



Paleoecology of the Rhinocerotidae (Mammalia, Perissodactyla) from Béon 1, Montréal-du-Gers (late early Miocene, SW France): Insights from dental microwear texture analysis, mesowear, and enamel hypoplasia

Manon Hullot, Yves Laurent, Gildas Merceron, Pierre-Olivier Antoine

► To cite this version:

Manon Hullot, Yves Laurent, Gildas Merceron, Pierre-Olivier Antoine. Paleoecology of the Rhinocerotidae (Mammalia, Perissodactyla) from Béon 1, Montréal-du-Gers (late early Miocene, SW France): Insights from dental microwear texture analysis, mesowear, and enamel hypoplasia. *Palaeontologia Electronica*, 2021, 24 (2), pp.a27. 10.26879/1163 . hal-03455891

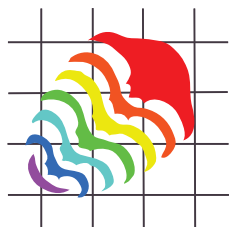
HAL Id: hal-03455891

<https://hal.science/hal-03455891>

Submitted on 29 Nov 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Paleoecology of the Rhinocerotidae (Mammalia, Perissodactyla) from Béon 1, Montréal-du-Gers (late early Miocene, SW France): Insights from dental microwear texture analysis, mesowear, and enamel hypoplasia

Manon Hullot, Yves Laurent, Gildas Merceron, and Pierre-Olivier Antoine

ABSTRACT

The late early Miocene Béon 1 locality has yielded an abundant vertebrate fauna with more than 5,000 remains assigned to four rhinocerotid species: the stem rhinocerotine *Plesiaceratherium mirallesi*, the teleoceratines *Prosantorhinus douvillei* and *Brachypotherium brachypus*, and the early-diverging elasmotheriine *Hispanotherium beonense*. Such a profusion of closely related large herbivore species co-occurring raises questions about habitat capacity and niche partitioning. To investigate potential niche partitioning of Béon 1 rhinocerotids, we studied their ecology through texture microwear (short-term diet proxy), mesowear (long-term diet proxy), and enamel hypoplasia (environmental stress proxy). The mesowear revealed no significant differences between the species, with low scores suggesting browsing preferences, while microwear suggested subtle dietary variations. We concluded that *Plesiaceratherium mirallesi* and *Prosantorhinus douvillei* were browsers probably consuming a lot of leaves, while *Brachypotherium brachypus* and *Hispanotherium beonense* were mixed-feeders. Concerning hypoplasia, the overall prevalence was high at Béon 1, with more than 25 % of the teeth affected (216 teeth out of 832). There were, however, huge discrepancies depending on the species, tooth loci, or type of hypoplasia considered. Interestingly, *H. beonense* had the greatest dietary flexibility and the lowest hypoplasia prevalence (13.04 %). Both teleoceratines were very affected by hypoplasia, suggesting whether a comparable high sensitivity to stress (phylogeny) or similar habitat-related stresses (environment). The most affected loci were p4 (48.15 %), m3 (46.81 %), and D4 (46.15 %), reflecting vulnerability periods around weaning, environment-related, and near birth, respectively.

Manon Hullot. Institut des Sciences de l'Évolution, UMR5554, Université de Montpellier, CNRS, IRD, EPHE, Place Eugène Bataillon, CC064, 34095 Montpellier, France. manon.hullot@gmail.com
Yves Laurent. Muséum de Toulouse, 35 Allées Jules Guesde, F-31000 Toulouse, France.
Yves.LAURENT@toulouse-metropole.fr

Hullot, Manon, Laurent, Yves, Merceron, Gildas, and Antoine, Pierre-Olivier. 2021. Paleoecology of the Rhinocerotidae (Mammalia, Perissodactyla) from Béon 1, Montréal-du-Gers (late early Miocene, SW France): Insights from dental microwear texture analysis, mesowear, and enamel hypoplasia. *Palaeontologia Electronica*, 24(2):a27. <https://doi.org/10.26879/1163>
palaeo-electronica.org/content/2021/3433-beon-1-rhinocerotids-ecology

Copyright: August 2021 Palaeontological Association.

This is an open access article distributed under the terms of Attribution-NonCommercial-ShareAlike 4.0 International (CC BY-NC-SA 4.0), which permits users to copy and redistribute the material in any medium or format, provided it is not used for commercial purposes and the original author and source are credited, with indications if any changes are made.
creativecommons.org/licenses/by-nc-sa/4.0/

Gildas Merceron. PALEVOPRIM, UMR 7262, CNRS, Université de Poitiers, 86073 Poitiers, France.
gildas.merceron@univ-poitiers.fr

Pierre-Olivier Antoine. Institut des Sciences de l'Évolution, UMR5554, Université de Montpellier, CNRS, IRD, EPHE, Place Eugène Bataillon, CC064, 34095 Montpellier, France. pierre-olivier.antoine@umontpellier.fr

Keywords: environmental stress; feeding ecology; General Linear Mixed Models (GLMM); niche partitioning; rhinocerotids; ungulates, rhinoceros, France

Submission: 21 April 2021. Acceptance: 6 August 2021.

INTRODUCTION

Using multiple approaches for addressing a question proves to be useful in paleontology because it allows for yielding more robust results and broader insights. Despite that fact, many studies remain centered around one method only, which may lead to misinterpretations (Jones and DeSantis, 2017). For instance, both microwear and mesowear are diet-related proxies, but they are relevant at different time scales (Davis and Pineda-Munoz, 2016). The study of mesowear alone, which is a cumulative over-life dietary signal (Ackermans et al., 2020), might miss seasonal variations or near-death modifications of the diet, while microwear alone, which documents the diet during the last days to weeks (Grine, 1986; Winkler et al., 2020), might only reveal a subset of dietary prefer-

ences (Schulz et al., 2007; Jones and DeSantis, 2017; Rivals et al., 2020).

The locality of Béon 1 (Montréal-du-Gers, Occitanie, France; Figure 1) is dated from the late early Miocene, about 17 million years ago and corresponding to the upper part of the MN4 of the European Neogene Mammal zones system (Antoine and Duranthon, 1997; Rage and Bailón, 2005; Antoine et al., 2018). The locality landscape is reconstructed as an oxbow lake surrounded by a wooded savannah-like biome (Duranthon et al., 1999; Rage and Bailón, 2005). It has yielded an abundant and diverse vertebrate fauna, including five species of rhinocerotids (Crouzel et al., 1988; Antoine and Duranthon, 1997; Duranthon et al., 1999; Orliac et al., 2006; Antoine et al., 2018). Four of these species – *Plesiaceratherium mirallesi*

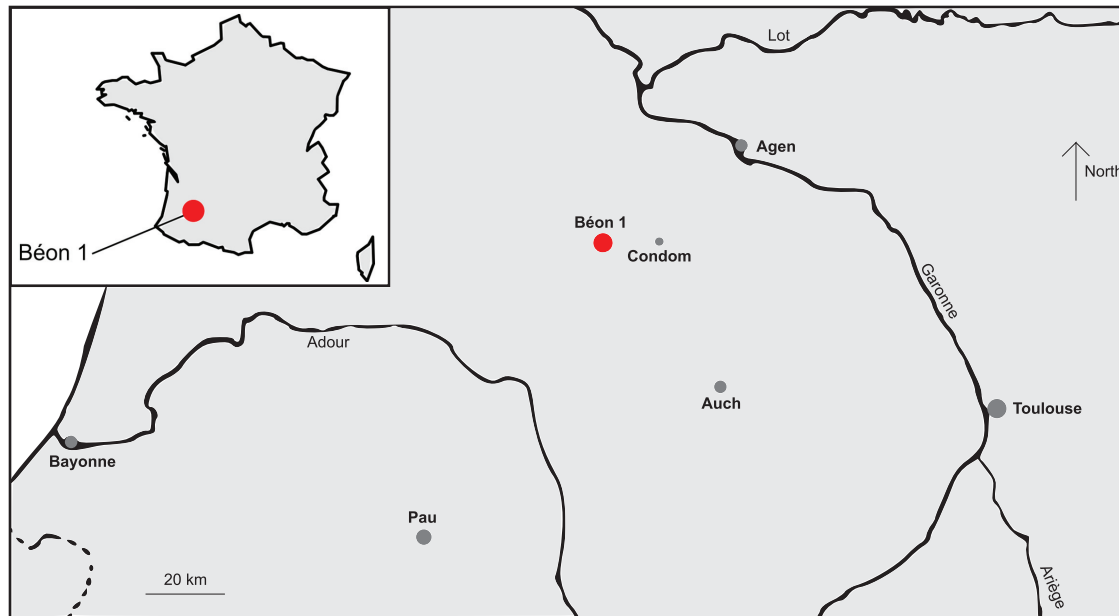


FIGURE 1. Location map of Béon 1 locality, Montréal-du-Gers (MN4; mid-Orleanian, late early Miocene, south western France). The locality of Béon 1 is located (red circle) on the map of France (upper left corner) and on the zoom of south western France. Main cities (grey circles; bold) and rivers are indicated on the zoomed map. Dashed line represents the Spain-France frontier. Modified from Antoine and Duranthon (1997).

(abbreviated *Pl. mirallesi* hereunder), *Prosantorhinus douvillei* (abbreviated *Pr. douvillei* hereunder), *Brachypotherium brachypus*, and *Hispanotherium beonense* – are documented by abundant dental samples, with over 1000 teeth found either isolated or in tooth rows (Antoine and Duranthon, 1997; Antoine et al., 2000; Hullot and Antoine, 2020). The postcranial material is also abundant (Antoine, 2002), and it allows for considering the following minimum number of individuals (MNI): five individuals of *B. brachypus*, 40 of *Pr. douvillei*, 38 of *Pl. mirallesi*, and seven of *H. beonense*, for a total MNI equaling 90. Regarding other herbivore mammals, Béon 1 locality yielded proboscideans, such as the deinotheriid *Prodeinotherium bavaricum* and the elephantoid *Gomphotherium* gr. *sylvaticum* (Antoine and Duranthon, 1997; Tassy, pers. comm. 2021), a wide array of ruminants (Duranthon et al., 1995) and suoids (Orliac, 2006; Orliac et al., 2006), and a three-toed equoid (*Anchitherium aurelianense*; Crouzel et al., 1988). This species-rich herbivore assemblage is interpreted as resulting from an attritional accumulation and most specimens originate from a single 15-50 cm thick dark clay layer, distributed over a 500 m² surface, in one piece (Antoine and Duranthon, 1997; Duranthon et al., 1999; Hullot and Antoine, 2020). Indeed, numerous marks of trampling are observed on cranio-dental and postcranial remains, most being compatible with three- and four-toed rhinocerotid feet (see Hullot and Antoine, 2020 for further details on geology and taphonomy of the locality). As such, it offers a perfect setting for testing resource availability and potential niche partitioning among associated rhinocerotids, even if direct ecological interaction could not be evidenced a priori between these four species.

Previous studies investigated the rhinocerotids from Béon 1 but remained evasive on their ecologies. Nevertheless, different habitats were inferred for these four species based on their morphology. Both teleoceratines (*B. brachypus* and *Pr. douvillei*) are considered swamp dwellers due to their short limbs and stout silhouette, while the slender-limbed hornless rhinocerotine *Pl. mirallesi* is interpreted as living in open woodland, and the early-diverging elasmotheriine *H. beonense* in savannah-like open environments (Antoine and Duranthon, 1997; Antoine et al., 2000; Bentaleb et al., 2006). Enamel carbonate oxygen isotope composition ($\delta^{18}\text{O}$) suggested that all species shared a common source of drinking water (Bentaleb et al., 2006), i.e., the oxbow lake itself or the main river, falling in the home range of all rhinocerotid species

(10-100 km² in extant species; Owen-Smith, 1988) without necessarily being their preferred habitat. Eventually the few differences between the $\delta^{18}\text{O}$ of both teleoceratine species, thought to live in the same habitat (swamp), might be due to different dietary preferences with different oxygen content (Bentaleb et al., 2006). All these previous findings point towards niche partitioning of Béon 1 rhinocerotids.

To investigate the ecology and the eventual niche partitioning of these four rhinocerotid species from the Miocene locality of Béon 1, we used several approaches on the abundant dental material. For this matter, multi-proxy approaches proved successful in revealing subtle niche partitioning in extant species (Merceron et al., 2021). Our approach combined dental microwear textures analyses (DMTA), mesowear, and enamel hypoplasia. Microwear (DMTA) and mesowear allowed for assessing dietary preferences at two different time scales (short term for microwear and long term for mesowear), and enamel hypoplasia revealed stress vulnerability. The objectives of this paper were to: i) characterize the dietary preferences of Béon 1 rhinos, ii) investigate intraspecific stress vulnerability, and iii) reveal subtle details of niche partitioning and competition.

ABBREVIATIONS

For teeth, lowercases refer to lower teeth and uppercases to upper teeth as follows: d/D – deciduous teeth, p/P – permanent premolars, m/M – permanent molars. *Prosantorhinus douvillei* is abbreviated as *Pr. douvillei* and *Plesiaceratherium mirallesi* as *Pl. mirallesi* to avoid confusion.

DMTA: Dental microwear texture analyses (here using scale-sensitive fractal analyses)/

Abbreviations of DMTA parameters as follows, epLsar: anisotropy, Asfc: complexity, FTfv: fine textural-fill volume, HAsfc: heterogeneity of the complexity (whether it is HAsfc9 or HAsfc81).

Other abbreviations, OR: occlusal relief, CS: cusp shape, LEH: linear enamel hypoplasia, MHNT: Muséum d'Histoire Naturelle de Toulouse.

MATERIALS AND METHODS

All material from Béon 1 is stored permanently at the MHNT. We studied all available teeth, both isolated and from tooth rows, for a total amount of around 1,000 teeth. However, depending on the constraints of the different methods used (microwear, mesowear, and hypoplasia) the dataset was restricted as detailed thereafter.

Dental Microwear Texture Analyses (DMTA)

Dental microwear is the study of microscopic scars left by food at the surface of the enamel. The shape and quantity of these scars are linked to the type of food items (and especially to their mechanical properties) consumed by the individual and the way to comminute these items, during the last few meals prior to its death (Hoffman et al., 2015; Winkler et al., 2020). Dental microwear is a short-term (Teaford and Oyen, 1989; Winkler et al., 2020) proxy of diet (Teaford, 1991; Rivals et al., 2014; Ramdarshan et al., 2016), widely used in paleontology for various mammalian clades (Grine, 1986; Solounias and Semprebon, 2002; Calandra et al., 2008; Jones and DeSantis, 2017; Berlioz et al., 2018; Rivals et al., 2020). In this study, we used Dental Microwear Texture Analyses (DMTA) with sensitive-scale fractal analyses to study microwear. This DMTA method was developed by Ungar et al. (2003), and we followed a protocol based on Scott et al. (2005).

We sampled one molar – preferentially the second (first or third otherwise) whether it was upper or lower, left, or right – displaying a good state of preservation and a sufficient wear, for each specimen. We analyzed two facets, one shearing and one grinding, from the same enamel band on the lingual part of the protocone on upper molars and distally to the protoconid or hypoconid (if the protoconid facet was unavailable) on lower molars (Figure 2). Sometimes the shearing facet was not developed and could not be sampled. Our dataset was also split according to the preparation method used on the specimens – hand-prepared or sand-prepared (using sand blasting) – to test the effect of sand blasting on microwear signals.

After selection, the facets of interest were cleaned twice with cotton-swabs soaked in acetone

or ethanol, and two silicone molds were produced (Regular Body President, ref. 6015 - ISO 4823, medium consistency, polyvinylsiloxane addition type; Coltene Whaledent). The second mold was used for further analyses. Scans of the molded facets were obtained with a Leica DCM8 confocal profilometer (“TRIDENT” profilometer housed at the PALEVOPRIM, CNRS, and University of Poitiers) using white light confocal technology with a Leica 100× objective (Numerical aperture: 0.90; working distance: 0.9 mm; Leica Microsystems). The .Plp files obtained were then pre-treated under Leica-Map v.8.2. (Leica Microsystems) as follows: surface inversion (scans produced on negative replicas), replacement of the missing points (i.e., non-measured, less than 1%) by the mean of the neighboring points, removal of aberrant peaks with automatic operators including a morphological filter (see Supplementary Information in Merceron et al., 2016 for details), and levelling of the surfaces. A 200×200-μm area (1551×1551 pixels) was selected and saved as a digital elevation model (.sur) to be used for DMTA. As some surfaces included Hunter-Schreger bands we ran a final removal of form (polynomial of degree 8) and a last levelling to temper the potential relief effects in the parameter calculations. These surfaces were then analyzed using the Scale-Sensitive Fractal Analysis with Toothfrax and Sfrax softwares (Surfract, www.surfract.com) following Scott et al. (2006) or using Sfrax and LeicaMap. For our study we selected five classical DMTA parameters:

- anisotropy (exact proportion of length-scale anisotropy of relief; epLsar), which measures the orientation concentration of surface roughness;

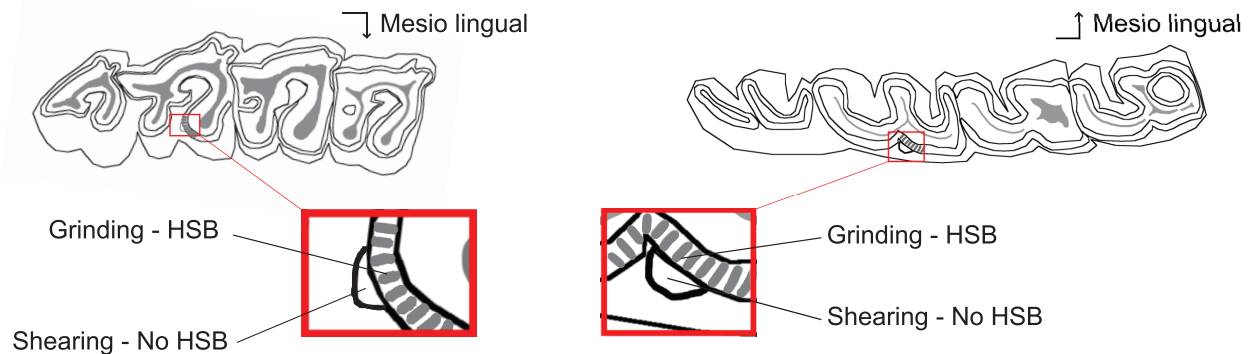


FIGURE 2. Localization of the microwear facets on rhinocerotid molars. Position of the two microwear facets (grinding and shearing) on the second upper molar (left) and second lower molar (right). Both facets are sampled on the same enamel band with (grinding) or without (shearing) Hunter-Schreger bands (HSB). Modified after Hullot et al. (2019).

- complexity (area-scale fractal complexity; Asfc) that estimates the roughness at a given scale;
- heterogeneity of complexity (heterogeneity of area-scale fractal complexity here at 3×3 and 9×9; HAsfc9 and HAsfc81) reflecting the variation of complexity within the studied zone;
- fine textural fill volume (here at 0.2 µm; FTfv) estimated by filling the surface with square cuboids of different volumes and is linked to surface relief (see Scott et al., 2006 for details).

We used a dataset modified from that of Hullot et al. (2019), with two new specimens as detailed below, and encompassing all five living rhinoceros' species to facilitate the interpretation of fossil textural microwear. The dataset consists of 17 specimens of *Ceratotherium simum* (white rhinoceros), four of *Dicerorhinus sumatrensis* (Sumatran rhinoceros), 21 of *Diceros bicornis* (black rhinoceros), 15 of *Rhinoceros sondaicus* (Javan rhinoceros; one new specimen), and five of *Rhinoceros unicornis* (Indian rhinoceros; one new specimen).

Mesowear

Dental mesowear studies ungulate molars occlusal morphology through relief and sharpness of cusp apices to categorize the gross dental wear observed into dietary preferences. Mesowear typically relies on two variables (Figure 3) scored originally on second upper molars as proposed by Fortelius and Solounias (2000):

- occlusal relief (OR) that can be high or low;
- cusp shape (CS) taking the values sharp, round, or blunt.

With this method, browsers are characterized by a high percentage of attrition-dominated specimens with high relief and sharp cusps, while grazers are abrasion-dominated with low relief and blunt cusps (Fortelius and Solounias, 2000). Nevertheless, as abrasion is cumulative over-life, there is an ontogeny effect in the mesowear signal (Fortelius and Solounias, 2000), which we took into account by using age classes as defined by Hillman-Smith et al. (1986).

In this study, we used three different mesowear scores to assess dietary preferences (Table 1):

- Score A: better known as 'Mesowear I & II—Expanded' (Solounias et al. 2014; Ackermans 2020) with refined categories for both OR and CS (e.g., high-high, high-low, round-round)

giving scores between 1 (high-high sharp) and 17 (low blunt), based on the work of Winkler and Kaiser (2011);

- Score B: or 'Mesowear II' which is a conversion of the classic original scoring ('Mesowear I') from Fortelius and Solounias (2000) as proposed by Rivals et al. (2007) into a score ranging from 0 (high relief sharp cusp) to 3 (low relief blunt cusp);
- Ruler: mesowear II ruler developed by Mihalbachler et al. (2011) on equid perissodactyls, giving scores from 0 to 6.

These methods were applied to all available upper permanent molars (M1, M2, and M3) and to the third and fourth upper deciduous teeth (D3 and D4). Mesowear was assessed on the paracone and not on the sharpest cusp (paracone or metacone) because significant differences between cusps have been reported in rhinoceros (Taylor et al., 2013). We studied deciduous teeth as they are molarized in rhinocerotids, which means that mesowear methods are easily transposable, and as they might give insights of the diet of juvenile and subadult individuals, and not only of adult individuals. Examples of scoring using the three different methods are provided in Figure 3.

Enamel Hypoplasia

Enamel hypoplasia is a permanent, sensitive, but non-specific and individual marker of stress. Causes of this enamel defect can be very diverse (around 100 factors interfering with normal amelogenesis in human are listed by Small and Murray, 1978), such as environmental stress (e.g., drought, food shortage; Skinner and Pruetz, 2012; Upex and Dobney, 2012), physiological stress (e.g., disease, parasitism; Suckling et al., 1986; Rothschild et al., 2001; Niven et al., 2004), and psychological stress (e.g., depression; Guatelli-Steinberg, 2001).

All cheek teeth available (deciduous and permanent) were included in the study of hypoplasia, with the exception of teeth too damaged, too worn, with not enough enamel visible (e.g., tooth unerupted in bone, sediment occluding), or not properly identified. This exclusion was considered because absence of hypoplasia in worn or damaged teeth might generate false negatives, and because hypoplasia on unidentified teeth is difficult to link to potential causes. This exclusion concerned 81 teeth: 10 permanent in *B. brachypus*, 19 (two deciduous – 17 permanent) in *H. beonense*, 28 (one deciduous – 27 permanent) in *Pl. mirallesi*, and 24 (four deciduous – 20 permanent) in *Pr. douvillei*. The analyzed sample represents a total of

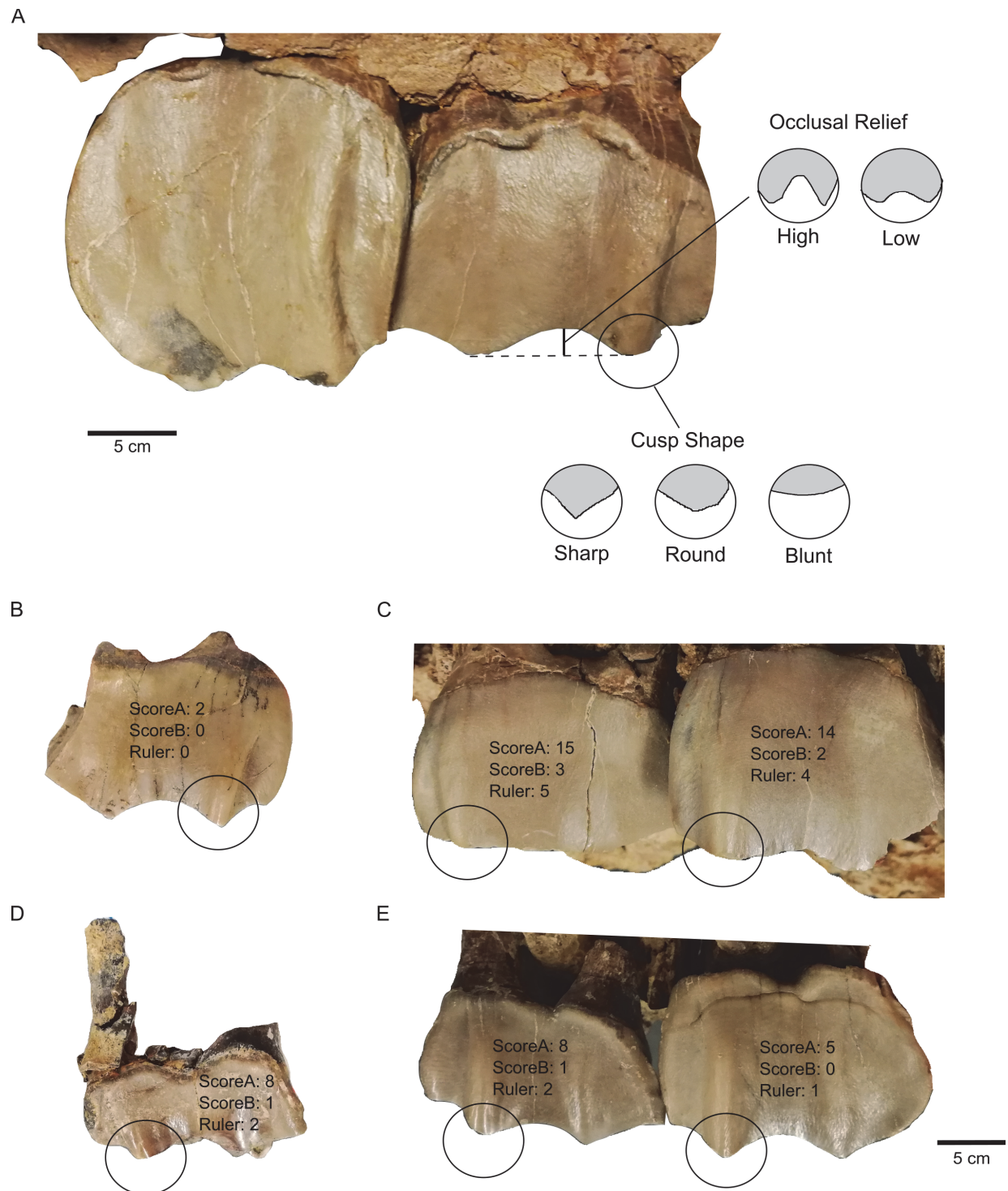


FIGURE 3. Principle of mesowear scoring with the main variables illustrated (occlusal relief and cusp shape) and examples on rhinocerotid teeth. A- Typically two parameters are studied in mesowear: cusp shape and occlusal relief. Cusp shape can be sharp, round or blunt, while occlusal relief is whether high or low. Illustration on the upper right M1 of the specimen MHNT.PAL.2004.0.58 (*H. beonense*). Examples of mesowear scores using the three methods tested in this study (ScoreA, ScoreB, Ruler) are provided on the paracone of the following specimens: B- Right D4 of MHNT.PAL.2015.0.1204 (G2 685; *Pl. mirallesi*), C- Left M1 and M2 MHNT.PAL.2015.0.277 (*Pr. douvillei*), D- Left D4 of MHNT.PAL.2015.0.1204 (Béon F2 193; *Pl. mirallesi*), E- Left D3 and D4 of MHNT.PAL.2015.0.2796 (*Pr. douvillei*). ScoreA: mesowear score based on Winkler and Kaiser (2011); B- ScoreB: mesowear score adapted from Fortelius and Solounias (2000); C- Ruler: mesowear score based on Muhlbachler et al. (2011).

TABLE 1. Conversion of occlusal relief (OR) and cusp shape (CS) into a mesowear score depending on the method. ScoreA: extended mesowear (refined categories for CS and OR) giving scores between 1 and 17, based on the work of Winkler and Kaiser (2011); ScoreB: conversion of the classic original scoring from Fortelius and Solounias (2000) as proposed by (Rivals et al., 2007) into a score ranging from 0 to 3; Ruler: mesowear ruler developed by Muhlbachler et al. (2011), giving scores from 0 to 6.

Occlusal relief	Cusp shape	ScoreA	ScoreB	Ruler
High-high	Sharp	1	-	-
High	Sharp	2	0	0
High-low	Sharp	3	-	-
High-high	Round-sharp	4	-	-
High	Round-sharp	5	-	1
High-low	Round-sharp	6	-	-
High-high	Round	7	-	-
High	Round	8	1	2
High-low	Round	9	-	-
High	Round-round	10	-	-
High-low	Round-round	11	-	3
Low	Sharp	12	-	-
Low	Round-sharp	13	-	-
Low	Round	14	2	4
Low	Round-round	15	-	5
High-low	Blunt	16	-	-
Low	Blunt	17	3	6

832 teeth (197 deciduous molars and 635 permanent premolars and molars).

To our knowledge, no standard protocol nor threshold is available to differentiate normal from pathological enamel in any given species. Here, we have chosen to investigate enamel hypoplasia with the naked eye. The approach consisted in the macroscopical identification of defects, following the Fédération Dentaire Internationale index (1982) and associated caliper measurements. In this paper, we distinguished three types of hypoplasia as illustrated in Figure 4:

- linear enamel hypoplasia (LEH), defect forming a line around the crown;
- pitted hypoplasia, restricted rounded defect; and
- aplasia, zone entirely missing enamel.

Figure 4 also details the measurements taken: distance to enamel-dentine junction (linked to the age at which the defect occurred) and width of the defect when applicable (linked to the severity and

duration of the stress). Other qualitative parameters were recorded: the type of tooth affected (e.g., second upper right molar), the position of the defect on the crown (labial or lingual, near which cusp), and its severity.

Statistics and Figures

All statistics were conducted in R (R Core Team, 2018. R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL <https://www.R-project.org/>). Figures were done using R packages ggplot2 (Wickham, 2016), cowplot (Wilke, 2020), and gridExtra (Auguie, 2017), as well as Inkscape v.0.91.

General Linear Mixed Models (GLMM) were applied to our data. They were built using a R code modified from Arman et al. (2019), based on the package lme4 (Bates et al., 2015), and adapted to each tested response variable. For DMTA, response variables were the five DMTA parameters (epLsar, Asfc, FTfv, HASfc9, and HASfc81) and factors were: specimen (random factor), species, tooth (e.g., first molar, fourth premolar), position (upper or lower), side (left or right), cusp (protocone, protoconid, hypoconid), and facet (grinding or shearing). For mesowear, the variables were ScoreA (from 1 to 17; Winkler and Kaiser, 2011), ScoreB (from 0 to 3; Rivals et al., 2007), and Ruler (from 0 to 6; Muhlbachler et al., 2011), and the factors were: specimen (random factor), species, tooth (e.g., first molar, fourth premolar), category (deciduous or permanent), side (left or right), wear (low = stages 1 to 4, average = stages 5 to 7, high = stages 8 to 10 according to scale defined by Hillman-Smith et al., 1986), ontogeny (adult, subadult, juvenile; as defined in Hullo and Antoine, 2020), and cusp (sharpest or paracone). For hypoplasia, response variables were Hypo (1 or 0 for presence or absence of hypoplasia), Defect (e.g., LEH, Pits, Aplasia; converted to numbers), Multiple (number of defects), Severity (0 to 4), and factors were: specimen (random factor), species, tooth (e.g., first molar, fourth premolar), position (upper or lower), side (left or right), defect (converted to factor; for response variables Severity and Multiple), and wear (low, average, high; same categories as above). An example of this code applied to mesowear ScoreA is given in Appendix 1.

Model construction followed a bottom-up approach (i.e., adding factors incrementally to the model), starting with the factor specimen alone, which was the only random factor of our dataset. For the second model we added the factor species,

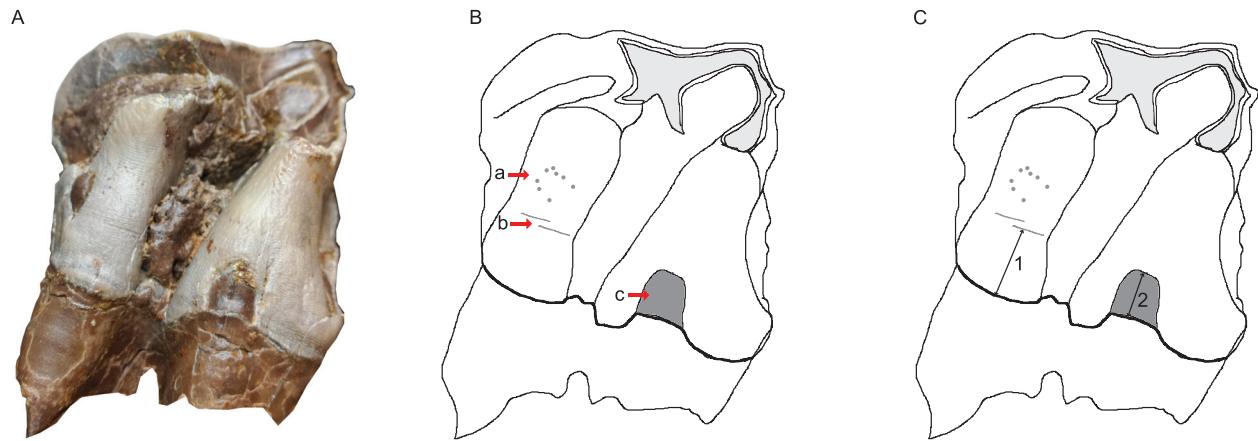


FIGURE 4. The three different types of hypoplasia considered in this study and the associated measurements. A- Lingual view of right M2 of the specimen MHNT.PAL.2004.0.58 (*H. beonense*) displaying three types of hypoplasia. B- Interpretative drawing of the photo in A illustrating the hypoplastic defects: a- pitted hypoplasia, b- linear enamel hypoplasia, and c- aplasia. C- Interpretative drawing of the photo in A illustrating the measurements: 1- distance between the base of the defect and the enamel-dentin junction, 2- width of the defect (when applicable).

as this was the principal interest of our analyses. We tested few interactions (e.g., Species x Facet for microwear) in the models as most factors were considered independent and to avoid unnecessarily complex models (models with interaction effects rarely selected after preliminary analysis; Arman et al., 2019). We created additional sets of models by adding a different factor (e.g., ontogeny, tooth, category, side, wear, cusp) to the second model (species + specimen) and selected the model that produced the lowest AIC score (Akaike's Information Criterion) as the base for the next set of models. We repeated this process until there was no improvement in AIC score. After selecting the best model (lowest AIC), we checked for over-dispersion (estimated through the ratio of deviance and degrees of freedom) and corrected the model by using quasi-Poisson or quasi-Binomial laws from the MASS package (Venables and Ripley, 2002) if necessary. In total, 200 models were compared across the 12 response variables (see Appendices 2 to 5).

RESULTS

DMTA: Modelling, Impact of Sand Preparation, and Dietary Preferences

Median, mean, and standard deviation of the mean were calculated for every DMTA parameter (anisotropy, complexity, FTfv, HASfc9, and HASfc81) by species, facet, and preparation type (hand or sand blasting). All results are detailed in Table 2. As for dietary interpretation, only hand-prepared

specimens are usable. Therefore, we will mainly refer to this subsample and compare it to the sand-prepared subsample to assess the influence of sand blasting on the microwear signal. In the hand-prepared subset, values of complexity were relatively similar among the four species on both facets, especially both teleoceratines, with most specimens plotting between 0.5 and 2 (Figure 5). Mean and median of complexity were low to moderate in all four species on both facets, ranging from 0.72 to 1.69. Within this sample, *Pl. mirallesi* had the highest values of anisotropy on the shearing facets (mean and median above 5×10^{-3}) but values similar to those of *Pr. douvillei* and *H. beonense* on the grinding ones ($2.5\text{--}3 \times 10^{-3}$; Figure 5, Table 2). The lowest values of anisotropy were those of *B. brachypus*, mostly around 2×10^{-3} . The heterogeneities of complexity (HASfc9 and HASfc81) were low for *Pl. mirallesi* but moderate in other species on both facets (except shearing of *Pr. douvillei*). Eventually, FTfv had the highest value on the grinding facets of two teleoceratines, with a mean of 4.78×10^4 for *B. brachypus* and 5.21×10^4 for *Pr. douvillei*, and the lowest value on the shearing facets of *B. brachypus*, with a mean of 1.63×10^4 . Overall, the shearing facet seems to discriminate better between the species, especially through anisotropy (Figure 5), though both facets are useful for dietary studies, as they give insights into different moments of the mastication.

Sand-prepared specimens show lower values and less dispersion in anisotropy on both facets, but higher values and more dispersion in complex-

TABLE 2. Median, mean, and standard deviation of the mean of DMTA parameters by species, type of preparation and facet. lepLsar: anisotropy, Asfc: complexity, FTfv: fine textural-fill volume, HAsfc: heterogeneity of the complexity (whether it is HAsfc9 or HAsfc81). Gr: grinding facet; Sh: shearing facet.

Species Prep. Facet		<i>B. brachypus</i>				<i>Pr. douvillei</i>				<i>Pl. mirallesi</i>				<i>H. beonense</i>			
		Hand		Sand		Hand		Sand		Hand		Sand		Hand		Sand	
		Gr	Sh	Gr	Sh	Gr	Sh	Gr	Sh	Gr	Sh	Gr	Sh	Gr	Sh	Gr	Sh
N		7	5	6	3	15	4	13	8	14	12	19	12	6	3	2	1
Asfc	Median	1.20	0.72	2.72	0.63	1.24	0.77	2.00	1.96	0.98	0.96	2.23	1.31	0.76	0.92	0.73	1.47
	Mean	1.50	0.92	3.36	0.71	1.50	1.03	3.67	2.00	1.27	1.02	2.20	1.38	1.69	0.75	0.73	1.47
	SD	0.73	0.65	2.94	0.22	1.00	0.52	3.27	0.89	0.94	0.67	1.07	0.53	1.89	0.31	0.01	NA
epLsar (*10 ⁻³)	Median	1.70	1.83	1.90	2.96	2.60	2.53	2.32	1.94	2.46	5.60	2.39	2.06	3.01	3.33	1.46	1.33
	Mean	2.56	1.98	1.87	3.58	2.81	2.41	2.30	2.39	3.36	5.70	2.43	2.23	2.95	3.67	1.46	1.33
	SD	1.58	0.44	0.78	3.01	1.68	1.34	1.79	1.74	2.14	2.21	1.06	1.09	0.93	2.02	0.12	NA
FTfv (*10 ⁴)	Median	4.76	1.98	6.58	2.38	5.39	2.66	4.27	4.29	4.05	2.34	5.27	3.13	3.73	4.23	2.65	2.76
	Mean	4.78	1.63	5.94	2.54	5.21	3.85	4.80	3.99	4.20	2.35	5.31	2.97	4.22	4.50	2.65	2.76
	SD	1.10	0.96	1.99	0.89	2.50	2.57	2.14	1.67	1.99	1.34	1.97	0.61	1.57	0.73	0.31	NA
HAsfc9	Median	0.27	0.23	0.32	0.24	0.32	0.18	0.30	0.35	0.21	0.16	0.31	0.29	0.27	0.22	0.25	0.18
	Mean	0.34	0.27	0.33	0.24	0.32	0.18	0.34	0.58	0.26	0.19	0.39	0.34	0.29	0.31	0.25	0.18
	SD	0.20	0.11	0.10	0.07	0.14	0.03	0.23	0.58	0.10	0.15	0.23	0.17	0.13	0.20	0.08	NA
HAsfc81	Median	0.49	0.46	0.69	0.52	0.56	0.37	0.70	0.78	0.40	0.40	0.72	0.59	0.45	0.53	0.46	0.76
	Mean	0.61	0.51	0.68	0.51	0.66	0.39	0.72	1.30	0.50	0.42	0.76	0.61	0.58	0.58	0.46	0.76
	SD	0.28	0.18	0.13	0.06	0.26	0.10	0.26	1.40	0.20	0.17	0.30	0.14	0.36	0.14	0.05	NA

ity, especially on the grinding facet (Figure 5). This is confirmed by mean and median values of Asfc for all species but *H. beonense* grinding and *B. brachypus* shearing, both exhibiting equivalent or lower values in the sand-prepared subset (Table 2). Concerning anisotropy, the shift is verified for species/facets with high values (*Pl. mirallesi* and *H. beonense*) but is less clear (shearing facet of *Pr. douvillei*) or inverted for other species (*B. brachypus*). The differences between hand- and sand-prepared surfaces are illustrated in Figure 6. Photo-simulations of the sand-prepared surfaces show bigger pits and coarser scratches, and the orientation of the scars is less obvious than in the hand-prepared surfaces. These observations are consistent with previous statements on anisotropy and complexity.

When compared to the living rhinoceros species dataset, we found that no species from Béon 1 was in the extant dietary space, except for the shearing facet of *Pl. mirallesi*, which clusters with the extant strict grazer *C. simum* and the folivore *D. sumatrensis* (Figure 7). The microwear pattern is clearly distinct at the species level, as the proportions of specimens with high anisotropy (epLsar > 0.005) and high complexity (Asfc > 2) change. Regarding sand preparation on the grinding facet, we observe more specimens above the complexity

threshold and sensibly less above the anisotropy one, for all species but *H. beonense*. These tendencies between hand prepared and sand prepared are more subtle on the shearing facet: *H. beonense* displays less anisotropic specimens but still no complex ones, *Pl. mirallesi* presents less anisotropic specimens and slightly more complex ones, *Pr. douvillei* has still no anisotropic specimens but much more complex ones, and *B. brachypus* has more anisotropic specimens but still no complex ones (Figure 7).

For all response variables (epLsar, Asfc, FTfv, HAsfc9, and HAsfc81), model support increased (lower AIC) when intraspecific factors (e.g., tooth loci, preparation type, facet) were included. The final models contained three to five factors. Alongside species and specimen, preparation type was in all final models but that of FTfv. Facet appeared in the final models of epLsar, Asfc, and FTfv. Both HAsfc had the same final model, and epLsar and Asfc shared the same factors but differ in the presence of interactions. Details and comparison of all models can be seen in Appendix 3.

Based on GLMMs results, we can infer the impact of sand preparation on the microwear pattern. Sand-prepared specimens display significantly higher mean complexity (p-value = 5×10^{-4}),

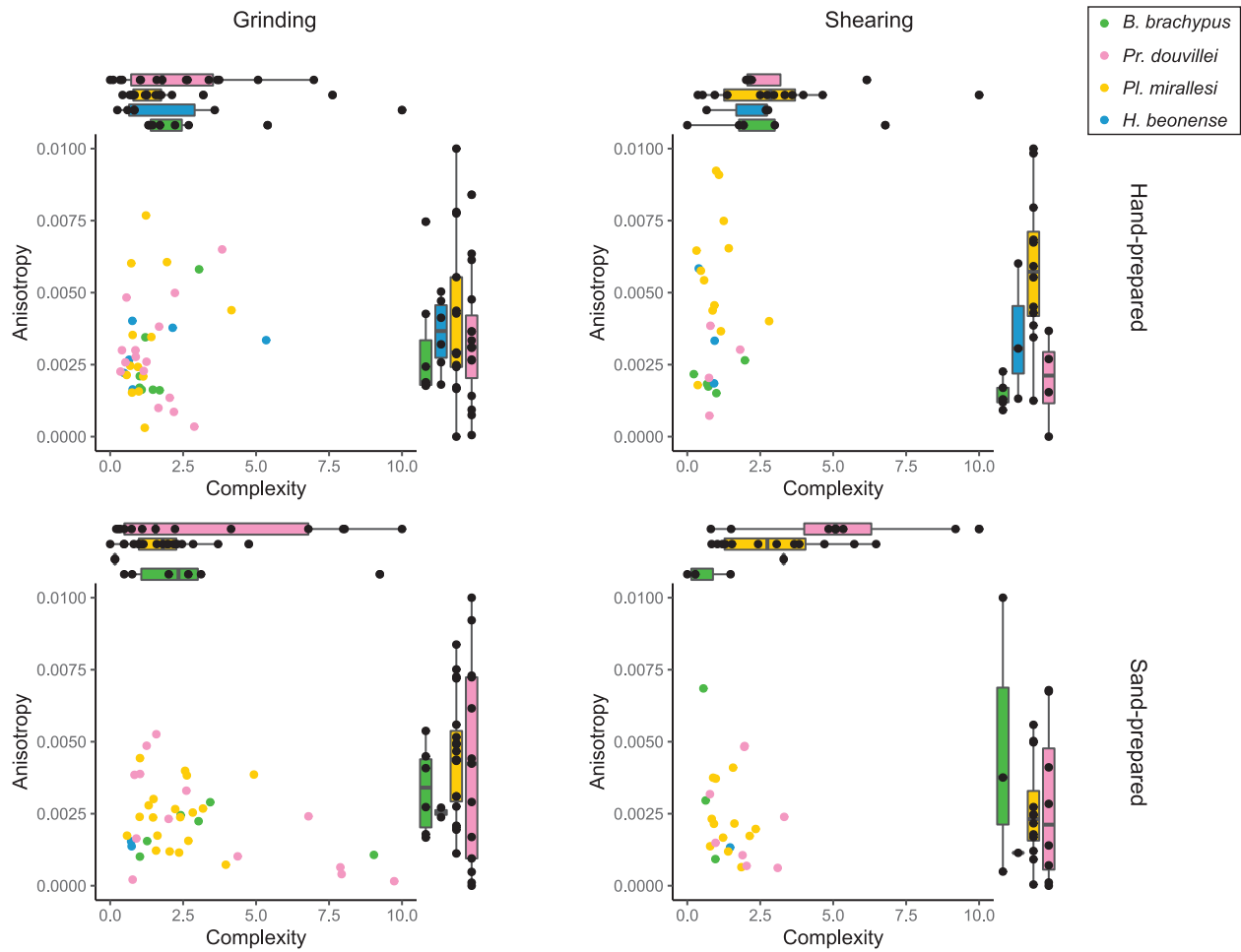


FIGURE 5. Comparison of the DMTA patterns by species, facet and preparation type. Upper graphs: hand-prepared specimens; lower graphs: sand-prepared specimens. Left graphs: grinding facet; right graphs: shearing facet. Box-plots of anisotropy and complexity were plotted along with the dotplots to facilitate graph interpretation.

HAsfc9, and HAsfc81 ($df = 122$, $\alpha = 0.95$, t -values > 2.5). The choice of facet also appears crucial as shearing facet has higher mean anisotropy (p -value $= 4 \times 10^{-4}$), and lower mean complexity (p -value ≈ 0.003) and FTfv (p -values $< 5 \times 10^{-4}$). Significant or nearly significant interspecific differences were observed relatively to *Pl. mirallesi* (set as reference for GLMMs): *Pr. douvillei* had higher mean Asfc (p -value ≈ 0.06) and HAsfc81 ($df = 122$, $\alpha = 0.95$, t -value $= 1.97$). The effects of interactions (species \times preparation type \times facet) were significant for anisotropy. The final model of FTfv includes tooth and position as factors, and it reveals that M3 has significantly lower mean FTfv than M2, and that upper teeth have significantly lower mean FTfv than the lower ones.

Mesowear: Modelling, Impact of Ontogeny, and Dietary Preferences

Out of the complete dataset used for GLMMs, we selected one tooth per specimen (preferentially M2) to calculate mean, standard deviation of the mean, and median of all mesowear scores (ScoreA, ScoreB, Ruler) by species. The results are given in Table 3. Mean of Ruler was around 2 for *Pr. douvillei* and *Pl. mirallesi*, lower in *H. beonense* (1.77), and higher in *B. brachypus* (3.5; Table 3). Median of Ruler was 2 for all species but *B. brachypus*. Mean ScoreA was around 7.5 and mean ScoreB between 0.77 and 0.98 for all species but *B. brachypus* for which they were considerably higher (10.5 and 2 respectively; Table 3). However, mesowear was assessed on only two molars for *B. brachypus*.

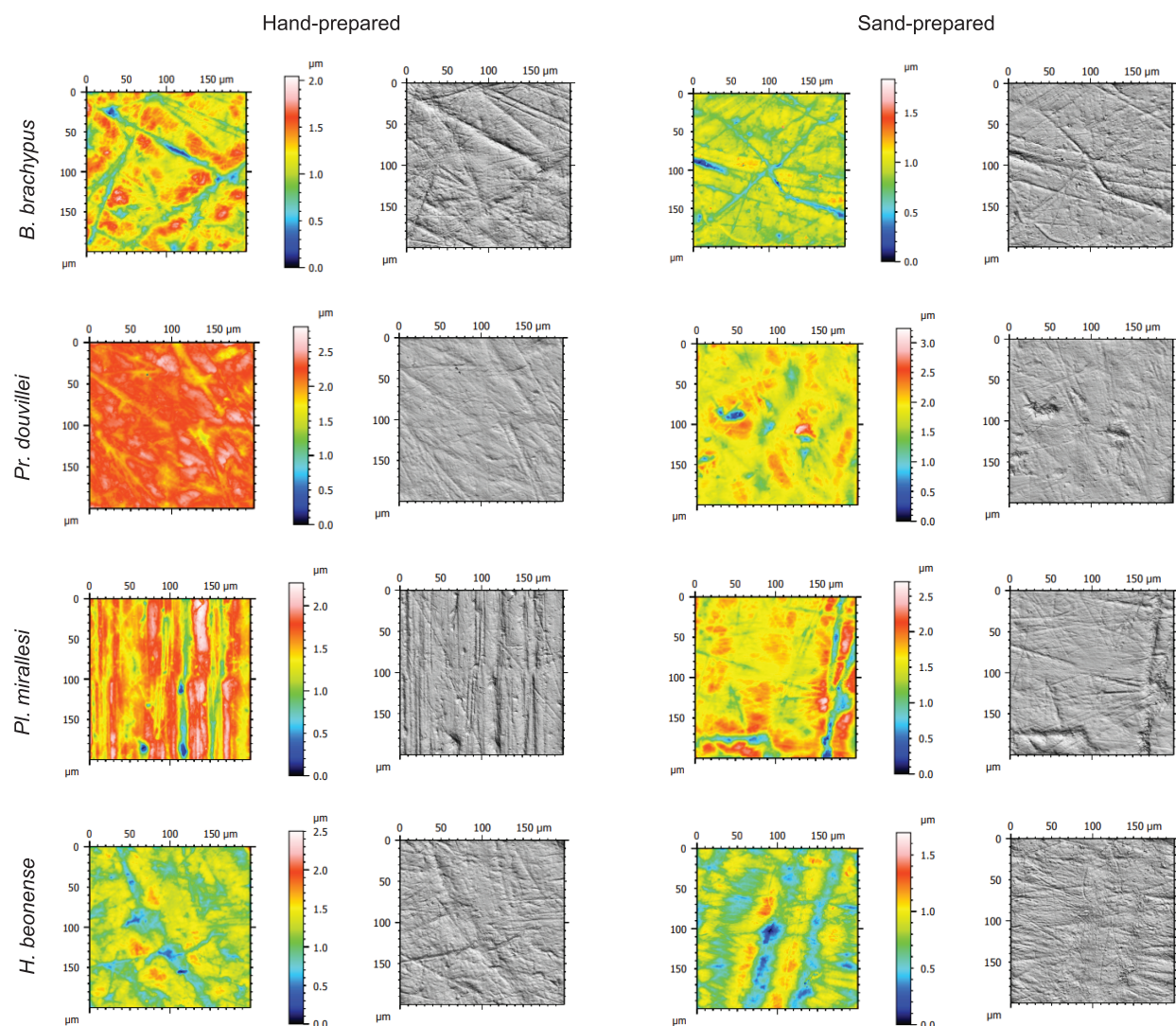


FIGURE 6. Comparison of hand- and sand-prepared DMTA surfaces (200x200 μm) by species. Topography and black and white photosimulation of the following specimens:

B. brachypus – hand-prepared MHNT.PAL.2015.0.1262 right m3 (protoconid, shearing facet) and sand-prepared MHNT.PAL.2015.0.2830 left m2 (hypoconid, shearing facet); *Pr. douvillei* – hand prepared MHNT.PAL.2015.0.1228 left m3 (protoconid, grinding facet) and sand-prepared MHNT.PAL.2015.0.2758 left m2 ptc (protoconid, grinding facet); *Pl. mirallesi* – hand-prepared MHNT.PAL.2015.0.1196 left m2 ptc (protoconid, shearing facet) and sand-prepared MHNT.PAL.2015.0.2794 (2002 E2 30) left m1 (hypoconid, shearing facet); *H. beonense* – hand-prepared MHNT.PAL.2015.0.1140 left m1 (hypoconid, grinding facet) and sand-prepared MHNT.PAL.2015.0.1136.1 right M3 (protocone, grinding facet).

ScoreB and Ruler had lower means when only deciduous teeth were considered and higher ones when permanent teeth alone were studied for all species (Table 3). Median of ScoreB was 1 in all species and sample (all teeth, deciduous only, or permanent only), while that of Ruler was 1 with deciduous teeth and 2 otherwise (all teeth and permanent only). Concerning ScoreA, deciduous teeth also had lower means and permanent teeth higher

ones for *Pl. mirallesi* and *H. beonense*, but it was the opposite for *Pr. douvillei* (Table 3).

Generalized linear mixed-models were built for each response variable (ScoreA, ScoreB, Ruler). For all variables, model support increased (lower AIC) when intraspecific factors (e.g., tooth loci, ontogeny, cusp) were included. The final models contained three to five factors, always including species, specimen, and ontogeny. Final models of ScoreA and ScoreB contained the same factors:

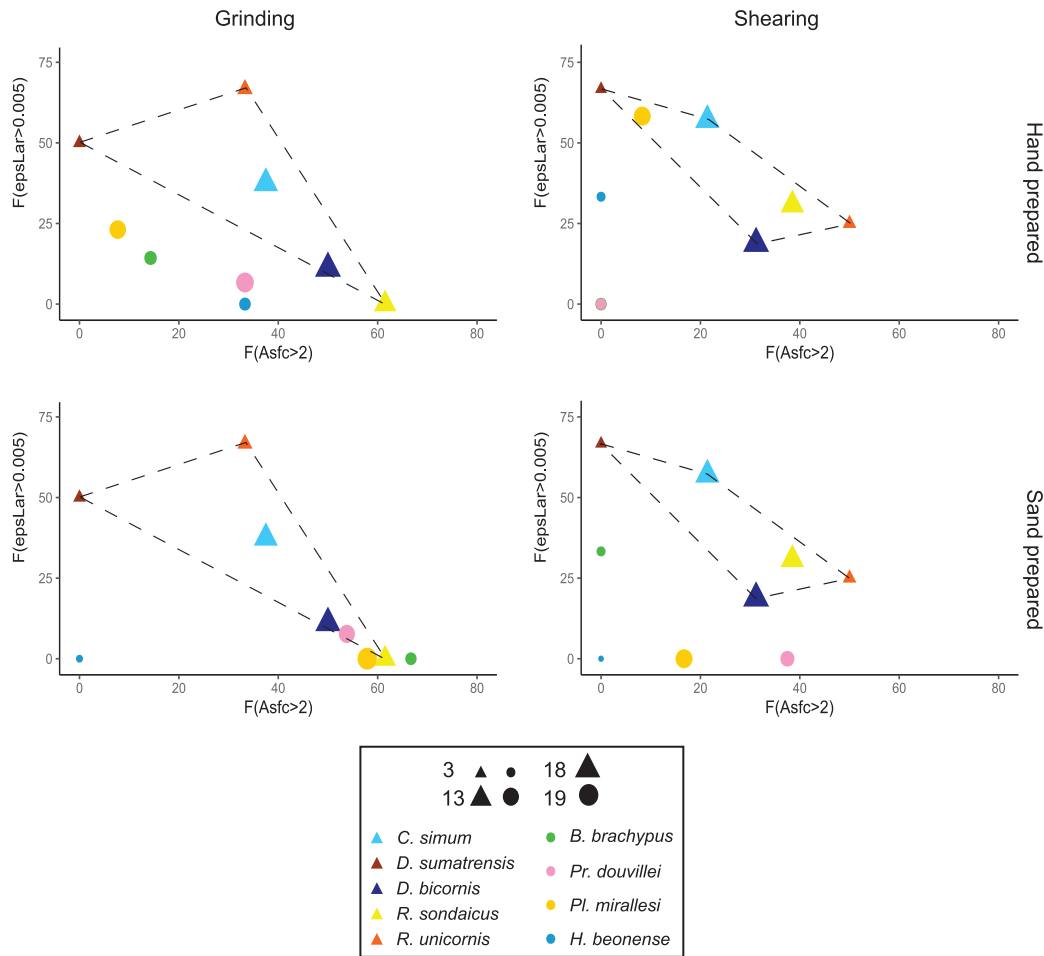


FIGURE 7. Percentages of specimens above anisotropy ($epLsar > 0.005$) or complexity ($Asfc > 2$) cutpoints by species, facet, and preparation type. Triangles: living rhinoceros' species; circles: Béon 1 fossil rhinocerotids; size proportional to the number of specimens.

specimens, species, tooth, cusp, and ontogeny. Details and comparison of all models can be seen in Appendix 4. Based on GLMMs results, we can infer the impact of ontogeny, tooth loci, or cusp sampled on the mesowear pattern. Ontogeny seems to play a crucial role in mesowear, as juveniles and subadults had significantly lower scores than adults for all methods ($p\text{-value} < 0.05$). Surprisingly, we found that D3, D4, M1, and M3 had higher mean mesowear ScoreA and ScoreB than M2. When scored on paracone, ScoreA and ScoreB yielded significantly higher mean values than with the sharpest cusp approach. Eventually, no significant interspecific differences were found in the mesowear score for any method (ScoreA, ScoreB, Ruler).

Despite the absence of significant differences between Béon 1 rhinocerotids, we observed differ-

ent mesowear score profiles (Figure 8). The only two teeth of *B. brachypus* studied yielded very different mesowear scores for all methods (ScoreA: 6 and 15; ScoreB: 1 and 3; Ruler: 2 and 5). Concerning ScoreA, many specimens displayed the average mesowear score of 8 in all three remaining species and most specimens had a mesowear score below 10 (Figure 8). ScoreB for *H. beonense* was 0 or 1 for all specimens, while all scores from 0 to 3 were observed for *Pl. mirallesi* and *Pr. douvillei*. ScoreB over 1 was however limited for *Pr. douvillei*. Eventually, Ruler was between 1 and 3 for *H. beonense* with most specimens having a score of 2. All scores but 5 were observed for *Pl. mirallesi* and *Pr. douvillei* but similarly to ScoreB, the profile was skewed to higher scores in *Pl. mirallesi* (Figure 8).

TABLE 3. Descriptive statistics (sample size, median, mean, standard deviation of the mean) of mesowear score by species and method. ScoreA: extended mesowear giving scores between 1 and 17, based on the work of Winkler and Kaiser (2011); ScoreB: conversion of the classic original scoring from Fortelius and Solounias (2000) as proposed by Rivals et al. (2007) into a score ranging from 0 to 3; Ruler: mesowear ruler developed by Mhielbachler et al. (2011), giving score from 0 to 6. Only one tooth per specimen was considered.

	<i>B. brachypus</i>			<i>Pr. douvillei</i>			<i>Pl. mirallesi</i>			<i>H. beonense</i>		
	ScoreA	Score B	Ruler	ScoreA	ScoreB	Ruler	ScoreA	ScoreB	Ruler	ScoreA	ScoreB	Ruler
All Teeth												
N	2	2	2	31	31	31	32	32	32	13	13	13
Median	10.5	2	3.5	8	1	2	8	1	2	8	1	2
Mean	10.5	2	3.5	7.42	0.90	1.90	7.61	0.98	2	7.19	0.77	1.77
SD	6.36	1.41	2.12	3.69	0.83	1.54	3.93	0.81	1.50	2.63	0.44	0.83
Deciduous only												
N	/	/	/	7	7	7	9	9	9	3	3	3
Median	/	/	/	8	1	1	8	1	1	9	1	1
Mean	/	/	/	7.71	0.71	1.71	6.67	0.67	1.22	6.33	0.67	1.33
SD	/	/	/	3.55	0.76	1.50	2.18	0.5	0.67	4.62	0.58	1.53
Permanent only												
N	2	2	2	27	27	27	24	24	24	9	9	9
Median	10.5	2	3.5	8	1	2	8	1	2	8	1	2
Mean	10.5	2	3.5	7.44	0.93	2	7.98	1.10	2.25	7.39	0.78	1.89
SD	6.36	1.41	2.12	3.94	0.87	1.62	4.32	0.86	1.62	2.15	0.44	0.60

Enamel Hypoplasia

At Béon 1, 216 teeth out of the 832 studied are affected by hypoplasia (all types of hypoplasia, all species, all dental loci), corresponding to an overall prevalence of 25.96 %. This prevalence corresponds to a minimum number of individuals affected by hypoplasia of 27 (Table 4) out of the 61 estimated at Béon 1 using tooth eruption incompatibilities (see Hullot and Antoine, 2020). There are, however, great variations depending on the type of hypoplasia, the species and the tooth locus considered.

As for species, the prevalence was between 13.04 % (12/92 teeth) for *H. beonense* and above 28 % for both *Pl. mirallesi* (98/340; 28.82%) and *B. brachypus* (13/46; 28.26%; Table 5). The prevalence by species must however be nuanced by the number of teeth available for each species: *Pl. mirallesi* and *Pr. douvillei* are equally abundant (340 and 354 teeth, respectively), while *B. brachypus* and *H. beonense* are scarce (46 and 92 teeth, respectively; Table 5).

Similarly, there are huge differences in the prevalence by tooth locus. Deciduous teeth are for instance less affected than permanent teeth with prevalences of 17.77 % (35/197) and 28.5 % (181/635), respectively, but permanent teeth are three times more numerous in the dataset (Table 5). This

difference is clear for *H. beonense* (milk: 1/30, 3.33 %; permanent: 11/62, 17.74 %) and *Pl. mirallesi* (milk: 12/76, 15.79 %; permanent: 86/264, 32.58 %), but less pronounced in proportions for *Pr. douvillei* (milk: 21/90, 23.33 %; permanent: 72/264, 27.27 %). For *B. brachypus* there is only one deciduous tooth – a D1 – and it presents hypoplasia (two LEHs). Multiple events of hypoplasia were twice as frequent on permanent teeth (68/635; 10.71 %) than on deciduous ones (11/197; 5.58 %). However, the ratio of multiplicity against the number of hypoplastic teeth is similar in deciduous (11/35, 31.43 %) and permanent (68/181; 37.57 %) teeth, around one third. Overall, upper teeth (100/446; 22.42 %) are also significantly less hypoplastic than lower teeth (116/386; 30.1 %). The difference between upper and lower teeth was similar for *Pr. douvillei* (lower: 46/148, 31.08 %; upper: 47/206, 22.82 %) and *Pl. mirallesi* (lower: 54/151, 35.76 %; upper: 44/189, 23.28 %), but inverted for *H. beonense*: upper teeth (8/44; 18.18 %) were more affected than lower ones (4/48; 8.33 %). For *B. brachypus*, very few upper teeth were available, making such comparison hazardous.

More precisely, d1/D1, d2/D2, and D3 were the least affected loci with less than 5% of the teeth presenting hypoplasia (even 0 % for d1 and D2). The most affected loci were p4 (48.15 %), m3 (46.81 %), and D4 (46.15 %). Hypoplasia preva-

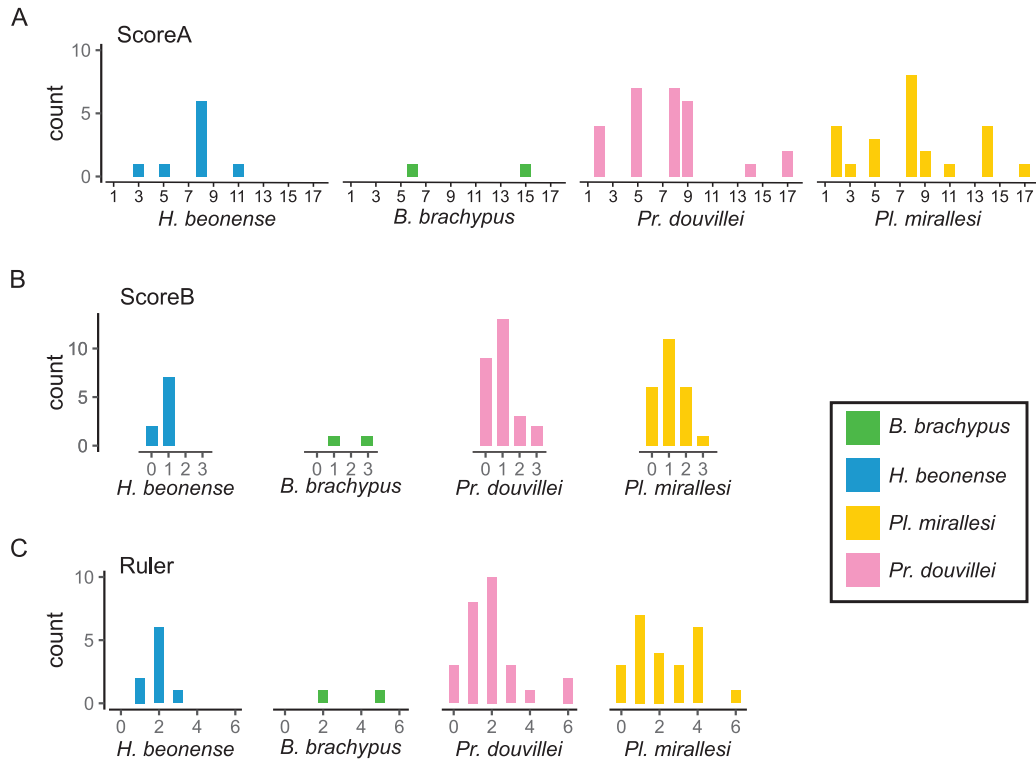


FIGURE 8. Barplots of mesowear scores on permanent teeth by method (ScoreA, ScoreB, Ruler) and by species. A- ScoreA: mesowear score based on Winkler and Kaiser (2011); B- ScoreB: mesowear score adapted from Fortelius and Solounias (2000); C- Ruler: mesowear score based on Mhlbachler et al. (2011). Only one tooth per specimen was considered.

TABLE 4. Minimum number of individuals (NI) at Béon 1 by species based on teeth. Corrected MNI based on dental eruption incompatibilities (e.g., no m3/M3 in wear associated with deciduous teeth but d1/D1) as proposed by Hullot and Antoine (2020).

	MNI	Tooth	Corrected	
			MNI	Teeth
<i>B. brachypus</i>	3	left m3	3	left m3
<i>Pr. douvillei</i>	7	right M2	12	right M2 + left D4
<i>Pl. mirallesi</i>	8	left p4	9	left P4 + left d2
<i>H. beonense</i>	3	left m1	3	left m1
Total	21		27	

TABLE 5. Hypoplasia (all type) and LEH prevalence by species and tooth type.

	Number of teeth		All types of hypoplasia						LEH only					
	Milk	Permanent	Milk		Permanent		All		Milk		Permanent		All	% of hypoplastic teeth
			N	%	N	%			N	%	N	%		
<i>B. brachypus</i>	1	45	1	100.00	12	26.67	28.26		1	100.00	5	11.11	13.04	46.15
<i>Pr. douvillei</i>	90	264	21	23.33	72	27.27	26.27		12	13.33	36	13.64	13.56	51.61
<i>Pl. mirallesi</i>	76	264	12	15.79	86	32.58	28.82		11	14.47	43	16.29	15.88	55.10
<i>H. beonense</i>	30	62	1	3.33	11	17.74	13.04		1	3.33	5	8.06	6.52	50.00
Total	197	635	35	17.77	181	28.50	4.21		25	12.69	89	14.02	13.70	52.78

lence is also high for d4 with 40 % of the teeth affected (Figure 9). Figure 9 highlights inter-specific differences in the most affected tooth loci. Few loci are affected for *B. brachypus* and *H. beonense*: for *B. brachypus* all lower permanent molars seem to be particularly affected, while for *H. beonense* M2 and m1/M1 seem more frequently hypoplastic. The other two species present a wider range of loci affected: for *Pr. douvillei* the most affected teeth are d4/D4, p4, and m3, while for *Pl. mirallesi* it is p4, m3, and p3 (Figure 9).

Concerning the type of hypoplasia, LEH (linear enamel hypoplasia) was the most frequent, affecting 17.55 % of the teeth (146/832) and corresponding to about the half of the defects encountered (Table 5). Pits were found on 7.45 % of the teeth (62/832) and aplasia on 2.4 % (20/832).

For all response variables (Hypo, Defect, Multiple, and Severity), model support increased (lower AIC) when intraspecific factors (e.g., tooth loci, wear stage) were included. The final models contained three to five factors. Besides species and specimens, tooth type was in the final models of the variables Hypo (i.e., presence or absence of hypoplasia) and Defect (i.e., type of defect), and defect (converted to factor) in that of variables Multiple and Severity. Details and comparison of all models can be seen in Appendix 5. Based on final GLMMs, we found significant differences (Appendix 5; p-values < 0.05) in species vulnerability to stress: *H. beonense* had suffered significantly less hypoplasia than *Pl. mirallesi* (set as reference for GLMMs), and it was more through single LEH than any other types of defects (e.g., aplasia, combination of defects). The prevalence and defect type did not differ significantly between *Pl. mirallesi* and the two teleoceratines. Similarly, mean severity was not different in all species. Upper teeth appeared less affected by hypoplasia but neither the defect types nor the multiplicity were different (p-values > 0.05). We also found significant differences depending on dental loci: p4/P4 had a higher prevalence of hypoplasia than d2/D2, d3/D3, p2/P2, and p3/P3. The defect types also differ between p4/P4 and d2/D2 or m3/M3 (p-values < 0.05). More surprisingly, we found that right teeth display significantly different hypoplastic patterns and that lightly worn teeth were presenting less hypoplasia. Eventually, our results suggest that pits were less severe and less frequently multiple than LEH, but this result can be imputed to how pits were recorded in the first place (i.e., several pits identified as a single hypoplasia event) and their aetiol-

ogy (less ameloblasts disrupted than for LEH or aplasia).

DISCUSSION

Impact of Sand Preparation on Microwear Signal

As suggested by Table 2 and Figures 5 to 7, the microwear signal is modified by sand blasting preparation. Before our study we had three hypotheses concerning the impact of sand blasting on the microwear signal observed: 1) no significant effect of sand blasting (null hypothesis), 2) generation of new features resulting in a different microwear signal, and 3) flattening of the surface and erasing of microwear features. Our results suggest a shift towards lower anisotropy and higher complexity values in all studied species, regardless of their diet. A decrease in anisotropy is likely to be explained by sand particles impacting randomly and erasing any eventual pre-existing preferential orientations, while an increase in complexity could be due to sand particles generating new features of various sizes and shapes primarily due to the wide range of angles of incidence while blasting sand. These results point towards our second hypothesis, with the apparition of a *de novo* microwear signal that may in turn bias microwear interpretation towards browsing or frugivory artificially. Sand blasting is a real issue for undertaking microwear studies *a posteriori*. Accordingly, not only this method should be used with care and parsimony for museum preparation of occlusal surfaces of fossil teeth, but it is also necessary to be particularly cautious when analyzing museum fossil specimens for microwear analyses.

Interest of GLMM

Generalized linear mixed-models are extremely powerful yet very complex tools in ecology. Their application and their interpretation require many precautions. Yet, the use of GLMMs is a good way to compare different methods and assess the impact of factors all at the same time. In our study, GLMMs revealed that the choice of the cusp for scoring mesowear was crucial in rhinocerotids as ScoreA and ScoreB yielded significantly different means depending on the cusp studied (sharpest cusp or paracone). This matter has already been emphasized in rhinocerotids (Taylor et al., 2013) as well as in equids (Taylor et al., 2016), probably due to the asymmetry of their teeth. This issue is apparently restricted to perissodactyls as no cusp differences are reported in rumi-

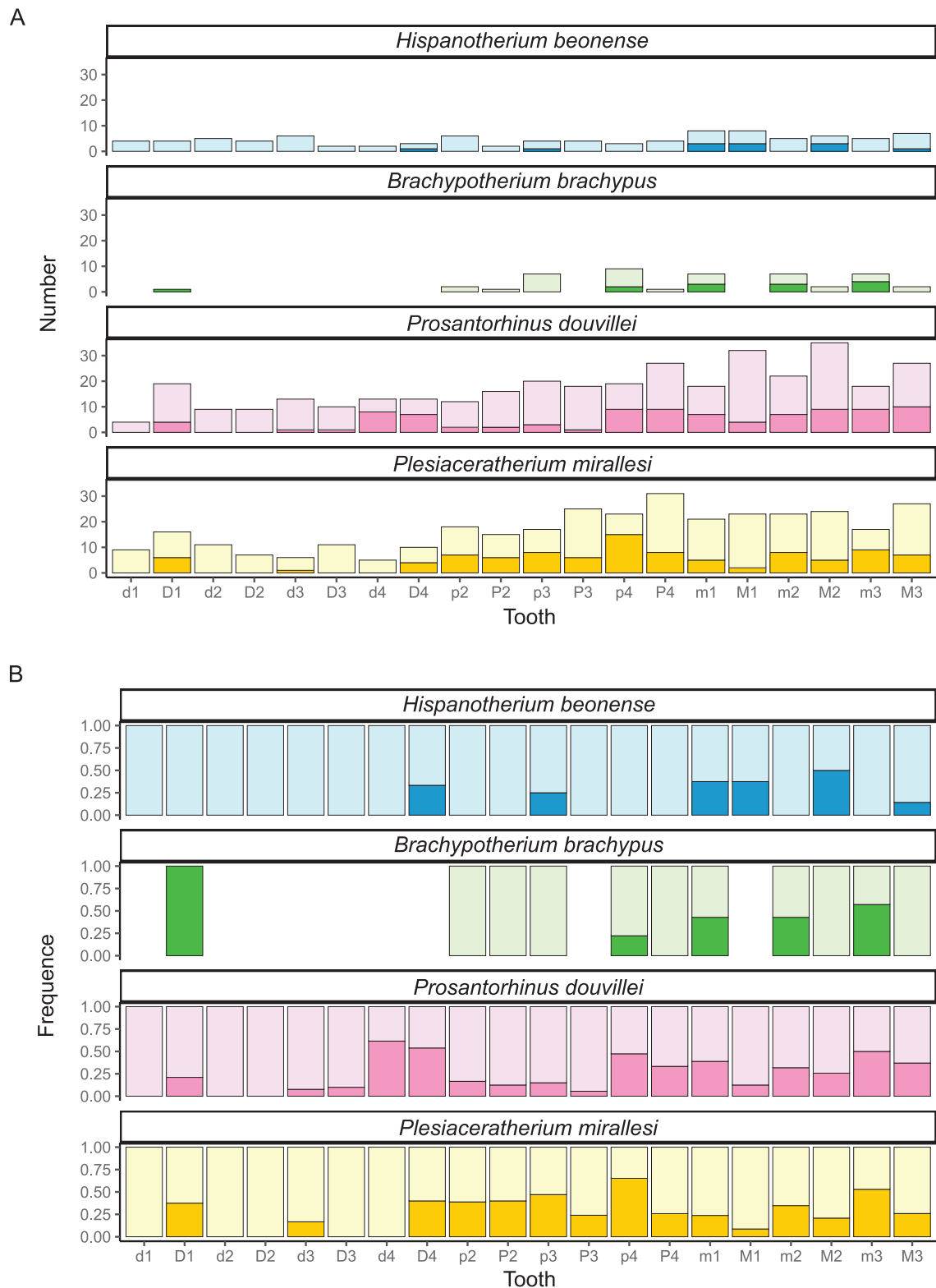


FIGURE 9. Prevalence of hypoplasia (all types) by species and tooth locus. A- Number of hypoplastic teeth (dark colors) compared to the number of healthy teeth (light colors). B- Frequency of hypoplastic teeth (dark colors) and healthy teeth (light colors). White stands for non-documented loci.

nant taxa, for instance (Fortelius and Solounias, 2000; Ackermans et al., 2018). GLMMs also highlighted differences due to ontogeny for mesowear. This result was expected, and it has been discussed before (Fortelius and Solounias, 2000; Rivals et al., 2007). However, the present study is the first investigation of mesowear on rhinocerotid deciduous teeth to our knowledge. These deciduous teeth were probably reacting differently to wear than would permanent teeth. Indeed, the enamel layer is thinner in deciduous teeth, which might lead to a distinct wear pattern (i.e., faster and/or more marked wear). Interestingly, the confounding factors – such as tooth loci/position in microwear, or left/right tooth (side) in hypoplasia or mesowear – were absent of most final models. This absence suggests that these factors might not play a crucial role in the variations observed, meaning that the most variation observed is due to actual interspecific differences.

Despite all these interests, our models are not perfect. First, we chose to study certain interactions and not others to simplify the models, which may have oriented our final results. Secondly, we based our model selection on AIC but several other criteria exist, with different sensitivities. For instance, when all models tested were ordered by Bayesian information criterion (BIC), the best candidate model differed for several response variables (Mesowear: Ruler, ScoreB; Microwear: Anisotropy, FTfv, HAsfc9, HAsfc81; Hypoplasia: Hypo, Defect). For FTfv the best model selected by BIC was based on different factors, but for the other variables, inter-individual variation (1|Specimen) was the only factor present. This could be due to the risk of over-fitting when using AIC and under-fitting with BIC. However, some biological parameters might also be responsible for this prevalence of inter-individual variation. For mesowear variables (ScoreB and Ruler), as no species differences were highlighted in the models, individual variability might be the greatest variation observed. As, for hypoplasia variables (Hypo and Defect), this could indicate that vulnerability is more at the individual level rather than at a species level.

Which Mesowear Method is best for Rhinocerotids?

In this study, we tested three different mesowear methods on our rhinocerotid sample. No critical difference was observed between the three methods (Figure 8; Table 3), however each of them relies on different scoring systems with their own advantages and disadvantages. For instance,

mesowear I (ScoreA) and extended mesowear I & II (ScoreB) are scored on the sharpest cusp, which is not fully applicable for rhinocerotid teeth as discussed in the previous section. This means that these methods may not be the most adapted to study mesowear for representatives of this clade or that they should be applied to the paracone, which in turn would impede comparison to other studies. Moreover, the very large scale (1-17) used in extended mesowear (ScoreB) is probably a source of intra- and inter-observer errors, instead of providing a finer discrimination between dietary preferences. Thus, we would recommend the use of Mesowear II – Mesowear Ruler (Ruler) to study mesowear on rhinocerotid teeth, especially since it has been specifically designed on equids (Mihlbachler et al. 2011), which are more closely related to rhinocerotids than artiodactyl ruminants.

Dietary Preferences

The dietary preferences of four associated rhinocerotid species from Béon 1 were investigated at two timescales: a short (days-to-weeks) scale with microwear, and a longer more cumulative, over-life scale, with mesowear. Previous work, based on gross dental and postcranial morphology, proposed a habitat partitioning for these rhinocerotids – swamps for the teleoceratines *B. brachypus* and *Pr. douvillei*, open woodland for the stem rhinocerotine *Pl. mirallesi*, and savannah-like open environments for the early-diverging elasmotheriine *H. beonense* (Antoine and Duranthon, 1997; Antoine et al., 2000; Bentaleb et al., 2006) – so we were expecting to find dietary differences.

Concerning mesowear, no significant differences were found between the four rhinocerotid species in any methods (ScoreA, ScoreB, and Ruler). The mesowear score profiles were different (Figure 8). In mesowear, leaf browsers have the lowest scores and grazers the highest, while mixed feeders and fruit browsers have intermediate overlapping scores. At Béon 1, all species had relatively low mean mesowear scores for all methods (ScoreA: 6.5-7; ScoreB: 0.7-1; Ruler: 1.2-2), excluding grazing as a dietary preference. Only *B. brachypus* had higher mean mesowear scores, which could suggest mixed-feeding, but as mesowear could be assessed on two teeth only, this remains an assumption. Accordingly, this species will not be further discussed. For the other associated species, ScoreB and Ruler suggest leaf browsing or mixed-feeding (Figure 8). Differences between deciduous and permanent teeth mesowear are highlighted with Ruler for all species

and for *Pl. mirallesi* only with ScoreB (Table 3). Permanent teeth display higher mesowear score than deciduous teeth, which might indicate a resource partitioning between the youngsters (weaned juveniles and subadults) and the adults. A less abrasive diet seems to be adopted by youngsters still possessing milk teeth, which might help to avoid competition with adults and to preserve deciduous teeth from excessive wear. ScoreA points toward a more abrasive diet than ScoreB and Ruler, but still coherent with browsing or mixed-feeding (Figure 8). Previous studies reported mesowear scores for extant rhinoceroses using different approaches (Fortelius and Solounias, 2000; Taylor et al., 2013; Muhlbachler et al., 2018; Rivals et al., 2020). The mean values reported by Rivals et al. (2020) for extant rhinos using Ruler are under 0.5 for all extant species but *C. simum* (4.67). Such values are very different from those observed at Béon 1 (between 1 and 2 for *Pl. mirallesi*, *Pr. douvillei*, and *H. beonense*), which could mean that modern rhinoceroses are not strict dietary analogues of the most abundant rhinocerotid species documented at Béon 1. However, the values reported in the study of Muhlbachler et al. (2018) were more contrasted, with mean mesowear scores around 2 for Asiatic browsing rhinoceroses (Javan + Sumatran rhinos), 3 for the Indian rhino, 4 for the black rhino, and 6 for the grazing white rhino. With these scores, Béon 1 rhinos appear closed to the Asiatic browsing rhinoceroses, known to live in close dense forest and relying mostly on leaf-browsing (Kahlke et al., 2011; Hullot et al., 2019).

Concerning microwear, the patterns observed can be summed up as in Table 6. Although there is a little overlap in microwear signals that could be due to sampling sites (see the discussion of Muhl-

bachler et al. 2016 for 2D microwear), we found subtle variations in the textures suggesting different feeding behaviors among the four rhinocerotid species. The relatively low values of Asfc for all species on both facets exclude the consumption of hard items, such as seeds, branches, or fruits with stone. This finding is consistent with the relatively open environment proposed from Montréal-du-Gers (oxbow lake surrounded by savannah and open woodlands; Duranthon et al., 1999; Hullot and Antoine, 2020). For the teleoceratines, the higher mean values of HAsfc for *B. brachypus* indicate that this species might have included more variety to its diet (Scott et al., 2006) compared to *Pr. douvillei*. The anisotropy and textural fill volume were lower in *B. brachypus* than in *Pr. douvillei*, which confirms that the investigated specimens of these closely related species had different dietary preferences. The microwear signature of *Pl. mirallesi* (Table 2) is suggestive of an abrasive and homogeneous diet especially on the shearing facet. This pattern is similar to what is observed in extant strict grazers such as the white rhinoceros or in strict selective folivores like the Sumatran rhinoceros (Scott, 2012; Hullot et al., 2019). Given the low-crowned dental pattern similar to that of *D. sumatrensis*, selective folivory seems to be more likely than strict grazing as a diet for *Pl. mirallesi*. Eventually, *H. beonense* exhibits moderate values of all parameters suggestive of a relative flexibility in the diet and probably of a mixed-feeding behavior (as inferred in early mesodont equids; Muhlbachler et al., 2011).

With all the previous results discussed in mind, we tried to provide feeding ecologies for Béon 1 rhinocerotids. Overall, microwear and mesowear concur, suggesting browsing or mixed-feeding diets. This congruence is especially veri-

TABLE 6. Summary of microwear pattern by species and DMTA parameter. epLsar: anisotropy; Asfc: complexity; FTfv: fine textural fill volume; HAsfc: heterogeneity of the complexity (9 and 81 cells).

	epLsar	Asfc	FTfv	HAsfc9	HAsfc81
<i>B. brachypus</i>	Low	Low-Moderate	Low-High	Moderate	Moderate
<i>Pr. douvillei</i>	Moderate	Low-Moderate	Moderate-High	Low-Moderate	Low-Moderate
<i>Pl. mirallesi</i>	Moderate-High	Low	Low-Moderate	Low	Low
<i>H. beonense</i>	Moderate	Low-Moderate	Moderate-High	Moderate	Moderate
<i>C. simum</i>	High	Low - Moderate	High	Low	Low
<i>D. sumatrensis</i>	High	Low	Low	Low	Low
<i>D. bicornis</i>	Low	Moderate - High	Moderate	Moderate-High	Moderate-High
<i>R. sondaicus</i>	Low	High	High	High	High
<i>R. unicornis</i>	Moderate	High	Moderate	Moderate	Moderate

fied for the two teleoceratines: *Pr. douvillei* appears to be a browser probably favoring leaves and *B. brachypus* has more variable feeding preferences that could suggest mixed-feeding. A browsing diet on low vegetation is compatible with *B. brachypus* morphology but dental wear suggests a mixed-feeding diet as proposed by Becker and Tissier (2020). The use of microwear and mesowear in tandem helped to reveal a subtle niche partitioning of the two teleoceratines through feeding ecology. This finding is consistent with the slight differences in the $\delta^{18}\text{O}$ of their teeth reported in a previous study (Bentaleb et al., 2006). Concerning *Pl. mirallesi* the microwear pattern pointed towards relatively abrasive and homogeneous food items, which is not retrieved in the mesowear score. This uncoupling, as well as low-crowned teeth, tends to exclude a grazing behavior and could suggest selective folivory similarly to what is observed in modern Sumatran rhinoceros instead. Eventually, *H. beonense* appears as a browser or a mixed-feeder with both proxies. This last result is quite surprising as Elasmotheriinae, to which *H. beonense* belongs, were in general hypothesized to have grazed on gramineous and abrasive plants from open environments (Inigo and Cerdeño, 1997; Cerdeño, 1998; Antoine, 2002; Becker and Tissier, 2020), due to their specialized morphology (i.e., high-crowned teeth with cement). However, hypsodonty does not necessarily imply obligate grazing and may instead allow for a wider range of dietary preferences, which is a key selective advantage (Janis, 1988; Semprebon and Rivals, 2007; Merceron et al., 2016).

Enamel Hypoplasia

The prevalence of hypoplasia affecting the rhinocerotids at Béon 1 is quite high, and comparable to what was observed in a Late Pleistocene woolly rhino sample (*Coelodonta antiquitatis*) from Fouvant-Sainte-Andoche, for which harsh cold conditions are reconstructed (France; Fourvel et al., 2015). There are, however, major differences depending on the species, tooth, or type of hypoplasia considered, as reported for tapirs and rhinoceroses found in association at Coc Muoi, a late Middle Pleistocene locality of Southeast Asia (Bacon et al., 2018). All prevalence values discussed in this paper must be nuanced in the light of species/tooth abundance and as some events might have been counted several times (defect on both opposing teeth from the same individual), because every isolated tooth was considered for counting. We found that deciduous teeth were less

affected than permanent teeth. This has been reported in several other studies focusing on various mammals (Dobney and Ervynck, 1998; Lukacs, 1999; Bacon et al., 2018; Towle and Irish, 2019; McGrath et al., 2021), and several factors might explain this difference. First, in rhinocerotids, deciduous teeth develop partially (d1/D1 and d4/D4) or totally (d2/D2 and d3/D3) *in utero*, which provides a relatively stable environment (Hitchins, 1978; Hillman-Smith et al., 1986; Böhmer et al., 2016). Such a stable environment might play a buffer role limiting stress, thus avoiding the formation of enamel hypoplasia on such dental loci. Also, deciduous teeth develop faster than their permanent counterparts (Dobney and Ervynck, 1998). This faster growth leaves less time to suffer from stress and record hypoplasia. Other factors were found to have an influence in the prevalence of hypoplasia observed but were difficult to explain. For instance, upper teeth present less enamel defects than their lower counterparts, whereas they have similar developmental timings, a pattern also observed in deciduous canines of great apes (Lukacs, 1999), but the underlying mechanisms of which has not been elucidated thus far. The higher frequency of hypoplastic right teeth with respect to their left counterparts might be due to a sampling bias as isolated teeth represent a great part of our sample.

Besides sampling biases, the marked interspecific differences in terms of prevalence could be due to specific sensibilities, distinct habitats, different preferred diet and flexibility, and/or contrasting life trajectories. Both species of teleoceratines exhibit similar levels of hypoplasia on their permanent teeth (*Pr. douvillei*: 27.27 %; *B. brachypus*: 26.67 %), out of which LEH is about as abundant in both species (*Pr. douvillei*: 13.64 %, *B. brachypus*: 11.11 %; Table 5). Such similarities are consistent with a comparable sensitivity to stress, perhaps linked to the phylogenetic closeness of these species. Moreover, as these two rhinocerotids were probably inhabiting the same habitat but presenting different feeding preferences (as highlighted here), we can hypothesize that environment, besides food resource, might be responsible for the observed stress. Indeed, previous works linked LEH with environmental stresses, such as drought or seasonal lack of food, in various mammals (Chollet and Teaford, 2010; Skinner and Pruett, 2012; Upex and Dobney, 2012). Periodic droughts are known to have happened at Béon 1 (Duranthon et al., 1999; Hullot and Antoine, 2020), and these events might have affected particularly the

teleoceratines that would have relied on water. Interestingly, the rhino with the least dietary flexibility had the highest hypoplasia prevalence (*Pl. mirallesi*; 28.82 %), whilst the most flexible (*H. beonense*, mixed feeder) had the lowest hypoplasia prevalence (13.04 %). The high-crowned teeth with cement of *H. beonense* might have allowed this species to consume a wider range of resources and avoid nutritional stress. On the contrary, the selective *Pl. mirallesi*, which probably relied on low quality food (fibrous and abrasive), would have been more prone to dietary stresses.

As stated in the results section, the most affected loci were p4 (48.15 %), m3 (46.81 %), D4 (46.15 %), and d4 (40 %). In *Teleoceras*, another Miocene rhinoceros, hypoplasias recorded on d4/D4 have been linked to birth related stresses, while that occurring on p4/P4 were related to cow-calf separation/weaning (Mead, 1999). These hypotheses are consistent with the mortality curves of Béon 1 rhinocerotids established by Hullot and Antoine (2020), that highlighted birth and weaning as vulnerability periods. At the Pleistocene site of Taubach (Germany), Bratlund (1999) suggests malnutrition or disease causing a fever to explain the nine hypoplasia defects detected on the permanent teeth (P2-M2) of *Stephanorhinus hemitoechus*. Concerning m3, it is the last tooth to develop and erupt. Hypoplasia on the last molars has been correlated with seasonality in sheep (Upex and Dobney, 2012) and fossil giraffe (Franz-Odendaal et al., 2004). The most affected loci are variable depending on the species (Figure 9). This suggests that the different species have distinct sensitivities to stress and are affected at various periods of their life. For *Pl. mirallesi* and *Pr. douvillei* we retrieved some vulnerability periods of their mortality curves (Hullot and Antoine, 2020): birth (d4/D4, m1/M1 and second and third premolars) and weaning (p4/P4). The last two molars are also frequently hypoplastic, suggesting that these rhinos experienced environmental stresses. Very few loci are affected for *B. brachypus* and *H. beonense* but they imply similar causes: stresses around birth (D4, m1/M1) and environmental stresses (m3, M2). It would be interesting to investigate the prevalence of hypoplasia in other taxa from Béon 1 to confirm the role of environment as a stressor and to compare sensitivity across more distant species, like what has been done for the Pliocene herbivores of Langebaanweg (South Africa; Franz-Odendaal et al., 2004).

CONCLUSIONS

In this study we reconstructed some ecological aspects of four rhinocerotid species from the late early Miocene locality of Béon 1 (Montréal-du-Gers, France). While mesowear did not discriminate between the species and suggested browsing habits, microwear (DMTA) revealed subtle dietary preferences: *Pl. mirallesi* and *Pr. douvillei* were browsers probably favoring leaves, while *B. brachypus* and *H. beonense* were mixed-feeders. The prevalence of hypoplasia was overall high (> 25 % of the teeth affected), although it was contrasted depending on species, tooth locus, and type of defect considered. This suggested a rather stressful environment, which is coherent with the periodical droughts that occurred at Béon 1. Interestingly, the rhinocerotid with the most flexible diet (*H. beonense*) was the least affected by hypoplasia. We also discussed three methodological points here: the impact of sand-preparation on microwear signatures (lower anisotropy and higher complexity), the interests and limits of Generalized Linear Mixed Models (powerful but complex tool), and the choice of mesowear approach for rhinocerotids (Mesowear II – Mesowear Ruler seems the most suitable).

ACKNOWLEDGEMENTS

This research has been partly funded by the foundation Yves Coppens (excavation at Béon 1) and a “Bourse de Mobilité Doctorale” granted by the Association Française des Femmes Diplômées des Universités.

We are indebted to F. Duranthon and P. Dalous (Muséum de Toulouse) who granted us the access to study the rhinocerotids from Béon 1. For the extant dataset we are grateful to S. Jiquel, B. Marandat, and A.-L. Charruault (University of Montpellier), D. Berthet and F. Vigouroux (Musée des Confluences de Lyon), F. Zachos and A. Bibl (Natuhistorisches Museum Wien), J. Lesur and V. Bouetel (Muséum National d'Histoire Naturelle, Paris), M. Lowe (University Museum of Zoology of Cambridge), E. Gilissen (Musée Royal d'Afrique Centrale, Tervuren), and L. Costeur (Natuhistorisches Museum Basel). We thank both reviewers, F. Rivals and M. Mhielbachler, for their constructive feedback that greatly improved an earlier version of the manuscript.

REFERENCES

- Ackermans, N.L. 2020. The history of mesowear: a review. *PeerJ*, 8. <https://doi.org/10.7717/peerj.8519>
- Ackermans, N.L., Winkler, D.E., Schulz-Kornas, E., Kaiser, T.M., Müller, D.W., Kircher, P.R., Hummel, J., Clauss, M., and Hatt, J.-M. 2018. Controlled feeding experiments with diets of different abrasiveness reveal slow development of mesowear signal in goats (*Capra aegagrus hircus*). *Journal of Experimental Biology*, 221:jeb186411. <https://doi.org/10.1242/jeb.186411>
- Ackermans, N.L., Martin, L.F., Codron, D., Hummel, J., Kircher, P.R., Richter, H., Kaiser, T.M., Clauss, M., and Hatt, J.-M. 2020. Mesowear represents a lifetime signal in sheep (*Ovis aries*) within a long-term feeding experiment. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 553:109793. <https://doi.org/10.1016/j.palaeo.2020.109793>
- Antoine, P.-O. 2002. Phylogénie et évolution des Elasmotheriina (Mammalia, Rhinocerotidae). *Mémoires du Muséum National d'Histoire naturelle*, 188:1-359.
- Antoine, P.-O. and Duranthon, F. 1997. Découverte de *Protaceratherium minutum* (Mammalia, Rhinocerotidae) dans le gisement Orléanien (MN 4) de Montréal-du-Gers (Gers). *Annales de Paléontologie (Vert.-Invert.)*, 83:201-213.
- Antoine, P.-O., Bulot, C., and Ginsburg, L. 2000. Les rhinocérotidés (Mammalia, Perissodactyla) de l'Orléanien des bassins de la Garonne et de la Loire (France): intérêt biostratigraphique. *Comptes Rendus de l'Académie des Sciences - Series IIA - Earth and Planetary Science*, 330:571-576. [https://doi.org/10.1016/S1251-8050\(00\)00174-9](https://doi.org/10.1016/S1251-8050(00)00174-9)
- Antoine, P.-O., Becker, D., Laurent, Y., and Duranthon, F. 2018. The Early Miocene teleoceratine *Prosantorhinus* aff. *douvillei* (Mammalia, Perissodactyla, Rhinocerotidae) from Béon 2, Southwestern France. *Revue de Paléobiologie*, 37:367-377.
- Arman, S.D., Prowse, T.A.A., Couzens, A.M.C., Ungar, P.S., and Prideaux, G.J. 2019. Incorporating intraspecific variation into dental microwear texture analysis. *Journal of the Royal Society Interface*, 16:20180957. <https://doi.org/10.1098/rsif.2018.0957>
- Auguie, B. 2017. gridExtra: Miscellaneous Functions for "Grid" Graphics. R package version 2.3. <https://CRAN.R-project.org/package=gridExtra>
- Bacon, A.-M., Antoine, P.-O., Huong, N.T.M., Westaway, K., Tuan, N.A., Düringer, P., Zhao, J., Ponche, J.-L., Dung, S.C., Nghia, T.H., Minh, T.T., Son, P.T., Boyon, M., Thuy, N.T.K., Blin, A., and Demeter, F. 2018. A rhinocerotid-dominated megafauna at the MIS6-5 transition: The late Middle Pleistocene Coc Muoi assemblage, Lang Son province, Vietnam. *Quaternary Science Reviews*, 186:123-141. <https://doi.org/10.1016/j.quascirev.2018.02.017>
- Bates, D., Mächler, M., Bolker, B., and Walker, S. 2015. Fitting linear mixed-effects models using lme4. *Journal of Statistical Software*, 67:1-48. <https://doi.org/10.18637/jss.v067.i01>
- Becker, D. and Tissier, J. 2020. Rhinocerotidae from the early middle Miocene locality Gračanica (Bugojno Basin, Bosnia-Herzegovina). *Palaeobiodiversity and Palaeoenvironments*, 100:395-412. <https://doi.org/10.1007/s12549-018-0352-1>
- Bentaleb, I., Langlois, C., Martin, C., Iacumin, P., Carré, M., Antoine, P.-O., Duranthon, F., Moussa, I., Jaeger, J.-J., and Barrett, N. 2006. Rhinocerotid tooth enamel $^{18}\text{O}/^{16}\text{O}$ variability between 23 and 12 Ma in southwestern France. *Comptes Rendus Geoscience*, 338:172-179. <https://doi.org/10.1016/j.crte.2005.11.007>
- Berlioz, É., Kostopoulos, D.S., Blondel, C., and Merceron, G. 2018. Feeding ecology of *Eucladoceros ctenoides* as a proxy to track regional environmental variations in Europe during the early Pleistocene. *Comptes Rendus Palevol*, 17:320-332. <https://doi.org/10.1016/j.crpv.2017.07.002>
- Böhmer, C., Heissig, K., and Rössner, G.E. 2016. Dental eruption series and replacement pattern in Miocene *Prosantorhinus* (Rhinocerotidae) as revealed by macroscopy and x-ray: Implications for ontogeny and mortality profile. *Journal of Mammalian Evolution*, 23:265-279. <https://doi.org/10.1007/s10914-015-9313-x>
- Bratlund, B. 1999. Taubach revisited. *Jahrbuch des Römisch-Germanischen Zentralmuseums Mainz*, 46:61-174. <https://doi.org/10.11588/jrgzm.1999.1.25776>
- Calandra, I., Göhlich, U.B., and Merceron, G. 2008. How could sympatric megaherbivores coexist? Example of niche partitioning within a proboscidean community from the Miocene of Europe. *Naturwissenschaften*, 95:831-838. <https://doi.org/10.1007/s00114-008-0391-y>

- Cerdeño, E. 1998. Diversity and evolutionary trends of the Family Rhinocerotidae (Perissodactyla). *Palaeogeography, Palaeoclimatology, Palaeoecology*, 141:13-34. [https://doi.org/10.1016/S0031-0182\(98\)00003-0](https://doi.org/10.1016/S0031-0182(98)00003-0)
- Chollet, M.B. and Teaford, M.F. 2010. Ecological stress and linear enamel hypoplasia in *Cebus*. *American Journal of Physical Anthropology*, 142:1-6. <https://doi.org/10.1002/ajpa.21182>
- Crouzel, F., Duranthon, F., and Ginsburg, L. 1988. Découverte d'un riche gisement à petits et grands mammifères d'âge Orléanien dans le département du Gers (France). *Comptes rendus de l'Académie des Sciences*, 307:101-104.
- Davis, M. and Pineda-Munoz, S. 2016. The temporal scale of diet and dietary proxies. *Ecology and Evolution*, 6:1883–1897. <https://doi.org/10.1002/ece3.2054>.
- Dobney, K. and Ervynck, A. 1998. A protocol for recording linear enamel hypoplasia on archaeological pig teeth. *International Journal of Osteoarchaeology*, 8:263-273. [https://doi.org/10.1002/\(SICI\)1099-1212\(199807/08\)8:4<263::AID-OA427>3.0.CO;2-P](https://doi.org/10.1002/(SICI)1099-1212(199807/08)8:4<263::AID-OA427>3.0.CO;2-P)
- Duranthon, F., Sola, S.M., Astibia, H., and Köhler, M. 1995. *Ampelomeryx ginsburgi* nov. gen., nov. sp. (Artiodactyla, Cervoidea) et la famille de Palaeomerycidae. *Comptes rendus de l'Académie des Sciences*, 321:339-346.
- Duranthon, F., Antoine, P.O., Bulot, C., and Capdeville, J.P. 1999. Le Miocène inférieur et moyen continental du bassin d'Aquitaine Livret-guide de l'excursion des Journées Crouzel (10 et 11 juillet 1999). *Bulletin de la Société d'histoire naturelle de Toulouse*, 135:79-91.
- Fédération Dentaire Internationale. 1982. An epidemiological index of development defects of dental enamel (DDE index). *International Dental Journal*, 42:411-426.
- Fortelius, M. and Solounias, N. 2000. Functional characterization of ungulate molars using the abrasion-attribution wear gradient: a new method for reconstructing paleodiets. *American Museum Novitates*, 2000:1-36.
- Fourvel, J.-B., Fosse, P., Fernandez, P., and Antoine, P.-O. 2015. Large mammals of Fouvent-Saint-Andoche (Haute-Saône, France): a glimpse into a Late Pleistocene hyena den. *Geodiversitas*, 37:237-266. <https://doi.org/10.5252/g2015n2a5>
- Franz-Odenaal, T., Chinsamy, A., and Lee-Thorp, J. 2004. High prevalence of enamel hypoplasia in an early Pliocene giraffid (*Sivatherium hendeyi*) from South Africa. *Journal of Vertebrate Paleontology*, 24:235-244. <https://doi.org/10.1671/19>
- Grine, F.E. 1986. Dental evidence for dietary differences in *Australopithecus* and *Paranthropus*: a quantitative analysis of permanent molar microwear. *Journal of Human Evolution*, 15:783-822.
- Guatelli-Steinberg, D. 2001. What can developmental defects of enamel reveal about physiological stress in nonhuman primates? *Evolutionary Anthropology: Issues, News, and Reviews*, 10:138-151. <https://doi.org/10.1002/evan.1027>
- Hillman-Smith, A.K.K., Owen-Smith, N.R., Anderson, J.L., Hall-Martin, A.J., and Selaladi, J.P. 1986. Age estimation of the white rhinoceros (*Ceratotherium simum*). *Journal of Zoology*, 210:355-377.
- Hitchins, P.M. 1978. Age determination of the black rhinoceros (*Diceros bicornis* Linn.) in Zululand. *South African Journal of Wildlife Research*, 8:71-80.
- Hoffman, J.M., Fraser, D., and Clementz, M.T. 2015. Controlled feeding trials with ungulates: a new application of in vivo dental molding to assess the abrasive factors of microwear. *The Journal of Experimental Biology*, 218:1538-1547. <https://doi.org/10.1242/jeb.118406>
- Hullot, M. and Antoine, P.-O. 2020. Mortality curves and population structures of late early Miocene Rhinocerotidae (Mammalia, Perissodactyla) remains from the Béon 1 locality of Montréal-du-Gers, France. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 558:109938. <https://doi.org/10.1016/j.palaeo.2020.109938>
- Hullot, M., Antoine, P.-O., Ballatore, M., and Merceron, G. 2019. Dental microwear textures and dietary preferences of extant rhinoceroses (Perissodactyla, Mammalia). *Mammal Research*, 64:397-409. <https://doi.org/10.1007/s13364-019-00427-4>
- Inigo, C. and Cerdeño, E. 1997. The *Hispanotherium matritense* (Rhinocerotidae) from Córcoles (Guadalajara, Spain): Its contribution to the systematics of the Miocene Iranotheriina. *Geobios*, 30:243-266. [https://doi.org/10.1016/S0016-6995\(97\)80232-X](https://doi.org/10.1016/S0016-6995(97)80232-X)
- Janis, C.M. 1988. An estimation of tooth volume and hypsodonty indices in ungulate mammals, and the correlation of these factors with dietary preferences. *Mémoires du Muséum National d'Histoire Naturelle*, 53:367-387.

- Jones, D.B. and DeSantis, L.R.G. 2017. Dietary ecology of ungulates from the La Brea tar pits in southern California: A multi-proxy approach. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 466:110-127. <https://doi.org/10.1016/j.palaeo.2016.11.019>
- Kahlke, R.-D., García, N., Kostopoulos, D.S., Lacombat, F., Lister, A.M., Mazza, P.P., Spassov, N., and Titov, V.V. 2011. Western Palaearctic palaeoenvironmental conditions during the Early and early Middle Pleistocene inferred from large mammal communities, and implications for hominin dispersal in Europe. *Quaternary Science Reviews*, 30:1368-1395. <https://doi.org/10.1016/j.quascirev.2010.07.020>
- Lukacs, J.R. 1999. Enamel hypoplasia in deciduous teeth of great apes: Do differences in defect prevalence imply differential levels of physiological stress? *American Journal of Physical Anthropology*, 110:351-363. [https://doi.org/10.1002/\(SICI\)1096-8644\(199911\)110:3<351::AID-AJPA7>3.0.CO;2-2](https://doi.org/10.1002/(SICI)1096-8644(199911)110:3<351::AID-AJPA7>3.0.CO;2-2)
- McGrath, K., Limmer, L.S., Lockey, A.-L., Guatelli-Steinberg, D., Reid, D.J., Witzel, C., Bocaegge, E., McFarlin, S.C., and El Zaatari, S. 2021. 3D enamel profilometry reveals faster growth but similar stress severity in Neanderthal versus *Homo sapiens* teeth. *Scientific Reports*, 11:522. <https://doi.org/10.1038/s41598-020-80148-w>
- Mead, A.J. 1999. Enamel hypoplasia in Miocene rhinoceroses (*Teleoceras*) from Nebraska: evidence of severe physiological stress. *Journal of Vertebrate Paleontology*, 19:391-397. <https://doi.org/10.1080/02724634.1999.10011150>
- Merceron, G., Ramdarshan, A., Blondel, C., Boissérie, J.-R., Brunetiere, N., Francisco, A., Gautier, D., Milhet, X., Novello, A., and Pret, D. 2016. Untangling the environmental from the dietary: dust does not matter. *Proceedings of the Royal Society B: Biological Sciences*, 283:20161032. <https://doi.org/10.1098/rspb.2016.1032>
- Merceron, G., Berlioz, E., Vonhof, H., Green, D., Garel, M., and Tütken, T. 2021. Tooth tales told by dental diet proxies: An alpine community of sympatric ruminants as a model to decipher the ecology of fossil fauna. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 562:110077. <https://doi.org/10.1016/j.palaeo.2020.110077>
- Mihlbachler, M.C., Rivals, F., Solounias, N., and Semprebon, G.M. 2011. Dietary change and evolution of horses in North America. *Science*, 331:1178-1181. <https://doi.org/10.1126/science.1196166>
- Mihlbachler, M.C., Campbell, D., Ayoub, M., Chen, C., and Ghani, I. 2016. Comparative dental microwear of ruminant and perissodactyl molars: Implications for paleodietary analysis of rare and extinct ungulate clades. *Paleobiology*, 42:98-116. <https://doi.org/10.1017/pab.2015.33>
- Mihlbachler, M.C., Campbell, D., Chen, C., Ayoub, M., and Kaur, P. 2018. Microwear–mesowear congruence and mortality bias in rhinoceros mass-death assemblages. *Paleobiology*, 44:131-154. <https://doi.org/10.1017/pab.2017.13>
- Niven, L.B., Egeland, C.P., and Todd, L.C. 2004. An inter-site comparison of enamel hypoplasia in bison: implications for paleoecology and modeling Late Plains Archaic subsistence. *Journal of Archaeological Science*, 31:1783-1794. <https://doi.org/10.1016/j.jas.2004.06.001>
- Orliac, M.J. 2006. *Eurolistriodon tenarezensis*, sp. nov., from Montréal-du-Gers (France): implications for the systematics of the European Listriodontinae (Suidae, Mammalia). *Journal of Vertebrate Paleontology*, 26:967-980. [https://doi.org/10.1671/0272-4634\(2006\)26\[967:ETSNFM\]2.0.CO;2](https://doi.org/10.1671/0272-4634(2006)26[967:ETSNFM]2.0.CO;2)
- Orliac, M.J., Antoine, P.-O., and Duranthon, F. 2006. The Suoidea (Mammalia, Artiodactyla), exclusive of Listriodontinae, from the early Miocene of Béon 1 (Montréal-du-Gers, SW France, MN4). *Geodiversitas*, 28:685-718.
- Owen-Smith, N.R. 1988. *Megaherbivores: The Influence of Very Large Body Size on Ecology*. Cambridge University Press.
- Rage, J.-C. and Bailón, S. 2005. Amphibians and squamate reptiles from the late early Miocene (MN 4) of Béon 1 (Montréal-du-Gers, southwestern France). *Geodiversitas*, 27:413-441.
- Ramdarshan, A., Blondel, C., Brunetiere, N., Francisco, A., Gautier, D., Surault, J., and Merceron, G. 2016. Seeds, browse, and tooth wear: a sheep perspective. *Ecology and Evolution*, 6:5559-5569. <https://doi.org/10.1002/ece3.2241>
- Rivals, F. and Semprebon, G.M. 2006. A comparison of the dietary habits of a large sample of the Pleistocene pronghorn *Stockoceros onusrosagris* from the Papago Springs Cave in Arizona to the modern *Antilocapra americana*. *Journal of Vertebrate Paleontology*, 26:495-500. [https://doi.org/10.1671/0272-4634\(2006\)26\[495:ACOTDH\]2.0.CO;2](https://doi.org/10.1671/0272-4634(2006)26[495:ACOTDH]2.0.CO;2)

- Rivals, F., Solounias, N., and Muhlbachler, M.C. 2007. Evidence for geographic variation in the diets of late Pleistocene and early Holocene *Bison* in North America, and differences from the diets of recent *Bison*. *Quaternary Research*, 68:338-346. <https://doi.org/10.1016/j.yqres.2007.07.012>.
- Rivals, F., Takatsuki, S., Albert, R.M., and Macià, L. 2014. Bamboo feeding and tooth wear of three sika deer (*Cervus nippon*) populations from northern Japan. *Journal of Mammalogy*, 95:1043-1053. <https://doi.org/10.1644/14-MAMM-A-097>
- Rivals, F., Prilepskaya, N.E., Belyaev, R.I., and Pervushov, E.M. 2020. Dramatic change in the diet of a late Pleistocene *Elasmotherium* population during its last days of life: Implications for its catastrophic mortality in the Saratov region of Russia. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 556:109898. <https://doi.org/10.1016/j.palaeo.2020.109898>
- Rothschild, B.M., Martin, L.D., Lev, G., Bercovier, H., Bar-Gal, G.K., Greenblatt, C., Donoghue, H., Spigelman, M., and Brittain, D. 2001. *Mycobacterium tuberculosis* complex DNA from an extinct bison dated 17,000 years before the present. *Clinical Infectious Diseases*, 33:305-311. <https://doi.org/10.1086/321886>
- Schulz, E., Fahlke, J.M., Merceron, G., and Kaiser, T.M. 2007. Feeding ecology of the Chalicotheriidae (Mammalia, Perissodactyla, Ancylopoda). Results from dental micro- and mesowear analyses. *Verhandlungen des Naturwissenschaftlichen Vereins zu Hamburg*, 43:5-31.
- Scott, J.R. 2012. Dental microwear texture analysis of extant African Bovidae. *Mammalia*, 76. <https://doi.org/10.1515/mammalia-2011-0083>
- Scott, R.S., Ungar, P.S., Bergstrom, T.S., Brown, C.A., Grine, F.E., Teaford, M.F., and Walker, A. 2005. Dental microwear texture analysis shows within-species diet variability in fossil hominins. *Nature*, 436:693-695. <https://doi.org/10.1038/nature03822>
- Scott, R.S., Ungar, P.S., Bergstrom, T.S., Brown, C.A., Childs, B.E., Teaford, M.F., and Walker, A. 2006. Dental microwear texture analysis: technical considerations. *Journal of Human Evolution*, 51:339-349. <https://doi.org/10.1016/j.jhevol.2006.04.006>
- Semprebon, G.M. and Rivals, F. 2007. Was grass more prevalent in the pronghorn past? An assessment of the dietary adaptations of Miocene to Recent Antilocapridae (Mammalia: Artiodactyla). *Palaeogeography, Palaeoclimatology, Palaeoecology*, 253:332-347. <https://doi.org/10.1016/j.palaeo.2007.06.006>
- Skinner, M.F. and Pruett, J.D. 2012. Reconstruction of periodicity of repetitive linear enamel hypoplasia from perikymata counts on imbricational enamel among dry-adapted chimpanzees (*Pan troglodytes verus*) from Fongoli, Senegal. *American Journal of Physical Anthropology*, 149:468-482. <https://doi.org/10.1002/ajpa.22145>
- Small, B.W. and Murray, J.J. 1978. Enamel opacities: prevalence, classifications and aetiological considerations. *Journal of Dentistry*, 6:33-42. [https://doi.org/10.1016/0300-5712\(78\)90004-0](https://doi.org/10.1016/0300-5712(78)90004-0)
- Solounias, N. and Semprebon, G. 2002. Advances in the reconstruction of ungulate ecomorphology with application to early fossil equids. *American Museum Novitates*, 3366:1-49. [https://doi.org/10.1206/0003-0082\(2002\)366<0001:AITROU>2.0.CO;2](https://doi.org/10.1206/0003-0082(2002)366<0001:AITROU>2.0.CO;2)
- Solounias, N., Tariq, M., Hou, S., Danowitz, M., and Harrison, M. 2014. A new method of tooth mesowear and a test of it on domestic goats. *Annales Zoologici Fennici*, 51:111-118.
- Suckling, G., Elliott, D.C., and Thurley, D.C. 1986. The macroscopic appearance and associated histological changes in the enamel organ of hypoplastic lesions of sheep incisor teeth resulting from induced parasitism. *Archives of Oral Biology*, 31:427-439. [https://doi.org/10.1016/0003-9969\(86\)90016-6](https://doi.org/10.1016/0003-9969(86)90016-6)
- Taylor, L.A., Kaiser, T.M., Schwitzer, C., Müller, D.W.H., Codron, D., Clauss, M., and Schulz, E. 2013. Detecting inter-cusp and inter-tooth wear patterns in Rhinocerotids. *PLOS ONE*, 8:e80921. <https://doi.org/10.1371/journal.pone.0080921>
- Taylor, L.A., Müller, D.W., Schwitzer, C., Kaiser, T.M., Castell, J.C., Clauss, M., and Schulz-Kornas, E. 2016. Comparative analyses of tooth wear in free-ranging and captive wild equids. *Equine Veterinary Journal*, 48:240-245. <https://doi.org/10.1111/evj.12408>
- Teaford, M.F. 1991. Dental microwear: what can it tell us about diet and dental function, p. 341-356. In Kelley, M. and Larsen, C. (eds.), *Advances in Dental Anthropology*. Wiley-Liss, New York.
- Teaford, M.F. and Oyen, O.J. 1989. In vivo and in vitro turnover in dental microwear. *American Journal of Physical Anthropology*, 80:447-460. <https://doi.org/10.1002/ajpa.1330800405>

- Towle, I. and Irish, J.D. 2019. A probable genetic origin for pitting enamel hypoplasia on the molars of *Paranthropus robustus*. *Journal of Human Evolution*, 129:54-61.
<https://doi.org/10.1016/j.jhevol.2019.01.002>
- Ungar, P.S., Brown, C.A., Bergstrom, T.S., and Walker, A. 2003. Quantification of dental microwear by tandem scanning confocal microscopy and scale-sensitive fractal analyses. *Scanning*, 25:185-193. <https://doi.org/10.1002/sca.4950250405>
- Upex, B. and Dobney, K. 2012. Dental enamel hypoplasia as indicators of seasonal environmental and physiological impacts in modern sheep populations: a model for interpreting the zooarchaeological record. *Journal of Zoology*, 287:259-268.
<https://doi.org/10.1111/j.1469-7998.2012.00912.x>
- Venables, W.N. and Ripley, B.D. 2002. *Modern Applied Statistics with S*. Springer, New York.
- Wickham, H. 2016. *ggplot2: Elegant Graphics for Data Analysis*. Springer-Verlag, New York.
- Wilke, C.O. 2020. cowplot: Streamlined Plot Theme and Plot Annotations for “ggplot2”. R package version 1.1.0. <https://CRAN.R-project.org/package=cowplot>
- Winkler, D.E. and Kaiser, T.M. 2011. A case study of seasonal, sexual and ontogenetic divergence in the feeding behaviour of the moose (*Alces alces* LINNÉ, 1758). *Verhandlungen des naturwissenschaftlichen Vereins zu Hamburg*, 46:331-348.
- Winkler, D.E., Schulz-Kornas, E., Kaiser, T.M., Codron, D., Leichliter, J., Hummel, J., Martin, L.F., Clauss, M., and Tütken, T. 2020. The turnover of dental microwear texture: Testing the “last supper” effect in small mammals in a controlled feeding experiment. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 557:109930.
<https://doi.org/10.1016/j.palaeo.2020.109930>

APPENDICES

All appendices are supplied as a zipped file at <https://palaeo-electronica.org/content/2021/3433-beon-1-rhinocerotids-ecology>.

APPENDIX 1.

Example of GLMM R Script applied to ScoreA mesowear (based on Arman et al., 2019).

APPENDIX 2.

Summary of GLMMs results for DMTA, mesowear, and hypoplasia variables. Best candidate model is tested, corrected for over-dispersion if needed, and significant effects are precised.

APPENDIX 3.

Details on specimens studied (RData) and the GLMMs tested for all DMTA variables. Selection of the best candidate models is proposed by AIC and BIC (lowest score).

APPENDIX 4.

Details on specimens studied (RData) and the GLMMs tested for all mesowear variables. Selection of the best candidate models is proposed by AIC and BIC (lowest score).

APPENDIX 5.

EDetails on specimens studied (RData) and the GLMMs tested for all Hypoplasia variables. Selection of the best candidate models is proposed by AIC and BIC (lowest score).