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Evaluation of Floss Remnants After Implant Flossing in Three Different Implant Conditions: A Preclinical Study

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(Article begins on next page)

1 TITLE:

2 Evaluation of Floss Remnants After Implant Flossing in Three Different Implant Conditions: A
3 Preclinical Study

4

5 ABSTRACT AND KEYWORDS:

6 Purpose. The aim of this preclinical study is to evaluate whether implant flossing could leave floss7 residues in three different implant-prosthetic conditions.

Materials and Methods. By mean of an anatomical model three different condition have been 8 9 studied: correct connection between implant and abutment and complete insertion of the implant 10 threads into the plaster (control group); misfit of about 220-230 µm between implant platform and abutment in absence of any threads exposure (misfit group); partial exposure of implant threads 11 12 but absence of misfit (threads group). Twenty-one micro-structured tapered threaded implants were divided among the three groups. Each sample was subjected to a flossing procedure using 13 14 spongy floss, standardized in terms of movement, frequency, time and pressure. Subsequently, a stereomicroscope examination with a standardized magnification of 10x was performed in order to 15 16 highlight the possible presence of floss residues on implant surface.

Results. No floss residue was ever detected for the control group. Both misfit and thread groups
showed floss residues discernible in two different types: microfilaments and amorphous particles.
Statistical analysis showed a significant difference for the presence of floss remnants between the
control group and the other two experimental groups (p = 0.005). No difference was observed
between misfit and threads group.

22	Conclusion. This study shows that exposed threads and misfit can induce the release of floss
23	residues during maintenance procedures.
24	Oral Hygiene, Dental Implants, Implant Flossing, Spongy Floss, Implant maintenance
25	
26	
27	ARTICLE TEXT:
28	INTRODUCTION
29	The clinical use of dental implants has become highly predictable in recent decades, improving the
30	quality of life in patients by restoring both functional and aesthetic support. ^{1,2}
31	Between the high reliability, it is important to realize that not all implants that survive are
32	necessarily successful. Successful implants are those that remain fully functional and healthy within
33	the oral cavity.
34	Peri-implantitis is pathological condition associated with implant failure and is becoming rather
35	prevalent. Several studies have shown that patients with poor plaque control and erratic
36	maintenance display an increase risk of developing peri-implantitis. ^{3,4,5} It is therefore prudent to
37	prevent bacterial colonization by having a very accurate oral hygiene. After implant placement, a
38	strict follow-up regime with a dental professional should be implemented in order to monitor
39	the implant and surrounding teeth for inflammatory disease. ⁶
40	The dental professional should continually encourage the patient to adhere to consistent home oral
41	care in order to prevent any peri-implant, inflammation and in turn increase the success of

42 their implants.⁷

43 It is very important to underline that implant rehabilitation implicates new anatomic conditions.

44 The relation among the implant and surrounding tissues, as well as the prosthetic rehabilitation,

45 require specific considerations about hygienic care.

46 One of the most frequently prescribed hygienic devices for home oral care is the dental floss, in

47 particular the stiffened-end spongy floss is quite commonly used thanks to its adaptability.

48 Narrow interproximal spaces, tilted structures and sub-marginal areas are only some of the

49 anatomical situations where specialists suggest its use.

50 Very few studies have been published supporting this ordinary prescription, generally mutualized by

51 the dental care or supported by strictly personal experience.^{8,9} The evidence regarding implant care

52 is indeed still sparse especially when compared to that for natural teeth.¹⁰

53 Some authors have recently raised some concerns about implant flossing. ^{11,12} Cases of trapped

54 floss fibers around implants with clear signs of peri-implant disease have been reported suggesting

a potential role for the clinical manifestation.

The presence of retained floss fibers could favor plaque retention acting like floss ligatures for 56 experimental peri-implantitis on animal studies.¹³ On the other side, eventual floss remnants can 57 58 also induce a direct immune reaction by the host. Condition that potentially correlates to the periimplantitis theory described by Albrektsson et al.¹⁴ The authors emphasize the primary role of the 59 immune system imbalance in peri-implant marginal bone loss. By this theory, the dental implant is 60 perceived by the immune system as a foreign body, and consequently the presence of additional 61 foreign bodies, such as prosthetic cement or eventually floss remnants can lead to greater bone 62 63 loss.

64 Several physical-mechanical aspects have been implicated in this clinical circumstance about
65 implant flossing, such as the incongruous implant-abutment connectionor the implant rough
66 surface exposure.

According to these considerations, the aim of this in-vitro study was to evaluate whether the generally recommended use of spongy floss around dental implants could represent a dangerous procedure in unexposed or exposed implant surfaces and in implants with a wrong fixture to abutment connection. The null hypothesis is that no difference in remnants on the 3 implantconditions herein evaluated is observed, i.e. the spongy floss works in the same way.

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73

74 METHODS

75 In this in-vitro pilot study twenty-one micro-structured tapered threaded implants with an internal

76 trilobate connection (Replace Select Tapered TiUnite, NP 3.5 x 10 mm, Nobel Biocare®) were

selected. This implant was chosen for its macro and microstructural characteristics, which can be

78 considered representative of the most commonly used implants.

79 Implants were equally and randomly divided into three groups. According to the study group in

80 which they were assigned, implants were differently immersed in dental plaster (picodent[®] quadro-

81 rock[®] plus) contained into plastic boxes (6x5x3 cm).

82 Seven implants with the corresponding abutments correctly inserted were completely fixed in the

- 83 plaster with the exception of the smooth neck (control group) (Fig. 1-A). Seven implants with
- 84 corresponding abutments positioned to create a small misfit (inadequate adaptation of the
- abutment on the implant) of approximately 220-230 μm, with the limit of the stereomicroscope

resolution, were completely fixed in the plaster with the exception of the smooth neck (misfit
group) (Fig. 1-B). Seven implants with a correct implant-abutment connection were fixed in the
plaster leaving four threads exposed (threads group) (Fig. 1-C).

89 Subsequently, the implants of each group were exposed to the cleaning movement of the spongy part of the multifilament floss (Superfloss[®], Oral-B). In particular, the spongy floss was manually 90 91 adapted by forming a loop with the crossed ends at the implant-abutment connection or the 92 eventual exposed implant surface. In this position, the circular criss-cross movement was induced 93 by applying a controlled pressure of 150-200 Newton, which was measured with an appropriate 94 stress and tension gauge (stress and tension gauge 25-250 g, Dentaurum, Germany) connected to 95 one end of the floss. Standardized movements were made both in terms of speed, with a cadence dictated by a professional metronome (metronome Taktell small, Wittner, Germany), and in 96 97 number of movements for a total of 10 tractions in a total time of 10 seconds carried out by a single 98 researcher. Particular attention was placed during the floss movement in not approaching the 99 plaster.

100 After this, in order to evaluate the possible presence or absence of floss remnants, an experimental stereomicroscope analysis (with a standard magnification of 10x) was carried out. The 101 102 stereomicroscope (Carl Zeiss stereomicroscope Stemi 2000-C, FL S configuration with KL 1500 electronic) connected to AXIO CAM MC system was used to collect pictures of eventual remnants of 103 104 spongy floss fibres on the study samples after the simulated hygiene procedures. The outcome of 105 the study, i.e. the floss residues, were dichotomously detected (presence\absence) and described on the basis of their shape. Measurements were performed by using a specific software (ImageJ1). 106 Sample size 107

108	A pilot sample of 3 implants reproducing the experimental conditions was used to estimate the						
109	non-inferiority margin.						
110	At a significance level α = 0.05 for a one-sided test, an 80% power, a non-inferiority margin equal to						
111	40% and an allocation ratio equal to 1:1, a total sample size of 14 implants was needed (7 in each of						
112	the two experimental groups). A control group of 7 implants was also created.						
113							
114	Statistical Analysis						
115	Cross tabulations were used to describe the observed results. Chi-square test and Fisher test were						
116	performed aiming to evaluate differences in material presence respectively among the three groups						
117	and between floss and amorphous floss particles in the two experimental groups. The level of						
118	significance α was a priori set at 0.05.						
119							
120							
121	RESULTS						
122	After flossing, all implants were analysed (Table 1).						
123	All dental implants in control group did not show any floss remnants both on the smooth surface of						
124	the neck than next to the implant-abutment connection.						
125	Misfit group implants showed the presence of floss material remnants in 85.7% of cases. In						
126	particular, in the misfit space, spongy floss microfilaments were found only in 1 implant, another 1						
127	implant showed amorphous particles (ranging from 50 to 600 μ m), whereas 4 implants presented						

both microfilaments and amorphous particles (Fig. 2). Only 1 implant was free of any spongy flossmaterial.

130	Threads group showed remnants of spongy floss in 57.1% of the implants. In particular, spongy						
131	floss microfilaments were observed in 1 implant, whereas amorphous particles were present in						
132	other 2 (Fig. 3). Both microfilaments and amorphous particles of spongy floss fibres were observed						
133	on 1 implant. Differently, no microfilaments or amorphous particles were detectable on exposed						
134	surfaces, around the smooth neck or at the abutment-implant connection of 3 implants.						
135	The observed differences about spongy floss remnants (microfilaments and amorphous particles)						
136	among the three groups were statistically significant (Chi-square test =10.691, p=0.005). The null						
137	hypothesis was therefore rejected.						
138	No significant difference was found between samples of misfit and threads groups (Fisher's exact						
139	Test shows a value of $p = 0.2861$).						
140							
140							
140	DISCUSSION						
	DISCUSSION From the present study no floss remnants have been detected on control group implants.						
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141 142 143 144	From the present study no floss remnants have been detected on control group implants. Therefore, presupposing a potential pathogenicity of this remnants to peri-implant surrounding tissues, the use of spongy floss seems to represent a safe procedure in a "regular state" implant.						

148 inconsistency and anatomical impediment. In particular, it is represented by the incomplete

insertion of the abutment to the implant, as well as the absence of an ideal matching between theabutment and the prosthetic crown.

151 In order to limit the misfit occurrence, a combination of visual, tactile and radiographic

152 examinations should be performed.¹⁵

153 The scientific literature about implant-supported fixed denture reports a wide range of values (from

154 10µm to 160µm) to consider the misfit technically acceptable. However, as clearly stated by two

recent systematic reviews,^{15,16} it is still lacking the effective role of misfit on clinical outcome.

156 Therefore, the extent at which it is considered tolerable remains unclear.

157 The present work used a standardized value of 220-230 μm and put in evidence a new negative role

158 of the misfit. It is quite rational to think that, not only the presence of it, but also its dimension can

influence on the effective ability to trap floss remnants. Further studies focused on different gap

160 dimensions are consequently strongly recommended.

161 Threads exposure is the other clinical aspect investigated in this study. Its presence can be a

162 consequence of either para-physiological or pathological conditions. The eventual bone remodelling

163 (following implant placement)¹⁷ as well as the bone resorption (caused by inflammatory peri-

164 implantitis or foreign body response) ^{14,18} are the prevalent conditions.

165 Irrespective by its origin, the threads exposure is also the result of a clinical condition able to trap

166 floss remnants. In this specific circumstance, macro and micro-texture of implant can play a role on

the final result. It is therefore rational to suggest future investigations on this field.

168 In the present work a clear distinction between two kinds of floss remnants has been made:

amorphous particles and microfilaments. This distinction comes for the assumption that the two

170 forms of residues may have different clinical implications. The first consideration is that amorphous

171 particles can be more prone to a spontaneous expulsion in respect to microfilaments. To support

the above consideration, it is necessary to remind that all clinical cases up to date described were
characterized by the detection of filamentous remnants.^{11,12} On the other hand, it must be taken
into account that this is probably due to the fact that microfilaments display a more direct and
intuitive correlation to the dental floss. Consequently, it is not known which is the real role of this
particles, and investigations about this topic are strongly advocated.

177 In accordance to the manufacturer statements, the spongy floss used for this research is composed178 by nylon fibres covered by coloured and aromatized wax.

Amorphous particles herein detected could probably correspond to wax remnants, and this
consideration is supported by the findings of van Velzen et al. (2016).¹² In a preclinical set, the
authors analysed the implant surface after flossing and detected the presence of organic material
through spectrophotometry, plausibly wax.

In this study, where the prevalence of microfilament and amorphous particles between the two investigated conditions has been observed, it could be speculated that flossing on exposed threads may tend to release more amorphous residues. Otherwise, sliding floss against misfit may tend to cause tearing and the consequent release of microfilaments. The absence of a statistically supported difference limits any further consideration, but it can be rational to deepen this aspect in further studies on broader samples. Other aspects that advocate for future investigations are the floss typology and the flossing movement.

To the best of our knowledge, the etiopathogenetic role of these residues has not yet been
discovered. However, the presence of trapped floss remnants in subgingival area may be a fertile
environment for bacterial invasion and proliferation. From this consideration, it can be assumed
that dental floss residues can promote the retention of bacteria and have an action quite similar to
that of experimental ligatures¹³ or biofilm-retaining factor like are luting cementum residues.^{19,20}

Among the pathogenic hypothesis, it can't be neglected the host response to those remnants acting
as foreign bodies. This aspect finds an interesting correlation with the peri-implantitis theory
described by Albrketsson et al.¹⁴

From the cases reported on the literature, it can be observed that the removal of these residues 198 generally leads to clinical improvements. Van Velzen et al. (2016)¹² report that in the last 3 years of 199 200 clinical practice they have encountered 10 patients with persistent peri-implantitis, previously 201 treated with a non-surgical and surgical protocol for the maintenance of implants (to which all of their patients with peri-implantitis are treated). Among those 10 patients, after exploratory surgery, 202 203 all implants showed the presence of remnants of dental floss adhering to the rough part of them. Interestingly, after a careful removal and debridement of the peri-implant site, nine of ten cases 204 resulted in an improvement in peri-implant conditions 6 months later (i.e. absence of bleeding on 205 206 probing and reduction of probing pocket depth).

These results are supported by long-term observations of a clinical case report describing a periimplantitis treated with an endoscopic access. ¹¹ Trapped filaments around several implants were thoroughly removed leading to a complete resolution of the peri-implantitis with a 6-years stable result.

211 CONCLUSION

This preclinical study has clearly demonstrated how implant treads exposure and abutment-implantmisfit can both favour the release of remnants from a spongy floss.

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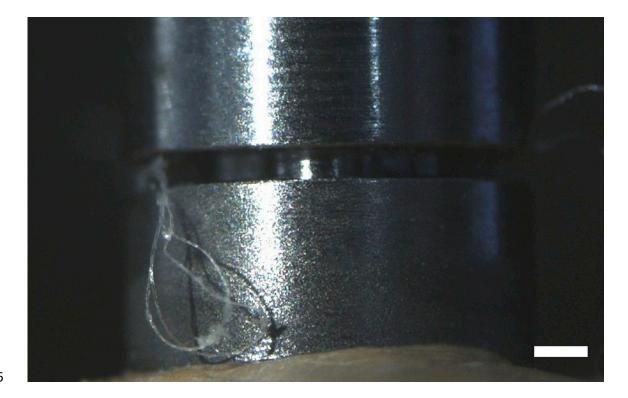
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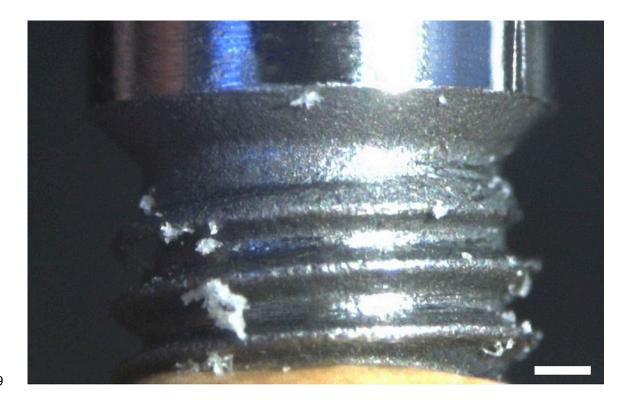
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- Fig. 1 Image of the three implant conditions studied that reproduce A) control group; B) misfit
- 285 group; C) threads group



- Fig. 2 Microscopic image (10x) showing both microfilaments and amorphous particles at the misfit
- 288 level. White bar = 400 μ m



- 290 Fig. 3 Microscopic image (10x) showing amorphous particles on the exposed threads. White bar =
- 400 μm
- **TABLE**:
- Table 1. Presence/absence percentage of spongy floss material in the three groups.

			Material		
			Presence	Absence	—Total
Group	Control	Count	0	7	7
		% in group	0.0%	100.0%	100.0%
	Misfit	Count	6	1	7
		% in group	85.7%	14.3%	100.0%
	Threads	Count	4	3	7
		% in group	57.1%	42.9%	100.0%