



Distribution and chronological framework for Iberian variscite mining and consumption at Pico Centeno, Encinasola, Spain



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ARTICLE INFO

Article history:

Received 8 May 2015

Available online 8 January 2016

Keywords:

Variscite
Mining
OSL
14C-AMS
Iberia
Pico Centeno

ABSTRACT

AMS radiocarbon and OSL dating, and profiling were used to directly delimit periods of variscite production at Pico Centeno Mine 2. These results were integrated with analysis of other well-dated periods of variscite production to establish an Iberian-wide chronological framework. Variscite production at Pico Centeno Mine 2 began at ~5200 BC, coincident with alpine jade production or Casa Montero Iberian flint production. Variscite was only used occasionally, together with other greenstones, during the 5th and 6th millennia BC. During the 4th millennium BC, variscite use began to increase to its apogee in the first half of 3rd millennium BC when it appeared in nearly every Iberian burial site. This increase in variscite production and use coincided with decline in the popularity of alpine jade. By the end of the 3rd millennium BC, new resources began to be valued such as Asian and African Ivory, Baltic and Sicilian amber, and copper-based metal products. The variscite cycle thus started with the decline of jade in the 5th–4th millennium BC, and ended with the appearance of copper, ivory and extra-peninsular amber by the end of the 3rd millennium BC.

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Introduction

Archaeological literature devoted to green body ornaments in Prehistoric Europe has thematically focused, almost exclusively, on the quest for the origin of these artefacts. Since the early 20th century, interpretations of the geographical origin of these ‘*perles du calais*’ traversed continents, from Middle East turquoise mines to European variscite mines, initially pointing to a French origin at Montebrias (Balagny, 1939), then later to the Pannacé aluminophosphate mines (Massé, 1971; Forestier et al., 1973a, 1973b; Lheur, 1993), and finally to a Spanish origin at the Palazuelo de las Cuevas (Arribas et al., 1970, 1971) and Can Tintorer (Alonso et al., 1978; Bosch and Estrada, 1995; Villalba, 2002) variscite mines.

Since the 1970s, research devoted to ‘*calaite*’ beads has focussed on locating and characterising further new variscite sources. Source areas have been discovered at Bragança in Northeast Portugal (Meireles et al., 1987), at the Sarrabus deposit in Sardinia (Marini et al., 1989), at the variscite and turquoise outcrops of Punta Corveiro in Spain (Moro et al., 1995), and at the Pico Centeno variscite mines in Spain (Nocete and Linares, 1999; Fig. 1). This focus on identifying geological mineral

sources has been driven by the view that their relationships to archaeological bead assemblages can be established by the intercomparison of their chemical compositions (Edo i Benaiges et al., 1995a; Dominguez Bella, 2004; Odriozola, 2014; Odriozola et al., 2010; Querré et al., 2014, 2008).

To date, most papers devoted to beads continue to use calaite and variscite as synonyms. However, the increase in geochemical analyses of green mineral sources was paralleled by an increase in the numbers of analyses of ‘*calaite*’ beads, and thus in the knowledge of the minerals used in beadmaking. Beads were found to include green mica, steatite, turquoise, talc and chlorite (Dominguez Bella, 2012). Therefore, calaite and variscite should not be considered synonyms, and the validity of the traditional analysis of variscite flows and consumption patterns (e.g., Dominguez Bella, 2012) is called into question. As the number of analysed beads increases, it becomes more apparent that Neolithic, Copper Age, and Bronze Age communities used nearly any available green mineral for beadmaking.

The current consensus on the geographic focus of European variscite body ornament production points to Palazuelo de las Cuevas (Aliste, Zamora), Can Tintorer (Gavá, Barcelona) and Pico Centeno (Encinasola, Huelva; Dominguez Bella, 2004; Herbaut and Querré, 2004; Querré et al., 2008, 2014; Odriozola et al., 2010; Odriozola, 2014). The chronological span of variscite production is less clear, however. Arribas et al.

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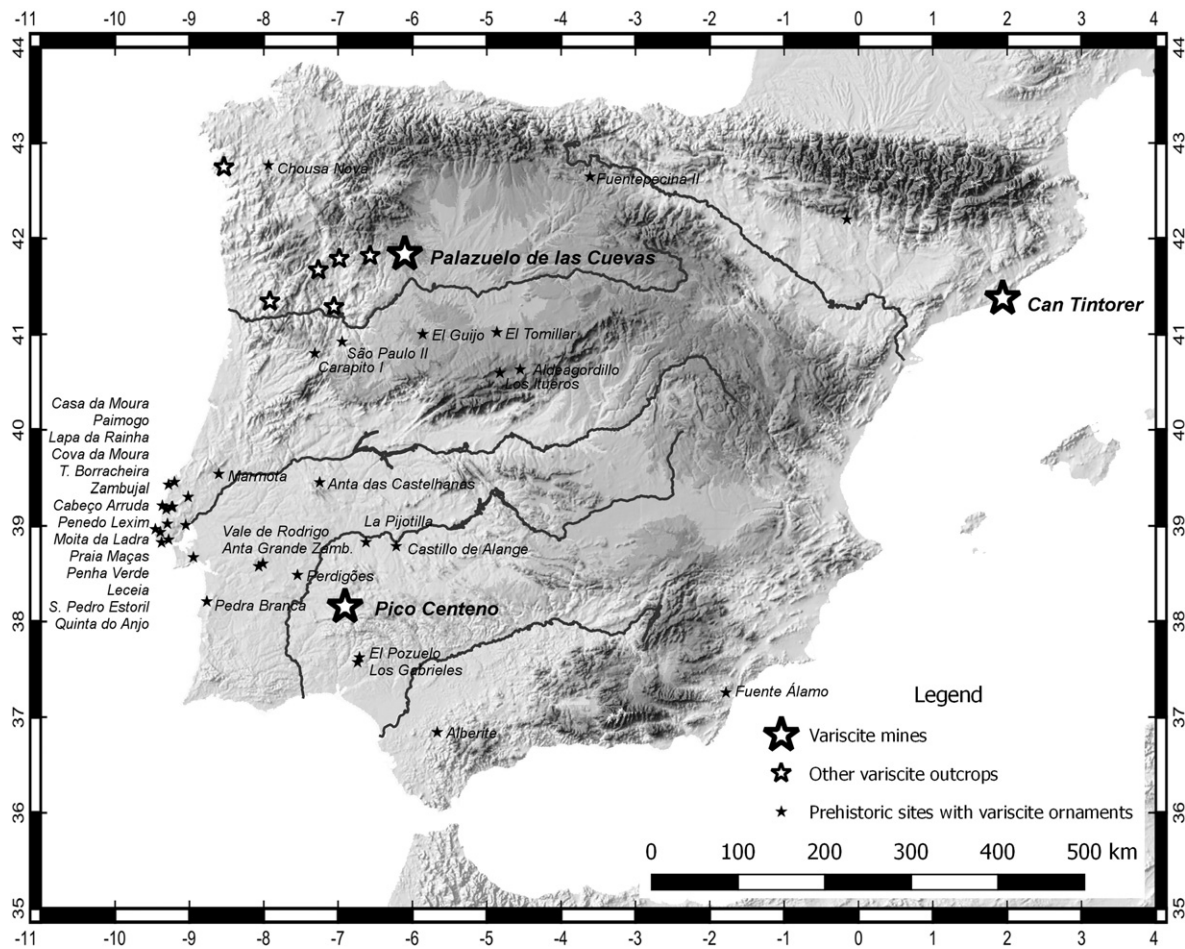


Figure 1. Location of known Iberian variscite sources.

(1970, 1971) proposed that variscite production occurred during the Arabic period at Palazuelo de las Cuevas, based on the assumption that the city of Zamora was named after the Arabic word for emerald (zamarrad). Virgilio Sevillano (1978) and Campano Lorenzo et al. (1985) instead found that Roman pottery and prismatic blanks were associated with bead production at Las Cercas, and Copper Age variscite mining has been proposed at La Mazada (Sanz Mínguez, et al., 1990; Esparza Arroyo and Larrazabal Galarza, 2000). Both Las Cercas and La Mazada form part of the Palazuelo de las Cuevas complex. Meanwhile, Can Tintorer has been extensively dated to the Late Neolithic, between ~4500 and 3500 BC, based on burials, charcoals and seeds found in direct association with the mine galleries (Table 1).

Jiménez Gómez (1995) attempted to chronologically order variscite production patterns at the Zambujal site (Torres Vedras, Central Portugal), based on radiocarbon ages and on mineralogical analysis. They concluded that production was mainly prehistoric; initially a mixture of greenstones was used, then variscite use dominated the second half of the 3rd millennium BC. Villalobos García (2012) has shown that this preference for variscite over other greenstones during the second half of the 3rd millennium BC also occurred in the northern Spanish Meseta.

Analyses of variscite exchange patterns have concentrated nearly exclusively on source and/or bead geochemistry (Alonso et al., 1978; Blasco et al., 1990; Dominguez Bella, 2004, 1996; Edo i Benaiges and Fernández Turiel, 1997; Edo i Benaiges et al., 1995a, 1995b; Edo i Benaiges et al., 1990; Fernández Turiel et al., 1996; Querré et al., 2008). Provenance analysis and consumption patterns often pinpoint a geological source that lacks not only evidence of production but also of any local human activity during periods relevant to the beads in question. No evaluation

has been undertaken of the beads' archaeological relevance to the socio-economic likelihood for exploitation of a source in a given locale during a given period.

In contrast to settlements, mines usually do not constitute well-stratified sites; instead, they constitute a complex system of use, re-use and re-location of products (Frumkin et al., 2014). Little, if any, datable material survives in direct stratigraphic association with the mined surface. In addition, mines usually experience several periods of production. Therefore, dating mining sites is challenging and is usually based on: 1) time period-specific mine typology and/or mining technology (tool marks and debris); 2) artefacts typology; and 3) delimitation of production by dating materials found in spoil and/or immediately pre-production contexts. Dating prehistoric mines based on typology and technology is therefore difficult. In Iberia, dating has resulted in chronological frameworks that span Late Prehistory without a clear delineation into different late prehistoric periods (Domergue, 1990; Hunt, 2003).

The production chronology at Pico Centeno is controversial. Nocete and Linares (1999) identified three trench mines and Pérez Macías (2011, 2008) considered them to represent the Bronze Age and Roman copper exploitation. However, Pérez Macías (2008) argues that marks left on the extraction face by metal tools at Pico Centeno may represent soundings made in 1883 by the Mina de Cobre Santo Tomás (Jubes and Carbonell, 1920). Also, a Neolithic–Chalcolithic variscite production period has been proposed recently based on the Pico Centeno Mine 2 (PCM2) typology, the typo-technological marks on the surface of the mine, the typology of the mining tools, and indirectly by dating of contexts in which worked variscite beads, thought to originate from Pico Centeno, were found (Odriozola et al., 2010; Odriozola, 2014).

Table 1
Available radiocarbon ages for European mining resources. Calibrated ages with 2 σ uncertainty.

Lab. code	Site	¹⁴ C age	Uncertainty	$\delta^{13}\text{C}$	Calibrated age	Mineral	Reference
		yr BP	+/- yr	per mil	yr BC		
Can Tintorer	I-12730	4310	150	0	3365–2497	variscite	Villalba et al. (1986)
Can Tintorer	I-12731	5350	190	0	4594–3713	variscite	Villalba et al. (1986)
Can Tintorer	I-11786	5070	100	0	4142–3645	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-41	4970	100	0	3973–3534	variscite	Villalba et al. (1986)
Can Tintorer	CSIC-488	4710	50	0	3634–3371	variscite	Villalba et al. (1986)
Can Tintorer	CSIC-489	4940	50	0	3912–3639	variscite	Villalba et al. (1986)
Can Tintorer	I-12158	4880	100	0	3943–3379	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-42	4820	100	0	3891–3366	variscite	Villalba et al. (1986)
Can Tintorer	I-13099	4820	100	0	3701–3350	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-49	4740	90	0	3766–3091	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-30	4710	130	0	3658–3105	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-48	4690	100	0	3633–3036	variscite	Villalba et al. (1986)
Can Tintorer	UBAR-47	4610	90	0	3365–2497	variscite	Villalba et al. (1986)
Can Tintorer	I-12730	4310	150	0	3654–3039	variscite	Villalba et al. (1986)
Can Tintorer	Beta-61491	4660	110	0	3943–3538	variscite	Bosch and Estrada (1994)
Can Tintorer	Beta-72551	4930	70	0	3946–3662	variscite	Bosch and Estrada (1994)
Can Tintorer	Beta-72552	5000	60	0	4038–3715	variscite	Bosch and Estrada (1994)
Can Tintorer	Beta-72553	5100	60	0	4325–3791	variscite	Bosch and Estrada (1994)
S. Ferreres	Beta-155686	5220	110	0	3943–3674	variscite	Borrell et al. (2009)
S. Ferreres	Beta-250402	5000	40	0	3935–3657	variscite	Borrell et al. (2009)
S. Ferreres	Beta-250403	4980	40	0	3935–3657	variscite	Borrell et al. (2009)
S. Ferreres	Beta-250405	4980	40	0	3943–3704	variscite	Borrell et al. (2009)
S. Ferreres	Beta-250406	5010	40	0	3943–3704	variscite	Borrell et al. (2009)
S. Ferreres	Beta-250404	5010	40	0	5471–5322	variscite	Borrell et al. (2009)
Casa Montero	Beta-206512	6410	40	–24.2	5324–5077	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-206513	6270	40	–26.2	5468–5229	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232884	6360	40	–25.4	5358–5080	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232885	6280	40	–24.9	5466–5225	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232886	6350	40	–25.6	5367–5208	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232887	6290	40	–22.2	5310–5066	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232888	6240	40	0	5367–5208	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232889	6290	40	–22.3	5533–5371	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232890	6500	40	–25.6	5458–5215	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232891	6320	40	–26.2	5324–5077	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232892	6270	40	–26.2	5463–5217	silex	Díaz del Río and Consuegra Rodríguez (2011)
Casa Montero	Beta-232893	6330	40	–25.6	6009–5727	silex	Díaz del Río and Consuegra Rodríguez (2011)
Defensola	UTC-1342	6990	80	0	5887–5568	flint	Díaz del Río et al. (2008)
Defensola	Beta-71143	6820	80	0	4683–4362	flint	Díaz del Río et al. (2008)
Defensola	Beta-71144	5670	70	0	5666–5475	flint	Díaz del Río et al. (2008)
Defensola	Beta-80604	6630	70	0	5616–5376	flint	Díaz del Río et al. (2008)
Defensola	Beta-80603	6540	60	0	5626–5492	flint	Díaz del Río et al. (2008)
Defensola	UTC-1411	6630	40	0	5295–4860	flint	Díaz del Río et al. (2008)
Tomaszów	GrN-7594	6145	70	0	5300–4746	flint	Díaz del Río et al. (2008)
Krzemionki	Gd-1425	6090	110	0	4612–4367	flint	Díaz del Río et al. (2008)
Porco	AA-62119	5665	47	0	4829–4559	jadeite	Petrequin et al. (2006)
Porco	AA-62120	5847	47	0	5212–4910	jadeite	Petrequin et al. (2006)
Porco	AA-62121	6110	48	0	4960–4721	jadeite	Petrequin et al. (2006)
Porco	AA-62123	5959	49	0	5218–4953	jadeite	Petrequin et al. (2006)
Porco	AA-62122	6146	49	0	4934–4712	jadeite	Petrequin et al. (2006)
Porco	AA-62125	5931	48	0	5309–5056	jadeite	Petrequin et al. (2006)
Porco	AA-62124	6231	48	0	265–535 AD	jadeite	Petrequin et al. (2006)
Bule	AA-66511	1644	36	0	4340–4068	jadeite	Petrequin et al. (2006)
Bule	AA-66512	5393	42	0	4517–4353	jadeite	Petrequin et al. (2006)
Bule	AA-66513	5605	42	0	4517–4353	jadeite	Petrequin et al. (2006)
Bule	AA-66514	5662	71	0	4682–4357	jadeite	Petrequin et al. (2006)
Bule	AA-66515	6222	44	0	5303–5056	jadeite	Petrequin et al. (2006)
Bule	AA-66516	5963	61	0	4996–4716	jadeite	Petrequin et al. (2006)
Bule	AA-66517	6212	71	0	5323–4986	jadeite	Petrequin et al. (2006)
Cabrieries	Beta-156929	3830	40	0	2458–2148	copper	Ambert (2002)
Cabrieries	Beta-156928	3900	40	0	2480–2212	copper	Ambert (2002)
Araico	Beta-312351	5640	40	0	4545–4367	Silex	Tarriño et al. (2011)
Araico	Beta-312352	6050	40	0	5054–4838	Silex	Tarriño et al. (2011)
Tomaszów		6220	120	0	5467–4852	Silex	Tarriño et al. (2011)
Tomaszów		6145	70	0	5295–4860	Silex	Tarriño et al. (2011)
Tomaszów		5700	70	0	4707–4371	Silex	Tarriño et al. (2011)
Blackpatch	BM-290	5090	130	0	4230–3643	silex	Tarriño et al. (2011)
Church Hill	BM-181	5340	150	0	4460–3800	silex	Whittle et al. (2011)
Cissbury	BM-183	4720	150	0	3892–3023	silex	Whittle et al. (2011)
Cissbury	BM-184	4650	150	0	3700–2932	silex	Whittle et al. (2011)
Cissbury	BM-185	4730	150	0	3907–3026	silex	Whittle et al. (2011)
Cissbury	BM-3082	5100	60	–19.2	4038–3715	silex	Whittle et al. (2011)
Cissbury	BM-3086	4710	60	–22.1	3634–3370	silex	Whittle et al. (2011)
Harrow Hill	BM-182	4930	150	0	4040–3370	silex	Whittle et al. (2011)

(continued on next page)

Table 1 (continued)

Lab. code	Site	¹⁴ age	Uncertainty	δ ¹³ C	Calibrated age	Mineral	Reference
		yr BP	+/- yr	per mil	yr BC		
Harrow Hill	BM-2099R	5040	120	-23.1	4225–3536	silex	Whittle et al. (2011)
Harrow Hill	BM-2097R	5140	150	-25.2	4319–3651	silex	Whittle et al. (2011)
Harrow Hill	BM-2071R	4900	120	-26.7	3960–3377	silex	Whittle et al. (2011)
Harrow Hill	BM-2075R	5020	110	-26.4	4044–3539	silex	Whittle et al. (2011)
Harrow Hill	BM-2124R	5060	90	-24.9	4037–3657	silex	Whittle et al. (2011)
Harrow Hill	BM-2098R	5350	150	-25.7	4486–3802	silex	Whittle et al. (2011)
Harrow Hill	BM-3084	4880	30	-21.9	3706–3638	silex	Whittle et al. (2011)
Harrow Hill	BM-3085	5070	50	-23.4	3970–3715	silex	Whittle et al. (2011)
Long Down	OxA-1152	5050	100	0	4041–3647	silex	Whittle et al. (2011)
Spiennes	Lv-1566	5510	55	0	4458–4259	silex	Collet (2004), Collet et al. (2008)
Spiennes	GrN-4674	5420	75	0	4444–4046	silex	Collet (2004), Collet et al. (2008)
Spiennes	Lv-1598	5100	65	0	4039–3713	silex	Collet (2004), Collet et al. (2008)
Spiennes	KN-I.16	5110	40	0	3980–3797	silex	Collet (2004), Collet et al. (2008)
Spiennes	OxA-3196	4830	80	0	3778–3376	silex	Collet (2004), Collet et al. (2008)
Spiennes	Beta-194770	4580	40	0	3499–3105	silex	Collet (2004), Collet et al. (2008)
Spiennes	Beta-194771	4550	40	0	3484–3100	silex	Collet (2004), Collet et al. (2008)
Spiennes	Beta-110683	4500	50	0	3361–3027	silex	Collet (2004), Collet et al. (2008)
El Milagro (Asturias)	OxA-3005	3990	90	0	2865–2210	copper	Collet (2004), Collet et al. (2008)
El Milagro (Asturias)	OxA-3006	3850	90	0	2567–2036	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-33207	3785	35	0	2338–2051	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-33209	3775	35	0	2331–2043	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-24538	3630	40	0	2133–1892	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-24537	3520	40	0	1950–1704	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-24550	3355	45	0	1747–1527	copper	de Blas Cortina (2011)
El Milagro (Asturias)	Ua-33206	3285	35	0	1643–1460	copper	de Blas Cortina (2011)
La Profunda (León)	Ua-35778	3865	35	0	2464–2209	copper	de Blas Cortina (2011)
La Profunda (León)	Ua-35779	3950	35	0	2570–2310	copper	de Blas Cortina (2011)
La Profunda (León)	Ua-35780	4075	35	0	2858–2490	copper	de Blas Cortina (2011)
El Aramo (Asturias)	OxA-1833	4090	70	0	2872–2488	copper	de Blas Cortina (2011)
El Aramo (Asturias)	OxA-1926	3810	70	0	2466–2040	copper	de Blas Cortina (2011)
El Aramo (Asturias)	OxA-3007	3900	90	0	2623–2057	copper	de Blas Cortina (2011)
El Aramo (Asturias)	OxA-6789	3995	50	0	2833–2345	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18629	3775	65	-21.4	2456–2028	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18630	3365	60	-20.1	1872–1504	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18631	3310	65	-20.4	1743–1446	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18632	3825	60	-20.5	2467–2062	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18633	3940	60	-20.1	2580–2210	copper	de Blas Cortina (2011)
El Aramo (Asturias)	Ua-18634	3215	55	-22	1623–1327	copper	de Blas Cortina (2011)
Chiflón	BM-1600	4840	50	0	3756–3389	copper	Acosta (1995)
Chiflón	BM-1599	4780	50	0	3654–3378	copper	Acosta (1995), Castro Martínez et al. (1996)
Chiflón	OxTL-200e3(II)	4000	300	0	3353–1744	copper	Acosta (1995)
Río Tinto	BM-2337R	2650	140	0	1121–406	copper	Castro Martínez et al. (1996)
Chiflón	BM-1529	3320	130	0	1939–1297	copper	Burleigh et al. (1982), Rothenberg and Frejeiro (1980)
Chiflón	BM-1528	2650	60	0	972–590	copper	Burleigh et al. (1982), Rothenberg and Frejeiro (1980)
Chiflón	BM-1601	2520	210	0	1190–108	copper	Burleigh et al. (1982), Rothenberg and Frejeiro (1980)

The dating of variscite mines is a crucial step to evaluate variscite production, consumption, and its socio-cultural significance. As such, our paper focuses on directly dating the exploitation of PCM2 by means of the combined luminescence and radiocarbon dating of three features. One of the features is an extraction face in which we found charcoal in the base level. This may relate to the setting of fire against a rock face to weaken it and so facilitate mining. The second feature is a pit that we suggest corresponds to a later phase of mining activity, and the third consists of tailings located adjacent to the face and pit. We also use indirect dating of PCM2 through the analysis of well-dated contexts containing variscite artefacts. Both the direct and indirect chronological datasets will be at the centre of a debate focussed on the production of a tentative model for Iberian variscite production and consumption.

PCM2 direct dating

The outcrop of the Pico Centeno aluminophosphate deposit includes three opencast trench mines. The typology of the mines resembles those of prehistoric mining activity (Shepherd, 1980; Domergue, 1990; Craddock, 1995; Hunt, 1996, 2003).

We performed an excavation in PCM2, where we conducted 5 test cuts (A-to-D) in 2011 (Fig. 2). In addition to early extraction activity,

indicated by small cavities following variscite veins, and tool-marking scars created by mallets and hammerstones, we also detected an excavation made by metallic picks that could date from Roman times to the 20th century 'copper fever' (Pérez Macías, 2008, after Jubes and Carbonell, 1920).

In summary, prehistoric features at PCM2 include an access ramp cut into the rock, a central transit area, and an extraction face with small cavities. These are characterised by concave marks left by the impact of rounded-edged tools (Odriozola and Villalobos García, 2015; and see Craddock, 1995, for a detailed summary of the mining technology). A 2 m deep pit near the access area suggests additional non-prehistoric production, which cut the prehistoric facies, and has marks left on the extraction face by metal tools.

In addition to the possibility that PCM2 was mined in recent times, the mining tools found during excavations, including quartzite cobbles, chisels and picks, point to prehistoric use. This most likely relates to an early stage of variscite exploitation during Late Prehistory.

Stratigraphy

Pico Centeno, similar to many other mines in Iberia, exhibits a long history of use, re-use, and re-location of products and debris, which results in a complex stratigraphy (Fig. 3).

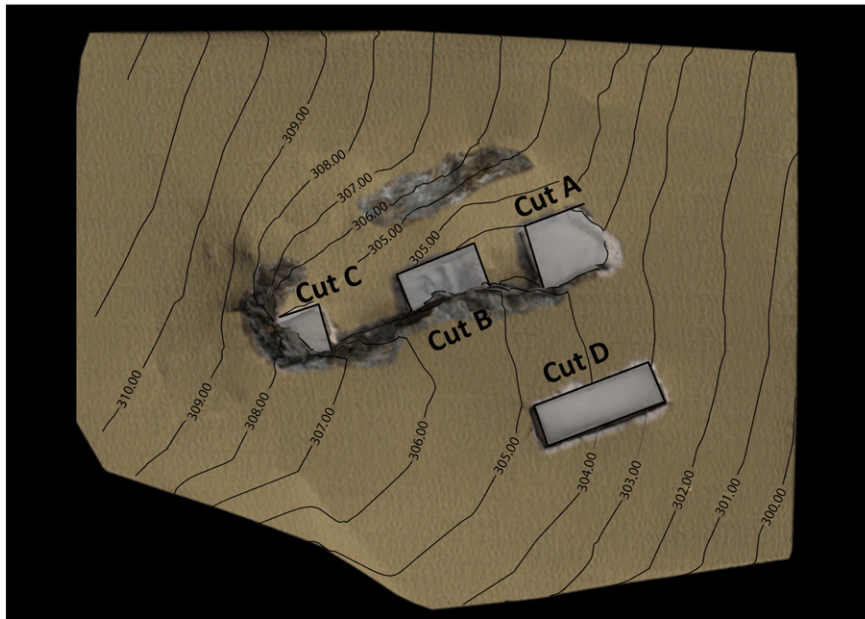


Figure 2. Test cut position and 3D model. Pico Centeno's PCM2 extraction face location is 38.161534738°N/6.949741517°W (WGS84, EPSG: 4326).

In cut B at PCM2 (Fig. 3), horizontal sedimentary units were identified through the stratigraphy. These are interpreted as floor units that are associated with the mining activities and movement of materials to the exterior of the mine. A deep excavated pit was recorded to cut these floor units (cut B, Fig. 3).

The last floor unit (test cut A SU 9, test cut B SU 11 and test cut C SU 12/13) used before the abandonment of the mine contained numerous stone tools such as picks and wedges that show strong use wear, small production debris with concave marks, and charcoal remains that adhere to both the extraction faces and the floor. We argue that the

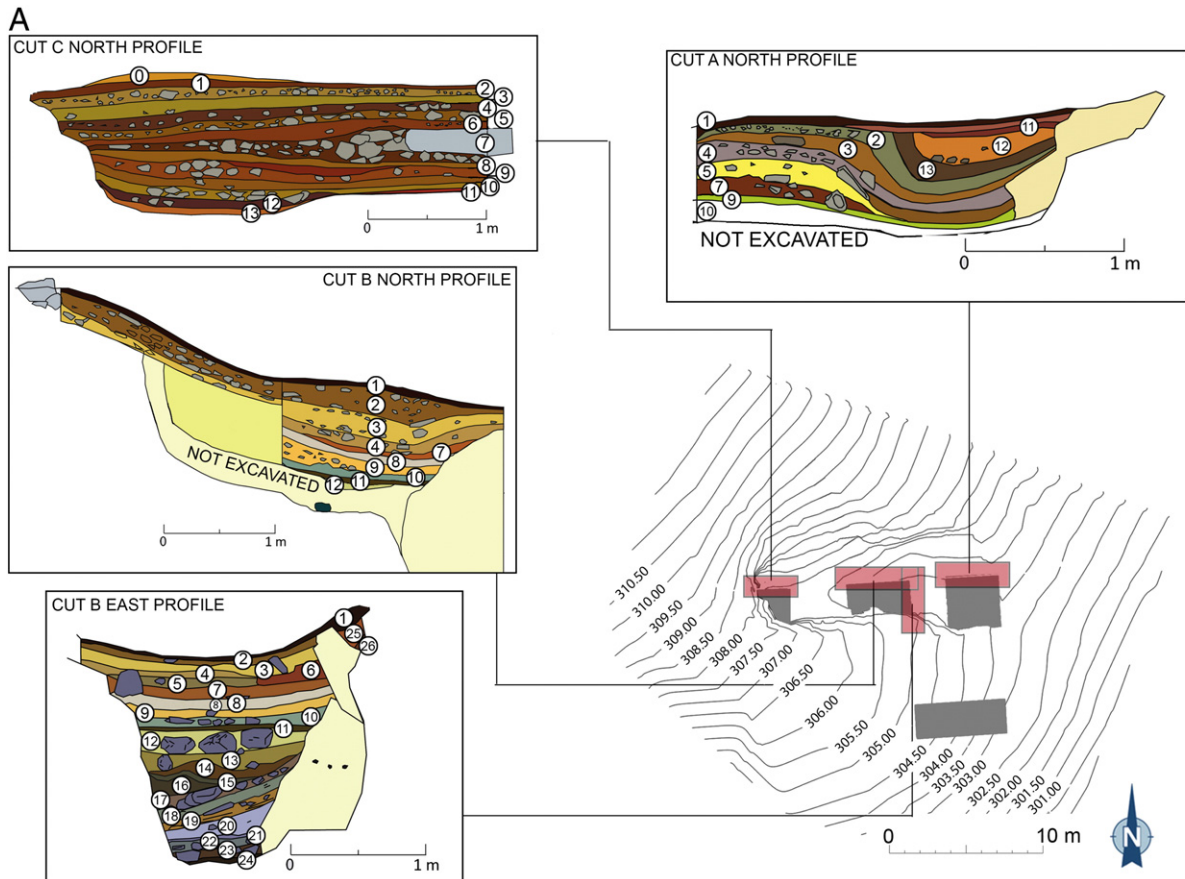


Figure 3. A) PCM2 planimetry and cuts A, B and C North profiles and cut B East profile stratigraphies; B) Harris matrix of cut B (software matrix Harris composer).

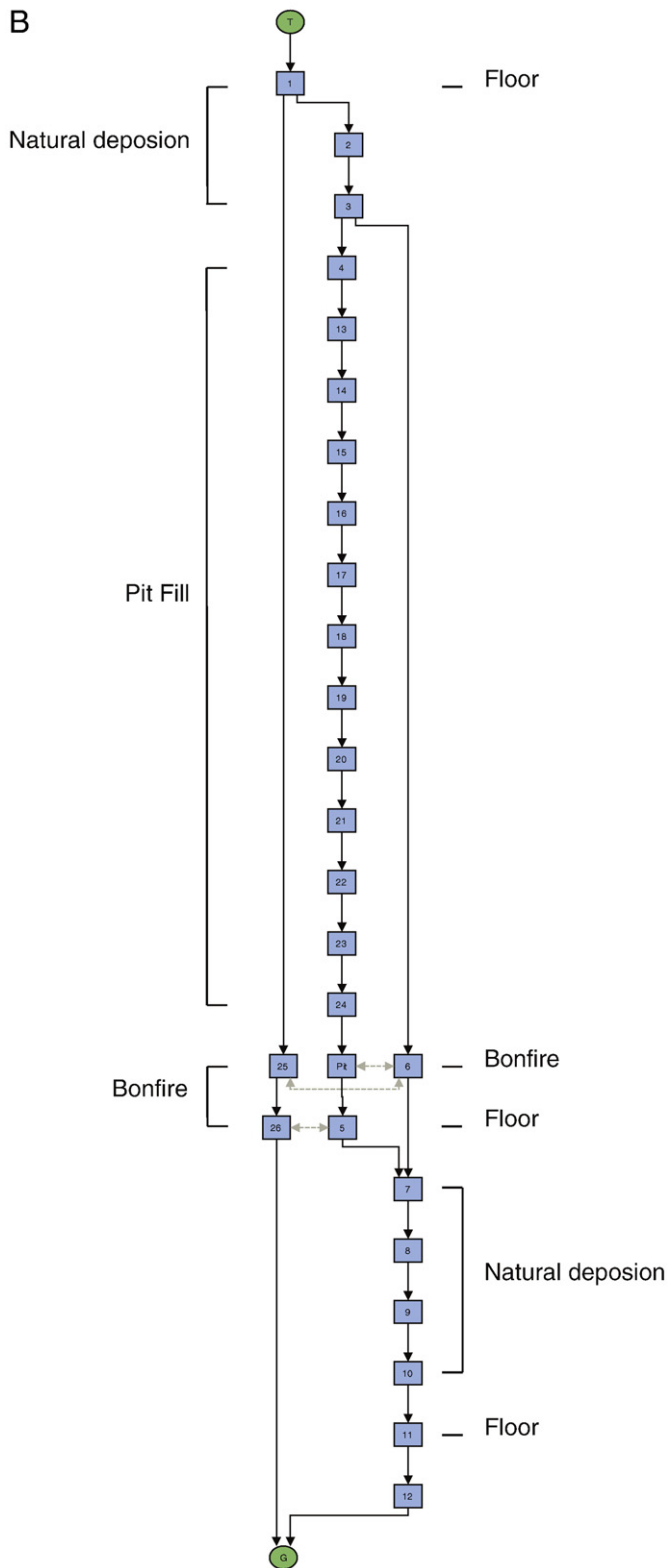


Figure 3 (continued).

abundance of charcoal in the base level and on the surface indicates the use of fire-setting technology (Willies, 1994; Craddock, 1995; Weisgerber and Willies, 2000). The floor units were sealed by apparently rapid accumulations of large and medium size rock blocks, which indicates human infilling with debris. The rubble was sealed by several apparently slower, natural, accumulation units, which were in turn

sealed by a recent compact ground surface layer that is similar to the floor units mentioned above.

A pit that cuts all units from the approximate actual surface level was encountered in test cut B. The base of this pit contained several horizontal fill units (SU 21-to-24) indicative of gradual infill by slopewash after abandonment. The mid sequence contained a series of oblique stony units that may indicate an episode of intentional human infill [SU 16–20], and this infill were capped by several relatively fine-grained sub-horizontal units indicative of gradual (natural) deposition [SU 13–15] (Figs. 3 and 4).

Luminescence dating

In this study, two complementary approaches to luminescence dating analysis were used. Semi-quantitative luminescence profiling combines simple sample preparation with multi-signal measurement: infrared- (IRSL), optically- (OSL) and thermally- (TSL) stimulated luminescence, of numerous small samples from around a site. This enables evaluation of dominant luminescence signals to guide subsequent sample preparation and fully quantitative analysis; it provides indications of whether material has been heated or not, and efficiently produces semi-quantitative estimates of absorbed dose with elevated stratigraphic resolution (Sanderson et al., 2001; Burbidge et al., 2007; Rodrigues et al., 2013; Odriozola et al., 2014). Thirteen profiling samples were taken using stainless steel tubes 2 cm diameter and 5 cm long from the cut B section (Figs. 2 and 4), plus two from the base of cut C (Figs. 2 and 3). The unexcavated volume of material remaining in cut C was too small for a quantitative dating tube: quantitative dating analyses of a lesser number of larger samples were made using OSL. These comprised three tubes, 3.8 cm in diameter and 15 cm long, of sediment and a piece of rock from cut B (units 6, 11, 13 & 18 in Fig. 4; Table 2), plus a tube of sediment from cut D (ITNLUM 701).

The east section of test cut B was chosen for OSL dating and profiling because it was the deepest available stratigraphic section at the site (Fig. 4) and was initially expected to include both prehistoric facies and the more modern pit fills. Samples for quantitative OSL dating were taken from the least stony layers (Table 2). In addition, one rock was collected to test whether the tailings included material to which fire had been set to facilitate mining. The remaining stratigraphic units were sampled for semi-quantitative luminescence profiling to test the severity of the residual signals in the stonier layers and/or to help delimit the phases of accumulation. The two additional profiling tubes were extracted from the remaining regolith at the base of cut C, close to the ^{14}C sample locations, to test the chronological relations between the cuts and between the OSL and ^{14}C dating results. The sample taken for quantitative analysis from the south section of cut D (ITNLUM 701) was designed to help evaluate the chronological relation between the fills in cut B and the layers of tailings spread around the mine site.

Luminescence and dosimetric measurements

In luminescence dating, the dose of ionising radiation absorbed by a crystal (absorbed dose, measured in Gy) since a heating or light exposure event of sufficient severity, is evaluated by comparing TSL or OSL signal from the as-prepared sample with those from laboratory irradiations (Burbidge, 2012, 2015). Comparison of doses from TSL and OSL can thus indicate whether a sample was heated or exposed to light. Assuming that the absorbed dose resulted from long-lived radionuclides in a fixed geometry, the dose rate (measured in Gy/ka) may be calculated from parent radionuclide concentrations of the sample or in situ dose rate measurements. Luminescence measurements were made on 3 Risø DA-15 and DA-20 readers using integrated $^{90}\text{Sr}/^{90}\text{Y}$ irradiators calibrated relative to the primary ^{60}Co standard of LPSR and CTN metrology laboratory (Reader 1, 75 ± 4 mGy/s β ; Reader 2, 95 ± 3 mGy/s β ; Reader 3, 111 ± 3 mGy/s β).

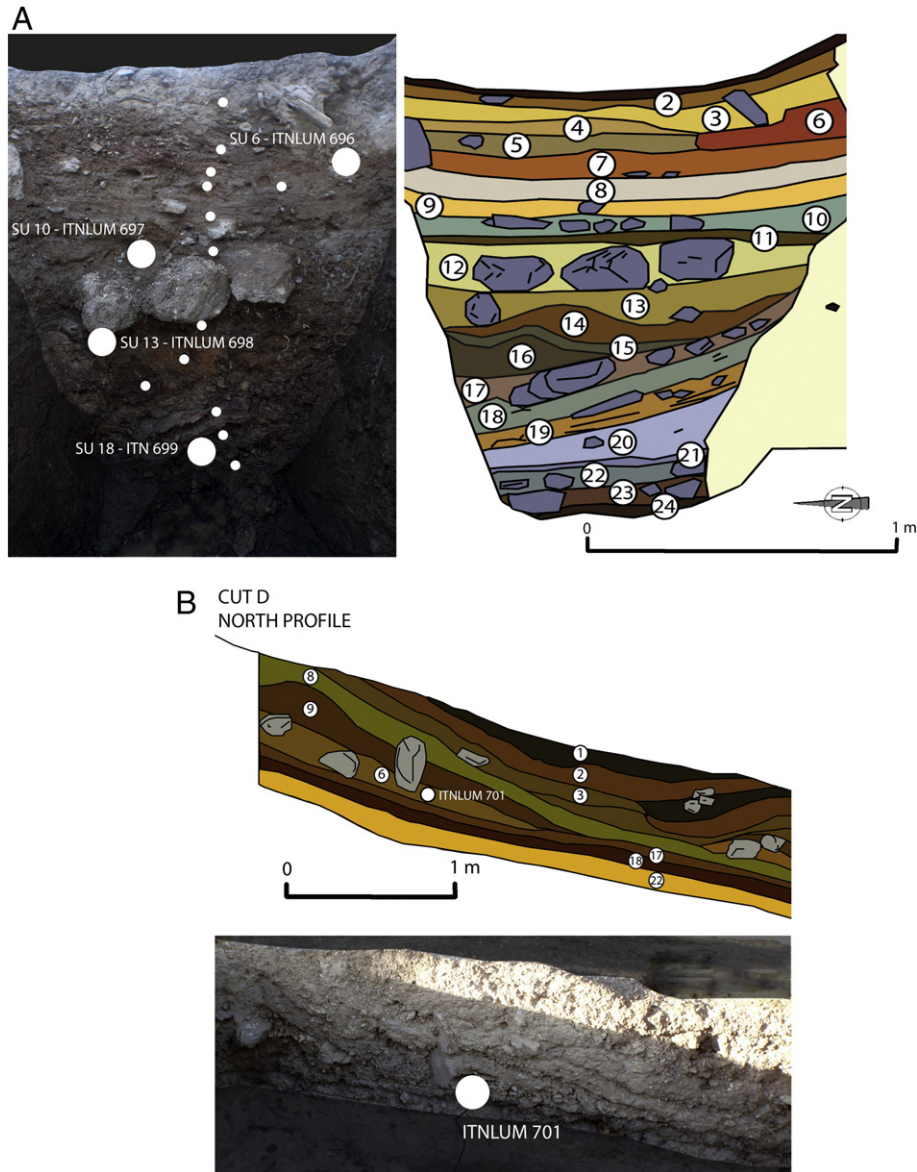


Figure 4. A) Cut B East profile stratigraphy and B) cut D North profile stratigraphy, with the detailed quantitative OSL sample position (big dots) and semi-quantitative OSL profiling (small dots).

For semi-quantitative luminescence profiling (Sanderson et al., 2001), a basic series of preparatory treatments was used to separate coarse (90–250 μm) 40% HF etched fractions enriched in quartz; they were measured using a simple multi-stimulation protocol (Rodrigues et al., 2013; Odriozola et al., 2014). The profiling approaches referred

to above have calibrated signals from the as-prepared material on an aliquot-to-aliquot basis, using signals that resulted from a single regenerative dose in the quasi-linear region of the sample's dose response. In the present case, the as-prepared signals from the profiling samples varied strongly and were often much greater than those produced by the calibration dose (e.g. Burbidge et al., 2007). The extrapolation of the quasi-linear calibration dose response would tend to overestimate the signal that resulted from doses in the range of 10–200 Gy, in which the effects of saturation in the dose response, particularly from quartz, would be expected to be evident. To help explain the signal saturation and to permit comparison between all results, the absorbed doses of the profiling samples were estimated by using a common saturating exponential dose response characteristic (DRC). This was obtained directly from the profiling measurements. It was defined by using the average of the standardised (Roberts and Duller, 2004) post-IR OSL responses to 50 sβ, with test dose of 10 sβ used in all analyses (Burbidge et al., 2006), all measured on Reader 3. The DRC is described by a single saturating exponential function, $I = I_{\infty}(1 - \exp(-D/\bar{D}))$ using a signal at saturation (I_{∞}) equal to the average dose of signal saturation (\bar{D}) (Burbidge, 2015), which was found to be 40 Gy. Semi-

Table 2
Brief description of the units sampled for OSL dating.

Sample #	Cut	Unit	Description
ITNLUM 696	B	6	A bonfire nearby the extraction surface
ITNLUM 697	B	10	Rock sample. This unit seems to be directly related with the last episode of prehistoric exploitation. This unit accounts for the 15% of the stone tools recovered. Units are cut by the pit.
ITNLUM 698	B	13	It is interpreted as a modern depositional unit filling the pit.
ITNLUM 699	B	18	Take part of the oblique depositional units that fills the pit with big rock blocks.
ITNLUM 701	D	6	Depositional unit, believed to be part of production debris accumulation.

quantitative age values were estimated by interpolating the total dose rates calculated for the quantitative dating samples.

In this study, sampling and analysis for quantitative luminescence dating were undertaken using the methods of Burbidge et al. (2014), which are based on a combination of instrumental neutron activation analysis (INAA, Dias et al., 2013; Dias and Prudêncio, 2007; Gouveia and Prudêncio, 2000; Prudêncio et al., 2006), high resolution gamma spectrometry (HRGS, Trindade et al., 2013), field gamma spectrometry (FGS; Trindade et al., 2014), water absorption and retention under free drainage, and OSL measurements. In the present case FGS was conducted using both Target Nanospec and HPI Rainbow Multi Channel Analysers, each with 2" × 2" NaI probes.

Water content was measured as a fraction of dry sample mass (50°C) with the sample in its as-received or 'field' state, once saturated, and following free drainage for 1 h, and 1, 2, 4 and 8 days (W_f , W_s , W_{D0-8} ; Burbidge et al., 2014). One end of each sediment sample tube was sealed with tape and the other end was closed with nylon mesh; the rock was brushed clean of loose material and weighted (W_r). Inverted tubes and the rock were soaked overnight in deionised water and weighed after removal of standing water (W_s). Tubes were then unsealed and set to drain on an inclined board; the rock was drained on an inclined sieve (W_{D0-8}). Fills that were rich in weathered pelitic host rock were highly water retentive; the clast-rich mine-waste and the sample of rock drained more completely. Differences between K and Th concentrations estimated by FGS, HRGS and INAA (Tables 3a and 3b) were explainable using attenuation by W_f for the more deeply buried samples. Representative drained values were used for samples from superficial, i.e., drier layers (ITNLUM 696, 701). Sampling was conducted in late autumn, in days following rain: well-drained ground had not been subject to prolonged wetting and was expected to retain a slightly greater than minimum water content, whereas samples from the poorly drained pit had been subject to prolonged wetting and were expected to retain greater than average water contents. On this basis the time averaged burial values for the fill and rock samples (ITNLUM 696–9) were estimated as the average of W_f for each sample and that of the driest sample (ITNLUM 701); the upper value in the average calculation for ITNLUM 701 was chosen as W_{D8} .

The mineralisation associated with the precipitation of aluminophosphates to produce variscite at PCM2 apparently also resulted in the presence of moderately elevated levels of U (Tables 3a and 3b). Therefore, to test for disequilibrium in the upper U-series, HRGS was performed for each quantitative dating analysis. 23–31 g of milled material was sealed and equilibrated in polystyrene Petri dishes. Twenty five emission lines from ^{40}K and the ^{235}U , ^{238}U and ^{232}Th decay series were mass-normalised and compared with the reference samples GSS1, GSS5, GSR6 used for INAA. Significant disequilibrium in the upper series was not apparent, and the weighted mean results over all emissions yielded similar results to the INAA (Tables 3a and 3b). However, after accounting for in situ water content, U dose rates obtained from (unsealed) FGS measurements were 10%–30% lower than INAA (parent ^{238}U) or the weighted mean HRGS (sealed). Given the relatively wet conditions in the pit at the time of sampling, the low FGS results are considered to

indicate minimum ^{222}Rn loss in the field. Since this appeared to better approximate burial conditions in the fills and mine wastes, the FGS U values were chosen for age calculations for samples ITNLUM 696, 697, 699 and 701.

The cosmic dose rate was estimated by averaging values calculated based on as-sampled burial depth and the height of the adjacent rock. Calculations were based on Prescott and Stephan (1982) and a fit to the data of Prescott and Hutton (1988).

With respect to the OSL analyses per se, the pelitic host rock produced abundant fine silica, the agglomerates of which exhibited slow OSL signal decay, poor recycling and strong recuperation in the single aliquot regenerative (SAR) OSL protocol (Murray and Wintle, 2000). Relatively small quantities of 90–160 µm size quartz were obtained for quantitative OSL analyses: this required repeated disaggregation, sieving, cleaning with HCl, density separation, HF dissolution (40%, 40 min), and re-sieving. In the analyses ITNLUM 696, 697, 699, 701, the calibration curve used the following radiation exposures: 0 (As-prepared), 20, 0, 5, 10, 40, 80, 0, 20, 20 (IR) sβ; test dose $D_T = 10$ sβ (the reader used to measure each sample is listed in Tables 3a and 3b), and the results were fitted with a single saturating exponential. Initial tests indicated that the quartz grains from sample PCM3 yielded relatively low OSL signals but high absorbed dose values. Thus, in the analysis ITNLUM 698 the calibration curve used exposures of: 0 (As-prepared), 200, 0, 800, 1600, 3200, 6400, 0, 200, 200 (IR) sβ; $D_T = 50$ sβ, and the results were fitted with a saturating exponential plus linear function. In all cases, preheats were made at 180, 200, 220, 240, 260 and 280°C/30 s to test for differences in the absorbed dose estimates as a function of the relative filling of, and/or transfer of the charges between, electron and hole traps during calibration and test irradiations and OSL measurements. All measurements that used the 180 and 280°C/30 s preheats were rejected in the analyses of ITNLUM 696, 697, 699 and 701, since systematic deviations or increased scatter in absorbed dose values were commonly observed. For ITNLUM 698, scatter in the absorbed dose estimates was great for all preheats, and the test-normalised OSL signal from the as-prepared material did not intercept the calibration curve in two cases (Tables 3a and 3b). For the accepted measurements in each analysis, the average recycling ratios ranged from 0.93 to 1.01, and from 0.96 to 1.00 (0.81, ITNLUM 698) after exposure to infrared light. The average zero dose responses were <6% of the average absorbed doses. The OSL signal per unit dose was relatively low for samples from the pit (Tables 3a and 3b), and for ITNLUM 698 it declined by 50% during the measurement sequence. Signal integrals that used the majority of the initial OSL decay as signal, with 'late' background subtraction, were therefore applied for dating calculations in all cases. Use of the initial signal gradient, i.e., counts in the third and fourth channels subtracted from those in the first two channels measured during OSL, resulted in highly dispersed datasets.

The weighted mean (1/variance) appeared consistent with the main grouping of accepted absorbed dose measurements in the analyses of ITNLUM 696, 697, 699, and 701, and was used to calculate central absorbed dose estimates for use in age calculation. However, individual

Table 3a

Luminescence dating measurements: radionuclide concentrations and estimates of water content.

PCMI	ITNLUM	Sample type ^a	Depth ^b (cm)	FGS in situ ^c			H ₂ O in situ (g/g)	HRGS lab, dry				INAA lab, dry			H ₂ O Time averaged (g/g)
				K (%)	Th (ppm)	U (ppm)		Ref.	K (%)	Th (ppm)	U (ppm)	K (%)	Th (ppm)	U (ppm)	
1	696	T	30	1.5	9.1	7.6	0.37	256	2.4	13	16	2.2	12	14	0.10
2	697	T	53	1.7	9.6	7.9	0.36	257	2.3	14	15	2.3	12	14	0.19
3	698	R	86	1.5	8.7	7.6	0.37	258	1.8	11	13	1.9	11	11	0.19
4	699	T	132	1.8	7.3	14	0.38	259	2.6	14	23	2.6	12	23	0.20
5	701	T	47	1.7	11	8.1	0.08	260	1.8	12	14	1.7	12	13	0.09
Average uncertainty				0.1	1.2	1.1	0.02		0.1	1.8	0.6	0.2	0.7	0.4	0.15

^a Tube; rock.

^b Below pre-excavation ground level.

^c As measured, i.e. not corrected for in situ water content.

Table 3b

Luminescence dating measurements: summary dose rate, absorbed dose, and calendar date estimates.

PCMII	ITNLUM	Sample Type ^a	Depth ^b (cm)	Dose rate					OSL			Absorbed dose			Calendar age		
				\dot{D}_{Cosmic}		\dot{D}_{total}		$\sigma_{\dot{D}}$	Reader	Aliquots	\bar{I}_{T1}^d	D	σ_D	Date	σ_{age}		
				(mGy a ⁻¹)	(mGy a ⁻¹)	=/ </ > / ? ^c	(Gy ⁶⁰ Co)									=/ </ > / ? ^c	
1	696	T	30	0.14	0.04	5.60	0.29	=	2	16	420	0.63	0.09	=	1900	20	AD
2	697	T	53	0.13	0.04	5.25	0.38	=	2	16	215	0.73	0.09	=	1870	20	AD
3	698	R	86	0.12	0.03	4.79	0.31	=	1	22	186	55	11	?	9000	2000	BC
4	699	T	132	0.12	0.03	6.97	0.59	=	3	16	362	1.34	0.19	>	1820	30	AD
5	701	T	47	0.22	0.01	4.79	0.27	=	3	16	2380	1.08	0.12	>	1790	30	AD

^a Tube; rock.^b Below pre-excavation ground level.^c Summary assessment of dosimetry and luminescence results. =/ </ > / ?^c: equal to/overestimates/underestimates “true” value, within quoted uncertainties. ?: potentially without uncertainties.^d Signal intensity measured in response to the first test dose in the SAR measurement sequence, averaged over all aliquots.

absorbed doses measured by OSL from the rock (ITNLUM 698) exhibited scatter to very high values, so that the weighted mean for this sample is considered merely to indicate a minimum estimate. These OSL results, and comparison of OSL and TSL in test measurements indicate that the rock was not heated: finite OSL measurements are thought to have been due to the aforementioned poor behaviour within the SAR protocol.

For age estimation, the different measurements were combined in accordance with the approach outlined in Burbidge et al. (2014). Alpha, beta and gamma dose rates from the environment that surrounded the sample location were estimated from FGS measurements, after correcting for measurement geometry and the difference between the in situ and the time averaged water contents. The attenuated environmental dose rates were combined with the self-dose rate of a volume of representative size and density for the adjunct samples taken from the holes excavated for FGS measurement. This was based on the dimensions of the holes, radionuclide concentrations from INAA and HRGS (U from FGS) on material from the adjunct sample, and the time-averaged water contents estimated from measurements on the tube samples. The attenuated dose rates from the environment and the adjunct were then combined with similarly estimated values for the samples themselves; for the tube samples the water content and radionuclide concentrations were assumed equal to the adjunct. The dose rates to the etched cores of the quartz grains measured by OSL were calculated from these values, and all were combined with calculated cosmic dose rates to estimate the total dose rates for the grains measured by luminescence. For each sample, the weighted mean OSL absorbed dose estimate from the grains was divided by the dose rate to estimate the age, which was converted an estimate the calendar age.

AMS-radiocarbon dating

Radiocarbon measurements were performed using a 1 MV accelerator mass spectrometer (AMS) at the facilities of the University of Sevilla (Centro Nacional de Aceleradores) on 5 charcoal samples recovered at the base level of test cut C (unit 12/13).

The chemical preparation of the samples followed standard procedures (Santos Arevalo et al., 2009). A Soxhlet extraction was applied using hexane, acetone and ethanol before treating the samples with the Acid–Alkali–Acid cleaning procedure (AAA). For the AAA procedure, 0.5 M HCl and 0.1 M NaOH were used, and time was carefully controlled to avoid severe losses by dissolution.

Between 7–10 mg of clean and dry charcoal was combusted at 950°C for 3 h in a vacuum-sealed quartz tube with CuO and Ag powder. The quartz tubes had been previously baked at 950°C to eliminate possible organic matter. The CO₂ produced was then reduced to graphite by adding excess H₂ and using cobalt as a catalyst. The resultant mixture of graphite and cobalt was pressed into aluminium cathodes and retained in a vacuum until measurement (Santos Arevalo et al., 2009).

The PCM2 site had little organic material suitable for radiocarbon dating: no materials from short-lived organisms materials were recovered from the excavations, and the only charcoal of sufficient size were obtained from cut C level 12/13. AMS radiocarbon ages are reported in conventional radiocarbon years (Stuiver and Polach, 1977) and calibrated using Calib 7.1 with Intcal'13 (Reimer et al., 2013) in cal. BC (2 σ uncertainty) and as BP (Table 4).

Anthracological analysis was performed on the charcoal fragments after removal of the samples for AMS-radiocarbon dating. The small size and poor preservation of the charcoal samples limited the anthracological determinations to angiosperm, and in some cases the proposal of a taxon (Table 4).

Indirect dating

Ideally, evidence should be combined from the excavations of several well-dated sites that contained beads, pendants or charm-assemblages made from variscite. Unfortunately, accurate identification of bead mineralogy is lacking for most ‘green bead’ contexts in Iberia, where many green minerals other than variscite were used for beadmaking. To realistically produce a chronological framework for variscite consumption we need first to identify bead mineralogy.

The mineralogical classification of beads by means of portable analytical devices is not straightforward and deserves a full-length paper of its own. However, we have developed a simplified approach for the purpose of helping to evaluate bead chronologies. This methodology includes mineralogical identification of beads (1392 samples from 42 different sites along the Iberian Meseta and Atlantic Façade) based on the chemical composition using an Oxford Instrument XMET-7500 portable X-ray spectrometer with a Rh tube, a silicon drift detector (SDD), and an automatic 5-position filter changer.

The identification of aluminophosphate is relatively straightforward based on Al-to-P atomic ratios. X-ray diffraction has confirmed variscite as the main crystallographic phase in almost all samples, either geological or bead-worked, with an Al-to-P atomic ratio in the compositional range of variscite [[MPO₄·2H₂O], where M = Al³⁺, Fe³⁺, Cr³⁺, V³⁺ (Larsen, 1942)], from ~1 to 1.8 (refer to Odriozola et al., 2010, and Odriozola, 2014). Thus, here we use the Al-to-P atomic ratio as an indicator of variscite as the raw material of beads. Nevertheless, turquoise, crandallite or aheylite may occur separately or as minor crystallographic phases together with the variscite (Larsen, 1942). In these cases, the Al-to-P atomic ratios need to be combined with Ca, Cu and Fe values to differentiate between minerals.

Differentiating green stones formed by sheet silicates, e.g., micas, talc-steatite, chlorite and serpentine, is more complicated. We conservatively classified beads as aluminophosphate, K-aluminosilicates, Mg-aluminosilicates, Mg-silicates or other silicates, based on their major element composition obtained by XRF: dominant P + Al = aluminophosphate; dominant K + Al + Si = K-aluminosilicate;

Table 4
Results of the AMS-radiocarbon dating and the anthracological analysis of sampled charcoals from PCM2. According to the calibration curves IntCal13 (samples of the terrestrial biosphere) of Reimer et al. (2013), and using CALIB rev 7.1 (Stuiver and Reimer, 1993) programme.

Lab. code	Specie	Sample	Context	Cut	Unit	Radiocarbon date	Calibrated date (2 σ)
CNA-2144	–	Charcoal	PCM2	C	12	5950 \pm 40	4033–4727 BC
CNA-2145	angiosperma	Charcoal	PCM2	C	12/13	2215 \pm 35	379–198 BC
CNA-2146	Cf. Cistus sp.	Charcoal	PCM2	C	12	1010 \pm 35	AD 969–1152
CNA-2147	Cf. Cistus sp., Cf. Pistacia lentiscus	Charcoal	PCM2	C	12	585 \pm 35	AD 1298–1417
CNA-2148	Cf. Quercus suber	Charcoal	PCM2	C	13	6205 \pm 40	5295–5051 BC

dominant Mg + Al + Si = Mg-aluminosilicate; dominant Mg + Si = Mg-silicate. This methodology appears adequate for the purposes of this work, where the objective is to evaluate whether green beads are made from variscite or another greenstone.

Results

Direct dating

The AMS radiocarbon dating indicates a palimpsest of organic remains at the base of cut C in the Neolithic (CNA 2144, 2148), Iron Age (CNA 2145), and Mediaeval periods (CNA 2146, 2147). The quantitative OSL results from fills indicate an accumulation of deposits in cut D and lower cut B in the late 18th to early 19th centuries AD (ITNLUM 696, 697) and in the late 19th century in upper cut B (ITNLUM 699, 701) (Tables 3a and 3b). In our study, profiling was solely conducted in the laboratory, and thus was not able to inform the sampling strategy (c.f. Burbidge et al., 2008; Sanderson and Murphy, 2009); instead, it was applied in parallel with quantitative dating to aid in the interpretation of fill phases and the intensity of accumulation mechanisms that operated during the infill of the pit, and to evaluate the relations between the sediment layers from around the site that were not amenable to sampling for quantitative analysis (e.g., Odrizola et al., 2014). A frequency plot of the profiling results (f_p , Fig. 5) shows how the mineral grains in most of the sampled layers were last exposed to light at ~7 ka (Neolithic), 0.6 ka (late Mediaeval) and 0.12 ka (Post Mediaeval). These phases correspond with the quantitative results obtained by OSL or AMS radiocarbon dating. Thus, many profiling samples that are out of stratigraphic sequence appear to contain redeposited material from the earlier phases of activity at the site, in which the OSL signal has not been reset. Such records are still useful in the interpretation of the history of the site, even though they have been removed from their original context, in the same manner as a survey or recovery of redeposited potsherds (e.g., Deckers et al., 2005; Burbidge et al., 2014).

The OSL profiling results (Fig. 5) through the stratigraphy of cut B exhibit two phases of elevated semi-quantitative age values, from samples taken in the more stony layers (120–70 and 35–25 cm). These are considered indicative of redeposition without (complete) liberation of the trapped charge that is the source of the OSL signal, and hence of rapid redeposition. The quantitative and semi-quantitative results from less stony layers associated with these, either immediately below or interleaved at approximately the same depth, indicate that their accumulation in their present location corresponds with early and late 19th century infill phases. The results from the first phase (120–70 cm) are scattered, e.g., the difference between the results from the two aliquots measured per sample is large relative to the geometric average value that is indicated by the trend line in Fig. 5; however, many of the semi-quantitative age values approximate that of the reproducible (tightly grouped pair) from the sample at the base of cut B (at 145 cm). This profiling sample, and one of the samples from cut C, both indicate a late Mediaeval accumulation age contemporaneous with the youngest AMS radiocarbon result from cut C (CNA 2147). The other profiling sample from cut C provided results that are consistent with the oldest, Neolithic, AMS radiocarbon ages (CNA 2144, 2148).

The late 19th century OSL age likely relate to soundings made in 1883 by the Mina de Cobre Santo Tomás, during the ‘copper fever’

(Pérez Macías, 2008, after Jubes and Carbonell, 1920). However, fills in the lower layers in cut B and the tailings in cut D unit 3, both relate to an earlier phase of accumulation.

The mixture of charcoals from different periods in the same unit, the base level (12/13), indicates phases of repeated use of the cut C space, until the Mediaeval period. This is also reflected by the semi-quantitative OSL results from the bases of cut C and cut B. The space surrounding the void created by the first miners was last used ~600 years ago (bonfires at the border of the pit). PCM2 is in the middle of a copper mining belt of the Ossa Morena Zone, which has been exploited since the Copper Age and was reopened at the end of the 19th century with the arrival of British mining companies.

Indirect dating

We have developed a new dataset of dated sites, combined with a bead mineralogical classification (Tables 1 and 5), which creates an ideal framework for the analysis of variscite consumption during the Late Prehistory across Iberia. Most of the contexts are burials, though the Zambujal and Penedo do Lexim sites also contain evidence for prolonged domestic occupation (Jiménez Gómez, 1995; Sousa, 2010). If the direct dating of mine contexts has limitations inherent to a continued use of the space, the dating of bead burial contexts (necropolis or settlements) is subject to difficulties of association. Most of the inventoried variscite adornments were collected in early archaeological excavations, without well-defined stratigraphic contexts. However, several bead consumption trends are evident.

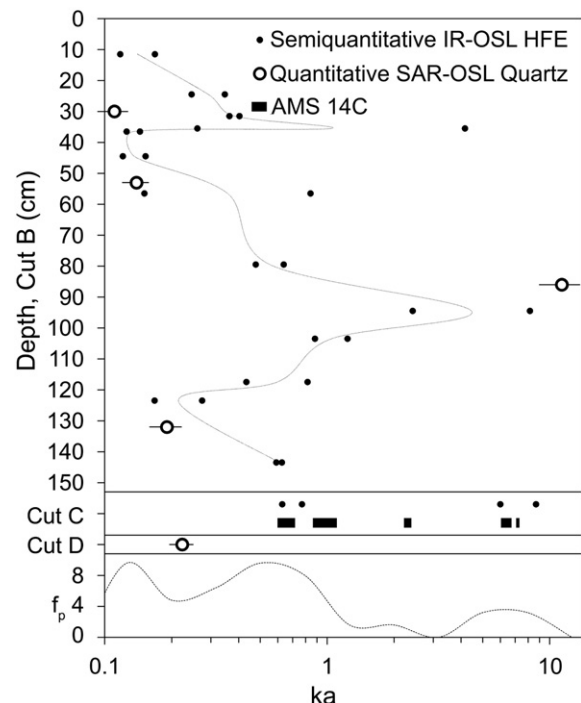


Figure 5. Chronological results from PCM2; note log time axis.

Table 5

Studied sites with available radiocarbon ages. Calibrated ages with 2 σ uncertainty.

Lab. code	Site	^{14}C age	Uncertainty	Calibrated age	$\delta^{13}\text{C}$	Reference
		yr BP	+/- yr	yr BC	per mil	
Beta-83084	ALDEAGORDILLO	4320	70	3325–2698	0	Fabián García (2006)
GrN-19168	ALDEAGORDILLO	4115	20	2859–2581	0	Fabián García (2006)
Beta-83085	ALDEAGORDILLO	4100	80	2879–2483	0	Fabián García (2006)
Beta-194313	ANTA DA HORTA	4480	40	3348–3026	–19.7	Oliveira (2010)
ICEN-1264	ANTA DAS CASTELHANAS	6360	110	5524–5054	0	Oliveira (2000)
Beta-243693	ANTA GRANDE DE ZAMBUJEIRO (corridor)	3910	40	2549–2234	0	Soares and Silva (2010)
Beta-123363	CABEÇO DA ARRUDA I	4370	70	3332–2883	0	Waterman (2012)
Beta-132975	CABEÇO DA ARRUDA I	4240	50	2999–2635	0	Waterman (2012)
GrN-5110	CARAPITO 1	4850	40	3707–3528	0	Senna-Martínez and Quintã Ventura (2000)
OxA-3733	CARAPITO 1	5125	70	4052–3712	0	Senna-Martínez and Quintã Ventura (2000)
TO-3336	CARAPITO 1	5120	40	3990–3798	0	Senna-Martínez and Quintã Ventura (2000)
OxA-5506	CASA DA MOURA	4600	90	3631–3030	0	Cardoso et al. (1996)
TO-953	CASA DA MOURA	5990	60	5019–4725	–19.6	Cardoso et al. (1996)
TO-2092	CASA DA MOURA	4850	100	3933–3372	–19.3	Cardoso et al. (1996)
TO-2093	CASA DA MOURA	5070	70	4031–3702	–19.2	Cardoso et al. (1996)
TO-2094	CASA DA MOURA	5020	70	3960–3663	–19.6	Cardoso et al. (1996)
ICEN-802	CASA DA MOURA (1A)	6100	70	5217–4841	0	Cardoso et al. (1996)
Beta-68667	CASTILLO DE ALANGE (c/ Umbria 3/ N. II)	3080	90	1520–1056	0	Pavón Soldevila (1994)
CNA-346	CASULLO	4410	50	3330–2909	0	Linares Catela and García Sanjuán, (2010)
Beta-277240	CHOUA NOVA	5450	40	4359–4237	0	Dominguez Bella and Bóveda (2011)
UBAR-593	COVA DA MOURA	4715	50	3634–3372	0	Silva (2003)
UBAR-536	COVA DA MOURA	3950	60	2619–2213	0	Silva (2003)
ICEN-1040	DOLMEN DA PEDRA BRANCA (chamber)	4620	60	3628–3106	–19.7	Soares (2010)
ICEN-1041	DOLMEN DA PEDRA BRANCA (corridor)	4120	60	2880–2497	–20.06	Soares (2010)
Beta-80602	DOLMEN DE ALBERITE	5320	70	4327–3991	0	Stip and Tamers (1996)
Beta-80600	DOLMEN DE ALBERITE	5110	140	4245–3641	0	Stip and Tamers (1996)
Beta-80598	DOLMEN DE ALBERITE	5020	70	3960–3663	0	Stip and Tamers (1996)
Poz-55021	EL GUIJO	4695	35	3629–3370	0	Villalobos García (2014)
Teledyne 19080	EL POZUELO (tomb 6)	3580	120	2283–1625	0	Linares Catela and García Sanjuán (2010)
GrN-18875	EL TOMILLAR	3925	40	2563–2291	0	Fabián García (2006)
GrN-18669	FUENTEPECINA II	5375	45	4333–4056	0	Delibes de Castro and Rojo Guerra (1997)
GrN-16073	FUENTEPECINA II	5170	100	4239–3714	0	Delibes de Castro and Rojo Guerra (1997)
Beta-176899	HERDADE DOS CEBOLINHOS (ANTA 2/ chamber)	3900	40	2480–2212	0	Gonçalves (2003)
Beta-177471	HERDADE DOS CEBOLINHOS (ANTA 2)	3840	40	2461–2154	0	Gonçalves (2003)
Beta-121143	LA PIJOTILLA (T3 UE 15)	4130	40	2872–2581	0	Odriozola et al. (2008)
ITN A6/209	LA PIJOTILLA (T3/ layer 16/ vessel 15)	2716	96	1188–561	0	Odriozola et al. (2008)
CNA-034	LA PIJOTILLA (T3/ layer 18)	4168	55	2890–2585	0	Odriozola et al. (2008)
UGAMS 8455	LAPA DA RAINHA	4080	25	2851–2498	0	Waterman (2012)
GrN-5628	LAPA DO BUGIO	4850	45	3711–3523	0	Silva and Wasterlain (2010)
OxA-5507	LAPA DO BUGIO	4420	110	3495–2780	0	Silva and Wasterlain (2010)
Beta-142035	LAS MINITAS (tomb 15)	3430	50	1881–1628	0	Pavón Soldevila (1994)
ICEN-674	LECEIA	4370	60	3326–2888	0	Cardoso (2014)
Wk-34424	LECEIA	3833	26	2455–2155	0	Cardoso (2014)
ICEN-95	LECEIA (C2)	4370	60	3326–2888	1.34	Cardoso (2014)
ICEN-89	LECEIA (C2)	4200	70	2915–2579	–19.91	Cardoso (2014)
ICEN-92	LECEIA (C2)	4120	80	2887–2489	–24.56	Cardoso (2014)
ICEN-102	LECEIA (C2)	3970	70	2838–2210	1.68	Cardoso (2014)
ICEN-1217	LECEIA (C2)	4020	80	2868–2307	–22.64	Cardoso (2014)
ICEN-1220	LECEIA (C2)	4030	70	2867–2348	–20.05	Cardoso (2014)
ICEN-737	LECEIA (C2)	3920	70	2579–2155	–19.56	Cardoso (2014)
ICEN-315	LECEIA (C2)	3730	170	2618–1668	–21.19	Cardoso (2014)
ICEN-1213	LECEIA (C2)	3970	70	2838–2210	–23.21	Cardoso (2014)
ICEN-1218	LECEIA (C2)	3910	60	2567–2206	–23.27	Cardoso (2014)
ICEN-1211	LECEIA (C2)	3900	80	2578–2141	–25.05	Cardoso (2014)
ICEN-1215	LECEIA (C2)	3900	70	2571–2150	–20.9	Cardoso (2014)
ICEN-1216	LECEIA (C2)	3880	80	2572–2136	–21.22	Cardoso (2014)
ICEN-1214	LECEIA (C2)	3840	110	2578–1972	–26.21	Cardoso (2014)
ICEN-314	LECEIA (C2)	3770	130	2571–1783	–25.75	Cardoso (2014)
ICEN-91	LECEIA (C3)	4130	60	2883–2501	–20	Cardoso (2014)
ICEN-673	LECEIA (C3)	4130	100	2915–2465	–24.9	Cardoso (2014)
ICEN-675	LECEIA (C3)	4100	90	2890–2469	–25.4	Cardoso (2014)
ICEN-1173	LECEIA (C3)	4170	50	2890–2589	–20.5	Cardoso (2014)
ICEN-1175	LECEIA (C3)	4090	80	2876–2476	–19.85	Cardoso (2014)
ICEN-1176	LECEIA (C3)	4090	60	2872–2489	–20.02	Cardoso (2014)
ICEN-1177	LECEIA (C3)	4050	50	2858–2469	–21.12	Cardoso (2014)
LY-4205	LECEIA (C3)	4030	120	2889–2210	–21.2	Cardoso (2014)
ICEN-1160	LECEIA (C4)	4630	60	3631–3114	0	Cardoso (2014)
ICEN-312	LECEIA (C4)	4530	100	3515–2924	0	Cardoso (2014)
ICEN-316	LECEIA (C4)	4520	70	3496–2940	0	Cardoso (2014)
ICEN-312	LECEIA (C4)	4530	100	3515–2924	–20.22	Cardoso (2014)
ICEN-313	LECEIA (C4)	4520	130	3626–2903	–22	Cardoso (2014)
ICEN-316	LECEIA (C4)	4520	70	3496–2940	–22.02	Cardoso (2014)
ICEN-1275	LECEIA (EN)	3950	90	2858–2149	0	Cardoso (2014)

(continued on next page)

Table 5 (continued)

Lab. code	Site	¹⁴ C age	Uncertainty	Calibrated age	$\delta^{13}\text{C}$	Reference
		yr BP	+/- yr	yr BC	per mil	
ICEN-1241	LECEIA (EN)	3950	90	2852–2149	0	Cardoso (2014)
Beta-260295	LECEIA (EN)	3840	40	2461–2154	0	Cardoso (2014)
Beta-260296	LECEIA (EN)	3980	40	2617–2347	0	Cardoso (2014)
Sac-1317	LECEIA (FM)	4220	50	2914–2634	0	Cardoso (2014)
Sac-1317	LECEIA (FM)	4220	50	2914–2634	0	Cardoso (2014)
Beta-260297	LECEIA (FM)	4140	40	2876–2585	0	Cardoso (2014)
Beta-260299	LECEIA (FM)	4100	40	2869–2498	0	Cardoso (2014)
Sac-1317	LECEIA (FM)	4220	40	2907–2675	0	Cardoso (2014)
Beta-260297	LECEIA (FM)	4140	40	2876–2585	0	Cardoso (2014)
Beta-260299	LECEIA (FM)	4100	40	2869–2498	0	Cardoso (2014)
ICEN-1160	LECEIA (layer 4)	4630	45	3624–3139	–21.81	Cardoso (2014)
ICEN-1161	LECEIA (layer 4)	4440	50	3336–2924	–23.39	Cardoso (2014)
ICEN-1159	LECEIA (layer 4)	4430	50	3334–2919	–21.35	Cardoso (2014)
ICEN-1158	LECEIA (layer 4)	4320	50	3090–2877	–21.45	Cardoso (2014)
Beta-185650	LOS GABRIELES (dolmen 4/ chamber 2)	3700	50	2273–1945	0	Linares Catela and García Sanjuán (2010)
Beta-185648	LOS GABRIELES (dolmen 4/ chamber 3)	3850	40	2460–2205	0	Linares Catela and García Sanjuán (2010)
Beta-185649	LOS GABRIELES (dolmen 4/ chamber 4)	3920	50	2567–2213	0	Linares Catela and García Sanjuán (2010)
I-16150	LOS ITUEROS	4120	130	3011–2300	0	Fabián García (2006)
I-83088	LOS ITUEROS	3960	90	2857–2155	0	Fabián García (2006)
I-16149	LOS ITUEROS	3850	100	2574–2028	0	Fabián García (2006)
OxA-5535	MARMOTA 2	4600	55	3520–3104	0	Gonçalves (2002)
Sac-2122	MOITA DA LADRA	3700	50	2273–1945	0	Cardoso and Caninas (2010)
Sac-2123	MOITA DA LADRA	3700	50	2273–1945	0	Cardoso and Caninas (2010)
Sac-2370	MOITA DA LADRA	3930	80	2831–2146	0	Cardoso and Caninas (2010)
Sac-2371	MOITA DA LADRA	3810	60	2463–2049	0	Cardoso and Caninas (2010)
ICEN-957	OLIVAL DA PEGA 2B (phase VF1/OP4)	4130	60	2883–2501	0	Gonçalves (2002)
ICEN-956	OLIVAL DA PEGA 2B (phase VF3/OP2)	4180	80	2918–2497	0	Gonçalves (2002)
ICEN-955	OLIVAL DA PEGA 2B (phase VF3/OP3)	4290	100	3328–2584	0	Gonçalves (2002)
Sac-1556	PAIMOGO	4259	90	3262–2575	0	Gonçalves (2002)
UBAR-539	PAIMOGO	4139	90	2900–2488	0	Gonçalves (2002)
Beta-186854	PENEDO DE LEXIM (locus 1/ layer 19)	4080	50	2865–2483	–20.5	Sousa (2010)
Beta-142451	PENEDO DE LEXIM (locus 1/ layer 19)	3820	40	2456–2142	0	Sousa (2010)
Sac-2067	PENEDO DE LEXIM (locus 1/ layer 19)	3820	50	2458–2140	–20.74	Sousa (2010)
Sac-2156	PENEDO DE LEXIM (locus 1)	3640	40	2291–2036	0	Sousa (2010)
Beta-186855	PENEDO DE LEXIM (locus 3/ layer 19)	3760	40	2291–2036	–19.6	Sousa (2010)
Beta-175775	PENEDO DE LEXIM (locus 3b/ layer 10)	4080	40	2862–2489	–21.2	Sousa (2010)
Sac-2158	PENEDO DE LEXIM (locus 3b/ layer 7)	3880	60	2550–2148	0	Sousa (2010)
Sac-2069	PENEDO DE LEXIM (locus 3b/ layer 7b)	3930	30	2550–2300	–21.2	Sousa (2010)
Sac-2168	PENEDO DE LEXIM (locus 5/ layer 8)	3760	50	2341–2026	0	Sousa (2010)
Beta-175774	PENEDO DE LEXIM (locus 3b/ layer 16)	4100	40	2869–2498	–20.2	Sousa (2010)
W-656	PENHA VERDE	3420	200	2289–1233	0	Cardoso (2010)
W-656	PENHA VERDE	3420	200	2289–1233	0	Cardoso (2010)
Beta-260300	PENHA VERDE (22/064)	4000	40	2830–2369	0	Cardoso (2010)
Beta-276400	PENHA VERDE (ditch HUT 2)	3970	40	2578–2346	0	Cardoso (2010)
Beta-276399	PENHA VERDE (HUT 1)	3890	40	2473–2211	0	Cardoso (2010)
Beta-296578	PENHA VERDE (HUT 2)	3700	30	2198–1981	0	Cardoso (2010)
Beta-296580	PENHA VERDE (HUT 3)	3680	40	2196–1948	0	Cardoso (2010)
Beta-276398	PENHA VERDE (pavement HUT 2)	3830	40	2458–2148	0	Cardoso (2010)
Beta-327750	PERDIGÕES (Tomb 1)	4030	40	2834–2468	0	Valera et al. (2014)
Beta-327748	PERDIGÕES (Tomb 1)	4060	30	2839–2483	0	Valera et al. (2014)
Beta-327747	PERDIGÕES (Tomb 1)	4130	30	2871–2583	0	Valera et al. (2014)
Beta-308789	PERDIGÕES (Tomb 2)	3840	30	2456–2202	0	Valera et al. (2014)
Beta-308791	PERDIGÕES (Tomb 2)	4090	30	2859–2499	0	Valera et al. (2014)
Beta-308792	PERDIGÕES (Tomb 2)	3890	30	2468–2291	0	Valera et al. (2014)
Beta-308793	PERDIGÕES (Tomb 2)	3970	30	2574–2350	0	Valera et al. (2014)
OxA-5533	POÇO VELHO	4245	55	3010–2634	0	Gonçalves (2008)
OxA-5532	POÇO VELHO	4090	55	2871–2490	0	Gonçalves (2008)
Beta-244396	POÇO VELHO (F-2)	4090	40	2865–2493	–19.1	Gonçalves (2008)
Beta-245137	POÇO VELHO (F-2)	4030	40	2834–2468	–10.2	Gonçalves (2008)
Beta-244397	POÇO VELHO (F-2)	3920	40	2562–2289	–9.8	Gonçalves (2008)
Beta-244394	POÇO VELHO (F-3)	4520	40	3360–3097	–19.1	Gonçalves (2008)
Beta-245138	POÇO VELHO (F-3)	4500	40	3355–3034	–19.1	Gonçalves (2008)
Beta-244395	POÇO VELHO (F-3)	4030	40	2834–2468	–18.5	Gonçalves (2008)
OxA-5533	POÇO VELHO (SIF)	4245	55	3010–2634	–19.4	Gonçalves (2008)
Beta-244393	POÇO VELHO (SIF)	4160	50	2886–2585	–19.1	Gonçalves (2008)
Beta-178464	POÇO VELHO (SIF)	4150	40	2879–2589	–19.3	Gonçalves (2008)
Beta-244390	POÇO VELHO (SIF)	4150	40	2879–2589	0	Gonçalves (2008)
OxA-5532	POÇO VELHO (SIF)	4090	55	2871–2490	–19.6	Gonçalves (2008)
Beta-244392	POÇO VELHO (SIF)	3970	40	2578–2346	–18.8	Gonçalves (2008)
Beta-178463	POÇO VELHO (SIF)	3960	40	2575–2341	–19.7	Gonçalves (2008)
Beta-244391	POÇO VELHO (SIF)	48090	1200	–	–20.1	Gonçalves (2008)
H-2049/148	PRAIA DAS MAÇAS (chamber)	4260	60	3076–2637	0	Gonçalves (2002)
H-2048/1458	PRAIA DAS MAÇAS (chamber)	3650	60	2200–1884	0	Gonçalves (2002)
OxA-5509	PRAIA DAS MAÇAS (west chamber)	4410	75	3338–2903	0	Gonçalves (2002)
OxA-5510	PRAIA DAS MAÇAS (west chamber)	4395	60	3331–2899	0	Gonçalves (2002)

Table 5 (continued)

Lab. code	Site	¹⁴ C age	Uncertainty	Calibrated age	$\delta^{13}\text{C}$	Reference
		yr BP	+/- yr	yr BC	per mil	
Leisner & Frreira/ 1963	PRAIA DAS MAÇAS (west chamber)	4160	110	3012–2467	0	Gonçalves (2002)
Leisner & Frreira/ 1963	PRAIA DAS MAÇAS (west chamber)	3650	100	2334–1706	0	Gonçalves (2002)
CNA-342	PUERTO DE LOS HUERTOS	4070	50	2863–2475	0	Linares Catela and García Sanjuán (2010)
CNA-344	PUERTO DE LOS HUERTOS	3940	50	2572–2291	0	Linares Catela and García Sanjuán (2010)
CNA-341	PUERTO DE LOS HUERTOS	3680	50	2202–1930	0	Linares Catela and García Sanjuán (2010)
GrN-10744	QUINTA DO ANJO (H-3)	4040	70	2872–2351	0	Gonçalves (2002)
OxA-5508	QUINTA DO ANJO (H-3)	4050	60	2866–2465	0	Gonçalves (2002)
UBAR-629	S. PAULO II	3960	190	2920–1921	0	Silva (2003)
UBAR-630	S. PAULO II	3870	70	2562–2139	0	Silva (2003)
Beta-188390	S. PEDRO DE ESTORIL 1	4720	40	3634–3374	–19	Gonçalves (2005)
Beta-178467	S. PEDRO DE ESTORIL 1	3830	40	2458–2148	–19.4	Gonçalves (2005)
Beta-178468	S. PEDRO DE ESTORIL 1	3790	40	2401–2045	–19.6	Gonçalves (2005)
UGAMS 8455	THOLOS DA BORRACHEIRA	4420	25	3309–2923	0	Waterman (2012)
UGAMS 8454	THOLOS DA BORRACHEIRA	3720	25	2198–2035	0	Waterman (2012)
Ua-10831	VALE DE RODRIGO (chamber)	3905	75	2576–2146	0	Boaventura (2011)
Ua-10830	VALE DE RODRIGO (chamber)	4905	60	3910–3532	0	Boaventura (2011)
KIA-31381	VALE DE RODRIGO 3 (layer 8)	4996	29	3935–3701	0	Boaventura (2011)
KIA-27559	ZAMBUJAL (ant. phase 1)	4238	29	2909–2705	0	Kunst and Lutz (2011)
KIA-7260	ZAMBUJAL (ant. phase 1a)	4134	43	2874–2582	0	Kunst and Lutz (2011)
KIA-27563	ZAMBUJAL (ant. phase 3b)	4065	37	2854–2485	0	Kunst and Lutz (2011)
KIA-28669	ZAMBUJAL (ant. phase 4)	4001	28	2574–2471	0	Kunst and Lutz (2011)
KIA-27557	ZAMBUJAL (ant. phase 4)	3996	23	2571–2470	0	Kunst and Lutz (2011)
KIA-27558	ZAMBUJAL (phase 1)	4129	31	2871–2582	0	Kunst and Lutz (2011)
KIA-7259	ZAMBUJAL (phase 1a)	3801	43	2456–2058	0	Kunst and Lutz (2011)
KIA-7258	ZAMBUJAL (phase 1a)	3891	43	2475–2209	0	Kunst and Lutz (2011)
KIA-4509	ZAMBUJAL (phase 1a)	3960	44	2577–2308	0	Kunst and Lutz (2011)
KIA-7256	ZAMBUJAL (phase 1b)	3951	55	2617–2236	0	Kunst and Lutz (2011)
GrN-7009	ZAMBUJAL (phase 1c)	4200	40	2899–2638	0	Kunst and Lutz (2011)
KIA-7257	ZAMBUJAL (phase 1c)	3836	39	2459–2153	0	Kunst and Lutz (2011)
GrN-6671	ZAMBUJAL (phase 2)	4170	55	2891–2585	0	Kunst and Lutz (2011)
GrN-7002	ZAMBUJAL (phase 2)	4050	40	2849–2472	0	Kunst and Lutz (2011)
KIA-27561	ZAMBUJAL (phase 2)	4155	32	2878–2629	0	Kunst and Lutz (2011)
KIA-27562	ZAMBUJAL (phase 2)	4049	25	2832–2486	0	Kunst and Lutz (2011)
KN-4989	ZAMBUJAL (phase 2)	3917	50	2566–2212	0	Kunst and Lutz (2011)
KN-4990	ZAMBUJAL (phase 2)	3934	51	2573–2236	0	Kunst and Lutz (2011)
KN-4988	ZAMBUJAL (phase 2)	3980	40	2617–2347	0	Kunst and Lutz (2011)
KIA-7261	ZAMBUJAL (phase 2b/c)	3842	37	2458–2202	0	Kunst and Lutz (2011)
KIA-27564	ZAMBUJAL (phase 3)	3992	32	2578–2464	0	Kunst and Lutz (2011)
KN-I.115	ZAMBUJAL (phase 3/4)	3530	65	2029–1692	0	Kunst and Lutz (2011)
GrN-7003	ZAMBUJAL (phase 3b)	4055	40	2851–2474	0	Kunst and Lutz (2011)
GrN-7004	ZAMBUJAL (phase 3b)	3995	35	2618–2459	0	Kunst and Lutz (2011)
GrN-7005	ZAMBUJAL (phase 3c)	4055	40	2851–2474	0	Kunst and Lutz (2011)
GrN-7008	ZAMBUJAL (phase 3c)	3980	35	2579–2350	0	Kunst and Lutz (2011)
GrN-6670	ZAMBUJAL (phase 3c/4a)	4150	105	3009–2464	0	Kunst and Lutz (2011)
KIA-28668	ZAMBUJAL (phase 4)	3999	29	2574–2470	0	Kunst and Lutz (2011)
GrN-7006	ZAMBUJAL (phase 4a/c)	4090	40	2865–2493	0	Kunst and Lutz (2011)
GrN-6669	ZAMBUJAL (phase 4b)	4025	95	2874–2300	0	Kunst and Lutz (2011)
GrN-7007C	ZAMBUJAL (phase 4b)	3950	65	2622–2209	0	Kunst and Lutz (2011)
GrN-6668	ZAMBUJAL (phase 4c/d)	3625	65	2197–1776	0	Kunst and Lutz (2011)
KN-4507	ZAMBUJAL (phase 5)	3466	53	1917–1645	0	Kunst and Lutz (2011)
KN-4506	ZAMBUJAL (phase 5)	3847	34	2457–2205	0	Kunst and Lutz (2011)
KIA-27565	ZAMBUJAL (post. phase 3b)	4445	31	3333–2936	0	Kunst and Lutz (2011)
KIA-27556	ZAMBUJAL (post. phase 3b)	3965	32	2573–2348	0	Kunst and Lutz (2011)
KIA-27566	ZAMBUJAL (post. phase 5)	3467	36	1885–1692	0	Kunst and Lutz (2011)
KIA-27641	ZAMBUJAL (post. phase 5)	2381	40	740–385	0	Kunst and Lutz (2011)

The calibrated age plotted with the mineralogical diversity of bead assemblages for each site show that variscite began in the 5th millennium BC (Fig. 5). Figure 6 shows that variscite is not the main mineral used in the 4th millennium BC and that variscite in greenstone ornaments became dominant during the 3rd millennium BC, as has previously been stated for Portuguese Estremadura (Jiménez Gómez, 1995) and the northern Spanish Meseta (Villalobos García, 2012). By the end of the 3rd millennium, the use of variscite suddenly ceases in favour of the use of green micas.

Discussion

In general, variscite was exploited for beadmaking during the Late Prehistory (Villalba et al., 2001) and during Roman times, for both beadmaking and *tesselae* production (Gutiérrez Pérez et al., 2015).

Weak evidence points to an occasional use during the Bronze Age (Schubart et al., 2004) and in the 18th century (García-Guinea et al., 2000). This resource was unlikely to have been exploited during the medieval times. Pico Centeno is most likely to have been exploited during the Neolithic and Roman periods, when the archaeological record indicates that variscite was used most intensively for beadmaking.

The ages for Pico Centeno's PCM2 variscite mine together with the typological and technological criteria provide strong evidence of intermittent exploitation during specific periods during Late Prehistory, the Iron Age, the Mediaeval Age, and the Modern Era.

For the Iberian Peninsula, a generalised use of variscite is known at the beginning of Late Prehistory (Villalba et al., 2001) and for Roman times (Gutiérrez Pérez et al., 2015); its use was sporadic in modern times (García-Guinea et al., 2000). Metal mining was also of historical importance in Southwest Iberia, specifically at Encinasola and its

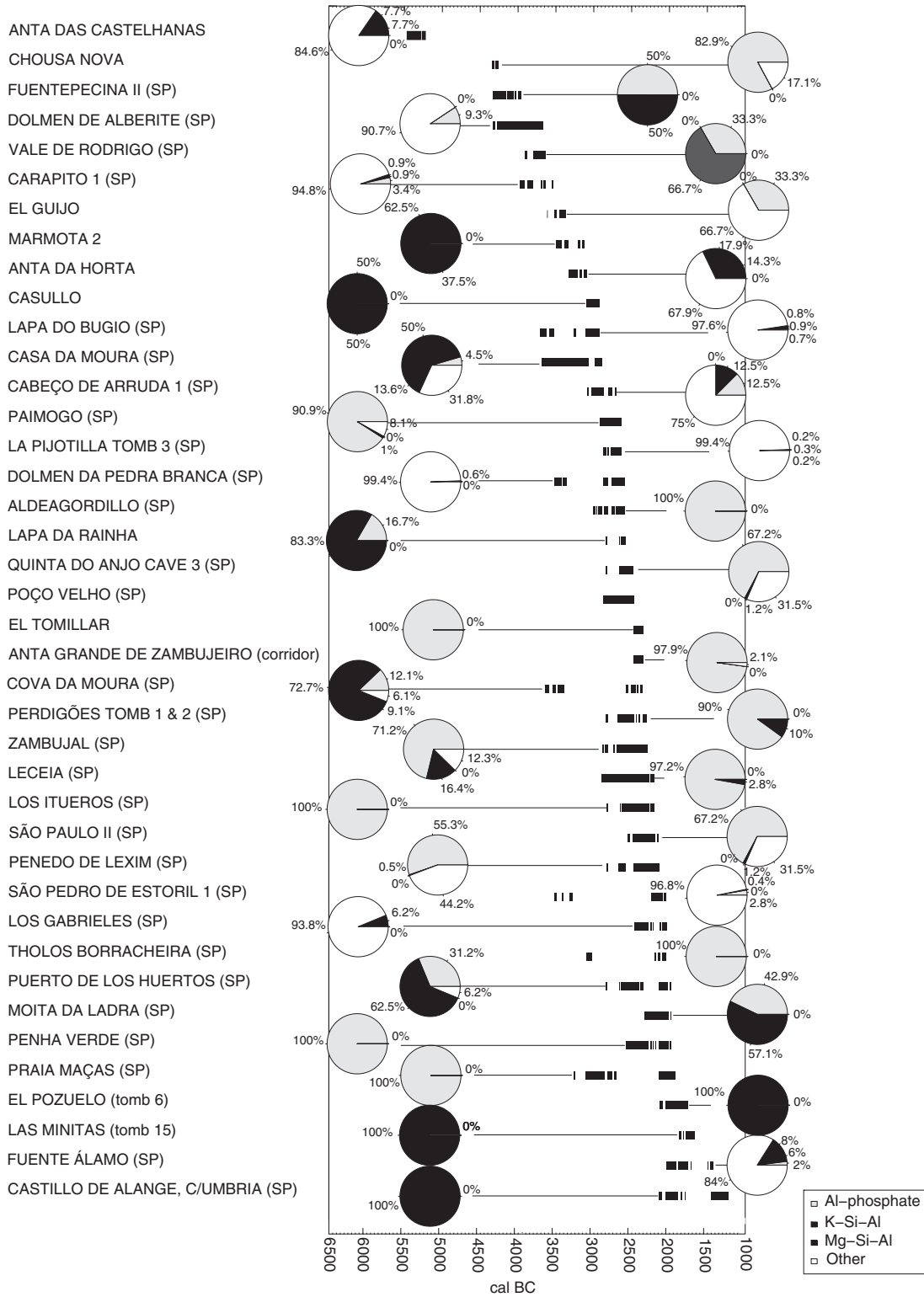


Figure 6. Radiocarbon calibrated age sum probability plot and a pie chart of mineral classification (Al-phosphate: variscite, K-Si-Al: potassium silicoaluminates, most probably green micas, Mg-Si-Al: magnesium silicates and silicoaluminates, most probably steatites and serpentinites) for studied sites.

surroundings (Pérez Macías, 2008). The 3rd century BC and 9–10th, 14th, and 18–19th centuries AD mining activities at Pico Centeno are therefore likely related to the production of metals rather than variscite.

The oldest ages recorded at Pico Centeno for Late Prehistory (5300–5000 and 4900–4700 BC) point to Early and Middle Neolithic exploitation. However, charcoal radiocarbon ages may be biased towards excessive antiquity by the old wood effect, i.e., the age of the tree and/or

the time between its death and its use as fuel and/or the time between its use as fuel and its deposition in the excavated context (Schiffer, 1986). The lack of recorded settlements from late 6th millennium BC in the Pico Centeno surroundings would, to a degree, support an old wood effect on the ages. Conversely, a closer examination of the available chronologies of Iberian sites would support an early exploitation of variscite, e.g., Cueva del Moro and Cueva de Chaves (Baldellou et al., 2012) and

likely Gruta do Caldeirão (Real, 1992) during 6th millennium BC, and Chousa Nova (Dominguez-Bella and Bóveda, 2011), Dolmen de Alberite (Sttip and Tamers, 1996), or Fuentepecina II (Rojo Guerra et al., 1996; Delibes de Castro and Rojo Guerra, 1997) during the 5th millennium BC.

According to PCM2 radiocarbon ages, quarrying and spread of the green alpine jade axe around Western Europe, with its widest spread and maximum exchange intensification at ~4500 BC, occurred coeval

in time with the initial process of discovery, production and network exchange of variscite (Pétrequin et al., 2006) (Fig. 6 and Table 1). The main difference appears to be one of scale.

From 4500 BC onwards, mainly in the 4th and first half of 3rd millennia BC, variscite mining activity greatly increased (Blasco et al., 1992), and variscite products become conspicuous in archaeological sites of the Iberian Peninsula (Guitán Rivera and Vázquez-Varela,

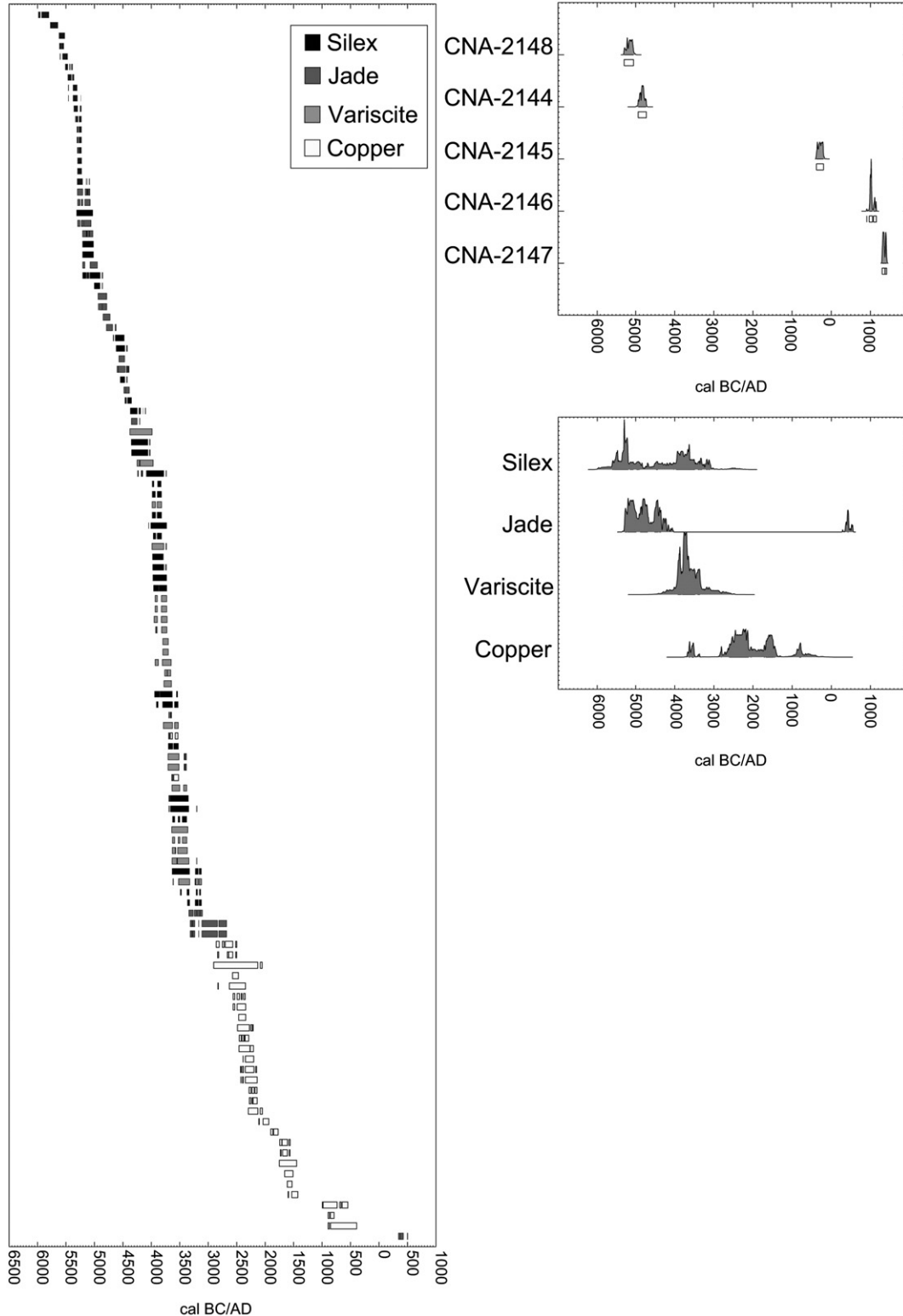


Figure 7. Calibrated data for European flint, jade, variscite and copper mining. With a detailed calibrated multi-sample probability plot of PCM2 radiocarbon ages and a sum probability plot of flint, jade, variscite and copper European radiocarbon ages.

1975; Gonçalves and Reis, 1982; Blasco et al., 1997; Bueno Ramírez et al., 2005; Costa et al., 2011; Villalobos García, 2012). Therefore, the use of variscite beads became extremely popular; achieving a period of maximum spread and use during the first half of the 3rd millennium BC, whereas the use of other green stones became rare (Fig. 7 and Table 5).

From ~2500 BC onwards variscite use began to decline, but not in favour of other greenstones, the use of which had already declined in favour of variscite (Villalobos García, 2012). Rather, this coincides with increased availability of copper-based metals (Murillo-Barroso and Montero Ruiz, 2012), and new 'exotic' resources such as Asian and African Ivory (Schuhmacher et al., 2009; Schuhmacher, 2012), and Baltic and Sicilian amber (Murillo-Barroso and Martín-Torres, 2012).

Conclusions

AMS radiocarbon and OSL dating of PCM2 indicate a long history of use, from the end of the 6th millennium BC Neolithic exploitation of variscite to the 19th century AD copper soundings made by Mina de Cobre Santo Tomás. The OSL dating indicates that fills and tailings accumulated in their present positions in the late 18th and late 19th centuries. The AMS radiocarbon dating of apparently in situ material set at the rear of the mine excavation indicated a palimpsest from the Neolithic, Iron Age and Mediaeval periods. The semi-quantitative OSL profiling results, from small samples obtained from stony layers and the remnants of excavated fills that were not amenable to sampling for fully quantitative OSL analysis, corroborate the chronological indications from both quantitative OSL and from AMS-radiocarbon. Thus, radiocarbon and OSL dating provide complementary information on different phases of site usage, which is linked by luminescence profiling.

Our new datasets support an interpretation of intermittent low intensity mining activity over a prolonged period. The intensity appears similar to that of 3rd millennium BC North Iberian copper mining, which has been calculated as 35 person-days of labour over 800 years (de Blas Cortina, 1998). The present dataset locates the beginning of variscite consumption coeval in time to the decline of jade in the 5th–4th millennium BC; the end coincides with the appearance of other signifying items in the second half of the 3rd millennium BC such as copper, ivory and extra-peninsular amber. Variscite consumption achieves its apogee in ~3000 BC, when it appears in nearly every Iberian burial (Jiménez Gómez, 1995; Villalobos García, 2012).

Acknowledgments

The authors acknowledge the Ministerio de Economía y Competitividad (HAR2012-34620) for their financial support. Odriozola acknowledges Universidad de Sevilla for a postdoctoral grant. Burbidge acknowledges PTDC/AAC-AMB/121375/2010 for implementation of gamma spectrometry.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <http://dx.doi.org/10.1016/j.yqres.2015.11.010>.

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