Introduction

An archaeological survey at the church of Nossa Senhora da Anunciada (Setúbal, Portugal) uncovered a burial site occupied between the 16th and 19th centuries. A male skeleton was retrieved, displaying a set of dry bone and radiological features compatible with a diagnosis of multiple osteochondromas (MO). This work reports the first case of MO recovered from a Portuguese archaeological site, contributing to a better portrayal of the chronological and geographic distribution of MO in the past. Moreover, the importance of the specimen herein described is the extent and number of lesions, which is in some respects analogous to modern clinical cases.

Osteochondroma (OC) is a “cartilage capped bony projection arising on the external surface of bone containing a marrow cavity that is continuous with that of the underlying bone” (Khurana et al., 2002: 234). It results from excessive chondrocyte proliferation and subsequent cartilaginous ossification at the growth plate (near the groove of Ranvier). As such, OCs develop during the first decade of life and cease after skeleton maturation (Porter and Simpson, 1999; Unni, 2001; Bovée, 2008; Khurana et al., 2002; Porter and Simpson, 1999; Fletcher et al., 2002; Unni, 2001; Bovée, 2008; Waldron, 2009). Osteochondromas can be solitary or multiple (c. 15% of patients), with this latter form termed MOs, a hereditary and autosomal dominant disorder associated with mutations on the genes EXT1 (8q24) and EXT2 (11p11–p12) (Unni, 2001; Bovée, 2008; Wuyts et al., 2014). Nowadays the global incidence of MO is approximately 1:50000 to 1:100000, showing a male predominance (Murphey et al., 2000; Unni, 2001; Khurana et al., 2002; Bovée, 2008; Wuyts et al., 2014).

The medical history of MO can be traced back to 1786, when John Hunter performed the first clinical description of this disease (Stieber et al., 2001), though paleopathological research has provided much earlier evidence of this condition, namely through a skeleton from the middle Bronze Age (c. 1700 BC) in ancient Jericho (Lyall and Mann, 1993). Murphy and McKenzie (2010) recently reviewed the paleopathological publications of MO, compiling 16 cases from a wide geographic (Africa, Americas, Asia and Europe) and chronological range (from the middle Bronze Age to the post-medieval period). However, according to this literature review (Murphy and McKenzie, 2010), the skeleton described herein is the first evidence of this disease in southern Europe.

Materials and Methods

Skeleton 37 was recovered without a coffin or other grave goods from a primary burial at the church of Nossa Senhora da Anunciada.
da Anunciada, Setúbal, Portugal in 2006. The chronology of the site was established between 1531 and 1839, based on historical and archaeological data. This site was occupied by a population of low socioeconomic status who were predominantly fishermen.

Among the 92 skeletons exhumed, consisting of 11 subadults and 81 adults (26 males, 46 females, and 9 of unknown sex), skeleton 37 was the only one inhumed in ventral decubitus (orientation east–west). This skeleton presented a fairly good degree of completeness, lacking the left fibula, left patella and left foot bones (Figure 1), yet due to the degree of fragmentation the vertebrae and sacrum were not observable. The right foot had 19.2% ($n = 5$) of the bones present, while this value was 87.0% ($n = 47$) for the hands. A total of 11 (45.8%) ribs were observable. Overall, the external bone surface was well preserved and observable.

The sex of skeleton 37 was determined from the morphological characteristics of the skull and pelvis (Buikstra and Ubelaker, 1994; Bruzek, 2002; Walker, 2008), indicating a male individual. The age at death assessment suggests that the individual was older than 30 years of age, because of the closure of all epiphysis, including the sternal extremity of the clavicle (Buikstra and Ubelaker, 1994). The analysis of the sternal end of the 4th rib points to an age interval of 28–52 years (İşcan et al., 1984). The lower limb lesions have the potential to produce gait alterations, precluding the use of the methods for age at death estimation based on the auricular surface and pubic symphysis metamorphosis. The physiological length of the left femur (386 mm) indicates an approximate stature of 150 cm (Mendonça, 2000).

Paleopathological analysis of the skeleton was performed by macroscopic observation (aided by a magnifying lens) and measurements using a sliding caliper. Digital radiographic analysis (Orthophos XG5 DS/Ceph from Sirona, 62 kV, 8 mA) of the affected bones was performed at the Clínica de Medicina Dentária, Escola Superior de Ciências da Saúde Egas Moniz.

**Results**

Skeleton 37 exhibited multiple bony projections, characterized by dense and rounded outgrowths on the external bone surface, mainly located on the metaphyseal and adjacent diaphyseal regions of the long bones (Figure 2, Figure 3, Figure 4, Table 1). The key radiological feature is the presence of well-delimited surface lesions, with both the cortex and marrow cavity showing continuity between the outgrowth and the host bone (Figure 5). These features are characteristic and compatible with the ones observed in OCs (Greenspan and Remagen, 1998; Unni, 2001; Khurana et al., 2002; Ortner, 2003; Richardson, 2005; Bovée, 2008;
Waldron, 2009). The radiographic analysis also revealed the typical flattening of the upper portion of the lesion, accompanied by an area of increased density, which is particularly noticeable on the distal right femur (Figure 5). The dry bone observation of this femoral area shows an irregular defect with darker coloration (Figure 3), which could indicate the

Figure 3. Right ((a) anterior and (c) posterior views) and left ((b) anterior and (d) posterior views) femora, exhibiting osteochondromas on multiple locations (arrows). Close up of the affected areas: (e) distal and anterior osteochondromas; (f) proximal and posterior osteochondromas, and (g) distal and posterior osteochondromas.

Figure 4. Proximal area of the right tibia ((a) posterior view) and proximal area of the right fibula ((c) lateral view) exhibiting osteochondromas (arrows). Close up of pedunculated osteochondroma on the tibia (b).
Table 1. Distribution, type, and size of the osteochondromas per anatomical area observed on skeleton 37

<table>
<thead>
<tr>
<th>Bone</th>
<th>No. OC</th>
<th>Location (no.)</th>
<th>Type</th>
<th>Size (mm)*</th>
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</table>
| Right radius     | 2      | Proximal (1)  
Anterolateral, neck of the radius | Sessile  | $6.3 \times 6.3 \times 1.1$ |
|                  |        | Distal (1)  
Medial, diaphysis             | Sessile  | Undetermined   |
|                  |        | Distal (1)  
Medial, diaphysis             | Sessile  | $18.0 \times 7.3 \times 7.4$ |
| Left radius      | 1      | Distal, medial, diaphysis             | Sessile  | $7.6 \times 4.5 \times 1.0$ |
| Right ulna       | 1      | Distal, posterior, diaphysis          | Sessile  | Undetermined   |
| Hand             | 3      | Distal, posterior, diaphysis          | Sessile  | $1.7 \times 5.7 \times 1.6$ |
|                  |        | Distal, posterior, diaphysis          | Sessile  | $2.5 \times 2.6 \times 0.8$ |
|                  |        | Distal, posterior, diaphysis          | Sessile  | $1.1 \times 1.5 \times 0.5$ |
| Right rib (3rd)  | 1      | Distal, posterior, diaphysis          | Sessile  | $18.0 \times 7.3 \times 7.4$ |
| Right os coxae   | 1      | Ischium, posterior surface (1)        | Sessile  | $16.5 \times 10.9 \times 5.5$ |
| Right femur      | 9      | Proximal (4)  
Anterolateral, adjacent to the femoral neck | Sessile  | $2.7 \times 10.5 \times 0.3$ |
|                  |        | Anterolateral, adjacent to the greater trochanter | Sessile  | $8.0 \times 26.0 \times 0.5$ |
|                  |        | Anterolateral, adjacent to the greater trochanter | Sessile  | $12.9 \times 10.4 \times 4.1$ |
|                  |        | Anterolateral, adjacent to the lesser trochanter | Sessile  | $15.9 \times 31.2 \times 9.1$ |
|                  |        | Distal (5)  
Anterolateral, diaphysis             | Sessile  | $9.5 \times 7.7 \times 0.3$ |
|                  |        | Anterolateral, metaphysis             | Sessile  | $3.8 \times 5.1 \times 1.9$ |
|                  |        | Posterolateral, adjacent to the lateral condyle | Sessile  | $23.8 \times 40.9 \times 10.9$ |
|                  |        | Posteromedial, adjacent to the lateral condyle | Sessile  | $25.6 \times 32.2 \times 2.7$ |
| Left femur       | 6      | Proximal (2)  
Anterior, femoral neck             | Sessile  | $10.6 \times 17.4 \times 5.7$ |
|                  |        | Posterior, adjacent to the lesser trochanter | Sessile  | $11.3 \times 13.3 \times 9.9$ |
|                  |        | Distal (4)  
Anterior, metaphysis             | Sessile  | $9.5 \times 7.7 \times 0.3$ |
|                  |        | Anteromedial, metaphysis             | Sessile  | $3.8 \times 5.1 \times 1.9$ |
|                  |        | Posterolateral, adjacent to the lateral condyle | Sessile  | $23.8 \times 40.9 \times 10.9$ |
|                  |        | Posteromedial, adjacent to the medial condyle | Sessile  | $25.6 \times 32.2 \times 2.7$ |
| Right tibia      | 9      | Proximal (8)  
Anteromedial, metaphysis             | Sessile  | $6.5 \times 7.2 \times 0.6$ |
|                  |        | Medial, metaphysis,                 | Sessile  | $7.4 \times 11.5 \times 5.4$ |
|                  |        | Medial, metaphysis,                 | Sessile  | $7.9 \times 11.7 \times 4.2$ |
|                  |        | Lateral, metaphysis                 | Sessile  | $14.1 \times 12.1 \times 7.3$ |
|                  |        | Posterolateral, metaphysis          | Sessile  | $12.9 \times 15.5 \times 9.1$ |
|                  |        | Posteromedial, metaphysis           | Sessile  | $13.7 \times 38.6 \times 9.3$ |
|                  |        | Medial, diaphysis                   | Sessile  | $17.2 \times 17.1 \times 7.2$ |
|                  |        | Posterolateral, diaphysis           | Pedunculated | $23.8 \times 29.8 \times 9.8$ |
| Left tibia       | 5      | Proximal (3)  
Medial, metaphysis                 | Undetermined | Undetermined |
|                  |        | Medial, metaphysis                 | Sessile  | $6.6 \times 6.3 \times 1.1$ |
|                  |        | Medial, metaphysis                 | Sessile  | $8.8 \times 14.9 \times 5.9$ |
| Right fibula     | 5      | Proximal (5)  
Medial, metaphysis                 | Sessile  | $1.3 \times 22.5 \times 6.4$ |
|                  |        | Medial, metaphysis                 | Sessile  | $5.4 \times 7.2 \times 3.8$ |
|                  |        | Anterior, metaphysis                | Sessile  | $19.2 \times 17.9 \times 5.9$ |
|                  |        | Posteromedial, metaphysis           | Sessile  | $12.8 \times 20.3 \times 7.3$ |
| Right foot       | 3      | Proximal, proximal phalanx, dorsal (1)  | Sessile  | $9.2 \times 1.3 \times 2.3$ |
|                  |        | Distal, proximal phalanx, dorsal (1)  | Sessile  | $2.5 \times 2.3 \times 1.3$ |
|                  |        | Distal, proximal phalanx, dorsal (1)  | Sessile  | $4.4 \times 3.5 \times 2.2$ |

no., number; OC, osteochondromas.

* Dimensions: lateral-medial × proximal-distal × height.
presence of the cartilage cap in vivo, as also noted by Auf­
derheide and Rodriguez-Martín (1998) in a description of
bone OC. There was also an absence of endosteal scalloping
(Figure 5).

A total of 46 OCs were observed in the skeleton 37
(Table 1). Two types of OCs were identified, i.e. the sessile
type \( n = 44 \), which was characterized by a broad base at­
tached to the cortex; and the pedunculated type \( n = 1 \),
where a narrow bone forms a pedicle (Greenspan and
Remagen, 1998; Murphey et al., 2000; Waldron, 2009;
Wuyts et al., 2014). The sole example of pedunculated OC
was detected on the right tibia (Figure 4, Table 1).

The upper limbs had a total of seven tenuous to moderate
OCs, i.e. 15.2% \( 7/46 \) of the total OCs registered, and their
distribution was bilateral for the radius (right: \( n = 2 \), left:
\( n = 1 \)), unilateral for the ulna (right: \( n = 1 \) ) and unknown for
the hands (\( n = 3 \)) (Table 1). On the upper limb eburnation
were observed at the distal articular surfaces of the left
radius and right ulna, on the left and right trapezium, the
proximal articulation of the first metacarpals and on the dis­
tal articular surface of the third metacarpals, suggesting the
presence of degenerative joint disease. Although the frag­
mentation of the bones precluded a metric analysis, there
were no apparent length asymmetries.

The lower limbs account for 80.4% \( 37/46 \) of OCs and
their severity was particularly marked on the bones that
make up the knee joints: 26.1% \( 12/46 \) and 15.2% \( 7/46 \) for
the right and left knees, respectively). The femora alone ac­
count for 15 OCs (Table 1). Both extremities of the femora
were deformed with widening of the metaphyseal region,
being more accentuated on the distal extremity (Figure 3).
All of the lesions were expressive in their dimensions. Fur­
thermore, thickening of the left femoral neck with an angle
of c. 137° was observed, indicating coxa valga deformity.
The fragmentation of the right femur precluded the assess­
ment of length asymmetries. The tibiae were the second an­
tomical elements with the highest number of OCs \( n = 14 \).
The preferential location was the proximal region. The tibial
lesions were moderate to expressive in their size (Figure 4,
Table 1). There is also evidence of genu valgum on the right
knee. On the right fibula all of the OCs \( n = 5 \) were located
on the proximal metaphysis (Figure 4, Table 1). On the right
foot, the OCs were detected on three proximal phalanges
(Table 1). Thus, the OC distribution was bilateral for the
femora (right: \( n = 9 \), left: \( n = 6 \) ) and tibiae (right: \( n = 9 \), left:
\( n = 5 \) ), being more expressive on the right side, and undeter­
mmed for the fibula \( n = 5 \) ) and foot \( n = 3 \).

Besides the limbs, the third right rib and the right ischium
were affected by OC (Table 1).

Discussion and Conclusion

The paleopathological diagnosis entails careful analyses
of the morphology and pattern of distribution of the lesions,
the age and sex of the individual, and the context from which
the specimen was retrieved (Ortner, 2003; Marques et al.,
2013).

The lesional pattern of skeleton 37 is characterized by
bony projections on the external surface of multiple bones.
The most remarkable radiological feature of these out­
growths is the continuity between the cortex and cancellous
portion of the host bone with the one of the bony projection.
This is a typology of lesion considered as pathognomonic
(Murphey et al., 2000) or highly suggestive (Greenspan and
Remagen, 1998; Unni, 2001; Richardson, 2005; Wuyts et
al., 2014) of OC. Nonetheless, other elements support the
diagnosis of MO as the most likely etiology. The external morphology of the lesions is compatible with the presentation of OC. Moreover, on the right femur the lesion exhibits an irregular defect of darker coloration, which was noted by other authors (e.g., Aufderheide and Rodríguez-Martín, 1998) as indicative of the presence of a cartilaginous cap in vivo, a distinctive element of OC (Richardson, 2003; Wuyts et al., 2014). The preferential location of the outgrowths on the metaphyseal and adjacent diaphyseal areas of long bones also supports this diagnosis. These lesions develop on the metaphyseal regions during the individual’s growth period and can migrate to the diaphysis along the bone growth (Greenspan and Remagen, 1998; Murphey et al., 2000; Ortner, 2003; Waldron, 2009). In our specimen the severity of lesions was predominant on the lower limb, particularly on the knee. According to the clinical literature, the OC most commonly occurs in the distal femur, proximal tibia and proximal fibula, with these areas affected in c. 70–90% of cases (Greenspan and Remagen, 1998; Murphey et al., 2000; Unni, 2001; Ortner, 2003; Richardson, 2005; Waldron, 2009; Clement and Porter, 2014). The cases detected on the paleopathological record suggests that the bones of the knee are most frequently affected (Murphy and McKenzie, 2010), and the pattern tends to be bilateral or symmetrical (Ortner, 2003; Wuyts et al., 2014), as observed in our case. Most of the OCs observed in the skeleton 37 were of sessile nature, which is compatible with the clinical (Richardson, 2005) and paleopathological data (Murphy and McKenzie, 2010).

Modern clinical studies suggest that the size (1–10 cm) and number of OCs vary significantly (Murphey et al., 2000; Waldron, 2009; Wuyts et al., 2014). We observed a total of 46 OCs, with a maximum size of 23.8 mm × 40.9 mm × 10.9 mm recorded in the distal part of the left femur. The size, location, number of lesions, and their sessile nature can be correlated with the presence of further skeletal alterations (Porter and Simpson, 1999; Murphey et al., 2000; Clement and Porter, 2014), such as bone deformity and hypertrophy, bone curving and length reduction, shortened stature, coxa valga, knee and ankle valgus, thickening and abnormalities of femoral neck, bone synostosis, and other clinical problems (Murphy et al., 2000; Stieber et al., 2001; Unni, 2001; Noonan et al., 2002; Richardson, 2005; Egha and Khurana, 2010; Wuyts et al., 2014). The deformities shown in skeleton 37 consisted of widening of the extremities of bones, particularly on the femora, coxa valga and genu valgum. Clinical studies refer to a prevalence of up to 25% of coxa valga and genu valgum in patients with MO (Murphey et al., 2000; Clement and Porter, 2014). Murphy and McKenzie (2010) listed 28.6% (n = 2/7) of coxa valga and 44.0% (n = 4/9) of genu valgum in the cases compiled from the paleopathological record. Shortened stature is a feature often mentioned on both clinical and paleopathological reports, which can occur in c. 40% of the individuals (Richardson, 2005; Murphy and McKenzie, 2010; Wuyts et al., 2014). The 150 cm of stature of skeleton 37 is shorter than the mean of the remaining males of the sample, which is 163.8 cm (SD = 6.4 cm, n = 10). It is also shorter than the male means of 164.3 cm (n = 21) obtained by Cardoso and Gomes (2009) from a compilation of Portuguese samples from the post-medieval period (15th–19th centuries AD) and of 163.2 cm (n = 27783) reported by Padez (2003) for the early 20th century in Portugal.

The sex and age at death profiles in our case do not seem to diverge from epidemiological data (Greenspan and Remagen, 1998; Murphey et al., 2000; Richardson, 2005; Bovée, 2008; Wuyts et al., 2014).

Although the morphological features of skeleton 37 are consistent with a diagnosis of MO, it is important to perform a differential diagnosis by analyzing other conditions that produce multiple bony projections (Unni, 2001; Richardson, 2005). Both periosteal chondromas and multiple enchondromas display bony projections; however, with these conditions there is a discontinuity between the lesion and the medullary cavity of the host bone through the intervening cortex. Moreover, enchondromas tend to be located centrally in the bone (Greenspan and Remagen, 1998; Murphey et al., 2000; Silve and Jüppner, 2006; Egha and Khurana, 2010). Therefore, these conditions are less likely the cause of the alterations observed on skeleton 37. The epiphyseal location of the exostoses on the dysplasia epiphysealis hemimelica (Trevor’s disease) also leads us to discount this cause (Murphy et al., 2000; Richardson, 2005; Glick et al., 2007). Myositis ossificans is also ruled out because the bone formations arise at the sites of insertion of muscles, tendons and ligaments, due to localized trauma, and are often found separated from the bone or still joined to it in an irregular and distorted shape. Moreover, the radiological examination reveals no continuity between the cortical and trabecular bone and the lesional tissue (Aufderheide and Rodríguez-Martín, 1998; Ortner, 2003; Richardson, 2005; Waldron, 2009).

A final note can be made regarding skeleton 37’s ventral deposition, which is atypical when compared with the remaining individuals from this site or with the archetypal Christian burials (Gonçalves and Santos, 2005). The paleopathological record demonstrated the existence of unusual burials associated with diverse skeletal pathologies or deformities (Gonçalves and Santos, 2005; Tsaliki, 2008). Even if we do not have clear evidence that the lesions of skeleton 37 were apparent during the individual’s life or that they induced a social stigma, we can hypothesize that the deformity led to the deviant burial, since MO produce osseous deformities (Murphy et al., 2000; Richardson, 2005) that can impact on the physical, social, and psychological well-being of individuals (Goud et al., 2012).

In sum, we propose MO as the diagnosis of the present specimen based on the morphological and radiological analysis. The present case report is very significant because it extends the known geographical distribution of MO to embrace the Portuguese territory and southern Europe, according to a literature review (Murphy and McKenzie, 2010). Due to the strong genetic and inherited character of MO a better assessment of its populational distribution would contribute to a better understanding of this disease.

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References


