



RESEARCH ARTICLE

CLINICAL FEATURES, AGE AND SEX DISTRIBUTIONS, RISK FACTORS AND THE TYPE OF BACTERIA ISOLATED IN PERIODONTITIS PATIENTS IN SANA'A, YEMEN

Khaled A AL-Haddad¹ , Mohammed Mohammed Ali Al-Najhi² , Al-Kasem Mohammed Abbas³ , Ameen Abdullah Yahya Al-Akwa¹ , Hassan Abdulwahab Al-Shamahy⁴ ,
 Mohammed A Al-labani¹ 

¹Orthodontics, Pedodontics and Prevention Department Faculty of Dentistry, Sana'a University, Yemen

²Orthodontics, Pedodontics and Prevention Department Faculty of Dentistry, Genius University for Sciences and Technology, Dhamar city, Republic of Yemen.

³Department of Maxillo-Facial, Faculty of Dentistry, Sana'a University, Republic of Yemen.

⁴Department of Basic Sciences, Faculty of Dentistry, Sana'a University, Republic of Yemen.

Article Info:



Article History:

Received: 6 December 2020

Reviewed: 8 January 2021

Accepted: 11 February 2021

Published: 15 March 2021

Cite this article:

AL-Haddad KA, Al-Najhi MMA, Abbas AKM, Al-Akwa AAY, Al-Shamahy HA, Al-labani MA. Clinical features, age and sex distributions, risk factors and the type of bacteria isolated in periodontitis patients in Sana'a, Yemen. Universal Journal of Pharmaceutical Research 2021; 6(1):1-8.

<https://doi.org/10.22270/ujpr.v6i1.532>

*Address for Correspondence:

Dr. Hassan A. Al-Shamahy, Department of Basic Sciences, Faculty of Dentistry, Sana'a University, Republic of Yemen, Tel- +967-770299847; E-mail: shmahe@yemen.net.ye

Abstract

Background: Periodontitis is an inflammation caused by plaque in the surrounding dental structures. It is a major factor in adult tooth loss. There is lack of information on associated clinical features, risk factors and microbial etiology of periodontitis in Sana'a, Yemen.

Aim: The study focused on associated clinical features, risk factors and the separation and classification of bacteria in periodontitis and associated risk factors amongst patients attending dental clinics in Sana'a city.

Methods: First, 296 patients were admitted to the dental clinic at the Republican University Hospital and private dental clinics in Sana'a during a period of nearly one year, which began in December 2019 AD and ended in November 2020 AD, when they were diagnosed with dental diseases. Standard culture and biochemical techniques were used for the isolation and identification. Structured questionnaires were used to record clinical features, demographic variables and other risk factors of periodontitis.

Results: A total of 130 microorganisms were isolated from 49 patients with periodontitis. Male patients accounted for 14.4% and females 18.9% of the all 296 patients who attended the clinics. There was no important association between sex and periodontitis occurrence while there was significant association the younger age groups (45.4% in <26 years of age). The most common signs and symptoms were swollen or puffy gums (91.9%), bleed easily gums (96%), halitosis (96%), painful chewing (87.8%), pus between teeth and gums (71.4%), loose teeth or loss of teeth (44.9%), gingival recession (83.7%), spitting out blood when brushing or flossing teeth (79.6%), and tender gums (93.9%).

Conclusion: This study is new in Sana'a city. The clinical features of preidentitis in Yemen and the risk factors are similar to those reported in the literature elsewhere, but the isolated bacteria differ in frequency from those reported elsewhere, as some upper respiratory tract pathogens such as *Streptococcus pyogenes* are commonly isolated in this study. Knowledge of the clinical features, bacterial causes of gum disease, and risk factors is the key to successful periodontal therapy.

Keywords: bacteria agents, clinical features, periodontitis, risk factors, Sana'a, Yemen.

INTRODUCTION

There is a significant spread of periodontal disease worldwide, and identify the etiology is the solution to controlling it. Periodontal disease is characterized by a

chronic bacterial infection with persistent inflammation, breakdown of connective tissue and destruction of the alveolar bone¹. Generally periodontal disease is classified into periodontitis and gingivitis. Gingivitis is inflammation of the gums initiated by the proliferation

of dental plaque and is reversible. Gingivitis is possibly common in children up to 5 years old². It is the result of inappropriate oral hygiene practices³. Periodontitis is a chronic inflammatory disease that begins with a buildup of dental biofilm plaque and continues through a disorganized immune response and is commonly started by gingivitis leading to irreversible demolition of the supportive tissues adjoining the tooth, together with the alveolar bone^{1,2,4}. Periodontal disease is a multi-microbial, multifactorial disease, with numerous host factors involved in influential an individual's susceptibility to disease⁵. Several reports have been description that the onset and sequence of the disease is not only similar to the existence of pathogenic bacterial strains in the gums but also due to the absence or minimum levels of beneficial equivalents commensals in the susceptible host⁵⁻⁷. A limited number of periodontal pathogens have been reported in the complex biofilms to initiate periodontal disease. Data clarified that some bacterial strains in the gingival environment can cause gingivitis and bone destruction. These bacterial strains are known as periodontal pathogens^{8,9}.

It is known that these periodontal disease pathogens possess, when present in even very small quantities, the capability to damage the gingival structure⁵. It is known that most periodontal pathogens are anaerobes, but biofilm can also accommodate facultative aerobes, capnophiles, and microaerophiles whose number depends on the environment in the developing biofilms and gingival pocket⁵. It became known that of 800–1000 species that colonized the oral cavity, 50 species have been identified with strong links to periodontal disease¹⁰. These complexes were classified Socratic and Hafidian into 5 complexes, which are the yellow or early colonized complex, the green or secondary colonized complex, then the orange, purple and red complexes. The red compound is the secondary colony that is the major pathogens related with bleeding upon investigation¹¹. There is strong bacterial progression in oral cavity infection that may be predisposed by age, diet, or site of infection¹². Nevertheless, in addition to periodontal disease pathogens, genetic and environmental factors predispose to disease progression. The risk of gum disease is determined by several factors including any health condition that leads to bacterial defense mechanisms defect such as diabetes, human immunodeficiency virus (HIV) and neutropenia. Tobacco smoking, Obesity, poor diet, and a inactive lifestyle are associated with an increased risk of periodontitis¹³. In the early stages, periodontitis has few symptoms, and the disease has progressed significantly in many individuals before they seek treatment. Symptoms may include redness or bleeding of the gums while brushing the teeth, the use of dental floss or gnawing on solid food, frequent swelling of the gums, gingival recession gingival recession, halitosis, deep pockets between the teeth and the gums, loose teeth, and drifting of incisors^{13,14}.

Periodontitis is the leading cause of tooth loss in adults universally and these people are at risk of edentulism, multiple tooth loss and masticatory impairment as a result of which a negative impact on nutrition, quality

of life and self-esteem and thus a significant societal imposition - the economic impact and the cost of health care^{3,14,15}. Most of the information about the causes of periodontitis emerged from studies conducted in Europe and the United States of America, and some third world countries. Although there are some studies on oral and dental problems in Yemen¹⁶⁻²⁵, no study has been conducted in Yemen on periodontitis²⁶. So this study focused on associated clinical features, risk factors and the isolation and identification of bacteria in periodontitis and associated risk factors among patients attending some dental clinics in Sana'a city.

SUBJECTS AND METHODS

Patients

This study included 49 patients suffering from periodontitis, who were admitted to the dental clinic at the Republican University Hospital and private dental clinics (Al-Mortadda dental clinics, Al-Abany dental clinics and Al-Kahara dental clinics) in Sana'a, during a period of about one year, which started in December 2019 and ended in November 2020, of whom 22 were males and 27 were females.

Data collection and microbial processing

A questionnaire was filled out for each patient with the patient's personal, clinical data and risk factors. This included age, gender, occupation and relevant clinical information regarding bacterial oral infections. Also risk factors of contracting periodontitis. Cultures were obtained from the collected pocket by probes in order to isolate the various bacterial causative agents. First, the supragingival plaque was removed (without disturbing the subgingival plaque) and a bacterial sample was collected from the deepest periodontal pockets with a sterile probe. The samples were then placed in a vial containing 2 ml of liquid thioglycolate enriched medium, sealed immediately and transported to the laboratory within 30 minutes. Bacteriological procedures were performed within one hour of sample collection. For germ cultures, the following media and conditions were used: Tryptic Soy Agar (TSA) with blood (5%) and MacConkey agar plates - incubated at 35°C under 5% CO₂ and examined at 24 and 48 hours; Brucella agar enriched with Vitamin K1 and CDC + amikacin blood agar - incubated at 35°C anaerobically in a Gaspak jar (Oxoid Ltd). Cultures were examined for the presence of bacteria at 48 and 96 h. Plates showing bacterial growth were retained until final processing and organism identification by classical standard techniques including culture colonies morphology, microscopy staining methods, and biochemical tests^{13,27}.

Data analysis

Clinical, personal, and risk factors data as well as sample culture results entered into the questionnaire were analyzed by Epi Info, Version 6. All subjects with pockets less than 2 mm were considered to have periodontitis. To correlate the clinical features and potential risk factors for periodontitis, the data were examined in the form of case-control studies. For people with periodontitis, people with other dental diseases have been matched. Differences in categorical

variables were assessed using Fisher's exact tests as appropriate. Ninety-five percent confidence intervals for odds ratios were calculated according to the Cornfield limits and 95% confidence intervals were calculated for simple ratios by an exact binomial method. The significance of the difference in the ratio and the odds ratio was analyzed, and a chi-square (χ^2)

greater than 3.84 and a probability value (p) less than 0.05 were considered statistically significant.

Ethical approval

Ethical approval was obtained from the Medical Research and Ethics Committee of the Faculty of Medicine and Health Sciences at Sana'a University. All data, including patient identification were kept confidential.

Table 1: The age and sex distribution of patients suffering from periodontitis.

Sex	Periodontitis n=49		OR	95% CI	χ^2	p
	No.	%				
Male n=153	22	14.4	0.7	0.3-1.3	1.1	0.29
Female n=143	27	18.9	1.3	0.7-2.5	1.08	0.29
Age in years						
< 16 n=36	8	22.2	1.5	0.6-3.5	0.9	0.32
16-25 n=56	13	23.2	1.7	0.8-3.5	2.2	0.13
26-35 n=64	10	15.6	0.91	0.4-1.9	0.05	0.82
36-45 n=50	6	12	0.68	0.3-1.5	0.9	0.34
> 45 n=90	12	13.3	0.7	0.34-1.4	0.9	0.32
Total n=296	49	16.6				

RESULTS

Male patients accounted for 14.4% of the total periodontitis patients, and the female percentage was 18.9% from the total dental patients attending to the clinics. Table 1 show the age and gender distribution of patients with periodontitis. There was no significant association between gender and periodontitis occurrence. When age was considered a dependent factor for periodontitis, the rate of periodontitis was highest in the younger age groups (45.4% in <26 years of age), while the rate in > 45 years was 13.3%. Table 2 shows the importance periodontitis signs and symptoms among dental patients in Sana'a, Yemen. When compared periodontitis with other oral diseases, swollen or puffy gums occurred in 91.9% of periodontitis patients and occurred more than 30.2 times (CI=10.4-87, $p<0.001$) of other oral diseases. Bleed easily gums occurred in 96% of

periodontitis patients and occurred more than 67 times (CI=15-284, $p<0.001$) of other oral diseases. Halitosis occurred in 96% of periodontitis patients and occurred more than 88 times (CI=20-374, $p<0.001$) of other oral diseases. Painful chewing occurred in 87.8% of periodontitis patients and occurred more than 13.4 times (CI=5.4-32, $p<0.001$) of other oral diseases. Pus between teeth and gums occurred in 71.4% of periodontitis patients and occurred more than 15.3 times (CI=7.6-32, $p<0.001$) of other oral diseases. Loose teeth or loss of teeth occurred in 44.9% of periodontitis patients and occurred more than 5.2 times (CI=2.6-10.3, $p<0.001$) of other oral diseases. Other symptoms and signs occurred 83.7% for gingival recession, 79.6% for spitting out blood when brushing or flossing teeth, 93.9% for tender gums with significant occurrence for all these signs as compared with other oral diseases. Table 3 shows the risk factors associated with periodontitis among dental patients in Sana'a, Yemen.

Table 2: The significance of signs and symptoms of periodontitis among dental disease patients in Sana'a Yemen.

Signs and symptoms	Periodontitis n=49		OR	95% CI	χ^2	p
	No.	%				
Swollen or puffy gums n=112	45	91.9	30.2	10.4-87	72	<0.001
Bright red, dusky red or purplish gums n=145	44	89.8	12.7	4.8-32	39	<0.001
Tender gums n=195	46	93.9	10	3.1-331	20	<0.001
Bleed easily gums n=111	47	96	67	15-284	85	<0.001
Bleeding during toothbrush or after brushing n=143	41	83.7	7.2	3.2-16	29	<0.001
Spitting out blood when brushing or flossing teeth n=89	39	79.6	15.3	7.1-32	68	<0.001
Halitosis n=99	47	96	88	20-374	102	<0.001
Pus between teeth and gums n=69	35	71.4	15.6	7.6-32	76	<0.001
Loose teeth or loss of teeth n=55	22	44.9	5.2	2.6-10.3	26	<0.001
Painful chewing n=129	43	87.8	13.4	5.4-32	46	<0.001
Gingival recession n=188	41	83.7	3.4	1.5-7.7	10.2	<0.001
Total n=296	49	16.6				

OR=odds ratio, CI=confidence limit 95%, χ^2 =Chi-square ≥ 3.84 (significant), p =probability value < 0.05 (significant)

Table 3: The associated risk factors of periodontitis among dental disease patients in Sana'a Yemen.

Risk factors	Periodontitis n=49		OR	95% CI	χ^2	p
	No.	%				
Low Education level n=95	26	27.3	2.9	1.5-5.4	11.8	<0.001
Repeated History of Gingivitis n=95	29	59.2	3.2	1.7-6.1	14.4	<0.001
Poor oral health habits n=151	16	32.6	0.4	0.21-0.77	7.9	0.004
Smoking or chewing tobacco n=45	12	24.4	0.74	0.2-1.8	0.39	0.52
Chewing Qat n=91	31	63.2	5.3	2.8-10.2	29	<0.001
*Hormonal changes n=45	6	12.2	0.74	0.2-1.8	0.39	0.52
*Recreational drug use n=6	1	2	1.01	0.11-8.8	0.001	0.99
Obesity n=57	17	34.7	2.7	1.3-5.4	8.9	0.002
Inadequate nutrition, including vitamin C deficiency n=71	10	20.4	1.2	0.5-2.5	0.23	0.6
*Certain medications that cause dry mouth or gum changes n=12	8	16.3	11.8	3.4-41	22	<0.001
*Conditions that cause decreased immunity n=9	4	8.2	4.3	1.1-16.6	5.2	0.02
Certain diseases n=34	16	32.7	6.1	2.8-13.2	25.8	<0.001
Total n=296	49	16.6				

Hormonal changes=such as those related to pregnancy or menopause.

Recreational drug use =such as smoking marijuana or vaping.

Certain medications that cause dry mouth (xerostomia): Antihistamines, Antidepressants, Antipsychotics, Lung inhalers.

Conditions that cause decreased immunity=such as leukemia, HIV/AIDS and cancer treatment.

Certain diseases, such as diabetes, rheumatoid arthritis and Crohn's disease.

OR=odds ratio, CI=confidence limit 95%, χ^2 =Chi-square ≥ 3.84 (significant), p=probability value< 0.05 (significant)

There was a significant correlation between a frequent history of gingivitis (OR=3.2, CI=1.7-6.1, $p<0.001$), qat chewing (OR=5.3, CI=2.8-10.2, $p<0.001$), obesity (OR=2.7, CI=1.3-5.4, $p=0.004$), some drugs that cause dry mouth or gingival changes (OR=11.8, CI=3.4-41, $p<0.001$), conditions that cause decreased immunity (OR=4.3, CI=1.1- 16.6, $p=0.02$), and some diseases, such as diabetes, rheumatoid arthritis, and Crohn's disease (OR=6.1, CI=2.8-13.2, $p<0.001$); and the occurrence of periodontitis. Table 4 shows the number and percentage of the cultivated microorganisms from the 49 patients suffering from periodontitis. Multi-infections occurred in 89.8% of the periodontitis patients and the most common bacteria isolated were *Actinobacillus actinomycetemcomitans* (79.6%), followed by *S. pyogens* (73.5%) and *Staphylococcus aureus* (53.1%). While the *Bacteriodes* species (20.4%), *S. mutans* (16.3%) and *Anaerobic lactobacillus* (4%) were less isolated from the

periodontitis patients. *Candida albicans* was isolated in 4 cases (10.2%).

DISCUSSION

Periodontal disease is an increasing health problem in Yemen. Currently, no work has been done to determine the clinical features, etiological and risk factors for periodontitis in Sana'a City, but little works have been done, they dealt with the spread of oral and dental diseases; and some dental and oral disorders¹⁶⁻²⁵. In the current study, there was no significant association between sex and the incidence of periodontitis, and many other researchers appear to favor the female preference^{28,29}. Also, current work contrasts with that of Ababneh and others, who reported a predisposition to males but current result is similar to that reported by Susin and Albander, which reported an equal distribution^{30,31}.

Table 4: The number and percentage of the cultivated microorganisms from the 49 patients suffering from periodontitis.

Bacterial isolates	Total n=49		Male n=22		Female n=27	
	No.	%	No.	%	No.	%
<i>Streptococcus pyogens</i>	36	73.5	17	77.3	19	70.3
<i>Staphylococcus aureus</i>	26	53.1	14	63.6	12	44.4
<i>Bacteriodes</i> species	10	20.4	7	31.8	3	11.1
<i>Actinobacillus actinomycetemcomitans</i>	39	79.6	19	86.4	20	74
<i>Streptococcus mutans</i>	8	16.3	3	13.6	5	18.5
<i>Anaerobic lactobacillus</i>	2	4	2	9	0	0
<i>Enterobacteriaceae</i>	4	8	2	9	2	7.4
<i>Candida albicans</i>	4	8	2	9	2	7.4
Multi-infections (more than 2 isolates)	44	89.8	20	90.9	24	88.9
No growth of potential pathogens	5	10.2	2	9.1	3	11.1
Total significant growth	130					

Age was a dependent factor for periodontitis in the current study, with periodontitis rate being higher in the younger age groups (45.4% in <26 years), while it was lower at > 45 years (13.3%) (Table 1). Several authors believe that age is not a determining factor but a lifetime accumulation of disease³⁰⁻³². For people over the age of 31, the probability of developing gingivitis increased by 5.17 times, and the likelihood of developing periodontitis increased by 2.28 times³⁰⁻³². In the current study, when comparing the clinical features of periodontitis patients with other oral diseases, the clinical signs and symptoms in Table 2 such as swollen or puffy gums, bleed easily gums, halitosis, painful chewing, pus between teeth and gums, loose teeth or loss of teeth, gingival recession, spitting out blood when brushing or flossing teeth, and tender gums occur more frequently in periodontitis patients than in patients with other oral diseases. This finding is similar to that reported in the literature in which previous signs and symptoms appeared in periodontitis more than other dental diseases³³⁻³⁵. In the current study, bleeding gums occurred in 96% of periodontitis patients and occurred 67 times more (CI =15-284, $p<0.001$) than other oral diseases. In a previous study by Maduakor *et al.*,³⁶ patients with bleeding gums showed an odds increase of 38.41 times the incidence of gingivitis and 2.58 times the incidence of periodontitis compared to patients without bleeding gums. It has also been reported that bleeding gums is one of the early signs of developing gum disease. This confirms the effect of maintaining good oral health and hygiene associated with preventing inflammation^{37,38}.

In the current study higher prevalence of periodontitis among subjects with low education (27.3%) with a significant correlation between a low education level and periodontitis, in which subjects with low school education were 2.9 times more likely to have periodontitis than subjects with a higher level of education (OR=2.9 times, CI=1.5- 5.4, $p<0.001$) (Table 3). A high prevalence of periodontitis has been reported among low-education patients in Nigeria, Jordan and Thailand^{30,36,39}. This confirmed that in the United States of America, people with a low school education were three times more likely to have periodontitis than people with a higher education level⁴⁰ and it has been reported in many studies that there is a correlation between gum disease and educational level. In the current study, patients with a previous history of recurrent gingivitis were approximately 3.8 times more likely to have periodontitis than those without a previous history of gingivitis (OR=3.2, CI=1.7-6.1, $p<0.001$) (Table 3). A previous frequent positive history of gingivitis has been reported by several researchers as a risk factor for developing gingivitis^{30,36,39}. In the current study, poor oral health habits were not risk factors for periodontitis (odds ratio =0.4). This finding differs from most of the reported studies in that poor oral health habits (oral hygiene) lead to a risk factor for gingivitis and periodontitis³⁶⁻³⁸. This finding may be because in current study, patients were compared with other oral disorders and not with healthy individuals. Khat or qat (*Catha edulis*) is a flowering plant inhabitant to

Ethiopia. Khat contains the alkaline cathinone, which is a stimulant that causes excitement, loss of appetite and euphoria. Chewing khat has a history as a social habit going back thousands of years similar to the use of coca leaves in South America and betel nuts in Asia⁴¹. It is estimated that up to 90% of adult males chew qat three to four hours per day in Yemen. There was a significant association between Khat chewing and periodontitis (OR=5.3, CI=2.8-10.2, $p<0.001$). These results are similar to previous studies in Yemen where Khat chewing is a risk factor for oral diseases²⁴, and better research on Khat chewing and its potential association with oral and dental disorders should be conducted on a large scale. Aside from the physiological causes of xerostomia, the iatrogenic effects of medications are the most common cause⁴². A drug recognized to cause dry mouth can be called xerogenic⁴³. More than 400 drugs are associated with dry mouth. Although dry mouth caused by medications is usually reversible, the situations for which these medications are prescribed are often chronic⁴⁴. The likelihood of developing a xerostomia increases compared to the total number of drugs taken, whether or not individual drugs are dehydrating⁴⁵. The sensation of dehydration usually begins shortly after starting the offending drug or after increasing the dose⁴². There was a significant association between the use of drugs that cause xerostomia or gingival changes and the development of periodontitis (OR=11.8, CI=3.4-41, $p<0.001$). These results are similar to previous studies that reported several medications associated with dry mouth, which is a risk factor for periodontitis³⁶.

There was a significant correlation between obesity and develop periodontitis, (OR=2.7, CI=1.3-5.4, $p=0.004$). This result is similar to previous studies in which obesity is predisposing factors for dental and gums disorders³⁶. There was a significant association between conditions causing decreased immunity and the development of periodontitis (OR=4.3, CI=1.1-16.6, $p=0.02$); and some diseases such as diabetes, rheumatoid arthritis, and Crohn's disease (OR=6.1, CI =2.8-13.2, $p<0.001$); with the occurrence of periodontitis. These disorders can be described in several different ways: by the component (s) of the affected immune system, whether the immune system is overactive or inactive, and whether the condition is congenital or acquired⁴⁶. These conditions usually make people more susceptible to dental and other local or systemic infections⁴⁶. The polymicrobial pattern that is a feature of periodontal disease was obvious in this study, 89.8% in periodontitis, and this is in agreement with the work of many other researchers^{12,47-49}. The occurrence of polymicrobial infection has important inferences for management as it changes the clinical course of disease, influences the choice of antimicrobial therapy and the expected response to treatment especially when it comes to pathogens that commonly exhibit antimicrobial resistance such as *S. aureus*⁵⁰. *Aggregatibacter actinomycetemcomitans* was the most prevalent facultative anaerobe in periodontitis, 79.6%. Of all the microorganisms in biofilm, three are said to be important in the initiation and progression of periodontal disease: *A. actinomycetemcomitans* is named

key pathogens or “red complex” bacterium^{2,51}. *A. actinomycetemcomitans* in aggressive chronic periodontitis patients⁵². It is reported to be strongly associated with destructive periodontal lesions⁵. It possesses many virulence factors including protease, leukotoxin, endotoxin, collagenase, fibroblast inhibition factor inducing bone resorption⁵. *S. aureus* was isolated from 53.1% of used periodontitis patients, as did several researchers^{52,53}. These microbes are known to easily become resistant to antibiotics, and may reach climax with super-infection. The capability to form biofilm has enabled *S. aureus* to survive in this environment also^{49,54}. *S. pyogenes* was isolated from 73.5% and *S. mutans* in 16.3% of our patients, and this result is similar to that previously reported in which *Streptococcus* species was detected in large numbers by several researchers^{12,52}. Some *streptococci* are useful to the host as colonization of the pocket in large numbers can delay the periodontal disease process⁵⁵.

In current study, *Enterobacteriaceae* was isolated in 8% of patients with periodontitis (Table 4). *Enterobacteriaceae* is unusual in patients with periodontitis⁵⁶. Several studies have linked enteric bacilli to periodontal disease⁵². According to Botero and colleagues, their role in periodontitis is not clear but is thought to indicate super-infection⁵⁷. They are thought to be opportunists that thrive after periodontal treatment. The drug of choice for the treatment of periodontal disease includes tetracycline, doxycycline, amoxicillin, and metronidazole. The gut bacteria illustrate resistance to these drugs and may consequently persist after taking them. Further studies are needed to explain its presence in the plaque biofilm and explain its role in periodontal infection⁵⁷. *C. albicans* was isolated in current study in 8% of patients with periodontitis, and several researchers including Daniluk *et al.*, have reported that *C. albicans* could have a role in the ultrastructure of gingival microbial plaques and in their attachment to periodontal tissue^{25,58,59}.

CONCLUSIONS

This study is new in Sana'a city. The clinical features of periodontitis in Yemen and the risk factors are similar to those reported in the literature elsewhere, but the isolated bacteria differ in frequency from those reported elsewhere, as some upper respiratory tract pathogens such as *S. pyogens* are commonly isolated in this study. Knowledge of the clinical features, bacterial causes of gum disease, and risk factors is the key to successful periodontal therapy.

ACKNOWLEDGEMENTS

The authors extend their thanks and appreciation to Genius University of Science and Technology, Dhamar City, Republic of Yemen, which supported this work, in particular Dr. Mohammed Mohammed Ali Al-Najhi, the generous scholar who usually supports medical education and research in Yemen.

AUTHOR'S CONTRIBUTION

AL-Haddad KA: writing original draft, literature survey. **Al-Najhi MMA:** methodology, formal analysis, conceptualization. **Abbas AKM:** formal analysis, review. **Al-Akwa AAY:** investigation, conceptualization. **Al-Shamahy HA:** critical review, supervision. **Al-labani MA:** data curation, investigation. All authors revised the article and approved the final version.

DATA AVAILABILITY

The datasets generated during this study are available from the corresponding author upon reasonable request.

CONFLICT OF INTEREST

No conflict of interest associated with this work.

REFERENCES

1. Yamamoto M, Kobayashi R, Kono T, Bolerjack B, Gilbert R. Induction of IL-10-producing CD4 T-cells in chronic Periodontitis. *J Dental Res* 2011;90(5):653-658. <https://doi.org/10.1177/0022034510397838>
2. Segura EP, Mendez L, Marquez E, Vargas A, Gregorio K, Martinez JL, Ilyna A. Effect of *Carya Illinoensis*, *Quercus rubra* and *Smilax glycyphylla* extracts, pectin and papain on the dental biofilm microorganisms. *J Pharm Pharmacog Res* 2015; 3(5): 118-129. <https://doi.org/10.5580/IJMB.54104>
3. Tonetti MS, Jepsen S, Jin L, Otomo-Corgel J. Impact of the global burden of periodontal diseases on health, nutrition and wellbeing of mankind: A call for global action. *J Clin Periodont* 2017; 44:456-462. <https://doi.org/10.1111/jcpe.12732>
4. Suvan J, D'Aiuto F, Moles DR, Petrie A, Donos N. Association between overweight/obesity and periodontitis in adults. A systematic review. *Obesity Rev* 2011; 12(5):e381-404. <https://doi.org/10.1111/j.1467-789X.2010.00808.x>
5. Popova C, Dosseva-Panova V, Panov V. Microbiology of Periodontal diseases- A review. *Biotech Biotechnol Equip* 2013; 27(3), 3754-3759. <https://doi.org/10.5504/BBEQ.2013.0027>
6. Haffajee AD, Socransky SS. Periodontal Microbial Ecology 2005: *Periodontol* 2000. 2005; 38, 9-12.
7. Newman MG, Takei H, Klokkevold PR, Carranza FA. *Carrianza's clinical Periodontology* 10th ed, Middle East and African Edition: 2006.
8. Paster BJ, Olsen I, Aas JA, Dewhirst FE. The breadth of bacteria diversity in the human periodontal pocket and other oral sites. *Periodontol* 2006; 42: 80 – 87. <https://doi.org/10.1111/j.1600-0757.2006.00174.x>
9. Socransky SS, Haffajee AD. In: *Clinical Periodontology and Implant Dentistry*, 4th Ed. (J. Lindhe. T. Karring. N. Lang, Eds.) 2003: 106-149.
10. Colombo AP, Boches SK, Cotton SL, *et al.* Comparisons of subgingival microbial profiles of refractory periodontitis, severe periodontitis, and periodontal health using the humanoral microbe identification microarray. *J Periodontol* 2009; 80:1421-1432. <https://doi.org/10.1902/jop.2009.090185>
11. Socransky SS, Haffajee AD. Dental biofilms: difficult therapeutic targets. *Periodontol* 2002, 28; 12-55.
12. Egwari LO, Obisesan B, Nwokoye NN. Microbiological status of periodontal diseases in Lagos, Nigeria. *West Indian Med J* 2009; 58 (4): 392-397. *PMID: 20099785*

13. Chapple IL, Bouchard P, Cagetti MG, et al. Interaction of lifestyle, behaviour or systemic diseases with dental caries and periodontal diseases: consensus report of group 2 of the joint EFP/ORCA workshop on the boundaries between caries and periodontal disease. *J Clin Periodontol* 2017; 44(Suppl 18), S39-S51. <https://doi.org/10.1111/jcpe.12685>
14. Chapel IL, Van der Weijden F, Doerfer C, et al. Primary Prevention of Periodontitis: Managing gingivitis. *J Clin Periodontol* 2015; 45(Suppl 16), S71-S76. <https://doi.org/10.1111/jcpe.12366>
15. Peterson PE, Ogawa HM. The global burden of periodontal disease: Towards integration with chronic disease prevention and control. *Periodontol* 2000, 2012;60, 15-39. <https://doi.org/10.1111/j.1600-0757.2011.00425.x>
16. Al-Sharani AA, Al-Hajj W, Al-Shamahy HA, Jaadan BM. The effect of nanosilver and chlorhexidine mouthwash on anaerobic periodontal pathogens counts. *Universal J Pharm Res* 2019; 4(5): 1-6. <https://doi.org/10.22270/ujpr.v4i5.309>
17. Alhasani AH, Ishag RA, Yahya Al-Akwa AY, Al Shamahy HA, Al-labani MA. Association between the *Streptococcus mutans* biofilm formation and dental caries experience and antibiotics resistance in adult females. *Universal J Pharm Res* 2020; 5(6):18-23. <https://doi.org/10.22270/ujpr.v5i6.507>
18. Abbas AM, Al-Kibsi TAM, Al-Akwa AAY, AL-Haddad KA, Al-Shamahy HA, Al-labani MA. Characterization and antibiotic sensitivity of bacteria in orofacial abscesses of odontogenic origin. *Universal J Pharm Res* 2020; 5(6):36-42. <https://doi.org/10.22270/ujpr.v5i6.510>
19. Al- deen AAS, Al-deen HMS, Abbas AKM, Al-Akwa AAY, AL-Haddad KA, Al-Shamahy HAW, Al-Sharani HM, Al-labani MA. Knowledge and perception of molar incisor hypomineralization among dental practitioners in Sana'a city - Yemen. *Universal J Pharm Res* 2020; 5(5):4-11. <https://doi.org/10.22270/ujpr.v5i5.479>
20. AL-Awadi TAM, AL-Haddad KA, Al-labani MA, Al-Shamahy HA. Prevalence of malocclusion among Yemeni children of primary schools. *Universal J Pharm Res* 2019; 5(1): 1-6. <https://doi.org/10.22270/ujpr.v5i1.329>
21. Ullahman MAASA, Yahya A, Al-Shamahy HA, Abbas AKMA. Occurrence of retromolar canal among a sample of Yemeni adults obtained from cone-beam computed tomography. *Int Res J Med Med Sci* 2020; 8(2): 35-41.
22. Alhadi Y, Rassem AH, Al-Shamahy HA, Al-Ghaffari KM. Causes for extraction of permanent teeth in general dental practices in Yemen. *Universal J Pharm Res* 2019; 4(2): 1-5. <https://doi.org/10.22270/ujpr.v4i2.231>
23. Mutaher NJA, AL-Haddad KA, Al-Akwa AAY, Al-labani MA, Al-Shamahy HA, Zabara AQMQ, Al- deen HMS. Prevalence and causes of traumatic dental injuries to anterior teeth among primary school children in Sana'a city, Yemen. *Universal J Pharm Res* 2020; 5(3):38-43. <https://doi.org/10.22270/ujpr.v5i3.329>
24. Al-Kebsi AM, Arwa M Othman, Al-Kasem M A Abbas, Ebtihal M Madar, Hassan A. Al-Shamahy, Khaled M Al-Gaffari, Samera M. Naser Daname, Fuad L. Motareb. Oral c. albicans colonization and non-candida albicans candida colonization among university students, Yemen. *Universal J Pharm Res*. 2017; 2(5): 5-10. <http://doi.org/10.22270/ujpr.v2i5.R2>
25. Al-Haddad KA, Al-dossary OAE, Al-Shamahy HA. Prevalence and associated factors of oral non-Candida albicans Candida carriage in denture wearers in Sana'a city-Yemen. *Universal J Pharm Res* 2018; 3(4): 7-11. <https://doi.org/10.22270/ujpr.v3i4.176>
26. Joshi VM, Vandana KL. The detection of eight putative periodontal pathogens in adult and rapidly progressive periodontitis patients: An institutional study. *Indian J Dental Res* 2007; 18: 6-10. <https://doi.org/10.4103/0970-9290.30914>
27. Cheesbrough M. District laboratory practice in tropical countries. Cambridge: Cambridge University Press; 2010. <https://doi.org/10.1017/CBO9780511581304>
28. Page RC, Altman LC, Ebersole JL, Vandesteen GE, Dahlberg WH, Williams BL, Bowen T. Rapidly progressive periodontitis. A distinct clinical condition, 1983: *J Periodontol* 1983;54: 197-209. <https://doi.org/10.1902/jop.1983.54.4.197>
29. Albandar JM. Periodontal disease in North America. *Periodontol* 2000 2002a 29: 31-69. <https://doi.org/10.1034/j.1600-0757.2002.290103.x>
30. Ababneh KT, Hwajj Z.M., Khaders YS. Prevalence and risk indicators of gingivitis and periodontitis in a multicentre study in North Jordan: a Cross sectional study. *Biomedcentral Oral Health* 2012; 12:1. <http://www.biomedcentral.com/1472-6831/12/1>
31. Susin C, Albandar JM. Aggressive periodontitis in an urban population in Southern. Brazil *J Periodontol* 2005; 76: 468-475. <https://doi.org/10.1902/jop.2005.76.3.468>
32. Genco RJ. Current view of risk factors for periodontal diseases. *J Periodontol* 1996; 67;1041-1049. <https://doi.org/10.1902/jop.1996.67.10.1041>
33. Cisar JO, Kolenbrander PE, McIntire FC. Specificity of coaggregation reactions between human oral streptococci and strains of *Actinomyces viscosus* or *Actinomyces naeslundii*, *Infection and Immunity* 1989; 24(3); 742-752. <https://doi.org/10.1128/IAI.24.3.742-752.1979>
34. Johnson, JL, Moore LV, Kaneko B, Moore WE. *Actinomyces georgiae* sp. Nov., *Actinomyces gerencseriae* sp. Nov., designation of two genus species of *Actinomyces naeslundii* and inclusion of *A. naeslundii* serotypes 11 and 111 and *Actinomyces viscosus* serotype 11 in *A. naeslundii* genospecies 2. *Int J Syst Bacteriol* 1990; 40(3): 273-286.
35. Kumar PS, Griffen AL, Moeschberger ML, Leys, EJ. Identification of candidate periodontal pathogens and beneficial species by quantitative 16S clonal analysis. *J Clin Microbiol* 2005; 43(8): 3944-3955. <https://doi.org/10.1128/JCM.43.8.3944-3955.2005>
36. Maduakor UC, Onyemelukwe NF, Maduakor SN, Azubuike NC, Onyemelukwe AO, Nnede EB. Bacterial etiology and risk factors of periodontal diseases in enugu metropolis, South East Nigeria. *The Int J Microbiol* 2019; 16(1). <https://doi.org/10.5580/IJMB.54104>
37. Khader YS, Rice JC, Lefante JJ. Factors associated with periodontal diseases in a dental teaching clinic population in northern Jordan. *J Periodontol* 2003; 74:1610-1617. <https://doi.org/10.1902/jop.2003.74.11.1610>
38. Axelsson P, Nystrom B, Lindhe J The long term effect of plaque control program on tooth mortality, caries and periodontal disease in adults. Results after 30 years of maintenance. *J Clin Periodontol* 2004; 31:749-757. <https://doi.org/10.1111/j.1600-051X.2004.00563.x>
39. Torrungruang K, Tamsailom S, Rojanasomsith K, et al. Risk indicators of periodontal disease in older Thai adults. *J Periodontol* 2005; 76:558-565. <https://doi.org/10.1902/jop.2005.76.4.558>
40. Borrel LN, Butt BA, Warren RC, Neighbors HW. The role of individual and neighborhood social factors on periodontitis: the third National Health and Nutrition Examination Survey. *J Periodontol* 2006; 77:444-453.
41. Al-Mugahed, Leen. Khat Chewing in Yemen: turning over a new leaf: khat chewing is on the rise in yemen, raising concerns about the health and social consequences. *Bullet World Health Org* 2008; 86 (10): 741-42. <https://doi.org/10.2471/BLT.08.011008>
42. Scully, Crispian. Oral and maxillofacial medicine : the basis of diagnosis and treatment (2nd ed). Edinburgh: Churchill Livingstone 2008; 79-85. ISBN 978044306818 8.
43. Tyldesley, Anne Field, Lesley Longman in collaboration with William R. Tyldesley's Oral medicine (5th ed.). Oxford: Oxford University Press 2003; 90-93. ISBN 978-0192631473.

44. Furness, S; Worthington, HV; Bryan, G; Birchenough, S; McMillan, R. Furness, Susan (ed.). Interventions for the management of dry mouth: topical therapies. Cochrane Data System Rev 2011; (12): CD008934.
<https://doi.org/10.1002/14651858.CD008934.pub2>
45. Bouquot, Brad W. Neville, Douglas D. Damm, Carl M. Allen, Jerry E. Oral & maxillofacial pathology (2nd, ed.). Philadelphia: W.B. Saunders 2002; 398–399. ISBN 978-0721690032.
46. Geha RS, Notarangelo LD, Casanova JL, et al. Primary immunodeficiency diseases: an update from the International Union of Immunological Societies Primary Immunodeficiency Diseases Classification Committee. J Allergy Clin Immun 2007; 120 (4): 776–94.
<https://doi.org/10.1016/j.jaci.2007.08.053>
47. Saini S, Aparna N. Gupta N, Mahajan A. Microbial flora in Oro-dental infections. Indian J Med Microbiol 2003; 21 (2) 111-114. PMID: 17642993
48. Mane AK, Karmarkar AP, Bharadwaj RS. Anaerobic bacteria in subjects with chronic periodontitis and in periodontal health. J Oral Health Comm Dent 2009; 3(3):49-51. <https://doi.org/10.5005/johcd-3-3-49>
49. Salari MH, Kadhoda, Z. Rate of cultivable sub-gingival periodontopathogenic bacteria in chronic periodontitis. J Oral Sci 2004; 46(3):157-161.
<https://doi.org/10.2334/josnusd.46.157>
50. Peter BM, Jabra-Rizk MA, Graeme AO, Costerton JW, Shirtliff ME. Polymicrobial interactions: impact on pathogenesis and human disease. Clin Microbiol Rev 2012; 25(1): 193-213.
<https://doi.org/10.1128/CMR.00013-11>
51. Sanz M, van Winkelhoff AJ, Herrera D, Dellemjinkippu N, Simon R, Winkel E. Differences in the composition of the subgingival microbiota of two periodontitis populations of different geographic locations. A comparison between Spain and the Netherlands. European J Dental Sci 2000; 108:383-92.
<https://doi.org/10.1034/j.1600-0722.2000.108005383.x>
52. Amel Y, Bouziane D, Leila M, Ahmed, B. Microbiological study of periodontitis in the West of Algeria 2009: Advances in Medical and Dental Sciences. 2009 Sep 1:80-6.
53. Loberto J, Martins C, Santo S, Cortelli J, Jorge, A. *Staphylococcus* spp in the oral cavity and periodontal pockets. Brazilian J Microbiol 2004; 35:64-68.
<http://dx.doi.org/10.1590/S1517-83822004000100010>
54. Cuesta AI, Jewtuchowicz G, Brusca ML, Rosa A.C. Prevalence of *Staphylococcus* spp and *Candida* spp in the oral cavity and periodontals pockets of periodontal disease patients. Acta Odontologica Latinoam 2010; 23(1):20-26. PMID: 20645638
55. Baehni PC, Guggenheim B. Potential of diagnostic microbiology for treatment and prognosis of dental caries and periodontal diseases. Critical Rev Oral Biol Med 1996; 7(3):259-277.
<https://doi.org/10.1177/10454411960070030401>
56. Betancourth M, Arce R, Botero J, Jaramillo A, Cruz C, Contreras A. Unusual microorganisms in gingival sulcus and periodontal pockets 2006. Colombia Medica 2006; (37): 1-5.
57. Botero JE, Contreras A, Lafeaurie G, Jaramillo A, Betancourt M, Arce RM. Occurance of periodontopathic and super-infecting bacteria in chronic and aggressive periodontitis subjects in Colombian population. J Periodontol 2000; 78: 696-704.
<https://doi.org/10.1902/jop.2007.060129>
58. Daniluk T, Tokajuk G, Cylwik-Rokicka D, Rozkiewicz D, Zaremba ML, Stolowska W. Aerobic and anaerobic bacteria in subgingival and supragingival plaques of adult patients with periodontal disease. Adv Med Sci 2006; 51 Suppl 1:81-5. PMID: 17458065.
59. Jarvesivu A, Hietanen J, Rautemaa R, Sorsa T, Richardson M. *Candida* yeasts in chronic periodontitis tissues and subgingival microbial biofilms *in vivo*. Oral Diseases 2004; 10: 106-112.
<https://doi.org/10.1046/j.1354-523x.2003.00978.x>