Interrelationship of Smoking, Lip and Gingival Melanin Pigmentation, and Periodontal Status

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Abstract

Background: Cigarette smoking is a significant risk factor for periodontal disease. It also causes pigmentation of oral mucosa. The present study was aimed to assess the effects of smoking on lip and gingival pigmentation and periodontal status and the relationship between pigmentation and periodontal parameters.

Methods: A total of 109 smokers and an equal number of non-smoker controls (mean age: 35.9 years, range: 35-44 years) comprised the study sample. All the participants were assessed for pigmentation on lip and gingiva and overall periodontal status (gingival bleeding, probing depth, and loss of attachment at six points in each tooth).

Findings: All the smokers in this study had lip and gingival pigmentation. Two-third of non-smokers had no pigmentation. The mean scores of lip and gingival pigmentation in smokers were seven and four times higher than those of non-smokers, respectively. Pigmentation and periodontal parameters (except gingival bleeding) were found to be positively related with exposure to smoking exposure. Probing depth and loss of attachment were the highest in subjects with pigmented lips and grade three pigmented gingiva.

Conclusion: Smoking influenced lip and gingival pigmentation and periodontium. All individuals with lip pigmentation presented some form of gingival pigmentation. Probing depth and loss of attachment were more severe in subjects with lip and gingival pigmentation.

Keywords: Oral mucosa, Pigmentation, Smoking, Periodontium

Citation: Multani S. Interrelationship of Smoking, Lip and Gingival Melanin Pigmentation, and Periodontal Status. Addict Health 2013; 5(1-2): 57-65.

Received: 31.07.2012 Accepted: 11.11.2012
Introduction

Smoking seems to have many adverse effects on the body. It is evident from the past literature that cigarette smoking is a significant risk factor for periodontal disease which is demonstrated by an increased loss of attachment, development and progression of periodontal inflammation, and increased gingival recession. Epidemiological evidence suggests that cigarette smoking is a stronger risk factor for the presence of periodontitis compared to the presence of certain suspected periodontal pathogens. However, experimental studies on plaque-induced gingivitis in humans suggest that clinical signs of gingival inflammation, e.g. redness, bleeding, and exudation, are not as prominent in smokers as in non-smokers. Decreased inflammation and gingival crevicular fluid volumes in smokers compared to non-smokers indicates the impairment of gingival blood flow by smoking.

In addition to periodontal destruction, a probable adverse effect of smoking on oral cavity is the pigmentation of oral mucosa. Apart from smoking, oral pigmentation has been associated with a variety of endogenous and exogenous etiologic factors. Pigmentation is mostly caused by five primary pigments including melanin, melanoid, oxyhemoglobin, hemoglobin, and carotene. Meanwhile, a benign focal pigmentation of the oral mucosa, smoker’s melanosis, has been attributed to smoking behavior. Melanin production in gingival tissue is stimulated as a result of the high content of nicotine and benzpyrene in tobacco smoke. On the other hand, disappearance of gingival pigmentation was observed following the reduction in smoking. Clinically, the lesion usually presents as multiple brown pigmented macules less than 1 cm in diameter, localized mainly at the attached anterior labial gingiva and the interdental papillae of the mandible.

Axel and Hedin first described oral pigmentation including lip pigmentation in 1982. Since then, there were no reports except for a recent study by Haresaku et al. who observed an association between lip and gingival pigmentation. In contrast to the sparse studies on lip pigmentation, extensive attention has been paid to gingival pigmentation during the past half a century. As the relationships between pigmentation and periodontal parameters have not been evaluated before, the present study sought to establish and compare the effects of smoking on lip and gingival pigmentation and periodontal status and to assess the relationship between pigmentation and periodontal parameters.

Methods

The present study was conducted on 218 individuals attending the outpatient department of public health dentistry (Chhattisgarh Dental College and Research Institute, Rajnandgaon, India). All the subjects were randomly selected from the outpatient register during August 2012. Totally, 109 smokers and an equal number of non-smoker control subjects aging 35-44 years (mean age: 39.5 years) were selected as the final sample. The controls were matched with the smokers in terms of age and sex. Individuals suffering from nutritional deficiencies and systemic disorders that would cause oral pigmentation were excluded from the study. The inclusion criteria for control subjects were not using tobacco products and not having a history of periodontal therapy.

The study protocol was approved by the ethics committee of the mentioned institute. Written informed consent was obtained from all participants. All the measurements were carried out by a single examiner who had been calibrated with a senior examiner in a one-week pilot study of 30 participants. The method of examination and scoring was standardized in the department of public health dentistry of Chhattisgarh Dental College and Research Institute and an intra-examiner reliability of 90% was obtained. The questionnaire was presented in a pilot survey and kappa (κ) and weighted kappa (κw) were used to evaluate the test-retest reliability of the questionnaire. Internal consistency was assessed by Cronbach's alpha (α) (κ = 0.86, κw = 0.9, and α = 0.78). Each examination took about 6-10 minutes.

Clinical examination was conducted by a single examiner who assessed the existence of lip and gingival pigmentation and periodontal problems. The absence and presence of blackish or brownish lip pigmentation were scored as zero and one, respectively. Each lip was divided into three sections and scores from each section were evaluated. The total score for each individual was
calculated as the sum of scores from the three sections.16

Gingival pigmentation in each jaw was scored based on the classification of melanin index proposed by Hedin.17 Accordingly, pigmentation was scored as zero for no pigmentation, one for one or two solitary unit(s) of pigmentation in papillary gingiva without formation of a continuous ribbon between solitary units, and two for more than three units of pigmentation in papillary gingiva without formation of continuous ribbon. Scores three and four corresponded to one or more short continuous ribbons of pigmentation and one continuous ribbon including the entire area between canines, respectively. Total score of upper and lower arches were taken and the final score was calculated by summing the scores. Examination of upper and lower gingiva was done with the help of a plane mouth mirror.

A full periodontal examination including measurements of gingival bleeding, probing depth, and loss of attachment in each tooth was performed. Data about the probing depth and loss of attachment was obtained from the measurements conducted by a senior clinician. A calibrated Williams graduated periodontal probe (Hu-Friedy, US) with circumferential lines at 1-3, 5, and 7-10 mm was used to assess mesiobuccal, mid-buccal, distobuccal, mesiolingual, midlingual and distolingual measurements of each tooth. The presence of gingival bleeding and the percentage of sites with bleeding was also evaluated for each individual.

Data regarding the smoking behavior (i.e. duration and frequency) was obtained through individual interviews with the participants. Then, the subjects were classified as non-smokers, current smokers, occasional smokers, and ex-smokers. The smokers were further categorized based on the duration of tobacco use (less than five years, 6-10 years, 11-20 years, and more than 20 years). Number of cigarettes smoked in a regular day was also classified as 1-2/day, 3-5/day, 6-10/day, and more than 10/day.

Current smokers constituted individuals who smoking at least once a day at the time of study. Non-smokers were subjects who had never smoked cigarettes. Occasional smokers smoked at least three consecutive days a week and former smokers had not used tobacco products for at least one year.

Statistical analysis
All analyses were performed using SPSS for Windows 15.0 (SPSS Inc., Chicago, IL, USA). Descriptive data was presented as means and standard deviation. Chi-square test was executed to find statistical differences in the distribution of pigmentation based on smoking status. Dunnett’s test was used for multiple comparisons where mean differences in pigmentation and periodontal scores between non-smokers (as the reference category) and other groups were evaluated. Spearman correlation assessed the strength of the relationships of smoking status with pigmentation and periodontal status. A contingency table was constructed to observe the concurrence of scores of lip and gingival pigmentation. Statistical differences in periodontal parameters based on lip and gingival pigmentation grades in maxillary and mandibular gingiva were evaluated using Mann-Whitney and Kruskal-Wallis tests, respectively.

Results
Current smokers constituted a major proportion (58%) among the smokers. It is clear from table 1 that melanin pigmentation on lips and gingiva were observed in all the individuals except for one occasional smoker who did not exhibit gingival pigmentation. Chi-square test revealed significant differences in the presence of pigmentation between the smokers and non-smokers. While nearly two-thirds of the non-smokers had no lip pigmentation, only 26.6% did not exhibit gingival pigmentation.

On the other hand, the above-mentioned parameters of smoking were inversely related with bleeding and positively related with probing depth and loss of attachment. According to Dunnett’s test, probing depth and loss of attachment were significantly higher and bleeding was significantly lower in smokers than in non-smokers (Table 2).

None of current smokers reported lack of pigmentation. All individuals with lip pigmentation had a clear presence of gingival pigmentation. While
### Table 1. Frequency of lip and gingival pigmentation in the participants according to smoking status

| Smoking status       | Lip pigmentation | Gingival pigmentation | Total |
|----------------------|------------------|-----------------------|-------|
|                      | Absent | Present | Absent | Present |          |
| Non-smokers          | 75 (68.8%) | 34 (31.2%) | 29 (26.6%) | 80 (73.4%) | 109     |
| Ex-smokers           | 0 | 10 (100%) | 0 | 10 (100%) | 10      |
| Occasional smokers   | 0 | 36 (100%) | 1 (2.8%) | 35 (97.2%) | 36      |
| Current smokers      | 0 | 63 (100%) | 0 | 63 (100%) | 63      |
| Statistical parameters | $\chi^2 = 113.561$ | $\chi^2 = 30.183$ |          |          |          |

| Lip pigmentation Mean ± SD | Gingival pigmentation Mean ± SD | Bleeding Mean ± SD | Probing depth Mean ± SD | Loss of attachment Mean ± SD |
|----------------------------|---------------------------------|-------------------|------------------------|-----------------------------|
| Non-smoker                 | 0.10 ± 0.17                     | 0.42 ± 0.07       | 0.07 ± 0.08*           | 2.06 ± 0.16*                | 0.02 ± 0.04                 |
| Ex-smoker                  | 0.44 ± 0.20*                    | 1.77 ± 0.30       | 0.30 ± 0.14*           | 1.84 ± 0.47*                | 0.47 ± 0.51*                |
| Occasional smoker          | 0.48 ± 0.30*                    | 0.91 ± 0.28       | 0.33 ± 0.18*           | 1.67 ± 0.56*                | 0.50 ± 0.30*                |
| Current smoker             | 0.77 ± 0.26*                    | 2.76 ± 0.45       | 0.45 ± 0.23            | 1.11 ± 0.32                 | 0.55 ± 0.42*                |
| Cigarettes per day         |                                 |                   |                        |                            |                           |
| 1-5                       | 0.45 ± 0.40                     | 1.50 ± 0.57       | 0.09 ± 0.03*           | 2.08 ± 0.73*                | 0.45 ± 0.35*                |
| 6-10                      | 0.62 ± 0.28*                    | 2.00 ± 0.82*      | 0.34 ± 0.16*           | 1.69 ± 0.46*                | 0.60 ± 0.37*                |
| > 10                      | 1.00 ± 0.10*                    | 3.00 ± 0.00       | 0.50 ± 0.17*           | 1.77 ± 0.49*                | 0.72 ± 0.29*                |
| Statistical parameters    | r = 0.746                       | r = 0.746         | r = 0.699              | r = -0.497                  | r = 0.573                   |
| Duration of smoking (years)|                                 |                   |                        |                            |                           |
| 1-5                       | 0.56 ± 0.30*                    | 1.93 ± 0.83*      | 0.21 ± 0.03            | 1.77 ± 0.54*                | 0.44 ± 0.12*                |
| 6-10                      | 0.72 ± 0.20*                    | 2.4 ± 0.56*       | 0.23 ± 0.14            | 1.85 ± 0.09*                | 0.52 ± 0.37*                |
| > 10                      | 0.92 ± 0.09*                    | 2.85 ± 0.60*      | 0.40 ± 0.21*           | 1.49 ± 0.23*                | 0.53 ± 0.25*                |
| Statistical parameters    | r = 0.674                       | r = 0.689         | r = 0.776              | r = -0.601                  | r = 0.678                   |
| Unit years                |                                 |                   |                        |                            |                           |
| 1-199                     | 0.63 ± 0.29*                    | 1.86 ± 0.90*      | 0.40 ± 0.24*           | 1.93 ± 0.50*                | 0.34 ± 0.30*                |
| 200-399                   | 0.50 ± 0.29*                    | 2.13 ± 0.87*      | 0.41 ± 0.14*           | 1.48 ± 0.54*                | 0.47 ± 0.18*                |
| > 399                     | 0.81 ± 0.27*                    | 2.64 ± 0.64*      | 0.19 ± 0.21*           | 1.59 ± 0.35*                | 0.61 ± 0.40*                |
| Statistical parameters    | r = 0.718                       | r = 0.682         | r = 0.494              | r = -0.417                  | r = 0.409                   |

* Significantly higher than that of non-smokers

### Table 2. The mean scores of pigmentation and periodontal parameters based on smoking status, duration, frequency, and lifetime exposure

| Smoking status | Lip pigmentation Mean ± SD | Gingival pigmentation Mean ± SD | Bleeding Mean ± SD | Probing depth Mean ± SD | Loss of attachment Mean ± SD |
|----------------|---------------------------|---------------------------------|-------------------|------------------------|------------------------------|
| Non-smoker     | 0.10 ± 0.17               | 0.62 ± 0.42                     | 0.07 ± 0.08*      | 2.06 ± 0.16*           | 0.02 ± 0.04                  |
| Ex-smoker      | 0.44 ± 0.20*              | 1.66 ± 0.77*                    | 0.30 ± 0.14*      | 1.84 ± 0.47*           | 0.47 ± 0.51*                 |
| Occasional smoker | 0.48 ± 0.30*              | 2.05 ± 0.91*                    | 0.33 ± 0.18*      | 1.67 ± 0.56*           | 0.50 ± 0.30*                 |
| Current smoker | 0.77 ± 0.26*              | 2.34 ± 0.76*                    | 0.45 ± 0.23       | 1.11 ± 0.32            | 0.55 ± 0.42*                 |
| Cigarettes per day |                   |                                 |                   |                        |                              |
| 1-5            | 0.45 ± 0.40               | 1.50 ± 0.57                     | 0.09 ± 0.03*      | 2.08 ± 0.73*           | 0.45 ± 0.35*                 |
| 6-10           | 0.62 ± 0.28*              | 2.00 ± 0.82*                    | 0.34 ± 0.16*      | 1.69 ± 0.46*           | 0.60 ± 0.37*                 |
| > 10           | 1.00 ± 0.10*              | 3.00 ± 0.00                     | 0.50 ± 0.17*      | 1.77 ± 0.49*           | 0.72 ± 0.29*                 |
| Statistical parameters | r = 0.746               | r = 0.746                      | r = 0.699         | r = -0.497             | r = 0.573                    |

### Table 3. Contingency table with gingival and lip pigmentation scores for regular smokers and non-smokers

| Gingiva | Regular smokers | Non-smokers | Total |
|---------|-----------------|-------------|-------|
| Lip     |                 |             |       |
| 0       | -                | -           | 23    | 52   | 75    |
| 1-2     | -                | 1           | 11    | 12   | 15    |
| 3-6     | -                | 6           | 45    | 51   | 6     |
| Statistical parameters | $\chi^2 = 42.148$ | $\chi^2 = 46.753$ |          |        |       |
| Total   | -                | 7           | 56    | 63   | 70    | 96    |

Contingency coefficient: 0.043
30.6% of non-smokers without lip pigmentation lacked gingival pigmentation, 85.7% of those with lip pigmentation had gingival pigmentation as well (Table 3).

It is apparent from Table 4 that all periodontal parameters differed significantly based on the presence of gingival and lip pigmentation. Gingival bleeding was found to be the highest in subjects without lip pigmentation and those with grade-one gingival pigmentation. However, it was the lowest in subjects with lip pigmentation and those having grade three pigmented gingiva. On the other hand, measurements of periodontal probing depth and loss of gingival attachment had the highest values in subjects with pigmented lips and those having grade-three gingival pigmentation. Their lowest values were detected in subjects without lip pigmentation and those having grade-one gingival pigmentation.

**Discussion**

The present study, aiming at exploring the effects of smoking on lip and gingival pigmentation and periodontal status, was of high importance since despite the great number of studies on smoking and periodontal disorders, no such study has been conducted in Indian subcontinent.

This study compared current, occasional, and ex-smokers with non-smokers. Although previous studies had excluded occasional and ex-smokers, they were considered in this research to obtain the most precise results by analyzing the effects of smoking cessation on pigmentation and periodontal status. Lower scores of pigmentation and periodontal indicators were found in occasional and ex-smokers which is in good accordance with disappearance of lip and gingival pigmentation and reduced severity of periodontal diseases on cessation of smoking evident from past studies.

In this study, smokers and non-smokers significantly differed in presence of pigmentation, i.e. all smokers had lip and gingival pigmentation. This could be attributed to nicotine and benzpyrene content of tobacco smoke which stimulates melanin production from the melanocytes. Even children of smoking parents have been reported to exhibit greater gingival pigmentation compared to children of non-smoking parents.
While lip and gingival pigmentations were present in all of our smoking participants, the mentioned problems were seen in 33% and 27% of a Japanese population, respectively. On the other hand, 54.2% of Turkish adult smokers showed gingival pigmentation. Previous studies have reported that 15% of Europeans vs. 80% of Asians have oral pigmentation. Such a difference might have been caused by ethnic and skin color differences.

Lip and gingival pigmentations were observed in 31% and 73% of non-smokers, respectively. These high values might have been due to ethnic pigmentation. Oral pigmentation has been previously reported among 96% of the Indian population but only in 15% of Europeans and 37% of Turkish subjects.

Smokers exhibited a clear periodontal deterioration compared to non-smokers who mostly presented healthier periodontium. A detrimental effect of smoking on periodontium was alteration in gingival bleeding, which was found to be more among non-smokers than in smokers. This change in normal physiology underlies in the fact that the innate immune response of smokers is hampered by an increment in the number of neutrophils and a decreased functionality in peripheral circulation. In fact, neutrophils show decreased chemotaxis, phagocytosis, and adherence and the action of T-lymphocytes is also affected. These decrease the individual's hemorrhagic response and gingival blood flow.

Other effects of smoking included increased periodontal probing depth and loss of attachment. It has been found that the strong reducing agents (like carbon monoxide) present in tobacco smoke reduce the redox potential of periodontal mucosa. This alteration enhances the growth of anaerobic microorganisms such as Porphyromonas gingivalis and Aggregatibacter actinomycetemcomitans and leads to subgingival infection, pocket formation, and finally loss of clinical attachment.

Several studies have shown a relationship between the amount of smoking and the prevalence and severity of periodontitis. A relationship has been established between the prevalence of moderate to severe periodontal disease and number of cigarettes smoked per day, numbers of years that a patient has smoked, and unit years (total number of cigarettes smoked by an individual in one year). Even exposure to smoking has been related to the severity of pigmentation. A similar relationship was observed in the present study where pigmentation and periodontal deterioration had directly proportional relationships with smoking duration, frequency, and unit years. On the other hand, when smoking status was considered, the highest and lowest degrees of pigmentation and periodontal deterioration were recorded in current smokers and non-smokers, respectively.

Also, in this study, the highest values of probing depth and loss of clinical attachment were found in subjects with lip pigmentation and grade-three gingival pigmentation. However, those with no pigmentation had the lowest scores. On the other hand, gingival bleeding was inversely related to pigmentation. Similar observations were reported by an earlier study in Turkey. The observed association between pigmentation and periodontal deterioration could be attributed to the indirect relationship of these two entities which have a common etiological agent, smoking.

The present study had several limitations. The accuracy of reporting is not known. No biomarkers such as cotinine levels or exhaled carbon monoxide were tested to validate tobacco exposure either through self-use or environmental exposure. This study was conducted on subjects who were present on the day of the survey and hence other eligible but absent individuals were not included. If the absents had been smokers, the obtained prevalence might have underestimated the actual level of smoking. On the other hand, if the absents had been non-smokers, we might have overestimated the actual smoking levels.

Further research has to be designed and implemented to evaluate the efficiency of comprehensive smoking cessation programs and their effects on periodontal health. Public awareness about the dangers of smoking should be promoted through public education campaigns. Policy efforts are also required to address the problems in this field. Furthermore, smoking cessation programs and anti-tobacco advertisements need to be implemented. Increased professional help for smoking cessation should be made available to persons who want to quit as all may positively impact the periodontal health status of this population.
Conclusion

Smoking was observed to influence lip and gingival pigmentation and periodontium. Periodontal status and melanin pigmentation were influenced by duration, frequency, and unit years of cigarette smoking. All individuals with lip pigmentation presented some form of gingival pigmentation. Probing depth and loss of attachment were more severe in subjects with lip and gingival pigmentation while the percentage of bleeding sites had an inverse relationship with pigmentation.

Conflict of Interest

The Authors have no conflict of interest.

Acknowledgements

This research was self-funded. I sincerely thank the participants. The author reports no conflicts of interest.

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ارتباط سیگار کشیدن با پیگماتاسیون ملایم لب، لثه و وضعیت پریودنتال

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مقاله پژوهشی

چکیده

مقدمه: سیگار کشیدن یکی از عوامل حاوی مهم بیماری پریودنتال می‌باشد که منجر به پیگماتاسیون موقع دهان می‌شود. این مطالعه با هدف ارزیابی ارتباط و مقایسه تأثیرات سیگار کشیدن روی لثه و پیگماتاسیون لثه و وضعیت پریودنتال و ارزیابی ارتباط پیگماتاسیون با پارامترهای پریودنتال انجام شد.

روش‌ها: ۱۰۹ فرد سیگار و ۱۰۹ فرد غیر سیگاری با سن ۲۵-۴۴ سال در این مطالعه مصوبه شدند. متوسط سن افراد ۲۹/۱۵ سال بود. همه شرکت کنندگان جهت پیگماتاسیون لثه و لثه و وضعیت پریودنتال با اندازه‌گیری خونریزی لثه، عمق پریوبینگ و از دست رفتن چسبندگی در ۶ نقطه از هر دندان مورد بررسی قرار گرفتند.

بافته‌ها: همه افراد سیگاری شرکت کننده در این مطالعه، پیگماتاسیون لثه و لثه داشتند. در حالی که، در افراد غیر سیگاری ۷ برای سیگاری ۴ و ۷ برای سیگاری بود. پارامترهای پیگماتاسیون و پریودنتال به جز خونریزی لثه به طور مستقلی با دیگر عوامل سیگاری و وضعیت پریودنتال نداشتند. عمق پریوبینگ و از دست رفتن چسبندگی در افراد با پیگماتاسیون لثه و ورد ۳ لتن پیگماتی بهتر بود.

نتیجه‌گیری: سیگار کشیدن در پیگماتاسیون لثه و لثه و به علاوه بلافاصله پریودنتال تأثیر دارد. این اثرب را دارند، معنی‌دار بود و از دست دادن چسبندگی در افراد با پیگماتاسیون لثه و لثه بهتر دیده شد.

واژگان کلیدی: موقع دهان، پیگماتاسیون، سیگار کشیدن، پریودنتال

ارجاع: مولتانا، سوراج. ارتباط سیگار کشیدن با پیگماتاسیون ملایم لب، لثه و وضعیت پریودنتال. مجله اعتیاد و سلامت ۱۳۹۲؛ ۵(۲): ۶۷-۶۵.

تاریخ پذیرش: ۹۱/۸/۲۱

تاریخ دریافت: ۹۱/۵/۱۰

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http://ahj.kmu.ac.ir, ۴ April

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Addict Health, Winter & Spring 2013; Vol 5, No 1-2