Multi-analytical identification of pigments and pigment mixtures used in 17th century Portuguese azulejos

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Abstract

A multi-analytical approach was used to study the pictorial layers of a set of 17th century historic glazed tiles (azulejos) of Portuguese manufacture. The pictorial layer was studied by μ-EDXRF, μ-Raman, SEM-EDS and OM. Although the established palette comprises few pigments, it was found that the tiles were enhanced by the use of pigment mixtures, which were identified by μ-Raman and μ-EDXRF. As expected, the blue colour derives from cobalt oxide, purple from manganese oxide and emerald-green from copper oxide. Regarding the yellow pigment, μ-Raman results show it is consistent with the composition of a ternary oxide, whereas μ-EDXRF shows a high intensity of Zn in this colour, indicating a composition close to a Pb–Sb–Zn ternary oxide. Some compounds from the original ores were also indentified: in two very dark blue samples, cobalt and nickel olivines (α-Co 2 SiO 4 , α-Ni 2 SiO 4 ), and in a dark brown sample, braunite (a manganese silicate).

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Keywords: 17th century majolica; Glazed tiles (azulejos); Pigments; Glaze analysis

1. Introduction

When visiting Portugal, one cannot overlook the ever-present azulejo panels decorating many buildings and public spaces. Azulejos are considered the most important historic Portuguese decorative art, having been used without interruption for more than five centuries.1,2 Azulejos are the Portuguese name for ceramic glazed tiles. Their local use may have started as early as the 13th century entering Portugal through Moorish influence.3 The majolica technique was adopted in the second half of the 16th century and, possibly through the activity of Flemish and Italian artists in Portugal, provided the ground for the creation of impressive figurative panels which can still be seen today.1 In the first half of the 17th century, the country was undergoing a political transition derived from the coronation of Philip II of Spain as the Portuguese king by right of inheritance. The court was now set in Madrid and that caused an economic and artistic decline in Portugal. Consequently the azulejo evolved, possibly because of its gaudiness, as the chosen material to decorate an otherwise plain architecture, substituting the rich oriental embroider draperies that once hanged from the walls of churches.2 Being a relatively cheap but durable material, decorative tiles could be manufactured and painted by unskilled craftsmen in large quantities to cover entire walls. Pattern tiles (often called “tapestry” tiles) were especially important in this context, first mimicking tapestries and then evolving gradually over time towards more complex and novel patterns.2 The decorative effect was striking and monumental.

It was also in the 17th century that the chromatic palette of the Portuguese tile makers reached a peak before decreasing progressively to the blue-and-white period that had set by the end of the century. Colours were applied over a white lead–tin majolica glaze and comprised blue, yellow, oranges, greens, purple, browns and a very dark, almost-black colour used for contours. The lead–tin glaze was applied over the previously
fired ceramic body and the decorative motifs were painted before firing the glaze. Colours were obtained through metal oxides (copper, manganese, cobalt, iron) or synthesized pigments (Naples Yellow), which were used singly or mixed with other pigments.

Despite the importance of the azulejo heritage, relatively few scientific studies concerning its materials have been published. However, studies on other sorts of majolica have also been a valuable source of information about the materials and techniques used in the 16th and 17th centuries. The yellow pigment has been widely investigated in the last decade after the identification of a Naples Yellow variant – the Pb–Sb–Sn ternary oxide – by Roy and Berrie. Cobalt blue has been also the subject of several provenance studies.

This work aimed at the identification of the pigments and pigment mixtures used to produce the colours of 17th century azulejos, as a first step to understand the production technology of the Portuguese azulejo manufacturers. A multi-analytical approach was used to chemically characterize the pigments responsible for the observed colours. Non-destructive analytical techniques, such as μ-Raman and μ-EDXRF, were favoured. Results are systematized by the most representative colours, and compared with the production technology described in the well-known majolica treatise from the 16th century, “The Books of the Potter’s Art” by Cipriano Piccolpasso.

2. Experimental

2.1. Samples

A set of 28 fragments of 17th century Portuguese glazed ceramic tiles was studied. The samples were provided by the National Tile Museum (Museu Nacional do Azulejo) in Lisbon, and a few samples came from private collections. Fig. 1 shows some samples where all the representative colours are depicted. Table 1 includes a full list of the analysed samples, with the colours observed for each and the respective analytical techniques used on each colour.

2.2. Analytical techniques

2.2.1. SEM-EDS

A JEOL JSM-6400 with an Oxford Instruments micro-analysis unit equipped with a Si(Li) X-ray detector was used. Polished cross-sections were prepared by impregnating samples with an epoxy resin, polishing them to 3 μm and applying a graphite coating.

2.2.2. μ-EDXRF

Analyses were carried out using an ArtTAX spectrometer (Intax GmbH), with a Mo anode and an XFlash® detector. The measurement parameters were: 40 kV of voltage and 600 μA of current under a He atmosphere. Spectra of white and blue glazes were acquired in three different locations for 360 s, whilst for the other colours the measurement time was of 180 s and the number of spectra varying between two to three per colour.

Semi-quantitative analysis of the white glaze was performed using WinAxil and WinFund softwares. A reference lead–tin glass was synthesized as a reference. The accuracy of the method was calculated using the glass standard CMOG C (The Corning Museum of Glass). The accuracy of the method was <5% for SiO₂ and PbO, <10% for K₂O and SnO₂, <20% for MnO, CaO, CoO, NiO, CuO and ZnO, and <50% for Al₂O₃, TiO₂, Fe₂O₃ and As₂O₃. Relative standard deviation is <10% for SiO₂, K₂O and PbO, <15% for MnO, CaO and Fe₂O₃, <25% for Al₂O₃, As₂O₃, CuO and SnO₂, and <40% for TiO₂, CoO and NiO. The analyses of the blue, yellow and orange colours were carried out calculating the peak intensities of the elements (Kα for Co, Ni, Fe, As, Zn and Lα for Sn, Sb and Pb), using the WinAxil software.

2.2.3. μ-Raman

The equipment used was a Labram 300 Jobin Yvon spectrometer, equipped with a He–Ne laser of 17 mW power operating at 632.8 nm and a solid state laser of 500 mW power operating at 532 nm. The laser beam was focused with a 50× Olympus objective lens. The laser energy was filtered at 10% using a neutral density filter for all analyses.
Table 2
Chemical composition of the white glaze (wt%) determined by μ-EDXRF.

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<tr>
<th></th>
<th>Al₂O₃</th>
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</table>

Fig. 1. Analysed colours in 17th century Portuguese azulejos, exemplified with four samples. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

2.2.4. Optical microscopy
A Zeiss Axioplan 2 optical microscope, equipped with a high resolution camera Nikon DXM 1200F.

2.2.5. Binocular lens
Polished cross-sections were observed through an Olympus SZH binocular lens, with digital image capture.

3. Results and discussion

3.1. White glaze

The study of the white glaze was very important to understand the Portuguese tile production technology. SEM observations of polished cross-sections of the tiles showed a heterogeneous glaze with several mineral inclusions and gas bubbles. Mineral inclusions can be relatively large (ca. 20–100 μm), and consist mainly of quartz and potassium feldspars, according to results obtained from SEM-EDS (Fig. 2a and b) and μ-Raman analysis (Fig. 3). They may be associated with sand used as a raw material for the glaze manufacture. The coarse granulometry may be explained through economic factors: a faster production would be cheaper but simultaneously, those inclusions contribute to the opacification of the glaze and hence reduce the amount of SnO₂ needed to cloak the biscuit. Cassiterite (SnO₂) crystals were identified in situ by μ-Raman microscopy and observed through SEM as small agglomerates.

The high lead content of the glaze gives it a much brighter appearance than the ceramic body in backscattered SEM images (BSE) and a clear interface between the two components of the tile, resulting from the double firing, is evident. The glaze thickness of the 28 samples varies between 300 and 500 μm with some in-tile spread due to irregularities in the surface of the ceramic body.

Semi-quantitative chemical analysis by μ-EDXRF revealed relatively high contents of SiO₂ (58.0–72.0 wt%) and relatively low contents of PbO (13.0–32.2 wt%) when compared to other international studies on majolica. Variations in SnO₂ content (1.60–7.10 wt%) reflect the glaze appearance: samples with >5 wt% SnO₂ content show a more homogeneous and opaque
Fig. 2. Backscattering images of cross-sections of the majolica glazes: (a) white glaze of sample SCT27, showing large gas bubbles and several mineral inclusions. At the surface, the lead-rich yellow pigment layer stands out; (b) enlarged image, showing a gas bubble and cassiterite crystals.

Fig. 3. μ-Raman spectra of (a) potassium feldspar, (b) anatase, (c) cassiterite and (d) quartz identified in situ on the white glaze. Laser excitation 632.8 nm; 50× objective ULWD; laser power was kept at 1 mW.

Table 3
Chemical composition of each colour.

<table>
<thead>
<tr>
<th>Colours</th>
<th>Chemical composition</th>
</tr>
</thead>
<tbody>
<tr>
<td>“Simple” colours</td>
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<tr>
<td>Blue</td>
<td>Cobalt oxide</td>
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<tr>
<td>Purple/brown</td>
<td>Manganese oxide</td>
</tr>
<tr>
<td>Emerald-green</td>
<td>Copper oxide</td>
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<tr>
<td>Yellow</td>
<td>Yellow ternary oxide of Pb–Sb–Zn (?)</td>
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<tr>
<td>“Mixed” colours</td>
<td></td>
</tr>
<tr>
<td>Orange</td>
<td>Yellow ternary oxide of Pb–Sb–Zn (?) + α-Fe₂O₃</td>
</tr>
<tr>
<td>Olive-green</td>
<td>Yellow ternary oxide of Pb–Sb–Zn (?) + cobalt oxide</td>
</tr>
<tr>
<td>Dark-brown</td>
<td>Manganese oxide + α-Fe₂O₃</td>
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</tbody>
</table>

3.2. Blue colour

μ-EDXRF analysis of the blue decoration revealed, as expected, cobalt oxide as the colouring agent. Cobalt ions are well diffused throughout the glaze matrix revealing a very homogenous layer (Fig. 4a). μ-Raman analyses on several of the blue samples, and collected at different depths, returned the Raman signatures of SnO₂ and α-quartz, in the glaze matrix. None of the spectra collected could be associated with the blue pigment. On the other side, the Si–O stretching and bending envelopes typical of glass structures were often identified.

To gather further information on the chemical composition of the blue pigment, the μ-EDXRF spectrum of the white glaze was subtracted from that of the blue glaze. This procedure enables isolating the contribution of the blue colour, and allows the comparison between peak intensities without the interference of the chemical composition of the white glaze matrix.

Our results show a blue pigment characterized by an association Co–Fe–Ni–As. A small peak intensity of Mn was also detected and, in some samples, Zn and Cu. The Co–Fe–Ni–As association is consistent with saffre exported from the district of Schneeberg (in the Erzgebirge region in Germany), to the rest of Europe since the 16th century. There is a documented evidence of Portuguese trade of saffre from Germany, dating from the beginning of the 16th century. However, other possible sources of the pigment cannot be excluded. In the Iberian Peninsula, a well-documented site of cobalt production is Teruel, in Spain, although its mines were of limited size. The most common cobalt minerals include arsenic and sometimes sulphur in their composition. In sulpho-arseniates, Co can be replaced

\(^{A}\) A calcinated cobalt ore.
Another and more often recorded spectra, with two strong bands at ca. 826 cm$^{-1}$, identifies a nickel olivine structure (\(\alpha\)-Ni$_2$SiO$_4$) (Fig. 7b). The first band corresponds to the Si–O symmetric stretching band \(\Delta_g(Si-O)_{s\text{-str}}\), and the second one to the Si–O asymmetric stretching band \(\Delta_g(Si-O)_{a\text{-str}}\). From the few studies performed on Co- and Ni-olivines there is proximity in wavenumber of the asymmetric \(\Delta_g(Si-O)_{a\text{-str}}\) band but a decrease in wavenumber of the symmetric \(\Delta_g(Si-O)_{s\text{-str}}\) from Ni to Co, even in a series of solid solutions. The Raman spectra obtained are in agreement with a Ni olivine structure and a higher content of this element corroborates this identification.

3.3. Yellow colour

As observed in the cross-sections, the yellow colour was generally painted over the other colours and remains as a layer at the surface of the glaze (Fig. 4a). The thickness of the yellow pigment layer was measured as being approximately ca. 50–100 \(\mu m\) for all the measured samples. Under the microscope the yellow pigment is quite heterogeneous, consisting of very light-yellow crystals, yellowish-orange crystals and some red crystals (Fig. 6b). On some samples it was possible to observe clear yellow hexagonal crystals (Fig. 6c). According to previous studies on the pigment Naples Yellow, its crystal morphology can help estimate the firing temperature of the pigment, with hexagonal crystals forming above ca. 1100 °C. \(^{32}\)

\(\mu\)-EDXRF analyses of our samples identified, as expected, Sb and Pb in every yellow sample, consistent with the expected use of the Naples Yellow pigment. The strong intensities of the peaks in the collected spectra indicate a higher content of Zn, when compared to the white glaze matrix. Spectra collected for the yellow pigment were not subject to subtraction of the white spectra, because of the thickness of the yellow layer.

Tin was also present, but sometimes difficult to detect, due to the weak intensities of the element peaks and also to the overlapping with Sb and K peaks (Fig. 8). The presence of K in the yellows is consistent with coeval recipes for the Naples Yellow pigment, \(^{18,33}\) which indicate the use of potash ashes as a flux.

Several studies regarding Naples Yellow and its variants have been published in the last decade, \(^{14,32–39}\) with particular attention given to the identification of a Pb–Sb–Sn triple oxide yellow pigment in oil painting and majolica. Piccolpasso describes two yellows, a “light yellow” and a “yellow” pigment. \(^{18}\) The former corresponds to the yellow colour, and therefore to the pigment Naples Yellow (Pb$_2$Sb$_2$O$_7$) and the latter corresponds to the orange colour, observed in our 17th century majolica azulejos. There are three relevant references to the production of antimony yellow: Piccolpasso’s treatise, an early 17th century Italian treatise on miniature painting technique, by Valerio Mariani da Pesaro (1568–1625?), and the Darduin manuscript, a glass recipe.

Fig. 4. Ocular lens photograph of majolica cross-sections (a) blue pigment layer, where the pigment is well diffused into the glaze, contrary to the yellow pigment layer, which remains at its surface; (b) a layer showing the olive-green colour. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)
book from the 15th century from the workshop of the Oudin family. The three references mention the addition of a ingredient called *tuttia allessandrina* (or *tuccia*), but there is still debate on whether it refers to zinc or tin oxide. The Pb–Sb–Zn triple yellow has been identified in majolica plates dated from the 16th century and it is usually described as a darker yellow with an orange overtone. In our samples, Zn is a constant presence in all the analysed yellows.

The pigment Naples Yellow has a cubic pyrochlore structure (A$_2$B$_2$O$_6$O$^+$, with possible combinations of A$^{2+}$, B$^{5+}$ or A$^{3+}$ and B$^{4+}$) and is isostructural with the occurring mineral bindheimite (Pb$_2$Sb$_2$O$_6$(O,OH)). In 1998 Roy and Berrie disclosed the usage of a modified Naples Yellow with a ternary oxide structure including lead, antimony and tin (Pb$_2$Sb$_2$Sn$_x$O$_{6−x}$), in 17th century painting. Prior to this, Cascales et al. report, in 1986, the identification of a set of new pyrochlores, including the ternary oxide Pb$_2$(SnSb)O$_6$. Our Raman spectra are consistent with a modified pyrochlore structure identified by several authors as either a Pb–Sb–Sn or Pb–Sb–Zn triple oxide, where Sn or Zn entered the pyrochlore structure substituting Sb. Spectra shown in Fig. 9 exemplify the results obtained for the yellow pigment, distinguishing three types of spectra, with the following main characteristics: very strong signals from the Pb–O lattice vibration at low wavenumbers (120–139 cm$^{-1}$), weaker to medium bands around between 200 and 400 cm$^{-1}$ and a medium A$_1g$ band at ca. 510 cm$^{-1}$. When examining Fig. 9 the striking difference between spectra concern the nature and position of the strong band(s) arising from the Pb–O lattice vibration. We found indiscriminately in every yellow analysed a doublet between ca. 120 and 139 cm$^{-1}$ (Fig. 9a), a strong band at ca.

![Image](image1.png)  
*Fig. 5. Ratios of As/Co, Ni/Co and Fe/Co in the blue pigment, obtained from the peak intensities in μ-EDXRF analysis.*

![Image](image2.png)  
*Fig. 6. Optical micrograph of polished cross-sections of majolica glazes (a) dendritic crystals observed in the surface of the dark blue pigment in sample SCT20; (b) surface of the yellow pigment in sample SCT20 showing light yellow, yellowish-orange and red crystals dispersed in the glaze matrix; (c) surface of the yellow pigment layer observed in darkfield mode in sample SCT05 showing clear hexagonal crystals; (d) surface of the orange colour layer where yellow and red crystals can be observed; (e) surface of the olive-green colour, where it can be seen that the morphology is very similar to the one of the yellow pigment; (f) surface of the purple colour where dark and red crystals can be observed. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)*
Fig. 7. µ-Raman spectra of blue pigment in sample SCT20: (a) cobalt olivine ($\alpha$-Co$_2$SiO$_4$) and the Raman signatures of the glass structure; (b) nickel olivine ($\alpha$-Ni$_2$SiO$_4$). Laser excitation 632.8 nm; 50× objective ULWD; laser power was kept at 1 mW.

Fig. 8. Overlapping µ-EDXRF spectra for the yellow and orange pigments in sample SCT05. Differences are notorious in Zn and Fe peaks.

Fig. 9. µ-Raman spectra of the yellow pigment showing three types of obtained spectra: (a) sample SCT 17, Raman spectrum showing a doublet at ca. 120–136 cm$^{-1}$; (b) sample SCT 27, spectrum showing a strong band at ca. 135 cm$^{-1}$ (Fig. 9b) or a strong band at ca. 135 with a shoulder (Fig. 9c). Previous studies assert that this band is not good for discrimination between the ternary oxides, since its intensity on the unmodified Naples Yellow also varies significantly, and can also change with the firing temperature of the pigment. Nonetheless, Rosi et al. observed in mock-up samples that Zn-modified Naples Yellow shows a strong band at ca. 145 cm$^{-1}$ and Sn-modified Naples Yellow shows a doublet at 125 and 142 cm$^{-1}$. We observed the three situations in every sample analysed and at different points in one sample, being unable to find a pattern for our results. The A$_{1g}$ band at ca. 510 cm$^{-1}$, corresponding to the symmetric elongation of the SbO$_6$ octahedra is considered more advantageous to discriminate the nature of the ternary oxides. Whilst on the unmodified Naples Yellow it is a strong band, it collapses and shows the emergence of a well-resolved band at ca. 450 cm$^{-1}$ for the ternary oxides. For Zn-modified Naples Yellow this effect is stronger. On our samples this band shifts ca. 5 cm$^{-1}$ within one sample, and the 450 cm$^{-1}$ band can emerge more or less resolved, also within one sample. Finally the appearance of a band at ca. 640 cm$^{-1}$ has been assigned, on one hand, as a Sn-modified Naples Yellow with silica added, and on the other as a binary lead antimonate with Pb excess. In sum, the constant alteration on the observed main spectral signatures of the yellow pigment, even within the same sample, does not lead to an identification of which ternary yellow oxide is responsible for the yellow colour, by Raman microscopy. In contrast, µ-EDXRF analyses show an intense Zn signal for every sample, with a small intensity signal for Sn, suggesting a ternary Zn yellow oxide.

3.4. Orange colour

Observation of the orange coloured areas reveals a very similar appearance to the yellow-coloured ones. The colour is opaque, sometimes rough to the touch, and it remains at the surface of the glaze (also confirmed with cross-sectional observation). Through OM, the surface of the pigment is also very similar to the yellow one, the exception being the very dark particles intertwining the yellow and orange crystals (Fig. 6d). In the orange colour, seemingly the *giallo* pigment described by Piccolpasso, we find our first true mixture colour, i.e., the mixture of two compounds to make a new colour.

The yellow and orange crystals were identified by µ-Raman as the previously discussed yellow pigment, with a ternary oxide structure, indicating the use of the same yellow pigment in both colours. Two spectra of the yellow and orange pigments in sample SCT05 are represented in Fig. 9d, showing an identical structure. The dark particles, on the other hand, were identified as hematite ($\alpha$-Fe$_2$O$_3$).

Similarities are also visible in the results from the chemical analysis through µ-EDXRF and µ-Raman. µ-EDXRF spectra collected in the orange colour identified Sb and Pb, as well as Zn, but show a higher Fe peak intensity when compared to the spectra collected for the yellow. Fig. 8 shows two overlapped µ-EDXRF spectra collected on the yellow and orange pigments for the same tile. The two spectra are identical, with the exception of
the Fe and Zn peak intensities. The Fe peak intensity is always higher in the orange colour, whereas the Zn content can vary significantly; when observing the spectra in Fig. 8, one can see there is a large difference in the intensity of the Zn peak, being much higher in the yellow pigment, but in other samples this difference was not notorious. Whilst the Zn is always a very important element in the yellow pigment, it sometimes shows only a very small peak in the μ-EDXRF spectra for the orange colour.

The orange pigment is thus a mixture of pigments, obtained by adding iron oxide to a yellow pigment. However, in some samples, it seems that it was not the final ternary yellow pigment that was used as a basis for the orange colour. Instead, the results of the chemical analyses indicate that there was a yellow pigment used as a basis for the recipes, to which was added either iron oxide to achieve the orange tone, or and zinc oxide to obtain the final yellow pigment.

3.5. Olive-green colour

A green colour of an olive tone and opaque appearance is often found in Portuguese 17th century azulejos. When observed through the optical microscope, its surface reveals a bluish-green glaze matrix in which crystals identical to the ones observed for the yellow pigment are dispersed (Fig. 6e). Cross-sections show a blue layer in the glaze matrix, corresponding to Co ions dissolved and diffused in the glaze, and a yellow layer on the surface, corresponding to the yellow pigment (Fig. 4b).

μ-EDXRF analysis identified both cobalt and antimony peaks as characteristic of this green and, in some cases, a high zinc peak. As for the yellow colour, the μ-EDXRF spectra collected for the olive-green is very similar to that of the yellow pigment (Fig. 10). The variation in the intensity of the Zn peak occurs in the same way as explained for the orange colour: in some samples it fits perfectly with the yellow spectrum and in other samples it is much weaker. Cobalt was identified in every olive-green along with its associated elements: Fe, Ni and As (vide Section 3.2).

μ-Raman analysis on this colour returned spectra consistent with the yellow pigment (as examples see Fig. 9). It was not possible to collect a spectrum of the blue compound.

The olive-green colour is thus a mixture of the yellow and blue pigments, previously described. As happened with the orange colour, some samples show a difference in Zn content, indicating that the pigment used for pure yellow is different from the basis used for the mix resulting in the olive-green colour.

Picolpasso describes a “mixed green” as a mixture of a copper oxide (ramina) with the yellow pigment. Although this seems to be a commonly used green in Italian majolica, copper was not identified in the Portuguese olive-green. However, a mixed green resulting from blue and yellow was also used by Della Robbia’s workshop.

3.6. Emerald-green colour

Besides the already mentioned olive-green, another green, of an emerald tone and transparent appearance, was observed in samples SCT10 and SCT29 (Fig. 1). This emerald-green colour was identified both by μ-EDXRF and μ-Raman analysis to derive from a copper oxide (CuO) dissolved in the lead–tin glaze (Fig. 11a and b). This pigment dissolves and diffuses easily in the glaze matrix, often causing the green to spread beyond the limits of the decorative motifs. This undesirable characteristic may justify its limited use in Portuguese azulejos.

3.7. Purple/brown/dark-brown contours

A set of tones between a dark brown and purple are found on Portuguese 17th century azulejos both in areas of colour and in figure and motif contours (Fig. 1). The colour itself is difficult to define, due to its variability and imprecise nomenclature used in Portuguese literature concerning azulejos studies. Optical microscopy reveals a morphological distinction between the purple and the brown/dark brown colours. For the purple colour the pigment is in solution in the glaze matrix, whereas for the brown/dark brown the colour is heterogeneously distributed although mostly at the surface of the glaze. Moreover it was observed that this colour is composed of areas of a dark metallic shade and areas of a reddish colour (Fig. 6f).

SEM-EDS and μ-EDXRF analysis show a manganese oxide responsible for the purple colour, as expected, and the pigment is distributed throughout the thickness of the glaze. According to μ-EDXRF analysis the brown/dark brown colour not only has a high Mn peak intensity, but also contains Fe and Ba. The latter was found in association with manganese in every μ-EDXRF analysis whenever the Mn signal was relatively high. The presence of barium could suggest the use of the mineral psilomelane (BaMn2+Mn4+8O16(OH)4) as a source of manganese, and typically the absence of it hinted at the use of pyrolusite (β-MnO2). These two manganese minerals correspond to two important and worldwide distributed manganese sources. However, Tite et al. pointed out that this assumption represents an oversimplification, pointing to the wide range of Mn minerals in nature, together with veins of baryte (BaSO4) and also Ba-feldspars. Raman analyses of samples SCT 36, SCT
Fig. 11. (a) μ-EDXRF spectrum of the emerald-green colour, with a very intense peak of copper; (b) μ-Raman spectrum of copper oxide, identified in the same colour. Laser excitation, 632.8 nm; 50× objective ULWD; laser power was kept at 1 mW.

10 and SCT 27 revealed a different kind of manganese source to make the brown/dark pigment. The samples analysed show a distinct thick dark layer and under the microscope it is possible to observe bright hexagonal crystals. Raman spectra of those crystals show the characteristic signature of braunite, a manganese silicate \((\text{Mn}^{2+}, \text{Mn}^{3+})_6\text{O}_8\text{SiO}_4\) (Fig. 12a).\(^{44}\) Braunite is an important manganese ore along with psilomelane, pyrolusite and manganite \((\text{MnO(OH)})\). This Mn silicate usually appears in metamorphosed manganese oxide ores or in hydrothermal deposits.\(^{45}\) It is a common mineral in the Portuguese south region of Alentejo (where most manganese in Portugal is extracted), namely in a region known as Iberian Pyrite Belt, of hydrothermal origin. Braunite emerges here as a secondary mineral following psilomelane and pyrolusite, alongside with iron oxides such as hematite, goethite and lepidocrocite, and also baryte. In conclusion, the manganese source could be a local one. Comparing the spectra obtained for braunite with the reference spectrum from RRUFF database, our spectra is shifted to a lower wavenumber.

Fig. 12. (a) μ-Raman spectra of braunite (a manganese silicate) identified in samples (i) SCT 36, (ii) SCT 10, (iii) SCT 27 and (iv) Reference spectrum of braunite from RRUFF project database.\(^{44}\) (b) μ-Raman spectrum of hausmannite obtained from sample SCT 36. (c) μ-Raman spectra on sample SCT 36 of (i) hematite mixed with goethite or a heated goethite structure; (ii) heated goethite structure; (iii) lepidocrocite. Laser excitation 632.8 nm; 50× objective ULWD; laser power was kept at 1 mW.
Fig. 13. Two types of purple/brown/dark-brown pigments: μ-EDXRF spectra of colour areas and contours in samples SCT10 and SCT31.

cia. 6 cm\(^{-1}\) for every Raman band (Fig. 12a(iii)). This could be due to cation substitution in the structure of braunite, where Mn\(^{3+}\) can be substituted by Fe\(^{3+}\), and Mn\(^{2+}\) by Mg\(^{2+}\), Ca\(^{2+}\).\(^{45}\) Since thermal absorption after the laser-induced heating, may lead to the formation of new mineral phases or amorphous materials,\(^{46}\) the beam power was kept low and the consistency of the spectra was always checked at different beam powers. Analyses were performed on small particle aggregates, a procedure preferentially used for dark materials, due to strong the coupling between dark materials and the laser light.\(^{47,48}\)

On sample SCT 36 the characteristic Raman signature of hausmannite (Mn\(^{2+}, Mn^{3+}\)\(_2\)O\(_4\)) was found, a spinel mineral with a strong A\(_{1g}\) band at ca. 659 cm\(^{-1}\), and two other weaker bands at 370 and 317 cm\(^{-1}\) (Fig. 12b).\(^{49}\) It is also found associated with other manganese minerals such as pyrolusite and psilomelane.

Some iron oxides were as well identified by μ-Raman (Fig. 12c), specifically hematite (α-Fe\(_2\)O\(_3\)), and lepidocrocite (γ-FeO\(_2\)OH), with the typical Raman bands for each compound.\(^{50}\) Hematite was often identified by itself (in the reddish colour areas) or in association with goethite (Fig. 12c(i)). This particular spectrum could also be assigned to a heated goethite structure, having a disordered hematite structure. Goethite can suffer dehydration to a hematite structure at low temperatures (ca. 260–280 °C), this dehydration process could have occurred here as an artefact resulting from a laser-induced heating or from the firing of the pigment in the glaze.\(^{51,52}\) Fig. 12c(ii) could also be assigned to the same structure of a heated goethite. Nonetheless this disordered hematite can be obtained from other factors namely grinding, biodegradation or weathering.\(^{53}\) Finally, lepidocrocite, found in sample SCT 27, is a common mineral found in iron ores (Fig. 12c(iii)).

μ-EDXRF analysis of the colour areas and the contours identified Fe, Mn and Ba (associated with the Mn) as the main elements in the purple, brown and dark-brown colours. In four of seven analysed tiles, the very dark colour of the contours was achieved by a higher content of a manganese-rich pigment, whilst for the rest of the samples, the contour showed a very intense peak of iron, considerably higher than the manganese one. To the naked eye, some samples exhibit a reddish colour in the iron-rich contour, easily observed under the optical microscope as red hexagonal crystals, later identified by μ-Raman as crystals of hematite. The chemical composition of the coloured areas is identical in all samples analysed, both in the purple and the brownish areas, suggesting the use of the same manganese-rich pigment. Thus, there are two types of purple/brown/dark-brown pigments used in these tiles: a manganese-rich pigment, used both for colour areas and for contours, and also an iron-rich pigment identified only in contours (a mixture of the manganese pigment and iron oxide in the form of hematite). Fig. 13 exemplifies the two cases: sample SCT10, where the same pigment was used in the colour area and in the contour, and sample SCT31, where different pigments were used.

With a multi-analytical approach it was possible to characterize the pictorial layers of 17th century Portuguese glazed tiles (azulejos) and determine the components used to create specific colours and tones. The chemical composition of the identified colours is summarized in Table 3.

4. Conclusions

Colours in 17th century Portuguese azulejos were varied in tones and chromatic effects, though obtained with a limited number of compounds: cobalt oxide, copper oxide, manganese oxide, iron oxide (as hematite) and a yellow pigment, possibly a Pb–Sb–Zn ternary oxide.

Two types of colours were distinguished based on their chemical composition:

1) the “simple” colours, where one single compound was employed, namely a metallic oxide dissolved in the glaze matrix (cobalt blue, copper green and manganese purple/brown) or, for the yellow colour, a yellow ternary oxide, possibly a Pb–Sb–Zn structure.

2) the “mixed” colours, which were made by mixing at least two compounds, such as the cobalt oxide mixed with the yellow synthesised pigment to give an olive-green; the yellow pigment mixed with hematite to give orange; and a manganese oxide (or silicate) with hematite to give a brown/dark colour.

Although the number of basic components was somewhat limited, the combination of at least two compounds to achieve different colours and shades lead to a richer palette available to the Azulejo painter. Yellow itself, as used in areas of pure colour by some shops, may actually have been obtained from a shop admixture of binary Naples Yellow with zinc oxide. This is a new result that may be important for provenance studies.
Studying the pigments in Portuguese azulejos was a first step in the understanding of the techniques used by the local tile makers. We hope that in a near future a broader study will make it possible to allocate tile panels to their regions of provenance and even to specific workshops.

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References


44. RRUFF project database. [http://rruff.info/braunite/display=default/R050385](http://rruff.info/braunite/display=default/R050385) [accessed 24.02.11].


