# **Original Article**

# Preprocedural Anxiety and Pain Perception Following Root Surface Debridement in Chronic Periodontitis Patients

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# Abstract

**Background:** The aim of this study was to evaluate and compare preprocedural dental anxiety levels and postprocedural pain perception in chronic periodontitis patients during conventional-staged root surface debridement (RSD) and single-stage RSD. **Materials and Methods:** Thirty-seven adult generalized chronic periodontitis patients requiring RSD were recruited in this study. Preprocedural anxiety levels were assessed using a self-reported questionnaire and postprocedural pain perceptions were assessed using 0–10 cm visual analog scale. The subject population was divided into two groups: staged RSD (n = 18) and single-stage RSD (n = 19). Staged RSD patients visited four times as opposed to single-stage RSD patients. Data were subjected to Pearson Chi-square test, Mann–Whitney U-test, and Spearman's rank correlation. **Results:** There was no statistically significant difference in dental anxiety levels between visit 4 and visit 3 (P = 0.037) and pain perception between visit 3 and visit 1 (P = 0.005), visit 4 and visit 1 (P = 0.002), and visit 4 and visit 2 (0.04) was statistically significant. There was a positive correlation of anxiety questionnaire (Q1–Q4) to the pain score in Group 1 which was statistically significant and in single-stage RSD. **Conclusion:** Conventional quadrant-wise RSD tends to cognitively condition the anxiety experience thus influencing pain experience.

Keywords: Chronic periodontitis, dental anxiety, pain, patient-centered outcomes, root surface debridement, visual analog scale

## INTRODUCTION

Dental anxiety and fear are strong negative feelings associated with dental treatment. Dental anxiety was described by Klingberg and Broberg as a state of apprehension that something dreadful is going to happen in relation to dental treatment or certain aspects of dental treatment.<sup>[11]</sup> Dental anxiety is a multidimensional construct that consists of somatic, cognitive, and emotional elements and describes a general state that is not stimulus specific. This trait in an individual may result in avoidance of dental treatment.<sup>[2]</sup> The experience of pain during dental procedures is a concern to many individuals. Hence, all members of the dental team including periodontists or dental hygienists must aim to minimize the degree of discomfort during periodontal procedures such as scaling and root surface debridement (RSD).<sup>[3]</sup>

There are reports in periodontal literature related to patients' perception of pain and discomfort during periodontal probing,<sup>[4]</sup> scaling,<sup>[5]</sup> RSD,<sup>[3]</sup> periodontal surgery,<sup>[6]</sup> and maintenance

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treatments.<sup>[7]</sup> However, the impact on patients in regard to single-visit or multi-visit RSD has yet to be fully investigated.

The treatment of chronic periodontitis primarily involves the reduction or elimination of bacteria present in the plaque biofilm.<sup>[8]</sup> The periodontal pathogens can establish not only in periodontal pockets but also on the tongue, tonsils, or on the other oral mucous membranes. These sites may represent a potential reservoir for the reinfection of adjacent sites following active periodontal treatment.<sup>[9,10]</sup>

To avoid the risk of intraoral bacterial translocation to recently instrumented and healing periodontal sites, Leuven group

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introduced the concept "one visit full mouth disinfection." This concept utilizes instrumentation of periodontal pockets and use of antiseptic disinfection for the remaining sites in the oral cavity within 24 h.<sup>[11]</sup>

The studies on the clinical efficacy of the full mouth disinfection versus conventional multi-visit approach suggest only minor differences between the two protocols.<sup>[12-14]</sup> Nevertheless, these conclusions reflect the therapeutic outcomes clinically based rather than patient-centered outcomes which must also be considered while selecting a treatment.<sup>[15]</sup> Nonsurgical periodontal therapies are often perceived as stressful and painful by the patient.<sup>[3]</sup> However, to our knowledge, there are very few studies in the literature comparing patient perceptions of single-visit RSD versus conventional multi-visit debridement in terms of anxiety and pain. Hence, this study aimed to assess the difference in preprocedural anxiety and postoperative pain levels in multi-visit and single-visit RSD approaches.

The objectives of this study were:

- To assess and compare the preprocedural anxiety levels in chronic periodontitis patients who were undergoing conventional multi-visit and single-visit RSD
- To assess and compare the postoperative pain perception in chronic periodontitis patients who underwent conventional multi-visit and single-visit RSD
- To correlate the anxiety levels to pain perception in patients undergoing multi-visit and single-visit RSD.

# **MATERIALS AND METHODS**

#### **Patient sample**

Seventeen male and twenty female patients with generalized chronic periodontitis (GCP) were included in this study. All subjects were recruited from the Department of Periodontics, SRM Dental College, Ramapuram, Chennai, India (Ethical Committee Approval number SRMU/MandHS/SRMDC/2010/M.D.S-Staff/103). Patients with GCP with the clinical evidence of probing depths  $\geq$ 4 mm, presence of more than 24 teeth with a minimum of four molars, no previous history of periodontal therapy, and with good systemic health (as assessed by the recruiting periodontist) were included in this study.<sup>[16]</sup> The assessment criteria for generalized chronic periodontitis was according to American Academy of Periodontology 1999 classification.<sup>[16]</sup>

Patient who were on antibiotics and analgesics in the last 6 weeks, patients with the history of systemic diseases which interfere with the pain perception, such as neurological or psychiatric disorders and who were on medications which interfere with pain perception were also excluded from the study. In addition, patients presenting with acute dental, periapical/periodontal pain, dentinal hypersensitivity, and subjects with orthodontic and prosthetic appliances were excluded from the study. Finally, patients who did not attend all appointments were excluded from the study.

The literature suggests that age, sex, and socioeconomic status (SES) of an individual can influence fear and anxiety. Tuba TaloYildirim 2016 in their study suggested statistically significant difference between the levels of dental anxiety and sociodemographic status; hence, we selected individuals belonging to similar SES so as to eliminate any bias incorporation.<sup>[17]</sup> The Kuppuswamy scale is recognized as a tool to measure SES in the urban population. This validated scale utilizes an online tool which was used in this study and utilized SES based on 2013 criteria.<sup>[18]</sup>

#### Study design

This is a randomized, prospective, single-blind, controlled clinical study. This study was conducted from January 2013 to December 2013. This study was approved by the Research Ethics Committee of the SRM Dental College, Ramapuram, Chennai, India. Patients were recruited based on the selection criteria and were randomly assigned to one of the following groups [Figure 1].

Group 1: Conventional multi-visit RSD, Group 2: Single-visit RSD.

The sample size was calculated based on the primary outcome measure, i.e., effect of RSD on preprocedural anxiety levels and pain perception obtained from the results of pilot study. Thus, a minimum sample size was determined to be 16 in each group ( $\alpha = 0.05$  power 90%). The proposed sample size of 16 was adjusted to 18 based on hospital records of the institute which indicated an average 15% drop out rate. An initial pilot study using 16 subjects was performed to help standardize the reliability of examiner in terms of clinical parameters and consistency of operator for the treatment provided.

Independent randomisation for allocation to Group 1 or Group 2 was performed using the flip of a coin. Written informed consent was obtained from each subject. Participants' social information such as education, income, previous visits, and smoking status were also recorded. Further, the SES of individuals in the Chennai urban population was subjected to 2013 Kuppuswamy tool.

#### **Clinical protocol**

Eighteen patients were treated with conventional multi-visit approach and 19 patients with single-visit RSD approach. The procedures were performed under local anesthesia: 2% lignocaine, 1:80,000 adrenaline. Buccal and palatal infiltration was given for maxillary quadrants and inferior alveolar nerve block with long buccal and lingual nerve block were given for mandibular quadrants. All the patients were given a similar dose of anesthesia. A combination of site-specific Gracey curettes (Hu-friedy, USA) and ultrasonic piezoelectric scaler (EMS, Piezon<sup>®</sup>) were used for RSD procedure followed by polishing with prophylactic paste (Proxyt<sup>®</sup>prophy paste, Ivoclar, Vivadent). Patients were asked to take analgesics (paracetomol 500 mg two tablets four times a day for 5 days) only if they felt necessary following wearing off the anesthesia as the patients were required to record their pain scores on visual analog scale (VAS). Only four

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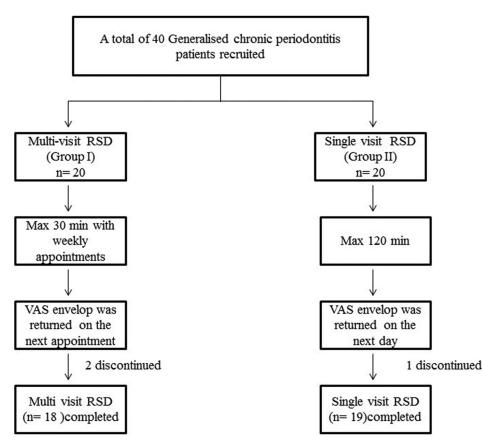


Figure 1: Flow chart of subject recruitment

patients out of thirty-seven reported the use of analgesics. Oral hygiene instructions were given for each patient on their first visit including Bass technique of toothbrushing and interdental brushes as appropriate and were prescribed a standard anticalculus and antigingivitis-formulated toothpaste (Colgate total). All patients were treated in a controlled clinical atmosphere by a single-experienced periodontist (VKN). The Group 1 patients were treated quadrant based in 4 visits, at an interval of 1 week and Group 2 patients were treated in a single visit.

#### **Clinical parameters**

Probing pocket depth and clinical attachment level were recorded by a single operator (AB). Assessment of anxiety and pain was performed by single operator (DA). The periodontist (VKN) was blinded to the anxiety and pain scores to avoid the incorporation of bias.

#### **Probing pocket depth**

This was assessed using UNC-15 probe (Hu-friedy, USA) from the gingival margin to the base of the pocket at 6 sites (mesiobuccal, midbuccal, distobuccal, mesiolingual, mid lingual, and distolingual).

#### **Clinical attachment level**

This was assessed using UNC-15 probe (Hu-friedy) from the cementoenamel junction to the base of the pocket at 6 sites (mesiobuccal, midbuccal, distobuccal, mesiolingual, midlingual, and distolingual).

#### Assessment of anxiety

All subjects were given information on the treatment to be provided and were assessed for their anxiety levels at the beginning of appointment before the start of the procedure using a self-reported questionnaire adopted by Chung et al. 2003<sup>[19]</sup> and Guzeldemir et al. 2008.<sup>[20]</sup> This questionnaire consisted of 4 questions from Corah's dental anxiety scale (DAS) and 3 questions from dental fear survey (DFS) [Table 1]. In this study, we replaced the word "drill" with "instruments" in Q6 and "cleaning of your teeth" with "deep cleaning of your teeth" in Q7 to suit the treatment protocol of this study. The validity of this questionnaire was assessed before the study by administering the questionnaire to thirty patients at two different time points at 1 week interval. Group 1 patients responded to this questionnaire at each treatment visit, while Group 2 patients responded once before the treatment.

#### Assessment of pain intensity

The subjective perception of pain was assessed postoperatively using a 0–10 cm VAS. Patients were given guidance about how to score the VAS; score "0" being no pain or discomfort, and score "10" being worst pain or discomfort. Patients were discharged posttreatment with the VAS in an envelope, so subjects could complete their pain scores once the anesthetic had worn (minimum 4 h) off. Subjects received verbal reminder over the phone 4 h after the procedure and asked to complete

#### Table 1: Dental Anxiety Questionnaire

- I. How much anxiety/fear or discomfort does each of these cause you?
  - 1. Being seated in dental chair
- a. None at all
- b. A little
- c. Somewhat
- d. Much
- e. Very much
- 2. Having your teeth cleaned
- a. None at all
- b. A little
- c. Somewhat
- d. Much
- e. Very much
- 3. All things considered, how fearful are you of having dental work done? a None at all
- b. A little
- c. Somewhat
- d. Much
- e. Very much

II. If you had to go to the dentist tomorrow, how would you feel about it?

- a. I would look forward to it as a reasonably enjoyable experience
- b. I would not care one way or another
- c. I would be a little uneasy about it
- d. I would be afraid that it would be unpleasant and painful
- e. I would be very frightened of what the dentist might do

III. When you are waiting in the dentist's office for your turn in the chair, how do you feel?

- a. Relaxed
- b. A little uneasy
- c. Tense
- d. Anxious

e. So anxious that I sometimes break out in a sweat or almost feel physically sick

IV. When you are in the dentist's chair waiting while she gets the instruments ready to begin working on your teeth, how do you feel?

- a. Relaxed
- b. A little uneasy
- c. Tense
- e. Anxious

f. So anxious that I sometimes break out in a sweat or almost feel physically sick

V. You are in the dentist's chair to have your teeth deep cleaned. While you are waiting and the dental assistant is getting out the instruments that the periodontist will use to clean your teeth around the gums, how do you feel? a. Relaxed

- b. A little uneasy
- c Tense
- c. rense
- d. Anxious

e. So anxious that I sometimes break out in a sweat or almost feel physically sick

Question I originate from the DFS and Questions II through V are from Corah's DAS. DFS: Dental fear survey, DAS: Dental anxiety scale

the VAS. Subjects were asked to return the completed VAS envelope at the next visit.

#### Treatment time

The time taken to perform the multi-visit RSD was maximum 30 min for each quadrant and for single-visit RSD maximum of 120 min.

A flow chart of data collection and treatment provided in Group 1 and Group 2 is represented in Figure 1.

#### **Statistical analysis**

At the outset, data were presented descriptively and compared between the two groups noting the characteristics of the subjects. Data distribution pattern of the study population was assessed using Shapiro–Wilk test. Based on data distribution, parametric and nonparametric tests were used for statistical analysis. Thus, for statistical analysis, the Pearson Chi-square test and Mann–Whitney U-test were used for comparison of data and Spearman's rank correlation was used for correlation analysis between dental anxiety scores to the pain scores. SPSS, Version 20 (IBM Corp., Chicago, Illinois, USA) was used to carry out statistical analysis. P < 0.05 was considered statistically significant for analysis.

## RESULTS

#### Characteristics of the study subjects

A total of 37 individuals (20 females, 17 males) with chronic periodontitis were recruited in the current study and randomly divided into Group 1 (n = 18) (conventional multi-visit debridement) and Group 2 (n = 19) (24 h RSD). Overall, the mean age in the study population was 40.97 ± 10.32 years and age ranged from 25 to 64 years. Table 2 presents the descriptive characteristics of the subjects recruited. There was no significant correlation between clinical parameters and anxiety levels and pain perceptions.

There was no significant difference in the demographic and clinical variables between Group 1 and Group 2 (P > 0.05) [Table 2]. No statistical significant difference was found between the Group 1 and Group 2 with regard to mean age, education levels, previous visits to the dentist, probing pocket depth, and clinical attachment level.

Based on the scores of the Kuppuswamy scale, the population was divided into upper, upper middle, lower middle, upper lower, and lower socioeconomic class. In Group 1, 61% of subjects and in Group 2, 58% belonged to upper lower class. Thus, the majority of the population belonged to similar socioeconomic class, eliminating any influence in the anxiety state [Table 3a and b].

#### **Anxiety scores**

The preprocedural anxiety levels were assessed using DAS and DFS questionnaire and were validated, before the start of the study. The validity of this questionnaire was assessed by distributing to 30 patients at two different time points 1 week apart. The Cronbach's  $\alpha = 0.87$  indicated excellent internal consistency of the questionnaire.

Table 4 represents the comparison of anxiety scores of both the groups at visit one at baseline which showed no statistical significance.

#### Visual analog scale scores for pain

Pain scores were assessed using VAS of 0–10 cm. Mean VAS scores for entire study population was  $1.8 \pm 1.5$  cm ranging

Table 2: Descriptive statistics						
Variables	Group 1	Group 2	Р			
Number of subjects	18	19				
Male	10	7	0.467			
Female	8	12	0.371			
Age (mean±SD)	41.72±2.69	40.26±2.16	0.893			
Smokers	2	4	0.414			
Education						
Secondary school	7	9	0.229			
Primary school	8	7				
UG	1	3				
PG	2	0				
Income per month (INR)						
5000	6	4	0.869			
10,000	7	13				
>10,000	5	2				
Previous visit						
1 <sup>st</sup> visit	0	0	0.223			
2 <sup>nd</sup> visit	9	9				
>2 visits	9	10				
Clinical parameters						
PD	3.71±0.22	3.6±0.25	0.578			
CAL	4.14±0.31	4.13±0.27	0.822			

SD: Standard deviation, INR: Indian rupees, PD: Probing depth, CAL: Clinical attachment loss

Table 3a: Kuppuswamy scale representing the	
socioeconomic status of recruited individuals	

Groups	Kuppasamy score					
	Upper	Upper- middle	Lower middle	Upper lower		
Group 1						
Count	0	2	5	11	18	
Percentage within groups	0	11.1	27.8	61.1	100.0	
Group 2						
Count	1	3	4	11	19	
Percentage within groups	5.3	15.8	21.1	57.9	100.0	
Total						
Count	1	5	9	22	37	
Percentage within groups	2.7	13.5	24.3	59.5	100.0	

from 0 to 5 cm. With regard to gender, the mean VAS scores for males were  $1.65 \pm 1.7$  cm ranging from 0 to 5 cm and for females were  $1.87 \pm 1.3$  cm, ranging from 0 to 4 cm. The comparison of VAS scores between Group 1 and 2 at baseline/visit one did not show any statistical significance with a P = 0.239.

#### Intragroup analysis within Group 1

The comparison of anxiety responses within Group 1 across 4 visits is presented in Table 5. The question 2 "having your teeth cleaned" showed a statistical significant difference in the anxiety scores with a P = 0.02.

 Table 3b: Comparison of socio-economic status of group

 1 and group 2 individuals based on Kuppuswamy scale

Chi-square tests						
	Value	df	Asymptotic significant (two-sided)			
Pearson Chi-square	1.285	3	0.733			
Likelihood ratio	1.672	3	0.643			
Linear-by-linear association	0.453	1	0.501			
Number of valid cases	37					

Table 4: Comparison of responses to anxiety questionnaire between groups at visit 1 (at baseline)

Question number	Group 1 versus Group 2 (Pearson $\chi^2$ )	Р
1	2.181	0.5
2	4.62	0.3
3	1.86	0.3
4	1.21	0.8
5	1.25	0.5
6	1.51	0.4
7	2.84	0.4

The results of intragroup comparison with relation to anxiety showed a statistically significant difference between visit 3 and 4. Further, the pain scores showed highly statistically significant difference between visit 3 and visit 1, visit 4 and visit 1, and between visit 2 and visit 4 as represented in Table 6.

#### **Correlation of anxiety levels to pain score**

Finally, the anxiety scores were correlated with the pain scores in both Group 1 and 2 at visit 1. Table 7 represents the correlation values along with the P values. In Group 1, question 1, 2, 3, and 7 showed a statistically significant positive correlation between anxiety and pain scores. On the contrary, in Group 2, none of the anxiety questions showed statistical significant correlation with the pain scores.

## DISCUSSION

The factors that have been shown to influence anxiety are age, gender, educational status, and SES. In this study, the subjects recruited showed similar demographics such as age, gender, educational status and SES as determined by Kuppuswamy scale and none of the subjects were completely new patients to dental treatment. Since subjects who are anxious are likely to experience more pain<sup>[21]</sup> than those who are not anxious, at the outset, all efforts were made in this study to avoid confounding factors that may influence dental anxiety levels.

The plaque biofilm is a community of microorganisms which are spatially organized into three dimensional structure and is supported by extracellular matrix.<sup>[22]</sup> The results of this prospective, randomized, blinded-controlled clinical study did not show differences in preprocedural dental anxiety levels and postoperative pain perceptions between single-visit RSD and conventional multi-visit RSD groups. This observation is in agreement with Santuchi *et al.* 2015 who conducted a

Table 5: Comparison	of	anxiety	responses	within	Group	1
across visits						

Question number	Pearson $\chi^2$	Р
1	11.5	0.07
2	23.2	0.02
3	13.6	0.13
4	11.08	0.5
5	6.7	0.3
6	9.9	0.3
7	7.7	0.8

Table 6: Intragroup comparison of anxiety and pain within Group 1

Anxiety	A2-A1	A3-A1	A4-A1	A3-A2	A4-A2	A4-A3
Р	0.673	0.564	0.08	1.0	0.07	0.037*
Pain	P2-P1	P3-P1	P4-P1	P3-P2	P4-P2	P4-P3
Р	0.496	0.005*	0.002*	0.254	0.04*	0.08

\*Statistical significance P<0.05

 Table 7: Correlation of anxiety questionnaire to the pain

 scores at first visit (1 time point) of Group 1 and Group 2

Gro	oup 1	Group 2		
r	Р	r	Р	
0.73	0.001*	0.24	0.30	
0.60	0.008*	0.33	0.16	
0.70	0.001*	0.33	0.16	
0.46	0.053*	0.02	0.92	
0.140	0.580	0.13	0.58	
0.40	0.093	0.25	0.29	
0.55	0.017*	0.25	0.29	
	<b>r</b> 0.73 0.60 0.70 0.46 0.140 0.40	0.73         0.001*           0.60         0.008*           0.70         0.001*           0.46         0.053*           0.140         0.580           0.40         0.093	r         P         r           0.73         0.001*         0.24           0.60         0.008*         0.33           0.70         0.001*         0.33           0.46         0.053*         0.02           0.140         0.580         0.13           0.40         0.093         0.25	

\*P < 0.05

6-month randomized controlled clinical trial to observe the clinical effects and patients' based outcomes with full mouth disinfection and scaling and root planing using a quadrant-based approach.<sup>[23]</sup> Further, authors in Santuchi *et al.* 2015 analyzed DAS and DFS independently as well as combined questionnaire and suggested that the scores of patients' based outcomes in full mouth disinfection (FMD) groups were superior to the quadrant-based group. However, they concluded that there were no statistically significant differences observed between both groups. On the contrary, in this study, we used DAS and DFS together as a single questionnaire as previously used by Sanikop *et al.* 2011 in an Indian population<sup>[5]</sup> and Guzeldemir *et al.* 2008.<sup>[20]</sup> In addition, the questionnaire was evaluated for internal consistency before the start of our study.

The pain perception by each individual is very subjective and can pose problems in recording a precise degree and amount of an individual's perceived pain.<sup>[24]</sup> There are several tools to evaluate pain perception. Self-reported tools used in dentistry to evaluate pain are often unifacial as they are easier to apply. Examples of unifacial pain assessment tools are VAS, verbal rating scale, numerical rating scale, computer graphic scale, and picture scales.<sup>[25]</sup> The examples of multidimensional pain assessment tools such as McGill pain questionnaire have shown to be highly consistent and supposedly the best tool to evaluate pain in research.<sup>[25,26]</sup> However, VAS is a simple measure to use, especially in research, where the purpose is to just record the pain perceived and not to evaluate all dimensions of pain characteristics.<sup>[26]</sup> Furthermore, VAS has been shown to be reliable and has been previously used to record pain levels following periodontal treatment.<sup>[27-29]</sup>

Perceived pain will differ, dependent on treatment provided by different clinicians and in different clinical environments. To exclude this as confounding factor, a single-experienced periodontist performed RSD on all patients in a standard clinical setting.

In this study, patients with  $\geq 4$  cm VAS score reported the use of analgesics on the day of the procedure, similarly, Karadottir et al. 2002 based on their study results suggested  $\geq$ 40 mm VAS score to be painful. Thus, the arbitrary thresholds of pain experience was set at  $\geq 4$  cm and overall about 10.5% of the recruited population showed  $\geq 4$  cm of arbitrary pain threshold and this is slightly lower percentage in comparison to Karadottir et al. 2002,<sup>[7]</sup> who reported 15% of patients experiencing pain. This most likely can be attributed to different study design, study population, use of hand instruments, and number of operators. Canakçi and Canakçi 2007 suggested that periodontal therapies were perceived as painful by GCP patients. Further postoperative pain was higher with surgical procedures which involved exposure of bone and increased time duration than with RSD.<sup>[27]</sup> Overall, our results are in agreement with Canakçi and Canakçi 2007 who also used a single-experienced operator to carry out all procedures. Similarly, Mei et al. 2016 suggested mild pain following periodontal and implant surgeries. Further, the duration, complexity of surgery, and additional anesthetic volume used increases local tissue expansion and the production of pro-inflammatory mediators, which stimulates the nociceptors influencing pain perceived.<sup>[30]</sup> However, we did not administer additional local anesthetic to any of our study subjects and standardized the volume that was used.

In the current study, in Group 1, patients came four times to complete the RSD and there was significant reduction in the anxiety levels between visit 4 and visit 3. Whereas Heaton *et al.* 2007 did not report any difference in anxiety levels with the past experience of a particular treatment or with the same clinician. However, they concluded that the results of earlier research in this regard had been conflicting.<sup>[31]</sup>

We observed a positive correlation of Q1–Q4 anxiety questions to the pain scores at the visit 1 in Group 1: Q1 being seated in the dental chair, Q2 having your teeth cleaned, Q3 all things considered, how fearful are you of having dental work done? And Q4 if you had to go to the dentist tomorrow, how would you feel about it? Showed a positive correlation to the VAS scores which were statistically significant. Whereas Karadottir *et al.* 2002<sup>[7]</sup> showed positive correlation of anxiety responses for Q4–Q7 (Corah's DAS) to the pain scores during instrumentation which were statistically significant. Both our study and Karadottir *et al.* 2002<sup>[7]</sup> showed similar correlation for Q4 which suggests patients experienced more anxiety and pain when they had to come back for the treatment.

Interestingly, in Group 2, although the anxiety questions positively correlated with the pain scores, they were not statistically significant. This suggests that Q4, if you had to go to dentist tomorrow, how would you feel about it? Clearly did not apply for Group 2 population, as they were assured that the RSD will be completed within 24 h.

Although dental anxiety levels progressively reduced in Group 1, dental anxiety was still a significant predictor for pain experience as per the correlation analysis. Dental anxiety is suggested to be cognitively acquired negative-conditioned response to undesirable stimuli in dental environment influenced by family relatives, peer groups, and information media.<sup>[32]</sup> Thus, Group 1 RSD patients were possibly conditioned or cognitively learned the feeling of anxiety at each visit of their procedure, but this was not observed in Group 2 single-visit RSD patients as they possibly did not have a chance to cognitively condition their anxious experience.

Furthermore, the pain scores did not differ between the two groups. However, as with the anxiety scores, the pain scores also significantly reduced between visit 3 and visit 1; between visit 4 and visit 1 and visit 4 and visit 2 in Group 1. This observation is similar to that of Van Steenberghe *et al.* 2004, who attributed these results to the familiarity of the clinical environment, operator, and the procedure itself by the patients.<sup>[3]</sup>

Further, our findings differed from Kocher *et al.* 2005 who suggested the pain scores be zero. This can be implied to a different piezo-driven ultrasonic device used (Vector System<sup>®</sup>) in their study.<sup>[33]</sup> Thus, in this study, the use of piezoelectric scaler may have given different results compared to Kocher *et al.* 2005<sup>[33]</sup> and Karadottir *et al.* 2002.<sup>[7]</sup>

In the current study, the pain experienced during subsequent visits in Group 1 significantly reduced from visit 1. This is in agreement with Van Steenberg *et al.* 2004 who suggested that a series of factors could be responsible for such an observation.<sup>[3]</sup> The main reason being the familiarity of the clinical setting, the procedure itself, and/or the operator.

In our study, we used standardized and controlled clinical environment for all the subjects as environmental factors can influence pain perception. We scheduled all our patients during the same time of the day (Forenoon) and thus we have possibly reduced the impact of any external factors influencing pain perception. In addition, we recorded pain scores 4 h after the procedure so as to reduce any different memory effects on pain perception with longer time lapse.

In this study, we did not measure physiological aspect of anxiety such as blood pressure and heart rate which would have been an interesting additional measure to support the self-reported anxiety data.

# CONCLUSION

This study aimed to investigate and compare preprocedural anxiety levels and postprocedural pain perception in single multi-visit and conventional quadrant-wise multi-visit RSD procedures. However, the results suggested both procedures did not show any significant differences in dental anxiety levels or pain perception. Based on the results of this study, it can be concluded that in Group 1 patients, anxiety can be a significant predictor for pain. Within the limitations of this study, single-visit RSD appears to be a favorable option in highly anxious generalized chronic periodontitis subjects requiring RSD. Further, explaining the procedure seems to ease the patient's anxiety levels and pain experienced. Hence, all efforts should be made to ease their anxiety during RSD procedures.

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Nil.

## **Conflicts of interest**

There are no conflicts of interest.

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