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Article

Exploratory Analysis of Objective Outcome Measures for the Clinical Assessment of Erosive Tooth Wear

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Abstract: This study proposed using enamel surface texture and thickness for the objective detection and monitoring of erosive tooth wear (ETW), comparing them to the standard subjective Basic Erosive Wear Evaluation (BEWE). Thirty-two subjects (n = 597 teeth) were enrolled in this longitudinal observational clinical study. Enamel thickness (by cross-polarization optical coherence tomography, CP-OCT) and 3D dental microwear parameters, i.e., area-scale fractal complexity (Asfc), anisotropy (Str), and roughness (Sa) (by white-light scanning confocal profilometry), were obtained from buccal surfaces. Buccal, occlusal, and lingual surfaces were scored for BEWE and the maximum score per tooth (BEWE_{Max}) was determined at baseline and 12 months (M12). Data outcome relationships were evaluated (alpha = 0.05). Enamel thickness decreased (p < 0.001), BEWE scores, Sa, and Str increased (p < 0.001), while Asfc did not change at M12. Baseline BEWE_{Buccal} correlated strongly with $BEWE_{Max}$ (r = 0.86, p < 0.001) and moderately with $BEWE_{Lingual}$ (r = 0.42, p < 0.001), but not with enamel thickness (r = 0.03, p = 0.43). Change (Δ) in surface texture outcomes correlated poorly but significantly with $\Delta BEWE_{Buccal}$ (r = -0.15–0.16, p < 0.001) and did not correlate with $\Delta enamel$ thickness (r = 0.02–0.09, p > 0.06). Teeth with BEWE progression revealed a greater increase in Δ Sa and ΔStr. These findings suggest that enamel surface roughness can potentially determine ETW severity, and CP-OCT may be relevant for clinically monitoring enamel thickness.

Keywords: erosive tooth wear; dental enamel; optical coherence tomography; enamel surface texture; BEWE

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1. Introduction

Erosive tooth wear (ETW) is a dental condition that results in loss of tooth structure as a consequence of chemo-mechanical wear processes [1]. Acids that are present in

Diagnostics **2023**, 13, 2568 2 of 12

the oral cavity from intrinsic (gastric) and/or extrinsic (dietary) sources [2] can soften exposed enamel and dentin surfaces and make them vulnerable to wear from the abrasive forces of mastication and toothbrushing. Given the irreversible nature of ETW, tooth form, esthetics, and function can be compromised [3]. ETW has a high prevalence worldwide [4] and it affects approximately 46% of teenagers [5] and 80% of adults [6] in the United States, as reported by the National Health and Nutrition Examination Survey (NHANES). Despite these alarming numbers, specific diagnostic rules and evidence-based management guidelines are yet to be established. Currently, clinical assessment and monitoring of ETW are performed by visual examination using subjective indices [7]. This traditional approach limits the detection of ETW lesions to mostly advanced stages, wherein considerable destruction of the tooth has already occurred, resulting in pain and irreversible changes in dental form, function, and esthetics. In these circumstances, the required restorative treatments are complex and costly [8]. There is a clear need for objective methods to detect, diagnose, and monitor ETW early, allowing the implementation of personalized and evidence-based management plans focused specifically on preventive measures.

This exploratory analysis involved subjects previously diagnosed with hyposalivation. Saliva plays a crucial role in the development of ETW, and the lack thereof puts patients at higher risk for ETW development or progression [9,10]. Our first hypothesis in this investigation was that ETW lesions could be detected and differentiated by using scalesensitive, tridimensional dental surface texture analysis of point clouds generated by white-light scanning confocal profilometry (WSCP). Some integrated metrics of dental microwear analysis have proven to be useful for similar dental applications, including surface fractal complexity (or change in apparent roughness with the scale of observation) and anisotropy (or directionality of the wear pattern) [11]. Our previous in vitro and in situ data have shown the ability of these outcomes to identify and differentiate between the main dental wear mechanisms of ETW lesions with a good degree of certainty [12,13]. Our second hypothesis was that cross-polarization optical coherence tomography (CP-OCT) could provide objective clinical measures of ETW progression, based on the longitudinal monitoring of dental enamel thickness. CP-OCT allows for safe, non-destructive, and repetitive measurements of enamel thickness [14]. It also allows for the study of the tissues' polarization properties and provides higher-resolution images and better visualization of the enamel structure and dentin–enamel junction [15,16]. Our previous in vitro and in situ studies have demonstrated its potential as a clinical tool to monitor ETW [17,18].

This exploratory study aimed to compare the objective outcomes of enamel surface texture using WSCP and enamel thickness with CP-OCT to a standard subjective visual assessment (Basic Erosive Wear Evaluation—BEWE) and explore their potential to be used clinically for the detection and monitoring of naturally developed ETW lesions in a high-risk population.

2. Materials and Methods

2.1. Study Design

This study consisted of a single-site longitudinal observational clinical study, conducted at the Oral Health Research Institute of the Indiana University School of Dentistry (IUSD). Twenty-nine subjects previously diagnosed with dry mouth at The Center for Oral Diagnosis and Treatment, IUSD, or any other IUSD clinic, and three control (no dry mouth) subjects were enrolled to participate in the study. The study was carried out for about 20 months employing a 'rolling' subject recruitment that lasted for six months. Each recruited subject was then assessed periodically up to 12 months after their baseline visit.

A trained and calibrated examiner performed the clinical assessment using the BEWE index to classify the surfaces of all incisors, canines, premolars, and first molars according to the degree of severity of the ETW lesion (score 0: no erosion; 1: loss of enamel surface texture; 2: enamel loss of <50% of surface area; 3: enamel loss of >50%) and a self-completed questionnaire (diet and behavioral habits) was used to identify related risk factors. Subjects at high risk for ETW development were selected for the study based on these criteria.

Diagnostics 2023, 13, 2568 3 of 12

Further evaluation of the subjects' teeth by trained examiners was performed using the test objective outcomes of surface texture and enamel thickness. Outcomes were assessed at baseline and after 3 (M3), 6 (M6), 9 (M9), and 12 (M12) months.

2.2. Study Participants

The clinical study protocol followed the Declaration of Helsinki and was reviewed and approved by the Indiana University-Purdue University Indianapolis Institutional Review Board (protocol no. 1910664803). Subject recruitment and screening occurred between July 2020 and February 2021. Potential subjects previously diagnosed with dry mouth at The Center for Oral Diagnosis and Treatment (CODT, IUSD) were invited to participate in the study. The control subjects (no diagnosis of hyposalivation) were recruited from the IUSD dental clinics. These subjects were prescreened to meet age and health requirements using an IRB-approved phone script.

Prior to screening, subjects were asked to sign a written informed consent form. To be included in the study, subjects had to be 18–85 years old and generally healthy; have a minimum of eight BEWE-scorable teeth with at least one ETW lesion (BEWE \geq 1), except for control subjects; have indicated dietary acid exposure in the questionnaire; and have been previously diagnosed with dry mouth, except for controls. Control subjects needed to have a normal salivary flow rate of \geq 0.8 mL/min stimulated and \geq 0.2 mL/min unstimulated saliva and teeth without clinical signs of advanced ETW. Subjects were excluded if they were visually assessed by the study dentist to clinically have any untreated cavitated caries lesions or moderate to severe periodontal disease. Subjects who presented with caries lesions at screening were only allowed to continue in the study if they had their caries lesions appropriately treated before the baseline visit.

The sample size calculation indicated the need to recruit 60 subjects with a higher risk for ETW (dry mouth) and 8 controls (normal salivary flow). We assumed 8 teeth per subject, with baseline BEWE scores in high-risk subjects as follows: 10% of teeth with BEWE 3, 20% with BEWE 2, 30% with BEWE 1, and 50% with BEWE 0. The 95% lower confidence bound (LCB) for overall model accuracy to classify teeth by baseline BEWE score extends 3% from the estimated accuracy; within each BEWE score, the LCB extends up to 10%. The expectation was that at least 10% of teeth would show ETW progression, so the LCB for accuracy of the model predicting ETW progression extends up to 10% from the estimated accuracy.

2.3. Clinical Study Procedures

Study completion entailed subject participation during each of the six study visits: screening, baseline, M3, M6, M9, and M12. Study participants were asked for informed consent and their medical history as well as the completion of a self-administered diet and behavioral questionnaire. Oral soft and hard tissue examinations were conducted and clinical assessment using the BEWE index was performed by a trained/calibrated examiner to determine which subjects qualified for the study. Qualified subjects had alginate impressions (Jeltrate Plus, Dentsply Caulk, Milford, DE, USA) taken for both dental arches that were used for the fabrication of CP-OCT tray guides (described below) and modified custom trays. Unstimulated and stimulated saliva flow rates were measured during the initial visit for the control group subjects. Qualified subjects returned after 1 week (± 5 days) for the continuation of the baseline assessments. Salivary flow rates were then determined for the dry-mouth subjects during the subsequent visit after discontinuing their saliva-stimulating medications 24 h prior. The subjects' teeth were brushed using a prepasted toothbrush (ReadyBrush, ReadyBrush, Boca Raton, FL, USA), rinsed, and dried to remove the biofilm or pellicle. Dental impressions were taken for both dental arches of all the subjects using polyvinylsiloxane impression material (President Jet Regular and Light Body, Coltene/Whaledent AG, Altstätten, Switzerland) for enamel surface texture analysis. Three-dimensional CP-OCT images were then taken from the window of the CP-OCT imaging tray guide for the measurements. Each of the assessment methods for

Diagnostics 2023, 13, 2568 4 of 12

ETW (BEWE, diet and behavioral questionnaire, CP-OCT enamel thickness, and surface texture) was conducted during the succeeding visits at M3, M6, M9, and M12.

2.4. BEWE Scoring

ETW severity was scored and recorded according to the BEWE index [19]. A previously trained examiner scored the buccal, occlusal (for posterior teeth), and lingual surfaces of all incisors, canines, premolars, and first molars according to the degree of severity of the ETW lesion. Scoring was performed at baseline, M3, M6, M9, and M12.

2.5. Surface Texture Analysis

White-light scanning confocal profilometry (Neox, Sensofar LLC, Newington, CT, USA) was used to analyze the buccal surface of the impressions. The central area of the middle third (mesiodistally and inciso/occluso-cervically) of the buccal surface was chosen for analysis. The planimetric work envelope for each tooth sample was $242 \times 181~\mu m^2$ with a lateral point spacing of $0.17~\mu m$ in both x and y directions, a vertical step of $0.2~\mu m$, and a published resolution of <2 nm, measured in three locations, as previously reported [12]. Trained and calibrated examiners performed measurements on the central area of the buccal surfaces of specimens collected at baseline and M12. Point clouds for each surface were assessed by dental microwear texture analysis. Area-scale fractal complexity (Asfc) [20] and ISO 25178 standards for texture aspect ratio (Str) and arithmetical mean height (Sa) were calculated to characterize scale-sensitive complexity, anisotropy, and roughness of each surface, respectively. Analyses were conducted using MountainsMap 8 (Digital Surf, Besançon, France). These variables have been shown to consistently reveal aspects of the surface texture of value for distinguishing dental wear types [12,21,22].

2.6. Enamel Thickness Analysis

CP-OCT imaging tray guides (Figure 1a) were custom-made to fit the subjects' maxillary and mandibular dental arches using soft ethylene vinyl acetate sheets (125 mm round, 1 mm thick, Keystone Industries, Gibbstown, NJ, USA). The sheets were vacuum-formed to the casts of each dental arch using a pressure molding machine (Biostar® SCHEU-DENTAL GmbH, Iserlohn, Germany) and trimmed to extend half a tooth away from the most posterior tooth included in the study and about 2 mm away from the gingival margin to maintain good adaptation. Round window holes 2 mm in diameter were created on the middle third (mesiodistally and inciso/occluso-cervically) of the buccal surfaces. A thin layer of red nail varnish was applied onto the walls of the CP-OCT tray guide windows for easier localization while positioned in the subject's oral cavity. The same CP-OCT imaging tray guides with round windows were used for each subject over the different study visits (while observing infection control measures) to facilitate CP-OCT probe repositioning and imaging of the same area on the buccal surfaces of the tooth over time to be used for longitudinal enamel thickness analysis.

Three-dimensional enamel scans were acquired using a portable dental CP-OCT system with a handheld probe (Santec Inner Vision IVS-300-S-L-C; Santec Corp, Komaki, Japan). The device used a swept source laser light with a center wavelength of 1310 ± 30 nm and a high scan rate of 30 kHz with a maximum lateral probe scanning area of 5×5 mm and a working distance of 1 mm. Axial imaging in the air was >5.6 mm with a 3 mm depth of focus while axial and lateral resolutions in the air were $\leq\!12~\mu m$ and $30~\mu m$, respectively.

Before scanning, the buccal surfaces of the maxillary teeth were gently air-dried for ~10 s and isolated using cotton rolls as needed. The CP-OCT imaging tray guide was then fitted on top of the teeth and 3D CP-OCT scanning was carried out. During scanning, the probe was held horizontally perpendicular to the long axis of the tooth being scanned and positioned directly on top of the CP-OCT tray guide with the borders of the CP-OCT tray guide window contained within the scanning area. Three-dimensional tomograms were obtained from the buccal surfaces of each of the included maxillary teeth using a

Diagnostics 2023, 13, 2568 5 of 12

dedicated imaging and analysis software (Inner Vision IVS-300 ver5.3.2, Santec Corp., Komaki, Japan) where the refractive index was set at 1.6 for enamel. Once the maxillary arch was scanned, the maxillary CP-OCT imaging tray guide was removed, and the same methods were applied to the mandibular arch. Central B-scans (2D images) in the Y-direction were then selected from each 3D scan and saved for enamel thickness analyses. Enamel thickness measurements (from DEJ to the surface of the specimen) on the 2D images were performed using Santec Inner Vision IVS-300 software (Santec Corp., Komaki, Japan). The measurement position was identified at the center of the enamel width at the base of the CP-OCT tray guide window with the aid of a screen ruler (A Ruler for Windows v3.3, https://www.arulerforwindows.com accessed on 8 July 2022). The distance (mm) between the depths of the highest light intensity peaks at the enamel surface and DEJ areas was then calculated from the A-scan. CP-OCT scanning and enamel thickness measurements were performed at baseline, M3, M6, M9, and M12. A representative B-scan of a maxillary molar is shown in Figure 1c.

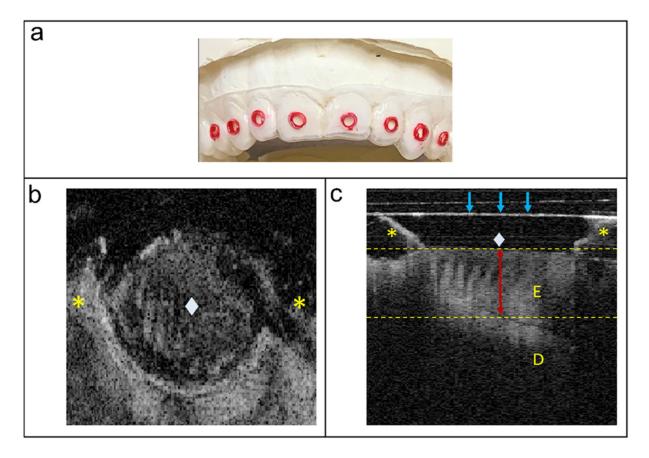


Figure 1. CP-OCT imaging of the buccal surface of a maxillary molar. (a) Anterior view of a maxillary CP-OCT imaging tray guide. (b) Top view. (c) B-scan. E: enamel; D: dentin. * CP-OCT tray guide; ♦ enamel surface within the 2 mm round CP-OCT tray guide window. Blue arrows: CP-OCT probe window with a plastic barrier; red arrow: area measured for enamel thickness which is the midline of the enamel width as measured from the base of the CP-OCT tray guide window.

2.7. Statistical Analyses

Data collected at baseline and M12 for all parameters were used for the analyses. Data for M3, M6, and M9 were excluded as surface texture parameters of Asfc (complexity), Str (anisotropy), and Sa (roughness) could not be obtained.

Spearman correlation coefficients were used to evaluate the associations among measurements. Using the longitudinal follow-up assessments, linear mixed-effects models evaluated changes in surface texture and enamel thickness (monitor ETW). These models

Diagnostics 2023, 13, 2568 6 of 12

included random effects to correlate data within a tooth over time, as well as allow different variances at each time point and correlate multiple teeth within a subject. Teeth with changes in BEWE scores were compared to those without such changes during follow-up for differences in changes in surface texture and enamel thickness parameters using linear mixed-effects models. A two-sided 5% significance level was used for all tests.

3. Results

Twenty-nine subjects with a history of diagnosed hyposalivation and three control subjects were enrolled in the study. One dry-mouth subject dropped out after the baseline visit. A maximum number of 597 teeth that were scoreable for BEWE on any of the surfaces at baseline were included. Among them, 584 teeth were used for surface texture analyses of their buccal surfaces, and 531 teeth were included for the enamel thickness measurements with CP-OCT. Teeth that were malposed or posteriorly located with the buccal mucosa not allowing proper CP-OCT probe positioning were excluded. Out of the 29 subjects who had a history of hyposalivation, 9 and 18 had normal unstimulated and stimulated saliva flow rates, respectively.

BEWE scores of the different surfaces (BEWE_{Buccal}, BEWE_{Occlusal}, BEWE_{Lingual}) at baseline and M12 of dry-mouth subjects are shown in Table 1. BEWE for all surfaces was greater at M12 compared to baseline (p < 0.001). For the same longitudinal comparison, enamel thickness decreased, Asfc showed no change, and Sa and Str parameters increased. Results for the different parameters for control subjects are shown in Table S1. Statistical analyses within the control group, as well as the comparison of the control and dry-mouth subjects, were not performed due to the small number of control subjects recruited.

Table 1.	Erosive tooth wea	ar (ETW) outcom	ies (mean and	l standard	deviation) a	t baseline	and
12 month	ns in dry-mouth sub	jects.					

ETW Outcomes	Baseline	12 Months	<i>p-</i> Value
Asfc (complexity) a	1.25 (0.80)	2.35 (5.94)	0.857
Sa (roughness) b	207.04 (121.24)	375.61 (213.62)	< 0.001
Str (anisotropy) c	0.48 (0.17)	1.23 (1.35)	< 0.001
Enamel Thickness ^d	1056 (247)	1042 (249)	< 0.001
$BEWE_{Buccal}$	0.88 (0.61)	1.08 (0.48)	< 0.001
BEWE _{Occlusal}	0.82 (0.69)	1.38 (0.52)	< 0.001
BEWE _{Lingual}	0.64 (0.56)	1.13 (0.46)	< 0.001
$BEWE_{Max}$	0.99 (0.63)	1.33 (0.56)	< 0.001

^a Asfc—area-scale fractal complexity (no unit); ^b Sa (nm); ^c Str—texture aspect ratio (no unit); ^d enamel thickness (μm).

The direct comparison between tested parameters (Table 2) revealed that BEWE $_{Occlusal}$ correlated poorly with enamel thickness as measured on the buccal surface of posterior teeth. BEWE $_{Buccal}$, on the other hand, did not correlate with enamel thickness measurements. BEWE $_{Buccal}$ and BEWE $_{Lingual}$ correlated moderately, while BEWE $_{Buccal}$ correlated strongly with the BEWE $_{Max}$. Comparisons of the longitudinal changes indicated that Δ Asfc and Δ Str correlated moderately, while Δ Asfc and Δ Sa, as well as Δ Sa and Δ Str, exhibited weak correlation (Table 3). Both Δ BEWE $_{Buccal}$ and Δ BEWE $_{Lingual}$ correlated moderately with Δ BEWE $_{Max}$. A similar correlation was observed between Δ BEWE $_{Occlusal}$ (posterior teeth) and Δ BEWE $_{Max}$. Only data at baseline and M12 were compared and analyzed.

Additional analyses comparing teeth with and without ETW progression—based on BEWE examination—revealed no significant differences in Δ Asfc and Δ enamel thickness over time. However, teeth with ETW progression (BEWE increase) showed greater changes in Δ Sa and Δ Str over time compared to teeth without ETW progression (Table 4).

Diagnostics **2023**, 13, 2568 7 of 12

 Table 2. Overall correlations between ETW outcome measurements at baseline and M12.

ETW Outcomes		Baseline			M12		
		п	r	<i>p</i> -Value	п	r	<i>p</i> -Value
	Sa	584	0.63	<0.001	440	0.04	0.464
A . C	Str	584	0.12	0.005	473	-0.42	< 0.001
Asfc	Enamel Thickness	506	-0.02	0.676	491	-0.08	0.090
	$BEWE_{Buccal}$	584	-0.11	0.006	506	0.09	0.046
	Str	584	0.27	< 0.001	391	0.22	< 0.001
Sa	Enamel Thickness	506	-0.01	0.889	417	0.09	0.071
	$BEWE_{Buccal}$	584	-0.14	0.001	428	-0.13	0.007
Str	Enamel Thickness	506	0.07	0.104	445	0.17	< 0.001
Str	$BEWE_{Buccal}$	584	-0.04	0.398	459	-0.14	0.002
Enamel	BEWE _{Buccal}	514	0.03	0.431	531	-0.01	0.859
Thickness	BEWE _{Occlusal}	120	-0.19	0.040	129	-0.28	0.002
	BEWE _{Occlusal}	150	0.18	0.025	135	0.05	0.533
$BEWE_{Buccal}$	$BEWE_{Lingual}$	568	0.42	< 0.001	523	0.21	< 0.001
	$BEWE_{Max}$	597	0.86	< 0.001	558	0.64	< 0.001
DEWE	BEWE _{Lingual}	157	0.58	<0.001	145	*	*
BEWE _{Occlusal}	$BEWE_{Max}$	157	0.50	< 0.001	146	0.75	< 0.001
BEWE _{Lingual}	$BEWE_{Max}$	595	0.55	<0.001	562	0.66	<0.001

^{*} Correlation was not computed because all observations with both measurements have $\overline{BEWE_{Lingual}} = 1$.

Table 3. Overall correlations between change (Δ) (M12–baseline) in ETW outcome measurements.

ETW O	n	r	<i>p</i> -Value	
	ΔSa	429	0.17	0.001
	$\Delta \mathrm{Str}$	460	-0.34	< 0.001
A A - C -	ΔEnamel Thickness	430	0.09	0.065
$\Delta Asfc$	$\Delta { m BEWE}_{ m Buccal}$	494	-0.15	0.001
	$\Delta \text{BEWE}_{\text{Occlusal}}$	117	0.10	0.302
	$\Delta BEWE_{Max}$	507	-0.10	0.018
	ΔStr	380	0.23	< 0.001
	ΔEnamel Thickness	373	0.09	0.075
ΔSa	$\Delta { m BEWE}_{ m Buccal}$	418	0.16	0.001
	$\Delta \text{BEWE}_{\text{Occlusal}}$	100	-0.18	0.066
	$\Delta \text{BEWE}_{ ext{Max}}$	428	0.03	0.531
	ΔEnamel Thickness	387	0.02	0.643
	$\Delta { m BEWE}_{ m Buccal}$	447	0.16	< 0.001
$\Delta \mathrm{Str}$	$\Delta \text{BEWE}_{\text{Occlusal}}$	103	-0.19	0.056
	$\Delta { m BEWE}_{ m Lingual}$	430	-0.08	0.120
	$\Delta ext{BEWE}_{ ext{Max}}$	458	-0.04	0.370
ΔEnamel Thickness	$\Delta BEWE_{Buccal}$	472	0.03	0.473
	$\Delta BEWE_{Occlusal}$	129	0.15	0.100
$\Delta \text{BEWE}_{ ext{Buccal}}$	$\Delta \text{BEWE}_{ ext{Lingual}}$	518	0.32	< 0.001
	$\Delta ext{BEWE}_{ ext{Max}}$	553	0.61	< 0.001
AREME	$\Delta ext{BEWE}_{ ext{Lingual}}$	141	0.47	< 0.001
$\Delta BEWE_{Occlusal}$	$\Delta BEWE_{Max}$	141	0.60	< 0.001
$\Delta \text{BEWE}_{\text{Lingual}}$	$\Delta BEWE_{Max}$	559	0.61	<0.001

Diagnostics 2023, 13, 2568 8 of 12

ETW Outcomes	BEWE Increase	п	Mean (Standard Deviation)	<i>p</i> -Value
ΔAsfc (complexity) ^a	No Yes	362 132	0.60 (4.28) 0.92 (6.97)	0.770
ΔSa (roughness) ^b	No Yes	309 109	137.32 (251.12) 234.42 (246.48)	0.006
ΔStr (anisotropy) ^c	No Yes	322 125	0.65 (1.28) 1.28 (1.45)	<0.001
ΔEnamel Thickness ^d	No Yes	364 108	-26.44 (83.84) -23.33 (67.97)	0.725

Table 4. Comparisons between teeth with vs without BEWE changes (M12–baseline).

4. Discussion

Our previous in vitro [12,18] and in situ studies [13,17] have demonstrated the potential of enamel thickness (by CP-OCT) and the dental microwear texture parameters Asfc, Str, and Sa (by WSCP) to detect and monitor ETW lesion progression over time. However, in vitro and in situ studies are limited as they do not fully replicate clinical conditions. In the present exploratory clinical study, these objective outcomes were compared to the BEWE index, a subjective but widely used clinical index for ETW. The study population consisted of patients previously diagnosed with hyposalivation, as they were considered at higher risk for ETW, and their clinical findings were to be compared to those of subjects who were never diagnosed nor had symptoms of hyposalivation (control). Unfortunately, our planned recruitment and part of the laboratory analyses (dental microwear) were negatively impacted by the restrictions imposed by the COVID-19 pandemic. Even after extending the recruitment phase from 3 to 6 months, it was not feasible to meet our total target subject numbers (n = 68). A decision was made to stop enrollment, considering that we had reached the number of teeth needed, as determined by our a priori sample size calculation (n = 544). This was possible as we had initially set the minimum number of scoreable teeth per subject at eight, but most subjects had a significantly higher number of teeth that were scoreable with BEWE. However, it is important to note that the lower number of enrolled subjects may have limited the individual subject variation representation in our data. Nevertheless, the varied nature of ETW per tooth surface within the same subject supported the shift to using the total number of teeth instead of the number of subjects as the basis for the sample size. In addition to the aforementioned limitation brought about by the smaller number of subjects, the initially planned comparison between hyposalivation and control participants (at n = 3), wherein ETW progression was less expected, was no longer applicable and is thus a limitation of this study. Tooth-based comparisons between teeth with and without ETW progression within the hyposalivation group were performed instead to see the trends and associations among the different ETW parameters (Table 4).

In the current study, all surfaces of each tooth included were scored for BEWE, and the BEWE_{Buccal} scores were compared to those of the other parameters. Progression of ETW over time was observed in the hyposalivation population, as shown by the significant increase in BEWE scores from baseline to M12 for all surfaces (p < 0.001). Nonetheless, the mean increase in the BEWE score was less than 1 and smaller than what we anticipated for high-risk subjects. In hindsight, a longer study duration could have been advantageous to better discern between subjects at high and low risk for ETW. Although recruited subjects had been previously diagnosed with hyposalivation, and were therefore at higher risk for ETW [9,10], 9 showed normal unstimulated salivary flow, while 18 out of the 29 subjects had normal stimulated flow rates during testing, despite temporarily discontinuing their saliva-stimulating medications at least 24 h prior to the measurements. This might indicate that for these subjects, dry mouth may not have been a major risk factor for ETW. Further

 $^{^{}a}$ Asfc—area-scale fractal complexity (no unit); b Sa (nm); c Str—texture aspect ratio (no unit); d enamel thickness (μ m).

Diagnostics 2023, 13, 2568 9 of 12

differentiation of the subjects based on their salivary flow rates was not performed since the related inclusion criterion was only a history of hyposalivation. Moreover, salivary flow rates could have varied during the length of the study due to the adoption of individual measures to control the clinical symptoms or changes in the severity of the condition [23]. Future studies ensuring a larger number of subjects and periodical measurement of actual salivary flow rates may provide better insights into the differences in effects of an earlier hyposalivation diagnosis and specific salivary flow rates on ETW progression. In terms of surface comparison, change in BEWE was greater for the occlusal and lingual surfaces than for the buccal surface, which means that ETW progression was more evident on both occlusal and lingual surfaces than the buccal. However, BEWE on all three surfaces correlated moderately with BEWE_{Max} (Table 2). This supports the use of buccal surfaces for comparison with the other test parameters in this study.

The presence of ETW progression was also supported by the decrease in enamel thickness from baseline to M12 as measured by CP-OCT. This corroborates CP-OCT's potential to monitor changes in enamel thickness both in vitro and in situ within the same tooth over time [17,18]. These prior studies [17,18] were carried out in ideal conditions, with enamel specimens of standardized dimensions and flat enamel surfaces allowing proper working distance and probe angulation during imaging. Despite the encouraging results observed in the current study, there were many challenges. Our measurements were limited to the buccal surfaces only as the wide and flat probe head configuration made the access of lingual surfaces challenging. Also, clinical CP-OCT imaging was limited by different surrounding anatomical structures, such as the oral soft tissues. As mentioned earlier, teeth that are malposed or with oral soft tissues precluding proper CP-OCT probe positioning were excluded, resulting in relatively fewer teeth examined for the enamel thickness parameter in the study. The natural curvature of teeth as well as the involvement of the whole tooth contributed to greater difficulty in probe positioning. Hence, the use of a CP-OCT tray guide with pre-established windows was essential in ensuring more accurate repositioning and imaging of the same area of each tooth during every visit. The scanning window of the CP-OCT probe was held as close as possible and parallel to the buccal surface of the CP-OCT tray guide during scanning. Nevertheless, it was inevitable to have some degree of variation during imaging. In such cases, there were slight differences in the angulation of the surface being scanned. This also could have had an effect on the variability and the ease of obtaining enamel thickness measurements at the center of the enamel width captured within the CP-OCT tray window from the enamel surface to the DEJ. For teeth in which some angulation variation was unavoidable, particularly in the molars and malposed teeth, uneven light attenuation was more evident with the unequal working distance from the enamel surface. A greater distance meant higher attenuation and a lower contrast of enamel from the underlying structures, which made the analysis of the central B-scans more difficult. Considering these limitations, it would be desirable if OCT systems for clinical use were further developed with a better probe head configuration and better light intensity to offset the attenuation effect with greater distances. Despite the imaging limitations, a 29 ± 72 µm mean change in enamel thickness, which was relatively small but significant, was detected over time, confirming the suitability of using CP-OCT to monitor ETW longitudinally. The detection of only a minimal change in enamel thickness was consistent with the relatively small change in BEWE scores.

Both BEWE and enamel thickness changed significantly over time, suggesting that both parameters could independently detect the clinical progression of ETW. However, in terms of correlation, neither enamel thickness nor change in enamel thickness correlated with BEWE or with a change in BEWE scores. There were also no significant differences in the changes in enamel thickness when comparing teeth that showed and did not show changes in BEWE (Table 4). This might be because a much larger area of the tooth was considered for the BEWE scoring while CP-OCT measured only a comparatively smaller area for enamel thickness measurements. Furthermore, the two parameters measured two different aspects of ETW severity on the enamel. BEWE scoring is performed depending

Diagnostics **2023**, 13, 2568 10 of 12

on the surface area involved, whereas enamel thickness measurements reflect surface loss as a function of depth into the enamel relative to the original surface. These results suggest that it would be beneficial to assess ETW severity not just with BEWE but also with the enamel thickness assessment. Future studies should focus on which of the two measures of ETW severity correlates better with the onset of clinical symptoms resulting from ETW progression.

Among the 3D surface texture outcomes, only Sa (roughness) significantly correlated with BEWE_{Buccal} both at baseline and at M12 (Table 2). BEWE_{Buccal} scores were lower at baseline than at M12 (Table 1), which means ETW was less severe at baseline than the latter. Enamel surface roughness (Sa) was also lower at baseline compared to M12 (Table 1), corroborating the results of a previous in situ study that found higher surface roughness in more severe ETW lesions [13]. Several limitations of this exploratory clinical study, however, precluded further differentiation of the degree of surface roughness according to BEWE scores. This could be a focus of future investigations to see if certain thresholds for Sa values could correspond with ETW severity according to BEWE. Both dental microwear texture parameters Sa and Str (anisotropy) increased over time from baseline to M12. The increase in roughness was consistent with the previous study [13], while the Str trend was different as it did not previously change significantly. Asfc did not show any significant change in this study, contrary to our previous in situ study [13], where it significantly increased in the severe ETW group. $\Delta Asfc$, ΔSa , and ΔStr all had a statistically significant but weak correlation with $\Delta BEWE_{Buccal}$ (Table 3), while none of the changes in all three surface texture parameters correlated with the change in enamel thickness. Teeth with an increase in BEWE had significantly greater Δ Sa and Δ Str than teeth that did not show BEWE progression. This was expected as both BEWE and surface texture parameters measure changes on the enamel surface, in contrast to enamel thickness measurements, which relate more to cumulative surface loss. Of the three microwear texture parameters, only roughness (Sa) consistently increased with ETW progression over time in the current and previous in situ study [13]. These results might support the use of the surface roughness (Sa) parameter as another objective measure of ETW progression. Surface texture assessment was originally planned to be conducted at baseline, M3, M6, M9, and M12. However, due to the limitations mentioned earlier, the baseline and M12 measurements were prioritized, with the intermediate time points dropped from this study.

5. Conclusions

Despite the several limitations of this exploratory study to objectively diagnose and monitor ETW clinically, the potential of surface roughness (Sa) as an objective outcome measure to identify ETW lesion severity and progression, comparable with the BEWE index, was confirmed. Moreover, longitudinal monitoring of enamel thickness using CP-OCT may be clinically relevant in addition to ETW severity determination by BEWE and or dental microwear parameters, accepting both tested hypotheses. Future studies resulting in a greater degree of ETW natural progression would be beneficial to further confirm our findings that surface roughness and enamel thickness are potentially suitable and appropriate objective measures for the clinical assessment and monitoring of erosive tooth wear.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/diagnostics13152568/s1, Table S1: Erosive tooth wear (ETW) outcomes (mean and standard deviation) at baseline and 12 months in control subjects.

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Diagnostics 2023, 13, 2568 11 of 12

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References

- 1. Mair, L. Wear in dentistry—Current terminology. J. Dent. 1992, 20, 140–144. [CrossRef] [PubMed]
- 2. Schlueter, N.; Amaechi, B.T.; Bartlett, D.; Buzalaf, M.A.R.; Carvalho, T.S.; Ganss, C.; Hara, A.T.; Huysmans, M.-C.D.; Lussi, A.; Moazzez, R.; et al. Terminology of Erosive Tooth Wear: Consensus Report of a Workshop Organized by the ORCA and the Cariology Research Group of the IADR. *Caries Res.* 2019, 54, 2–6. [CrossRef] [PubMed]
- 3. Lussi, A.; Carvalho, T.S. Erosive Tooth Wear: A Multifactorial Condition of Growing Concern and Increasing Knowledge. In *Erosive Tooth Wear: From Diagnosis to Therapy;* Monographs in Oral Science; Karger Publishers: Basel, Switzerland, 2014; Volume 25, pp. 1–15. [CrossRef]
- 4. Jaeggi, T.; Lussi, A. Prevalence, Incidence and Distribution of Erosion. Monogr. Oral Sci. 2006, 20, 44–65. [CrossRef] [PubMed]
- 5. Mcguire, J.; Szabo, A.; Jackson, S.; Bradley, T.G.; Okunseri, C. Erosive tooth wear among children in the United States: Relationship to race/ethnicity and obesity. *Int. J. Paediatr. Dent.* **2009**, *19*, 91–98. [CrossRef] [PubMed]
- 6. Okunseri, C.; Wong, M.C.M.; Yau, D.T.W.; McGrath, C.; Szabo, A. The relationship between consumption of beverages and tooth wear among adults in the United States. *J. Public Health Dent.* **2015**, *75*, 274–281. [CrossRef] [PubMed]
- 7. Ganss, C.; Lussi, A. Diagnosis of Erosive Tooth Wear. In *Erosive Tooth Wear: From Diagnosis to Therapy*; Karger Publishers: Basel, Switzerland, 2014; Volume 25, pp. 22–31. [CrossRef]
- 8. Peutzfeldt, A.; Jaeggi, T.; Lussi, A. Restorative Therapy of Erosive Lesions. In *Erosive Tooth Wear: From Diagnosis to Therapy*; Karger Publishers: Basel, Switzerland, 2014; Volume 25, pp. 253–261. [CrossRef]
- 9. Hara, A.T.; Zero, D.T. The Potential of Saliva in Protecting against Dental Erosion. In *Erosive Tooth Wear: From Diagnosis to Therapy*; Karger Publishers: Basel, Switzerland, 2014; Volume 25, pp. 197–205. [CrossRef]
- 10. Ramsay, D.S.; Network, O.B.O.T.N.P.; Rothen, M.; Scott, J.M.; Cunha-Cruz, J. Tooth wear and the role of salivary measures in general practice patients. *Clin. Oral Investig.* **2014**, *19*, 85–95. [CrossRef] [PubMed]
- 11. Ungar, P.S.; Sponheimer, M. The Diets of Early Hominins. Science 2011, 334, 190–193. [CrossRef] [PubMed]
- 12. Hara, A.; Livengood, S.; Lippert, F.; Eckert, G.; Ungar, P. Dental Surface Texture Characterization Based on Erosive Tooth Wear Processes. *J. Dent. Res.* **2016**, *95*, 537–542. [CrossRef] [PubMed]
- 13. Hara, A.; Elkington-Stauss, D.; Ungar, P.; Lippert, F.; Eckert, G.; Zero, D. Three-Dimensional Surface Texture Characterization of In Situ Simulated Erosive Tooth Wear. *J. Dent. Res.* **2021**, *100*, 1236–1242. [CrossRef] [PubMed]
- 14. Wilder-Smith, C.H.; Wilder-Smith, P.; Kawakami-Wong, H.; Voronets, J.; Osann, K.; Lussi, A. Quantification of Dental Erosions in Patients with GERD Using Optical Coherence Tomography Before and After Double-Blind, Randomized Treatment with Esomeprazole or Placebo. *Am. J. Gastroenterol.* 2009, 104, 2788–2795. [CrossRef] [PubMed]
- 15. Baumgartner, A.; Dichtl, S.; Hitzenberger, C.; Sattmann, H.; Robl, B.; Moritz, A.; Fercher, A.; Sperr, W. Polarization–Sensitive Optical Coherence Tomography of Dental Structures. *Caries Res.* 1999, 34, 59–69. [CrossRef] [PubMed]
- 16. Chen, Y.; Otis, L.; Piao, D.; Zhu, Q. Characterization of dentin, enamel, and carious lesions by a polarization-sensitive optical coherence tomography system. *Appl. Opt.* **2005**, *44*, 2041–2048. [CrossRef] [PubMed]

Diagnostics **2023**, 13, 2568 12 of 12

17. Romero, M.J.R.H.; Bezerra, S.J.C.; Fried, D.; Yang, V.; Lippert, F.; Eckert, G.J.; Zero, D.T.; Hara, A.T. Cross-polarization optical coherence tomographic assessment of in situ simulated erosive tooth wear. *J. Biophotonics* **2021**, *14*, e202100090. [CrossRef] [PubMed]

- 18. Romero, M.J.R.H.; Bezerra, S.J.C.; Fried, D.; Lippert, F.; Eckert, G.J.; Hara, A.T. Longitudinal assessment of dental erosion-abrasion by cross-polarization optical coherence tomography in vitro. *Braz. Oral Res.* 2022, *in press*.
- 19. Bartlett, D.; Ganss, C.; Lussi, A. Basic Erosive Wear Examination (BEWE): A new scoring system for scientific and clinical needs. *Clin. Oral Investig.* **2008**, *12*, 65–68. [CrossRef] [PubMed]
- 20. Brown, C.A.; Hansen, H.N.; Jiang, X.J.; Blateyron, F.; Berglund, J.; Senin, N.; Bartkowiak, T.; Dixon, B.; Le Goïc, G.; Quinsat, Y.; et al. Multiscale analyses and characterizations of surface topographies. *CIRP Ann.* **2018**, *67*, 839–862. [CrossRef]
- 21. Ungar, P.S.; Brown, C.A.; Bergstrom, T.S.; Walker, A. Quantification of Dental Microwear by Tandem Scanning Confocal Microscopy and Scale-Sensitive Fractal Analyses. *Scanning* **2006**, *25*, 185–193. [CrossRef] [PubMed]
- 22. Schulz, E.; Calandra, I.; Kaiser, T.M. Feeding ecology and chewing mechanics in hoofed mammals: 3D tribology of enamel wear. *Wear* **2013**, *300*, 169–179. [CrossRef]
- 23. Pijpe, J.; Kalk, W.W.I.; Bootsma, H.; Spijkervet, F.K.L.; Kallenberg, C.G.M.; Vissink, A. Progression of salivary gland dysfunction in patients with Sjogren's syndrome. *Ann. Rheum. Dis.* **2006**, *66*, 107–112. [CrossRef] [PubMed]

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