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Research paper



Stress reduction intervention and the initial periodontal therapy outcome in chronic periodontitis patients. a randomized controlled pilot study

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Abstract

Background: Psychosocial stress, have been implicated as risk indicators for periodontal disease. Minimal evidence in literature exists to assess the effect of stress reduction therapy (SRT) in the outcome of non-surgical periodontal therapy (NSPT).

Objective: Hence, we aimed to explore the possibility of employing SRT in stressed patients as an adjunctive intervention in the management NSPT.

Materials and Methods: Sixty male patients divided into Group 1 comprised of 20 unstressed chronic periodontitis patients receiving nonsurgical periodontal therapy alone, Group 2a (20 stressed chronic periodontitis patients) received intervention focused on SPT with NSPT and Group 2b (20 stressed chronic periodontitis patients) received only NSPT without SPT. Their stress level was evaluated using a standard questionnaire method and salivary cortisol levels, at baseline and 3 months along with periodontal parameters.

Results: There was significant reduction in stress parameters for Group 2a patients. The reduction of total periodontitis affected sites was highest in Group I patients (55.4%), followed by Group 2a patients (53.2%) and Group 2b patients (38%).

Conclusion: SRT may result in comparable treatment outcome to unstressed chronic periodontitis patients. This study opens new avenues to investigate the effects of stress management as an adjunctive to conventional NSPT, which should be explored further.

Keywords: Derogatis Stress Profile Scores; Non-Surgical Periodontal Therapy; Periodontal Disease; Psychological Stress; Stress Reduction Therapy; Salivary Cortisol.

1. Introduction

Periodontitis is a multifactorial disease where microbial dental biofilms are considered to be a key etiological agent for the initiation & progression of the inflammatory process (Genco, 1996; Page and Beck, 1997; Page. 1998; Elter et al., 1999). Apart from the microbial biofilms, several other risk factors have been associated with increased susceptibility, progression and severity of periodontal diseases such as systemic diseases, genetic polymorphisms, socio-economic or educational status, tobacco smoking, oral hygiene level and psychological stress (Reners and Breex, 2007; Breivik et al., 1996).

With respect to other possible factors influencing chronic periodontitis, more direct evidence has emerged that stress, depression and anxiety contribute to the development of periodontitis in odds ratio of 1:2. Further, it has been shown that patients with stress are more prone to develop periodontal disease than patients without stress (Seiffert et al., 2002; Breivik, 2001; Bartold and Kylstra, 1994; Kiecolt-Glaser et al., 2003). It is speculated that chronic stress contribute to the development of periodontitis by having a net negative effect on the immunological response of body or by health related risk behaviors such as smoking, over eating and lessen compliance with the preventive behavior or even both (Leresche and Dworkin, 2002). In addition, it has been found that patients experiencing stress were slower in recovery from periodontal treatment compared to patients who are not experiencing stress (Leresche and Dworkin, 2002).

Interestingly, the impact of stress on the pathogenesis and periodontal treatment outcomes depends upon the individuals coping ability. Literature evidence shows that emotional-focused coping individuals (defensive coping, resigned coping, distractive coping which are advantageous in the short term) have more advanced disease and poor response to non-surgical periodontal treatment when compared to problem focused coping (i.e. active coping) (Wimmer et al., 2002). Hence, assessment of a patient's stress level, their coping ability and stress management might be of value in understanding psychological effects on periodontal health and its disease process, which will be helpful in future preventive care. Considering these facts, we hypothesize that if stress is causally related to the worsening of parameters in a chronic periodontitis patient, its alleviation might result in an additive response to the conventional periodontal therapy. However, till date, there are no intervention studies on possibility of employing psychological background. Hence, considering this hypothesis, the present study is conducted which is first of its kind, to explore and evaluate, if intervention focused on stress management enhancement



training may serve as adjunctive role in non-surgical treatment of periodontal diseases by monitoring the improvements in periodontal condition through clinical parameters and correlating with stress marker like salivary cortisol levels and Derogatis stress profile (DSP) scores.

2. Materials and methods

2.1. Source of data

It is a randomized control parallel study. The study population consisted of randomly selected sixty, systemically healthy, male patients selected out of 150 screened patients (with exclusion of 90 patients) who were screened at the Border Security Force, Yelahanka, Bangalore. The study was conducted during the period from March 2015 to November 2015. They were recruited for the study when they satisfied the following inclusion and exclusion criteria.

Inclusion criteria includes, patients within the age group of 30 to 55 years, systemically healthy individuals with chronic periodontitis and who were co-operative and willing to attend follow up visits

Exclusion criteria includes, patients on corticosteroids and antipsychotic drug therapy, who had received any periodontal therapy, surgical or non- surgical within the past six months of baseline examination, Smokers, no prior history of non-surgical periodontal therapy within 6 months, with less than 20 permanent teeth remaining, history of psychiatric treatment within past 6 months, known systemic diseases and conditions.

All eligible patients, who volunteered, were informed of the nature, potential risks and benefits of their participation in the study and a written signed informed consent was obtained from whom who agreed to participate. The ethical clearance for the study was obtained from the institutional ethical committee of Krishnadevaraya College of Dental Sciences and Hospital, affiliated to the Rajiv Gandhi University of Health Sciences, Bangalore, India.

2.2. Criteria for subject grouping

Based on the clinical parameters like modified gingival index (GI; Loe and Silness, 1963), Plaque Index (PI; Silness and Loe, 1964), periodontal probing depth (PPD), clinical attachment level (CAL), bleeding on probing (BOP; Ainamo and Bay, 1975), salivary cortisol levels and Derogatis stress profile (DSP; Derogatis, 1995), sixty subject populations were categorized into two groups. The study population of chronic generalized periodontitis subjects were selected according to the criteria proposed by the 1999 International World Workshop for a Classification of Periodontal Diseases and Conditions: chronic generalized periodontitis subjects had moderate to severe alveolar bone loss (CAL \geq 3mm and PPD \geq 5mm) in multiple sites of all four quadrants of the mouth but with no evidence of rapid progression. Group 1: Comprises of 20 chronic generalized periodontitis unstressed patients receiving non-surgical periodontal therapy (scaling and

Group 1: Comprises of 20 chronic generalized periodontitis unstressed patients receiving non-surgical periodontal therapy (scaling and root planing) alone.

All patients who showed clinical signs of gingival inflammation and attachment loss, gingival index ≥ 2 , plaque index 2.0 - 3.0, gingival bleeding index : > 1, bleeding present on probing within 10 seconds.

Group 2: Comprises of 40 chronic periodontitis patients associated with stress and clinical signs of gingival inflammation and attachment loss (gingival index ≥ 2 , plaque index 2.0 – 3.0, gingival bleeding index: > 1, bleeding on probing within 10 seconds, PPD ≥ 5 mm, clinical attachment level ≥ 3 mm, Radiographic evidence of bone loss, Derogatis stress profile 77 questionnaire score 0.72 - 0.90, cortisol levels > 2.5 ng/ml (AM; stressed).

They were further subdivided by simple randomization of computer software program that generates the random sequence method into two sub groups of 20 each, based on the intervention (SRT): enclose assignments in sequentially numbered, sealed envelopes. Examiner (1) generated random allocation sequence and carried out post intervention clinical examination of periodontal parameters, second examiner (2) Enrolled the participants for the study when they satisfied the inclusion and exclusion criteria. Carried out case history taking, initial clinical examination of periodontal parameters and radiographic examination to group the participants, third examiner (3) assigned participant to intervention and examiner (4) performed scaling and root planning, examiner (5) performed supportive periodontal therapy and examiner (6) performed and evaluated the stress reduction therapy. It is a double blind study where all the examiners and participants, both were blinded.2a: Comprises of 20 stressed chronic periodontil therapy. 2b: Comprises of 20 stressed chronic stress, who received intervention focused on SRT along with non-surgical periodontal therapy (SRP) without intervention focused on SRT. A sequential flow chart for the study protocol is shown in Figure.1.



Figure. 1: The Sequential Flow Chart for Study Protocol.

2.3. Pre-sampling clinical evaluation of patients

Patients were selected for each group after a brief and precise case history recording that included patient's chief complaint, clinical examination and radiographic evaluation. Demographic characteristics, such as age, sex, diet and medical history were also recorded.

2.4. Systemic examination

Assessment of Blood pressure (BP), Body Mass Index (BMI) and hematological testing for Fasting Blood Sugar (FBS) was carried out for all the patients, to rule out the presence of unidentified systemic conditions such as cardiovascular diseases and diabetes mellitus, which can potentially influence cortisol levels. The BP was recorded by using a mercury sphygmomanometer. BMI was evaluated by use of standard height and weight scales (Misra et al., 2009).

2.5. Intra-oral examination

Clinical assessment of the oral hygiene and gingival status was done by plaque index, bleeding on probing index and gingival index. These indices were repeated at 90th day for assessment of oral hygiene maintenance and presence or absence of gingival inflammation. For each subject full-mouth periodontal probing and charting was recorded using UNC-15 probe. PPD and CAL were measured on six sites (mesio-buccal, mesio-lingual, mid-lingual and disto-lingual) per tooth using UNC-15.

The measurement of REC (recession), PPD and clinical attachment level (CAL) was done according to Pilgram et al., (2000). REC was measured from the CEJ to the gingival margin, with a positive value if there was recession and a negative value in the absence of recession; CAL was calculated by summation of PPD and REC. The clinical diagnosis of chronic periodontitis was made based on the criteria described by Armitage, 1999. CAL was stratified into 3 ordered categories: slight: 1 - 2 mm, moderate: 2 - 4, severe: > 5 mm.

2.6. Examiner calibration for periodontal examination

Before recording probing depths calibration exercise for clinical parameters was performed in five patients before the actual study. Examiners 1 and 2 conducted the recordings for clinical parameters at 1 week interval. The order of patients was changed in between the examinations, by a third examiner (3) who was blinded to the whole process. The periodontal probing depth estimation was judged to be reproducible if the intra and inter-examiner agreement within ± 1 mm between repeated measurements was at least 80%. The kappa value for

intra-examiner agreement for examiner 1 and 2, between the two measurements was recorded to be 0.91 and 0.94 respectively. The interexaminer calibration was recorded to be 0.84.

The above-mentioned parameters were recorded at baseline and at follow-up visits scheduled at 90th day, following initial therapy by prealigned and assessed examiners BVK.

2.7. Stress assessment

The Derogatis stress profile (DSP) is a self-report measure designed to assess chronic stress. This 77-item inventory assesses 11 components of stress, including vocational environment, domestic environment, health environment, time pressure, driven behavior, attitude posture, relaxation potential, role definition, hostility, anxiety, and depression. Participants rated each statement from 0 (not at all true of me) to 4 (extremely true of me). The DSP contains items such as, I take some time out almost every day just to relax, and I am usually worried about something.

Stressed and non-stressed patients were differentiated by Derogatis stress profile 77 questionnaire. The scale takes approximately 12-13 minutes to complete. Stress score range is 0.72 - 0.90. (< 0.72 - unstressed, 0.72 to 0.90 - stressed, 0.90 - severe stressed)

A professional recorded the clinical data before commencing stress reduction therapy. The stress questionnaires were issued by psychologist involved in psychological assessment & analysis and who is fluent in local language and English. Instructions were explained and patients were arranged to complete the questionnaires in a calm environment. The examiner 1 was standby to clarify any queries arising during completion of the questionnaires. All the questionnaires were designed in self-administered format. For those illiterate or marginally literate patients, who were mainly from the older age groups, their questionnaires were completed in an interviewer assisted format.

Patients were asked to complete a questionnaire including the following sections: 1) demographic and socio-economic details; 2) medical history – reporting symptom and diagnosed systemic diseases; 3) dental habits and dental care utilization; and 4) history of cigarette smoking and exposure to occupational hazards. Patients were then given a set of self-administered stress questionnaires in a face-to-face interview with examiner 1.

Salivary sample collection was done according to the passive drool method (Dawes et al., 2000) and samples were collected in plastic vials and centrifuged for 20 min at 3000 rpm to remove bacteria and cellular debris. The supernatant was pipetted out and stored at -20°C for further analysis.

2.8. Intervention therapy

2.8.1. Periodontal treatment

All patients of Group 1 and Groups 2a and 2b were first submitted to the hygiene phase of the periodontal therapy, which includes supragingival plaque and calculus removal, provisional restoration, and the removal of overhanging fillings. Scaling and root planing (SRP) was performed under local anesthesia (2% lignocaine hydrochloride with 1:200000 adrenaline) using area specific Gracey periodontal curettes and an ultrasonic device. Within the duration of the study, all patients received supportive therapy, which included professional plaque control and oral hygiene information.

The treatment was concluded within 7 days without the use of antibiotics or local antimicrobials. The patients were monitored by assessment of clinical parameters at baseline and at 90th day of therapy.

2.8.2. Stress reduction therapy

After the assessment of clinical, stress and coping parameters and completion of non- surgical periodontal therapy, intervention for stress reduction was commenced for Group 2a patients through transactional model protocols.

Transactional model is an adept framework for treatment and evaluation of stress through problem solving and emotional coping methods. It is considered as a treatment of choice for stressed patients confronting both unmanageable and manageable stressor events. The patient's evaluation of potential gravity of the situation (primary appraisal) and his/her ability to beneficially alter the situations and emotions (secondary appraisal), both play a crucial part in assessing the coping efforts which are required and their obtained outcomes. Since, the use of Transactional model adjudicates the degree, duration and type of coping efforts required, it was used to assess, evaluate and accordingly counsel the stressed patients.

As a part of the Transactional Model modus operandi, a course of seven sessions were held for Group 2a patients, distributed uniformly over a period of six months (two sessions per month, totally 7 sessions). These sessions of approximately 45 minutes duration, were held in a setting of calm and pleasant environment.

Session 1: Day 0

The subjects (5 patients at a time) were educated for better impact of our intervention regarding stress, its primary etiologic agent and its ill effects on periodontal & systemic health through lectures, images and video tapes (Powerpoint slides). Further, patients were motivated to employ stress reduction protocols such as assertion training via diary writing and relaxation techniques including breathing exercises, meditation, yoga, musical detours etc., as a daily practice. To keep a check on patient compliance, the patients were provided with a pamphlet, pictorially depicting the protocols they were supposed to follow. They were supposed to mark a tick against the stress reduction practices they followed each day. No compulsion was made for them to strictly adhere to these practices, to avoid additional stress. The patients were asked to bring back the marked pamphlet in the next session.

Session 2: Day 15

In the next session (15th day), the patient's ability to manage and alter the stressful situation was assessed through cope questionnaire individually along with explanation of the benefits of adequate coping with stress (Stress reduction techniques reinforcement). The patient compliance to the suggested protocol was evaluated by assessing their daily attempts towards stress reduction, through the pamphlet they were asked to bring back. Execution of even one stress reduction protocol daily was considered as a positive marker for subjective compliance and improvisation.

Subjects unwilling to follow the stress reduction protocols were excluded from the study. Motivational reinforcement of the coping measures was carried out for subjects, who were unable to adhere to the protocol due to personal problems. Telephonic communications with patients were also made to remotely observe and guide the subjects through the protocol. Session 3: Day 30

In manageable stressor incidences, coping strategies directed at changing a stressful situation were advised to the patient (problem solving coping). As an example, the subject A 36 years old man, had faced a problem of financial crisis to pay the hire for his vehicle and home because of his inability or severe illness and because of that situation he is worried and tensed about that situation which caused stress mentally to guide him towards smoking, drinking and other habits which are injurious to systemic health. Because of this financial stress, ignore concern towards his family, health and surrounding environment and he considered himself as he cannot solve his problem.

In cases of unmanageable stressor incidences, coping strategies directed towards an alternative solutions to their problems, such as obtaining advices from close associates, confiding in trustworthy acquaintances and availing assistance from supporters were suggested (emotional coping).Ex: As an example, the subject A 32 year's old man had faced a problem of sudden death of his father and mother in an accident. He felt that himself as an isolated from family and loving ones and went into depression along with ignoring health and addicted to bad habits such as drinking, smoking to forget that worst situation because of his inability to manage his rest of life without them. Sessions 4, 5 & 6: Day 45, 60 and 75

Reinforcement of the benefits of adequate coping with stress (Stress reduction techniques reinforcement). In addition, Oral Hygiene instructions were concreted to the patients.

Session 7: Day 90

In this session, Group 1, 2a and 2b patients were recalled back. The Group 2a and 2b patient's ability to manage and alter the stressful situation was assessed through cope questionnaire. The compliance of Group 2a subjects to stress reduction practices was judged through pamphlets, which they were supposed to bring back.

The reassessment for the stressed subjects was carried out after every 15 days by the examiner 1 within 13 - 15 minutes of time period and interpreting the values by the patient's selected answers in the followed scale and analyzing whether the patients are following or not about our therapy protocol and what actually there are doing. This included their evaluation for use of coping techniques and reinforcement of stress reduction protocols.

Biochemical assessment of cortisol

Samples were assayed for cortisol levels by using chemiluminescence immunoassay (CLIA) by microplateluminometers provides a sensitive, high throughput and economical alternative to conventional colorimetric methodologies, such as enzyme-linked immunosorbent assays (ELISA).. This method has a reported sensitivity of $< 0.5 \mu$ g/ml. Results were reported as the total amount in micrograms per milliliter of sample (μ g/ml).

2.9. Method of statistical analysis

One-way analyses of variance were used to test the difference between groups. Analysis of Variance is a technique by which the total variation is split into two parts one between groups and the other within the groups. The Student 't' test was used to determine whether there was a statistical difference between groups in the parameters measured. The association between clinical parameter and DSP and salivary cortisol were assessed by PERSONS CORRELATION coefficient. Normality of data was tested using Shapiro-Wilk test. The results were averaged (mean + standard deviation) for continuous data are presented in Table 1 to 4. In the entire above test p value less than 0.05 were taken to be statistically significant. The data was analyzed using SPSS package (ver 10.5).

3. Results

From the 150 subjects enrolled in this randomized control parallel study from March 2015 to November 2015, 150 were screened and 60 were included in the present study. Approximately 54% (N = 80) of the enrolled subjects were lost to not meeting the inclusion criteria, 6.7% (N = 10) were refused to participate, in included subjects 3.3% (N = 2) were lost to follow up and 3.3% (N = 2) were discontinued intervention (Fig. 1). Power of the study is 80% and the estimated effect size and precision was set at 95% confidence interval.

Included 40 patients who were randomly assigned into Group 2a (20) and Group 2b (20). All the participants received intended intervention (SRP) and where analyzed for the primary outcome (improvement in Clinical Parameters). When compared from the baseline, a significant reduction was noted at three months in the mean GI and PI scores for group 2a once only (p < 0.001). However, there was a significant reduction in BOP in all the groups from baseline to 3 months (p < 0.001).

Analytical assessment for the mean number of PD sites for all the groups revealed a significant reduction after 3 months of intervention (p < 0.001). At end of 3 months, the highest overall reduction in pocket depth was seen in Group 1 (2.32; p < 0.001), least in Group 2b (0.29; p > 0.05) and intermediate in Group 2a (1.95; p < 0.001). Likewise improvement in the CAL levels was significantly higher in Group 1 (1.16; p < 0.001) and least in Group 2b (0.15; p > 0.05) but Group 2a values (1.11; p < 0.001) were intermediate between the highest and lowest values. These values suggest that after periodontal and psychological intervention there is a significant improvement in the clinical parameters and stress levels of Group 2a chronic stressed periodontitis patients when compared with the Group 2b chronic stressed periodontitis patients without any stress reduction procedure.

When the mean gain in the CAL was assessed among Groups 1, 2a and 2b, using ANOVA test, the difference of mean was shown to be statistically significant at baseline with p value < 0.001. Further pairwise intergroup analysis by t-test revealed that at the end of 3 months, significant gain in the sites with CAL was obtained in all the groups except on ≥ 5 mm group of Group 2b.

A weak positive correlation between DSP scores and CAL found in the Group 2a of 3 - 4 mm, Group 1 and Group 2a of \geq 5 mm the at the base line and end of the third month and between the salivary cortisol levels and pocket depths at the base line and end of the third month, a weak positive correlation found in the Group 2b of 4 - 6 mm, Group 1 and Group 2b of > 8 mm. But a strong positive correlation found in the Group 2a of 4 - 6 mm, Group 1 and Group 2b of > 8 mm. But a strong positive correlation found in the Group 1 of 1-2 mm, Group 2b of \geq 5 mm. But a strong positive correlation found in the Group 1 of 1-2 mm, Group 2b of \geq 5 mm. But a strong positive correlation found in the Group 1 of \geq 5 mm at the base line and end of the third month.

In summary, there was a significant mean reduction in salivary cortisol levels, DSP scores, GI, PI, BOP, PD, CAL of Group 2a when stress reduction therapy with periodontal intervention were given when compared to other groups, where only periodontal intervention was given. This indicates in patients under psychological stress when SRT was used as an adjunctive, there was a statistically significant outcome to NSPT (Table 1-4 & Figures 2, 3). There was no associated untoward effects in all the study groups.

Table 1: The Mean Values, Standard Deviation (SD), Minimum and Maximum Values for Salivary Cortisol Levels and DSP Scores

Salivary Cortisol Levels	Baseline					3 months					
	Ν	Mean	SD	Min.	Max.	Ν	Mean	SD	Min.	Max.	
Group 1	20	2.12	0.25	1.65	2.46	20	1.98	0.20	1.51	2.31	
Group 2a	20	3.46	0.69	2.52	5.10	16	2.60	0.58	1.78	2.99	
Group 2b	20	3.26	0.50	2.55	4.50	20	2.85	0.31	2.35	3.54	
DSP Scores	Baselii	Baseline					3 months				
	Ν	Mean	SD	Min.	Max.	Ν	Mean	SD	Min.	Max.	
Group 1	20	0.48	0.13	0.25	0.71	20	0.46	0.12	0.28	0.70	
Group 2a	20	2.25	0.56	1.10	2.94	16	1.35	0.39	0.98	2.11	
Group 2b	20	1.95	0.60	0.90	2.73	20	1.84	0.57	0.91	2.72	

Table 2: The Mean Values, Standard Deviation (SD), Minimum and Maximum Values for PD, CAL

	Group		Mean	SD	Min	Max	"P" value	Mean diff
PD	Crown 1	Baseline	6.307	1.9172	.0	9.0	-0.001*	2.32
	Group I	3 month	3.980	1.4872	.0	7.5	<0.001*	
	Crown 2a	Baseline	5.377	1.6352	3.2	8.4	< 0.001*	1.95
	Group 2a	3 month	3.428	1.2045	1.5	5.9		
	Crown 2h	Baseline	6.982	1.3384	4.7	9.2	<0.001*	0.29
	Group 20	3 month	6.698	1.5379	3.4	9.1		
CAL	Crown 1	Baseline	3.174	1.7350	.0	6.6	<0.001	1.16
	Group I	3 month	2.013	1.4022	.0	5.0	<0.001	
	C	Baseline	3.672	1.6692	1.4	6.6	-0.001*	1 1 1
	Group 2a	3 month	2.468	1.4199	.3	4.3	<0.001**	1.11
	Course 2h	Baseline	3.763	1.7460	1.4	6.7	-0.001*	0.15
	Group 26	3 month	3.610	1.7886	1.2	6.6	<0.001*	0.15

• Statistically significant.

Table 3: The Mean Values, Standard Deviation (SD), Minimum and Maximum Values for GI, BOP, PI

GI	Baseline						3 months				
	Mean	SD	Median	Min.	Max.	Mean	SD	Median	Min.	Max.	
Group 1	1.224	0.355	1.115	0.77	1.91	1.042	0.355	1.000	0.53	1.81	
Group 2a	1.476	0.366	1.420	0.87	2.07	1.055	0.320	1.010	0.32	1.61	
Group 2b	1.116	0.160	1.075	0.97	1.66	1.045	0.127	1.010	0.87	1.46	
BOP	Baseline			3 months							
	Mean	SD	Median	Min.	Max.	Mean	SD	Median	Min.	Max.	
Group 1	4.433	1.045	4.895	2.10	5.70	1.695	0.768	1.610	0.69	3.33	
Group 2a	4.567	0.996	4.985	2.12	5.71	1.755	0.646	1.765	0.80	2.76	
Group 2b	4.531	0.933	4.895	2.12	5.71	3.414	1.046	3.670	1.28	5.09	
PI	Baseline			3 months							
	Mean	SD	Median	Min.	Max.	Mean	SD	Median	Min.	Max.	
Group 1	1.559	0.688	1.370	0.33	2.75	1.330	0.609	1.175	0.32	2.20	
Group 2a	1.747	0.546	1.720	0.91	2.89	1.211	0.374	1.025	0.71	1.89	
Group 2b	1.652	0.404	1.620	1.04	2.29	1.603	0.394	1.585	1.01	2.20	

 Table 4: Results of Pearson'S Correlation Coefficient (R) Test Compare Salivary Cortisol Level, Pocket Depths and Clinical Attachment Levels within the Groups at Baseline and End of Third Month

Pearson's correlation coefficient (r) test compare salivary cortisol level and pocket depths									
		Group 1	Group 2a	Group 2b					
Baseline	R	-0.122	0.093	-0.267					
	p' value	0.608	0.697	0.256					
3 month	R	-0.139	-0.445	-0.141					
	p' value	0.560	0.084	0.553					
Pearson's correlation coefficient (r) test compare salivary cortisol level and clinical attachment levels									
Baseline	R	-0.047	0.166	0.280					
	p' value	0.843	0.484	0.232					
3 month	R	-0.214	0.498	-0.018					
	p' value	0.366	0.049	0.940					

Pearson coefficient of correlation.

If the (r) value.

Is -1, there is a perfect negative correlation.

Falls between -1 and -0.5, there is a strong negative correlation.

Falls between -0.5 and 0, there is a weak negative correlation.

Is 0, there is no correlation.

Falls between 0 and 0.5, there is a weak positive correlation.

Falls between 0.5 and 1, there is a strong positive correlation.

Is 1, there is a perfect positive correlation between the 2 sets of data.



Fig. 2: Correlation of Pocket Depths and Clinical Attachment Levels (No. of Sites) in All the Groups from Baseline to 3 Months.





Fig. 3: Comparison of Gingival Index, Plaque Index and Bleeding on Probing in Study Groups before and after Treatment

4. Discussion

Psychosocial stress is being considered to play an important role in the pathogenesis of periodontal diseases and outcome of its treatment (Genco et al., 2008). Various mechanisms have been put forward to explain the role of stress in the development of periodontal disease. First, stress-induces a response that is transmitted to the hypothalamo-pituitary-adrenal (HPA) axis and promotes the release of corticotrophin-releasing hormone from the pituitary gland and glucocorticoid hormones from the adrenal cortex, which in turn induces the reduction of proinflammatory cytokines secretions (immunosuppressive effect). Second, exposure to stressor agents can induce the sympathetic nervous system to release adrenaline and noradrenaline which stimulates the formation and activity of prostaglandins and proteolytic enzymes, leading to periodontal tissue breakdown indirectly (Breivik and Thrane, 2001; Genco et al., 2008). Third, stress can induce the release of neuropeptides from sensory nerve fibers (neurogenic inflammation), whose presence has been implicated as a neurogenic promoter in various inflammatory processes modulating the activity of the immune system and the release of cytokines. Fourth, the behavioral model suggests that psychosocial stress may influence behavioral changes which affect health behaviors (i.e., negligent oral hygiene, intensification of smoking, poor compliance). Fifth, stress leads to overeating, especially high fat diets which increases cortisol production (Ross et al., 2014; Genco et al., 2008).

Contextually, studies to date imperatively suggest that if patients adapt successful coping strategies to reduce, control or tolerate the state of stress, there can be moderation on the impact of stress in the development of periodontal disease. Stressed periodontitis patients with maladaptive coping strategies (emotion focused coping-'avoidance') are at greater risk for developing severe periodontal disease and showed poorer response to a non-surgical periodontal treatment. Whereas, those with active coping strategies (problem solving coping-'make a plan of action), are associated with high levels of wellbeing and found to have less periodontal tissues destruction and showed

good response towards periodontal treatment. This suggests that a subject's responses to a stressor may be a key determinant in moderating the effect of stress on the progression of periodontal disease and outcome of the periodontal therapy (Wimmer et al., 2005).

If indeed stress and inadequate coping are important risk indicator for periodontal disease then to what extent intervention in stress reduction will have an effect in moderating these are unexplored till date. In view of this, the present study, which is first of its kind was undertaken to explore and evaluate intervention focused on employing psychological intervention (stress reduction therapy) as an adjunctive to the non-surgical periodontal therapy (NSPT). The patients were categorized as stressed or unstressed based on the salivary cortisol levels and DSP scores, as these are considered effective biological and psychological markers of stress (Vining et al., 1983). Studies have reported that the sources of stress and type of coping engaged by different genders and among the younger and elder individuals are categorically different; this might have an influence in the outcome of the study. Hence, to eliminate these confounding factors like age and gender differences, only the male, non-smoker individuals in the range of 30-55 years were recruited.

Since, the study participants were screened from Border Security Force (constitutes solely male population). the study group constituted only male participants.

In our study, transactional model of coping (Moss, 1999) was adapted, as a means of stress intervention. According to the Transactional Model, emotional and functional effects of primary and secondary appraisals are mediated by actual coping strategies. Original formulations of the model conceptualized coping efforts along two dimensions: (1) problem management and (2) emotional regulation. The problem-focused coping/problem-management strategies are directed at changing the stressful situation with the use of mechanisms such as active coping, problem solving, and information seeking. By contrast, emotion-focused coping efforts are directed at changing the way one thinks or feels about a stressful situation, by incorporating strategies such as seeking social support and venting feelings, as well as avoid-ance, and denial. The model predicts that problem-focused coping strategies will be most adaptive for stressors that are changeable, whereas emotion-focused strategies are most adaptive when the stressor is unchangeable or when this strategy is used in conjunction with problem-focused coping strategies. Irrespective of the coping style the patients were following, they were guided through the transactional model towards problem or emotional focused coping, as the situation warranted. Further, the patients were assessed periodically for their compliance and were directed towards effective stress management and oral hygiene behavior throughout the study period.

This indicates that the stressed patients undergoing NSPT + SRT demonstrated significantly higher associated reduction in the stress and clinical parameters when compared to stressed patients undergoing NSPT alone. It can also be observed that although the stress parameter reductions in Group 2a patients remained above the normal range, these patients with concomitant SRT showed a similar reduction in periodontitis affected sites as the unstressed patients. However, it was the unstressed Group 1 patients, who showed highest improvement in clinical parameters.

The mechanism whereby SRT may improve the outcome of periodontal treatment is not clear. The protocol of the present study included attempts to remodel patient's behavior and coping strategies along with non-surgical periodontal therapy, thus creating a three-sided approach towards disease management. Firstly, a more active participation in problem solving could have encouraged a more dynamic participation of the patients in day to day activities, including oral hygiene maintenance. Secondly, the decrease in oral inflammatory load after NSPT and periodic supportive periodontal therapy promoted decrease in periodontal inflammation. Thirdly, with the commencement of stress intervening programs, physiologic parameters favoring periodontal disease progression were controlled.

The results of our study are in similar lines with the previous studies, which propose that if the patient adapts to effective coping strategies, there could be moderation of the impact of stress on the periodontal tissues, thereby beneficially influencing the treatment outcomes as well. Wimmer et al. (2005) showed that passive coping strategies were more pronounced in advanced disease as well as in cases of poor response to a non-surgical periodontal treatment, whereas patients with active coping modes had milder disease and a more favorable course of treatment. Further, Gamboa et al. (2005) demonstrated that patients with good emotional intelligence (EI, a psychological construct) i.e., effective coping mechanism, is associated with favorable short-term changes in plaque and bleeding and the possibility of developing an intervention aimed at improving patients' EI was advised, which may improve response to initial treatment of periodontal disease. This suggests that subject's response to the stressors may be a key determinant of the overall effect of the stress.

Cortisol is a well-known stress-related hormone that can be detected in blood, saliva and gingival crevicular fluid (GCF). Salivary level of cortisol reliably reflects HPA axis activity and is used in human psychological studies as a biological marker of stress (Koh et al., 2007). In our study, salivary cortisol level was used as stress parameter because, it's an easy, non-invasive, rapid, stable, non-clotting medium, which requires less manpower and materials than other procedures. However, to compensate for diurnal variation, single-sample determination was done for all the patients at the same time of the day (Gharavi, 2008). The levels of cortisol in saliva were estimated by CLIA (glow-based chemiluminescent reaction) which provides a sensitive, high throughput, and economical alternative to conventional colorimetric methodologies, such as Enzyme-linked immunosorbent assays (ELISA). Further, it provides a broader dynamic assay range, superior low-end sensitivity, and a faster protocol than the conventional colorimetric methods (Gharavi, 2008). The DSP questionnaire is used to assess the degree of psychological stress, as it is one of the most comprehensive stress questionnaires designed to assess the chronic stress. The reliability of the questionnaire has been confirmed and it has also been proven that it requires no further random samples for standardization (Derogatis, 1995).

The three months follow up period adopted in the study was sufficient to demonstrate short-term changes in the clinical parameters. However, it is not sufficient to conclude for the long term and permanent changes. The results of the present study should thus, be interpreted with caution. The small, homogenous samples with short follow up duration, exclusion of female individuals, lack of random assignment to study groups, restriction to middle class or lower middle class families (those who were subjected to high strain and stress), limits the generalizability of the findings. Ideally, a study would use a set of biomarkers (Cortisol, DHEA, adrenaline, noradrenaline, dopamine and aldosterone) to measure stress response, while using questionnaires to measure stressor exposure and stress appraisal. In our study we used only a single stress biomarker (Cortisol) to objectively assess stress which could have led to some unknown confounding factors.

Future periodontal studies should be conducted with a larger sample size and longer follow up, including both male and female patients, with different lifestyles and family status, testing alternative psychological models of stress reduction in relation to adherence to treatment. Control groups should include lack of periodontal maintenance with various periodontal diseases. Integrated clinical, sociological and molecular based (biological markers, C-reactive protein) studies are needed very much for understanding the role of stress reduction therapy on the outcome of periodontal treatment.

5. Conclusion

To conclude, the result of our study is the first evidence to suggest that psychological intervention focused on stress reduction resulted in the same order of magnitude in the outcome as NSPT for the stressed chronic periodontitis patients when compared to unstressed chronic

periodontitis patients. Hence, the finding of our study raises the possibility of developing an adjunctive intervention based on SRT for the holistic management of stressed patients undergoing periodontal treatment. However, more intervention studies are needed before psychological stress can be firmly established as been importance in the treatment or prevention of periodontal disease.

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