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Archaeometric evidence attesting production of indigenous archaic pottery at Monte Polizzo (Western Sicily)

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Abstract

Excavations at the proto-urban indigenous settlement of Monte Polizzo (western Sicily) have not yielded so far any evidence of in-situ ceramic production (i.e. kiln structures). However several archaeological concerns put forward to consider it as a likely production centre of pottery during the Archaic age. In this paper a first attempt to check the compositional correspondence between ceramic fabrics and local clay sources has been made. A comprehensive archaeometric investigation of indigenous pottery, mainly composed of matte-painted table ware dated from the 7th to the 4th century B.C., recovered from the Acropolis of Monte Polizzo has allowed the identification of six distinct "Paste Compositional Reference Units" (PCRUs). The six PCRUs were independently identified and confirmed through textural and mineralogical characteristics (petrographic observations) and chemical characterization (XRF analysis).

A parallel investigation of clayey raw material of Monte Polizzo's natural resources revealed two geologic formations the Terravecchia and Marnoso-Areacea del Belice (MAB). Experimental ceramic briquettes were created and petrographically and chemically characterized. A comparison between the indigenous and experimental ceramics revealed that the local potters preferred clays from the Terravecchia Formation.

The mineralogical, textural and chemical composition of the raw clays of the Terravecchia Formation resulted to be well correlated with the 92% of the indigenous ceramic artefacts recovered from Monte Polizzo. The remaining 8% are interpreted to be imported from the Sicani Mounts area.

Key words: Western Sicily; Archaic age; indigenous ceramic production; incised and matte painted tableware; thin section petrography; XRF.

Introduction

Monte Polizzo was a proto-urban indigenous settlement, dating back approximately to the 9th

century B.C., active till the end of 4th century B.C., frequented again in the Middle Ages and occasionally in more recent times.

The current archaeological site lies only six kilometres northwest of Salemi, Trapani province, at 726 m above sea level, dominating a vast area of western Sicily and controlling the major trade routes to Selinunte, the greatest Greek colony in the area and Segesta, the most important Hellenised indigenous centre. The topography of the site shows clear analogies with other indigenous settlements in archaic western Sicily (Erice, Segesta, Entella, Monte Iato, Monte Adranone, Monte Maranfusa just to list the most important ones), as it is located on top of a hill in a suitable position for defence and territorial control (Figure 1). The correspondence concerns material culture as well testifying for their belonging to a unique and homogeneous civilization (Spatafora, 1996).

Investigations at Monte Polizzo can therefore spread new light on a wider area in Western Sicily that lies out of colonial (Greek and Phoenician) territories and that retained the peculiar features of indigenous civilization in archaic and classical times. First study on Monte Polizzo took place in the early 70's (Tusa, 1972). Later, several indepth field surveys and excavations were carried (1998-2010)within the "Sicilian Scandinavian Archaeological Project" (SSAP), which the Universities of Göteborg and Oslo, as well as the American Universities of Stanford, Northern Illinois and Northern Iowa are part of (Johansson and Prescott, 1999; Prescott et al., 2001; Morris et al., 2001; Kolb and Tusa, 2001; Mühlenbock and Prescott, 2001; Morris et al., 2002; Morris et al., 2003; Morris et al., 2004; Morris and Tusa, 2004). The excavations are being carried out not only on Monte Polizzo (Acropolis, House 1, Portella S. Anna) but also in the historical center of Salemi and in the site of Pizzo Mokarta (not far away).

Notwithstanding a considerable amount of ceramics imported from the Greek world (possibly through the colony of Selinus) and, on a lesser grade, from the Phoenician area, the main ceramic assemblage consists mostly of

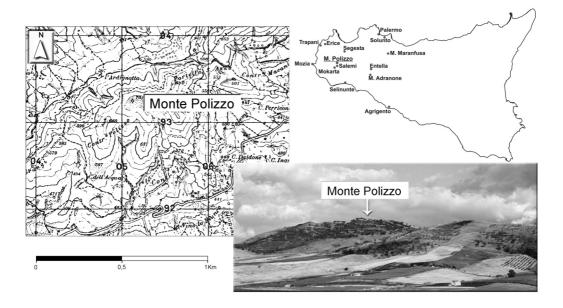


Figure 1. Map of western Sicily marking the positions of Monte Polizzo, Mokarta, Salemi, and some relevant archaeological sites.

storage and tableware decorated with incised or painted geometric patterns. This pottery is characteristic of indigenous settlements in central and western Sicily and it has even been used as marker of 'Elymian' culture, though this issue is still debated (Spatafora, 1996). The retrieving of these ceramics at Monte Polizzo is therefore obvious, but it does not give enough information about technical, productive and commercial aspects of the Monte Polizzo settlement. In fact, nowadays, nothing can be inferred about the existence of ceramic workshops at Monte Polizzo. For that reason, several important socio-economic questions, such as the capability in facing local needs or in spreading their ceramic items on a larger market, cannot be answered. Monte Polizzo differs therefore from sites like Entella (PA), where kilns, furnace separators and misfired sherds of archaic age were found, testifying for a local production of cooking ware, plain ware and vases decorated in a simple geometric style (Guglielmino, 2000). As a consequence, at Entella it was possible to clearly identify local products also following their spreading in the neighborhoods and to appreciate the ratio local vs. imported ceramics (Montana et al., 2011a).

An attempt in relating ceramics from Monte Polizzo to other fabrics in western Sicily was made in a preliminary archaeometric study involving several pottery samples collected at Monte Polizzo, published five years ago by Kolb Speakman (2005). They analyzed a consistent number of Archaic table ware sherds (9th to the 6th century B.C.) collected from 12 indigenous settlements all located in western Sicily by means of instrumental neutron activation analysis (INAA), with the aim of studying the ceramic production centers and the exchange patterns. The chemical data obtained from INAA were evaluated using an array of multivariate statistical procedures in order to facilitate the identification of compositional groups. Up to five *chemical groups* were clearly

identified suggesting an intra-regional organization of ceramic trade in western Sicily, however the source locations of raw clays and the location of the correspondent ceramic workshops remained uncertain (Kolb and Speakman, 2005). More recently, 20 ceramic samples from the areas A, B and C of Monte Polizzo Acropolis, dated to the mid 6th century B.C. and stylistically attributed to local indigenous ware were analyzed by thin section microscopy and ICP-AES (Brorsson, 2007). Up to 7 groups consisting of different temper materials and/or microfabric were described even if chemical analyses suggested only small compositional variations for the majority of the samples (Brorsson, 2007). Nevertheless, in this case the hypothesis of provenance was debated on the individual samples and the compositional and textural fingerprints of local ceramic products were only approximately described and not evaluated on a geo-lithological basis.

Such a 'knowledge plateau' concerning the Archaic ceramic production in the site of Monte Polizzo is not surprising considering that the assignment of a ceramic paste (macroscopic/microscopic fabric) to a specific workshop can be easily addressed only if kiln wasters are available. However, if we consider a consumption center the situation is radically different. In this case, a *paste compositional reference unit* or PCRU can be only formed. It can not be related to a particular workshop unless by association with already established compositional groups (Bishop et al., 1982; Rice, 1987; Buxeda et al., 1995)

Nevertheless, the comparison of the *paste* compositional reference units with local geology and clayey raw materials available in the territory can be an alternative way to establish if the settlement of Monte Polizzo - within a specific chronological interval (7th - 4th century B.C.) - was simply a centre of consumption, importing ceramics from other indigenous and coeval sites, or if - as it seems possible according

to the archaeological data acquired so far - it was itself a centre of production, and to discriminate between autochthonous or allochthonous ceramic productions.

Thus, the awareness of the ceramic raw materials available in a given territory may be a correct approach for an accurate definition of ceramic provenance, by determining the mineralogical and chemical composition and also some critical textural aspects of clays (i.e. packing and size distribution of aplastic grains) and comparing them with materials retrieved from archaeological excavations (e.g. Magetti, 1982; Picon, 1992; Degryse et al., 2003; Gliozzo and Memmi Turbanti, 2004; Hein et al., 2004; Gliozzo et al., 2005; Tite, 2008).

This paper provides a detailed petrographic and chemical investigation of ceramic finds recovered from excavations at Monte Polizzo. A set of 44 ceramic samples were carefully selected on an autoptical basis by the archaeologists of the American mission. The sherds were mostly representative of small amphorae, bowls, jugs, vessels and other less distinct forms that may be classified as 'tableware' from a functional point of view. These samples were studied by optical microscopy (thin sections) and X-ray fluorescence spectroscopy (XRF). archaeometric investigation of ceramic sherds was carried out in parallel with one on the raw clayey materials cropping out in the studied area (6 samples). The textural and compositional comparison between the potential raw materials and the ceramic artefacts is not easy because of the analogies, shown by the geo-lithologic contexts of the already recognized production centres situated in western Sicily (Montana et al., 2003; Montana et al., 2007; Montana et al., 2009; Montana et al., 2010; Montana et al., 2011b).

Geological outline and local raw clays

From a geologic point of view, the adjacent hinterlands surrounding Monte Polizzo and

Salemi are characterized by a discontinuous sedimentary record dating from the late Triassic to the present-day. Common bedrock types include limestone, calcareous marl, shale, siltstone, sandstone, conglomerate, calcarenite, evaporitic limestone, selenitic gypsum, loose sand and clay assemblages that reveal deep to shallow marine environments and younger transitional alluvial sedimentation (Catalano and D'Argenio, 1982; Tortorici et al., 2001). Clays beds suitable for making ceramics and outcropping close to the site of Monte Polizzo (Figure 2) are those belonging to the Upper Tortonian - Lower Messinian Terravecchia Formation (Ruggeri and Torre, 1984; Butler et al., 1999) and to the Middle-Upper Pliocene Marnoso-Arenacea del Belice (MAB) Formation (Roda, 1967; Ruggeri and Torre, 1974). The Terravecchia Formation is a sequences of fluvial deltaic deposits up to 2500 m thick forming a molassic package of clastic sediments. It represents a major sedimentation period throughout Sicily with some stratigraphic unconformities. The MAB Formation is a terrigenous succession that lies discordantly on older deposits forming a characteristic polyphase thrusting (piggy-back). The MAB also contains clay-beds that contain significant concentrations of calcareous microfauna.

Terravecchia and MAB clayey raw materials are abundant in western Sicily and they have been recently characterized in terms of grain size distribution, mineralogical and chemical composition (Montana et al., 2006; Montana et al., 2011b).

Materials and methods

Description of ceramic samples and clays

Forty-four ceramic samples were subjected to textural and compositional analytical methods. These samples were recovered from Stanford University's excavations, under the direction of Dr. Ian Morris, on Monte Polizzo's Acropolis.

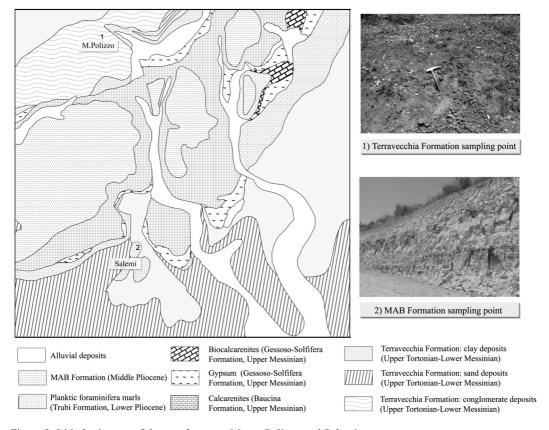


Figure 2. Lithologic map of the area between Monte Polizzo and Salemi.

Dr. Emma Blake (Tufts University) organized the excavation's recovered material culture. Dr. Blake by means of careful naked-eye observations identified three major macrofabric groups: F24, F7 and F54 (Table 1). These groups were schematically classified into chronological groups (detailed descriptions of the different chronological phases are provided in Morris et al., 2004), macrofabric, typology and surface treatments.

F24 macrofabric (3 samples) was by far the least common in the studied set of ceramic sherds (Blake, personal communication). F24 represents diagnostic undecorated/handmade large storage

vessels, *pithoi*, (average thickness between 15-20 mm). This macrofabric was also characterized by a temper frequency greater than 10% (area) as well as the presence of coarser sand grains (maximum grain size, MGS, up to 3 mm).

Macrofabric F7 (Figure 3a) was well-represented (18 samples) in the available set of ceramic samples. F7 samples represent smaller diagnostic wheel-made tableware forms (e.g. bowls) having the external surface decorated with incised geometric patterns (mainly horizontal lines). F7 is classified as a fine to very-fine paste (MGS around 0.5 mm) with a greyish surface colour, average thickness up to 5 mm and temper

Table 1. Chronology, macrofabric (see text), type, thickness and surface treatment of the studied ceramic samples from Monte Polizzo.

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	PLZ30		F 54	closed vessel		
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	PLZ45	7 th -6 th century B.C. (Phase I-II)	F 54	table ware	7	matte painted

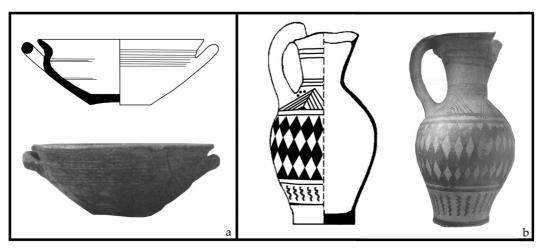


Figure 3. Examples of the indigenous matte painted ceramics found at Monte Polizzo (Acropolis) and others sites of western Sicily: a) bowl with incised decoration; b) table amphora with matte painted geometric decoration.

packing always less than 5% (area). The macrofabric F54 (Figure 3b) was the most frequent within the studied ceramics (23 samples) and characteristic of matte painted (geometric decoration) wheel-made tableware (e.g. table amphorae). It presents mainly very fine-fine sand grains (sporadic medium-coarse) with MGS up to 1 mm. F54 is also characterized by a pale reddishyellow surface and a greyish core. The average thickness was about 6 mm and the temper packing more abundant than the 5% (area).

Potential local ceramic raw materials representative of the Terravecchia Formation were found along the slopes of Monte Polizzo with complications mainly due to the rarity of exposed clay beds under soil horizons and the presence of thick vegetation. A silty-clay (sample SALT 11: sand content around the 13% wt) and a silty-clayey sand (sample SALT 10: sand content around the 55% wt), the latter utilizable in its natural state only for mixing and tempering purposes, were collected. Concerning the MAB Formation, because of the extensive occurrence of cultivated fields in its outcrop area, 4 uncontaminated samples composed of silty

clayey material (SALT1, SALT3, SALT4, SALT6: mean sand content around 15% wt) were collected in the urban centre of Salemi in a clean and deep excavation for a building foundation. We are aware that the number of the collected clay samples, even considering the above stated logistic difficulties, could be considered inadequate to represent well enough both chemical and mineralogical composition of local clays, specially if the analysis are aimed to make a distinction between handmade artefacts. However, it has to be considered that a large number of analysis have been carried out recently in the whole territory of western Sicily on both Terravecchia Formation (159 clay samples) and MAB Formation (48 clay samples), many of which refers to clay samples collected within a radius of 50 km respect the site of Monte Polizzo, being now available and actually significant for comparative purposes (Montana et al., 2011b).

Analytical methods

Clay briquettes of the locally sampled raw materials (small bars of 9 cm width, 2 cm length

and 0.5 cm thick) were moulded and fired in a muffle furnace at 900 °C with soaking time maintained for 3 hours. Furthermore, the Terravecchia Formation clay raw materials, due to their varying sand concentrations, were used to prepare an additional briquette (coded MIX1): 3 parts in weight of SALT11 were mixed with 2 parts of SALT10.

Microscopic observations of thin sections were performed by a Leica DM-LSP polarizing microscope equipped with a digital photographic camera. The ceramic artefacts were characterized in terms of the mineralogical and textural features of their aplastic inclusions and accordingly categorized into *fabrics* if showing apparent similarities. The experimental/fired briquettes were petrographically assessed using the same methods used to characterize the artefacts from the ceramic manufactures.

The chemical composition of ceramic samples and local raw clays were determined using a Rigaku ZSX Primus wavelength dispersive Xray fluorescence spectrometer (XRF), equipped with four diffracting crystals: LiF (200), LiF (220), PET (002), and TAP (100). A small portion of powdered and homogenised samples (around 2 g) was dried at 100°C for 24 h. Each specimen, previously mixed with two drops of a 3% water solution of Mowiol N50-98, was compressed on a base of ultrapure H₃BO₃ at about 5 tons/inches² by a hydraulic press to obtain a circular powder pellet (4 cm in diameter). A Rh tube anode routinely running at 4 kW was used for the major elements analysis as well as for minor and trace elements. The quantified elements (major and minor elements recorded as oxides, wt.% while trace elements given as ppm) were the following: Fe₂O₃ (as total Fe), Al₂O₃, MgO, MnO, P₂O₅, TiO₂, CaO, Na₂O, K₂O, SiO₂, V, Cu, Zn, Rb, Sr, Zr, Ba, La, Ce. The loss on ignition (LOI) was determined by firing at least 0.5 g of dried specimen, at 1000 °C for 3 h. Elemental concentrations were quantified using calibration lines up to 52

standard reference materials (SRM). The average relative standard deviations evaluated (RSD) on seven runs of the SRMs IAEA/Soil-7, IAEA/SL-1, NIST/BC-679 were: $SiO_2 = 0.64\%$, $TiO_2 = 0.97\%$, $Al_2O_3 = 0.89\%$, $Fe_2O_3 = 0.92\%$, MnO = 1.22%, MgO = 1.38%, CaO = 1.19%, $Na_2O = 1.91\%$, $K_2O = 0.83\%$, $P_2O_5 = 1.10\%$, V = 2%, Cu = 2%, Zn = 1%, Rb = 2%, Sr = 1%, Zr = 3%, Sr = 4%, Sr = 4%, Sr = 1%.

The analysis of each element was fully corrected for line interference and matrix effects of all the other analysed elements and the results were normalised on a dry-weight basis (Hein et al., 2002).

Multivariate statistic methods such as Principal Components Analysis (PCA) and Multiple Correspondence Analysis (MCA) were carried out through the commercial software S-Plus (Mathsoft 1999).

Results and discussion

Petrography of the ceramic samples

The petrographic analysis of the 44 ceramic samples from Monte Polizzo's Acropolis has facilitated the identification of six *fabrics* (FBRs). The essential characteristics: mineralogical composition, packing, sorting and grain size, for every FBR, were schematised and summarized.

FBR 1a (samples: PLZ1; PLZ4; PLZ15; PLZ18; PLZ19; PLZ22; PLZ25; PLZ28; PLZ29; PLZ31; PLZ33; PLZ34; PLZ37; PLZ44; PLZ46)

It shows a clear prevalence towards a sandy/silicoclastic temper rather than carbonatic, which is nearly absent or not significantly represented, like in the case of sample PLZ15 (Figure 4a). It is characterized by homogeneous texture and quite high packing estimated within 20-30% (area). Temper consists predominantly of fine sand (0.125-0.25 mm), very fine sand (0.06-0.125 mm), and coarse silt (< 0.06 mm). The sand grains are predominantly represented by sub-angular or angular monocrystalline

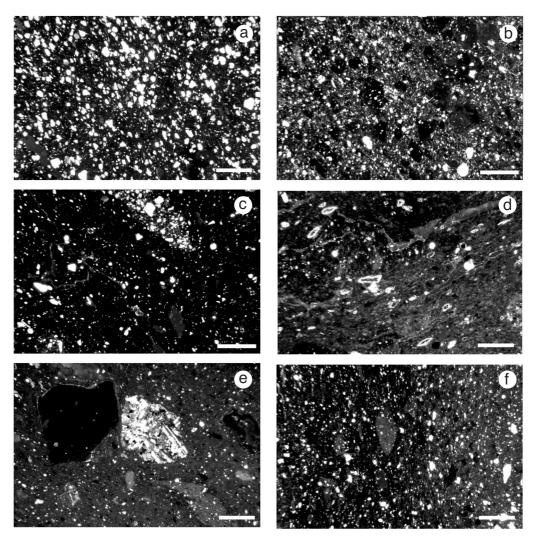


Figure 4. Representative thin section microphotographs of the defined fabrics (FBR): a) FBR 1a; b) FBR 1b; c) FBR 2; d) FBR 3; e) FBR 4; f) FBR 5 (crossed polars; scale bar width = 0.5 mm).

quartz (size < 0.2 mm), only sporadically subrounded (size > 0.2 mm). FBR 1a contained subordinate components including: K-feldspar, chert, quartzarenite fragments, mica flakes (fine fraction) and zircon grains. Micritic clots, formed from the thermal decomposition of calcite (Cau et al., 2002), were estimated

sporadic to rare. The groundmass generally shows weak birefringence except for the sample PLZ28. All samples exhibited a characteristic *black core* that indicates an abrupt passage from a reducing atmosphere, at least in the final part of the firing process, to an oxidant cooling phase atmosphere. The iron oxides particles are

diffused in the groundmass and they also assemble in small round or irregular clots (varying dimensions from 0.1 to 0.6 mm). The pores mainly exhibit an irregular or vesicular form range in size from 0.1 to 1 mm.

FBR 1b (samples: PLZ8; PLZ30; PLZ38; PLZ39)

It exhibited intermediary characteristics between FBRs 1a and 2. It was characterized by a medium temper packing (10-15%) with fine sand size (Figure 4b).

Sample PLZ8 primarily contained very fine subangular or angular grains of monocrystalline quartz (0.06-0.125 mm); while K-feldspar, polycrystalline quartz, quartzarenite fragments and tiny mica flakes and micritic clots were sporadic to rare components. Samples PLZ30, PLZ38, PLZ39 exhibited an increase in the relative abundance of calcareous grains although the majority of temper grains remained silicoclastic.

The groundmass appeared to be rather non-homogeneous (lumpy texture) and not birefringent except for sample PLZ39 (weak birefringence). Pores mainly expressed irregular shapes and sizes ranging between 0.1 and 1 mm.

FBR 2 (samples: PLZ2; PLZ3; PLZ6; PLZ10; PLZ12; PLZ14; PLZ16; PLZ17; PLZ20; PLZ21; PLZ23; PLZ24; PLZ27; PLZ40; PLZ41; PLZ42; PLZ43; PLZ45; PLZ47)

All of the samples grouped within FBR 2 were characterized by a distinctive lumpy texture and medium-low temper packing (ranging between 5-10%). Coarse silt (< 0.06 mm) and very fine sand grains (0.06-0.125 mm) are prevailing. Medium sand temper (0.25-0.5 mm) was rare or sporadic. Compositionally this microscopic fabric contained, subangular to angular monocrystalline quartz grains, while K-feldspar, polycrystalline quartz, quartzarenite fragments, and mica flakes were subordinate temper components (Figure 4c). It appears that the calcareous microfossils more or less decomposed after the firing process and micritic clots (in the samples fired at higher

temperature) are relatively more abundant respect FBRs 1a and 1b.

The FBR 2 groundmass was not birefringent. Pores exhibited an irregular or vesicular shape, with size widely ranging between 0.1 and 1 mm.

FBR 3 (samples: PLZ7; PLZ36)

This fabric is represented by only two samples. It is characterized by a low packing (around 3% area) and a heterogeneous distribution of the aplastic inclusions with a clear prevalence of coarse silt (0.04-0.06 mm) and very fine sand (0.06-0.125 mm) grains.

The temper was composed mainly of angular or subangular monocrystalline quartz, but calcareous microfossils, occasionally well-preserved and more frequently in the form of subrounded or vesicular cast, were identified as common constituents (Figure 4d). Feldspars grains (orthoclase and plagioclase) and mica flakes were sporadic.

It showed a birefringent groundmass except for the samples PLZ36 (weakly birefringent) and PLZ7 (not birefringent). Pores showed irregular or vesicular shape with average size widely spread between 0.1 and 1 mm. Each sample was characterized by a *black core*.

FBR 4 (samples: PLZ32; PLZ35)

This fabric is composed of only two samples which contain few aplastic inclusions, predominantly represented by coarse silt (0.04 - 0.06 mm) and very fine sand (maximum grain size or MGS = 0.15 mm). Temper packing was low (around 3% area) if referred merely to the detritic siliceous components, hence not considering the clay lumps and micritic clots similarly deriving from the manufacture process.

The temper was essentially composed of monocrystalline quartz with a lesser amount of mica and K-feldspars. Micritic clots were common only in sample PLZ35. Minor quantities of volcanic lithic fragments and minerals (clinopyroxene, alkaline feldspar, hornblende, and glass) were recognized in both samples and can be considered a distinctive

"marker" of this fabric (Figure 4e).

The groundmass showed a distinct birefringence typically yellowish brown to brown (crossed polars). Texture was manifestly non homogeneous with frequent clay lumps and a *black core*. Pores were mainly irregular or vesicular shaped and the size was highly variable from 0.1 to 1 mm.

FBR 5 (samples: PLZ3; PLZ26)

This fabric was once again represented by only two samples (PLZ3, PLZ26). Each sample was characterized by a medium temper packing (higher than10% area) with the aplastic inclusions distributed uniformly (Figure 4f). Temper size was predominantly fine-medium (0.2-0.5 mm) with tails toward coarse sand (0.5-1 mm) and silt (< 0.06 mm).

From the point of view of mineralogical composition each sample exhibited a prevalence of monocrystalline quartz, with subordinate to rare K-feldspar, polycrystalline quartz, chert, quartzarenite fragments and mica flakes. Each calcareous lithoclasts and microfossils are common constituents as well.

The groundmass was optically active and the texture homogeneous even with the *black core*. Pores show irregular or vesicular shape and many of them represent casts of decomposed calcareous microfossils. Pore size ranged widely between 0.1 and 1 mm.

Chemistry of the ceramic samples

The raw chemical data concerning the ceramic samples collected at Monte Polizzo are shown in Table 2. They have been carried out by XRF spectrometry and arranged in succession according to the above described *fabrics* recognized after petrographic observations. In fact, the first aim of the XRF analysis, in this case study, was to verify if the petrographically delineated *fabrics* (FBR 1a, FBR 1b, FBR 2, FBR 3, FBR 4, FBR 5) were still evident when viewed by chemical composition and categorizable just like *Paste Compositional*

Reference Unit (PCRUs). At first sight it is possible to notice that for the studied samples there is a very good correlation between petrographic classification of microscopic fabrics and chemical data. Therefore, PCRU 1a, PCRU 1b, PCRU 2, PCRU3, PCRU 4, PCRU 5 can be easily distinguished by means of major and trace elements concentrations, respectively corresponding to FBR 1a, FBR 1b, FBR 2, FBR 3, FBR 4 and FBR 5.

In particular, PCRUs 1a and 1b differ from all the other paste compositional reference units by a larger SiO₂ abundance (in particular PCRU 1a) and, in contrast, really lower amounts of CaO and Al₂O₃. On the contrary, PCRU 2 was characterized by a relatively higher CaO concentration with respect to PCRU 1a, and also by higher abundances of Al₂O₃, TiO₂, MgO, Fe₂O₃. Particularly, the inverse correlation between SiO₂ and CaO, distinguishing PCRU 1a and 1b from PCRU 2, can be interpreted like an effect due to greater relative occurrence of sand grains largely made of quartz and/or to a correspondent minor incidence of the calcareous bioclasts already detected under the petrographic characterization. The inverse correlation between SiO₂ and Al₂O₃, then again, reveals the different relative frequency of sand inclusions (packing assessed at the polarizing microscope and given in % area). In fact, Al₂O₃, TiO₂, MgO, Fe₂O₃ resulted relatively higher concentrated in the less tempered PCRU 2 reflecting the more relevant contribution of the mineralogical/amorphous constituents that compose the groundmass in comparison with the aluminium-silicates of the sand fraction. In addition, the samples petrographically attributed to the PCRU 1a showed a marked dispersion of concentration (from the 64% to around the 74% wt), causing also a sort of "dilution effect" on CaO abundance and of the geochemically correlated Sr which is fairly higher in PCRU 2. In fact, with the increase of quartz grains content - both in terms of their relative abundance

Table 2. Concentration of major elements (% weight, after normalization against LOI) and trace elements (ppm) of the studied ceramic samples from Monte Polizzo.

Zn PCRU	101 la	105 la	112 la	97 la	98 la	116 la	105 la	102 la	105 la	114 la	122 la	119 la	113 la	117 la	109	122	76	128 1b	135 1b		132 1b			
Cu	18	15	21	14	16	18	20	20	17	21	24	22	23	24	20	24	14	25	28	26		29	29 27	29 27 29
^	117	119	121	103	104	125	106	105	101	126	113	120	118	110	113	126	101	121	119	127		121	121	121 122 127
Ce	61	49	49	52	55	62	70	69	48	89	69	65	28	61	62	70	48	71	89	99		70	02	70 69 17
La	25	25	27	22	24	26	29	27	22	28	29	28	23	25	26	29	22	31	28	26		59	29	29 29 31
Ba	256	253	302	228	241	317	251	284	231	293	324	292	301	322	278	324	228	373	382	352		341	341	341 362 382
Zr	496	499	412	612	584	400	475	534	602	476	420	512	533	555	208	612	400	381	370	412		394	394	394 389 412
Sr	93	86	123	82	98	124	105	105	92	116	127	113	112	110	106	127	82	152	185	165		181	181	181 171 185
Rb	71	78	69	28	61	87	81	9/	2	82	96	85	95	91	78	96	28	110	108	108		26	97	97 106 110
K_2O	1.71	1.74	1.92	1.54	1.58	1.9	1.85	1.79	1.63	1.86	2.12	1.89	1.97	1.98	1.82	2.12	1.54	2.15	2.13	2.09		1.98	1.98	1.98 2.09 2.15
Na_2O	0.25	0.28	0.29	0.18	0.22	0.29	0.30	0.28	0.19	0.33	0.43	0.30	0.39	0.41	0.30	0.43	0.18	0.30	0.29	0.27		0.75	0.25	0.25 0.28 0.30
CaO	1.70	2.31	3.87	1.59	1.55	3.27	2.37	2.35	1.79	3.16	3.55	3.13	2.14	2.93	2.55	3.87	1.55	5.04	5.78	5.22		5.69	5.43	5.69 5.43 5.78
MnO	0.07	0.04	0.05	0.03	0.04	0.08	90.0	90.0	0.04	0.09	0.07	0.15	0.04	0.10	0.07	0.15	0.03	0.10	90.0	90.0	90	0.08	0.08	0.08
MgO	1.52	1.64	1.27	1.26	1.37	1.81	1.88	1.45	1.32	1.99	1.77	1.73	1.77	1.83	1.62	1.99	1.26	1.72	1.77	2.09		1.81	1.81	1.85
Fe_2O_3	7.84	7.60	7.75	5.97	6.77	7.50	6.49	6.33	5.93	8.08	8.09	8.25	7.79	7.59	7.28	8.25	5.93	8.02	7.40	8.05		7.84	7.83	7.83
P_2O_5	0.14	0.13	0.21	0.11	0.14	0.28	0.12	0.31	0.12	0.17	0.13	0.17	0.26	0.19	0.18	0.31	0.11	0.27	0.20	0.17		0.15	0.20	0.15 0.20 0.27
Al_2O_3	17.57	17.63	18.43	14.69	16.40	18.36	16.51	17.91	16.54	18.22	18.74	18.43	19.56	19.08	17.72	19.56	14.69	19.82	19.51	18.73	10.01	19.61	19.67	19.67 19.48 19.87
TiO ₂ /	0.91	0.92	1.01	0.79	88.0	68.0	98.0	98.0	0.84	86.0	1.14	0.94	1.08	66.0	0.94	1.14	62.0	1.00	1.04	1.06	701	1.04	1.04	1.04
SiO_2	68.29	67.71	65.19	73.84	71.06	65.62	85.69	29.89	71.60	65.12	63.96	65.02	64.98	64.90	67.54	73.84	63.96	61.58	61.82	62.25	61 20	77:17	61.74	61.74
Samples	PLZ 1 (PLZ 4 (PLZ 15 6	PLZ 18 7	PLZ 19 7	PLZ 22 (PLZ 25 (PLZ 28 (PLZ 29 7	PLZ 33 (PLZ 34 (PLZ 37 (PLZ 44 6	PLZ 46 6	Mean (Max 7	Min	PLZ 8 (PLZ 30 (PLZ 38 (PI 7 39 6			

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PCRU	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2				3	3				4	4				5	5			
Zn	127	122	149	134	140	161	134	149	134	131	168	133	128	137	126	129	138	168	122	190	179	185	190	179	146	140	143	146	140	156	159	158	159	156
Cn	27	26	34	29	32	39	30	36	33	34	38	26	25	59	28	28	31	39	25	22	32	27	32	22	79	74	77	62	74	20	21	21	21	20
>	155	145	122	146	173	129	130	121	137	162	113	125	152	141	145	143	140	173	113	96	104	100	104	96	120	117	119	120	117	06	98	88	06	98
Ce	73	29	69	71	99	50	54	57	70	52	59	48	61	57	57	61	61	73	48	70	73	72	73	70	44	39	42	44	39	82	96	68	96	82
La	32	28	31	31	27	23	26	24	31	24	26	22	27	25	28	25	27	32	22	28	29	29	29	28	17	14	16	17	14	35	40	38	40	35
Ba	370	376	449	388	411	452	391	463	391	379	427	368	356	379	371	368	396	463	356	785	757	771	785	757	493	527	510	527	493	599	646	623	949	599
Zr	397	376	382	349	372	365	393	384	393	330	361	374	383	319	388	372	371	397	319	185	187	186	187	185	206	221	214	221	206	189	177	183	189	177
$S_{\mathbf{r}}$	145	139	218	173	197	245	191	215	167	155	264	153	147	187	158	165	182	264	139	332	319	326	332	319	233	211	222	233	211	569	283	276	283	269
Rb	122	109	137	102	86	116	121	1111	124	131	120	141	127	130	148	142	124	148	86	130	115	123	130	115	128	178	153	178	128	102	91	76	102	91
K_2O	2.36	2.24	2.40	2.19	2.09	2.16	2.26	2.21	2.36	2.50	2.39	2.49	2.34	2.39	2.51	2.49	2.34	2.51	2.09	2.58	2.30	2.44	2.58	2.30	2.54	2.80	2.67	2.80	2.54	2.10	1.98	2.04	2.10	1.98
Na_2O	0.49	0.41	0.37	0.38	0.32	0.37	0.45	0.35	0.40	0.47	0.46	0.53	0.47	99.0	0.44	89.0	0.45	89.0	0.32	0.61	0.57	0.59	0.61	0.57	0.70	0.75	0.73	0.75	0.70	0.34	0.37	0.36	0.37	0.34
CaO	4.90	4.66	7.29	5.58	6.01	8.11	5.80	7.70	5.59	5.47	8.90	5.11	4.46	5.92	4.98	4.95	5.96	8.90	4.46	12.15	11.74	11.94	12.15	11.74	7.34	7.20	7.27	7.34	7.20	80.6	9.43	9.26	9.43	80.6
MnO	90.0	0.07	90.0	90.0	90.0	90.0	0.08	0.07	90.0	0.07	0.08	0.07	0.07	0.09	0.08	0.08	0.07	0.09	90.0	90.0	0.10	0.08	0.10	90.0	0.14	0.13	0.14	0.14	0.13	90.0	0.05	90.0	90.0	0.05
MgO	2.12	1.73	2.10	1.94	2.34	2.29	1.86	1.61	1.65	1.81	2.14	1.60	2.38	2.89	2.05	2.47	2.06	2.89	1.60	2.40	2.07	2.23	2.40	2.07	4.03	4.16	4.10	4.16	4.03	2.01	2.18	2.10	2.18	2.01
Fe_2O_3	9.03	9.45	8.24	9.65	9.07	8.31	8.93	8.14	8.50	9.40	8.08	9.18	69.6	9.22	8.77	8.88	8.91	69.6	80.8	7.59	8.42	8.00	8.42	7.59	9.01	8.52	8.77	9.01	8.52	7.88	7.37	7.63	7.88	7.37
P_2O_5	0.22	0.19	0.18	0.19	0.41	0.22	0.16	0.18	0.25	0.28	0.23	0.25	0.17	0.18	0.22	0.22	0.22	0.41	0.16	0.27	0.25	0.26	0.27	0.25	0.08	0.08	0.08	0.08	0.08	0.23	0.24	0.24	0.24	0.23
Al_2O_3	21.63	21.20	19.97	22.84	22.80	20.63	20.66	21.50	22.12	22.19	20.27	22.09	22.36	21.60	21.37	21.34	21.54	22.84	19.97	20.14	20.24	20.19	20.24	20.14		20.17	20.35	20.53	20.17	18.57	18.87	18.72	18.87	18.57
TiO_2	1.13	1.20	1.09	1.25	1.29	1.03	1.10	1.17	1.18	1.35	1.05	1.27	1.21	1.10	1.10	1.09	1.16	1.35	1.03	1.07	1.07	1.07	1.07	1.07	0.92	98.0	0.89	0.92	98.0	0.89	0.87	0.88	0.89	0.87
SiO_2	58.05	58.84	58.29	55.92	55.61	56.81	58.69	57.06	57.89	56.46	56.41	57.42	58.85	55.96	58.49	57.81		58.84	55.61	53.14	53.24	53.19	53.24	53.14	54.62	55.33	54.98	55.33	54.62	58.84	58.63	58.74	58.84	58.63
Samples	PLZ 2	PLZ 3	PLZ 6	PLZ 12	PLZ 14	PLZ 16	PLZ 17	PLZ 20	PLZ 21	PLZ 24	PLZ 27	PLZ 40	PLZ 41	PLZ 42	PLZ 45	PLZ 47	Mean	Max	Min	PLZ 7	PLZ 36	Mean	Max	Min	PLZ 32	PLZ 35	Mean	Max	Min	PLZ 9	PLZ 26	Mean	Max	Min

(packing) and/or average size - an equivalent variation of SiO₂ (wt%) is expected. On the other hand this aspect does not invalidate the consistency of PCRU 1a as it could be also considered a natural variation of the quartz sand content in the exploited clay sources rather than due to potter's premeditated technological actions. Concerning the other trace elements, Zr resulted clearly more abundant in the ceramic samples assigned to PCRU 1a being correlated to the higher detritic siliciclastic sand grains frequency characterising this paste, while Rb and Sr resulted relatively enriched in the samples belonging to PCRU 2 in agreement with the correspondent presence of feldspars and mica flakes in the very fine sand and silt fractions. Also V, Cu and Zn seem to be slightly more concentrated in PCRU 2 ceramic samples with respect to PCRU 1a, while no relevant differences were found concerning La and Ce abundances.

Few comments can be made for the remaining PCRUs 3, 4 and 5 due to their scarce representativeness in terms of number of samples. In general, they show compositional characteristics relatively closer on the whole to PCRU 2, apart of PCRU 3 that has a clearly higher CaO content.

To further test the reliability of the delineated PCRUs from Monte Polizzo, involving all the determined major, minor and trace elements, Principal Components Analysis (PCA) was applied to chemical data. Figure 5 exhibits the distribution of the considered ceramic samples in the plane defined by the first two principal components (C1 and C2), which collectively represent up to the 78% of total variance. Welldefined chemical groups, especially those constituted by the greatest number of samples, PCRUs 1a, 1b and 2, were clearly recognized. As it can be derived by the graph of factor loadings (histogram in the right part of Figure 5) the component C1 is positively influenced by the most part of the considered elements (especially

CaO, K₂O, Rb, Sr, Ba, Cu and Zn) while both SiO₂ and Zr have negative loadings. In addition, Zr is the only element having a noteworthy negative loading. C2 appears to have a negligible influence in the differentiation of the PCRUs 1a, 1b and 2. The consistent negative contribution on C2 of Cu separates the two samples classified in the PCRU 4 very well. Samples PLZ32 and PLZ35 (PCRU 4) were the only ones that contained a temper with volcanic granules and they had double the concentration of Cu in comparison with the other ceramic samples. Moreover, the positive contribution given to the same component C2 by P₂O₅ separates, in the upper right hand part of the graph, pastes 5 and 3.

Petrography and chemistry of local clays

Sample SALT11, an experimental briquette fired at 900 °C, represents the locally available clays of the Terravecchia Formation. Thin observations using a polarizing microscope revealed that SALT11 possessed a texture of coarse silt-very fine sand temper with a packing around the 15% (area). The siliceous granules were mainly composed monocrystalline quartz and to a lesser degree polycrystalline quartz and K-feldspar (Figure 6a). Mica flakes were also common constituents within the finest fraction, while calcareous microfossils (even if decomposed after firing) were sporadic.

The additional experimental briquette MIX1 (2 parts weight of SALT10 mixed with 3 parts of SALT11) was characterized by a greater incidence of the siliceous temper with a resulting average packing around the 25-30% area (Figure 6b). The experimental fired MAB Formation clay briquettes contained a greater relative abundance of calcareous microfauna (mainly planktonic foraminifera) with respect to the Terravecchia Formation. Investigating the fossil content within briquettes is not easy due to the rarity of well-preserved fossil remains as a consequence to the firing and soaking, 900 °C,

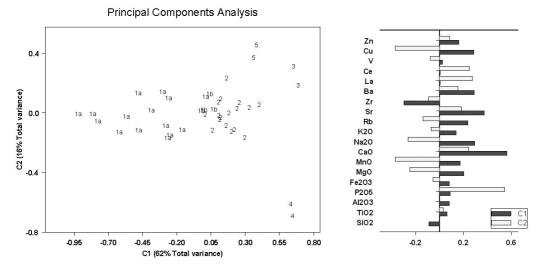


Figure 5. Plot of factor scores concerning all ceramic samples categorized within the PCRUs (1a, 1b, 2, 3, 4, 5) in the plane of the first two principal components C1 and C2. Histogram at the right shows factor loadings on chemical variables on the two considered principal components.

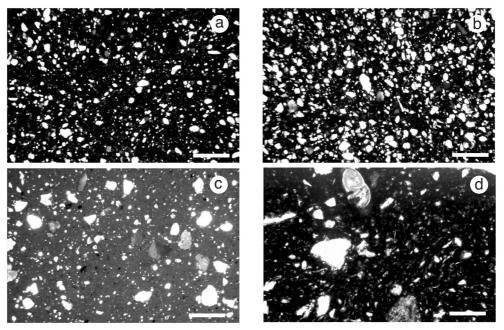


Figure 6. Thin section microphotographs of Monte Polizzo's raw materials after experimental firings at 900°C: (a) Terravecchia Formation (sample SALT 11: crossed polars; scale bar width = 0.5 mm); (b) Terravecchia Formation (sample MIX 1: crossed polars; scale bar width = 0.5 mm); (c) MAB Formation (sample SALT 1: crossed polars; scale bar width = 0.5 mm); (d) MAB Formation (sample SALT 1: parallel polars; scale bar width = 0.2 mm).

process (Figure 6c, d). Monocrystalline and polycrystalline quartz were still the most abundant temper components while feldspars and mica were sporadic to rare. Temper packing was between 15 and 20% (area) and fine sand grains (up to 0.25 mm) were quite common.

Table 3 describes the chemical composition of local clay samples obtained by XRF analysis. The maximum, minimum and average values concerning 47 additional clay (belonging to 7 different outcrops of the Terravecchia Formation in the territory of western Sicily within a radius of 50 Km from the settlement of Monte Polizzo were also reported for comparative purposes (data after Montana et al., 2010b). The outstanding discrepancy between the two representative local samples belonging to the Terravecchia Formation is undoubtedly due to the difference in the relative abundance of the sand fraction. In fact, SALT10 (sand = 55% wt) was characterized by a higher SiO₂ concentration (quartz grains) and lower contents of all the other oxides of major and minor elements (especially true for CaO). The same was observed in all of the trace elements apart from Zr (ZrSiO₄, zircon, was a common accessory mineral relatively enriched in the sand fraction of Neogene sedimentary deposits in Sicily). Elemental abundances of sample SALT11, on the contrary, were in good agreement with those detected for the clays of the Terravecchia Formation in other areas of western Sicily (Montana et al., 2006; Montana et al., 2011b). The Al₂O₃ content was also found to be relatively high and positively correlated to K₂O content. This can be explained by the abundance of mica flakes which were found especially enriched in the coarse silt fraction. CaO concentration was found to be the 7% weight (the average value found in western Sicily is a little bit higher = 7,68% wt), due to the relatively limited presence of calcareous microfauna. With reference to concentrations of Ba and Rb, both geochemically related to

potassium, mica flakes and K-feldspar characterize the very fine sand and coarse silt fractions.

The bulk chemistry of the MIX1 sample (Table 3) reflects precisely the relative amounts of SALT10 (2 parts) and SALT11 (3 parts) used in the experimental mixing procedure.

All the samples belonging to the MAB Formation (SALT1, SALT3, SALT4, SALT6) showed quite comparable bulk chemical compositions. They are interpreted as being 'very calcareous clays' with remarkable CaO concentrations, up to 25% (wt) which is very likely related to the abundant calcareous microfauna. From a comparative point of view, the CaO concentration, significant differences between major and trace elements in the local clayey materials representative of Terravecchia and MAB formations were recognized also in the relative abundances of Fe₂O₃, Al₂O₃, Na₂O, K₂O, Rb, Ba, Sr.

Petrographic comparison between ceramic samples and local clays

In order to ascertain the petrographic markers of Monte Polizzo's productions within the studied set of ceramic samples, already divided into the previously described fabrics, it was assumed important to make a quantitative comparison amongst significant textural and mineralogical features derived from thin sections observation both on ceramic samples and briquettes deriving from the experimental firing of representative samples of the exploitable clays deposits. Useful variables for this purpose are represented by sand inclusions packing (% area) and size, presence (with estimation of relative abundance) or absence of specific rock fragments, presence (with estimation of relative abundance) or absence of specific minerals. Then, the Multiple Correspondence Analysis (MCA), a methodology of multivariate statistical analysis (Baxter, 1994), was used with the aim of providing a quantitative

and MAB Formation (48 samples collected from 5 sites) in the territory of western Sicily within a radius of 50 Km from Monte Polizzo were also Table 3. Concentration of major elements (% weight, after normalization against LOI) and trace elements (ppm) of the considered raw clay samples. The maximum, minimum and average values concerning clay samples representative of Terravecchia Formation (52 samples collected from 8 sites) reported (after Montana et al., 2011b).

Ce V Cu Zn	75 15 78 MAB		52 18 84 MAB	18 84 14 72	18 84 14 72 13 71	18 84 14 72 13 71 14 92	18 84 14 72 13 71 14 92 35 148	18 84 14 72 13 71 14 92 35 148 30 127	18 84 14 72 13 71 14 92 35 148 30 127	18 84 14 72 13 71 14 92 35 148 30 127 24 108	18 84 14 72 13 71 14 92 35 148 30 127 24 108	52 18 84 79 14 72 62 13 71 97 14 92 154 35 148 135 30 127 154 24 108 99 14 78 194 41 138	52 18 84 79 14 72 62 13 71 97 14 92 154 35 148 135 30 127 154 24 108 99 14 78 194 41 138	52 18 84 79 14 72 62 13 71 97 14 92 154 35 148 135 30 127 154 24 108 99 14 78 194 41 138 88 16 59	52 18 84 79 14 72 62 13 71 97 14 92 154 35 148 135 30 127 154 24 108 99 14 78 194 41 138 88 16 59
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CaO Na ₂ O	22.68 0.05	22.40 0.13	20.02 0.15		22.01 0.19				0.53 0.1 0.53 0.1 7.01 0.8 4.66 0.4 Terrave	0.53 0.15 7.01 0.85 4.66 0.49 Terravecc 7.68 0.65	22.01 0.19 0.53 0.15 7.01 0.85 4.66 0.49 Terravec 7.68 0.65 3.35 0.21	22.01 0.19 0.53 0.15 7.01 0.85 4.66 0.49 Terravecc 7.68 0.65 3.35 0.21 13.04 1.50	22.01 0.1. 0.53 0.1. 7.01 0.8 4.66 0.4 Terrave 7.68 0.6 3.35 0.2 MA	0.53 0.15 7.01 0.85 7.01 0.85 4.66 0.49 Terravec 7.68 0.65 3.35 0.21 13.04 1.50 MABB	22.01 0.19 0.53 0.15 7.01 0.85 4.66 0.49 Terravecc 7.68 0.65 3.35 0.21 13.04 1.50 MAB 25.78 0.27
OIIIAI	0.07	0.00	0.13	0.08		17 0.02						0.06 0.06 0.06 0.06 0.004 0.004	0.06 0.06 0.06 0.06	0.06 0.06 0.06 0.087 0.087	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0
1 22 3 MgO	5.41 1.52	4.55 2.35	5.91 1.33	5.71 2.44		7.11 1.37									
1205 15	60.0	70.0	0.10	0.14		0.07									_
Al_2O_3	16.09	12.53	13.49	12.08		13.27									
7011	0.77	0.63	69.0	09.0		99.0	0.66	0.66	0.66	75.17 0.66 56.39 1.04 64.16 0.92 57.56 1.09	0.92 0.92 0.92 0.90 0.90	56.39 1.04 64.16 0.92 67.56 1.09 57.56 1.09 52.17 0.90	0.92 0.92 0.92 0.90 0.90 0.90	0.66 0.92 0.92 0.90 0.90 0.90 0.90	0.96 0.92 0.90 0.90 0.90 0.71 0.71
1	51.43	55.94	56.62	55,21	75 17										
	SALT1	SALT3	SALT4	SALT6	SALT10		SALT11	SALT11	SALTII	SALT11 MIX1 Mean	SALT111 MIX1 Mean Mean	SALT111 MIX1 Mean Min Min Max	SALT11 MIX1 Mean Min Max	SALT11 MIX1 Mean Min Max Meax	SALT111 MIX1 Mean Min Mean Mean Mean

TV = Terravecchia Formation; MAB = Marnoso-Arenacea del Belice Formation; MIX1 = experimental mixing (see text).

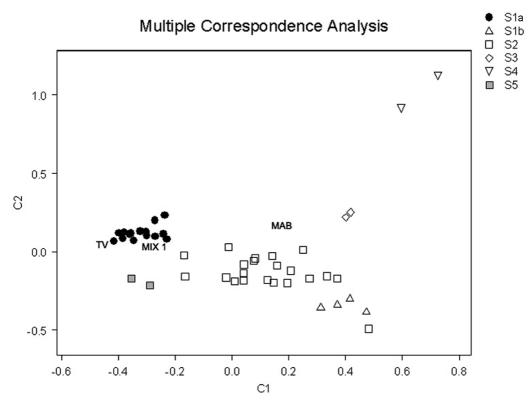


Figure 7. Biplot of MCA scores of transformed textural and compositional parameters derived from thin section microscopy (see text). Legend: S1a = FBR 1a; S1b = FBR 1b; S2 = FBR 2; S3 = FBR 3; S4 = FBR 4; S5 = FBR 5; TV = Terravecchia Formation; MIX1 = experimental mixing of two different samples of Terravecchia clays (see text); MAB = Marnoso-Arenacea del Belice Formation (average values obtained from the 4 samples analyzed).

and straight comparison between ceramic samples and local raw clays. This method was already successfully tested for the petrographic characterization of archaeological ceramics (Cau et al., 2004; Baxter et al., 2008; Montana et al., 2010; Montana et al., 2011b). The petrographic data (both compositional and textural) deriving from the analysis with the polarizing microscope of the ceramic samples and the selected raw materials (average values for the MAB Formation and Terravecchia Formation as well as experimental mixing MIX1) were thus categorized and converted in a binary form

(entries 0/1). A list with all the considered variables, categories and conversion values of "qualifiers" was reported in detail (Appendix A). The numerically transformed data (Appendix B) have been treated with the MCA.

Figure 7 projects the petrographic binary scores in the plane represented by the first two components (C1 and C2) that together represent a substantial portion of the total variance (70%). The interpreted FBRs seem to form homogeneous and separated clusters. A clean convergence between FBR 1a and both the experimental test-pieces from the Terravecchia

Formation materials (SALT11 and MIX1) is also recognizable. Samples belonging to FBRs 2 and 1b appear to be split from the FBR 1a cluster and the Terravecchia clays. Therefore, even if FBRs 1b and 2 are both thought to also represent local productions, following the archaeological classification (use type, form, nature and style of external ornamental patterns), they do not correspond exactly with the textural features of the raw materials supplied from local Terravecchia Formation's outcrops (specially quartz temper relative abundance and packing). On the other hand, is very clear that clays belonging to the MAB Formation are not petrographically similar to any of the specific fabric groups achieved from the petrographic observations. In fact, they do not fit FBR 1a or together **FBR** which represent mineralogical and textural features of the most part of the studied ceramic samples. Similar considerations can be made for the remaining FBRs (FBR 3, FBR 4 and FBR 5) that, in fact, could be considered imported ceramic products. FBR 4 is also clearly distinct from the others in the upper right part of the graph likely due to the peculiar presence of volcanic minerals and lithic fragments. Consequently, this approach demonstrated that the textural aspects like temper packing and average size could have a considerable effect in both the macroscopic and microscopic appearance of the fabric, and that, sometimes, they may also become misleading for classifying purposes in terms of provenance attribution.

Chemical comparison between ceramic samples and local clays

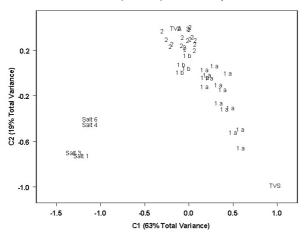
In the light of the complexities just verified with comparing only by textural and mineralogical features ceramic artefacts and clayey raw materials, it is important to reconsider the level of chemical compatibility between the Monte Polizzo's Archaic pottery and the local clay outcrops. The end goal remains to

identify the archaeological site at Monte Polizzo as a ceramic production centre versus a consumption centre.

Thus, the multivariate method of the Principal Components Analysis (PCA) was applied for a second time to further corroborate the estimated correspondence between the ceramic pastes and local clay materials. This analysis was only applied to the PCRUs 1a, 1b and 2 (respectively corresponding to FBR 1a, FBR 1b, FBR 2), because they contained the most part of individuals. Figure 8 projects the factor scores in the plane corresponding to the first two principal components (C1 and C2) together representing the 82% of the total variance. Once more it appears very evident that the clays of the MAB Formation are clearly separated from the PCRUs 1a, 1b and 2, which in turn are quite homogeneously clustered. This evident separation from MAB Formation clays may be explained by taking into account the histogram of factor loadings (in the right part of Figure 8), where the negative load on component C1 is strongly influenced by CaO and Sr. A good correlation was once again recognized between PCRUs 1a, 1b, 2 and the clays of the Terravecchia Formation. This PCA correlation also exhibits an opposing chemical "dilution effect" that can be interpreted as complementary to the relative abundances of SiO₂ and CaO along the line identifiable by the two clay samples representative of the Terravecchia Formation (SALT10 = TVS and SALT11 = TV), and clearly enhanced by the strong divergence of the C2 component.

Taking note of the large natural variability within clayey materials of the local beds of the Terravecchia Formation, it is necessary to obtain a further verification of their possible use for this specific ceramic production. Therefore, an additional comparison between PCRUs 1a, 1b and 2 against the briquette MIX1 produced by the experimental combination of the Terravecchia Formation samples SALT10 (2

Principal Components Analysis



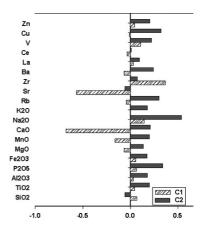


Figure 8. Plot of PCA scores in the plane of the first two principal components C1 and C2 concerning the ceramic samples belonging to the PCRU's 1a, 1b, 2, the MAB Formation clays (SALT 1, SALT 3, SALT 4, SALT 6) and the Terravecchia Formation clays (TVA= SALT 11; TVS = SALT 10). Histogram at the right shows the loadings of chemical variables on the two considered principal components.

parts weight) and SALT11 (3 parts weight) was made. The extremely different content of quartz sand (SALT10 reached 53% wt of quartz-bearing sand while SALT11 reached only 13% wt of quartz-bearing sand) plays indirectly for "dilution effect" mainly in the correspondent abundance of CaO. Figure 9, a composed binary graph, exhibits the values of SiO₂, Al₂O₃, CaO and K₂O found in the samples belonging to the PCRUs 1a, 1b and 2 projected together with the samples representative of the local beds of the Terravecchia Formation (SALT11 with a standard content of sand and SALT10 with an extreme content of sand) and the experimental mixture made with them. Through this representation it is possible to recognize that the very most part of the ceramic samples classified into PCRU 1a are delimited by the potential raw materials SALT10 and MIX1 (relatively lower CaO, Al_2O_3 , K_2O and relatively higher SiO_2), while the ceramic finds classified in the PCRUs 1b and 2 are delimited by MIX1 and SALT11

(relatively lower SiO₂ and increasing CaO, Al₂O₃, K₂O. This compositional features could be interpreted either as a result of technological choices inductively pursued by the local craftsmen (selection of specific clay beds and/or mixture of clays with different content in sand), to a pure chance linked to the natural variability of the clayey outcrops along the stratigraphic column (intra-deposit natural variability) or to a change over time in the exploitation of local clayey materials.

Concluding remarks

The mineralogical-petrographic and chemical study of 44 ceramic artefacts obtained from the Acropolis of Monte Polizzo allowed the identification of 6 primary different 'Paste Compositional Reference Units' (PCRUs). These PCRUs were recognized and independently confirmed by their textural and mineralogical characteristics (fabrics or FBRs by petrographic observations) and chemical composition (XRF

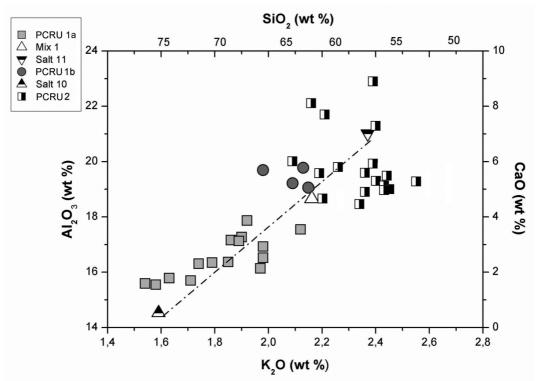


Figure 9. Composite binary plot (SiO₂, Al₂O₃, CaO and K₂O) that exhibits PCRUs (1a, 1b and 2) with clay samples from the Terravecchia Formation.

analysis). In order to support the identification of local production we chose to study also the available clay-sized raw materials in the area surrounding the archaeological site, belonging to the Terravecchia Formation (Upper Tortonian-Messinian) and to the Marnoso-Arenacea del Belice or MAB Formation (Middle-Upper Pliocene). Several ethnographic proofs recently acquired provide evidence that clays from Terravecchia and MAB Formations were utilized extensively for indigenous ceramic production in western Sicily (Montana et al., 2006; Montana et al., 2011b). Representative samples of these thus investigated were from mineralogical-petrographic point of view (by firing the test pieces of the original raw materials or of artificial mixtures in a muffle kiln) and

from the chemical one, and subsequently compared with the ceramic artefacts object of the study.

The cross comparison of the mineralogical-petrographic and chemical data allows the confirmation that, among the available raw materials near Monte Polizzo, the local workers preferred clays from the Terravecchia Formation. The mineralogical composition of the Terravecchia Formation correlated well with PCRUs 1a, 1b and 2 that represented 92% of the investigated ceramic finds.

Afterwards, the mineralogical, textural and chemical markers of the ceramic vessels for food consumption and storage, corresponding to macrofabric F7 (incised open or closed small pots), macrofabric F54 (matte painted small

amphoras and other nonspecific table ware) and macrofabric F24 (pithoi) produced in the indigenous settlement of Monte Polizzo in a wide chronological interval between the 7th and the 4th century B.C. were archaeometrically ascertained. Archaeometric investigations accordingly allowed to declare Monte Polizzo a ceramic production centre, together with other Sicilian anhellenic sites where positive evidence for pottery workshops was also retrieved. Clay resources used at Monte Polizzo lay in the neighbourhood of the settlement and do not need long range road transport.

The remaining 8% of the investigated ceramic findings, samples PLZ7 and PLZ36 (PCRU 3, macrofabric F54 and F7), PLZ32 and PLZ35 (PCRU 4, both macrofabric F54), PLZ3 and PLZ26 (PCRU 5, both macrofabric F7), have been interpreted to be most likely regional imports from other coeval production centres located in western Sicily. In fact, in the settlements of Entella and Adranone, both sited in the Sicani area, were recently established coeval productions of incised, impressed and matte painted tableware typologically fitting with those produced at Monte Polizzo (Montana et al., 2007; Montana et al., 2010; Montana et al., 2011a). PCRU 4 was characterized by minor quantities of basic volcanic rock fragments and minerals in the temper, nevertheless the stratigraphic successions of the Jurassic deep sea limestone outcropping in the area of Sicani Mounts often exhibit pillow lavas of basaltic nature. Though a wider selection could partly change these results, the limited amount of imported indigenous pottery at Monte Polizzo suggests that local workshops were able to yield the bulk of coarse and decorated pottery commonly used in the settlement.

In conclusion, new inferences useful for the archaeological/historical reconstructions of Archaic Sicily may possibly develop from this archaeometric study, concerning the trade patterns of incised and matte painted tableware

between Monte Polizzo and the main indigenous centres located in the Sicani Mountains area (e.g. Adranone, Entella and Maranfusa) as well as interactions with Selinunte and Segesta.

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