Beads excavated from Antsiraka Boira necropolis (Mayotte Island, 12th-13th centuries)
Colouring agents and glass matrix composition comparison with contemporary Southern Africa sites

Noémi Fischbach, Anh-Tu Ngo, Philippe Colomban and Martial Pauly
Beads Excavated from *Antsiraka Boira* Necropolis (Mayotte Island, 12th-13th centuries)

**Colouring Agents and Glass Matrix Composition**

**Comparison with Contemporary Southern Africa Sites**

*Agents colorants et matrices vitreuses des perles de la nécropole d’Antsiraka Boira (Île de Mayotte, xii-xiii siècles)*

*Comparaison avec les sites contemporains d’Afrique Australe*

**Noémi Fischbach**, **Anh-Tu Ngo**, **Philippe Colomban** and **Martial Pauly**

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**Abstract**: About hundred out of three hundred colored beads excavated from the necropolis of Antsiraka Boira (AB), in Mayotte Island (12-13th c.) were classified according to Wood’s morphological criteria and studied with a portable Raman spectrometer (532nm). Based on the recorded spectra, 22 beads were identified as representative and further analyzed in the laboratory with High-Resolution Raman spectrometers, using wavelengths of 458, 633 and 785nm. Additional SEM-EDS analysis was carried out on the surface and, sometimes, the bead cross-section. It turns out that white beads are made of aragonite and that almost all other beads have a soda glass matrix. Pyrochlore (yellow), amber/Fe-S (black), manganese oxide (black), copper metal nanoparticles (red), and Cu$^{2+}$ ions (turquoise) chromophores were identified. Some red, yellow, black and turquoise beads also show the signature of chromium-doped tin sphene that could therefore be used as a marker. Most beads from the AB site can be classified as “Indo-Pacific”, revealing a similarity with the contemporary South African site of K2 (close to Mapungubwe). However, some red and black beads are similar to molten ceramic beads from the Vohemar Islamic necropolis (13-17th century AD, Madagascar Island). The on-site Raman analysis appears sufficient for the identification of chromophores and glass types.

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**Keywords**: Mayotte, glass, beads, Indian Ocean, pigment, composition.

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1. INTRODUCTION

At the beginning of the first millennium A.D., the Indian Ocean became a region of long-distance trade between the Middle East, China, India, South Asia and Africa. Over the centuries, mostly thanks to monsoons, it was organized into a space around which the first World Trade system was built (Beaujard, 2009; Beaujard, 2012). Indeed, by its geographical situation, the Indian Ocean is subject to the phenomenon of monsoon winds whose seasonal fluctuation determined navigation and commerce in the area (Nativel & Rajaonah, 2007). At the time, it was possible to travel from Asia-to-Africa and up to the entrance of the Mozambique Channel in a single season; on the other side, trade to ports located more to the South needed a waiting period for the next cycle. Livestock, raw or processed materials like ivory, pearls, wood, metals, pottery and glass beads have travelled over the Indian Ocean trade routes. Glass beads, which were used as a currency, attest to international trade by their presence. They are among the most abundant objects found at excavations due to their very good preservation, even in the humid tropics.

Identifying the source of excavated beads is useful to understand the history of exchanges and the building of trade routes. Beads are present from the 7th and the 20th century and over this period morphological and compositional changes take place. These modifications can be associated to the onset of specific periods of production or to some geographical area (Dussubieux et al., 2008; Prinsloo & Colomban, 2008; Robertshaw et al., 2006; Robertshaw et al., 2009; Robertshaw et al., 2010; Tournié et al., 2011; Wood, 2005; Wood, 2011; Wood et al., 2009; Denbow et al., 2015). International trade to South/East Africa through the Indian Ocean during the Iron Age (~ 600-1750 A.D.) received considerable attention (Beaujard, 2009; Beaujard, 2012). The first study of glass beads discovered in South Africa followed the excavation in 1930 of two large sites in the Limpopo valley, namely Mapungubwe and K2. The chronology of these sites has long been debated; the abundance and the variety of beads found there naturally led to use them as markers. To this end, numerous studies attempted to classify beads according to morphological criteria (Beck, 1937; Van Riet Low, 1955; Schofield 1958; Gardner, 1963), elemental composition or traces of rare earths (REE) (Davidson, 1972; Davidson, 1974). Recently, Wood proposed a classification system for southern African beads from the precolonial period (700-1600 A.D.) (Wood, 2005; Wood, 2011; Wood et al., 2009). This classification, based on morphological and technological characteristics – which implies subjectivity – defines seven different sets of beads, each of them being related to a particular period determined by radiocarbon dating (RC) of the site from which they were excavated. By definition, this classification requires reliable radiocarbon dating of the associated sites. However, the 1970s’ RC dating of Mapungubwe and K2 sites was heavily influenced by the ideological context of the time, and questioned by the recent identification of chrono-technological markers (Tournié et al., 2011), which were more in line with the early work by Gardner (1963). To improve this ranking system, elemental analyzes were conducted by LA-ICP-MS (Robertshaw et al., 2006; Robertshaw et al., 2009; Robertshaw et al., 2010) so to provide information on glass production techniques and origins, including colouring elements (Robertshaw et al., 2010). Additionally, trace elements usually found in African glass beads compared to those found in India, Sri Lanka and East/South African Coast ones have enabled more accurate time correlations (Chaisuwan, 2011; Dussubieux et al., 2008; Dussubieux et al., 2010; Robertshaw et al., 2006; Robertshaw et al. 2009).

Meanwhile, Raman spectroscopy has emerged as the most effective non-invasive method to identify chromophores, pigments and opacifiers in glasses and enamels (Bonneau et al., 2013; Colomban et al., 2001; Colomban & Trepoz, 2001; Colomban et al., 2003; Ricciardi et al., 2009a; Ricciardi et al., 2009b; Prinsloo & Colomban, 2008; Prinsloo et al., 2011; Tournié et al., 2010, Tournié et al., 2011; Welter et al., 2007) as well as the types of glass matrix (Colomban et al., 2006a; Colomban et al., 2006b). This approach is widely used now (Baert et al., 2011; Caggiani et al., 2014; de Ferri et al., 2012; Gedzvičiūtė et al., 2009; Koleini et al., 2016a; Prinsloo et al., 2011; Prinsloo et al., 2012). Some pigments and chromophores are unique technological chrono-markers. In addition to confirming the first chronology of the site established by Gardner, Raman analysis of Mapungubwe and K2’s beads also confirmed the links that Africa had with the outside world thanks to the finding of: Indo-Asian black beads with Fe-S chromophore (Prinsloo & Colomban, 2008); Lapis Lazuli colored blue beads typical of Islamic glasses between the 13th and 15th century (Caggiani et al., 2014; Prinsloo et al., 2011). These studies concern the major sites of the upper valley of the Limpopo River (Mapungubwe / K2) but other sites were studied: such as further along the Limpopo in South Africa (Koleini et al., 2016c), Botswana (Koleini et al., 2016a), and Zimbabwe sites, rivals of Mapungubwe and K2 ones in a later date (Koleini et al., 2016b) and on the material collected in the early 20th century at Vohemar, a harbor occupied by Malagasy Islamized and in close rela-
tionship with the Swahili world in northeastern Madagascar (Colomban & Simsek, 2014).

This work concerns a selection of beads from the necropolis of Antsiraka Boira, Mayotte, one of the four islands in the Comoros Archipelago in the Indian Ocean. Mayotte is located at the northern entrance of the Mozambique Channel between the island of Madagascar and East Africa, and because of its position at the end of the road of Monsoon (going south imposes waiting for the next cycle of Monsoon) an important step in trade with southern Africa. Commercial networks with the other islands of the channel, Madagascar and Africa, are attested from the 8th to the 13th century (Beaujard, 2009; Beaujard, 2012; Pauly, 2014). Because of its position at the entrance of the Mozambique Channel, many Islamic traders chose to establish trading posts in the island of Mayotte ever since the 9th century, which led to the introduction of the Swahili culture and the Muslim religion into the Island. The spread of Islam across the Indian Ocean is closely related to commercial networks linking Africa to the Middle East (Pauly, 2014). Recent excavations (2012-2014) have uncovered a medieval necropolis dated to around 1100-1250 with a well-defined stratigraphy (Pauly, 2014). Nearly two thirds of the excavated tombs contained glass beads, for a total of about ten thousand. These burials revealed a phase of synergy between Muslim funeral rituals and pre-Islamic rituals. With the exception of the Islamic Necropolis of Vohemar (13th-17th centuries), northeast of Madagascar, Antsiraka Boira is the sole provider of bead adornments found in funerary context (table 1, the concordance between the labels used in this work and those of the funerary excavation). About a hundred beads of the necropolis of Antsiraka Boira, representative of 331 beads chosen on morphological criteria (shape, color, size and type of apparent production technology) were first Raman analyzed with a portable 532 nm set-up that can be used on site (Colomban, 2013). Twenty-two of them (Figure 1) were selected based on their Raman signature representative of the different groups for further analysis with a laboratory instrument. Indeed, this work also aims at establishing an appropriate on site selection methodology, in order to limit the number of objects requiring additional laboratory analysis.

2. MATERIALS AND METHODS

Beads

Figure 1 shows the 22 selected beads and the first classification according to the morphological criteria defined by Wood (Wood, 2005, Wood, 2011, Wood, 2012). The beads used in the discriminating criteria can be linked to a series associated with a chronological period. According to visual criteria, almost all beads belong to the Indo-Pacific category, divided into three series: K2-IP, East Coast (EC-IP), and Khami-IP (Figure 1 and Table 2). All the beads are monochrome, but of different colors: black (n), blue / turquoise (b), yellow (y), and red-brown (r). Most of the selected beads were made by drawing a glass gob and have a cylindrical shape. Two beads were made by winding in a more or less spherical fashion (10-n and 22-r). Beads 18-y and 17-b do not exhibit morphological criteria suitable to assign them to a particular series and bead #11 is not made of glass.

Analyses

A small spot on the bead surface was cleaned with a fine dry abrasive paper in order to remove surface residues and any loose layers of corrosion products, to obtain the best Raman signature on the uncorroded matrix. Micro-Raman analyzes were performed with 4 spectrometers: a portable HE 532 (HORIBA Jobin Yvon Scientific), a fixed HR 800 (HORIBA Jobin Yvon Scientific), and a transportable

<table>
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<th>Label</th>
<th>1</th>
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<td>SP10PRp4</td>
<td>SP02PJP3</td>
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<td>SP30PNICp1</td>
<td>SP09PNp11</td>
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<td>SP01PNp2</td>
<td>SP34PCp26</td>
<td>SP09PNp2</td>
<td>SP02PNp2</td>
<td>SP10PRp5</td>
<td>SP10PTp2</td>
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<td>SP34PRp2</td>
<td>SP30PNICp2</td>
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Table 1: Correspondence between archaeological labels (sepulture number (SP), bead number (P) color (red, R; …), loin cloth (p), necklace (c)).

Tableau 1: Correspondance entre les numéros d’inventaire archéologiques (numéro de sépulture (SP), de perle (P); symboles : couleur (rouge, R; …), pagne (p), collier (c)).
Infinity (Dilor-HORIBA Scientic) and a transportable SENTERRA Scope III (Bruker Optics). All are coupled with long working distance microscope objectives, magnification of x10, x50 and x100 that lead to 100, 500 and 1,000 magnifications. The use of different optics allows controlling the glass heterogeneity. For each record, Raman spectrometers were used with an acquisition time of 30 to 450 seconds, and if needed, several accumulations to improve the signal / noise ratio and clear the cosmic photons. The hundred beads were analyzed with the portable spectrometer (under excitation of ~2-5 mW for black beads because of the high absorption, and ~10-20 mW for colored beads). Those whose spectrum recorded with HE532 portable instrument was of low quality (yellow and blue) were also measured with the HR 800 spectrometer under laser excitation of 458 nm, blue excitation being well suited for the analysis of glasses. Infinity (633 nm, 35 mW) and SENTERRA (785 nm, 10-50 mW) spectrometers were used on black beads to evaluate and / or to overcome the resonance efect of the Fe-S chromophore.

Despite the lower spectral resolution of the portable instrument and the limited access to low wavenumber range (<100 cm⁻¹), most Raman spectra obtained with the portable apparatus are significant and exploitable (Figure 2a) and...
rather similar in quality to those obtained with the laboratory instruments (Figure 2b). On a single bead, depending on the analyzed spot, the results can differ (e.g. Figure 3c) due to the glass heterogeneity among beads.

Archaeological materials generally exhibit strong fluorescence due to organic residues on the surface or in the pores and cracks of the material. These residues are removed under energetic radiation (a few mW blue laser or tens of mW green laser, typically with x50 objective) or by examination of a fresh fracture. Thus, a bead of each colour (1-r, 15-b, 18-y, 21-n and 22-r) was cut parallel to the perforation by performing a cut with a diamond saw Minitom (Struers) and then mechanical fracturing to rate the heterogeneity of the matrix with fixed Raman spectrometers by performing five records in the section along a diagonal.

Subtracting the background of the Raman spectra was conducted using the multi-segment procedure validated in other glass studies (Colomban, 2008; Tourinié, 2009) with the Labspec software to compare the spectra obtained on various instruments.

The Raman spectra recorded with fixed instruments and different wavelengths have a resolution twice as good and in some cases a lower background than those recorded with the portable set-up. For crystalline compounds, pigments or secondary phases, the characteristic Raman bands are narrow, intense and thus easily detectable (Figure 3). In case of (pre)resonance, change of the excitation wavelength significantly modifies the respective intensities of the chromophore and its matrix allowing a better observation of one or the other (Faurel et al., 2003).

**Scanning electron microscopy (SEM-EDS)**

Elemental analyzes (by oxide weight %, Table 3) of Boira Antsiraka beads were performed with a SEM (JEOL JSM-5510LV) equipped with an EDS detector (Energy Dispersive Scattering). A 30kV acceleration voltage was used with a working distance of 20 mm. Fixed to a metal support with carbon adhesive recovery to best remove the charges created by electron irradiation, beads were examined without preparation. The validity of the measurements was monitored by applying the same procedure to certified glass-reference samples “Corning Museum B, C and D” and American “National Bureau of Standard (NBS 620)”, as usual (Vicenzi et al., 2002; Henderson et al., 2016). The error is below 1% for SiO₂, Na₂O and CaO, and below 5% for the other oxides. For each bead, several qualitative and quantitative measurements were performed, first at low magnification (100x, overall analysis), then at high magnification (up to 3000 x) to assess the material heterogeneity.
3. RESULTS

Identification of the chromophores

Raman spectroscopy has proven to be particularly effective in identifying colouring matters, except for the chromophore ions (Cu\(^{2+}\), Fe\(^{2+}\), Fe\(^{3+}\), Co\(^{2+}\), Mn\(^{n+}\), etc.) that are “simply” dissolved in the glass matrix, as their presence do not have a strong influence on the matrix spectrum (Colomban et al., 2001). The Raman signature of well crystallized compounds usually consists of narrow peaks that are easy to detect. This is the case of most pigments and opacifiers: highly crystalline compounds that are stable in the molten glass or have precipitated on cooling. Additionally, when the excitation wavelength (almost) corresponds to the coloration, the (pre-)resonance character of the spectrum improves its detectivity and small amounts of chromophores (<% wt) give intense spectra.

Yellow beads

For yellow beads 3-y and 4-y (Figures 2, 3 & 4) the Raman spectrum is characteristic of a pyrochlore pigment, a complex solid solution PbSb\(_{2-x}\)Sn\(_x\)Z\(_y\)O\(_{7-\delta}\), also called Naples Yellow. The spectrum of this pigment is characterized by bands at about 135, 330 and 453 cm\(^{-1}\) (Kirmizi et al. 2010; Rosi et al., 2009; Sakellariou et al., 2004). The exact position of these bands and the relative intensities vary according to both the composition (including the Sn / Sb ratio and the non-stoichiometry in oxygen resulting from oxidation state of the metal elements) and heat treatments, thus modifying the non-stoichiometry and order (Pereira et al., 2009). It is often difficult for determining the non-stoichiometry (Naples Yellow Type I [Sb-rich, usually recognized from its 510 cm\(^{-1}\) band] or II [Sn-rich with characteristic 450 cm\(^{-1}\) component]) only on the basis of the Raman spectrum. The yellow bead 18-y, does not exhibit the pyrochlore signature. The absence of lead is confirmed by SEM-EDS analysis, and high iron content (1.9 wt oxide %) rather indicates a colour due to Fe\(^{3+}\) ions, a known technique for enamels but rare for glass objects. A resonant spectrum (761-775 cm\(^{-1}\) and 942-963 cm\(^{-1}\) peaks followed by harmonics / combinations, Figure 4) characteristic of a sphene (malayite) containing tin doped with chromium and used for the pink to brown porcelain enamel since the 19th century (Faurel et al., 2003),

Figure 3: Raman spectra of the chromophore Cr-Sn / sphene recorded by a) the fixed HR800 instrument (458nm) and b) the portable HE532, (532nm); c) comparison of representative spectra recorded with 532 (not labelled) or 458 nm (labelled) excitation (LB: the baseline has been subtracted).

Figure 3 : Spectres Raman du chromophore Cr-Sn sphène obtenus avec a) le spectromètre fixe HR800 (458nm) et b) le spectromètre portable HE532, (532 nm). c) Comparaison de spectres typiques obtenus avec les excitations à 532 ou 458 nm (LB: une ligne de base a été soustraite).
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is obtained at points of the beads 18 and 4-y. The SEM-EDS analysis confirms the presence of chromium (0.02-0.04 wt%) and tin (0.2-0.3 wt%). The ratio between the two elements is consistent with this type of pigment. When this sphene pigment is combined with other chromophores, such as Naples yellow (bead 4-y) or the yellow iron (bead 18-y), it could only change the hue, not the colour. This very specific Raman signature can be used as a marker of bead production.

Red beads

Raman signatures vary according to the beads and the analysed spot (Figure 4a, spectrum 22-r and 1-r). Elemental analyses were able to detect the presence of copper (0.04 -0.2 wt%) and iron (0.4-0.8 wt%) in all the red beads, in accordance with a red colouring obtained by dispersing Cu⁰ metallic copper nanoparticles in the glass matrix (Colomban, 2009; Colomban et al., 2009). This is consistent with the complexity of Raman signatures (Colomban & Schreiber, 2005). The above mentioned Raman signature of the Cr-Sn chromophore was also identified for three beads, 2-r, 20-r and 22-r, with some variations as a function of the analysed spot (Figure 4a). Elemental analyses confirm the presence of tin (0.1-0.2 wt%) and chromium traces (0.01-0.1 wt%).

Turquoise Beads

No specific Raman signature was observed for turquoise beads (15-b, 17-b, 19-b) of Antsiraka Boira (e.g. Figure 2, the spectrum 17-b), in agreement with a dispersion of Cu²⁺ ion in an alkaline glass matrix (Eppler & Eppler, 2000; Colomban et al., 2001). The SEM-EDS elemental analysis confirmed the presence of copper (0.1-0.3% by weight of

Table 3: SEM-EDX oxide compositions of beads (wt%) measured using 100x magnification on the surface. Only major oxides are considered. Σ Flux = CaO + Na₂O + K₂O.

Tableau 3 : Compositions de surface (% pondéral en oxyde) des perles mesurées par MEB-EDX pour un grossissement de 100. Seuls les éléments majeurs sont pris en compte. Fondants : Σ Flux = CaO + Na₂O + K₂O.
oxide), and iron (0.3-0.8% by weight of oxide). The Fe$^{2+}$ ion gives a light blue colour whereas the Cu$^{2+}$ ion is the classic turquoise chromophore of alkaline glasses and enamels (Colomban & Schreiber, 2005). The Cr-Sn chromophore is detected in bead 19-b (Figure 3).

Black Beads

Representative Raman spectra of black beads are given in Figure 5. For almost all of the analyzed beads, the identified black pigment corresponds to the amber chromophore “Fe-S” used to produce amber glass (Schreurs and Brill, 1984; Prinsloo & Colomban, 2008). The resonant Raman spectrum of this pigment is characterized by a broad band around 415 cm$^{-1}$, which is superimposed on the spectrum of the glass. The position of this band and its components depends on the glass matrix, i.e. the direct environment of the Fe-S chromophore (Figure 5a) and of the laser line used (Figure 5b). In Figure 5b, examples of the Raman signatures of bead 21-n obtained with different excitation wavelengths are given. Depending on the wavelength of the laser, the intensity and position of the main Raman band of the chromophore Fe-S vary (Prinsloo & Colomban, 2008): the main peak at 413 cm$^{-1}$ under green excitation (532nm) moves to ~390 cm$^{-1}$ under red excitation (633 and 785 nm), and loses intensity (Prinsloo and Colomban, 2008; Prinsloo et al., 2011). The spectrum recorded with the 785 nm laser line allows access to the Raman signature of the bead’s vitreous matrix (SiO$_4$ $\nu_{max}$ = 1071 cm$^{-1}$, SiO$_4$ $\delta_{max}$ = 482 cm$^{-1}$, Fig 5b, spectrum 21-n at 785 nm).

Another black pigment has been detected in bead 13-n (Figure 5a), which corresponds to a manganese oxide characterized by the bands at 297, 480 and 633 cm$^{-1}$ (Julien et al., 2004) and confirmed by SEM-EDS analysis (1.8 wt% of MnO$_2$). A similar spectrum and a high Mn content was observed in Magoro black beads (Koleini et al., 2014; Koleini et al., 2016c). Precise identification of the oxide remains difficult, because of the high sensitivity of these (black) oxides to laser heating, as heating changing the degree of oxidation and allotrope. As with some yellow and red beads, the spectrum of Cr-Sn chromophore was identified in some spots of the black beads 13-n and 21-n. Analyses by SEM-EDS clearly illustrate chromium (0.02-0.2 wt%) and tin (0.09-0.1 wt%).

White beads

The Raman spectra of white beads (e.g. figure 6a) are typical of aragonite, an allotrope of calcium carbonate (CaCO$_3$) occurring in living matter, which exhibit characteristic peaks at 153, 208, 709 and 1084 cm$^{-1}$ (White, 2008).

Other Phases

Figure 6b presents the characteristic spectra of secondary crystalline phases encountered in glass matrices. The Raman spectrum 4-y, with peaks at 357, 440 and 1008 cm$^{-1}$ is cha-
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Figure 5: Representative Raman spectra of the black beads a) with 532 nm excitation and according to the excitation wavelength for the black bead 21-n.

Figure 5 : Spectres Raman représentatifs obtenus a) pour différentes perles noires sous 532 nm et b) pour la perle noire 21-n sous différentes excitations laser.

Figure 6: Raman spectrum (HE 532) of mineral phases present in a) shell bead (11-w) and b) glass matrices; 1-r: sodium-rich feldspar; 4-y: zircon.

Figure 6 : Spectres Raman représentatifs obtenus avec le spectromètre HE 532 des phases minérales : a) perle blanche (11-w, coquillage) et b) perles de verre, 1-r: feldspath sodique; 4-y: zircon.
racteristic of zircon, ZrSiO$_4$ (Raskovska et al., 2010). This mineral, a very stable nesosilicate, detected in the beads 4-y, 13-n and 21-n is present in many ancient rocks and products of their degradation (clays).

The Raman spectrum 1-r (290, 478, 506, 760 and 812 cm$^{-1}$) is caracteristic of a sodium-rich feldspar, albite NaAlSi$_3$O$_8$ (Freeman et al., 2008). It was identified in the beads 1-r and 21-n. The feldspars are commonly used as fluxes in the glass or ceramic production. All characteristic peak wavenumbers are summarized in Table 4.

**Glassy matrix**

The micro-Raman spectrometry of amorphous silicate phases has already been subjected of numerous studies for non-destructive identification of glass or enamelled objects (Colomban et al., 2003; Colomban et al., 2006a; Colomban & Treppoz, 2001; McMillan & Piriou, 1982 and references herein). The glass is a more or less polymerized SiO$_4$ network, with each tetrahedron sharing or not their oxygen ion. Ions lowering the melting point (mainly Na$^+$, K$^+$, Pb$^{2+}$ and Ca$^{2+}$) and the colouring ions are housed in the void spaces of the polymer network.

The Raman spectrum of an amorphous silicate is composed of two wide bands resulting from vibrations of the SiO$_4$ tetrahedron, a strong covalent entity and vibrationally well defined: in the middle frequencies the band centred around 500 cm$^{-1}$ corresponds to the symmetrical deformation vibrations ($\delta$ SiO$_4$); the second band around 1000 cm$^{-1}$ corresponds to the symmetric stretching vibrations of the tetrahedron (v SiO$_4$). Actually, the contribution of asymmetric stretching and bending modes can be neglected (Colomban & Prinsloo, 2009). The graphical representation of v SiO$_4$ vs. $\delta$ SiO$_4$ maximum wavenumber distinguishes types of glass (Figure 7) (Caggiani et al., 2014; Colomban et al., 2005b; Colomban et al., 2006a). It is noted that an increase in alumina (i.e. the network ionicity) or lead (i.e. the depolymerization) in the glass composition causes major

![Raman Spectra](image)

*Figure 7: (See colour plate X) a) representative Raman spectra of the main glass matrix: 18-y) depolymerized glass; 17-b) soda glass with a high alumina content (Al$_2$O$_3$ Na$_2$O); 10-n) soda glass; 20-r) soda corroded glass. b) Classification of glassy matrices of the beads according to the maximum position of the Raman stretching (v$_{\text{max}}$ Si-O) as a function of the maximum deformation position ($\delta_{\text{max}}$ Si-O). The areas delineated as typical of eras and backgrounds glass (see Figure 8). The colour of the beads and the specificities of each field are indicated. Symbols markers correspond to the classification of Wood.*

*Figure 7 : (Voir planche couleur X) a) Spectres Raman représentatifs des matrices vitreuses: 18-y) verre dépolymérisé; 17-b) verre sodique riche en alumine (noté Al$_2$O$_3$ Na$_2$O); 10-n) verre sodique; 20-r) verre sodique corrodé. b) Classification des matrices vitreuses à partir des positions des maxima du massif Raman des modes d’élongation (v$_{\text{max}}$ Si-O) et de déformation ($\delta_{\text{max}}$ Si-O). Les zones des principaux types de verre sont indiquées (cf. Figure 8). Les couleurs et les principales caractéristiques sont notées. Les symboles correspondent à la classification morphologique de Wood.*
wavenumber shift (Colomban et al., 2006b; Tournié et al., 2008). Corrosion phenomenon with the protonation/leaching may also be responsible for lowering the bending and stretching mode wavenumber (Tournié et al., 2008).

Consistent with the results of the elemental analysis (Figure 7, Table 3), almost all beads are made of soda glass. Na$_2$O content ranges between 12% and 18%, except for beads #7 (11.8%) and #9 (9.8%). K$_2$O content ranges between 1.5% and 2.8%, except for black beads #7, 10 and 21 exhibiting lower values (1.2%-1.4%). Only a spectrum corresponds to a more highly depolymerized glass (18-y, corrosion pits are visible in Figure 1) and a bead belongs to the soda-lime-type (13-n, stained with manganese oxide). The wound bead 10-n is located outside the dispersing zone of other soda beads in accordance with its particular process. The deformation band maximum ($\delta_{\max}$ SiO$_4$ at 470-500 cm$^{-1}$) and the elongation band maximum ($\nu_{\max}$ SiO$_4$ around 1080 to 1090 cm$^{-1}$) are typical of a sodium silicate glass rather rich in Al$_2$O$_3$. The 17-b spectrum (Figure 7a) is typical of a highly polymerized soda glass, with three bands at 335, 495 and 576 cm$^{-1}$ and two very intense stretching components at 1090 cm$^{-1}$ and another of medium intensity at 990 cm$^{-1}$. The spectrum 13-n of soda-lime type is characterized by a band at 554 cm$^{-1}$, and three stretching components at 945, 992 and 1082 cm$^{-1}$. The data is somewhat scattered but it is possible to identify subgroups. The dispersion of structural changes is due to compositional variations, linked to variable raw materials used, or to the implementation of technology (puriﬁcation / selection of raw materials). The beads "speciﬁcity" in terms of composition of the glass matrix is also found in terms of chromophore (13-n and 18-y) or manufacturing technique (10-n).

The examination of the Raman proﬁles brings a number of additional information. In Figure 7a, the representative Raman signatures of the main glass matrices, demonstrate that in spite of their belonging to the same group (17-b, 10-n and 20-r) differences are visible. The 20-r spectrum is characteristic of a corroded glass. This alteration is attributed to a chemical reaction between the glass surface and an acidic aqueous solution causing the exchange of K$^+$ and Na$^+$ / Mg$^{2+}$ ions by protonic species: moving the $\nu_{\max}$ SiO$_4$ to a lower wave number (1072 cm$^{-1}$) corresponds to an elongation of the Si-O bond due to the interaction of oxygen atoms with the protonic species (Tournié et al., 2008).

The analysis by SEM-EDS was performed to complete the classiﬁcation obtained by Raman spectroscopy, and identify chromophores which do not give speciﬁc Raman signatures. Most beads have a relatively high content of Al$_2$O$_3$ (6 to 9.5 wt%, Table 3, in agreement with the original Indo-Paciﬁc beads (Dussubieux et al., 2010; Robertshaw et al., 2010), reﬂecting the use of a ﬂux of mineral type. These compositions are shown graphically in ternary diagrams (Figure 8). The ternary diagram (SiO$_2$-Al$_2$O$_3$-Σ ﬂuxes) in the Figure 8a & a’, is sensitive to both raw materials but also to technology because of the direct correlation between the amount of ﬂuxes and of the operation temperature. This is to distinguish four sub-groups:

- A large majority of beads from Antsirak Boira consists of a ﬂux-rich glass (~ 20 wt%).
- Some black beads have clear inclusions and/or corrosion of red dots (9-n, 7-n, 21 n), and the blue bead 16-b contains less flux (~ 15 wt%).
- Bead 17-b appears isolated, with a very alumina-rich glass (Al$_2$O$_3$ ~ 13 wt%).
- Beads 13-n and 15-b are richer in ﬂuxes than all the other beads (~ 23 wt%). The main difference of this black bead with those recovered from South-African Magoro Hill site is its higher Al and less Mn content.

The ternary diagram of ﬂuxes (Na$_2$O-CaO-K$_2$O, Figure 8b & b’) refines the visualization of differences. The homogeneity of a glass notably depends on the temperature and heating time, the agitation of the melt, etc. Although all of the beads seem to be composed of a soda glass (Figure 7b), two sub-groups are emerging:

- The first consists of a glass matrix rich in sodium, which brings together the red beads (1-r, 2 r, 14 r, 20-r and 22-r) and a large majority of black beads (5-n, 6-n, 7-n, 8-n, 10-n, 12-n, 21-n).
- The second group consists of beads made of a glass richer in calcium (3-y, 9-n, 13-n, 17-b).

The comparison with literature data, presented in the ternary diagram of “Flux” (Figure 8 a’ & b’), conﬁrms the beads of this assemblage belonging to the group of “Indo-Paciﬁc” soda glass beads.

4. DISCUSSION

We will highlight the identiﬁed characteristics of each family of bead, those features that are useful in the discussion of origins and dating. Table 4 lists the elements identiﬁed, pigments or dyes, and highlights the speciﬁc characteristics.

Yellow beads

Raman spectroscopy demonstrates for yellow beads 3-y and 4-y (Figure 3c) the use of a pyrochlore solid solution based on Pb$_2$Sb$_2$Sn$_2$Z$_3$O$_{13}$ solid solution, also called Naples Yellow (Colomban et al., 2001, Kirmizi et al., 2010, Perreira et al., 2009; Rosi et al., 2009). The presence of lead is confir-
med by elemental analysis (0.1 to 0.2% by weight) with the exception of the 18-y bead which is colored by an iron yellow (~1.8 wt%) whose orange tone is due to the presence of the chromophore Cr-Sn, identified by Raman scattering. The elemental analysis showed the presence of tin traces (Sn 0.05 wt%). Naples Yellow solid solutions have been used in the Mediterranean area and Mesopotamia since Antiquity (Colomban, 2013) but the presence of Cr-Sn chromophore was not reported in the previous Raman studies.
**Red beads**

The element identified in red-brown beads is the metallic copper nanoparticles. The Cu content varies from 0.05 to 0.2 wt%. In agreement with this type of coloring agent, Raman spectra of the glass matrix are very variable and difficult to obtain, as expected for glasses containing metallic nanoparticles (Colomban & Schreiber, 2005). Since the Neolithic, the red color of the lead-based glasses is achieved in Europe by the dispersion of copper nanoparticles (Cu⁰) in the glass matrix (Colomban et al., 2009) in a reducing environment. This practice has been widely used in the Roman period (Ricciardi et al., 2009a) and in some Islamic productions. Production in China was also suspected (Beaujard, 2012). The joint observation of Cr-Sn chromophore is also a very special feature of the studied glass.

**Blue and turquoise beads**

No blue pigment (Co₃SiO₄, CaAl₂O₄, Lapis lazuli) was detected with Raman spectroscopy. The lack of the above mentioned signature is consistent with the color obtained by a small amount of cobalt dissolved in its ionic form in the glass. Elemental analyzes by SEM-EDS show a very low cobalt content, ~0.01 wt% for the bead 16-b, in agreement with the light blue colour of the bead (less than 0.5 wt% of cobalt is sufficient to give an intense blue colour). The turquoise color (beads 15-b, 17-b & 19-b) is the classical result of the dissolution of Cu²⁺ ions in an alkaline silicate glass matrix (Colomban & Schreiber, 2005). Elemental analyses indicate Cu content between 0.1 and 0.3 wt%. Iron detected by elemental analysis, is probably an impurity with no significant role in the color of matrix. So no specific markers for these beads were found, except the Cr-Sn chromophore present in the bead 19-b.

Table 4: Raman fingerprints of pigments, chromophores and crystalline phases were in Mayotte beads. (IP: Indo-Pacific. MO: Middle East).

<table>
<thead>
<tr>
<th>Pigments / Coloring agents / Phases crystalline</th>
<th>Bead color</th>
<th>Phases</th>
<th>Raman peak cm⁻¹</th>
<th>Chromo-phrase</th>
<th>Period Origin</th>
<th>Origin Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pyrochlore Solid solution</td>
<td>Yellow</td>
<td>Pb₂Sb₂ₓSnₓO 137-144</td>
<td>Pb</td>
<td>1293-1570 av. J.-C. MO / IP</td>
<td>Kirmizi et al., 2009</td>
<td></td>
</tr>
<tr>
<td>Amber</td>
<td>Black</td>
<td>Chromophore Fe-S 415-420</td>
<td>Fe. S</td>
<td>I-P</td>
<td>Prinsloo et Colomban, 2008</td>
<td></td>
</tr>
<tr>
<td>Albite (Na-feldspar)</td>
<td>Black Red-brown NaAlSi₃O₈ 290-478. 506</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Copper</td>
<td>Blue Red</td>
<td>Cu²⁺ Cu⁰ no</td>
<td>Cu</td>
<td>Europe. MO IP</td>
<td>Colomban et Schreiber, 2005</td>
<td></td>
</tr>
<tr>
<td>Pyrolusite</td>
<td>Red Black</td>
<td>β-MnO₂ 280-490. 630</td>
<td>Mn</td>
<td>Julien et al., 2004</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Iron ions</td>
<td>Blue Yellow</td>
<td>Fe²⁺ Fe³⁺ no</td>
<td>Fe</td>
<td>IP</td>
<td>Colomban, 2004</td>
<td></td>
</tr>
<tr>
<td>Sphene (+anastase)</td>
<td>Black Red Yellow Chromophore Sphène Sn (+TiO₂) 147-780</td>
<td>Ti. Sn (&lt; 0.2%)</td>
<td></td>
<td></td>
<td>Faurel et al., 2003</td>
<td></td>
</tr>
<tr>
<td>Calcium phosphate</td>
<td>Black Red Yellow P-O 950-960</td>
<td>/</td>
<td>MO</td>
<td>Ricciardi et al., 2009</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Zircon</td>
<td>Black Yellow ZrSiO₄ 358-440. 1008</td>
<td>Zr (21-n)</td>
<td></td>
<td></td>
<td>Tournié et al., 2009</td>
<td></td>
</tr>
<tr>
<td>Aragonite shell</td>
<td>shell CaCO₃ 155. 207. 704. 1085</td>
<td></td>
<td></td>
<td></td>
<td>White S.C., 2008</td>
<td></td>
</tr>
</tbody>
</table>

Black Beads

The main identified coloring agent by Raman scattering in black beads, is the “Fe-S” amber chromophore. In parallel with the morphological and chemical classifications, the presence of “Fe-S” chromophore is in agreement with black Indo-Pacific beads, which are supposed to originate from the south of India, Sri Lanka or South-East Asia (Dussubieux et al., 2008; Prinsloo & Colomban, 2008; Prinsloo et al., 2011). Accordingly, beads with variations in the position and intensity of the main Raman bands (Figure 5a) may indicate various origins or periods/production qualities.

The 13-n seems morphologically consistent with Indo-Pacific series. Nevertheless, and contrary to the morphological criteria, the lack of “Fe-S” chromophore shows another function of composition, but also depends on other factors (temperature, manufacturing process etc.). Therefore, the same material composition may not have the same structure and vice versa. Furthermore, the SEM-EDS analysis made with different magnifications shows a significant differences in the local composition.

Soda glass is the constituent of all Indo-Pacific beads (type K2, East-Coast or Khami) dated according to Wood between the 10th and the 17th century (Robertshaw et al., 2010; Wood, 2005; Wood, 2011; Wood et al., 2009). The dispersion within the Na2O group may result in variations between compositions K2 groups, EC-IP and Khami due to the variability of raw materials but also of the surface alterations of some beads. So the red beads (1-r, 2-r, 14-r, 20-r and 22-r) belong to the soda group but the majority of Raman spectra are typical of a slightly corroded glass (whatever the elimination of surface film by grinding). The easy oxidation of metallic copper used for colouring (Colomban & Schreiber, 2005; Ricciardi et al., 2009a) may be the source of significant corrosion for these red beads. The bead 17-b, outside of the main soda group presents a characteristic Raman signature similar to that of the K2 series (Prinsloo et al., 2011; Koleini et al., 2016a; Koleini et al., 2016c). The use of soda glasses is common in Southeast Asia/China (16th century enamels), Venice and Holland (16th and 17th century A.D.) (Ricciardi et al., 2009b; Robertshaw et al., 2010; Dussubieux et al., 2010; Kirmizi et al., 2010; Colomban et al., 2006; Tournié, 2009; Prinsloo et al., 2011; Tournié et al., 2011) and this criterion alone does not qualify objects.

The presence of albite is consistent with a poor refining of molten glass (albite, chalk and silicates form liquid phase below 900°C), and zircon a geological marker of raw materials. These minerals were detected in opaque beads, red-brown or black, which could then be categorized as very heterogeneous beads, closer to a molten ceramic. The same result was also obtained for some reddish beads of the Vohemar necropolis dated from 12th-17th century, north of Madagascar (Colomban & Simsek, 2014). SEM imaging supports this hypothesis for the red and black cut beads, also revealing a highly porous matrix typical of unrefined molten material (Fischbach, 2015).

White beads

The white beads are made of aragonite, the allotrope of most limestone shells or corals. Visually similar beads were found on the African coastal sites, Tanzania (the ruins of Kaole) (Wood, 2012) and at Chibuene, south of the Mozambique Channel (Wood, 2012). Other aragonite beads were also identified at Mahilaka and Sandrakatsy, Madagascar, two sites dated between the 9th and 15th century A.D. (Robertshaw et al., 2006). Shells of *Tridacna maximia* were found as offerings in burials on Antsiraka Boira. The inorganic compound of this shell is aragonite (Romaneck et al., 1987). Since it is the most abundant species found in the archaeological levels around Acoua, these beads may well be locally manufactured, although no contemporary Mayotte site show traces of this production. In contrast, bead polishers were found on the Lamu Archipelago’s Kenyan sites (Horton, 1996; Pauly, 2014) demonstrating this type of local production. These beads could thus be of African origin.

Glass matrices

Subgroups that were identified by Raman analysis coincide well with those identified by the ternary composition diagrams (Figure 8a & b), but dispersions within each groups is different. Indeed, the structure of the network of SiO₄ tetrahedron forming the glass skeleton is not only a
The beads’ compositional data for different sites in South Africa shows a good correlation between Raman signatures and compositions (Figure 8a’ & b’). The classification of Antsiraka Boira beads on the basis of Wood morphological classification system appears almost correct. The beads of Indo-Pacific series (East Coast-IP, K2-IP, Khami-IP), Mapungubwe Oblate and Islamic (South Africa), have a composition richer in Al₂O₃ than beads of Zhizo series. The series of Islamic, Mapungubwe Oblate and Zhizo have a high CaO content in comparison with Indo-Pacific beads. Dispersal areas of South Africa beads on the diagrams of Figures 7 and 8 belonging to the Indo-Pacific and Khami...
series match those beads of Antsiraka Boira, supporting the first conclusion of the Wood’s morphological classification (Figure 1). However, there are compositional variations within the series (Figure 8, compatible with an inherent inhomogeneous quality of a non-industrial production), and allocating a bead to a specific series on the basis of its morphological features or its composition alone is not always clear cut, especially for Indo-Pacific series. The Cr-Sn sphene Raman signature (Table 6) in many beads offers a tool to identify this subgroup of Indo-Pacific production.

A comparison with the beads of Mapungubwe and K2 sites

Figure 9 compares the distribution of many beads’ glass matrices from the Mapungubwe and K2 sites – two twin sites in the upper valley of the Limpopo River near the present meeting point of South Africa, Botswana and Zimbabwe (Prinsloo et al., 2011). The diagram constructed on the Raman parameters in analysing the structure of the polymeric network of SiO$_4$ tetrahedron. Almost all of the beads from the Antsiraka Boira and K2 sites are soda (Na$_2$O), while a significant portion of Mapungubwe beads belong to soda-lime group (Na$_2$O/CaO).

A wider dispersion was recorded between Na$_2$O and Na$_2$O/CaO groups for Mapungubwe beads, while most of K2 beads belong to the Na$_2$O group. The variety in glass structure can be explained by a much longer occupation of Mapungubwe (11$^{th}$-19$^{th}$ century), which could be in accordance with many supplies origins and multiple technologies used, compared to K2 site and its shorter period of occupation (9$^{th}$-13$^{th}$ century). Mayotte is much comparable to the contemporary K2 site for both short period of occupation and a more uniform glass structure. Mapungubwe Na$_2$O/CaO glass bead also have less consistency in composition than Na$_2$O group which is in accordance with the various sources and technologies used. K2 and Antsiraka Boira beads also have not much variety in pigments than those reported for Mapungubwe beads (Prinsloo et al., 2011; Tournié et al., 2011).

5. CONCLUSION

The use of parameters extracted from the Raman signature of matrices, allowed classifying Antsiraka Boira glass artefacts into a main group of Indo-Pacific soda glass, similar to that of the contemporary site of K2 in South Africa. Elemental analyses with SEM EDS certified the Raman results. Differences in the morphology of the beads are however clear. The peculiarity lies in red and some of the black beads, which seem to be closer to a molten ceramic than to a glass, akin to some beads from the Vohemar necropolis in the north of Madagascar (14$^{th}$ to 16$^{th}$ century necropolis). The used coloring agents are generally conventional: copper red, “Naples Yellow” – a pyrochlore solid solution, the amber/black Fe-S chromophore, and Cu$^{2+}$ ions in the turquoise color. Unexpectedly in the study of these beads, resonance Raman spectroscopy has identified the presence of a Cr-Sn sphene chromophore. This pigment has been detected in several different beads: yellow, black, red, and turquoise; the presence of chromium and tin elements being confirmed by SEM-EDS analysis. This Raman signature is a marker of the origin of these glasses. The chromophore was not used as a colouring agent.

The Indo-Pacific beads of the Antsiraka Boira necropolis open a long period of overlap with other sites to determine the location of production facilities. A comparative study is necessary, especially to investigate whether the Cr-Sn sphene chromophore is present.

Despite the lower spectral resolution of the portable device, and its strong complex background, most of the
measurements performed with the portable Raman spectrometer led to significant and usable Raman spectra (Table 4). For crystalline compounds such as pigments, the characteristic Raman bands are narrow, often intense, and easily detectable on the Raman spectrum (Figure 2, 3-y). Therefore, the spectra obtained with the portable device allow easy identification of almost all crystal phases. Regarding the glass matrices, portable Raman analysis of yellow and turquoise beads is more difficult but sufficient in numerous measures. However, for red and black beads the useful signal can be very small compared to the background, especially if fluorescence is superimposed. The latter is very common for porous archaeological samples due to the biological film’s degradation at the oxide surface. In this case, laboratory fixed instruments are better suited because of their better accuracy and overall performance. Blue, violet or UV energy excitation eliminates the biofilm causing fluorescence very easily and in a few minutes of irradiation.

The use of mobile equipment that can be implemented on-site overcomes the technical and ethical constraints related to the subject, offering the possibility of direct and rapid, in situ identification, without having to move / export objects. Thus, expanding the areas of intervention and the amount of accessible objects, and resulting in the acquisition of richer and more representative data. However, these portable devices are less efficient due to the devices’ miniaturization which leads to low resolution, continuous background… and may – by degrading the quality of data – cause inappropriate conclusions.

**Acknowledgements**

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Beads Excavated from Antsiraka Boira Necropolis (Mayotte Island, 12th-13th centuries)...


Figure 1: Noémi Fischbach et al., Beads Excavated from *Antsiraka Boira* Necropolis (Mayotte Island, 12th-13th centuries) Colouring Agents and Glass Matrix Composition Comparison with Contemporary Southern Africa Sites (p. 86)

Figure 7: Noémi Fischbach et al., Beads Excavated from *Antsiraka Boira* Necropolis (Mayotte Island, 12th-13th centuries) Colouring Agents and Glass Matrix Composition Comparison with Contemporary Southern Africa Sites (p. 92)
Figure 8: Noémi Fischbach et al., Beads Excavated from Antsiraka Boira Necropolis (Mayotte Island, 12th-13th centuries) Colouring Agents and Glass Matrix Composition Comparison with Contemporary Southern Africa Sites (p. 94)