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## Color stability of blue aluminates obtained from recycling and applied as pigments

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Aluminates have been used as synthetic inorganic pigments due to their structural stability. In this context, we report on the synthesis of blue aluminates obtained from the boehmite combination with  $\text{Co}^{2+}$  and  $\text{Ni}^{2+}$  ions and calcination at 1000 °C. The boehmite was obtained from recycling aluminum metal from can seals. The aluminates were characterized by X-ray diffractometry (XRD) and Raman spectroscopy. The oxidation state of the ions responsible for the color was evaluated by XPS and UV-vis, suggesting the presence of divalent Co and Ni ions. Colorimetry confirmed the blue coloration and different shades. The colorimetric stability of the blue aluminates in harsh environments for pigments was evaluated. Both pigments were stable after 240 hours of exposure, showing no strong color difference.

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### Sustainability spotlight

This work reports on the production of synthetic inorganic pigments from recycling aluminum can seals. In this way, it contributes to the circular economy because the can seals leave the beverage industry and become part of the pigment industry (goal 9). In addition, the development of our research has the potential to generate jobs (recyclable waste pickers) meeting the UN goals 1 and 8 of sustainable development, to preserve natural resources (goals 12 and 13), and to promote partnerships (goal 17), as it involved educational institutions in Brazil and Belgium. Therefore, the subject of our study aims at promoting the sustainable development.

## 1. Introduction

The circular economy aims at harmonizing the market, society and the environment, with an ecologically correct vision. For this purpose, it encourages the use of alternative secondary resources such as a recycled material to obtain products.<sup>1</sup> Within this context, a fully recyclable material is the aluminum can; for each kilo of secondary (recycled) aluminum, about 5 kilos of bauxite are saved. Furthermore, for a ton of aluminum recycled, 95% of the energy to produce the same amount of primary aluminum is saved.<sup>1</sup>

Synthetic inorganic pigments have applications in different areas due to their high hiding power, color stability, weather resistance, and surface protection.<sup>2</sup> The doping of metallic oxides with coloring ions has been the most used strategy for their synthesis.<sup>3,4</sup> Both structural and physicochemical properties of the pigments depend on the precursors and synthetic routes used.<sup>5</sup> Some metal oxides widely used as a matrix for pigments are titanium, aluminum, zirconium, and silicon oxide.<sup>6</sup> The attribution of color to these matrices occurs with the

insertion of chromophore ions; transition metals such as V, Cr, Mn, Fe, Co, Ni, and Cu are commonly used for this purpose.<sup>7</sup> In general, the use of inorganic compounds tends to increase the chemical and physical stability of the pigments,<sup>8</sup> ensuring greater durability.

The cobalt blue pigment with spinel structure ( $\text{CoAl}_2\text{O}_4$ ) is known to be an environmentally friendly blue pigment.<sup>9</sup> In addition, this pigment can be applied in various areas, such as ceramic pigment,<sup>10</sup> anti-corrosion inks,<sup>11</sup> in printing ink,<sup>12</sup> and as a near-infrared reflection pigment.<sup>13</sup> Another way to produce blue colored pigment is to use the transition ion  $\text{Co}^{2+}$  to  $\text{Ni}^{2+}$ , resulting in different shades of blue. However, it is necessary to find a stable structure for these ions.<sup>14</sup> In this context, there is the possibility of preparing aluminates.

Aluminates have been widely used as dyes and pigments due to their structural arrangement in the spinel form ( $\text{A}^{2+}\text{B}_2\text{O}_4$ ) that allows the insertion of chromophore ions, which guarantee intense color and high thermal stability, in addition to resistance to chemically aggressive environments.<sup>15,16</sup> The synthetic routes most used to obtain aluminates are coprecipitation,<sup>17</sup> mixed oxide calcination,<sup>18</sup> solid-state reactions,<sup>19</sup> and the sol-gel method.<sup>20</sup> In addition, there is the possibility of recycling aluminum to be used as a pigment precursor,<sup>17</sup> providing a more sustainable synthesis compatible with the circular economy concept, which is based on reducing, repairing, recycling, remanufacturing, and redirecting the materials life

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cycle,<sup>21,22</sup> contributing to the production of more sustainable materials.

Within the context of circular economy, the objective of this study is to evaluate the color stability in environments containing hydrochloric acid (HCl) and sodium hydroxide (NaOH) of blue aluminates obtained from recycled aluminum for their application as synthetic inorganic pigments.

## 2. Material and methods

### 2.1 Synthesis of boehmite: metallic aluminum acid digestion

Aluminum can seals were recycled through acid digestion using hydrochloric acid (HCl) at a concentration of 1.1 mol L<sup>-1</sup> and a ratio of 1 g/100 mL (Fig. 1). The reaction lasted about 24 hours, which is the time necessary for the stabilization of the reaction medium. The solution containing the Al<sup>3+</sup> ions had an acid pH of 0.35<sup>17</sup>.

### 2.2 Obtaining boehmite (γ-AlOOH)

To obtain the boehmite phase (Fig. 1), the pH of the solution containing Al<sup>3+</sup> ions was modified through the addition of sodium hydroxide (NaOH) by dripping until pH 8, referring to the formation of the phase of interest. After precipitation, the oxide-hydroxide was vacuum filtered and oven dried at 70 °C.<sup>17,23</sup>

### 2.3 Boehmite purification

For purification (Fig. 1), the obtained product was washed in hot water to solubilize and remove sodium chloride (secondary product of the reaction). After this step, the boehmite was filtered and dried at 70 °C.<sup>17,23</sup> To obtain the white sample, the boehmite was calcined at 1000 °C.

### 2.4 Aluminates synthesis

Transition metal chlorides (coloring ion): CoCl<sub>2</sub> and NiCl<sub>2</sub> were used to obtain blue aluminates (Fig. 2). First, the transition metal chloride (10% in relation to the total mass of boehmite powder (m m<sup>-1</sup>)) is added to 50 mL of water. Then, 3 g of boehmite is added and left under constant stirring (600 rpm) for 24 hours. After this period, the mixture is calcined at 1000 °C, macerated, and stored. The obtained materials were labeled CoAl<sub>2</sub>O<sub>4</sub> and NiAl<sub>2</sub>O<sub>4</sub>, respectively.



Fig. 1 Scheme of the synthetic route used to obtain boehmite. (Adapted from ref. 23).

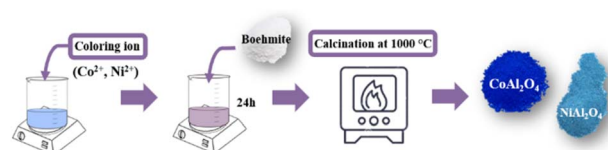


Fig. 2 Scheme of the synthetic route used to obtain aluminates.

### 2.5 Application as pigments

For application as a pigment, the aluminate samples were dispersed in white paint (10% aluminate, 90% white paint) prepared in the proportion of 2 : 1 paint : water.<sup>17</sup> The mixture was applied on plaster substrates and painted with two paint coats, according to the manufacturer's instructions.

### 2.6 Color stability

The color stability test was carried out in acidic and alkaline environments, using desiccators as a controlled experimental environment. Separately, Petri dishes with hydrochloric acid (HCl) or sodium hydroxide (NaOH) were placed in different desiccators in order to mimic aggressive environments with vapors.<sup>24</sup> The coated substrates, the plaster blocks, were added to the desiccators; further, these were sealed. Colorimetric measurements were performed before the test and after 120 and 240 hours in order to verify the colorimetric stability of the aluminates applied as synthetic inorganic pigments.

### 2.7 Characterization

The samples were characterized by X-ray diffraction (XRD), carried out in a Bruker D2 Phaser diffractometer, which uses a copper cathode with copper K $\alpha$  emission ( $\lambda = 1.5418$  Å) and equipped with a LynxEye high-performance detector, with a power of 300 W. A scanning electron microscope Hitachi SU8020 SEM was used to obtain morphology information. The oxidation state and chemical composition of the aluminates were evaluated by X-ray photoelectron spectroscopy (XPS) (Versaprobe PHI 5000, from Physical Electronics, Chanhassen, MN, USA), equipped with a monochromatic Al K $\alpha$  X-ray source. The spectra were analyzed using the CASA-XPS software, and the binding energy of the XPS spectra was calibrated using the C 1s peak at 284.6 eV.<sup>25</sup> Multipack version 9.8 software (ULVAC-PHI, 2017, Chigasaki, Japan) was used to evaluate the relative composition of the elements. Raman spectra were recorded using a micro-Raman system (Senterra Bruker Optik GmbH, Massachusetts, USA),  $\lambda = 532$  nm, laser power 10 mW. The visible spectra were obtained using the Ocean Optics spectrophotometer (USB 2000), equipped with optical fiber, tungsten-halogen source, and silicon (350–720 nm) and germanium (720–1050 nm) detectors.

The colorimetric analysis was performed on the pigments in the form of powder, and after application on the plaster substrate, a portable colorimeter (3nh, model NR60CP) with a D65 light source. The data representative of colorimetric (CIE  $L^*a^*b^*$ ) analyses are: the  $L^*$  parameter is the brightness and ranges from 0 to 100. The parameter  $a^*$  represents the variation between red and green, where  $+a$  tends to red and  $-a$



tends to green. The parameter  $b^*$  is the variation between blue and yellow, where  $+b^*$  tends to yellow and  $-b^*$  tends to blue.<sup>17</sup> The  $h^*$  parameter refers to the hue angle ( $\tan^{-1} b^*/a^*$ ).<sup>26</sup> The value of  $C^*$  ( $\sqrt{a^{*2} + b^{*2}}$ ) is the chroma and represents the color saturation of the samples.<sup>26</sup> The total color deviation is expressed by the value of  $\Delta E$ , which was created by the CIE to designate a metric distance and can be calculated using equation  $\Delta E = \sqrt{(\Delta L^*)^2 + (\Delta a^*)^2 + (\Delta b^*)^2}$ .

### 3. Results and discussion

#### 3.1 X-ray diffractometry (XRD)

The obtention boehmite phase was confirmed by the crystallographic chart [96-901-22-49] (Fig. 3a). The pristine  $\text{Al}_2\text{O}_3$  sample (Fig. 3b) shows the crystallographic phase  $\theta - \text{Al}_2\text{O}_3$ , this phase is obtained when boehmite is calcined above 900 °C.<sup>27</sup> The XRD pattern of the  $\text{CoAl}_2\text{O}_4$  sample (Fig. 3c), suggests the formation of the cobalt aluminate spinel [96-900-5204] crystallographic phase. This phase presents a cubic arrangement,<sup>28</sup> where the divalent cations ( $\text{Co}^{2+}$ ) occupy 1/8 of the tetrahedral sites of this structure.<sup>15</sup> Whereas in the XRD of the  $\text{NiAl}_2\text{O}_4$  sample (Fig. 3d), only the crystallographic phase of nickel aluminate [96-900-1433] was observed.<sup>29</sup> The homogeneous coloration observed in the samples is associated with the formation of monophasic aluminates since there is no presence of possible interferences such as  $\text{Al}_2\text{O}_3$ .<sup>15</sup> The aluminates obtention samples caused a decrease in crystallinity (Table 1) compared to the  $\text{Al}_2\text{O}_3$

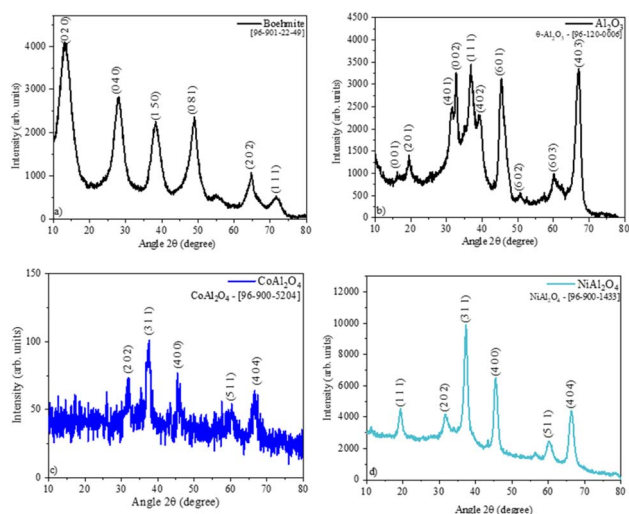


Fig. 3 Sample diffractograms: (a) boehmite, (b)  $\text{Al}_2\text{O}_3$ , (c)  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate), and (d)  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).

Table 1 Phase, crystallographic chart, crystallinity of the samples

| Sample                    | Phase                              | Chart         | Crystallinity (%) |
|---------------------------|------------------------------------|---------------|-------------------|
| $\text{Al}_2\text{O}_3$   | $\theta - \text{Al}_2\text{O}_3$   | [96-120-0006] | 63.9              |
| $\text{CoAl}_2\text{O}_4$ | Spinel – $\text{CoAl}_2\text{O}_4$ | [96-900-5204] | 30.8              |
| $\text{NiAl}_2\text{O}_4$ | Spinel – $\text{NiAl}_2\text{O}_4$ | [96-900-1433] | 58.7              |

sample due to the difference in the ionic radius of these metals that can cause deformations in the crystal lattice.<sup>30</sup>

#### 3.2 Raman spectroscopy

Raman spectroscopy allows the characterization of both crystalline as well as low-crystallinity materials such as  $\text{CoAl}_2\text{O}_4$ , therefore Raman spectroscopy was used to verify the suggested structures by XRD. The Raman spectra (Fig. 4) show bands at low frequencies (centered at 250  $\text{cm}^{-1}$  for  $\text{Al}_2\text{O}_3$  (Fig. 4a), 198  $\text{cm}^{-1}$  for  $\text{CoAl}_2\text{O}_4$  (Fig. 4b), 373  $\text{cm}^{-1}$  for  $\text{NiAl}_2\text{O}_4$  (Fig. 4c)) that were attributed to the Al–O octahedral vibration,<sup>31,32</sup> as well as a low-intensity band near 500  $\text{cm}^{-1}$  that was attributed to metal–oxygen bonding, specifically to the symmetrical Al–O stretching related to octahedral groups,<sup>17,33</sup> verifying the X-ray diffractometry results. Bands near 400  $\text{cm}^{-1}$  (Fig. 4b and c) refer to Al–O vibrations, indicating greater organization in normal spinel-like structures.<sup>15</sup> In the  $\text{NiAl}_2\text{O}_4$  sample, the signal extending from ~470 to 650 was associated with the normal cubic spinel  $Fd3m$  of  $\text{NiAl}_2\text{O}_4$ .<sup>34</sup> The low-intensity band centered at 780  $\text{cm}^{-1}$  can be associated with metal–Ni–oxygen bonds.<sup>17,35</sup> The effect of tetrahedral cation substitution on the Raman bands can be evaluated, through the analyses of the internal vibrations of the  $\text{AlO}_6$  octahedron and/or  $\text{A}^{2+}\text{O}_4$  tetrahedron. The Raman shift of the  $\text{F}_{2g}$  mode varies linearly with the radius of the tetrahedral cation, being 0.65 Å for cobalt and 0.69 Å for nickel. According to the literature, the  $\text{F}_{2g}$  mode is assigned to  $\text{AO}_4$  within the spinel structure. The  $\text{A}_{1g}$  mode Raman shift varies significantly less than in the other modes. This Raman mode reveals a minor dependence on the type of divalent cation, although it has been attributed to the  $\text{A}^{2+}\text{O}$  stretching vibration of the  $\text{AO}_4$  groups.<sup>36,37</sup>

#### 3.3 Scanning electron microscopy (SEM)

The  $\text{Al}_2\text{O}_3$  sample (Fig. 5a) is characterized by large particles with a rough and irregular surface, with an average particle size equal to 8.2  $\mu\text{m}$ . The SEM image of the cobalt aluminate sample (Fig. 5c) shows large particles with an average size value of 7.6  $\mu\text{m}$ . It can be observed that smaller particles (with a size of

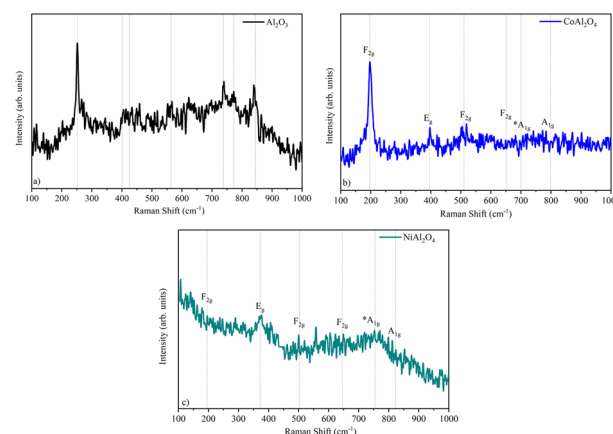


Fig. 4 Raman spectroscopy of samples: (a)  $\text{Al}_2\text{O}_3$  (alumina), (b)  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate), and (c)  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).

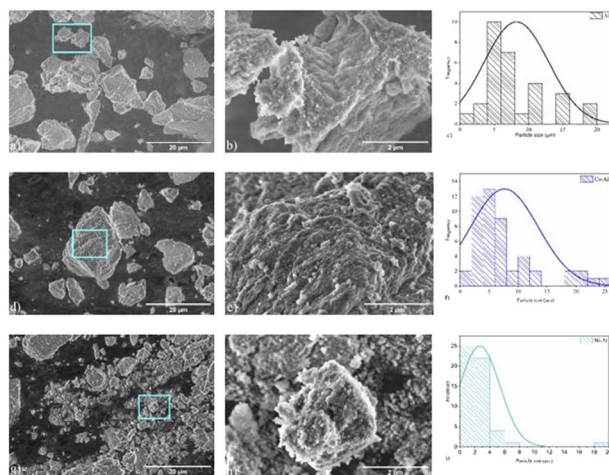


Fig. 5 Scanning electron microscopy images of samples: (a)  $\text{Al}_2\text{O}_3$ , (b)  $\text{Al}_2\text{O}_3$ , (c) histogram of  $\text{Al}_2\text{O}_3$  particle size distribution, (d)  $\text{CoAl}_2\text{O}_4$ , (e)  $\text{CoAl}_2\text{O}_4$ , (f) histogram of  $\text{CoAl}_2\text{O}_4$  particle size distribution, (g)  $\text{NiAl}_2\text{O}_4$ , (h)  $\text{NiAl}_2\text{O}_4$ , and (i) histogram of  $\text{NiAl}_2\text{O}_4$  particle size distribution. In the high magnification SEM images, we can observe the presence of grooves all over the surface of the particles.

approximately  $0.18 \mu\text{m}$ ) are distributed over the surface (Fig. 5d). For the nickel aluminate sample, the observed average particle size was  $\sim 2.6 \mu\text{m}$ . The difference in particle size among the samples can be attributed to structural disorder, resulting from different ionic radii of the cations and particle clustering.<sup>38</sup>

### 3.4 X-ray photoelectron spectroscopy (XPS)

The  $\text{Al}_2\text{O}_3$ ,  $\text{CoAl}_2\text{O}_4$ , and  $\text{NiAl}_2\text{O}_4$  XPS survey spectra are presented in Fig. 6, the chemical elements found are labeled, and the elemental quantification is presented in Table 2. The XPS spectra recorded in the Al 2p region (Fig. 7a, b and d) show a peak fitted with two components centered at 73.6 eV and 74.0 of binding energy, generated by photoelectrons emitted from

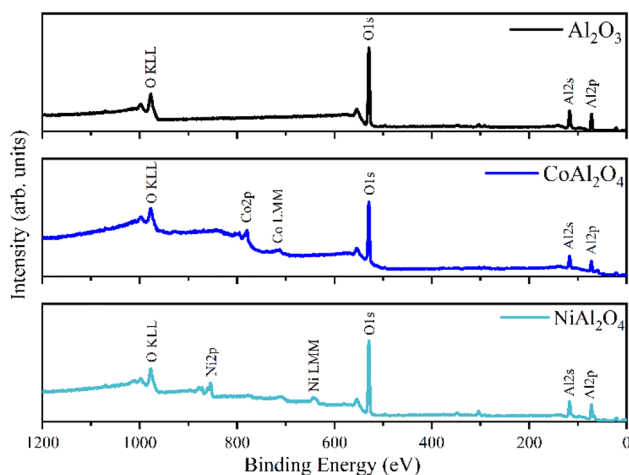


Fig. 6 XPS survey spectra of samples: (a)  $\text{Al}_2\text{O}_3$  (alumina), (b)  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate) and (c)  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).

Table 2 Aluminates composition obtained from XPS spectra

| Sample                    | Atomic (%) |      |     |     |
|---------------------------|------------|------|-----|-----|
|                           | Al         | O    | Co  | Ni  |
| $\text{Al}_2\text{O}_3$   | 32.6       | 67.4 | —   | —   |
| $\text{CoAl}_2\text{O}_4$ | 31.8       | 64.2 | 3.9 | —   |
| $\text{NiAl}_2\text{O}_4$ | 28.4       | 66.7 | —   | 4.9 |

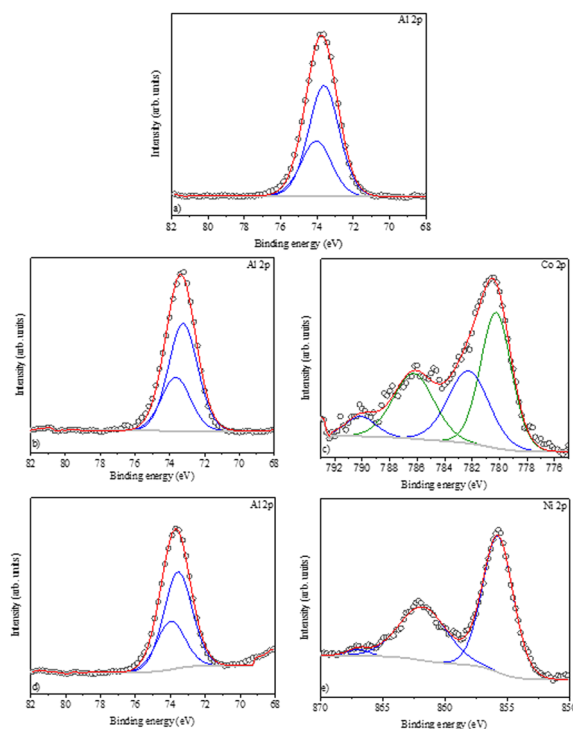


Fig. 7 XPS spectra: (a) Al2p –  $\text{Al}_2\text{O}_3$  (alumina), (b) Al2p –  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate), (c) Co2p –  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate), (d) Al2p –  $\text{NiAl}_2\text{O}_4$  (nickel aluminate), and (e) Ni2p –  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).

the Al  $2p_{3/2}$  and Al  $2p_{1/2}$  electronic levels respectively. These indicate the presence of  $\text{Al}_2\text{O}_3$  in all samples.<sup>35,40</sup>

The Co 2p spectrum (Fig. 7c) shows a peak centered at 780.7 eV (Co  $2p_{3/2}$ ) with a wide satellite at 786.2 eV, and the main peak can be reproduced by two components characteristic of cobalt aluminate, corroborating with the XRD data and confirming the presence of  $\text{Co}^{2+}$ ,<sup>41,42</sup> which is typical of high-spin divalent cobalt.<sup>42</sup> Furthermore, the high-resolution Ni 2p spectrum (Fig. 7e) is characteristic of  $\text{Ni}^{2+}$  with the Ni  $2p_{3/2}$  component centered at 855.8 eV and a satellite at 862.0 eV.<sup>43</sup> The distance between the main peak and its satellite is approximately 6.0 eV, which coincides with the reference values for nickel aluminates.<sup>39,44</sup>

### 3.5 UV-vis absorbance

The  $\text{Al}_2\text{O}_3$  sample (Fig. 8a) is a white powder and therefore shows no bands in the visible region.<sup>17</sup> The triplet band observed in the  $\text{CoAl}_2\text{O}_4$  sample (Fig. 8b) centered at  $\sim 540$ , 580, and 640 nm was reported to be related to the  $^4\text{A}_2(\text{F}) \rightarrow ^4\text{T}_1(\text{P})$

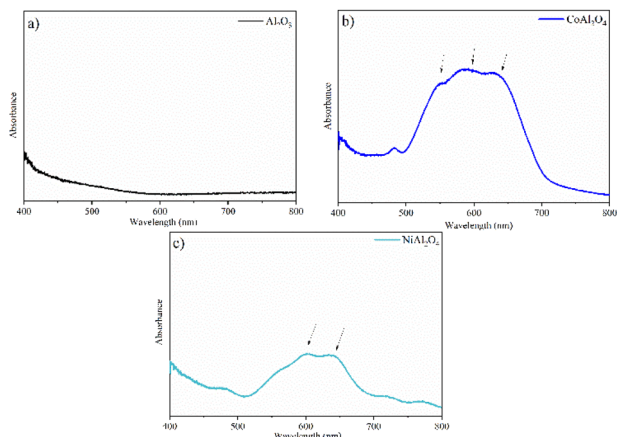


Fig. 8 Absorption spectra in the visible region: (a)  $\text{Al}_2\text{O}_3$  (alumina), (b)  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate) and (c)  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).

transition of cobalt ions in a tetrahedral arrangement, confirming the obtention of  $\text{CoAl}_2\text{O}_4$ .<sup>45</sup> The bands observed at approximately 590 and 640 nm in the  $\text{NiAl}_2\text{O}_4$  sample (Fig. 8c) were attributed to the  ${}^3\text{T}_1(\text{F}) \rightarrow {}^3\text{T}_1(\text{P})$  transitions, corresponding to tetrahedral nickel,<sup>46</sup> characteristic of the spinel phase of this aluminate, verifying the XRD results.

Table 3 Aluminates colorimetric parameters

| Sample                    | Colorimetric parameters |       |       |       |       |            | Photo |
|---------------------------|-------------------------|-------|-------|-------|-------|------------|-------|
|                           | $L^*$                   | $a^*$ | $b^*$ | $C^*$ | $h^*$ | $\Delta E$ |       |
| $\text{Al}_2\text{O}_3$   | 79.8                    | 0.5   | 9.4   | 9.4   | 87.2  | —          |       |
| $\text{CoAl}_2\text{O}_4$ | 13.0                    | 11.8  | -43.4 | 45.0  | 285.3 | 85.9       |       |
| $\text{NiAl}_2\text{O}_4$ | 42.5                    | -14.4 | -8.7  | 16.8  | 211.3 | 44.0       |       |

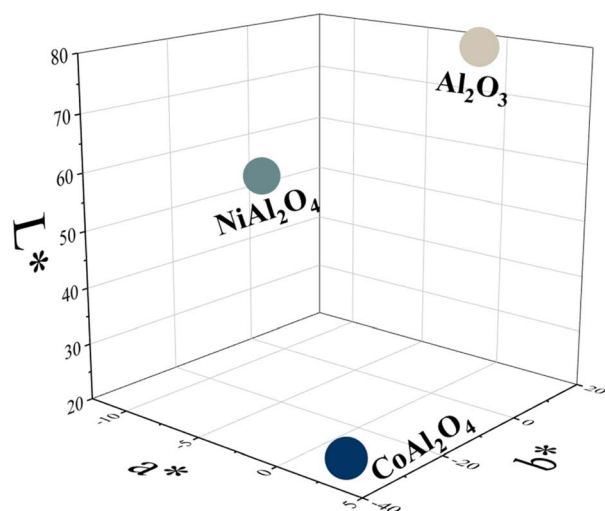


Fig. 9 Distribution of  $\text{Al}_2\text{O}_3$  (alumina),  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate) and  $\text{NiAl}_2\text{O}_4$  (nickel aluminate) in 3D CIE  $L^*a^*b^*$  color space.

### 3.6 Colorimetry

Regarding the colorimetric parameters (Table 3), the most luminous sample was the pristine  $\text{Al}_2\text{O}_3$  (79.75); this was also the sample with the lowest chromaticity ( $C^*$ ). The highest chromaticity was observed in  $\text{CoAl}_2\text{O}_4$ , and it presented the lowest luminosity (21.00). The addition of dopants made the color difference very strong in the samples compared to the  $\text{Al}_2\text{O}_3$  sample, varying the chromatic parameters and,

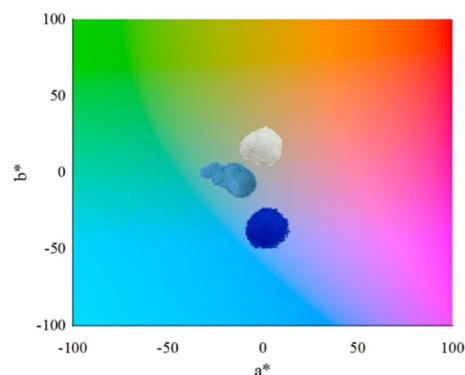


Fig. 10  $\text{Al}_2\text{O}_3$  (alumina),  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate) and  $\text{NiAl}_2\text{O}_4$  (nickel aluminate) distribution according to color quadrant.

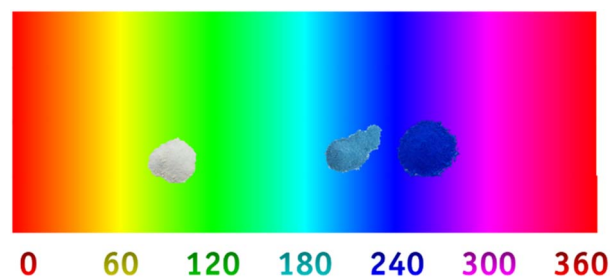


Fig. 11  $\text{Al}_2\text{O}_3$  (alumina),  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate) and  $\text{NiAl}_2\text{O}_4$  (nickel aluminate) distribution according to hue color.

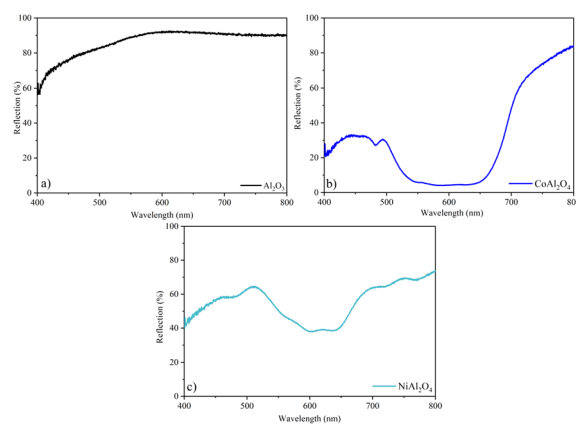


Fig. 12 Reflection spectra in the visible region: (a)  $\text{Al}_2\text{O}_3$  (alumina), (b)  $\text{CoAl}_2\text{O}_4$  (cobalt aluminate), and (c)  $\text{NiAl}_2\text{O}_4$  (nickel aluminate).



Table 4 Aluminates colorimetric parameters after application

| Sample                           | Colorimetric parameters |       |       |       |       |            | Photo                                                                             |
|----------------------------------|-------------------------|-------|-------|-------|-------|------------|-----------------------------------------------------------------------------------|
|                                  | $L^*$                   | $a^*$ | $b^*$ | $C^*$ | $h^*$ | $\Delta E$ |                                                                                   |
| White paint                      | 95.8                    | −0.5  | 1.5   | 1.6   | 109.2 | —          |  |
| Al <sub>2</sub> O <sub>3</sub>   | 93.7                    | −0.5  | 2.4   | 2.5   | 101.4 | 2.3        |  |
| CoAl <sub>2</sub> O <sub>4</sub> | 85.7                    | −3.4  | −8.3  | 8.9   | 247.9 | 14.8       |  |
| NiAl <sub>2</sub> O <sub>4</sub> | 89.5                    | −3.3  | −0.4  | 3.3   | 186.3 | 7.1        |  |

consequently, modifying the color,<sup>17</sup> as observed in the distribution of aluminates within the 3D color space (Fig. 9). The CoAl<sub>2</sub>O<sub>4</sub> sample is in the red/blue (+ $a$ /− $b$ ) quadrant, and NiAl<sub>2</sub>O<sub>4</sub> in green/blue (− $a$ /− $b$ ) (Fig. 10), corroborating the hue values ( $h^*$ ) (285.25 and 211.33) observed.<sup>47</sup> The hue refers to the pure color of the material, and its tonality, without considering white and black.<sup>47</sup> Aluminates are present in different shades of blue of the hue, as shown in Fig. 11, demonstrating the

possibility of obtaining pigments in different colors through simple synthesis and recycling.

### 3.7 UV-vis reflectance

The Al<sub>2</sub>O<sub>3</sub> sample (Fig. 12a) showed high reflectance, confirming the high luminosity observed in the colorimetric analysis. Furthermore, the yellowish hue of the Al pigment was confirmed by the broadband from ~500 nm. The bluish hue of CoAl<sub>2</sub>O<sub>4</sub> is confirmed by the broadband in the blue region of the visible spectrum (Fig. 12b). In the NiAl<sub>2</sub>O<sub>4</sub> sample, the shift towards red in this band (Fig. 12c) justifies its blue-green coloration, with negative colorimetric parameters  $a^*$  and  $b^*$ .

### 3.8 Pigments application

**3.8.1 Colorimetry.** The colorimetry performed after dispersion in paint and application in plaster presented the colorimetric parameters described in Table 4, and all samples obtained luminosity above 85; therefore, they are all luminous samples. The chromatic parameters ( $a^*$  and  $b^*$ ) decreased in relation to the colorimetry performed in the powder form, as well as the chromaticity ( $C^*$ ), due to the presence of the white matrix in the commercial paint.<sup>17</sup> The hue values ( $h^*$ ) decreased, however, the samples remained in the same hue color. Regarding the white paint, the colored samples showed a very strong total color difference ( $\Delta E$ ), while the white Al sample showed a clear difference.

**3.8.2 Color stability.** A comparative color study ( $\Delta E$ ) (Table 5) of the applied pigments was performed before, after 120, and

Table 5 Colorimetric parameters of aluminates after 120 and 240 hours in an acid and alkaline environment

| Environment | Sample                                   | Colorimetric parameters |       |       |       |        |            | Photo                                                                                 |
|-------------|------------------------------------------|-------------------------|-------|-------|-------|--------|------------|---------------------------------------------------------------------------------------|
|             |                                          | $L^*$                   | $a^*$ | $b^*$ | $C^*$ | $h^*$  | $\Delta E$ |                                                                                       |
| Acid        | Al <sub>2</sub> O <sub>3</sub> – 120 h   | 93.3                    | −0.5  | 2.2   | 2.20  | 103.1  | 0.5        |  |
|             | Al <sub>2</sub> O <sub>3</sub> – 240 h   | 94.3                    | −0.4  | 2.1   | 2.11  | 100.6  | 0.7        |  |
|             | CoAl <sub>2</sub> O <sub>4</sub> – 120 h | 85.6                    | −3.4  | −8.4  | 9.09  | 247.9  | 0.2        |  |
|             | CoAl <sub>2</sub> O <sub>4</sub> – 240 h | 87.2                    | −3.4  | −8.4  | 9.04  | 247.7  | 1.5        |  |
|             | NiAl <sub>2</sub> O <sub>4</sub> – 120 h | 91.1                    | −3.4  | −0.4  | 3.38  | 187.0  | 1.6        |  |
|             | NiAl <sub>2</sub> O <sub>4</sub> – 240 h | 92.2                    | −3.5  | −0.3  | 3.48  | 184.4  | 2.7        |  |
| Alkali      | Al <sub>2</sub> O <sub>3</sub> – 120 h   | 93.6                    | −0.4  | 2.4   | 2.38  | 99.2   | 0.2        |  |
|             | Al <sub>2</sub> O <sub>3</sub> – 240 h   | 94.4                    | −0.3  | 2.2   | 2.20  | 98.2   | 0.7        |  |
|             | CoAl <sub>2</sub> O <sub>4</sub> – 120 h | 86.2                    | −3.3  | −8.5  | 9.09  | 248.7  | 0.5        |  |
|             | CoAl <sub>2</sub> O <sub>4</sub> – 240 h | 86.6                    | −3.4  | −8.7  | 9.34  | 248.68 | 1.0        |  |
|             | NiAl <sub>2</sub> O <sub>4</sub> – 120 h | 91.2                    | −3.1  | −0.2  | 3.08  | 182.9  | 1.7        |  |
|             | NiAl <sub>2</sub> O <sub>4</sub> – 240 h | 86.5                    | −0.6  | 2.3   | 2.35  | 103.9  | 4.8        |  |



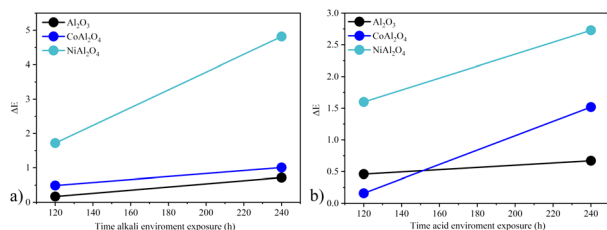


Fig. 13 Color stability  $\Delta E$  vs. sample exposure time in the different environments studied: (a) acid environment and (b) alkali environment.

240 hours of exposure to acid and alkaline environments. In general, it was observed that the color difference increased after 240 hours of exposure. However, none of the samples showed a very strong color difference under any conditions studied, demonstrating colorimetric stability.<sup>17</sup> The white sample  $\text{Al}_2\text{O}_3$  at 120 hours showed greater stability in the alkaline environment. However, the color variation increased by 0.55 after 240 hours, while in an acid environment, it varied by only 0.21, indicating greater stability after a longer period of exposure. For the  $\text{CoAl}_2\text{O}_4$  sample, the highest stability was observed in an alkaline environment, varying by only 0.52. In both environments, the observed color variation can be considered weak.<sup>17</sup> Whereas for the  $\text{NiAl}_2\text{O}_4$  sample, the acidic environment was more stable and had a variation of 1.13. The biggest difference was observed in the  $\text{NiAl}_2\text{O}_4$  sample after 240 h in alkali environment with  $\Delta E = 4.8$  and large variation in hue (from 182.9 to 103.9). In general, the color of the pigments was not drastically affected, demonstrating the good color stability of these compounds. This was due to the structural arrangement of aluminates in the spinel form, which is stable even in harsh environments.<sup>15,48</sup> Fig. 13 shows the graph  $\Delta E$  vs. exposure time of the samples in both environments studied, and it is possible to see that the least stable coloration in both environments was the one obtained with the  $\text{NiAl}_2\text{O}_4$  sample, indicating lower stability compared to cobalt aluminate. This may be due to the lower stability of the normal nickel spinel, which is more stable in the inverse spinel form,<sup>49</sup> or the chemical nature of this compound.

## 4. Conclusions

X-ray diffractometry data indicated that boehmite was successfully formed, and the addition of coloring ions followed by calcination at 1000 °C generated cobalt and nickel aluminates. The oxidation state (2+) of the ions responsible for the coloration was confirmed by UV-vis and XPS. The colorimetry indicates that through a synthesis strategy using recycling as a first step, blue aluminates can be obtained. Their color is distributed in different color quadrants, with  $\text{CoAl}_2\text{O}_4$  in the blue/red quadrant and  $\text{NiAl}_2\text{O}_4$  in the blue/green quadrant, as they have different shades of hue. The pigments show color stability in acid and alkaline environments, with minor color variation in the time range studied; this characteristic can be associated with the stability of spinel-like structures, which was confirmed by Raman spectroscopy.

## Author contributions

Conceptualization, D. F. L. H. and C. B.; methodology, D. F. L. H. and N. B.; validation, J. d. O. P. and D. F. L. H.; formal analysis, D. F. L. H.; investigation, D. F. L. H., and J. d. O. P.; resources, F. J. A. and C. B.; writing—original draft preparation, D. F. L. H., J. d. O. P. and C. B.; writing—review and editing, D. F. L. H., F. J. A. and C. B.; visualization, D. F. L. H. and C. B.; supervision C. B.; project administration, F. J. A. and C. B.; funding acquisition, F. J. A. and C. B. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## References

- S. Capuzzi and G. Timelli, *Metals*, 2018, 8.
- J. Zou, Y. Chen and P. Zhang, *Ceram. Int.*, 2021, 47, 12661–12666.
- J. Cao, T. Hasegawa, Y. Asakura, P. Sun, S. Yang, B. Li, W. Cao and S. Yin, *Adv. Powder Technol.*, 2022, 33, 103576.
- A. Riapanitra, Y. Asakura and S. Yin, *Tungsten*, 2019, 1, 306–317.
- E. Aslan, M. Bulut, A. Grijalbo and B. Çiçek, *Adv. Powder Technol.*, 2022, 33, 103421.
- M. C. F. Karlsson, Z. Abbas, R. Bordes, Y. Cao, A. Larsson, P. Taylor and B. M. Steenari, *Dyes Pigm.*, 2019, 162, 145–152.
- S. W. Kim, G. E. Sim, J. Y. Ock, J. H. Son, T. Hasegawa, K. Toda and D. S. Bae, *Dyes Pigm.*, 2017, 139, 344–348.
- G. R. S. Cavalcanti, F. Rodrigues, G. Zhuang, S. Balme, J. M. Janot, M. G. Fonseca and M. Jaber, *Dyes Pigm.*, 2021, 190, 109306.
- H. Yang, B. Mu, S. Li, N. Wang, A. Hui and A. Wang, *J. Ind. Eng. Chem.*, 2022, 112, 440–450.
- X. He, F. Wang, H. Liu, L. Niu and X. Wang, *J. Am. Ceram. Soc.*, 2018, 101, 2578–2588.
- A. Zhang, F. Teng, Q. Zhang, Y. Zhai, Z. Liu, Z. Liu, W. Gu, W. Hao, Z. U. Abideen and Y. Teng, *Appl. Surf. Sci.*, 2018, 428, 586–592.
- S. Zhao, J. Guo, W. Li, H. Guo and B. You, *Dyes Pigm.*, 2018, 151, 130–139.



- 13 C. M. Álvarez-Docio, J. J. Reinoso, A. del Campo and J. F. Fernández, *Dyes Pigm.*, 2017, **137**, 1–11.
- 14 G. Costa, M. J. Ribeiro, W. Hajjaji, M. P. Seabra, J. A. Labrincha, M. Dondi and G. Cruciani, *J. Eur. Ceram. Soc.*, 2009, **29**, 2671–2678.
- 15 S. Yaemphutchong, W. Wattanathana, K. Chansaenpak, S. Singkammo, P. Kanjanaboos, P. Siri-apai, S. Janejobsakonkit, P. Pipattanaporn, N. Suetrong, S. Wannapaiboon and Y. Hanlumuayang, *Ceram. Int.*, 2022, **48**, 18490–18501.
- 16 C. E. Housecroft and A. G. Sharpe, *Inorganic chemistry*, Pearson Education, 2008, vol. 1.
- 17 D. F. L. Horsth, J. O. Primo, M. Dalpasquale, C. Bittencourt and F. J. Anaissi, *Cleaner Engineering and Technology*, 2021, **5**, 100313.
- 18 R. Salomão, L. F. Amaral and v. C. Pandolfelli, *Cerâmica*, 2010, **56**, 135–140.
- 19 S. A. Ahmed, *Cryst. Res. Technol.*, 2017, **52**, 1600335.
- 20 B. K. Kwak, D. S. Park, Y. S. Yun and J. Yi, *Catal. Commun.*, 2012, **24**, 90–95.
- 21 S. Luthra, S. K. Mangla, J. Sarkis and M. L. Tseng, *Resour. Policy*, 2022, **77**, 102652.
- 22 H. Desing, G. Braun and R. Hischier, *Resour., Conserv. Recycl.*, 2021, **164**, 105179.
- 23 D. F. L. Horsth, J. d. O. Primo, N. Balaba, J. S. Correa, C. M. Zanette, D. K. Silva, C. Bittencourt and F. J. Anaissi, *Nanomaterials*, 2022, **12**, 2771.
- 24 L. C. B. Lima, F. C. Silva, E. C. Silva-Filho, M. G. Fonseca, G. Zhuang and M. Jaber, *Appl. Clay Sci.*, 2020, **191**, 105604.
- 25 S. Parres-Esclapez, I. Such-Basañez, M. J. Illán-Gómez, C. Salinas-Martínez De Lecea and A. Bueno-López, *J. Catal.*, 2010, **276**, 390–401.
- 26 B. B. Bible and S. Singha, *Canopy Position Influences CIELAB Coordinates of Peach Color*, 1993, vol. 28.
- 27 X. Sun, X. Jiang, Y. Shan, X. Han, J. Xu and J. Li, *Ceram. Int.*, 2022, **48**, 17471–17480.
- 28 R. Khalighi, F. Bahadoran, M. H. Panjeshahi, A. Zamaniyan and N. Tahouni, *Microporous Mesoporous Mater.*, 2020, **305**, 110371.
- 29 C. Venkataramana, S. M. Botsa, P. Shyamala and R. Muralikrishna, *Chemosphere*, 2021, **265**, 129021.
- 30 G. Raja, A. Nallathambi, A. Prakasam, S. Gopinath, C. Ragupathi, S. Narayanan, P. Tamizhdurai, R. Kumaran, N. S. Alsaiani, K. M. Abualnaja and M. Ouladsmame, *J. Saudi Chem. Soc.*, 2022, **26**, 101440.
- 31 M. Müller, J. C. Villalba, F. Q. Mariani, M. Dalpasquale, M. Z. Lemos, M. F. G. Huila and F. J. Anaissi, *Dyes Pigm.*, 2015, **120**, 271–278.
- 32 S. v. Stefanovsky, O. I. Stefanovsky and M. I. Kadyko, *J. Non-Cryst. Solids*, 2016, **443**, 192–198.
- 33 M. Dalpasquale, F. Q. Mariani, M. Müller and F. J. Anaissi, *Dyes Pigm.*, 2016, **125**, 124–131.
- 34 A. S. B. Neto, A. C. Oliveira, J. M. Filho, N. Amadeo, M. L. Dieuzeide, F. F. de Sousa and A. C. Oliveira, *Adv. Powder Technol.*, 2017, **28**, 131–138.
- 35 M. Bouchard and A. Gambardella, *J. Raman Spectrosc.*, 2010, **41**, 1477–1485.
- 36 V. D'Ippolito, G. B. Andreozzi, D. Bersani and P. P. Lottici, *J. Raman Spectrosc.*, 2015, **46**, 1255–1264.
- 37 O. N. Shebanova and P. Lazor, *J. Solid State Chem.*, 2003, **174**, 424–430.
- 38 C. Ragupathi, J. J. Vijaya, P. Surendhar and L. J. Kennedy, *Polyhedron*, 2014, **72**, 1–7.
- 39 A. Navabi, M. Vandadi, T. Bond, V. Rahmashin, J. Obayemi, R. Ahmed, J. E. Oghenevweta, V. Champagne, N. Rahbar and W. O. Soboyejo, *J. Mater. Sci. Eng. A*, 2022, **841**, 143036.
- 40 T. Tago, N. Kataoka, H. Tanaka, K. Kinoshita and S. Kishida, *Procedia Eng.*, 2017, **216**, 175–181.
- 41 N. Srisawad, W. Chaitree, O. Mekasuwandumrong, P. Praserttham and J. Panpranot, *J. Nanomater.*, 2012, **2012**, 108369.
- 42 X. Duan, M. Pan, F. Yu and D. Yuan, *J. Alloys Compd.*, 2011, **509**, 1079–1083.
- 43 M. Gil-Calvo, C. Jiménez-González, B. de Rivas, J. I. Gutiérrez-Ortiz and R. López-Fonseca, *Appl. Catal., B*, 2017, **209**, 128–138.
- 44 J. L. Rogers, M. C. Mangarella, A. D. D'Amico, J. R. Gallagher, M. R. Dutzer, E. Stavitski, J. T. Miller and C. Sievers, *ACS Catal.*, 2016, **6**, 5873–5886.
- 45 Y. el Jabbar, H. Lakhli, R. el Ouati, L. Er-Rakho, S. Guillemet-Fritsch and B. Durand, *J. Non-Cryst. Solids*, 2020, **542**, 120115.
- 46 V. Elakkiya, R. Abhishekram and S. Sumathi, *Chin. J. Chem. Eng.*, 2019, **27**, 2596–2605.
- 47 A. Flachot and K. R. Gegenfurtner, *Vision Res.*, 2021, **182**, 89–100.
- 48 M. Salavati-Niasari, M. Farhadi-Khouzani and F. Davar, *J. Sol-Gel Sci. Technol.*, 2009, **52**, 321–327.
- 49 J. N. Roelofsen, R. C. Peterson and M. Raudsepp, *Am. Mineral.*, 1992, **77**, 522–528.

