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1 The effect of captivity on craniomandibular and calcaneal ontogenetic trajectories in wild boar

2 Short running title: Captivity and ontogenetic trajectories

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Abstract

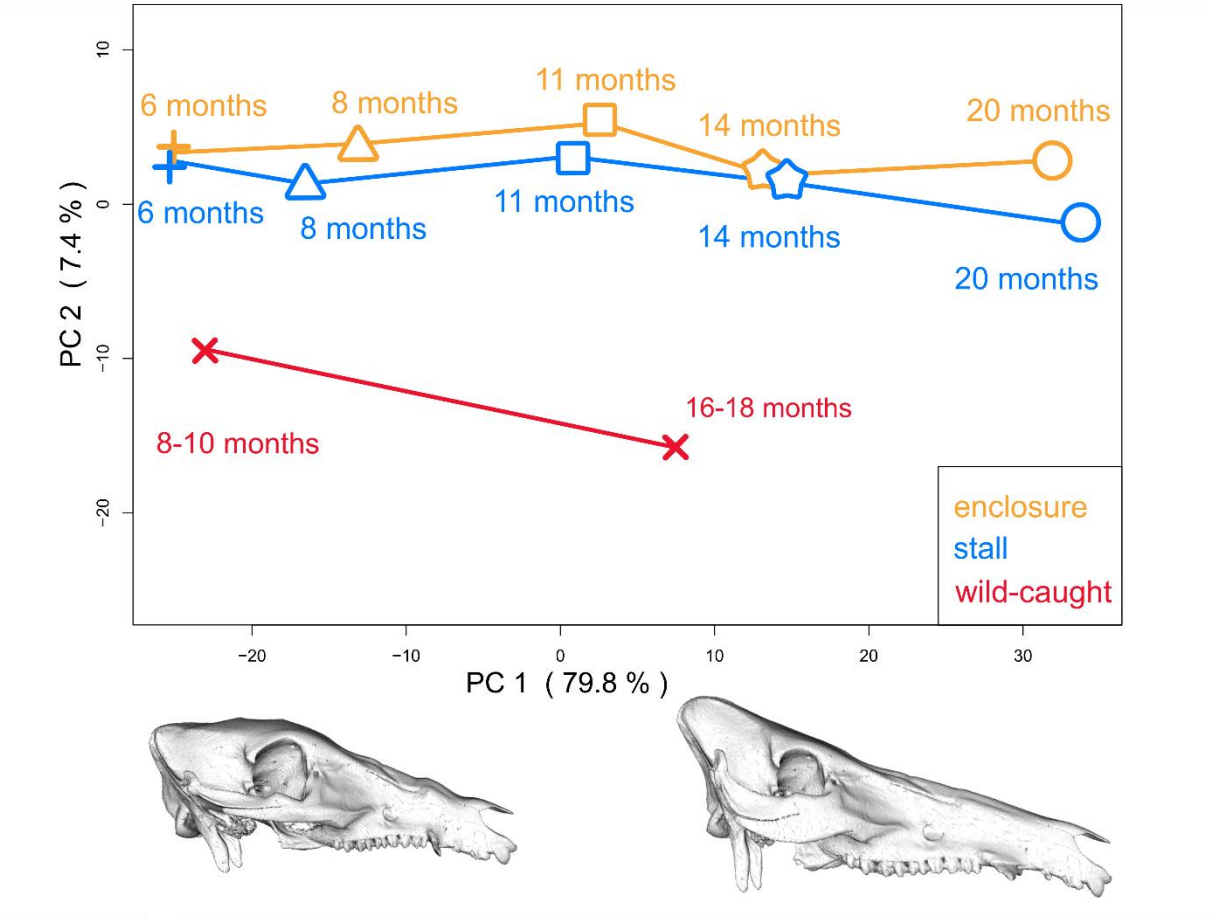
Deciphering the plastic (i.e. non-heritable) changes induced by human control over wild animals in the archaeological record is challenging. Previous studies detected morphological markers associated with captivity in the cranium, mandible, and calcaneus of adult wild boar (*Sus scrofa*) but the developmental trajectories leading up to these changes during ontogeny remain unknown. To assess the impact of growth in a captive environment on morphological structures during postnatal ontogeny, we used an experimental approach focusing on the same three structures and taxon. We investigated the form and size differences of captive-reared and wild-caught wild boar during growth using three-dimensional (3D) landmark-based geometric morphometrics. Our results provide evidence of an influence of captivity on the morphology of craniomandibular structures, as wild specimens are smaller than captive individuals at similar ages. The food resources inherent to anthropogenic environments may explain some of the observed differences between captive-reared and wild specimens. The calcaneus presents a different contrasted pattern of plasticity as captive and wild individuals differ in terms of form but not in terms of size. The physically more constrained nature of the calcaneus and the direct influence of mobility reduction on this bone may explain these discrepancies. These results provide new methodological perspectives for bioarchaeological approaches as they imply that the plastic mark of captivity can be observed in juvenile specimens in the same way it has been previously described in adults.

Keywords: ontogeny, growth, domestication, geometric morphometrics, phenotypic plasticity

49 **Research Highlights**

50 We showed the influence of captivity, an early step of domestication, on the morphology of
51 craniomandibular and postcranial structures during development. It underlines the plastic nature of
52 bony structures and their ability to change in a short time period.

53



Introduction

Animal domestication is an ongoing process (Vigne, 2011; Zeder, 2012) associated with substantial phenotypic changes that form part of the so-called domestication syndrome (Lord, Larson, Coppinger, & Karlsson, 2020; Sánchez-Villagra, Geiger, & Schneider, 2016; Zeder, 2012). Exploring the developmental mechanisms associated with the emergence of domestic phenotypes is crucial to document the roots of animal domestication over the last 15,000 years (Zeder, 2018). Zooarchaeologists previously considered that morphological changes observed in the archaeological record, such as bone shape and size modifications, were subsequent to the integration of animals into human society (Clutton-Brock, 1992) through adaptations to the new constraints of the anthropogenic environment (Price, 1999). Therefore, morphological markers have been deemed irrelevant to document the early processes of domestication (e.g. population control through captivity; Vigne, Carrère, Briois, & Guilaine, 2011), as they would only be detectable when genetic isolation and breeding selection are already in place (Frantz et al., 2015; Marshall, Dobney, Denham, & Capriles, 2014). Yet, a series of recent experimental studies with wild boar (*Sus scrofa*) have demonstrated that early domestication steps, such as the control of wild animals, can be detected and quantified. These studies further showed that a lifetime in captivity induces changes in the functional demands (e.g. locomotor, foraging or feeding behaviours), modifying the shape of craniomandibular (Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021; Neaux, Blanc, Ortiz, Locatelli, Schafberg, et al., 2021) and postcranial (Harbers, Neaux, et al., 2020; Harbers, Zanolli, et al., 2020) bony structures. More importantly, these studies showed that captivity leaves an anatomical imprint on the musculoskeletal system beyond the phenotypic variation range observed in animals in their natural habitat. These results have been confirmed by studies on reindeer (*Rangifer tarandus*) comparing wild and captive populations (Pelletier, Kotiaho, Niinimäki, & Salmi, 2020, 2021).

While previous studies detected morphological markers associated with captivity in the cranium, mandible (Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021; Neaux, Blanc, Ortiz, Locatelli, Schafberg, et al., 2021), and calcaneus (Harbers, Neaux, et al., 2020), and humerus (Harbers, Zanolli, et al., 2020) of adult specimens, the tempo of these changes during postnatal ontogeny remains unknown. Substantial differences in terms of shape and size between wild and captive animals have already been identified in mammals during growth but with contradictory results (O'Regan & Kitchener, 2005). While some studies found greater cranial dimensions in captive lion cubs (*Panthera leo*; Smuts, Anderson, & Austin, 1978) and captive-bred chinchillas (*Chinchilla lanigera*; Crossley & del Mar Miguélez, 2001), others showed a decrease in cranial dimensions in captive Indian rhinoceroses (*Rhinoceros unicornis*; Groves, 1982) and equids (*Equus* spp.; Groves, 1966). The morphological ontogenetic changes associated with captivity therefore remain to be understood.

To assess the impact of growth in a captive environment on morphological structures during postnatal ontogeny, we used an experimental approach focusing on the same bones and taxon on which morphological markers associated with captivity were detected, i.e. the cranium, mandible (Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021; Neaux, Blanc, Ortiz, Locatelli, Schafberg, et al., 2021), and calcaneus (Harbers, Neaux, et al., 2020) of wild boar. We collected weaned wild boar piglets from a genetically homogenous population and raised them in a captive anthropogenic environment from the age of 6 months. We scanned them *in vivo* at five different age classes and compared them with wild-caught wild boar populations.

To determine the influence of captivity on the growth and development of the wild boar skeleton, we compared ontogenetic changes in form (i.e. size and shape) and size between captive-reared and wild-caught specimens. We predicted that form and size differences should not differ significantly at an early age but should rather start diverging in later age classes, when the effect of captivity on morphological structures becomes more prominent. Next, we investigated differences in ontogenetic

allometry, i.e. the relationship between shape and size over the course of ontogeny at different ages (Alberch, Gould, Oster, & Wake, 1979; Klingenberg, 2016), as allometry has been shown to affect postnatal ontogenetic trajectories in domesticated clades when compared to their wild counterparts (Sánchez-Villagra et al., 2017; Wilson, 2018). We hypothesized that differences in ontogenetic allometry should be significant between captive-reared and wild-caught wild boar, indicative of a plastic effect of captivity upon developmental trajectories.

Methods

Experimental design

Captive wild boar groups (Appendix S1) consist of wild boar from the DOMEXP project: a multidisciplinary experiment aiming to assess the effect of captivity on the musculoskeletal system (<http://anr-domexp.cnrs.fr/>). They include the same specimens that were studied as adults by Neaux et al. (2021; 2021) and Harbers et al. (2020; 2020). We relied on a control population of wild boar living in a 100,000 m² fenced forest in Urciers (France). These specimens came from a wild boar farm, where human interactions are intentionally kept to a minimum in order to ensure that the behaviour of the wild boar remains as natural as possible. They are free to forage for food in the woods. From this population, we sampled 24 piglets that were divided into two groups of 12 specimens of equal sex ratio (6 males and 6 females). These groups were raised from the age of 6 months at the zoological reserve of La Haute-Touche (France) in two different contexts of mobility reduction: a 3,000 m² wooded pen ('enclosure' group) and an indoor stall of 100 m² ('stall' group). These space restrictions respectively represent a reduction of 97% and 99.9% of the range of the control population and do not allow the captive specimens to roam as freely as animals from natural populations of wild boar (Palencia et al., 2019; Russo, Massei, & Genov, 1997). We supplied individuals from both groups with processed dry food pellets including 15.5% of raw proteins adapted for domestic pig diet. These specimens were repeatedly scanned *in vivo* at the age of 6, 8, 11, 14, and 20 months, using a Computed Tomography

(CT) scanner with a spatial resolution of between 100 and 500 μm at the Chirurgie et Imagerie pour la Recherche et l'Enseignement (CIRE) platform of the Institut National de Recherche pour l'Agriculture, l'Alimentation et l'Environnement (INRAE). This experiment received ethics approval from the French Ministère de l'Enseignement Supérieur, de la Recherche et de l'Innovation (APAFIS#5353-201605111133847).

Comparative wild-caught wild boar samples

For comparison with the two captive groups, we collected free-ranging specimens ('wild-caught' group). This group included 6 individuals from the control free-ranging population of Urciers (100,000 m^2) mentioned previously. Also included in the 'wild-caught' groups are 15 free-ranging wild boar that were sampled in the forests of Chambord and Compiègne (approximated at 54,400,000 m^2 and 150,000,000 m^2 respectively; Harbers, Neaux, et al., 2020), belonging to similar geographic and climatic environment (i.e. temperate central France), to reduce the confounding effects of geographic and climate-induced morphological variation known to exist in *Sus scrofa* (Albarella, Dobney, & Rowley-Conwy, 2009; Groves, 2021; Iannucci, Sardella, Strani, & Mecozzi, 2020). Like most wild boar in western Europe, these free-ranging specimens likely had an omnivorous diet consisting mostly of plants (e.g. acorns, roots, and crops) supplemented with animal matter (e.g. insect and earthworms) as a primary source of protein (Schley & Roper, 2003). These specimens were wild-caught either between 8 and 10 months or between 16 and 18 months of age. We based age estimation on the mandibular tooth eruption and wear stages in occlusal view comparing our specimens with the charts developed by Grant (1982) and Horard-Herbin (1997). The selection of wild-caught specimens has been performed to match at best the age variation of captive specimens. Due to the inherent difficulty of collecting juvenile wild-caught specimens, the individuals from our study do not cover fully this variation, leading possibly to slight over-interpretation of results.

Data acquisition and analyses

We used homologous landmarks and semilandmarks placed on three-dimensional (3D) surfaces to describe the morphology. Digitisation and landmark definition were performed following published protocols (Appendix S2; Harbers, Neaux, et al., 2020; Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021; Neaux et al., 2020). We performed all the analyses in the R environment (R Core Team, 2019). Coordinates were aligned using a generalised Procrustes superimposition (Rohlf & Slice, 1990), implemented in the `procSym` function of the package ‘Morpho’ (Schlager & Jefferis, 2020).

We chose to work on form, i.e. the combination of size and shape (Dryden & Mardia, 1998), rather than shape as form is a more comprehensive description of an object than shape alone in the context of ontogenetic studies (Mitteroecker, Gunz, Windhager, & Schaefer, 2013). We constructed the form space by augmenting the Procrustes coordinates by the logarithm (log) of centroid size (CS; Mitteroecker, Gunz, Bernhard, Schaefer, & Bookstein, 2004). The exploratory approach of the major directions of variation in this form space relies on a principal component analysis (PCA) on form variables performed on the mean form of each age class of each of the three groups. We visualized the deformations between negative and positive scores on the first two principal components as well as the deformations between age classes for each of the three groups using heatmaps through the `meshDist` function of ‘Morpho’. The lack of congruence in age class among wild-caught groups and the two captive groups prevented us from directly comparing the ontogenetic trajectories (Adams & Collyer, 2009). We therefore tested the difference in form between captive-reared and wild-caught groups using a factorial MANOVA through the `procD.lm` function of ‘the package ‘geomorph’ (Adams, Collyer, & Kaliontzopoulou, 2019). We also tested the difference in CS and in body mass (Appendix S3) between captive-reared and wild-caught groups with a pairwise test and visualized it graphically with a bivariate plot. Body masses were measured on a scale before each scan *in vivo* for the captive-reared specimens and on uneviscerated specimens for the wild-caught ones.

To estimate and compare ontogenetic allometries between the three groups, we used a multivariate regression (Drake & Klingenberg, 2008) between shape (Procrustes coordinates) and size, computed as log CS (Collyer, Sekora, & Adams, 2015). We displayed graphically the difference in ontogenetic trajectories among our three mobility groups using a biplot of regression shape scores against log CS and tested it with a MANCOVA through the `procD.lm` function.

Results

Form

The factorial MANOVA (Table 1) showed that the ‘8-10 month wild-caught’ cranium, mandible, and calcaneus forms are not significantly different from the 6-month and 8-month old captive forms (i.e. stall and enclosure). Those structures are different from those of all the older captive groups. The ‘16-18 month wild-caught’ group is not significantly different from the 11-month and 14-month captive groups for any of the studied structures. The mandible and calcaneus of this age class do not differ from those of the 8-month ‘enclosure’ nor from the 8-month ‘stall’ groups for the mandible. They are different from those of all the other captive groups (see Appendix S4 for the Factorial MANOVA between all groups).

For the PCA (Fig. 1), PC1 for the cranium, mandible, and calcaneus show respectively 79.8%, 86.1%, and 75.5% of the total variance. For the three studied structures, PC1 displays mostly changes associated with growth as the younger specimens have the lower PC scores and the older ones have the higher scores. The cranium form changes associated with positive scores involve (1) an anteroposteriorly longer and more concave rostrum, (2) more robust zygomatic arches, and (3) smaller orbits relative to the overall cranium size (Fig 1.a). For the mandible, the associated form changes towards positive scores include (1) a reduction of the gonial angle, (2) a longer corpus and mandibular

symphysis, and (3) a mediolateral reduction of the space between the two mandibular rami in dorsal view (Fig 1.b). For the calcaneus, the changes towards positive scores consisted in (1) a shift of the sustentaculum tali and of the calcaneal sulcus towards the distal extremity, (2) a more dorsoplantarily curved calcaneus, and (3) a more elongated epiphysis that is orientated toward the plantar side (Fig 1.c). PC2 bear less variation in form for the cranium, mandible, and calcaneus (respectively 7.4%, 5.1%, and 9.1% of the total variance) but clearly distinguishes captive and wild-caught specimens. Cranial form changes along PC2 from wild-caught (low values) to captive individuals (high values) involved (1) a narrower cranium in dorsal view, specifically in the zygomatic region, (2) a less vertical occipital region, and (3) an anteroposteriorly shorter nasal and maxillary region (Fig 1.a). The mandible form changes along PC2 are characterized by (1) a mediolateral narrowing, (2) a higher ramus, and (3) a ventrally orientated ramus (Fig 1.b). The calcaneus form changes along PC2 mainly express (1) a downward shift of the sustentaculum tali and (2) an epiphysis more orientated toward the plantar side (Fig. 1.c). From 6 months to 20 months, the main changes for captive specimens are localised on the zygomatic arches and on the occipital region for the cranium (Fig. 1.d), on the ramus and on the symphysis for the mandible (Fig. 1.e), and on the epiphysis, the sustentaculum tali, and the cuboid facet for the calcaneus (Fig. 1.f). From 8-10 months to 16-18 months, the changes for wild-caught specimens are localised on the same areas but are less prominent.

Size

The size of the '8-10 month wild-caught' group is not significantly different from the 6-month and 8-month captive groups for the cranium, mandible, and calcaneus (Fig. 2; Table 2). It is different from all the older captive groups. The '16-18 month wild-caught' group is not significantly different from the 11- and 14-month captive groups for the cranium, the 8-, 11- and 14-month captive groups for the mandible, and all the captive groups from 8 to 20 months for the calcaneus. The body mass of the '8-10 month wild-caught' is not significantly different from the 6-, 8- and 11-month captive groups and

the '16-18 month wild-caught' group is not significantly different from the 11- and 14-month captive groups and the '20 months – enclosure' group. All the other groups are statistically different.

Ontogenetic allometry

We found overall ontogenetic allometry across all three structures in the three groups of mobility, which explains greater variation in calcaneus (26%) than in cranium (21%), and mandible (16%; Fig. 3). The ontogenetic allometric trajectories depicted graphically are parallel between enclosure and stall captive groups across the three structures. They differ from the wild-caught group with a more positive allometry, although the interaction term was only significant for the calcaneus, indicative of a difference in allometric growth among the three groups for this structure.

Discussion

For the cranium, mandible, and calcaneus, we found that the form and size of the '8-10 month wild-caught' wild boar are similar to those of younger captive specimens aged between 6 and 8 months, but are significantly different from the 11-month-old animals and older ones. This may suggest that the growth in captivity of a wild ungulate impacts the development of the three studied structures (O'Regan & Kitchener, 2005). Differences also exist later for the cranium and mandible, when the effect of the reduction of mobility on morphological structures becomes more prominent. Indeed, the form and size of the '16-18 month wild-caught' group are similar to the 11- and 14-month-old captive specimens but statistically different from the 20-month-old ones. The same results are observed when comparing body mass between wild-caught and captive specimens. These results are strengthened by the fact that the wild-caught control specimens, belonging to the same initial herd as the captive ones, follow the same trend as the other wild-caught wild boar (see Fig. 1.a and 1.b). Unfortunately, 6-month-old wild-caught wild boar were not available. Yet, since the '8-10 month wild-caught' group is similar to the 6-month-old captive specimens, we could speculate that 6-month-old wild-caught wild boar would be similar to

younger specimens. This suggests that captive growth in wild boar induces a morphological divergence driven by an acceleration of changes in the size and form of the skull. This developmental delay in wild-caught wild boar compared to captive ones is characterised by a retention of more juvenile traits observed in younger wild specimens as evidenced by the MANOVA. This implies an increase in the developmental rate of captive wild boar when compared to that of wild-caught specimens. These findings are in line with previous studies on postnatal growth assessing that, for a given age, captive specimens are often significantly larger than wild-caught individuals. Comparing skeletal measurements, Zihlman et al. (2007) found that the tempo of growth in wild chimpanzees (*Pan troglodytes*) contrasts sharply with the rate demonstrated for captive individuals that can mature as much as 3 years earlier. Cheverud et al. (1992) also found a significant increase of the growth rate in toque macaques (*Macaca sinica*) raised in the laboratory when compared to wild specimens. Finally, captive-raised lion cubs were reported as being nearly twice the size of wild cubs of the same age (Schaller, 1973). The more consistently available food is probably the main cause of these differences (Turner, Cramer, Nisbett, & Gray, 2016). Indeed, one of the consequences of the spatial control of wild animals by humans is the presence of more constantly available food resources. As it is the case in the context of our experimental study, animals in captivity are regularly provisioned and do not need to spend energy searching for food, hence removing intra-group feeding competition. In addition, intrasexual competition is removed in stall specimens and drastically reduced for the pen specimens. Differences in cranial size during the postnatal development were previously described between domestic pigs and wild boar, especially in the neurocranial region, surrounding and protecting cerebral structures (Evin et al., 2017). Our study stresses that this increase of the size of neurocranial structures is not necessarily a product of the long selective breeding leading to the morphology observed in domestic pigs. Indeed, we observe a similar increase as a plastic response to captivity, considered as one of the earliest domestication steps (Vigne, 2011), most likely due to an unrestricted access to food

leading to a faster growth (Kimura & Hamada, 1996). Furthermore, these findings are consistent with recent studies suggesting that the commonly assumed reduction of brain size associated with domestication should be questioned (Lord, Larson, Coppinger, et al., 2020; Lord, Larson, & Karlsson, 2020). Indeed, both early domestication steps, in the context of our study, and long term selective processes (Evin et al., 2017) result in a size increase and not a reduction of the structures surrounding the brain. Yet, although described previously (Finarelli, 2006; 2011), the presence of a clear positive relationship between brain size and cranial structures size is still unclear (Logan, & Palmstrom, 2015), highlighting the need for future studies directly addressing the evolution of endocranial shape and size in relation to domestication.

The calcaneus presents a different pattern when the specimens are older, i.e. when the effect of captivity becomes prominent as the ‘16-18 month wild-caught’ group is similar to the 11-, 14-, and 20-month-old captive specimens in size. This result is supported by the fact that the wild-caught control specimens follow the same trend as the captive ones (see Fig. 1.c). The more physically constrained nature of the calcaneus, articulating with both the talus and the cuboid bones, may partly explain this difference. In this sense, Hanot et al (2017) described a strong and significant morphological integration between the calcaneus and talus in horses (*Equus caballus*). In comparison, lower level of integration were found between the cranium and mandible of wild boar, specifically for captive individuals (Neaux, Blanc, Ortiz, Locatelli, Schafberg, et al., 2021). This comparatively loose integration in the skull may allow more size variation, and a greater and more rapid growth of these structures in captive individuals. Further covariation studies of the tarsus of less specialized taxa than horses, such as wild boar, will help untangle the role of morphological integration in the observed differences between the craniomandibular and postcranial structures.

The calcaneus is subjected to high tensile, bending and compressive forces (Su, Skedros, Bachus, & Bloebaum, 1999) and has often being described has a key proxy to assess terrestrial mammal locomotor

behaviours (Bassarova, Janis, & Archer, 2009; Ginot, Hautier, Marivaux, & Vianey-Liaud, 2016; Panciroli, Janis, Stockdale, & Martín-Serra, 2017). In this respect, the phenotypic plasticity in shape (but not the size) of the calcaneus has been shown to capture the direct influence of the anthropogenic control of wild boar locomotor behaviour (Harbers, Neaux, et al., 2020). Conversely, the morphological modifications associated with the mandible in captive animals are not the direct consequence of mobility reduction but may rather be related to functional demands resulting from the anthropogenic environment (Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021). They include the increase of feeding activity (Turner et al., 2016), the reduction of foraging behaviour (Mason & Mendl, 1997), and the appearance of stereotypy (Rhodes et al., 2005), i.e. repeated sequences of movements with no obvious purpose, particularly common in captive animals (Fraser & Broom, 1990). In this sense, in PC2 of the PCA on the form space for the calcaneus, from negative to positive values, there is a clear gradient from wild-caught, to enclosure, to stall specimens (Fig. 1.c), while for the cranium (Fig. 1.a) and mandible (Fig. 1.b) captive individuals (i.e. enclosure and stall) are similar. This result is consistent with the hypothesis that the calcaneus captures mainly locomotor behaviour as the habitats of the wild-caught group is the larger ($\geq 100,000 \text{ m}^2$), followed by the enclosure group ($3,000 \text{ m}^2$) and finally the stall group (100 m^2). The PC2 for the cranium and mandible form space primarily records changes between the captive specimens (i.e. enclosure and stall) under anthropogenic control on the one hand and the wild-caught group on the other. This result is in line with the hypothesis that craniomandibular changes are not directly associated with a reduced mobility but rather with functional requirements resulting from the anthropogenic environment (e.g. feeding activity, foraging behaviour, stereotypy). The difference in the way captivity affects the calcaneus (direct influence) and the cranium and mandible (indirect influence) may explain this distinction and should be further explored.

The slope of the ontogenetic allometry for the cranium and mandible is not significantly different between captive-reared and wild-caught specimens. This result is in line with previous findings that

ontogenetic allometry generally does not evolve on short evolutionary time scales and that modifications of ontogenetic trajectories are usually achieved by heterochronic shifts along a shared slope rather than directional changes (Voje, Hansen, Egset, Bolstad, & Pélabon, 2014; Wilson, 2018). Moreover, previous results (Sánchez-Villagra et al., 2017; Wilson, 2018) found no significant differences in the ontogenetic slopes between domestic pigs and wild boar contrary to dogs (*Canis lupus familiaris*) and wolves (*C. lupus*), and llamas (*Lama glama*) and guanacos (*L. guanicoe*). This is congruent with our results on wild boar captivity, as the control of mobility is considered one of the first steps of the domestication process leading to the morphology observed in modern pigs (Vigne, 2011). The differences in the direction of the slope for the calcaneus shows that some level of directional change can nevertheless occur on a short-time scale. The distinctions between the skull and the calcaneus may once again arise from how mobility reduction influences these structures, i.e. respectively indirectly and directly.

Conclusions

Our results provide evidence that captivity influences the morphology of craniomandibular and postcranial structures of wild boar, as wild specimens are significantly smaller than captive individuals of a similar age. Consistently available food resources and the reduction of stress associated with the search for food and intra-group feeding competition may explain the distinction between captive-reared and wild specimens. The calcaneus presents a different pattern as captive and wild individuals differ in terms of form but not in terms of size. Furthermore, it is the only structure presenting differences in ontogenetic allometry. The more physically constrained nature of the calcaneus and the direct influence of mobility reduction on this bone may explain these specificities. These results provide new methodological perspectives for bioarchaeological approaches as they imply that the plastic mark of captivity can be observed in juvenile specimens as well as in adults (Cucchi et al., 2021; Harbers, Neaux, et al., 2020; Neaux, Blanc, Ortiz, Locatelli, Laurens, et al., 2021; Neaux, Blanc, Ortiz, Locatelli,

341 Schafberg, et al., 2021). Further studies need to explore the morphological integration during growth
342 in captive conditions for both craniomandibular and postcranial structures in order to decipher the role
343 of developmental and functional correlates between structures in generating the differences observed
344 in our study.

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Conflict of Interest Statement

The authors declare no conflicts of interest.

Data Availability Statement

All analytical codes and data are freely available at: <https://zenodo.org/record/5547335> (Neaux, Harbers, Blanc, Ortiz, Locatelli, et al., 2021)

363 Adams, D. C., Collyer, M., & Kaliontzopoulou, A. (2019). geomorph: Geometric morphometric
 364 analyses of 2D/3D landmark data (Version 3.1.2).
 365 Adams, D. C., & Collyer, M. L. (2009). A general framework for the analysis of phenotypic
 366 trajectories in evolutionary studies. *Evolution; International Journal of Organic Evolution*,
 367 63(5), 1143–1154. <https://doi.org/10.1111/j.1558-5646.2009.00649.x>
 368 Albarella, U., Dobney, K., & Rowley-Conwy, P. (2009). Size and shape of the Eurasian wild boar
 369 (*Sus scrofa*), with a view to the reconstruction of its Holocene history. *Environmental*
 370 *Archaeology*, 14(2), 103–136. <https://doi.org/10.1179/146141009X12481709928283>
 371 Alberch, P., Gould, S. J., Oster, G. F., & Wake, D. B. (1979). Size and Shape in Ontogeny and
 372 Phylogeny. *Paleobiology*, 5(3), 296–317.
 373 Bassarova, M., Janis, C. M., & Archer, M. (2009). The Calcaneum—On the Heels of Marsupial
 374 Locomotion. *Journal of Mammalian Evolution*, 16(1), 1–23. [https://doi.org/10.1007/s10914-](https://doi.org/10.1007/s10914-008-9093-7)
 375 008-9093-7
 376 Cheverud, J. M., Wilson, P., & Dittus, W. P. J. (1992). Primate population studies at Polonnaruwa.
 377 III. Somatometric growth in a natural population of toque macaques (*Macaca sinica*). *Journal*
 378 *of Human Evolution*, 23(1), 51–77. [https://doi.org/10.1016/0047-2484\(92\)90043-9](https://doi.org/10.1016/0047-2484(92)90043-9)
 379 Clutton-Brock, J. (1992). The process of domestication. *Mammal Review*, 22(2), 79–85.
 380 <https://doi.org/10.1111/j.1365-2907.1992.tb00122.x>
 381 Collyer, M. L., Sekora, D. J., & Adams, D. C. (2015). A method for analysis of phenotypic change
 382 for phenotypes described by high-dimensional data. *Heredity*, 115(4), 357–365.
 383 <https://doi.org/10.1038/hdy.2014.75>
 384 Crossley, D. A., & del Mar Miguélez, M. (2001). Skull size and cheek-tooth length in wild-caught
 385 and captive-bred chinchillas. *Archives of Oral Biology*, 46(10), 919–928.
 386 [https://doi.org/10.1016/S0003-9969\(01\)00055-3](https://doi.org/10.1016/S0003-9969(01)00055-3)

387 Cucchi, T., Domont, A., Harbers, H., Leduc, C., Guidez, A., Bridault, A., ... Vigne, J.-D. (2021).
388 Bones geometric morphometrics illustrate 10th millennium cal. BP domestication of
389 autochthonous Cypriot wild boar (*Sus scrofa circeus* nov. ssp). *Scientific Reports*, 11(1),
390 11435. <https://doi.org/10.1038/s41598-021-90933-w>

391 Drake, A. G., & Klingenberg, C. P. (2008). The pace of morphological change: Historical
392 transformation of skull shape in St Bernard dogs. *Proceedings of the Royal Society B:*
393 *Biological Sciences*, 275(1630), 71–76. <https://doi.org/10.1098/rspb.2007.1169>

394 Dryden, I. L., & Mardia, K. V. (1998). *Statistical Shape Analysis*. New York, NY: John Wiley and
395 Sons.

396 Evin, A., Owen, J., Larson, G., Debiais-Thibaud, M., Cucchi, T., Vidarsdottir, U. S., & Dobney, K.
397 (2017). A test for paedomorphism in domestic pig cranial morphology. *Biology Letters*, 13(8),
398 20170321. <https://doi.org/10.1098/rsbl.2017.0321>

399 Finarelli, J. A., (2006). Estimation of endocranial volume through the use of external skull measures in
400 the carnivora (Mammalia). *Journal of Mammalogy*, 87(5), 1027–1036,
401 <https://doi.org/10.1644/05-MAMM-A-430R1.1>

402 Finarelli, J. A., (2011). Estimating endocranial volume from the outside of the skull in Artiodactyla.
403 *Journal of Mammalogy*, 92(1), 200–212, <https://doi.org/10.1644/09-MAMM-A-391.1>

404 Frantz, L. A. F., Schraiber, J. G., Madsen, O., Megens, H.-J., Cagan, A., Bosse, M., ... Groenen, M.
405 A. M. (2015). Evidence of long-term gene flow and selection during domestication from
406 analyses of Eurasian wild and domestic pig genomes. *Nature Genetics*, 47(10), 1141–1148.
407 <https://doi.org/10.1038/ng.3394>

408 Fraser, A. F., & Broom, D. M. (1990). *Farm Animal Behaviour and Welfare*. Wallingford, UK:
409 CABI Publishing.

410 Ginot, S., Hautier, L., Marivaux, L., & Vianey-Liaud, M. (2016). Ecomorphological analysis of the
 411 astragalo-calcaneal complex in rodents and inferences of locomotor behaviours in extinct
 412 rodent species. *PeerJ*, 4, e2393. <https://doi.org/10.7717/peerj.2393>

413 Grant, A. (1982). The use of tooth wear as a guide to the domestic ungulates. In B. Wilson, C.
 414 Grigson, & S. Payne (Eds.), *Ageing and Sexing Animal Bones from Archaeological Sites*
 415 (BAR British Series, pp. 991–108). UK.

416 Groves, C P. (1966). Skull-changes due to captivity in certain Equidae. *Zeitschrift Fuer*
 417 *Saugietierkunde*, pp. 44–66.

418 Groves, C. P. (1981). *Ancestors for the Pigs, Taxonomy and Phylogeny of the Genus Sus*. Canberra,
 419 Australia: Department of Prehistory Technical Bulletin, Australian National University Press.

420 Groves, C P. (1982). The skulls of Asian rhinoceroses: Wild and captive. *Zoo Biology*, 1(3), 251–
 421 261. <https://doi.org/10.1002/zoo.1430010309>

422 Hanot, P., Herrel, A., Guintard, C., & Cornette, R. (2017). Morphological integration in the
 423 appendicular skeleton of two domestic taxa: The horse and donkey. *Proceedings of the Royal*
 424 *Society B: Biological Sciences*, 284(1864), 20171241. <https://doi.org/10.1098/rspb.2017.1241>

425 Harbers, H., Neaux, D., Ortiz, K., Blanc, B., Laurens, F., Baly, I., ... Cucchi, T. (2020). The mark of
 426 captivity: Plastic responses in the ankle bone of a wild ungulate (*Sus scrofa*). *Royal Society*
 427 *Open Science*, 7(3), 192039. <https://doi.org/10.1098/rsos.192039>

428 Harbers, H., Zanolli, C., Cazenave, M., Theil, J.-C., Ortiz, K., Blanc, B., ... Cucchi, T. (2020).
 429 Investigating the impact of captivity and domestication on limb bone cortical morphology: An
 430 experimental approach using a wild boar model. *Scientific Reports*, 10(1), 19070.
 431 <https://doi.org/10.1038/s41598-020-75496-6>

432 Horard-Herbin, M.-P. (1997). *Le village celtique des Arènes à Levroux. L'élevage et les productions*
 433 *animales dans l'économie de la fin du second âge du Fer-Levroux 4* (Vol. 12). Fédération
 434 pour l'édition de la Revue archéologique du Centre de la France.

435 Iannuci, A., Sardella, R., Strani, F., Mecozzi, B. (2020). Size shifts in late Middle Pleistocene to
 436 Early Holocene *Sus scrofa* (Suidae, Mammalia) from Apulia (southern Italy):
 437 ecomorphological adaptations? *Hystrix, the Italian Journal of Mammalogy*, 31(1), 10–20.
 438 <https://doi.org/10.4404/hystrix-00258-2019>

439 Kimura, T., & Hamada, Y. (1996). Growth of wild and laboratory born chimpanzees. *Primates*,
 440 37(3), 237–251. <https://doi.org/10.1007/BF02381856>

441 Klingenberg, C. P. (2016). Size, shape, and form: Concepts of allometry in geometric
 442 morphometrics. *Development Genes and Evolution*, 226, 113–137.
 443 <https://doi.org/10.1007/s00427-016-0539-2>

444 Logan, C. J., & Palmstrom, C. R. (2015). Can endocranial volume be estimated accurately from
 445 external skull measurements in great-tailed grackles (*Quiscalus mexicanus*)? *PeerJ*,
 446 3:e1000. <https://doi.org/10.7717/peerj.1000>

447 Lord, K. A., Larson, G., Coppinger, R. P., & Karlsson, E. K. (2020). The History of Farm Foxes
 448 Undermines the Animal Domestication Syndrome. *Trends in Ecology & Evolution*, 35(2),
 449 125–136. <https://doi.org/10.1016/j.tree.2019.10.011>

450 Lord, K. A., Larson, G., & Karlsson, E. K. (2020). Brain Size Does Not Rescue Domestication
 451 Syndrome. *Trends in Ecology & Evolution*, 35(12), 1061–1062.
 452 <https://doi.org/10.1016/j.tree.2020.10.004>

453 Marshall, F. B., Dobney, K., Denham, T., & Capriles, J. M. (2014). Evaluating the roles of directed
 454 breeding and gene flow in animal domestication. *Proceedings of the National Academy of*
 455 *Sciences*, 111(17), 6153–6158. <https://doi.org/10.1073/pnas.1312984110>

- Mason, G., & Mendl, M. (1997). Do the stereotypies of pigs, chickens and mink reflect adaptive species differences in the control of foraging? *Applied Animal Behaviour Science*, 53(1), 45–58. [https://doi.org/10.1016/S0168-1591\(96\)01150-1](https://doi.org/10.1016/S0168-1591(96)01150-1)
- Mitteroecker, P., Gunz, P., Bernhard, M., Schaefer, K., & Bookstein, F. L. (2004). Comparison of cranial ontogenetic trajectories among great apes and humans. *Journal of Human Evolution*, 46(6), 679–698. <https://doi.org/10.1016/j.jhevol.2004.03.006>
- Mitteroecker, P., Gunz, P., Windhager, S., & Schaefer, K. (2013). A brief review of shape, form, and allometry in geometric morphometrics, with applications to human facial morphology. *Hystrix, the Italian Journal of Mammalogy*, 24(1), 59–66. <https://doi.org/10.4404/hystrix-24.1-6369>
- Neaux, D., Blanc, B., Ortiz, K., Locatelli, Y., Laurens, F., Baly, I., ... Cucchi, T. (2021). How changes in functional demands associated with captivity affect the skull shape of a wild boar (*Sus scrofa*). *Evolutionary Biology*, 48, 27–40. <https://doi.org/10.1007/s11692-020-09521-x>
- Neaux, D., Blanc, B., Ortiz, K., Locatelli, Y., Schafberg, R., Herrel, A., ... Cucchi, T. (2021). Constraints associated with captivity alter craniomandibular integration in wild boar. *Journal of Anatomy*, 239(2), 489–497. <https://doi.org/10.1111/joa.13425>
- Neaux, D., Harbers, H., Blanc, B., Ortiz, K., Locatelli, Y., Herrel, A., ... Cucchi, T. (2021). *The effect of captivity on cranial and postcranial ontogenetic trajectories in wild boar* [Data set]. Zenodo. <https://doi.org/10.5281/zenodo.5547335>
- Neaux, D., Sansalone, G., Lecompte, F., Haruda, A., Schafberg, R., & Cucchi, T. (2020). Examining the effect of feralization on craniomandibular morphology in pigs, *Sus scrofa* (Artiodactyla: Suidae). *Biological Journal of the Linnean Society*, 131(4), 870–879. <https://doi.org/10.1093/biolinnean/blaa156>

479 O'Regan, H. J., & Kitchener, A. C. (2005). The effects of captivity on the morphology of captive,
 480 domesticated and feral mammals. *Mammal Review*, 35(3–4), 215–230.
 481 <https://doi.org/10.1111/j.1365-2907.2005.00070.x>

482 Palencia, P., Vicente, J., Barroso, P., Barasona, J. Á., Soriguer, R. C., & Acevedo, P. (2019).
 483 Estimating day range from camera-trap data: The animals' behaviour as a key parameter.
 484 *Journal of Zoology*, 309(3), 182–190. <https://doi.org/10.1111/jzo.12710>

485 Panciroli, E., Janis, C., Stockdale, M., & Martín-Serra, A. (2017). Correlates between calcaneal
 486 morphology and locomotion in extant and extinct carnivorous mammals. *Journal of*
 487 *Morphology*, 278(10), 1333–1353. <https://doi.org/10.1002/jmor.20716>

488 Pelletier, M., Kotiaho, A., Niinimäki, S., & Salmi, A.-K. (2020). Identifying early stages of reindeer
 489 domestication in the archaeological record: A 3D morphological investigation on forelimb
 490 bones of modern populations from Fennoscandia. *Archaeological and Anthropological*
 491 *Sciences*, 12(8), 169. <https://doi.org/10.1007/s12520-020-01123-0>

492 Pelletier, M., Kotiaho, A., Niinimäki, S., & Salmi, A.-K. (2021). Impact of selection and
 493 domestication on hindlimb bones of modern reindeer populations: Archaeological
 494 implications for early reindeer management by Sámi in Fennoscandia. *Historical Biology*,
 495 0(0), 1–19. <https://doi.org/10.1080/08912963.2021.1947268>

496 Price, E. O. (1999). Behavioral development in animals undergoing domestication. *Applied Animal*
 497 *Behaviour Science*, 65(3), 245–271. [https://doi.org/10.1016/S0168-1591\(99\)00087-8](https://doi.org/10.1016/S0168-1591(99)00087-8)

498 R Core Team. (2019). R: A language and environment for statistical computing (Version 3.1.2).
 499 Vienna, Austria.

500 Rhodes, R. T., Appleby, M. C., Chinn, K., Douglas, L., Firkins, L. D., Houpt, K. A., ... Wills, R. W.
 501 (2005). A comprehensive review of housing for pregnant sows. *Journal of the American*

- Veterinary Medical Association*, 227(10), 1580–1590.
<https://doi.org/10.2460/javma.2005.227.1580>
- Rohlf, F. J., & Slice, D. (1990). Extensions of the Procrustes method for the optimal superimposition of landmarks. *Systematic Biology*, 39(1), 40–59. <https://doi.org/10.2307/2992207>
- Russo, L., Massei, G., & Genov, P. V. (1997). Daily home range and activity of wild boar in a Mediterranean area free from hunting. *Ethology Ecology & Evolution*, 9(3), 287–294.
<https://doi.org/10.1080/08927014.1997.9522888>
- Sánchez-Villagra, M. R., Geiger, M., & Schneider, R. A. (2016). The taming of the neural crest: A developmental perspective on the origins of morphological covariation in domesticated mammals. *Royal Society Open Science*, 3(6), 160107. <https://doi.org/10.1098/rsos.160107>
- Sánchez-Villagra, M. R., Segura, V., Geiger, M., Heck, L., Veitschegger, K., & Flores, D. (2017). On the lack of a universal pattern associated with mammalian domestication: Differences in skull growth trajectories across phylogeny. *Royal Society Open Science*, 4(10), 170876.
<https://doi.org/10.1098/rsos.170876>
- Schaller, G. B. (1973). *The Serengeti Lion: A Study of Predator-Prey Relations* (University of Chicago). Chicago.
- Schlager, S., & Jefferis, G. (2020). *Morpho: Calculations and visualisations related to geometric morphometrics*.
- Schley, L., & Roper, T. J. (2003). Diet of wild boar *Sus scrofa* in Western Europe, with particular reference to consumption of agricultural crops. *Mammal Review*, 33(1), 43–56.
<https://doi.org/10.1046/j.1365-2907.2003.00010.x>
- Smuts, G. L., Anderson, J. L., & Austin, J. C. (1978). Age determination of the African lion (*Panthera leo*). *Journal of Zoology*, 185(1), 115–146. <https://doi.org/10.1111/j.1469-7998.1978.tb03317.x>

526 Su, S. C., Skedros, J. G., Bachus, K. N., & Bloebaum, R. D. (1999). Loading conditions and cortical
 527 bone construction of an artiodactyl calcaneus. *Journal of Experimental Biology*, 202(22),
 528 3239–3254. <https://doi.org/10.1242/jeb.202.22.3239>
 529 Turner, T. R., Cramer, J. D., Nisbett, A., & Gray, J. P. (2016). A comparison of adult body size
 530 between captive and wild vervet monkeys (*Chlorocebus aethiops sabaues*) on the island of St.
 531 Kitts. *Primates; Journal of Primatology*, 57(2), 211–220. [https://doi.org/10.1007/s10329-015-](https://doi.org/10.1007/s10329-015-0509-8)
 532 0509-8
 533 Vigne, J.-D. (2011). The origins of animal domestication and husbandry: A major change in the
 534 history of humanity and the biosphere. *Comptes Rendus Biologies*, 334(3), 171–181.
 535 <https://doi.org/10.1016/j.crvi.2010.12.009>
 536 Vigne, J.-D., Carrère, I., Briois, F., & Guilaine, J. (2011). The Early Process of Mammal
 537 Domestication in the Near East: New Evidence from the Pre-Neolithic and Pre-Pottery
 538 Neolithic in Cyprus. *Current Anthropology*, 52(S4), S255–S271.
 539 <https://doi.org/10.1086/659306>
 540 Voje, K. L., Hansen, T. F., Egset, C. K., Bolstad, G. H., & Pélabon, C. (2014). Allometric
 541 Constraints and the Evolution of Allometry. *Evolution*, 68(3), 866–885.
 542 <https://doi.org/10.1111/evo.12312>
 543 Wilson, L. A. B. (2018). The evolution of ontogenetic allometric trajectories in mammalian
 544 domestication. *Evolution*, 72(4), 867–877. <https://doi.org/10.1111/evo.13464>
 545 Zeder, M A. (2012). Pathways to animal domestication. In P. Gepts, T. R. Famula, & R. L. Bettinger
 546 (Eds.), *Biodiversity in Agriculture: Domestication, Evolution, and Sustainability* (pp. 227–
 547 259). Cambridge, UK: Cambridge University Press.

548 Zeder, M A. (2018). Why evolutionary biology needs anthropology: Evaluating core assumptions of
549 the extended evolutionary synthesis. *Evolutionary Anthropology*, 27(6), 267–284.
550 <https://doi.org/10.1002/evan.21747>

551 Zihlman, A. L., Bolter, D. R., & Boesch, C. (2007). Skeletal and dental growth and development in
552 chimpanzees of the Taï National Park, Côte D’Ivoire. *Journal of Zoology*, 273(1), 63–73.
553 <https://doi.org/10.1111/j.1469-7998.2007.00301.x>

554

555 Table 1. MANOVA p -values of the form coordinates between captive (stall and enclosure) and wild-
556 caught groups of wild boar (*Sus scrofa*) computed for the cranium, mandible, and calcaneus. Significant
557 values ($p < 0.05$) are in bold. m.: months.

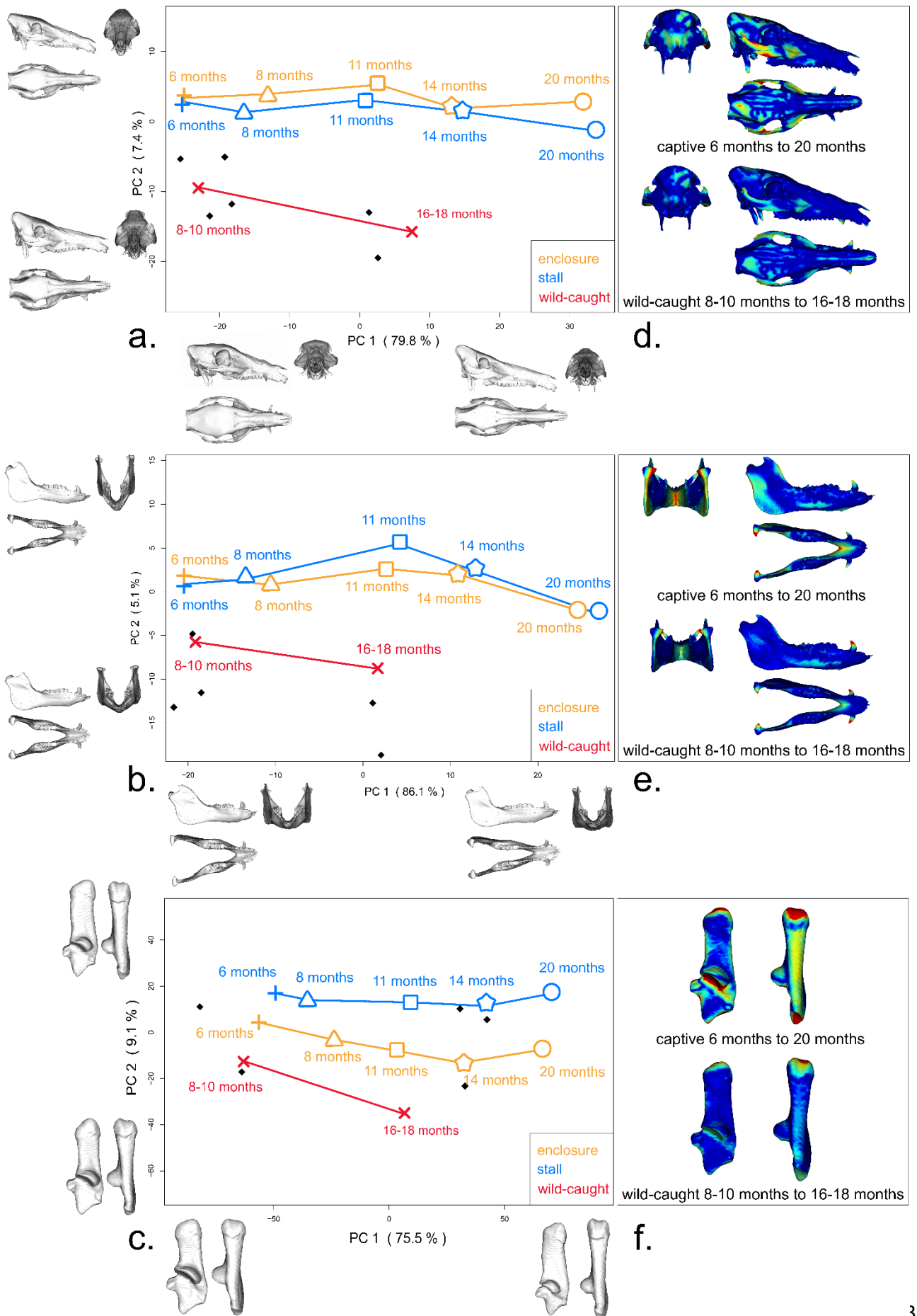
	Cranium		Mandible		Calcaneus	
	8-10 m.	16-18m.	8-10 m.	16-18 m.	8-10 m.	16-18 m.
	wild-caught	wild-caught	wild-caught	wild-caught	wild-caught	wild-caught
6 m. – enclosure	0.21	< 0.01	0.55	< 0.01	0.49	< 0.01
6 m. – stall	0.27	< 0.01	0.74	< 0.01	0.35	0.01
8 m. – enclosure	0.11	0.02	0.27	0.09	0.13	0.09
8 m. – stall	0.22	0.01	0.38	0.05	0.22	0.03
11 m. – enclosure	< 0.01	0.12	0.01	0.42	0.01	0.35
11 m. – stall	0.01	0.13	0.01	0.30	< 0.01	0.20
14 m. – enclosure	< 0.01	0.17	< 0.01	0.21	< 0.01	0.25
14 m. – stall	< 0.01	0.15	< 0.01	0.12	< 0.01	0.09
20 m. – enclosure	< 0.01	0.01	< 0.01	0.01	< 0.01	0.02
20 m. – stall	< 0.01	0.01	< 0.01	0.01	< 0.01	0.01

558

559 Table 2. ANOVA p -values of the centroid size (CS) between captive (stall and enclosure) and wild-
560 caught groups of wild boar (*Sus scrofa*) computed for the cranium, mandible, and calcaneus and
561 ANOVA p -values of body mass. Significant values ($p < 0.05$) are in bold. . m.: months.

	CS Cranium		CS Mandible		CS Calcaneus		Body mass	
	8 to 10	16 to 18	8 to 10	16 to 18	8 to 10	16 to 18	8 to 10	16 to 18
	m.	m.	m.	m.	m.	m.	m.	m.
	wild-	wild-	wild-	wild-	wild-	wild-	wild-	wild-
	caught	caught	caught	caught	caught	caught	caught	caught
6 m. – enclosure	0.75	< 0.01	0.54	< 0.01	0.93	< 0.01	0.50	< 0.01
6 m. – stall	0.67	< 0.01	0.58	0.01	0.78	0.01	0.45	< 0.01
8 m. – enclosure	0.37	0.03	0.39	0.09	0.18	0.12	0.81	0.01
8 m. – stall	0.60	0.00	0.61	0.05	0.32	0.06	0.64	< 0.02
11 m. – enclosure	0.01	0.42	< 0.01	0.79	0.01	0.62	0.20	0.12
11 m. – stall	0.02	0.37	0.02	0.81	< 0.01	0.95	0.16	0.22
14 m. – enclosure	< 0.01	0.71	< 0.01	0.49	< 0.01	0.63	< 0.02	0.60
14 m. – stall	< 0.01	0.60	< 0.01	0.33	< 0.01	0.28	0.01	0.99
20 m. – enclosure	< 0.01	0.04	< 0.01	0.02	< 0.01	0.12	< 0.01	0.06
20 m. – stall	< 0.01	0.02	< 0.01	0.01	< 0.01	0.08	< 0.01	0.03

562



564 Figure 1. Principal component analyses for the cranium (a) mandible (b), and calcaneus (c) of wild
565 boar (*Sus scrofa*) in the PC1-PC2 form space. Symbols represent the mean form for each group. Black
566 dots represent the “control” specimens from the wild-caught group, i.e. the specimens belonging to the
567 same initial population as the individuals from the stall and enclosure groups. Form changes are
568 depicted in lateral, dorsal and frontal views for the cranium and the mandible, and in medial and plantar
569 views for the calcaneus. Heatmap of the intensity of form variation between captive (i.e. enclosure and
570 stall) groups (6 months and 20 months) and between wild-caught groups (8-10 months and 16-18
571 months) for the cranium (d), mandible (e), and calcaneus (f). Blue indicates a low intensity of variation
572 and red indicates a high intensity of variation. Form changes are depicted in lateral, dorsal and frontal
573 views for the cranium and the mandible, and in medial and plantar views for the calcaneus

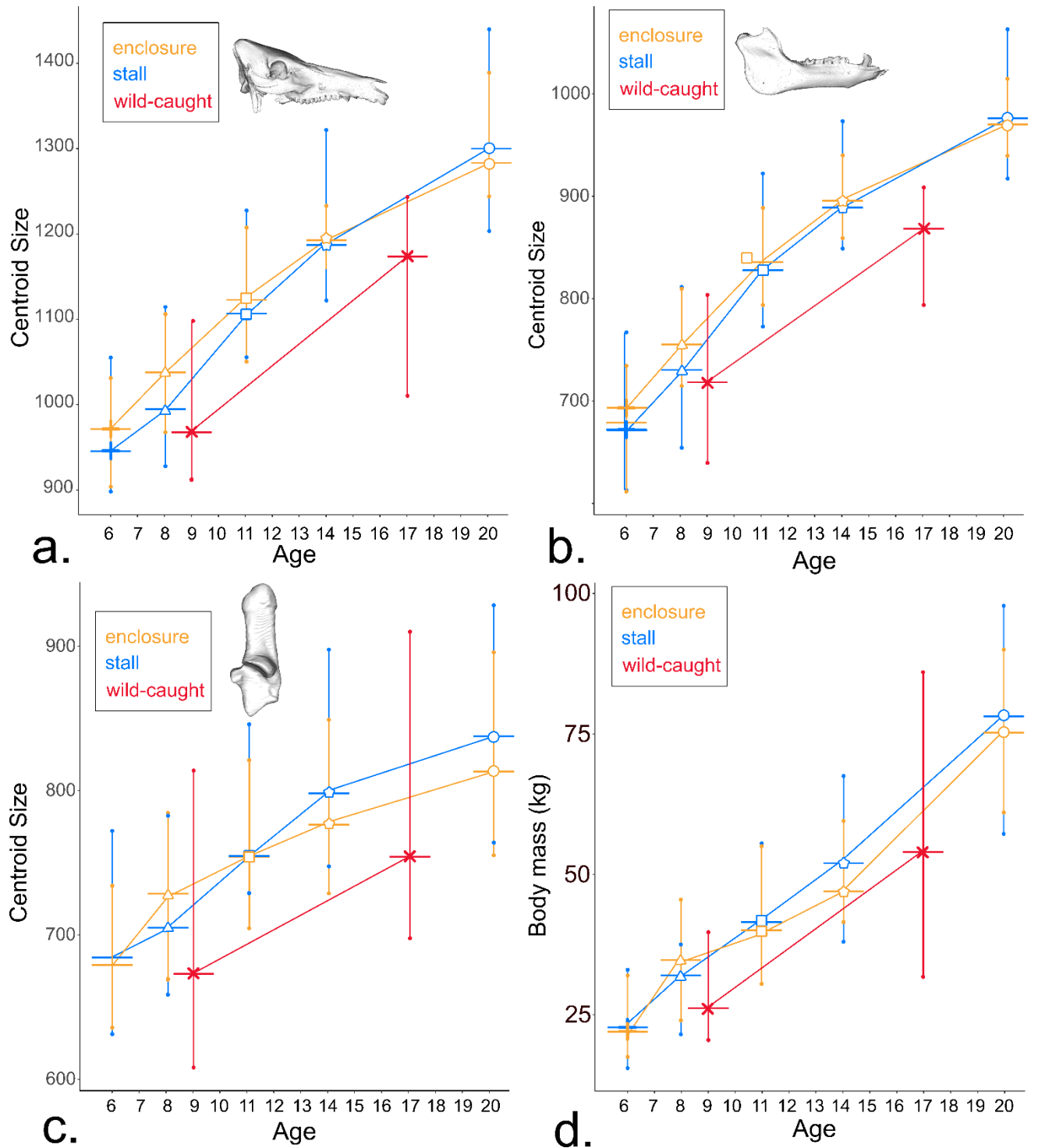


Figure 2. Boxplots of centroid size (CS) for the cranium (a), mandible (b), and calcaneus (c) and of body mass (d) of wild boar (*Sus scrofa*). The vertical lines represent all the values within 1.5 times of the interquartile range accounting for 50% of the data, from the 25th percentile to the 75th percentile. The horizontal lines are the median values.

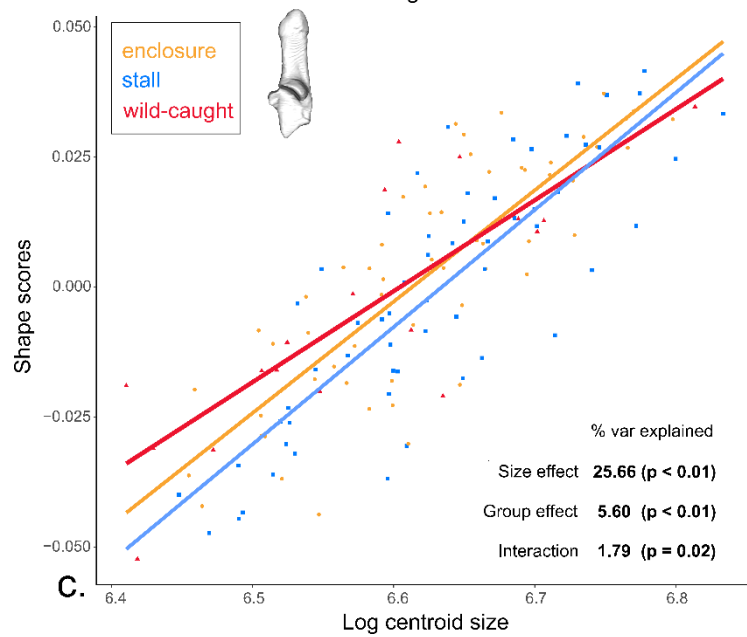
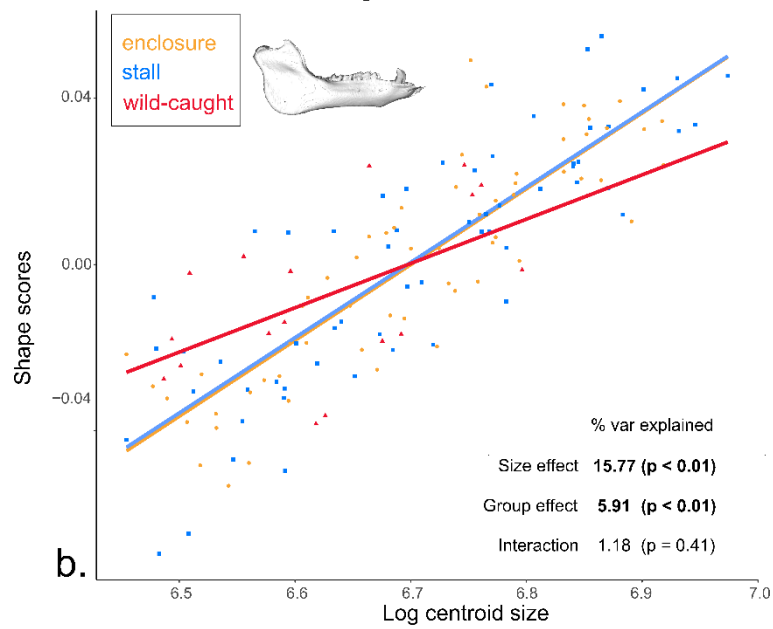
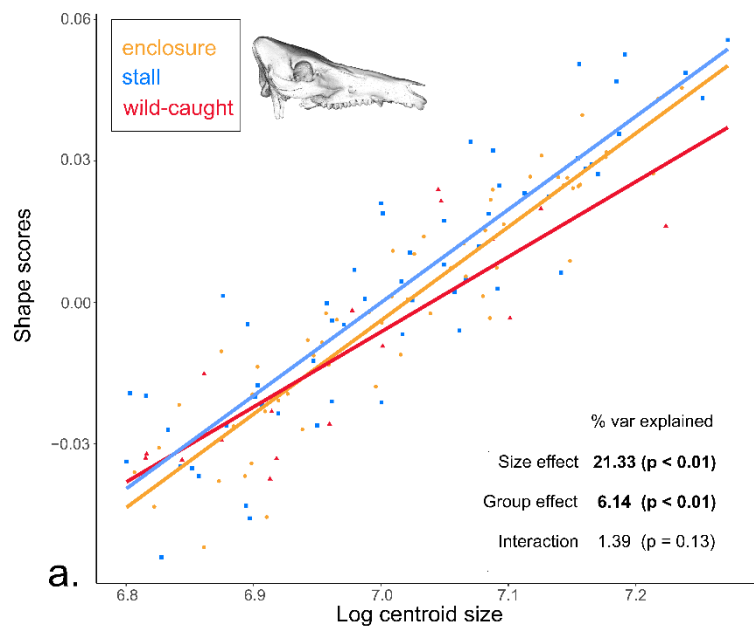


Figure 3. Regression of log centroid size (CS) on shape scores for the cranium (a), mandible (b) and calcaneus (c) of wild boar (*Sus scrofa*) and effects of size, group, and interaction between size and group on the regression of log CS on shape scores. Significant values ($p < 0.05$) are in bold.

589 The effect of captivity on craniomandibular and calcaneal ontogenetic trajectories in wild boar

590 Short running title: Captivity and ontogenetic trajectories

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608

609 SUPPORTING INFORMATION

610 **Appendix S1: Groups and specimens used**

611 a. List of groups included in the study and number of specimens

Cranium	Cranium	Mandible	Calcaneus
Enclosure	12	12	12
Stall	12	12	12
8-10 months wild-caught	10	12	10
16-18 months wild-caught	8	6	8
TOTAL	42	42	42

612

b. List of specimens. M: male, F: female, nd: not determined individuals. Specimens are localised at the Muséum national d'Histoire naturelle (Paris, France)

Catalogue number	Sex ¹	Age class	Status	Location	Cranium	Mandible	Calcaneus
2017-557	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
H285	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-560	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-562	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-555	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-556	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-569	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
H319	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-554	F		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-571	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-574	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes

2017-575	M		stall	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-558	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-559	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-561	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-563	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-564	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-565	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-566	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-567	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-568	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-570	F		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-572	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
2017-573	M		enclosure	Réserve de la Haute-Touche	Yes	Yes	Yes
PRA_186	nd	8-10 months	wild-caught	Urciers	Yes	Yes	No
PRA_172	F	8-10 months	wild-caught	Urciers	Yes	Yes	Yes

PRA_174	F	8-10 months	wild-caught	Urciers	Yes	Yes	Yes
2017-582	M	8-10 months	wild-caught	Urciers	Yes	Yes	Yes
2017-584	M	8-10 months	wild-caught	Urciers	Yes	Yes	Yes
CHA_S_77	F	8-10 months	wild-caught	Chambord	Yes	Yes	Yes
CHA_S_509	F	8-10 months	wild-caught	Chambord	Yes	Yes	Yes
CHA_S_581	M	8-10 months	wild-caught	Chambord	Yes	Yes	Yes
CHA_S_664	F	8-10 months	wild-caught	Chambord	Yes	No	Yes
CHA_S_577	nd	8-10 months	wild-caught	Chambord	No	Yes	No
COMP_2013-1262	nd	8-10 months	wild-caught	Compiègne	Yes	Yes	No
COMP_2013-1247	F	8-10 months	wild-caught	Compiègne	No	Yes	Yes
COMP_2013-1269	F	8-10 months	wild-caught	Compiègne	No	Yes	Yes
2017-583	M	16-18 months	wild-caught	Urciers	Yes	Yes	Yes
2017-581	F	16-18 months	wild-caught	Chambord	Yes	Yes	Yes
2017-577	M	16-18 months	wild-caught	Chambord	Yes	Yes	Yes
2017-579	F	16-18 months	wild-caught	Chambord	Yes	Yes	Yes

2017-580	F	16-18 months	wild-caught	Chambord	Yes	Yes	Yes
2017-578	F	16-18 months	wild-caught	Chambord	Yes	No	Yes
COMP_2013-1264	F	16-18 months	wild-caught	Compiègne	Yes	Yes	Yes
COMP_2013-1270	M	16-18 months	wild-caught	Compiègne	Yes	No	Yes

¹Sexes and ages for wild-caught specimens were estimated based on osteological observations, using respectively the morphology of canine cross section (Mayer & Brisbin, 1988) and the mandibular tooth eruption and wear stages in occlusal view (Grant, 1982).

Grant, A., 1982. The use of tooth wear as a guide to the domestic ungulates. In: Wilson, B., Grigson, C., Payne, S. (Eds.), *Ageing and Sexing Animal Bones from Archaeological Sites*. UK, pp. 991–108.

Mayer, J.M., & Brisbin, I.L., 1988. Sex identification of *Sus scrofa* based on canine morphology. *Journal of Mammalogy* 69:408–4

Appendix S2: Digitisation and definitions of landmarks

a. Digitisation protocol

All specimens were scanned using a Computed Tomography (CT) scanner with a spatial resolution of between 100 and 500 μm . The wild boar from Urciers were scanned as living specimens at the *Chirurgie et Imagerie pour la Recherche et l'Enseignement* (CIRE) platform of the *Institut National de Recherche pour l'Agriculture, l'Alimentation et l'Environnement* (INRAE). Other individuals were scanned as dry specimens using a CT scanner close to the collections they were housed in. We segmented the bones using the segmentation tools of the Avizo v8.0 software, and then converted the volumes into three-dimensional PLY surfaces format. We digitised the anatomical landmarks and semilandmarks using IDAV Landmark v3.0 software (Wiley et al., 2005). To remove variation related to their initial arbitrary position along the curves, the semilandmarks were slid along the tangent of the curves minimising bending energy (Gunz and Mitteroecker, 2013).

Gunz, P., Mitteroecker, P., 2013. Semilandmarks: a method for quantifying curves and surfaces. *Hystrix, the Italian Journal of Mammalogy*. 24, 103–109.

Wiley, D., Amenta, N., Alcantara, D., Ghosh, D., Kil, Y.J., Delson, E., Harcourt-Smith, W., Rohlf, F.J., St. John, K., Hamann, B., Motani, R., Frost, S., Rosenberger, A.L., Tallman, L., Disotell, T., O'Neill, R., 2005. Evolutionary Morphing. In: *Proceedings of IEEE Visualization 2005*. Presented at the VIS'05, IEEE, Minneapolis, MN, pp. 431–438.

b. Number of landmarks

	Homologous landmarks	Sliding semi- landmarks	Surface sliding semi- landmarks
Cranium	14	181	0
Mandible	70	28	0
Calcaneus	23	48	763

c. Definitions of cranial (1-70), mandibular (71-94), and calcaneus (95-108) homologous landmarks.

Landmark	Definition
1	Most anterior midline point of the nasals
2	Most anterior, dorsal midline point of the premaxillae
3, 4	Most anterior point of the nasal-premaxilla suture
5, 6	Most anterior, lateral point of the upper canine alveolus
7, 8	Suture at the meeting point of premaxilla, maxilla, and nasal
9, 10	Most anterior point of the infraorbital foramen
11, 12	Most posterior point of the infraorbital foramen
13, 14	Most anterior lateral point of the facial tuberosity
15, 16	Most ventral point of the zygomatic-maxilla suture
17, 18	Most anterior, lateral point of the orbit
19, 20	Most dorsal point of the lower lacrimal foramen
21, 22	Most posterior point of the supraorbital foramen
23, 24	Most dorsal point of the orbit
25, 26	Most ventral point of supraorbital process of the frontal bone
27, 28	Meeting point of the parietal-frontal suture and temporal line
29, 30	Most anterior, dorsal point of the zygomatic process of the squamosal bone
31, 32	Most posterior point of the zygomatic bone
33, 34	Most dorsal point of the zygomatic process of the squamosal bone
35, 36	Most anterior, lateral point of the nuchal crest
37, 38	Most anterior point of the palatine fissure
39, 40	Most posterior point of the palatine fissure

41, 42	Most anterior point of the cheek-tooth row (excluding P1)
43, 44	Most posterior point of the cheek-tooth row
45	Most posterior point of the posterior nasal spine on the palatine bone
46, 47	Most ventral, lateral point of the pterygoid process of the sphenoid
48, 49	Most posterior point of the pterygoid hamulus
50, 51	Meeting point of the pterygoid process with the ridge of the lateral pterygoid plate
52, 53	Meeting point of the pterygoid hamulus with the ridge of the medial pterygoid plate
54	Most posterior point of the vomer in contact with the sphenoid
55, 56	Most ventral, lateral, posterior point of the sphenoid-squamosal suture
57, 58	Most ventral, medial, posterior point of the sphenoid-squamosal suture
59, 60	Most posterior, medial point of the petro-occipital fissure
61, 62	Most lateral point of the occipital condyle
63	Most anterior, ventral midline point of the premaxilla
64	Most posterior midline point of the nuchal crest
65, 66	Most posterior, lateral point of the nuchal crest
67, 68	Most lateral point of the foramen magnum
69	Most posterior, dorsal point of the foramen magnum
70	Most anterior point, ventral of the foramen magnum
71, 72	Most anterior, lateral point of the lower canine alveolus
73, 74	Most anterior point of the cheek-tooth row (excluding P1)
75, 76	Most lateral point at the maximum of curvature between the mandibular ramus and corpus
77, 78	Most lateral point at the maximum of curvature between the coronoid process and the mandibular ramus

79, 80	Most dorsal point of the coronoid process
81, 82	Most lateral point of the mandibular condyle
83, 84	Most posterior point of the mandibular condyle
85, 86	Point at the maximum of curvature of the mandibular angle
87	Most ventral, posterior point of the mandibular symphysis
88	Most ventral, anterior point of the mandibular symphysis
89, 90	Most medial point of the mandibular condyle
91	Most dorsal, posterior point of the mandibular symphysis
92	Most dorsal, anterior point of the mandibular symphysis
93, 94	Most anterior point of the mandibular foramen
95	Distal end of the cuboid facet
96	Proximo-plantar end of the cuboid facet
97	End of the beak of the coracoid process
98	Maximum of curvature of the plantar bulge on the plantar margin
99	Dorso-proximal end of the calcaneal sulcus
100	Planto-lateral end of sustentaculum tali
101	Dorsal end of the sustentaculum tali
102	Medial end of sustentaculum tali
103	Plantar end of the epiphysis
104	Dorso-proximal end of the bulge of the proximal part (not on the epiphysis)
105	Proximal end of the lateral lobe of the epiphysis (secondary lobe)
106	Proximal end of the medial lobe of the epiphysis (main lobe)
107	Dorsal end of the epiphysis
108	Dorso-proximal end of the lateral part of the coracoid process

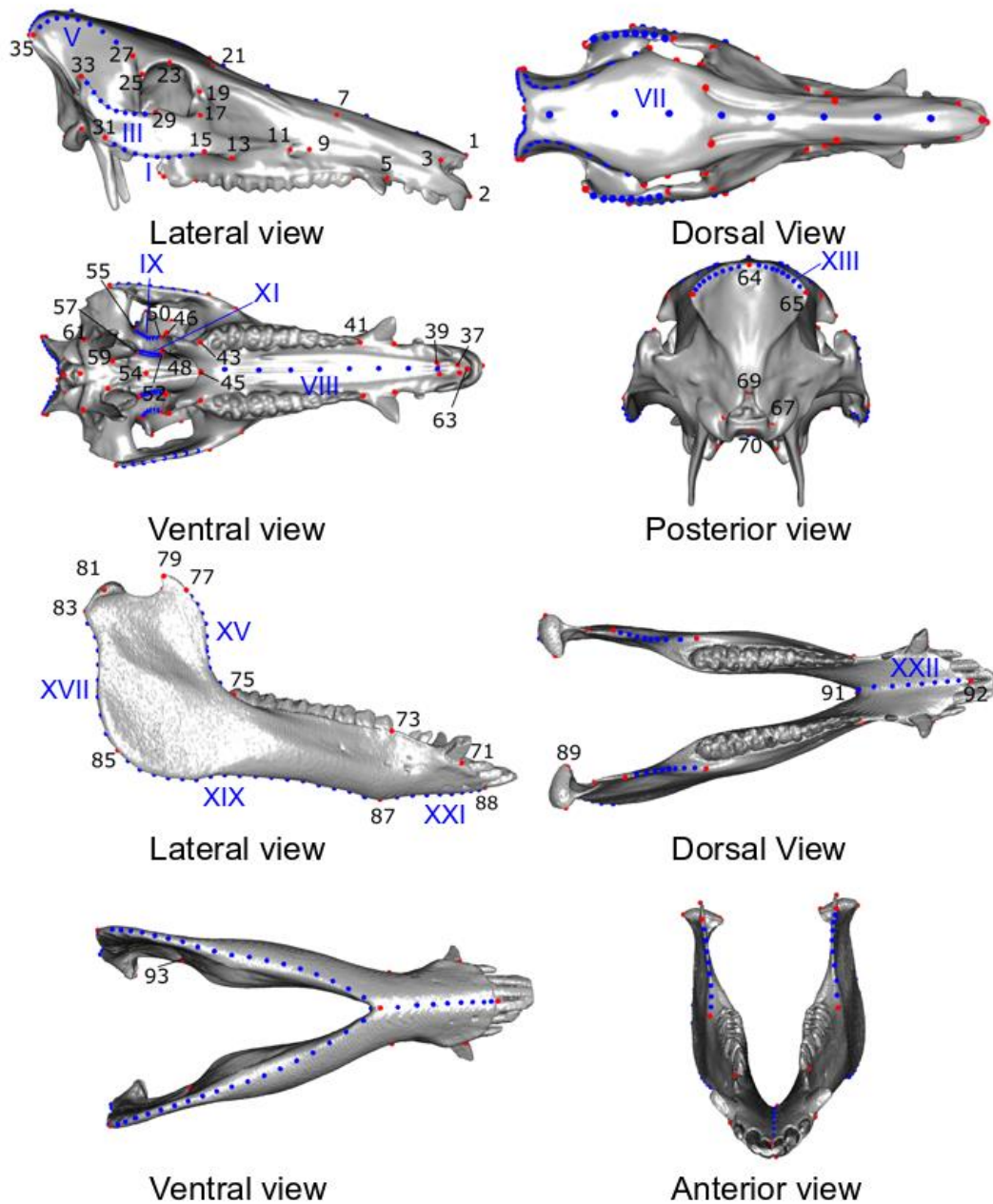
d. Definitions of cranial (I to XIV), mandibular (XV to XXII), and calcaneus (XVI to XXIX)

curves

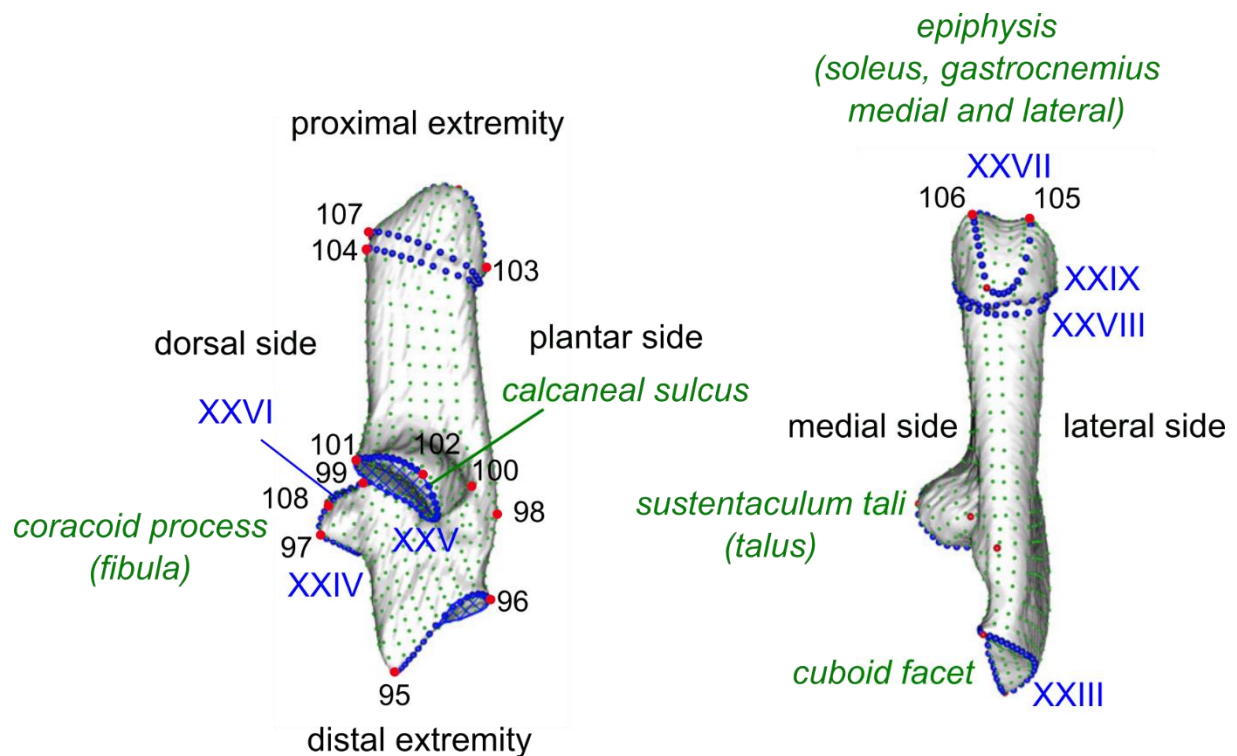
Curve	Definition
I	from LM 13 to LM31
II	from LM 14 to LM32
III	from LM 29 to LM33
IV	from LM 30 to LM34
V	from LM 27 to LM35
VI	from LM 28 to LM36
VII	from LM 1 to LM64
VIII	from LM 45 to LM63
IX	from LM 50 to LM55
X	from LM 51 to LM56
XI	from LM 52 to LM57
XII	from LM 53 to LM58
XIII	from LM 64 to LM65
XIV	from LM 64 to LM66
XV	from LM 75 to LM77
XVI	from LM 76 to LM78
XVII	from LM 83 to LM85
XVIII	from LM 84 to LM86
XIX	from LM 85 to LM87
XX	from LM 86 to LM87
XXI	from LM 87 to LM88

XXII	from LM 91 to LM92
XXIII	Edge of the articular surface of the cuboid facet
XXIV	Medial edge of the coracoid process
XXV	Edge of the articular surface of the sustentaculum tali
XXVI	Lateral edge of the coracoid process
XXVII	Edge of the attachment surface of the tendon on the epiphysis
XXVIII	Distal delineation of the junction zone between the epiphysis and the rest of the calcaneus
XXIX	Proximal delineation of the junction zone between the epiphysis and the rest of the calcaneus

e. Wild boar (*Sus scrofa*) cranium and mandible showing the homologous landmarks (red dots and Arabic numerals) and semilandmarks (blue dots and Roman numerals) used in the study.



f. Wild boar (*Sus scrofa*) calcaneus showing the homologous landmarks (red dots and Arabic numerals) and semilandmarks (blue dots and Roman numerals) used in the study.



Appendix S3: Body mass measurements (kg)

	Captive wild boar				
	6 months	8 months	11 months	14 months	20 months
2017-557	18.5	32.5	51.0	60.5	83.0
H285	24.0	27.0	30.5	38.0	59.0
2017-560	21.5	28.5	55.5	56.5	77.0
2017-562	29.0	37.5	40.5	50.0	76.0
2017-555	15.5	21.5	31.0	49.5	77.0
2017-556	27.0	34.0	35.0	40.0	59.0
2017-569	26.0	36.5	38.5	45.0	57.2
H319	19.5	29.5	51.5	67.5	97.8
2017-554	21.5	31.5	42.5	54.0	77.0
2017-571	18.0	24.0	31.0	44.5	68.0
2017-574	33.0	37.5	50.0	67.0	95.5
2017-575	30.5	35.0	42.5	57.5	78.5
2017-558	19.0	30.5	40.0	55.5	80.5
2017-559	18.0	24.0	36.0	43.0	62.0
2017-561	26.0	39.5	47.0	59.5	83.0
2017-563	31.0	45.0	56.5	73.5	89.0
2017-564	22.0	31.2	30.5	44.5	61.0
2017-565	17.5	29.0	33.0	41.5	61.5
2017-566	21.0	32.0	38.0	46.0	66.5
2017-567	32.0	38.0	50.0	53.5	81.0
2017-568	27.0	38.5	48.5	50.5	77.0

2017-570	19.5	30.0	40.0	45.0	67.0
2017-572	32.0	45.5	55.0	73.0	90.0
2017-573	24.0	37.5	36.5	47.0	73.5

wild-caught wild boar

	8-10 months	16-18 months
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PRA_186	30.0
PRA_172	23.0
PRA_174	28.0
2017-582	35.0
2017-584	52.0
CHA_S_77	20.9
CHA_S_509	24.3
CHA_S_581	20.5
CHA_S_664	36.0
CHA_S_577	25.0
COMP_2013-1262	NA
COMP_2013-1247	23.9
COMP_2013-1269	39.7

2017-583	53.0
2017-581	49.8
2017-577	35.3
2017-579	31.7
2017-580	68.5
2017-578	20.0

COMP_2013-1264	60.8
COMP_2013-1270	86.0

1 **Appendix S4: Pairwise analyses between all groups**

2 a. MANOVA p-values of the form coordinates between all groups computed for the cranium,
3 mandible, and calcaneus.

	8 to 10	16 to 18					11
Cranium	months wild	months wild	6 months	6 months	8 months	8 months	months
	caught	caught	enclosure	stall	enclosure	stall	enclosure
16-18 months wild caught	0.00						
6 months enclosure	0.21	0.00					
6 months stall	0.27	0.00	0.98				
8 months enclosure	0.11	0.02	0.24	0.20			
8 months stall	0.22	0.01	0.34	0.46	0.88		
11 months enclosure	0.00	0.12	0.00	0.00	0.11	0.03	
11 months stall	0.01	0.13	0.00	0.00	0.13	0.06	0.70
14 months enclosure	0.00	0.17	0.00	0.00	0.01	0.00	0.26
14 months stall	0.00	0.15	0.00	0.00	0.00	0.00	0.15
20 months enclosure	0.00	0.01	0.00	0.00	0.00	0.00	0.00
20 months stall	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	8 to 10	16 to 18					11
Mandible	months wild	months wild	6 months	6 months	8 months	8 months	months
	caught	caught	enclosure	stall	enclosure	stall	enclosure
16-18 months wild caught	0.02						
6 months enclosure	0.55	0.01					
6 months stall	0.74	0.01	0.89				
8 months enclosure	0.27	0.09	0.18	0.22			

8 months stall	0.38	0.04	0.25	0.39	0.73		
11 months enclosure	0.00	0.42	0.00	0.00	0.07	0.03	
11 months stall	0.00	0.30	0.00	0.00	0.04	0.02	0.95
14 months enclosure	0.00	0.21	0.00	0.00	0.00	0.00	0.33
14 months stall	0.00	0.12	0.00	0.00	0.00	0.00	0.16
20 months enclosure	0.00	0.01	0.00	0.00	0.00	0.00	0.00
20 months stall	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	8 to 10	16 to 18					11
Calcaneus	months wild	months wild	6 months	6 months	8 months	8 months	months
	caught	caught	enclosure	stall	enclosure	stall	enclosure
16-18 months wild caught	0.01						
6 months enclosure	0.49	0.00					
6 months stall	0.35	0.01	0.89				
8 months enclosure	0.13	0.09	0.18	0.21			
8 months stall	0.22	0.03	0.31	0.68	0.74		
11 months enclosure	0.01	0.35	0.01	0.03	0.37	0.09	
11 months stall	0.00	0.20	0.00	0.02	0.13	0.07	0.56
14 months enclosure	0.00	0.25	0.00	0.00	0.02	0.01	0.29
14 months stall	0.00	0.09	0.00	0.00	0.00	0.00	0.08
20 months enclosure	0.00	0.02	0.00	0.00	0.00	0.00	0.01
20 months stall	0.00	0.01	0.00	0.00	0.00	0.00	0.01

5 b. ANOVA p -values of the centroid size (CS) between all groups computed for the cranium,
6 mandible, and calcaneus and of body mass (kg).

	8 to 10	16 to 18					11	
Cranium	months wild	months wild	6 months	6 months	8 months	8 months	months	mo
	caught	caught	enclosure	stall	enclosure	stall	enclosure	st
16-18 months wild								
caught	0.00							
6 months enclosure	0.75	0.00						
6 months stall	0.67	0.00	0.86					
8 months enclosure	0.37	0.03	0.22	0.14				
8 months stall	0.60	0.00	0.37	0.30	0.70			
11 months enclosure	0.01	0.42	0.01	0.00	0.09	0.04		
11 months stall	0.02	0.37	0.00	0.00	0.11	0.04	0.96	
14 months enclosure	0.00	0.71	0.00	0.00	0.01	0.00	0.20	
14 months stall	0.00	0.60	0.00	0.00	0.00	0.00	0.14	
20 months enclosure	0.00	0.04	0.00	0.00	0.00	0.00	0.00	
20 months stall	0.00	0.02	0.00	0.00	0.00	0.00	0.00	
	8 to 10	16 to 18					11	
Mandible	months wild	months wild	6 months	6 months	8 months	8 months	months	mo
	caught	caught	enclosure	stall	enclosure	stall	enclosure	st
16-18 months wild								
caught	0.02							
6 months enclosure	0.54	0.00						
6 months stall	0.58	0.01	0.94					
8 months enclosure	0.39	0.09	0.15	0.18				

8 months stall	0.61	0.05	0.25	0.29	0.74		
11 months enclosure	0.01	0.79	0.00	0.00	0.09	0.03	
11 months stall	0.02	0.81	0.00	0.00	0.08	0.03	0.99
14 months enclosure	0.00	0.49	0.00	0.00	0.00	0.00	0.25
14 months stall	0.00	0.33	0.00	0.00	0.00	0.00	0.15
20 months enclosure	0.00	0.02	0.00	0.00	0.00	0.00	0.00
20 months stall	0.00	0.01	0.00	0.00	0.00	0.00	0.00

	8 to 10	16 to 18					11	
Calcaneus	months wild	months wild	6 months	6 months	8 months	8 months	months	mo
	caught	caught	enclosure	stall	enclosure	stall	enclosure	st

16-18 months wild								
caught	0.01							
6 months enclosure	0.93	0.00						
6 months stall	0.78	0.01	0.74					
8 months enclosure	0.18	0.12	0.15	0.27				
8 months stall	0.32	0.06	0.29	0.50	0.70			
11 months enclosure	0.01	0.62	0.01	0.02	0.25	0.12		
11 months stall	0.00	0.95	0.00	0.00	0.10	0.03	0.65	
14 months enclosure	0.00	0.63	0.00	0.00	0.02	0.01	0.28	
14 months stall	0.00	0.28	0.00	0.00	0.00	0.00	0.09	
20 months enclosure	0.00	0.12	0.00	0.00	0.00	0.00	0.02	
20 months stall	0.00	0.08	0.00	0.00	0.00	0.00	0.01	

	8 to 10	16 to 18					11	
Mass	months wild	months wild	6 months	6 months	8 months	8 months	months	mo
	caught	caught	enclosure	stall	enclosure	stall	enclosure	st

16-18 months wild							
caught	0.01						
6 months enclosure	0.50	0.00					
6 months stall	0.45	0.00	0.91				
8 months enclosure	0.81	0.01	0.33	0.28			
8 months stall	0.64	0.02	0.19	0.18	0.80		
11 months enclosure	0.20	0.12	0.04	0.03	0.27	0.39	
11 months stall	0.16	0.22	0.02	0.01	0.19	0.26	0.82
14 months enclosure	0.02	0.60	0.00	0.00	0.02	0.05	0.26
14 months stall	0.01	1.00	0.00	0.00	0.00	0.01	0.09
20 months enclosure	0.00	0.06	0.00	0.00	0.00	0.00	0.00
20 months stall	0.00	0.01	0.00	0.00	0.00	0.00	0.00

7

8

9