Association of oral manifestations in ulcerative colitis: A pilot study

Mohan Kumar KP¹, Nachiammai N², Madhushankari GS¹

¹Department of Oral Pathology and Microbiology, College of Dental Sciences, Davangere, Karnataka, ²Department of Oral and Maxillofacial Pathology, Chettinad Dental College and Research Institute, Tamil Nadu, India

INTRODUCTION

Inflammatory bowel disease (IBD) is a group of inflammatory conditions of the colon and small intestine, which principally includes Crohn’s disease and ulcerative colitis (UC). In genetically susceptible individuals, an inappropriate and continued inflammatory response by the complex interaction of environmental, genetic and immunoregulatory factors to gut microbiota is thought to result in UC. The current and more accepted theory is that intestinal inflammation is a consequence of an...
abnormal action of T-lymphocytes on enteric bacterial flora in genetically susceptible individual. This chronic inflammation of the colonic mucosa causes ulceration, edema, bleeding, diarrhea and fluid and electrolyte loss.\textsuperscript{[1]} UC is a relapsing and remitting disease. Incidence and prevalence rate of UC in India are reported to be $6.02/10^5$ per year and $44.3/10^5$ population, respectively.\textsuperscript{[2]}

UC exhibits intestinal and extraintestinal complications. Local complications include ischiorectal abscess, rectal prolapse, hemorrhoids, fibrous stricture, pseudopolypsis, perforation, massive hemorrhage and also carcinoma of colon. Extraintestinal complications are common in UC and may dominate the clinical picture. Some of these occur during relapse of intestinal disease others appear unrelated to disease severity. These include erythema nodosum, pyoderma gangrenosum, skin eruptions, uveitis, arthritis, transient hepatitis and venous thrombosis.\textsuperscript{[1]}

As oral cavity is the portal of entry and part gastrointestinal (GI) tract, oral manifestations do occur. Oral manifestations of UC occur as a systemic complication or extraintestinal manifestation that may precede, exacerbate or regress with alterations in severity of the disease. Pyostomatitis vegetans (PV) is considered as a specific oral indicator of UC, and other manifestations such as aphthous ulcers, lichenoid lesion, halitosis, dysgeusia, dry mouth, coated tongue, gingivitis and periodontitis are considered as nonspecific oral manifestation.\textsuperscript{[3‑6]}

The purpose of this study was to list and evaluate the oral manifestation in UC patients and also to evaluate whether oral manifestation can be a predictor for relapse and remission of UC.

**MATERIALS AND METHODS**

The study was conducted in clinics of gastroenterology, Davangere, Karnataka. Fifteen (8 males and 7 females) individuals diagnosed with UC [Figures 1 and 2 indicating active colitis, Figure 3 showing healing colitis] and their age- and sex-matched controls were included in the study by obtaining informed consent.

From the review of various studies and personal experience of oral manifestations of UC, a detailed case history format was prepared that included patient evaluation for PV, aphthous ulcer, lichenoid lesion, tongue coating, halitosis, dysgeusia, dry mouth, gingivitis and periodontitis. PV, aphthous ulcer (composite index),\textsuperscript{[7]} lichenoid lesion, tongue coating (Miyazaki et al’s index)\textsuperscript{[8]} were assessed by clinical examination. Halitosis and dysgeusia were evaluated by series of questions, dry mouth by estimation of unstimulated salivary flow rate. Periodontal status was assessed by gingival and periodontal index. All the above parameters were assessed in diagnosed UC.
patients. A questionnaire was prepared to evaluate these manifestations during the past remissions, which was explained to patients and was filled. All the parameters were compared with controls.

**Statistical analysis**

Mann–Whitney test was used for comparison of intergroup data. To evaluate the oral manifestations during previous episode and present status, Wilcoxon signed-rank test was applied. \( P < 0.05 \) was considered statistically significant.

**RESULTS**

The findings of the present study in the case group are depicted in Table 1. Of the 15 patients, 10 patients reported aphthous ulceration, one patient revealed PV, three patients exhibited lichenoid lesion, and 11–12 patients had halitosis and dry mouth and 4–5 patients showed coated tongue and dysgeusia. Salivary flow rate, tongue coating and periodontal status between study and control group are presented in Table 2. Salivary flow rate was significantly reduced, coated tongue was present and periodontal status was comparatively same in UC patients when compared to controls. Aphthous ulcer status during previous relapse and present status is statistically highly significant [Table 3]. Among all the manifestations, studied aphthous ulcer showed statistically significant values when compared between previous relapse of the disease and at present examination.

**DISCUSSION**

UC, an inflammatory bowel, is postulated that they result from unregulated and exaggerated immune response to commensal microbes in the gut in genetically susceptible individuals. The genetic susceptibility plays an important role, and the implicated genes regulate several important biologic functions including immunoregulation, mucosal barrier integrity and microbial clearance and/or homeostasis. The gene associated with UC MDR1 located on chromosome 7 physiologically protects cells from toxic substances or metabolites. Some genotypes of UC, HLADRB1*0103, B*27 and B*58 are linked with extraintestinal manifestations.\(^9\)

Oral lesions occur as an extraintestinal manifestation that may precede, exacerbate or regress with alterations disease. Thus, the present study was carried out to evaluate the oral manifestations and also to assess whether these manifestations can predict remission and relapse of UC.

Results of our study indicate that all the patients diagnosed with UC exhibited oral manifestations. Oral ulceration, halitosis, altered taste and dry mouth were frequently prevalent oral manifestations in patients with UC.

One of the male patients presented with PV on the lower anterior gingiva, a specific oral manifestation of UC, and reported exacerbation during relapse.\(^10\) PV is common in young middle-aged adults with male predominance (3:1), clinically characterized by initial formation of 2–5 mm in diameter, abscess on erythematous mucosa, erode and coalesce, and undergoes necrosis to form “snail track” appearance. Sometimes, they correspond to exophytic lesions, producing a vegetating appearance. The surface is yellow creamy color and is covered by a pseudomembrane that easily disintegrates, facilitating formation of small ulcers and superficial erosion.\(^11\) The most common site of occurrence are the labial and buccal mucosa, hard and soft palate, gums and less frequently floor of the mouth. Histopathologically PV shows hyperkeratosis, acanthosis, intraepithelial and subepithelial microabscess with prominent neutrophil and eosinophilic infiltrate.\(^12\)

Several authors hypothesized that PV results from an aberrant immune response to yet unidentified factors. As IBD is the most common underlying disorder, it is postulated that the cross-reacting antigens in the bowel and mucosa to be responsible for secondary mucocutaneous manifestation.\(^13\) Thus, numerous case reports and few studies have suggested PV as a specific indicator of UC.

In the present study, 10 patients had aphthous ulcers and reported exacerbation of the aphthous ulcers during the previous relapse compared too, suggesting that aphthous ulcer evaluation can be used as an indicator of the relapse and remission. Aphthous ulceration in UC is due

| Manifestation                  | Study group (n=15) |
|-------------------------------|-------------------|
| Pyostomatitis vegetans        | 1                 |
| Aphthous ulceration           | 10                |
| Lichenoid lesion              | 3                 |
| Halitosis                     | 12                |
| Dysgeusia                     | 5                 |
| Dry mouth                     | 11                |
| Tongue coating                | 4                 |

| Manifestation                  | Study group | Control group | \( P \) |
|-------------------------------|-------------|---------------|--------|
| Salivary flow rate (ml/min)   | 1.5         | 1.98          | 0.02(S) |
| Tongue coating                | 4.5         | 3             | 0.12(NS) |
| Periodontal status            |             |               |        |
| Gingival index (score)        | 1.2         | 1.3           | 0.23(NS) |
| Periodontal index (score)     | 1.0         | 1.4           |        |
| Loss of attachment (mm)       | 0.4         | 0.2           |        |

NS: Not Significant
to micronutrient malabsorption of iron-inducing iron deficiency anemia and Vitamin B₁₂ in pernicious anemia. Our study results stimulate us to support findings of Correll et al., who suggested that recurrent oral aphthous ulcers should be included as one of the factors in determining the occurrence, relapse and remission of UC.[¹⁴]

Among the evaluated UC patients, three patients exhibited oral lichen planus (OLP). The co-occurrence of OLP and UC can be attributed to the side effects of drugs used such as sulfasalazine and diltiazem. It seems that aminosalicylic acid moiety of sulfasalazine drug is responsible for the reaction and that LP is a true complication of mesalazine therapy.[¹³] Some authors indicated that OLP is a drug complication and discontinuation of sulfasalazine resolved the condition, whereas others contraindicated and reported that discontinuation did not resolve but instead became more extensive.[¹³] Still further on, studies should be carried to prove whether OLP is a manifestation due to UC or reactive lesions to unavoidable medications used for the treatment of UC.

Twelve of our UC patients reported halitosis; this finding of ours was in accordance with Elahli et al. This finding can be due to an increased amount of colonic sulfate-reducing bacteria in UC patients, resulting in higher concentrations of the toxic gas hydrogen sulfide. Human colonic mucosa is maintained by the colonic–epithelial barrier and immune cells in the lamina propria. N-butyrate, a short-chain fatty acid, gets oxidized through the beta-oxidation pathway into carbon dioxide and ketone bodies. It has been shown that N-butyrate helps supply nutrients to this epithelial barrier. Studies have proposed that hydrogen sulfide plays a role in impairing this beta oxidation pathway by interrupting the short chain acetyl Co-A dehydrogenase, an enzyme within the pathway. An unrelated study suggested that the sulfur contained in red meats and alcohol may lead to an increased risk of relapse for patients in remission.[¹⁶,¹⁷] Hence, red meat and alcohol should be avoided in UC patients.

Altered taste was reported by five UC patients, which was similar to the results of Elahli et al. It is considered to be due to long-term use of medication in the treatment of UC as sulfasalazine metabolizes to sulfapyridine. Serum sulfapyridine levels >50 µg/mL are associated with altered taste.[¹⁸]

In our study, salivary flow rate was decreased when compared to controls which was in accordance with other studies[¹⁹] and is thought to be due to long-term use of steroids and sulfasalazine therapy that decreases the secretory capacity of acinar cells. Four patients exhibited yellow-to-white discoloration with heavy thick coating and is suggested to be due to steroid therapy or superimposed candidal infection or stress-induced negligence of oral hygiene.

Periodontal status was comparable to that of controls and one of the cases exhibited severe gingivitis and periodontitis. Brito et al. suggested that IBD patients harbor higher levels of Treponema denticola and other bacteria that are related to opportunistic infections in inflamed subgingival sites that might be harmful for the crucial microbe–host interaction and responsible for severity of gingivitis and periodontitis.[²⁰]

Finally, UC patients are suggested to be prone to colonic cancer with increase in duration of colitis and relapse. Meta-analysis by Eden et al. suggested that with increase in duration of colitis, viz., 10, 20 and 30 years, the risk of colon cancer increases by 2%, 8% and 18%, respectively.[²¹] Thus, simple early detection of oral manifestations in UC patients can aid in the prognosis, monitoring of relapse and remission and possibly in the prevention of colorectal cancer.

**CONCLUSION**

From the present study, oral symptoms and lesions such as aphthous ulceration, PV and also other manifestation should alert a clinician to elicit history and suspect the possibility of UC even in the absence of full-blown GI manifestations.[²¹] Few patients of our study considered oral manifestation as the marker of disease severity and took adequate care.

Information regarding the oral manifestation in UC is less and is based mainly on case reports. The results of this study have been obtained based on a small number of patients; hence, studies on larger samples would provide more insight into the evaluation of oral manifestations in UC patients and their correlation with relapse and remission.
Acknowledgment
We would like to acknowledge Dr. Prakash M. G, Sujan Gastroenterology and Maternity Centre, Davangere - 577004.

Financial support and sponsorship Nil.

Conflicts of interest
There are no conflicts of interest.

REFERENCES

1. Friedman S, Blumberg RS. Inflammatory bowel disease. In: Kasper DL, Braunwald E, Fauci AS, Hauser SL, Jameson JL, editors. Harrison's Principles of Internal Medicine. 16th ed. USA: McGraw-Hill Companies, Inc.; 2005. p. 1776.
2. Gunisetty S, Tiwari SK, Bardia A Phanibhushan M, Satti V, Habeeb MA, et al. The epidemiology and prevalence of ulcerative colitis in the South of India. Open J Immunol 2012;2:144-8.
3. Paradowska A. Oral cavity at ulcerative colitis – Preliminary study. Dent Med Probl 2008;45:382-5.
4. Daley TD, Armstrong JE. Oral manifestations of gastrointestinal diseases. Can J Gastroenterol 2007;21:241-4.
5. Edwards FC, Truelove SC. The course and prognosis of ulcerative colitis. III. Complications. Gut 1964;5:1-22.
6. Ganapelli A, Ayinampudi BK, Pacha VB, Poduluturi SR, Gangula S, SaiKreddy S. Oral indicators of ulcerative colitis: A rare case report and review of literature. J Indian Aca Oral Med Radiol 2012;24:56-62.
7. Mumeu G, Sur H, Inane N, Karacayli U, Cimilli H, Simsan N, et al. A composite index for determining the impact of oral ulcer activity in Behcet's disease and recurrent aphthous stomatitis. J Oral Pathol Med 2009;38:785-91.
8. Van Tornout M, Dadamio J, Coucke W, Quirynen M. Tongue coating: Related factors. J Clin Periodontol 2013;40:180-5.
9. Rothfuss KS, Stange EF, Herrlinger KR. Extraintestinal manifestations and complications in inflammatory bowel diseases. World J Gastroenterol 2006;12:4819-31.
10. Lobkovicz F, Eckert F, Braun-Falco O. Pyostomatitis vegetans: A specific marker of Crohn disease and ulcerative colitis. Hautarzt 1991;42:92-5.
11. Knapp N, Albuquerque R, Khan Z, Richards A, Brown RM. Pyostomatitis vegetans in ulcerative colitis: management with topical tacrolimus and systemic azathioprine in a 10-year-old boy (case report and review of the literature). J Int Oral Health 2016;8:132-6.
12. Ruiz-Roca JA, Berini-Aytes L, Gay-Escoda C. Pyostomatitis vegetans: Report of two cases and review of the literature. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 2005;99:447-54.
13. Femiano F, Lanza A, Buonaiuto C, Perillo I, Dell’Ermo A, Cirillo N, et al. Pyostomatitis vegetans: A review of the literature. Med Oral Patol Oral Cir Bucal 2009;14:E114-7.
14. Correll RW, Wescott WB, Jensen JL. Recurring, painful oral ulcers. J Am Dent Assoc 1981;103:497-8.
15. Alstead EM, Wilson AG, Farthing MJ. Lichen planus and mesalazine. J Clin Gastroenterol 1991;13:335-7.
16. Davies PT, Shadforth MF. Sulphasalazine induced oral lichen planus. Br Med J 1984;288:194.
17. Tannock GW. The bowel microbiota and inflammatory bowel diseases. Int J Inflam 2010;2010:954051.
18. Tilg H, Kaser A. Diet and relapsing ulcerative colitis: Take off the meat? Gut 2004;53:1399-401.
19. Brito F, Zaltman C, Carvalho AT, Fischer RG, Persson R, Gustafsson A, et al. Subgingival microflora in inflammatory bowel disease patients with untreated periodontitis. Eur J Gastroenterol Hepatol 2013;25:239-45.
20. Eaden JA, Abrams KR, Mayberry JF. The risk of colorectal cancer in ulcerative colitis: A meta-analysis. Gut 2001;48:526-35.
21. Elahi M, Telkabadi M, Samadi V, Vakili H. Association of oral manifestations with ulcerative colitis. Gastroenterol Hepatol Bed Bench 2012;5:155-60.