° steps at 0.6 s/step. The data were analyzed using Diffraction Plus software, integrated with JCPDS – International Center for Diffraction Data (ICDD) version 2001.

The purpose of the elemental analysis was to identify elements commonly found in organic compounds, such as carbon, hydrogen and nitrogen, and was performed in a Perkin Elmer 2400 CHN elemental analyzer. The elemental analysis was performed in duplicate.

The infrared spectrum of adsorption was obtained by means of Fourier transform infrared spectroscopy (FTIR), using a Perkin Elmer Frontier FTIR spectrometer.

The PGA21Ca was thermally analyzed in the temperature range of 10–1000 °C, applying a heating rate of 20 °C min⁻¹. The samples were degraded in a TA Instruments 50 thermogravimetric analyzer (TGA 50) in an air oxidizing atmosphere and nitrogen atmosphere, using a nitrogen flow of 50 cm³ min⁻¹ and a synthetic air flow of 100 mL min⁻¹. A sample of about 12 mg was used in all the experiments.

Structural information about PGA21Ca was obtained using a conventional scanning electron microscope (SEM). The micrograph was magnified by up to 5000 X after coating the surface of the sample with Au. The analysis was performed using a scanning electron microscope (LEO 440i) coupled to an energy dispersive X-Ray spectrometer (EDS) (Oxford Si (Li) detector).

3. Results and discussion

3.1. Physicochemical characterization of PGA21Ca

The specific surface area (S BET) of PGA21Ca was determined by the Brunauer-Emmet-Teller (BET) method through physical adsorption of N₂ at low temperatures, using measurements in the relative pressure range (P/Po) of 0.05–0.1. The pore volume was calculated by the Barrett-Joyner-Halenda (BJH) method [17] and by means of the N₂ adsorption isotherm. PGA21Ca was found to have a specific surface area of 1799 m² g⁻¹ and a pore volume of 0.004 cm³ g⁻¹.

The physicochemical analysis of PGA21Ca was performed via EDXRF, which revealed the percentage of oxides in the commercial product (Table 1). The results were normalized and expressed as Na and U oxides, excluding light elements such as H, He, Li, Be, B, C, N, O, F, and Ne.

The elemental analysis of PGA21Ca indicated CHN percentages in the composition of the commercial product, which are listed in Table 2. According to Ho et al. [18], an elemental analysis of γ-polyglutamic acid typically shows 44.86% (C), 5.91% (H) and 10.49% (N), which is consistent with the composition calculated here as 46.51% (C), 5.43% (H) and 10.85% (N). The γ-polyglutamic acid content in PGA21Ca was estimated considering that the formula of γ-polyglutamic acid is (C₅H₁₂NO₃)ₙ, while that based only on the monomer is 46.51% (C), 5.46% (H), 10.84% (N) and 37.17% (O). However, it is prudent to deduce the γ-PGA content from the

<table>
<thead>
<tr>
<th>Sample</th>
<th>SiO₂</th>
<th>Al₂O₃</th>
<th>SO₃</th>
<th>MgO</th>
<th>CaO</th>
<th>Na₂O</th>
<th>K₂O</th>
<th>Cl₂O</th>
<th>Fe₂O₃</th>
<th>TiO₂</th>
</tr>
</thead>
<tbody>
<tr>
<td>PGA21Ca</td>
<td>6.63</td>
<td>2.97</td>
<td>47.33</td>
<td>0.49</td>
<td>40.41</td>
<td>0.09</td>
<td>0.13</td>
<td>0.99</td>
<td>0.71</td>
<td>0.05</td>
</tr>
</tbody>
</table>

*Table 1. Chemical composition of PGA21Ca analyzed by X-ray fluorescence (in mass%).*
carbon analysis, based on the assumption that this element is present only in \( \gamma \)-PGA, since hydrogen may be present in other salts or as hydration water and the nitrogen content was negligible. Owing to this low \( \gamma \)-PGA content in the sample, the nitrogen results were not considered sufficiently reliable to deduce its content by the analytical combustion technique used here. In this case, considering the carbon content in \( \gamma \)-PGA, the commercial product contained 5.26%.

### 3.2. X-ray diffraction

The mineralogical composition of the sample was determined qualitatively based on an X-ray diffraction analysis. The XRD analysis showed peaks characteristic of gypsum \( \text{CaSO}_4 \cdot 2\text{H}_2\text{O} \) and anhydrite \( \text{CaSO}_4 \). The identified compounds are summarized in Fig. 1 and Table 3. No mineralogical species typical of silicates were found, but the chemical analysis revealed silicon, which may be present as amorphous silica.

### 3.3. PGa21Ca morphology

The microstructure of the sample was examined by SEM. To this end, the powder sample was spread on an adhesive, double-stick carbon tape which was then pressed onto a 0.5 inch diameter stub. Various areas of the sample spread on the stub were examined, and after ensuring that its morphology and morphometry did not vary, images were recorded by a secondary electron detector.

The SEM micrographs revealed morphologies typical of gyspum (Fig. 2). The mineral appeared in the form of thick platelets ranging in size from 10 to 30 \( \mu \)m, and at another moment, as aggregates of up to 30 \( \mu \)m in size, composed of particles of approximately one micron. The representativeness of the sample can be shown in other micrographs. Chemical mapping of its composition by EDS revealed the presence of Ca, S, Si and Al.

### 3.4. Infrared spectroscopy

The FTIR-ATR technique was used to identify the typical chemical groups of \( \gamma \)-PGA in PGa21Ca, by means of the characteristic bands. The infrared spectra were obtained from FTIR grade potassium bromide pellets in the range of 4000–4000 cm\(^{-1}\) for observation of the functional groups (Fig. 3).

The FTIR analysis of PGa21Ca in Fig. 3 shows vibrations characteristic of hydroxyl groups at a wavelength of 3400 cm\(^{-1}\), as well as vibrations in the range of 900 cm\(^{-1}\) corresponding to the Al-OH\(_2\) group. Standing out between the bands characteristic of aluminum sulfate are \( v_1 \) at around 990 cm\(^{-1}\), \( v_2 \) at around 1113–1089 cm\(^{-1}\) and \( v_3 \) at around 620–553 cm\(^{-1}\) [19].

The position of the 3392 and 3495 cm\(^{-1}\) bands in some gypsum infrared spectra may occur close to 3400 and 3540 cm\(^{-1}\) [20]. One of the four vibrational modes of SO\(_4^{2-}\) was observed in the spectrum of the PGa21Ca sample, in the \( v_2 \) SO\(_4\) stretching mode, which occurs at around 1120 cm\(^{-1}\) for gypsum [21]. The gyspum showed two bands at 1685 and 1620 cm\(^{-1}\), which were attributed to the presence of two types of water molecules. The band at 1685 cm\(^{-1}\) is characteristic of weak molecular bonds, while the band at 1620 cm\(^{-1}\) is characteristic of strong molecular bonds [21].

The PGa21Ca spectrum showed bands characteristic of gyspum at 3400 cm\(^{-1}\). The gyspum also presented three absorption bands in the stretching region of the OH group (\( \nu_{\text{OH}} \)) at 3245, 3392 and 3495 cm\(^{-1}\) [20].

The FTIR analysis also revealed vibration bands characteristic of \( \gamma \)-PGA at wavelengths of 1620, 1113 and 675 cm\(^{-1}\). Ho et al. [18] described the FTIR absorption for calcium \( \gamma \)-polyglutamate in KBr pellets as: Amide I, N–H bending band at 1622 cm\(^{-1}\); Amide II, stretching band not observed presumably due to strong ionic complexation; C=O symmetric stretching band at 1412 cm\(^{-1}\);
Fig. 2. SEM images of PGa21Ca (500, 1000, 2500 and 5000X magnification).

Fig. 3. Infrared spectra of PGa21Ca absorption bands in the range of 400–4000 cm⁻¹.
C—N stretching band at 1116 cm⁻¹; N—H bending band at 669 cm⁻¹; and O—H stretching band at 3415 cm⁻¹.

3.5. Thermal degradation of PGeo21Ca

PGA was analyzed by TGA/DTG and DSC thermoanalytical techniques in synthetic air and N₂ atmosphere at a heating rate of 20 °C min⁻¹ (Fig. 4). The TGA/DTG curves indicated that the sample was thermally stable up to 97.93 °C and decomposed in three main stages, between 152.4, 334.2 and 801.3 °C. The three well-defined thermal events in both synthetic air and nitrogen atmospheres were clearly represented by successive reactions involving thermal characteristics of calcium sulfate and aluminum sulfate. In the case of PGeo21Ca, the thermogravimetric analysis revealed a slight mass loss of 0.6192 mg, with a calculated residual mass of 89.10%, during the decomposition process. Çilgi and Cetisli [22] stated that the thermal decomposition of aluminum sulfate hydrate occurs in four major stages, all endothermic. The peak temperatures for the dehydration reaction occur in Stage I (51.85-80.85 °C) and II (80.85-184.85 °C). The aluminum sulfate decomposition step begins at 691 °C.

Gypsum showed two peaks in the range of 150–200 °C, representing the stepwise removal of water. At low concentrations, gypsum yielded a single peak at about 150–160 °C. Free water had an endothermic effect at 100 °C. The hexagonal phase exhibited an endothermic peak at about 160–175 °C, the cubic form of gypsum at 300–325 °C [23]. The DTG curve of PGeo21Ca showed well-defined thermal events characteristic of gypsum decomposition at 152.7 and 178.3 °C. The decomposition of gypsum occurs in two stages, with the formation of hemihydrate at around 130 °C and of anhydrite at around 170 °C.

\[
\text{CaSO}_4 \cdot 2\text{H}_2\text{O} \rightarrow \text{CaSO}_4 \cdot 1/2\text{H}_2\text{O} + 1.5\text{H}_2\text{O} \rightarrow 2\text{CaSO}_4 + \text{H}_2\text{O}
\]

Ho et al. [18] made a thermal analysis of calcium γ-polyglutamate and found 20% hydrated water, a dehydration temperature of 110 °C, and a decomposition temperature of 335.7 °C, but did not observe the melting point.

3.6. Coagulation-flocculation-sedimentation tests

One of the approaches of this study focused on coagulation-flocculation-sedimentation experiments of raw water samples from the Salto de Pirapora WTP (Latitude 23°39’22.55" S and Longitude 47°34’27.66" W), in the state of São Paulo, Brazil. The raw water analyses were performed no more than 24 h after collecting the samples. Sampling was carried out in 2015 during rainy and dry periods. A sample was frozen immediately after filtering to determine nitrogen and phosphorus compounds. Filtration was performed using a vacuum pump and hydrophilic membrane filters (Millipore) with 0.45 μm pores. Parameters such as alkalinity, anions, cations, and trace metals were also analyzed, and are described in Tables 4 and 5. The geology of the municipality of Salto de Pirapora favors mining of limestone rock belonging to the São Roque Group, in the form of limestone lenses, and dolomitic phyllite and calc-schist. This lithological substrate, allied to the region’s mining operations, leads to substantial calcium carbonate content in the raw water.

The commercial coagulants used in the coagulation-flocculation-sedimentation tests were aluminum sulfate (16% Al₂O₃) and PAC (9.59% Al₂O₃), as well as PGeo21Ca imported from Japan. The levels of coagulants were established based on preliminary tests and recommended by the WTP, but the parameters of pH and temperature of the raw water were not changed. The optimum operating conditions for bioflocculant treatment were determined by the jar test procedure and, in this case, the PGeo21Ca was used directly in its powder form for the jar test.

The experiments were performed on a laboratory scale (jar testing), whose main purpose was to vary the velocity gradient during the simultaneous stirring of six square 2-l acrylic test jars. In these jar test experiments, we applied rapid mixing times of 10 s, an average rapid mixing velocity gradient of 1000 s⁻¹, flocculation time of 30 min, and an average flocculation velocity gradient of 15 s⁻¹. Lastly, we studied the influence of the settling velocity of 1 cm min⁻¹ with simultaneous collection of samples from the sampling port located 7 cm below the 2 L marking on the six jars.
Table 4
Analytical results of raw water from the Salto de Pirapora WTP pertaining to the main dissolved chemical species (cations and anions), in mg L⁻¹.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Na⁺</th>
<th>Ca²⁺</th>
<th>K⁺</th>
<th>Mg²⁺</th>
<th>Al</th>
<th>Fe</th>
<th>Mn</th>
<th>Cl⁻</th>
<th>HCO₃⁻</th>
<th>NO₃⁻</th>
<th>SO₄²⁻</th>
<th>PO₄³⁻</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dry season</td>
<td>8.56</td>
<td>20.11</td>
<td>4.62</td>
<td>2.43</td>
<td>0.19</td>
<td>0.23</td>
<td>LQ</td>
<td>14.77</td>
<td>54.68</td>
<td>3.72</td>
<td>7.24</td>
<td>0.13</td>
</tr>
<tr>
<td>Rainy season</td>
<td>5.63</td>
<td>12.95</td>
<td>4.54</td>
<td>2.70</td>
<td>16.29</td>
<td>5.77</td>
<td>LQ</td>
<td>9.06</td>
<td>45.32</td>
<td>3.26</td>
<td>7.12</td>
<td>LOQ</td>
</tr>
</tbody>
</table>

Note: results below the limit of quantification of the method (LOQ=0.01 mg L⁻¹).

Table 5
Physicochemical parameters of raw water from the Salto de Pirapora WTP in the dry and rainy seasons.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Unit</th>
<th>Dry</th>
<th>Rainy</th>
</tr>
</thead>
<tbody>
<tr>
<td>pH</td>
<td>–</td>
<td>7.50</td>
<td>6.92</td>
</tr>
<tr>
<td>color</td>
<td>HU</td>
<td>720</td>
<td>750</td>
</tr>
<tr>
<td>turbidity</td>
<td>NTU</td>
<td>458</td>
<td>633</td>
</tr>
<tr>
<td>alkalinity</td>
<td>mg L⁻¹</td>
<td>42</td>
<td>43</td>
</tr>
<tr>
<td>EC</td>
<td>μS cm⁻¹</td>
<td>157.9</td>
<td>260.17</td>
</tr>
<tr>
<td>T</td>
<td>°C</td>
<td>25</td>
<td>24.8</td>
</tr>
</tbody>
</table>

Note: Hu, Hazen units; EC, electrical conductivity; NTU, nephelometric turbidity unit; T, temperature.

After the tests, the residual turbidity and percent removal of apparent color of the best settling results (1 cm min⁻¹) were calculated, as illustrated in Figs. 5–7. Fig. 8 shows the results of the tests performed on water collected during the rainy season, and in this case, using only the commercial product with the highest efficiency in the previous phase, i.e., PGo21Ca.

Fig. 5 shows the residual turbidity of 90.55% and the percent removal of apparent color of 9.67 NTU obtained with PAC, while Fig. 6 shows the residual turbidity of 85.49% and percent removal of apparent color of 12.61 NTU obtained with aluminum sulfate. The best values obtained with PAC were at the level of 90 mg L⁻¹. The use of PAC in place of aluminum sulfate reportedly offers economic and environmental advantages because it significantly reduces neutralizing agents such as hydrated lime and sodium carbonate. PAC also promotes coagulation in any pH range and can be applied at two to eight times lower levels than aluminum sulfate, generating less sludge, providing better water clarification, and facilitating filtration by reducing the filter cleaning frequency [24,25].

Fig. 7 illustrates the use of PGo21Ca at a level of 80 mg L⁻¹, with removal of apparent color and residual turbidity of 89.74% and 5.78 NTU, respectively. Note that PGo21Ca was effective even at a low level of 55 mg L⁻¹ (81.24% and 11.81 NTU), approaching or even surpassing the results of residual turbidity of the other coagulants applied at higher levels in this study.

According to Pan et al. [26], the flocculation efficiency of glutamic acid is dosage-dependent. Research has shown that, in kaolin suspensions, γ-PGA shows a flocculation rate of 95.8%, in the conditions of 300 mg L⁻¹ of PGA, pH 7, a temperature of 25 °C and a 350 rpm agitation rate. Carvajal-Zarrabal et al. [11], who studied the use of PGA in the treatment of tequila vinasse, a waste generated in ethanol distilleries, reported that the use of γ-PGA (250–300 mg L⁻¹) combined with sodium hypochlorite and sand filtration successfully removed about 70% of turbidity and reduced the chemical oxygen demand (COD) by 79.5%, with the additional benefit of color removal. The authors, who called PGo21Ca a harmless biodegradable biopolymer flocculant, report the effectiveness of the product, due to the high turbidity removal rate even at the natural pH of the vinasse (3.5), highlighting its remarkable effect in removing turbidity.

Fig. 8 shows variations in residual turbidity of 14.67–198 NTU and apparent color removal rates of 86.65–96.95% at levels of 60–100 mg L⁻¹. As can be seen, at levels starting from 70 mg L⁻¹, turbidity was reduced to below 5 NTU, the maximum value permitted for drinking water in Brazil [27], while a level of 90 mg L⁻¹ resulted in residual turbidity of 1.93 NTU (removal of 99.74%) and apparent color of 96.89%. The results indicate the satisfactory performance of PGo21Ca in the removal of apparent color and residual turbidity in the treatment of raw water in the rainy season, demonstrating flexibility when varying the water quality parameters.

The chemical analysis of the composition of PGo21Ca indicated the presence of aluminum sulfate and γ-PGA. The former is responsible for agglomerating suspended or colloidal impurities and the latter for forming bonds between the colloids, leading to the formation of denser particles and heavier flakes. According to Huang [28], chemical coagulants such as aluminum salts, iron salts, polyaluminum chloride, polyferric chloride and polyacrylamide are still widely applied in the treatment of drinking water due to their outstanding coagulation performance and low cost [29]. However, these common coagulants still induce several health-related and environmental problems. The potential neurotoxic hazard posed by aluminum is generally accepted [30], as is the genetic toxicity of acrylamide monomer [31].

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**Fig. 5.** Residual turbidity and apparent color removal rate of water treated with PAC (Vs 1 cm min⁻¹) in the dry season.

**Fig. 6.** Residual turbidity and apparent color removal rate of water treated with aluminum sulfate (Vs 1 cm min⁻¹) in the dry season.
The excellent performance PGo21Ca in the removal of turbidity and apparent color is associated with the functions of its components, i.e., the inorganic fraction neutralizes colloidal charges owing to the presence of Ca$^{2+}$ and Al$^{3+}$ cations [32], and γ-PGA acts as a chelator, forming bridges between the small particles. Table 1 shows the presence of aluminum in the formulation of PGo21Ca. However, the ability to form residual aluminum in water treated with PGo21Ca can be considered low, since γ-PGA can bind to multiple metal ions, such as Al$^{3+}$ [32].

4. Conclusion

Conventional water treatment tests were carried out using the jar test with six square beakers (2 L), which provided velocity gradients in the range of 10–1200 s$^{-1}$. PAC showed better performance than aluminum sulfate in the removal of apparent color and turbidity from raw water from Salto de Pirapora. However, PGo21Ca, whose formulation contains an inorganic portion and another of γ-PGA, showed better results in the removal of apparent color and turbidity from raw water than the other coagulants, and also caused little change in the water pH level.

At 90 mg L$^{-1}$, PAC presented an apparent color removal rate of 90.55% and residual turbidity of 9.67. In contrast, at an initial level of 90 mg L$^{-1}$, aluminum sulfate removed 85.49% of apparent color while residual turbidity was 12.61 NTU. In the dry season, 80 mg L$^{-1}$ of PGo21Ca removed 89.74% of apparent color and residual turbidity was 5.78 NTU. In the rainy season, at 90 mg L$^{-1}$, PGo21Ca removed 96.89% of apparent color and residual turbidity was 1.93 NTU.

PGo21Ca is basically composed of calcium sulfate (87%), γ-polyglutamic acid (5.26%), and other chemical compounds. The chemical analysis of PGo21Ca indicated that the product is essentially inorganic. In fact, PGo21Ca can be considered a hybrid coagulant that combines aluminum sulfate and γ-PGA. In this case, we understand that the presence of gypsum and anhydrite as excipients is justified since they facilitate dosing and also contribute to the formation of a water-insoluble compound, because γ-polyglutamic acid can react with calcium, i.e., the solubility of CaSO$_4$ in water is 2 g L$^{-1}$, thus enabling it to react with γ-PGA. In addition to containing calcium sulfate as excipient, the formulation of PGo21Ca includes a conventional coagulant, aluminum sulfate, calcium carbonate as the alkalinizing agent, and γ-polyglutamic acid as an adjuvant compound, i.e., an anionic polymer that is able to form bridges between coagulated particles and thus create larger flakes.

In the process of coagulation-flocculation and separation of solids from the liquid phase by settling, PGo21Ca showed significant results in the removal of apparent color and in the residual turbidity of water samples collected in the dry and rainy seasons, especially with respect to the aluminum sulfate coagulants. However, it should be kept in mind that this product is inorganic and may produce large quantities of raw sludge, which would have to be managed.

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