



Zurich Open Repository and Archive University of Zurich Main Library Strickhofstrasse 39 CH-8057 Zurich www.zora.uzh.ch

Year: 2013

Dietary abrasiveness is associated with variability of microwear and dental surface texture in rabbits

Schulz, Ellen; Piotrowski, Vanessa; Clauss, Marcus; Mau, Marcus; Merceron, Gildas; Kaiser, Thomas M

Abstract: Dental microwear and 3D surface texture analyses are useful in reconstructing herbivore diets, with scratches usually interpreted as indicators of grass dominated diets and pits as indicators of browse. We conducted feeding experiments with four groups of rabbits (Oryctolagus cuniculus) each fed a different uniform, pelleted diet (lucerne, lucerne oats, grass oats, grass). The lowest silica content was measured in the lucerne and the highest in the grass diet. After 25 weeks of exposure to the diets, dental castings were made of the rabbit's lower molars. Occlusal surfaces were then investigated using dental microwear and 3D areal surface texture analysis. In terms of traditional microwear, we found our hypothesis supported, as the grass group showed a high proportion of (long) "scratches" and the lucerne group a high proportion of "pits". Regardless of the uniform diets, variability of microwear and surface textures was higher when silica content was low. A high variability in microwear and texture analysis thus need not represent dietary diversity, but can also be related to a uniform, low-abrasion diet. The uniformity or variability of microwear/texture analysis results thus might represent varying degrees of abrasion and attrition rather than a variety of diet items per se.

DOI: 10.1371/journal.pone.0056167

Posted at the Zurich Open Repository and Archive, University of Zurich ZORA URL: http://doi.org/10.5167/uzh-73564 Published Version



Originally published at: Schulz, Ellen; Piotrowski, Vanessa; Clauss, Marcus; Mau, Marcus; Merceron, Gildas; Kaiser, Thomas M (2013). Dietary abrasiveness is associated with variability of microwear and dental surface texture in rabbits. PLoS ONE, 8(2):e56167. DOI: 10.1371/journal.pone.0056167

Dietary Abrasiveness Is Associated with Variability of Microwear and Dental Surface Texture in Rabbits

Ellen Schulz¹*, Vanessa Piotrowski¹, Marcus Clauss², Marcus Mau³, Gildas Merceron⁴, Thomas M. Kaiser¹

1 Biocenter Grindel and Zoological Museum, University of Hamburg, Hamburg, Germany, 2 Clinic for Zoo Animals, Exotic Pets and Wildlife, University of Zurich, Zurich, Switzerland, 3 Dental Institute, King's College London, Guy's Hospital, London, United Kingdom, 4 Institut de Paléoprimatologie, Paléontologie Humaine, Evolution et Paléoenvironnements, Unité Mixte de Recherche Centre National de la Recherche Scientifique 7262, Université de Poitiers, Poitiers, France

Abstract

Dental microwear and 3D surface texture analyses are useful in reconstructing herbivore diets, with scratches usually interpreted as indicators of grass dominated diets and pits as indicators of browse. We conducted feeding experiments with four groups of rabbits (*Oryctolagus cuniculus*) each fed a different uniform, pelleted diet (lucerne, lucerne & oats, grass & oats, grass). The lowest silica content was measured in the lucerne and the highest in the grass diet. After 25 weeks of exposure to the diets, dental castings were made of the rabbit's lower molars. Occlusal surfaces were then investigated using dental microwear and 3D areal surface texture analysis. In terms of traditional microwear, we found our hypothesis supported, as the grass group showed a high proportion of (long) "scratches" and the lucerne group a high proportion of "pits". Regardless of the uniform diets, variability of microwear and surface textures was higher when silica content was low. A high variability in microwear and texture analysis thus need not represent dietary diversity, but can also be related to a uniform, low-abrasion diet. The uniformity or variability of microwear/texture analysis results thus might represent varying degrees of abrasion and attrition rather than a variety of diet items per se.

Citation: Schulz E, Piotrowski V, Clauss M, Mau M, Merceron G, et al. (2013) Dietary Abrasiveness Is Associated with Variability of Microwear and Dental Surface Texture in Rabbits. PLoS ONE 8(2): e56167. doi:10.1371/journal.pone.0056167

Editor: Alistair Robert Evans, Monash University, Australia

Received July 25, 2012; Accepted January 8, 2013; Published February 6, 2013

Copyright: © 2013 Schulz et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This work was funded by DFG KA 1525/8-1 and a Humboldt research fellowship for postdoctoral researchers (to GM). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: ellen.schulz@uni-hamburg.de

Introduction

Plant phytoliths (opaline silica and plant silica) are considered defenses against herbivory because they decrease the digestibility of plant components and abrade mammalian teeth due to the greater hardness of phytoliths compared to enamel [1,2], although the question whether they are actually harder than enamel has not been resolved conclusively to date [3,4]. Herbivory by voles, for example, induced silica incorporation in grasses [5] and reduced growth rates and digestion efficiency of the voles [6]. Field studies indicate that ingested soil abrades teeth in ungulates [7,8]. Therefore, the importance of external abrasive particles such as soil, dust, sand, and grit [9] as causative factors for tooth abrasion in ungulates has received significant attention. Phytoliths might have a greater role in causing variation in tooth wear [8]. But until now, apart from a feeding experiment of Baker et al. [1] testing for the wear of sheeps' incisors due to soil ingestion, controlled feeding experiments testing the role of phytoliths on tooth wear, and in particular on the microwear patterns of molar surfaces, are still missing.

In most mammalian groups it is assumed that the consumption of grass, which is rich in phytoliths, causes a two-dimensional microwear pattern of mainly scratches, while ingestion of browse, and hard items like seeds, produces pits [10–12]. These patterns were also observed in rodents, with typical grazers presenting an abundance of scratches, while insectivores and frugivores display more pits [13–16]. Since the traditional dental microwear analysis is limited to two dimensions, surface texture analysis, a more quantitative threedimensional (3D) method, has become established for dental dietary reconstruction [17–20]. Most such studies are based on museum specimens and information on the natural diet of a species is usually taken from the literature. Comprehensiveness, accommodation of effects of seasonality and habitat conditions covered, as well as methods applied result in inconsistency of data provided. Only a few reports are based on long term observation and provide data with strong statistical evidence. For most species, it is nevertheless possible to assign a general dietary strategy that allows understanding occlusal surface features that are related to this dietary strategy. However, we do not know how much variability in diet is reflected by occlusal surface textures, nor the exact mechanism of texture formation.

This gap of knowledge in relating textures to actual diets led us to conduct feeding experiments with controlled diets of known silica content, and relate them to occlusal parameters. The study thus aims at quantifying surface textures and to relate them to the proportions of abrasives (silica) in the diet. This should enhance our understanding of the equilibrium between abrasion and attrition (e.g. [21]) in mammal post canine teeth. Abrasion is due to food-tooth or particle-tooth contact while attrition is due to (most closely) tooth-tooth contact [22,23]. Teaford and Walker [24] showed that tooth-tooth wear produces featureless surface in stillborn guinea pigs. We would not expect to see the extreme featureless surface in the rabbits. Since Rensberger [25] recognised that food is involved in most or perhaps all natural wear and Fortelius [26] concluded that most dental wear resulted from abrasion, we used the domestic rabbits as a model species to measure high abrasion- as well as low abrasion-induced textures.

If theoretical attritional tooth-tooth contacts happen, we assume that surface structures like spiky summits would be moved against each other and it is likely that plateau-like surface structures will result, which can be measured. The fact that the ever-growing molars of the rabbits have high wear rates as compared to ungulates and primates [27] reduces the risk of cumulative wear signatures. The ISO/FDIS parameters (ISO/FDIS25178, [28]) applied to texture models allow quantification of aspects of the basic geometry of surface textures and comprehensively indicate their biomechanical properties in ungulates [19] and primates [17]. We expect our results to contribute to the better understanding of surface textures of lagomorphs and occlusal surface functionality of other species with hypselodont check teeth. Furthermore, we expect new insights into biomechanical constraints of the general process of tooth wear in mammals.

As compared to high silica diets, we assume that low-silica diets have a lower probability to induce surface lesions. Also, from microwear experience we expect point lesions ("pits") to prevail over linear lesions ("scratches"). Overall, random effects should increase in surface textures. We thus test the following hypotheses: A higher proportion of silica particles in the grass meal feed causes a scratch-dominated pattern of comparatively low variability; a lower proportion of silica particles in the lucerne feed causes a pitdominated and more variable surface texture.

Materials and Methods

Thirty-two New Zealand white rabbits were kept in 4 groups receiving exclusively 4 different pelleted compound feeds consisting of grass meal (G, n = 7), grass meal with crushed oats (GO, n = 6), lucerne with crushed oats (LO, n = 7) and lucerne (L=6) [17,19,29]. Water was available ad libitum. Pellets are different from a natural diet in terms of physical properties and lower water content than fresh plant material. This might have an influence on the wear characteristics, but to date, no information on the influence of feeding the same diet fresh (as forage in the wild), dried (as hay) or pelleted is available. Nevertheless, pelleted food offers the advantage of large batches of a uniform feed of consistent composition, where the influence of selective feeding on the part of the animal is excluded. Oats were used as an additive to create diets of intermediate silica content.

Coprophagy was not prevented, in order to allow wear to develop as would be representative for free-ranging animals. Gidenne and Lebas [30] observed that rabbits just swallow the soft pellets (cecotrophs) without chewing. This observation is supported by the frequent observation that cecotrophs are found in the stomach of slaughtered rabbits intact, i.e. without indication of mechanical disruption (an observation also made at the end of this experiment; M. Clauss, pers. obs., see Fig. S1 in File S1). We therefore do not expect coprophagy in rabbits to influence dental surface textures. New Zealand white rabbits (Leporidae, Lagomorpha) are a breed of the common rabbit *Oryctolagus cuniculus*. They are essentially intermediate feeders in the wild [31]. After 25 weeks animals were sacrificed, skulls were macerated, and moulds were made from the lingual part of the mesial enamel ridge of the lower first molars.

Diets were analyzed for silica concentration using a dry ash method [32]. Dental microwear analysis [20] was conducted following the protocol by Merceron et al. [33] as adapted to small mammal teeth by Gomes Rodrigues et al. [13]. A stereomicroscope (Leica MZ 16) with a spot CCD camera (Leica DC 300) at $\times 100$ magnification was used. A $300 \times 300 \ \mu m$ square was outlined on the centre of the dental facet. Microwear features \mathcal{N}_{p} (number of pits), $\mathcal{N}_{p}10$ (number of pits >10 μ m), $\mathcal{N}_{p}5$ (number of pits >5 μ m), \mathcal{N}_{s} (number of scratches), $\mathcal{N}ws5$ (number of scratches wider than 5 μ m), $\mathcal{N}ws10$ (number of scratches wider than 10 μ m), L_{s} (length of scratches) were counted by Vanessa Piotrowski (one observer only) using Optimas 6.2 software (Media Cybernetics, Rockville, U. S.).

As a second method of dental wear analysis at micro scale the 3D areal surface texture analysis was conducted using the confocal disc-scanning system µsurf custom (Nanofocus AG, Germany) according to Schulz et al. [19]. Threshold of recorded points was set to 80% as compared to a threshold of 90% in Schulz et al. [19]. For both methods, the measuring area is the lingual enamel facet of the primary shearing blade of the first lower molar. Before 3D surface texture parameters can be applied to surface data, however, filtering operators should be employed after ISO/FDIS 25178 [28]. The default operator is the set of S-Filters. As a default, the areal Gaussian filter (one of the S-Filters) is applied, which excludes the smallest scale elements from the surface resulting in the primary surface. In order to suppress form alterations (e.g., the curvature of a cylinder), the F operator is applied, which results in the S-F surface [28]. Subsequently, the L-Filter removes the low frequency alterations. The final product is the S-L surface. In accordance with Schulz et al. [19], Calandra et al. [17] and Winkler et al. [34] we apply the 3D ISO/FDIS 25178 texture parameters on the S-F surface. In order to determine whether and in what way form and waviness of the tooth enamel influences our results, we additionally apply the texture parameters on the primary as well as the S-L surface.

The 3D ISO/FDIS 25178 texture parameters (Table 1) employed to quantify texture were originally developed to classify 3D areal surface textures [28] to get an understanding of how the topography was influenced by industrial manufacturing processes and how the topography influences its function [35]. In ungulates, the ISO/FDIS 25178-2 parameters Sq (height amplitude of the surface), Sal (autocorrelation length), Shv (closed hills volume), Spd (density of peaks) and Std (texture direction) have been demonstrated to be indicative of dental functional traits [19]. Purnell et al. [36] demonstrated that the ISO/FDIS 25178-2 parameters Sa (mean surface roughness), Sk (roughness depth of the core), Spk (roughness depth of the peaks), Vmp (material volume of the peaks), Vmc (material volume of the core), and Vvc (void volume of the core) reflected diet in cichlid fishes. In primates, the ISO/FDIS 25178-2 parameters Sq (height amplitude of the surface), S5v (depth of the valleys), Vm (material volume), Spd (density of peaks), Sha (closed hill area), and closed dale area (Sda) allows inference on functional interaction of food and tooth enamel during mastication [17]. Generalised schematic models of the surface textures with high and low values of closed dale area (Sda) or height amplitude of the surface (Sp) are given in Figure 1.

Statistical analyses were performed using the software R 2.12.1 [37]. The Welch-Yuen heteroscedastic omnibus test (WY [38,39]) was coupled with a heteroscedastic pair-wise comparison test (analog to Dunnett's T3 test, PW [40]) to detect differences between trimmed means (15% trimming). Additionally the heteroscedastic rank-based test after Cliff (CM [40]) was applied. Only texture patterns found significant ($p \le 0.05$) in both robust approaches were described and discussed. Discriminant analysis was applied in SYSTAT 12 (SYSTAT Software, Inc., Chicago, U.S.). The automatic stepping forward algorithm was employed using the values $F_{to enter} = 1$ and $F_{to remove} = 0.9$. Group centroids were calculated with confidence intervals of 90%.



Figure 1. Schematic models. Schematic models of hypothetical surface textures indicating the parameter value of the ISO/FDIS 25178 parameter closed dale area (*Sda*) or maximum peak height (*Sp*) having high (left) or low values (right). doi:10.1371/journal.pone.0056167.g001

Results

Silica contents differed significantly between all groups (p < 0.001; Fig. 2A, Table 2), ranging from 11.96 mg/g dry matter in the grass to 0.10 mg/g dry matter in the lucerne group. Group G had the lowest number of pits (Np) and the longest scratches (Ls)

(Fig. 2B–C, Table 2 and Table S1, S2, S5A and S6A in File S1). The reverse was found for Group L. The variation in number of pits (Fig. 2B) and length of scratches (Fig. 2C) indicated by the interquartile range of the box plots was lower when dietary silica content was higher (L>LO>GO>G). The differences in the

Table 1. Description of the ISO/FDIS 25178 parameters.

Parameter	Description (condition)	Unit
\$10z	ten-point height	μm
S5p	five-point peak height	μm
S5v	five-point valley height	μm
Sa	arithmetic mean height or mean surface roughness	μm
Sal	auto-correlation length (s = 0.2)	μm
Sda	closed dale area	μ m²
Sdq	root mean square gradient	no unit
Sdr	developed interfacial area ratio	%
Sdv	closed dale volume	μ m³
Sha	closed hill area	μm²
Shv	closed hill volume	μm³
Sku	kurtosis of the height distribution	no unit
Smc	inverse areal material ratio (p = 10%)	μm
Smr	areal material ratio, bearing area ratio at a given height (c = 1 μm under the highest peak)	%
Sp	maximum peak height, height between the highest peak and the mean plane	μ m
Spc	arithmetic mean peak curvature	1/µm
Spd	density of peaks	1/µm²
Sq	standard deviation of the height distribution, or RMS surface roughness	μm
Ssk	skewness of the height distribution	no unit
Std	texture direction	0
Str	texture aspect ratio (s = 0.2)	no unit
Sv	maximum pit height, depth between the mean plane and the deepest valley	μm
Sxp	peak extreme height difference in height between p% and q% (p = 50%, q = 97.5%)	μ m
Sz	maximum height, height between the highest peak and the deepest valley	μ m
Vm	material volume at a given material ratio (p=10%)	μm³/μm²
Vmc	material volume of the core at given material ratio ($p = 10\%$, $q = 80\%$)	μm³/μm²
Vmp	material volume of peaks (p = 10%)	$\mu m^3/\mu m^2$
Vv	void volume at a given material ratio (p = 10%)	$\mu m^3/\mu m^2$
Vvc	void volume of the core ($p = 10\%$, $q = 80\%$)	μm³/μm²
Vvv	void volume of the valley at a given material ratio (p=80%)	μ m³/μm²

Description and units of the applied parameters are indicated according to ISO/FDIS 25178 analysis. The most effective parameters that are found to discriminate rabbit diets are set in bold.

doi:10.1371/journal.pone.0056167.t001

means of Np values between grass-fed (Np_{grass-fed lagomorphs} = 21.7) and lucerne-fed lagomorphs (Np_{lucerne-fed lagomorphs} = 45.9) are similar to the ones between grazing and browsing ungulates (cf. mean values of Np_{grazing ungulates} = 15, Np_{browsing ungulates} = 31, [41]). However, the differences in Ls values between grass-fed (Ls_{grass-fed lagomorphs} = 59.1 µm) and lucerne-fed lagomorphs (Ls_{kenne-fed lagomorphs} = 38.9 µm) are slightly smaller than the ones reported in ungulates (Ls_{grazing ungulates} = 170.4 µm, Ls_{browsing ungulates} = 139.6 µm, [41]).

Group G had large areas of surface lesions (high *Sda*) on the primary surface and lower and less variable mean peak heights values on the S-L surface (low *Sp*, Fig. 2D–E, Table 2 and Tables S1, S3, S4, S5B, S6B in File S1). Group L was characterised by low *Sda* on the primary surface and higher variability in *Sp* on the S-L surface (Fig. 2D–E). Groups GO, LO and L group had similar mean *Sda*, indicating that both lucerne and the inclusion of oats



Figure 2. Box plots and discriminant analysis plot. Box plots indicating (a) the silica concentration (SC, %) in animal feeds, the microwear parameters (b) number of pits (*Np*), (c) length of scratches (*Ls*), the ISO/FDIS 25178-2 parameters: (d) closed dale area (*Sda* on primary surface, μ m²), (e) maximum peak height (*Sp* on S-L surface, μ m) within the feeding groups (G=grass meal, GO=grass meal with crushed oats, LO=lucerne with crushed oats, and L=lucerne), *=p≤0.05. Box plots showing the median (middle line), the interquartile range (IQR, box) and the minimum/maximum values 1.5×IQR (whiskers), extreme values are excluded. Discriminant analysis (f) using microwear (*Np*, *Ls*) and texture parameters (*Sda* on primary surface, *Sp* on S-L surface) indicating significant group differences (Table 2). The canonical discriminant function coefficients are *Np*_{score1/2}=0.038/0.015, *Ls*_{score 1/2}=-0.128/ -0.014, *Sda*_{score1/2}=-0.532/0.015, *Sp*_{score 1/2}=-0.076/-0.226.

Table 2. WY-test statistics.

	Parameter	Ft	p	nu1	Nu2
silica	SC	386.714	< 0.001	3	61.145
microwear (A)	Np	8.894	0.004	3	9.166
	Ls	6.571	0.011	3	9.547
texture (A)	Sda	4.359	0.037	3	8.971
	Sdv	5.971	0.016	3	9.128
texture (B)	Sz	2.972	0.095	3	8.213
texture (C)	S5v	5.474	0.021	3	8.868
	Sal	4.445	0.035	3	9.043
	Sp	4.192	0.043	3	8.629
	Ssk	4.498	0.034	3	9.169
	Sv	3.862	0.048	3	9.424
	Sz	4.282	0.042	3	8.432

Test statistics from WY-tests with 15% trimming for the silica concentration, microwear (primary surface only) and texture analyses of the primary surface (A), S-F surface (B), and S-L surface (C).

Values in bold indicate a significant difference ($p \le 0.05$). Ft = test statistics, nu1 and nu2 = 1st and 2nd degree of freedom, <math>p = significance level, SC = silica concentration [%] in animal feeds, Np = number of pits, Ls = length of scratches (µm), Sda = closed dale area [µm²], Sdv = closed dale volume [µm³], S5v = five point pit height [µm], Sal = auto correlation length [µm], Sp = maximum peak height [µm], Ssk = skewness, Sv = maximum pit height [µm], Sz = maximum height [µm].

doi:10.1371/journal.pone.0056167.t002

lowered the probability of surface lesions (Fig. 3). Table S1 in File S1 summarises the group-wise and pair-wise comparisons, and detailed test results for all variables are given in Tables S4, S5, S6 in File S1. Silica, microwear, and 3D texture parameters for the primary, S-F, and S-L surface indicate that beside silica concentration (SC) and microwear parameters (Ls, Np) the 3D texture parameters Sda (primary surface) and Sp (S-L surface) are most significant in differentiating the feeding group GO from G and L. Apart from Sda and Sp, the parameters Sdv (primary surface), Sdq, Sdr, Sz (S-F surface) and S5v, Sal, Ssk, Sdr, Sdq, Sv, Sxp, Sz and Vvv (S-L surface) describe single group differences (Table S1 in File S1). In the ISO/FDIS 25178 the parameters are pooled in height, spatial, hybrid, function and segmentation parameters [28]. We found that on the primary surface, parameters (Sda, Sdv) derived from segmentation parameters dividing the surface into motifs (dales and hills) yield prominent differentiation in our case, while on the S-F surface, a height (Sz) as well as two hybrid parameters (Sdr, Sdq) related to the spatial shape of the surface yield such differentiation. Only on the S-L surface all parameter groups (height (Sp, Ssk, Sz, Sv), spatial (Sal), hybrid (Sdr, Sdq) and function (S5v, Sxp, Vvv) yield such differentiation.

The discriminant analysis using the most significant microwear (Ls, Np) and texture parameters (*Sda* on primary surface, *Sp* on S-L surface) indicates that G and L occupy distinct factor spaces, while groups GO and LO overlap (Fig. 2F). The factor space of both groups which include grass meal (G, GO) was smaller than that for lucerne, which indicates a specific, consistent texture pattern, while the groups with lucerne (LO, L) showed a larger factor space which indicates higher variability in surface texture.

Discussion

Two-dimensional microwear [1,12,13,16,42–47] as well as the three-dimensional microwear texture (scale sensitive fractal analysis, [11]) method represent established approaches for dietary



Figure 3. Meshed 3D models. Meshed axiomatic 3D models of tooth enamel surfaces (primary surface) of the second upper molar (160×160 μm) of animals feeding on grass (UZH-G8), grass/oats (UZH-G04), lucerne/oats (UZH-L06), or lucerne (UZH-L2). A deeper red indicates a top of a plateau or hill, whereas a green towards dark blue indicates deeper areas. doi:10.1371/journal.pone.0056167.g003

reconstructions in mammals [11,20,48–51]. More recently, ISO parameters became available for such dietary reconstruction [21] and are used in dietary discrimination of fishes [36] and functional analysis of complex dental surfaces in ungulates [19] and primates [17]. Our controlled feeding experiments in rabbits confirm that scratching results from high silica content in the monocotyledonous dominated diet (grass), while pitting prevails in the dicotyle-donous dominated diet (lucerne). In the case of our study, where the same animal species was used, an influence of differences in dental morphology or chewing mechanisms can be largely excluded as an explanation of these differences. Because of the use of pelleted diets, the observed differences also cannot be contributed to differences in physical growth forms between monocotyledonous and dicotyledonous plants, or different cropping mechanisms required by the animals.

Our findings are consistent with the general assumption underlying the process of 2D microwear formation in ungulates, with browsers having more pits and grazers having more scratches [15,41,43,45,46,48,52]. Our results are partly in accordance with Gomes et al. [13] who found that grazing murids have a high abundance of scratches, while insectivorous and frugivorous murids have a coarser microwear pattern with a higher number of pits. We detected a similar difference between a grassdominated diet and another (non-grass) diet. Townsend and Croft [16] pointed out that 2D microwear patterns between feeding types are more subtle among cavimorph rodents than reported for ungulates and primates. In contrast, we found microwear patterns (Np, number of pits) of grass and lucerne-fed lagomorphs similar to patterns reported in grazing and browsing ungulates [41]. The higher absolute *Ls* value (length of scratch) is related to the smaller in body size in rabbits and hence the size of the molar surface.

The 3D microtexture parameters describe the surface texture pattern of lagomorphs in great detail. The ISO/FDIS parameters quantify the aspects of the basic geometry of texture and can be interpreted as a comprehensive representation of textures and its functional trait [17]. However, because the diets in the feeding experiment did not include seeds or other large and abrasive particles, the frequently stressed hypothesis that seeds are required to cause pit formation is not corroborated in this study. Instead, our results indicate that pits develop without such substrates. In particular, we found that the fewer silica particles are in the diet, the higher the variability of 3D textures. We relate this observation to a generally lower probability of abrasive wear when comminuting dicots such as lucerne or browse.

We found more significant surface texture parameters of the primary and S-L surface compared to the S-F surface. We link this to our very homogeneous sample which consists of domestic white rabbits only, when compared to interspecific datasets using the S-F surface only [17,19,34]. It is obvious that the segmentation parameters are prominent on the primary surface, before further filtering processes are applied. Height, hybrid, spatial and function parameters are dependent on the height distribution, which will be less distinct if a regular long wavelength pattern like form and waviness covers the short wavelength roughness signal. Therefore, these parameters become more prominent, after additional filtering processes, on the S-F and S-L surface.

A very interesting and, to our understanding of functional aspects of dental wear, very important question is why lowabrasion wear results in a more variable texture pattern dominated by plateau like structures and less areas of surface lesions. In particular, the biomechanical properties of teeth and diet responsible for the difference in the orientation of surface features (isotropic for high-abrasion diets, anisotropic for low-abrasion diets) remain to be identified experimentally. As our findings demonstrate, anisotropic orientation is associated with a less consistent, more randomly distributed texture pattern. Due to the lower frequency of abrasion on low-abrasion diets, microwear patterns on such diets must represent signals that accumulated over longer periods of time. For example, plateau-like structures may be present in higher proportions on low-abrasion diets because the lower wear rate does not remove them from the surface. "Overwriting", so to speak, takes longer and is less uniform on low-abrasion diets.

In contrast, processing high-abrasion diets causes a constant "overwriting" of previous signals with a uniform, isotropic pattern. This leads to the interesting hypothesis that those processes that lead to the microwear signal on low-abrasion diets occur on highabrasion diets as well, yet are not detected because their traces are constantly overwritten. Because grass eating rodents and ungulates often are hypsodont [53], it appears reasonable to assume that this overwriting is associated with increased wear rates. For the same reason, it is plausible to assume that a specific abrasion-dominated texture forms over a shorter period of time (the 'last supper' equivalent), whereas the texture on low-abrasion diets (as often found on low-crowned teeth) represents a variety of textureshaping incidents that are distributed over a longer series of meals.

In classical microwear research variability of scratch and pit counts on fossil teeth has often been used to infer dietary variability such as seasonality in foraging behaviour [54,55]. The variability signal in our sample of lucerne-fed rabbits, however,

References

- Baker G, Jones LHP, Wardrop ID (1959) Cause of wear in sheep's teeth. Nature 184: 1583–1584.
- McNaughton SJ, Tarrants JL (1983) Grass leaf silicification: Natural selection for an inducible defense against herbivores. Proceedings of the National Academy of Sciences 80: 790–791.
- Merceron G, Schulz E, Kordos L, Kaiser TM (2007) Paleoenvironment of Dryopithecus brancoi at Rudabánya, Hungary:evidence from dental meso- and micro-wear analyses of large vegetarian mammals. Journal of Human Evolution 53: 331–349.
- Sanson GD, Kerr SA, Gross KA (2007) Do silica phytoliths really wear mammalian teeth? Journal of Archaeological Science 34: 526–531.
- Massey FP, Ennos AR, Hartley SE (2007) Grasses and the resource availability hypothesis: the importance of silica-based defences. Journal of Ecology 95: 414– 424.
- Massey FP, Hartley SE (2006) Experimental demonstration of the antiherbivore effects of silica in grasses: impacts on foliage digestibility and vole growth rates. Proceedings of the Royal Society B: Biological Sciences 273: 2299–2304.
- Healy WB, Ludwig TG (1965) Wear of sheep's teeth: The role of ingested soil. New Zealand Journal of Agricultural Research 8: 737–752.
- Mayland HF, Shewmaker GE, Bull RC (1977) Soil ingestion by cattle grazing crested wheatgrass. Journal of Range Management 30: 264–265.
- Kaiser TM, Müller DWH, Fortelius M, Schulz E, Codron D, et al. (2011) Hypsodonty and tooth facet development in relation to diet and habitat in herbivorous ungulates: implications for understanding tooth wear. Mammal Review.

cannot be related to dietary variation, because the diet was constant throughout the experiment. These findings therefore caution against equating variation in the microwear signal with dietary variation, and emphasize that variation in the microwear signal might rather be linked to the overall abrasiveness of the diet.

For future studies, it will be particularly intriguing whether the variable pattern created by low-abrasion diets follows specific rules that are not detected yet, but that will allow additional conclusions for dietary reconstructions. For our understanding of the functional process of tooth wear, the exact conditions that lead to texture patterns associated with low-abrasion diets, such as differential forces due to uneven distributions of food between tooth antagonists, or subtle local differences in enamel hardness, remain to be investigated.

Supporting Information

File S1 Supporting figures and tables. Figure S1 Examples of cecotrophs in the stomach of rabbits. Table S1 Groupwise comparison of microwear and 3D texture parameters. Table S2 Descriptive statistics of microwear parameters. Table S3 Descriptive statistics of 3D texture parameters. Table S4 Statistics from WY-tests for the 3D texture parameters. Table S5 Statistics from pair-wise comparison for the microwear and 3D texture parameters. Table S6 Statistics for the microwear and 3D texture parameters. (PDF)

Acknowledgments

We thank Hanspeter Renfer and the Farmers' School Strickhof Lindau for supporting this project, Bettina Burger, Olga Martin Jurado and Carmen Füglistaller for help with rabbit handling, Bruno Gerzner for slaughtering the rabbits, and Klaus Zwonarz (University of Hamburg) for assistance in maceration and preparation of the rabbit skulls. This research is publication no. 45 of the DFG Research Unit 771.

Author Contributions

Conceived and designed the experiments: TMK MC ES GM. Performed the experiments: ES VP GM MM. Analyzed the data: ES VP GM. Contributed reagents/materials/analysis tools: ES GM MM MC TMK. Wrote the paper: ES GM MM MC TMK.

- Solounias N, Semprebon G (2002) Advances in the reconstruction of ungulate ecomorphology and application to early fossil equids. American Museum Novitates 3366: 1–49.
- Ungar PS, Brown CA, Bergstrom TS, Walker A (2003) Quantification of dental microwear by tandem scanning confocal microscopy and scale-sensitive fractal analyses. Scanning 25: 185–193.
- Walker A, Hoeck HN, Perez L (1978) Microwear of mammalian teeth as an indicator of diet. Science 201: 908–910.
- Gomes Rodrigues H, Merceron G, Viriot L (2009) Dental microwear patterns of extant and extinct Muridae (Rodentia, Mammalia): ecological implications. Naturwissenschaften 96: 537–542.
- Hopley PJ, Latham AG, Marshall JD (2006) Palaeoenvironments and palaeodiets of mid-Pliocene micromammals from Makapansgat Limeworks, South Africa: A stable isotope and dental microwear approach. Palaeogeography, Palaeoclimatology, Palaeoecology 233: 235–251.
- Nelson S, Badgley C, Zakem E (2005) Microwear in modern squirrels in relation to diet. Palaeontologia Electronica 8: 14A.
- Townsend KEB, Croft DA (2008) Enamel microwear in caviomorph rodents. Journal of Mammalogy 89: 730–743.
- Calandra I, Schulz E, Pinnow M, Krohn S, Kaiser TM (2012) Teasing apart the contributions of hard Items on 3D dental microtextures in primates. Journal of Human Evolution 63: 85–98.
- Merceron G, Escarguel G, Angibault J-M, Verheyden-Tixier H (2010) Can dental microwear textures record inter-individual dietary variations? PLoS ONE 5: e9542.

- Schulz E, Calandra I, Kaiser TM (2010) Applying tribology to teeth of hoofed mammals. Scanning 32: 162–182.
- Scott RS, Teaford MF, Ungar PS (2012) Dental microwear texture and anthropoid dets. American Journal of Physical Anthropology 147: 551–579.
- Kaiser TM, Brinkmann G (2006) Measuring dental wear equilibriums the use of industrial surface texture parameters to infer the diets of fossil mammals. Palaeogeography, Palaeocclimatology, Palaeoecology 239: 221–240.
- Butler PM (1972) Some functional aspects of molar evolution. Evolution 26: 474–483.
- Stones HH (1948) Oral and dental diseases. Edinburgh: E. and S. Livingstone.
 Teaford MF, Walker A (1983) Dental microwear in adult and still-born guinea pigs (*Cavia porcellus*). Archives of Oral Biology 28: 1077–1081.
- Rensberger JM (1978) Scanning electron microscopy of wear and occlusal events in some small herbivores. In: Butler PM, Joysey KA, editors. Development, Function and Evolution of Teeth. New York: Academic Press. pp. 415–438.
- Fortelius M (1985) Ungulate check teeth: developmental, functional, and evolutionary interrelations. Acta Zoologica Fennica 180: 1–76.
 Königswald Wv, Golenishev FN (1979) A method for determining growth rates
- Kongswald WV, Golensnev FN (1979) A method for determining growth rates in continuously growing teeth. Journal of Mammalogy 60: 397–400.
- International Organization for Standardization (2010) ISO/FDIS 25178-2 Geometrical product specifications (GPS) – Surface textureAreal – Part 2: Terms, definitions and surface texture parameters.
- Leiber F, Meier J, Burger B, Wettstein H-R, Kreuzer M, et al. (2008) Significance of coprophagy for the fatty acid profile in body tissues of rabbits fed different diets. Lipids 43: 853–865.
- Gidenne T, Lebas F (2006) Feeding behaviour in rabbits. In: Bels V, editor. Feeding in Domestic Vertebrates: From structure to behaviour. Wallingford: CAB International. pp. 179–194.
- Rogers AM, Arthur CP, Soriguer RC (1994) The rabbit in continental Europe. In: Thompson HV, King CM, editors. The European rabbit: the history and biology of a successful colonizer. Oxford: Oxford University Press. pp. 22–63.
- Parr JF, Lentfer CJ, Boyd WE (2001) A comparative analysis of wet and dry ashing techniques for the extraction of phytoliths from plant material. Journal of Archaeological Science 8: 875–886.
- Merceron G, Blondel C, De Bonis L, Koufos GD, Viriot L (2005) A new method of dental microwear analysis: application to extant primates and *Ouranopithecus* macedoniensis (Late Miocene of Greece). Palaios 20: 551–561.
- Winkler DE, Schulz E, Calandra I, Gailer JP, Landwchr C, et al. (2012) Indications for a dietary change in the extinct Bovid genus *Myotragus* (Plio-Holocene, Mallorca, Spain). Geobios. In press.
- Williams J (2005) Engineering tribology. Cambridge: Cambridge University Press. 488 p.
- Purnell M, Sechausen O, Galis F (2012) Quantitative three-dimensional microtextural analyses of tooth wear as a tool for dietary discrimination in fishes. Journal of the Royal Society Interface 9: 2225–2233.
- R Development Core Team (2010) R: A language and environment for statistical computing. Vienna, Austria: R Foundation for Statistical Computing.
- Welch BL (1938) The significance of the difference between two means when the population variances are unequal. Biometrika 29: 350–362.

- 39. Yuen KK (1974) The two-sample trimmed t for unequal population variances. Biometrika 61: 165–170.
- Dunnett CW (1980) Pairwise multiple comparisons in the unequal variance case. Journal of the American Statistical Association 75: 796–800.
- Merceron G, Costeur L, Maidet O, Ramdarshan A, Göhlich UB (2012) Multproxy approach detects heterogeneous habitats for primates during the Miocene climatic optimum in Central Europe. Journal of Human Evolution 63: 150–161.
- 42. Calandra I, Göhlich UB, 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.
- 43. Merceron G, Ungar P (2005) Dental microwear and palaeoecology of bovids from the Early Pliocene of Langebaanweg, Western Cape Province, South Africa. South African Journal of Science 101: 365–370.
- Semprebon G, Godfrey LR, Solounias N, Sutherland MR, Jungers WL (2004) Can low-magnification stereomicroscopy reveal diet? Journal of Human Evolution 47: 115–144.
- Solounias N, Dawson-Saunders B (1988) Dietary adaptations and paleoccology of the Late Miocene ruminants from Pikermi and Samos in Greece. Palaeogeography, Palaeoclimatology, Palaeoccology 65: 149–172.
- Solounias N, Moelleken SMC (1993) Tooth microwear and premaxillary shape of an archaic antelope. Lethaia 26: 261–268.
- Teaford MF (1988) A review of dental microwear and diet in modern mammals. Scanning Microscopy 2: 1149–1166.
- Ungar P, Merceron G, Scott R (2007) Dental microwear texture analysis of Varswater Bovids and Early Pliocene paleoenvironments of Langebaanweg, Western Cape Province, South Africa. Journal of Mammalian Evolution 14: 163–181.
- Scott JR, Godfrey LR, Jungers WL, Scott RS, Simons EL, et al. (2009) Dental microwear texture analysis of two families of subfossil lemurs from Madagascar. Journal of Human Evolution 56: 405–416.
- Scott RS, Ungar PS, Bergstrom TS, Brown CA, Childs BE, et al. (2006) Dental microwear texture analysis: technical considerations. Journal of Human Evolution 51: 339–349.
- Scott RS, Ungar PS, Bergstrom TS, Brown CA, Grine FE, et al. (2005) Dental microwear texture analysis shows within-species diet variability in fossil hominins. Nature 436: 693–695.
- Schubert BW, Ungar PS, Sponheimer M, Reed KE (2006) Microwear evidence for Plio–Pleistocene bovid diets from Makapansgat Limeworks Cave, South Africa. Palaeogeography, Palaeoclimatology, Palaeoecology 241: 301–319.
- Williams SH, Kay RF (2001) A comparative test of adaptive explanations for hypsodonty in ungulates and rodents. Journal of Mammalian Evolution 8: 207– 229.
- Merceron G, Viriot L, Blondel C (2004) Tooth microwear pattern in roe deer (*Capreolus capreolus* L.) from Chizé (Western France) and relation to food composition. Small Ruminant Research 53: 125–138.
- Rivals F, Deniaux B (2005) Investigation of human hunting seasonality through dental microwear analysis of two Caprinae in late Pleistocene localities in Southern France. Journal of Archaeological Science 32: 1603–1612.