Reporting quality of randomized controlled trials published in prosthodontic and implantology journals

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Reporting quality of randomized controlled trials published in prosthodontic

and implantology journals

Running head: Reporting quality of RCTs in prosthodontics and implantology **Article Category:** Original article

D. KLOUKOS^{*}, S.N. PAPAGEORGIOU[†], I. DOULIS[‡], H. PETRIDIS[§] & N. PANDIS[¶]

Author affiliations

*Department of Orthodontics and Dentofacial Orthopedics, Faculty of Medicine, University of Bern, Bern, Switzerland,

[†]Department of Orthodontics, School of Dentistry, University of Bonn, Bonn, Germany; Department of Oral Technology, School of Dentistry, University of Bonn, Bonn, Germany; Clinical Research Unit 208, University of Bonn, Bonn, Germany,

[‡]Department of Orthodontics and Dentofacial Orthopedics, 251 Hellenic Air Force V.A. General Hospital, Athens, Greece,

[§]Department of Restorative Dentistry, UCL Eastman Dental Institute, London, UK,

[®]Department of Orthodontics and Dentofacial Orthopedics, Faculty of Medicine, University of Bern, Bern, Switzerland; Private Practice, Corfu, Greece

CORRESPONDENCE Dr. Haralampos Petridis, Program Director, Master in Conservative Dentistry, Senior Clinical Lecturer, Department of Restorative Dentistry, UCL Eastman Dental Institute, Prosthodontic Unit, 256 Gray's Inn Road, London WC1X 8LD, UK.

E-mail: c.petridis@ucl.ac.uk

SUMMARY The purpose of this study was to examine the reporting quality of randomized controlled trials (RCTs) published in prosthodontic and implantology journals. Thirty issues of 9 journals in prosthodontics and implant dentistry were searched for RCTs, covering the years 2005-2012: The Journal of Prosthetic Dentistry, Journal of Oral Rehabilitation, The International Journal of Prosthodontics, The International Journal of Periodontics & Restorative Dentistry, Clinical Oral Implants Research, Clinical Implant Dentistry & Related Research, The International Journal of Oral & Maxillofacial Implants, Implant Dentistry and Journal of Dentistry. The reporting quality was assessed using a modified CONSORT statement checklist. Data were analyzed using descriptive statistics followed by univariable and multivariable examination of statistical associations (α =0.05). A total of 147 RCTs were identified with a mean CONSORT score of 69.4 (SD = 9.7). Significant differences were found among journals with the Journal of Oral Rehabilitation achieving the highest score (80.6, SD= 5.5) followed by Clinical Oral Implants Research (73.7, SD= 8.3). Involvement of a statistician/methodologist was significantly associated with increased CONSORT scores. Overall, the reporting quality of RCTs in major prosthodontic and implantology journals requires improvement. This is of paramount importance considering that optimal reporting of RCTs is an important prerequisite for clinical decision-making.

KEYWORDS: randomized clinical trials, prosthodontics, dental implants

Introduction

Reliable evidence is more likely consequent to sound design and methodology (1, 2). Among the various study designs the randomized controlled trial (RCT) is considered as the "gold standard" for assessing the effectiveness and safety of medical interventions. Nevertheless, RCTs are also prone to inadequacies and there is a substantial body of evidence in the biomedical literature, which indicates that the quality of many RCTs is suboptimal (3-6).

Accurate and transparent reporting of RCTs is prerequisite for the assessment of their internal validity and the clinical translation of their results (7). In an effort to improve and standardize reporting of RCTs the CONSORT (Consolidated Standards of Reporting Trials) guidelines were developed by the CONSORT group and are continuously being updated. The main CONSORT document consists of 25 items and sets standards on how and what should be included in an RCT report (2).

The CONSORT guidelines have been endorsed by over 580 journals (8) and there is evidence of a positive impact on RCT reporting (9). In dentistry reporting quality of RCTs has been assessed in a number of general and dental specialty journals (10-18), indicating that there is room for improvement. However, there is a lack of studies comparatively evaluating the completeness of reporting of recently published RCTs in prosthodontic and implantology using the CONSORT guidelines (2).

Therefore, the primary objective of this study was to evaluate the completeness of reporting of RCTs in prosthodontic and implantology journals using the CONSORT statement. A secondary aim was to identify factors associated with better reporting of RCTs.

Materials and methods

Four dental journals with emphasis on prosthodontics (*The International Journal of Prosthodontics*, *Journal of Oral Rehabilitation, The Journal of Prosthetic Dentistry, The International Journal of Periodontics & Restorative Dentistry*), 4 dental implantology journals (*Clinical Implant Dentistry & Related Research, Clinical Oral Implants Research, Implant Dentistry, The International Journal of Oral & Maxillofacial Implants*) and 1 general dental journal with a predilection for prosthodontics (*Journal of Dentistry*) were included in the study. The selected journals had the highest impact factors of prosthodontic and implant dentistry journals based on 2009 data.

The contents of 30 issues of each journal from June 2012 backwards were searched for RCTs on humans. Supplemental issues were included in the search, but were not counted as an issue. Initially the abstract was read and any trials that were clearly RCTs were included. Other articles that used

terminology in the title or abstract such as "prospective", "comparative", "efficacy" or an indication was given that a comparison of treatment groups was assessed prospectively, were further investigated to examine whether randomization was implemented. Studies that did not involve humans and studies, where it was concluded that no true randomization was implemented, were excluded. Screening and selection of studies were conducted independently by two authors (DK, SNP).

The information extracted from each article included journal and year of publication, region of publication (Europe, Americas or other region, based on the first author), ethical approval, statistical significance of main finding, number of authors, involvement of a statistician or methodologist, and whether the study was single- or multicenter. Involvement of a statistician or methodologist was ascertained by checking author affiliations (public health or epidemiology departments were considered as providing statistical assistance), author degrees (where provided), and information in the methods or acknowledgement sections of each paper.

A modified CONSORT checklist as presented by Tiruvoipati *et al.* (6) was used to evaluate the reporting completeness of RCTs. This checklist has 30 questions related to the CONSORT items excluding the first item of the CONSORT checklist (title and abstract), since the authors have to follow the instructions of the journal in preparing the abstract. The given score per item ranged from 1 to 3, with 1=no description, 2=inadequate description and 3=adequate description. The scores for the 30 items were added, and a percentage score was calculated for each trial, whereas non- applicable items were not scored. A trial with adequate descriptions (score 3) for all items would receive a score of 90. All scores were converted to a percentage scale and therefore a score of 90 was equivalent to 100% in the percentage scale. When non-applicable items were identified (for example inability to blind the treatment provider) only the applicable items were considered for the calculation of the percentages. Therefore, a trial with only 28 applicable items, but adequate descriptions (score 3) for these, would receive a maximum score of 84, corresponding to a percentage of 100%.

Each RCT was also scored using the Jadad scale (19), allocating trials a score between zero (very poor) and five (rigorous). The Jadad scale includes three questions and each one of them is answered with either yes (1 point) or no (no point): (1) "Is the study described as randomized?"; (2) "Is the study described as double blinded?"; (3) "Is there a description of withdrawals and dropouts?". Two additional points, to reach a maximum score of 5, are given (i) if the method of randomization is clearly described and appropriate or (ii) if the method of blinding is clearly described and appropriate. One or

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two points are subtracted if the method of randomization or the method of blinding is described, but is inappropriate.

Each included RCT was scored independently by 2 authors (DK, SNP), and subsequently results were compared and modified in order to arrive to a mutually agreed score. Discrepancies between the 2 authors (DK, SNP) were resolved by discussion. Before data extraction, a calibration exercise was performed between the two authors responsible for it (DK, SNP) with 80 randomly selected studies. Inter-rater agreement was evaluated for all extracted data with Cohen's kappa and any disagreements were resolved with discussion.

Statistical Analysis

Descriptive statistics were calculated for the modified CONSORT scores and tabulated by trial characteristics. The modified CONSORT scores were approximately normally distributed. Data were analyzed through linear regression modeling; univariable analysis was utilized to determine articles' characteristics associated with the modified CONSORT scores, whereas multivariable analysis was employed to adjust for possible confounders. A two-tailed P-value of 0.05 was considered statistically significant with a 95% confidence interval. Analyses were performed with the STATA[®] version 13.0 software (Stata Corporation, College Station, Texas, USA).

Results

From the 3667 articles that were originally screened 3520 were excluded for not adhering to the inclusion criteria, leaving 147 RCTs for detailed assessment (Appendix 1). Inter-rater agreement was found to be excellent (kappa 0.88, 95% CI: 0.87-0.89). The included RCTs reported on a wide selection of topics ranging from surgical implant procedures and techniques, survival of implants and prostheses, biological responses, clinician's perspective of esthetics and patient satisfaction.

Table 1 displays the 147 RCTs tabulated by their characteristics. The journals contributing with the most RCTs were *Clinical Oral Implants Research* (31.3%), followed by *The International Journal of Oral and Maxillofacial Implants* (16.3%) and *Journal of Dentistry* (14.3%). The majority of RCTs originated from Europe (58.5%), were approved by an ethical committee (72.1%) and reported statistically significant findings (59.2%). Concerning the number of authors, most RCTs included four to six authors (64.6 %), whereas a statistician/methodologist was involved in 37.4% of the RCTs. Finally, the majority of RCTs were a multi-center effort (71.4%).

Table 2 displays the scores for all items on the modified CONSORT checklist. Description of pre-study sample size calculation was absent in the majority of the trials (64.0%). No description of the random number generation was also evident in 32.7% of the trials. Allocation concealment was not reported in 61.9% of the studies, whereas details of personnel involved in sequence allocation, enrollment, and assignment were not described in 51.0% of those. Blinding of participants and treatment providers was not reported in 36.7% and 37.4% of the sample respectively. No description of blinding of assessors and analysts reached 61.9% and 89.8% respectively. Absence of a flow chart describing patient numbers at different stages of a study was apparent in 76.9% of the sample. Trial limitations and generalizability of the trial results were not reported in 55.8% and 52.4% of the studies respectively.

The modified CONSORT scores per study characteristic are presented in Table 3. The highest modified CONSORT score was found for the *Journal of Oral Rehabilitation* (80.6%), which, however, contributed only 2 RCTs, followed by the *Clinical Oral Implants Research* (73.7%). The highest modified CONSORT scores chronologically were found in the years 2010-2011. Increased CONSORT scores were also found for RCTs with ethical approval and RCTs with involvement of a statistician/methodologist.

Table 4 presents the results of the univariable and multivariable linear regression analyses. In the univariable analysis, the journal of publication and the involvement of a statistician/methodologist were significantly associated with the CONSORT scores. Similar associations were observed in the multivariable analysis.

Table 5 displays the Jadad scores for the 147 RCTs tabulated by journal. The median Jadad scores ranged from 1.0 (*Journal of Prosthetic Dentistry*) to 3.5 (*Implant Dentistry*).

Discussion

In this study the reporting quality of RCTs in the fields of prosthodontics and implant dentistry was assessed using a modified CONSORT statement (6) and the Jadad scale (19). The mean modified CONSORT scores ranged from 60.9% to 80.6 % among the journals included in the study, a finding similar to the scores reported in medical journals (6,20). The Jadad score ranged from 1.0 to 3.5; this finding is comparable to other fields in medicine (6). Although all quality score scales have inherent limitations and caution should be used when evaluating reporting quality, the overall score indicates that there is room for improvement.

Pre-study sample size calculation is an important part of designing a trial, and guards against underpowered trials that may result in waste (21-24). In the present study 64.0% of the RCTs did not report sample size calculation at all, while, 8.2% of them reported it inadequately. Chan and Altman (25) reported that 73% of the 519 medical trials published in PubMed in December 2000 did not report sample size calculation. It seems that problematic reporting of pre-study sample size calculations in RCTs is a common finding in the literature (11, 14, 26-30). Trials with insufficient sample size can be considered unethical, wasteful (21-24) and less credible compared to trials of sufficient size.

The reporting of the randomization process should, ideally, include details about both the methods used to generate the random allocation sequence and any restrictions used during the process. Terms such as "patients were randomly assigned" or "two groups were formed at random" are considered inadequate. The current study showed that the generation of the unpredictable allocation sequence was reported inadequately in 11.6% of the cases or not at all in 32.7% of the cases. Altman and Dore (31) studied 80 medical trials published in four leading medical journals and concluded that in 30% of trials there was no clear evidence that the groups had been randomised.

In dentistry, Montenegro *et al.* (15) found that only 17% of the trials published in periodontal journals reported the randomization process adequately. Koletsi *et al.* (32) found that from 112 clinical trials in the orthodontic literature labeled as RCTs, only 29.5% were indeed identified as RCTs based on clear descriptions of appropriate random number generation.

Allocation concealment ensures that neither the investigators nor the patients know which treatment the next patient will be allocated to and guards against confounding. Although allocation concealment is always feasible, the results showed that 61.9% of the included RCTs did not report allocation concealment at all, while 17.7% of them reported it inadequately. These results are in accordance with previous studies; Pandis *et al* (18) reported 22% adequacy in reporting allocation

concealment among dental journals and Montenegro *et al.* (15) only 7% among three periodontal journals.

Another key element in RCT reporting is the description of blinding. Blinding is important to the validity of a trial, as it prevents performance and detection bias, and protects the sequence after allocation. Often the concepts of allocation concealment and blinding are confused. Blinding is especially important for subjective outcomes (e.g. pain scores), as these are more prone to bias. Blinding of the patients and the treatment providers may not always be possible, however, blinding of the assessors and the analysts is (33, 34). The results of this study showed that blinding of the various groups was not reported at all in 36.7% to 89.8% of the cases. Pandis *et al.* (18) using a similar scale reported adequate description of blinding in RCTs published in leading dental journals in the range of 0 to 26%.

Statistical methods used for data analysis were not described in 6.1% whereas 59.2% of the RCT reports provided an adequate description. These results are similar to a previous assessment in dentistry, which reported that 3% of the studies provided no description and 51% provided adequate description of statistical methods (18). Analyses should be pre-specified and ideally described in the trial protocol. Pre-specification allows for the assessment of selective reporting and data driven analysis which can be misleading. A common statistical pitfall is the conduct of multiple tests, which leads to increased type I error (false positive) that can be misleading when associated with selective reporting. It is recommended that subgroup analyses should be pre-specified and kept to the minimum (35, 36). The results of the present study showed that 30.6% of the trials did not describe how this issue was handled, while 34.0% of the reports described it adequately. Pocock *et al.* (37) studied 45 medical trials published in three high impact factor medical journals and they reported that multiple endpoints were analyzed without being pre-specified as primary endpoints.

Finally, in the present study 6.1% and 80.2% of the trials lacked complete description of estimates and confidence intervals, respectively. Previous studies found inadequate results' reporting in leading medical journals (37). Pandis *et al.* (18) found that dental trials also suffered from problematic reporting in this area of interest, with lack of description in 3% and 80% of the studied trials and adequate description in 62% and 20% of them for the complete reporting of the results and for the reporting of confidence intervals respectively. Reporting of estimates and confidence intervals facilitates interpretation in relation to clinical importance. P-values and statistical significance are based on arbitrary cut-off points (i.e. 0.05) and are sensitive to sample size and variance. Small P-

values are often misinterpreted as showing a clinically important effect and vice versa as trivial and clinically unimportant differences can be statistically significant when sample size is large and variance is small (38).

This study is not free of limitations. A limitation might be that the scoring of trials is always susceptible to some degree of subjectivity. Nevertheless, considerable efforts were made to compensate for inter-rater subjectivity by calibration exercises before study commencement and strict adherence to applied CONSORT guidelines. RCT assessment was limited to high impact factor prosthetic journals and therefore published RCTs in lower impact factor journals or even non-published RCTs were excluded. However, we believe that the selected journals constitute a representative or best case scenario sample of the reporting status in the specialty. It should be, also, underlined that incomplete reporting of trials does not necessarily infer low quality of conducting or false methodology (39). Researchers might have designed and conducted a study ideally, but they might have omitted reporting accurately all stages and aspects of their trial due to, for instance, space limitations. Even though RCTs are pivotal for evidence-based dentistry and medicine, they are not free of shortcomings. It is important that they are designed properly, implemented and reported well.

Numerous journals have adopted the CONSORT guidelines and very few have implemented active compliance. The American Journal of Orthodontics and Dentofacial Ortopedics, for instance, has recently implemented a novel approach which includes assessment of compliance at the editorial level and specific recommendations for the authors in order to improve RCT reporting. A preliminary study indicated that this approach has increased dramatically reporting quality (40). In addition the journal has recently adopted a structured report which diverges from the standard IMRaD (Introduction, Methods, Results, Discussion) structure and includes 17 subheadings that lead the report (41). Similar initiatives have been proposed elsewhere (42). The results of the present study indicate that adherence to the CONSORT statement of RCTs in major prosthodontic and implantology journals can be improved.

Conflict of interest

No funding was obtained for the current study. No conflict of interest declared.

Tables

Table 1. Characteristics of the	ne 147 included randomized controlled trials
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Characteristic	Category	Ν	%
Journal	Clin Implant Dent Relat Res	13	8.8
	Clin Oral Implants Res	46	31.3
	Implant Dent	4	2.7
	Int J Oral Maxillofac Implants	24	16.3
	Int J Periodontics Restor Dent	15	10.2
	Int J Prosthodont	20	13.6
	J Dent	21	14.3
	J Oral Rehabil	2	1.4
	J Prosthet Dent	2	1.4
Year	2007	4	2.7
	2008	12	8.2
	2009	13	8.8
	2010	41	27.9
	2011	42	28.6
	2012	35	23.8
Continent	Europe	86	58.5
	Americas	30	20.4
	Asia/Other	31	20.4
		51	21.1
Ethics committee approval	No	41	27.9
	Yes	106	72.1
Statistical significance of main finding	No	60	40.8
	Yes	87	59.2
Number of authors	<4	35	23.8
	4-6	60	40.8
	6≤	52	35.4
Statistician/methodologist involvement	No	92	62.6
	Yes	55	37.4
Number of centers	Single-center	42	28.6
	Multicenter	105	71.4

 Table 2. Distribution of consensus scores for the items in the modified CONSORT checklist (n=147)

Item	No description – n (%)	Inadequate – n (%)	Adequate – n (%)
1. Justification for the trial for the trial	15 (10.2)	16 (10.9)	116 (78.9)
2.Explicit definition of eligibility criteria	9 (6.1)	26 (17.7)	112 (76.2)
3.Detailed description of setting/location of recruitment and data collection	21 (14.3)	27 (18.4)	99 (67.3)
4.Details of intervention studied	0 (0.0)	14 (9.5)	133 (90.5)
5. Clear statement of hypothesis or objectives	2 (1.4)	29 (19.7)	116 (78.9)
6.Identification and definition of outcome measures	2 (1.4)	24 (16.3)	121 (82.3)
7.Description of pre-study sample size calculation	94 (63.9)	12 (8.2)	41 (27.9)
8.Description of the generation of unpredictable allocation sequence	48 (32.7)	17 (11.6)	82 (55.8)
9. Details of any restriction used in randomization	85 (57.8)	13 (8.8)	49 (33.3)
10.Description of allocation concealment	91 (61.9)	26 (17.7)	30 (20.4)
11.Details of personnel involved in sequence allocation, enrollment, and assignment	75 (51.0)	35 (23.8)	37 (25.2)
12.Details of blinding of participants	54 (36.7)	9 (6.1)	84 (57.1)
13.Details of blinding of treatment providers	55 (37.4)	4 (2.7)	88 (59.9)
14.Details of blinding of assessors	91 (61.9)	18 (12.2)	38 (25.9)
15.Details of blinding of analysts	132 (89.8)	6 (4.1)	9 (6.1)
16.Details of measurement of success of blinding	143 (97.3)	1 (0.7)	3 (2.0)
17.Description of statistical methods	9 (6.1)	51 (34.7)	87 (59.2)
18.Flow chart describing patient numbers at different stages	113 (76.9)	6 (4.1)	28 (19.1)
19.Clear description of protocol deviations	55 (37.4)	19 (12.9)	73 (49.7)
20.Description of dates of recruitment	89 (60.5)	8 (5.4)	50 (34.0)
21.Details of follow-up	12 (8.2)	17 (11.6)	118 (80.3)
22.Description of baseline characteristics	19 (12.9)	64 (43.5)	64 (43.5)
23.Reporting of intention-to-treat principle	135 (91.8)	3 (2.0)	9 (6.1)
24.Complete reporting of results	9 (6.1)	16 (10.9)	122 (83.0)
25.Reporting of confidence intervals	118 (80.3)	2 (1.4)	27 (18.4)
26.Multiple testing and corrections	45 (30.6)	52 (35.4)	50 (34.0)
27.Description of side effects/adverse effects	33 (22.5)	19 (12.9)	95 (64.6)
28. Trial limitations and weaknesses	82 (55.8)	31 (21.1)	34 (23.1)
29.External validity of trial results	77 (52.4)	49 (33.3)	21 (14.3)
30.Literature review	1 (0.7)	9 (6.1)	137 (93.2)

Clin Implant Dent Rel Res (n=13) Clin Oral Implants Res (n=46) Implant Dent (n=4) Int J Oral Maxillofac Implants (n=24) Int J Periodontics Restor Dent (n=15) Int J Prosthodont (n=20) J Dent (n=21) J Oral Rehabil (n=2) J Prosthet Dent (n=2) 2007 (n=4) 2008 (n=12) 2009 (n=13) 2010 (n=41) 2011 (n=42)	64.1 73.7 65.8 71.0 60.9 68.0 69.3 80.6 62.2 70.0 64.8 67.0	5.8 8.3 4.5 10.6 9.3 10.9 7.5 5.5 12.6 11.4 10.1
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U Dent (n=21) U Oral Rehabil (n=2) U Prosthet Dent (n=2) 2007 (n=4) 2008 (n=12) 2009 (n=13) 2010 (n=41)	69.3 80.6 62.2 70.0 64.8 67.0	7.5 5.5 12.6 11.4 10.1
U Dent (n=21) U Oral Rehabil (n=2) U Prosthet Dent (n=2) 2007 (n=4) 2008 (n=12) 2009 (n=13) 2010 (n=41)	80.6 62.2 70.0 64.8 67.0	5.5 12.6 11.4 10.1
V Oral Rehabil (n=2) V Prosthet Dent (n=2) 2007 (n=4) 2008 (n=12) 2009 (n=13) 2010 (n=41)	62.2 70.0 64.8 67.0	5.5 12.6 11.4 10.1
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2008 (n=12) 2009 (n=13) 2010 (n=41)	64.8 67.0	10.1
2009 (n=13) 2010 (n=41)	67.0	
2010 (n=41)		11.0
		11.9
	70.5	10.0
	70.5	8.6
2012 (n=35)	69.1	9.5
Europa (n=86)	69.6	10.1
	69.8	8.8
Asia/Other (n=31)	68.4	9.6
No (n=41)	67.5	10.0
Yes (n=106)	70.1	9.5
No (n=41)	70.1	9.7
Yes (n=106)	68.9	9.7
<4 (n=35)	70.4	9.8
4≤n< 6 (n=60)	68.9	10.0
≤6 (n=52)	69.2	9.4
$N_{\rm D}$ (p=02)	68 1	8.3
		8.5 11.5
res (n=33)	/1.1	11.5
Single-center (n=42)	68.3	9.1
		9.9
Total (n=147)	69.4	9.7
	2011 (n=42) 2012 (n=35) Europa (n=86) Americas (n=30) Asia/Other (n=31) No (n=41) Ves (n=106) Ves (n=106) Ves (n=106) Ves (n=52) No (n=92) Ves (n=55)	2011 (n=42) 70.5 $2012 (n=35)$ 69.1 Suropa (n=86) 69.6 Americas (n=30) 69.8 Asia/Other (n=31) 68.4 No (n=41) 67.5 Yes (n=106) 70.1 No (n=41) 70.1 Yes (n=106) 70.1 No (n=41) 70.1 Yes (n=106) 68.9 $54 (n=35)$ 70.4 Yes (n=52) 69.2 No (n=92) 68.4 Yes (n=55) 71.1 Single-center (n=42) 68.3 Aulticenter (n=105) 69.8

Table 3. Modified CONSORT scores of the 147 included randomized controlled trials

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Table 4. Univariable and multivariable linear regression-derived coefficients (β) and 95% Confidence Intervals (CIs) for

modified CONSORT score as dependent variable for the 147 included randomized controlled trials

		Univariable analysis			Multivariable analysis			
		β	95% CI	Р	β	95% CI	Р	
Journal		•						
	Clin Implant Dent Relat Res	3.21	(-3.53, 9.95)	0.35	2.75	(-3.96,9.46)	0.42	
	Clin Oral Implants Res	12.76	(7.47,18.05)	< 0.001	12.42	(7.08,17.76)	< 0.00	
	Other	7.72	(-0.06,15.51)	0.05	8.37	(0.62, 16.11)	0.04	
	Int J Oral Maxillofac Implants	10.08	(4.23, 15.94)	0.001	10.21	(4.27, 16.15)	0.001	
	Int J Prosthodont	7.11	(1.04, 13.19)	0.02	6.37	(0.28, 12.46)	0.04	
	J Dent	8.26	(2.25, 14.28)	< 0.01	8.21	(2.12, 14.32)	< 0.01	
Year								
		0.64	(-0.57,1.85)	0.30	0.55	(-0.66,1.75)	0.37	
Country								
	Europe	1.21	(-2.82,5.25)	0.55				
	America	1.39	(-3.54,6.32)	0.56				
	Asia/Other	Baseline						
	Asia/Other	(reference)						
Ethics committee								
approval								
	No	Baseline						
	NO	(reference)						
	Yes	2.56	(-0.95,6.07)	0.15	0.42	(-3.02,3.86)	0.81	
Statistical								
significance of								
main finding								
	No	1.17	(-2.05,4.39)	0.47				
	Yes	Baseline						
	105	(reference)						
Number of authors								
	<4	1.44	(-2.66,5.53)	0.49				
	4-6	Baseline						
		(reference)						
	6≤	0.26	(-3.38,3.91)	0.87				
Statistician/								
methodologist								
involvement								
	No	Baseline						
		(reference)						
	Yes	2.77	(-0.48,6.01)	0.09	3.44	(0.29,6.59)	0.03	
Number of centers								
	Single-center	Baseline						
	•	(reference)						
	Multicenter	1.54	(-1.96,5.04)	0.39				

N/A

Table 5. Descriptive statistics for the Jadad score of the 147 included RCTs by journal

Journal (n)	Median	IQR	
Clin Implant Dent Relat Res (n=13)	3.0	1.0	
Clin Oral Implants Res (n=46)	3.0	1.0	
Implant Dent (n=4)	3.5	2.0	
Int J Oral Maxillofac Implants (n=24)	3.0	2.0	
Int J Periodontics Restor Dent (n=15)	2.0	1.0	
Int J Prosthodont (n=20)	2.0	1.0	
J Dent (n=21)	3.0	0.0	
J Oral Rehabil (n=2)	3.0	2.0	
J Prosthet Dent (n=2)	1.0	2.0	

3.0 2.0 1.0 2.0 23-Q1).

Appendix 1. List of scored papers

ID	<mark>Yea</mark> r	Journal	<mark>Yea</mark> r	<mark>lssue(volum</mark> e)	First Author	<mark>Jada</mark> d	CONSO RT	CONSOR <mark>%</mark>
<mark>18</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(5)</mark>	<mark>Sisti A</mark>	4 4	<mark>67</mark>	74.4
<mark>22</mark>	201 2	Clin Oral Implants Res	<mark>201</mark> 2	<mark>23(5)</mark>	Canullo L	<mark>4</mark>	<mark>58</mark>	<mark>64.4</mark>
<mark>30</mark>	201 2	Clin Oral Implants Res	<mark>201</mark> 2	<mark>23(5)</mark>	Van Assche N	<mark>3</mark>	<mark>66</mark>	<mark>73.3</mark>
<mark>31</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(5)</mark>	Quirynen M	<mark>3</mark>	<mark>65</mark>	<mark>72.2</mark>
<mark>32</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(5)</mark>	Chongcharoen N	<mark>3</mark>	<mark>66</mark>	<mark>73.3</mark>
<mark>38</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(4)</mark>	Trombelli L	<mark>5</mark>	<mark>69</mark>	<mark>76.7</mark>
<mark>46</mark>	<mark>201</mark> 2	Clin Oral Implants Res	201 2	<mark>23(4)</mark>	Krennmair G	<mark>2</mark>	<mark>60</mark>	<mark>66.7</mark>
<mark>47</mark>	201 2	Clin Oral Implants Res	<mark>201</mark> 2	<mark>23(4)</mark>	Romano M	<mark>5</mark>	<mark>61</mark>	<mark>67.8</mark>
<mark>48</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(4)</mark>	Elsyad M	<mark>3</mark>	<mark>73</mark>	<mark>81.1</mark>
<mark>58</mark>	2 <mark>01</mark> 2	Clin Oral Implants Res	201 2	<mark>23(3)</mark>	Lorenzo R	<mark>5</mark>	<mark>83</mark>	<mark>92.2</mark>
<mark>78</mark>	201 2	Clin Oral Implants Res	201 2	<mark>23(2)</mark>	Hammerle C	<mark>5</mark>	<mark>76</mark>	<mark>84.4</mark>
<mark>79</mark>	<mark>201</mark> 2	Clin Oral Implants Res	201 2	<mark>23(2)</mark>	<mark>Urban T</mark>	<mark>3</mark>	<mark>63</mark>	<mark>70.0</mark>
<mark>143</mark>	201 1	Clin Oral Implants Res	201 1	<mark>22(11)</mark>	<mark>den Hartog L</mark>	<mark>5</mark>	<mark>75</mark>	<mark>83.3</mark>
<mark>155</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(10)</mark>	Cordaro L	<mark>3</mark>	<mark>70</mark>	<mark>77.8</mark>
<mark>162</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(10)</mark>	Enkling N	<mark>3</mark>	<mark>67</mark>	<mark>74.4</mark>
<mark>173</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(8)</mark>	Karabuda Z	<mark>3</mark>	<mark>71</mark>	<mark>78.9</mark>
<mark>175</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(8)</mark>	Galindo-Moreno P	<mark>3</mark>	<mark>58</mark>	<mark>64.4</mark>
<mark>182</mark>	2 <mark>01</mark> 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(7)</mark>	Sakalioglu U	<mark>2</mark>	<mark>54</mark>	<mark>60.0</mark>
<mark>193</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(6)</mark>	<mark>van Brakel</mark>	<mark>2</mark>	<mark>61</mark>	<mark>67.8</mark>
<mark>201</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(6)</mark>	Nissan J	<mark>2</mark>	<mark>54</mark>	<mark>60.0</mark>
<mark>202</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(6)</mark>	Bressan E	<mark>1</mark>	<mark>55</mark>	<mark>61.1</mark>
<mark>209</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(5)</mark>	Jokstad A	<mark>5</mark>	<mark>80</mark>	<mark>88.9</mark>
<mark>211</mark>	<mark>201</mark> 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(5)</mark>	Chackartchi T	3	<mark>63</mark>	<mark>70.0</mark>
<mark>222</mark>	<mark>201</mark> 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(5)</mark>	Heberer S	<mark>3</mark>	<mark>58</mark>	<mark>64.4</mark>
<mark>234</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(4)</mark>	Mardas N	4	<mark>79</mark>	<mark>87.8</mark>
<mark>239</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(3)</mark>	Heitz-Mayfield L	5	<mark>69</mark>	<mark>76.7</mark>
<mark>241</mark>	<mark>201</mark> 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(3)</mark>	Rickert D	<mark>4</mark>	<mark>65</mark>	<mark>72.2</mark>
<mark>252</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(3)</mark>	Alsabeeha N	<mark>3</mark>	<mark>74</mark>	<mark>82.2</mark>
<mark>272</mark>	<mark>201</mark> 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(1)</mark>	Tan W	<mark>3</mark>	<mark>64</mark>	<mark>71.1</mark>
<mark>278</mark>	201 1	Clin Oral Implants Res	<mark>201</mark> 1	<mark>22(1)</mark>	Galucci G	<mark>4</mark>	<mark>70</mark>	<mark>77.8</mark>
<mark>298</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(12)</mark>	Felice P	<mark>5</mark>	<mark>81</mark>	<mark>90.0</mark>
<mark>302</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(11)</mark>	Van Der Bilt A	<mark>1</mark>	<mark>53</mark>	<mark>58.9</mark>
<mark>304</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(11)</mark>	Van de Velde T	<mark>3</mark>	<mark>66</mark>	<mark>73.3</mark>
<mark>309</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(11)</mark>	<mark>Urban T</mark>	<mark>3</mark>	<mark>61</mark>	<mark>67.8</mark>
<mark>313</mark>	201 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(11)</mark>	Koch F	<mark>3</mark>	<mark>64</mark>	<mark>71.1</mark>
<mark>318</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(9)</mark>	Pineiro A	<mark>2</mark>	<mark>59</mark>	<mark>65.6</mark>

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	U.

<mark>343</mark>	<mark>201</mark>	Clin Oral Implants Res	<mark>201</mark>	<mark>21(7)</mark>	Degidi M	<mark>5</mark>	<mark>73</mark>	<mark>81.1</mark>
344	0 201	Clin Oral Implants Res	0 201	21(7)	Mardas N	<u>4</u>	72	80.0
	0 201		0 201					
373	0 201	Clin Oral Implants Res	0 201	<mark>21(5)</mark>	Elsyad M	3	<mark>68</mark>	<mark>75.6</mark>
<mark>374</mark>	0 201	Clin Oral Implants Res	0 201	<mark>21(5)</mark>	Zembic A	<mark>4</mark>	<mark>69</mark>	<mark>76.7</mark>
<mark>377</mark>	0	Clin Oral Implants Res	0	<mark>21(5)</mark>	Thone-Muhling M	<mark>3</mark>	<mark>59</mark>	<mark>65.6</mark>
<mark>381</mark>	201 0	Clin Oral Implants Res	201 0	<mark>21(5)</mark>	Pelegrine A	2	<mark>60</mark>	<mark>66.7</mark>
<mark>426</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(2)</mark>	Park J-C	<mark>4</mark>	<mark>76</mark>	<mark>84.4</mark>
<mark>428</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(2)</mark>	<mark>Jofre J</mark>	<mark>2</mark>	<mark>64</mark>	<mark>71.1</mark>
<mark>431</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(1)</mark>	Sanz M	<mark>3</mark>	<mark>69</mark>	<mark>76.7</mark>
<mark>444</mark>	<mark>201</mark> 0	Clin Oral Implants Res	<mark>201</mark> 0	<mark>21(1)</mark>	Canullo L	<mark>4</mark>	<mark>61</mark>	<mark>67.8</mark>
<mark>467</mark>	<mark>201</mark> 2	Clin Implant Dent Relat Res	<mark>201</mark> 2	<mark>14(s1)</mark>	<mark>Al-Zubeidi M</mark>	<mark>2</mark>	<mark>59</mark>	<mark>65.6</mark>
<mark>503</mark>	<mark>201</mark> 2	Clin Implant Dent Relat Res	201 2	<mark>14(2)</mark>	Enkling N	<mark>3</mark>	<mark>49</mark>	<mark>54.4</mark>
<mark>509</mark>	201 2	Clin Implant Dent Relat Res	2 <mark>01</mark> 2	<mark>14(1)</mark>	Lindgren C	<mark>2</mark>	<mark>58</mark>	<mark>64.4</mark>
<mark>515</mark>	201 2	Clin Implant Dent Relat Res	201 2	<mark>14(1)</mark>	Ortorp A	1	<mark>61</mark>	<mark>67.8</mark>
<mark>521</mark>	2 201 2	Clin Implant Dent Relat Res	201 201 2	<mark>14(1)</mark>	Enkling N	2	<mark>50</mark>	<mark>55.6</mark>
<mark>534</mark>	<mark>201</mark>	Clin Implant Dent Relat Res	<mark>201</mark>	<mark>13(3)</mark>	Wenneberg A	<mark>4</mark>	<mark>69</mark>	<mark>76.7</mark>
<mark>550</mark>	1 201	Clin Implant Dent Relat Res	1 201	<mark>13(2)</mark>	Visser A	3	<mark>61</mark>	<mark>67.8</mark>
<mark>586</mark>	1 201	Clin Implant Dent Relat Res	1 201	12(2)	Cehreli M	<mark>4</mark>	<mark>53</mark>	58.9
<mark>597</mark>	0 201	Clin Implant Dent Relat Res	0 201	12(1)s	Turkyilmaz I	3	60	<mark>66.7</mark>
<mark>634</mark>	0 200	Clin Implant Dent Relat Res	0 200	11(3)	Mericske-Stern R	2	<mark>60</mark>	<mark>66.7</mark>
649	9 200	Clin Implant Dent Relat Res	9 200	11(1)s	Canullo L	- 3	59 59	65.6
695	9 200	Clin Implant Dent Relat Res	9 200	10(1)	Guncu G	<mark>3</mark>	55	61.1
	8 200	Clin Implant Dent Relat Res	8 200		Hall J		56	62.2
726	<mark>7</mark> 201		<mark>7</mark> 201	9(1)		3		
819	2 201	Int J Oral Maxillofac Implants	2 201	27(2)	Wohlfahrt J	4 4	70 22	77.8
<mark>823</mark>	2 201	Int J Oral Maxillofac Implants	2 201	<mark>27(2)</mark>	Ramel C		<mark>63</mark>	<mark>70.0</mark>
<mark>873</mark>	1 201	Int J Oral Maxillofac Implants	1 201	<mark>26(6)</mark>	Taguchi T	2	<mark>55</mark>	<mark>61.1</mark>
<mark>946</mark>	1	Int J Oral Maxillofac Implants	1	<mark>26(3)</mark>	Krennmair G	1	<mark>47</mark>	<mark>52.2</mark>
<mark>948</mark>	201 1 201	Int J Oral Maxillofac Implants	201 1 201	<mark>26(3)</mark>	Canullo L	<mark>4</mark>	<mark>67</mark>	<mark>74.4</mark>
<mark>949</mark>	201 1	Int J Oral Maxillofac Implants	201 1	<mark>26(3)</mark>	Fung K	<mark>4</mark>	<mark>66</mark>	<mark>73.3</mark>
<mark>972</mark>	<mark>201</mark> 1	Int J Oral Maxillofac Implants	<mark>201</mark> 1	<mark>26(2)</mark>	Heberer S	2	<mark>53</mark>	<mark>58.9</mark>
<mark>975</mark>	<mark>201</mark> 1	Int J Oral Maxillofac Implants	<mark>201</mark> 1	<mark>26(2)</mark>	<mark>De Kok I</mark>	<mark>2</mark>	<mark>62</mark>	<mark>68.9</mark>
<mark>989</mark>	<mark>201</mark> 1	Int J Oral Maxillofac Implants	<mark>201</mark> 1	<mark>26(1)</mark>	Salihoglu U	<mark>2</mark>	<mark>55</mark>	<mark>61.1</mark>
<mark>998</mark>	201 1	Int J Oral Maxillofac Implants	<mark>201</mark> 1	<mark>26(1)</mark>	Pieri F	<mark>5</mark>	<mark>71</mark>	<mark>78.9</mark>
<mark>102</mark> 2	<mark>201</mark> 0	Int J Oral Maxillofac Implants	<mark>201</mark> 0	<mark>25(6)</mark>	Jofre J	<mark>4</mark>	<mark>65</mark>	<mark>72.2</mark>
<mark>107</mark> 8	201 0	Int J Oral Maxillofac Implants	201 0	<mark>25(4)</mark>	Merli M	<mark>5</mark>	<mark>85</mark>	<mark>94.4</mark>
109 9	201 0	Int J Oral Maxillofac Implants	201 0	<mark>25(3)</mark>	van Kesteren C	2	<mark>60</mark>	<mark>66.7</mark>
9 <mark>121</mark> 3	200 9	Int J Oral Maxillofac Implants	200 9	<mark>24(5)</mark>	Aimetti M	<mark>3</mark>	<mark>63</mark>	<mark>70.0</mark>
<mark>127</mark>	<mark>200</mark>	Int J Oral Maxillofac Implants	<mark>200</mark>	24(2)	Prosper L	5	80	88.9
8 133	9 200	Int J Oral Maxillofac Implants	9 200	23(5)	Covani U	3	<mark>58</mark>	<mark>64.4</mark>

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								17
6 400	8		8					
133 9	200 8	Int J Oral Maxillofac Implants	200 8	<mark>23(5)</mark>	Cannizzaro G	<mark>5</mark>	<mark>79</mark>	<mark>87.8</mark>
134 7	<mark>200</mark> 8	Int J Oral Maxillofac Implants	200 8	<mark>23(5)</mark>	<mark>Shahidi P</mark>	<mark>3</mark>	<mark>60</mark>	<mark>66.7</mark>
137 3	<mark>200</mark> 8	Int J Oral Maxillofac Implants	200 8	<mark>23(4)</mark>	Schropp L	<mark>3</mark>	<mark>56</mark>	<mark>62.2</mark>
137 5	<mark>200</mark> 8	Int J Oral Maxillofac Implants	<mark>200</mark> 8	<mark>23(4)</mark>	<mark>Crespi R</mark>	<mark>2</mark>	<mark>57</mark>	<mark>63.3</mark>
<mark> 38</mark> 7	<mark>200</mark> 8	Int J Oral Maxillofac Implants	<mark>200</mark> 8	<mark>23(3)</mark>	Schincaglia G	<mark>3</mark>	<mark>69</mark>	<mark>76.7</mark>
<mark> 40</mark> }	200 8	Int J Oral Maxillofac Implants	<mark>200</mark> 8	<mark>23(2)</mark>	Morneburg T	<mark>2</mark>	<mark>57</mark>	<mark>63.3</mark>
146 9	200 7	Int J Oral Maxillofac Implants	<mark>200</mark> 7	<mark>22(5)</mark>	Oates T	<mark>2</mark>	<mark>57</mark>	<mark>63.3</mark>
147 3	<mark>200</mark> 7	Int J Oral Maxillofac Implants	200 7	<mark>22(5)</mark>	Testori T	<mark>5</mark>	<mark>78</mark>	<mark>86.7</mark>
1 <mark>51</mark> 5	201 2	Implant Dent	2 <mark>01</mark> 2	<mark>21(3)</mark>	Gadallah A	<mark>5</mark>	<mark>65</mark>	<mark>72.2</mark>
154 7	201 2	Implant Dent	2 <mark>01</mark> 2	<mark>21(3)</mark>	Gadallah A	<mark>4</mark>	<mark>56</mark>	<mark>62.2</mark>
171 9	<mark>201</mark> 0	Implant Dent	201 0	<mark>19(2)</mark>	Basha A	<mark>2</mark>	<mark>57</mark>	<mark>63.3</mark>
1 <mark>79</mark>	200 9	Implant Dent	200 9	<mark>18(1)</mark>	<mark>Guncu G</mark>	<mark>3</mark>	<mark>59</mark>	<mark>65.6</mark>
1 <mark>91</mark>	201 2	J Dent	201 2	<mark>40(5)</mark>	<mark>de Sousa Barbosa</mark> R	<mark>2</mark>	<mark>54</mark>	<mark>60.0</mark>
196 7	201 2	J Dent	201 2	<mark>40(1)</mark>	West N	<mark>4</mark>	<mark>79</mark>	<mark>87.8</mark>
1 <mark>98</mark>)	201 1	J Dent	201 1	<mark>39(s3)</mark>	Moffa E	<mark>3</mark>	<mark>58</mark>	<mark>64.4</mark>
2 <mark>00</mark>	201 1	J Dent	201	<mark>39(11)</mark>	Lopez-Jornet M	<mark>3</mark>	<mark>60</mark>	<mark>66.7</mark>
1 200	<mark>201</mark>	J Dent	201	<mark>39(11)</mark>	Kitasako Y	<mark>4</mark>	<mark>65</mark>	<mark>72.2</mark>
8 2 <mark>01</mark>	1 201	J Dent	1 201	<mark>39(10)</mark>	Ren Y-F	3	<mark>58</mark>	<mark>64.4</mark>
203	1 201	J Dent	1 201	39(7)	Huth K	4	<mark>69</mark>	<mark>76.7</mark>
3 204	1 201	J Dent	1 201	39(7)	Nelson-Filho P	3	<mark>63</mark>	70.0
2 204	1 201	J Dent	1 201	<u>39(7)</u>	Shen P	3	<mark>70</mark>	77.8
205	1 201	J Dent	1 201	<mark>39(5)</mark>	Wirsching E	2	<mark>53</mark>	<mark>58.9</mark>
5 2 <mark>11</mark>	1 201	J Dent	201	38(12)	Meireles S	5	70	77.8
2 <mark>12</mark>	0 201	J Dent	0 201	38(12)	Huth K	3	67	74.4
2 <mark>12</mark>	0 201	J Dent	0 201	38(11)	Hyde T	5	66	73.3
213	0 201	J Dent	0 201	38(11)	Pan S	3	<mark>62</mark>	68.9
1 217	0 201	J Dent	0 201	<mark>38(7)</mark>	Syrek A	3	54	60.0
3 218	0 201	J Dent	0 201	38(6)	dos Santos M	3	<mark>64</mark>	71.1
1 218	0 201	J Dent	0 201	38(6)	Banerjee A	<mark>З</mark>	58	<mark>64.4</mark>
4 218	0 201	J Dent	0 201	38(6)	Mcdonald E	3 3	58 58	64.4
9 221	0 201	J Dent	0 201	38(3)	Emami E	2 2	<mark>64</mark>	71.1
9 225	0 201	J Dent	0 201	38s3	Mason S	2 3	52	57.8
2 225	0 201	J Dent	0 201		Maggio B		52 63	57.8 70.0
3 235	0 201		0 201	38s3		2		
2 242	1 201	J Prosthet Dent	<mark>1</mark> 201	106(1)	Burns D	2	64 48	71.1
4 261	0 201	J Prosthet Dent	0 201	104(6)	Damodara E	0 D	48 60	53.3
8 275	1 201	J Oral Rehab	1 201	38(10)	Nilsson H	2	69	76.7
9 285	0 201	J Oral Rehab	0 201	<mark>37(7)</mark>	Kimoto S	4	<mark>76</mark>	<mark>84.4</mark>
2 2	2	Int J Prosth	2	<mark>25(4)</mark>	Gjengedal H	<mark>2</mark>	<mark>53</mark>	<mark>58.9</mark>

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<mark>286</mark> 5	<mark>201</mark> 2	Int J Prosth	<mark>201</mark> 2	<mark>25(3)</mark>	Stober T	<mark>2</mark>	<mark>53</mark>	<mark>58.9</mark>
286 7	201 2	Int J Prosth	201 2	<mark>25(3)</mark>	Sagirkaya E	<mark>2</mark>	<mark>63</mark>	<mark>70.0</mark>
286 8	201 2	Int J Prosth	201 2	<mark>25(3)</mark>	Volpato Sanita P	<mark>3</mark>	<mark>70</mark>	<mark>77.8</mark>
0 288 6	201 2	Int J Prosth	201 2	<mark>25(2)</mark>	Elsyad M	<mark>2</mark>	<mark>54</mark>	<mark>60.0</mark>
0 288 7	201 201 2	Int J Prosth	<mark>201</mark>	<mark>25(2)</mark>	Machado de	1	<mark>49</mark>	<mark>54.4</mark>
2 <mark>94</mark>	<mark>201</mark>	Int J Prosth	2 201	<mark>24(4)</mark>	<mark>Andrade I</mark> Zicari F	3	<mark>72</mark>	80.0
7 299 3	1 201 1	Int J Prosth	1 201	<mark>24(1)</mark>	Cehreli M	2	<mark>62</mark>	<mark>68.9</mark>
300 7	201 0	Int J Prosth	201 0	<mark>23(6)</mark>	Larsson C	0	<mark>49</mark>	<mark>54.4</mark>
3 <mark>03</mark>	<mark>201</mark>	Int J Prosth	<mark>201</mark>	<mark>23(4)</mark>	Cune M	0	52	<mark>57.8</mark>
) 305 -	0 201	Int J Prosth	0 201	<mark>23(3)</mark>	Klat-amnuay S	3	79	87.8
5 306	0 201	Int J Prosth	0 201	23(2)	Kimoto S	4	76	84.4
) 308	0 200	Int J Prosth	0 200	22(6)	Sailer I	2	57	63.3
6 812	9 200	Int J Prosth	9 200	22(4)	Walton J	4	78	86.7
) 312	9 200	Int J Prosth	9 200	22(4)	Pradies G	2	62	68.9
6 314 -	9 200	Int J Prosth	9 200	22(3)	Cannulo L	3	<mark>69</mark>	76.7
3 314	9 200	Int J Prosth	9 200	22(3)	Haim M	2	57	63.3
3 3 <mark>21</mark>	9 200	Int J Prosth	9 200	21(4)	Berg E	2	58	64.4
) 3 <mark>22</mark>	8 200	Int J Prosth	<mark>8</mark> 200	21(4)	Luthardt R	2	50 50	55.6
1 329	8 200	Int J Prosth	<mark>8</mark> 200	20(5)	Naumann M	2	<mark>61</mark>	67.8
3 333	7 201	Int J Periodontics Restorative	7 201	<u>32(4)</u>	Cardaropoli D	<mark>-</mark> 4	67	74.4
) 3 <mark>34</mark>	2 201	Dent Int J Periodontics Restorative	2 201	<u>32(3)</u>	Riza Certin A	0	51 51	56.7
7 335	2 201	Dent Int J Periodontics Restorative	2 201	32(3)	Griffiths G		56	62.2
1 335	2 201	Dent Int J Periodontics Restorative	2 201	32(2)	Jankovic K	2	60 60	66.7
3 336	2 201	Dent Int J Periodontics Restorative	2 201	32(2)	Margossian P	- 3	57	63.3
1 337	2 201	Dent Int J Periodontics Restorative	2 201	32(1)	Cordaro L	2	73	81.1
7 338	2 201	Dent Int J Periodontics Restorative	2 201	<u>31(6)</u>	Froum S		<mark>59</mark>	65.6
3 344	<mark>1</mark> 201	Dent Int J Periodontics Restorative	<mark>1</mark> 201		Rasperini G	2 1	50	55.6
3 350	<mark>1</mark> 201	Dent Int J Periodontics Restorative	<mark>1</mark> 201	31(2)				63.3
4 351	0 201	Dent Int J Periodontics Restorative	0 201	30(3)	Rasperini G	1	57	
5 353	0 200	Dent Int J Periodontics Restorative	0 200	30(2)	Wu S-Y	0	51 40	56.7
5 357	9 200	Dent Int J Periodontics Restorative	9 200	29(6)	Trammel K	3	<mark>49</mark>	54.4
9 358	9 200	Dent Int J Periodontics Restorative	9 200	29(2)	Haghighati F	0	44	48.9
9 360	9 200	Dent Int J Periodontics Restorative	9 200	29(1)	Cardaropoli D	2	47	52.2
7 361 361	8 200	Dent Int J Periodontics Restorative	8 200	28(5)	Merli M	2	<u>59</u>	<mark>65.6</mark>
9 9	<mark>200</mark> 8	Dent	200 8	<mark>28(4)</mark>	<mark>Jung R</mark>	<mark>1</mark>	<mark>42</mark>	<mark>46.7</mark>

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