

Type 1 Diabetes Mellitus Patients' Self-perception of Periodontal Diseases

Zaridah Zainal Abidin¹⁰ Erni Noor²⁰ Noor Shafina Mohd Nor^{3,40} Nor Shafina Mohamed Nazari⁵⁰ Azriyanti Anuar Zaini⁶⁰ Nurul Zeety Azizi⁷⁰ Shahrul Aiman Soelar⁸⁰ Marshah Mohamad Shahrizad⁹⁰ Rohaida Abdul Halim¹⁰

¹Centre of Paediatric Dentistry and Orthodontics Studies, Faculty of Dentistry, Universiti Teknologi MARA, Sungai Buloh, Selangor, Malaysia

² Centre of Studies for Periodontology, Faculty of Dentistry, Universiti Teknologi MARA, Sungai Buloh, Selangor, Malaysia

³ Department of Paediatrics, Faculty of Medicine, Universiti Teknologi MARA, Sungai Buloh, Selangor, Malaysia

⁴Institute for Pathology, Laboratory and Forensic Medicine (I-PPerForM), Faculty of Medicine, Universiti Teknologi MARA, Sungai Buloh, Selangor, Malaysia

⁵ Department of Restorative Dentistry, Faculty of Dentistry, Universiti Malaya, Kuala Lumpur, Malaysia

Eur J Dent 2024;18:534-543.

Abstract

Address for correspondence Rohaida Abdul Halim, BDS (Malaya), DClinDent Paediatric Dentistry (Otago), DipClinHyp (LCCH), AM, FIAPD, Centre of Paediatric Dentistry and Orthodontics Studies, Faculty of Dentistry, Universiti Teknologi, MARA 47000 Sungai Buloh, Selangor, Malaysia (e-mail: dr_rohaida@uitm.edu.my).

⁶ Paediatric Department, Faculty of Medicine, Universiti Malaya, Kuala Lumpur, Malaysia

- ⁷ Department of Paediatric Dentistry and Orthodontics, Faculty of Dentistry, Universiti Malaya, Kuala Lumpur, Malaysia
- ⁸ Clinical Research Centre, Hospital Sultanah Bahiyah, Alor Setar, Kedah, Malaysia
- ⁹Kuching Division Dental Office, Sarawak State Dental Health Department, Braang Bayur Dental Clinic, Sarawak, Malaysia

Objectives The study aimed to evaluate type 1 diabetes mellitus (T1DM) patients' self-perceived periodontal health status and to identify the association between periodontal disease (PD) and DM.

Materials and Methods This cross-sectional study included 113 T1DM children between 3 and 18 years old from the Universiti Teknologi MARA and the University of Malaya. Periodontal health parameters, including plaque index, gingival index, probing pocket depth, simplified basic periodontal examination, and clinical attachment loss, were recorded. Self-perceived periodontal health status was assessed with questionnaires.

Statistical Analysis Statistical analysis was performed to evaluate the sensitivity of the questionnaire and the relationship between T1DM and periodontal parameters. **Results** The median age was 11.4 years. Half of them (50.4%) were females. A total of

83.5% rated their oral condition as good, whereas 27.5% reported a history of gingival

bleeding. Clinical examination revealed that 48.7% had healthy gingiva, whereas 47.8%

had gingivitis. The question "Do you have bleeding when brushing, flossing, or eating

Keywords

- periodontal disease
- type 1 diabetes mellitus
- diabetes mellitus
- ► oral health
- self-perception

food?" showed good accuracy in the evaluation of PD (p < 0.001). **Conclusion** The questionnaire has a high potential to be used by medical professionals in identifying T1DM patients at risk of PD to guide nondental health care providers in making appropriate referrals to dental services.

article published online December 4, 2023 DOI https://doi.org/ 10.1055/s-0043-1772777. ISSN 1305-7456. © 2023. The Author(s).

This is an open access article published by Thieme under the terms of the Creative Commons Attribution License, permitting unrestricted use, distribution, and reproduction so long as the original work is properly cited. (https://creativecommons.org/licenses/by/4.0/) Thieme Medical and Scientific Publishers Pvt. Ltd., A-12, 2nd Floor, Sector 2, Noida-201301 UP, India

Introduction

Diabetes mellitus (DM) is a metabolic disease characterized by the dysregulation of carbohydrate metabolism. It commonly manifests as hyperglycemia due to diminished insulin secretion, impaired insulin action, or both.¹ Type 1 diabetes mellitus (T1DM) makes up 5 to 10% of all DM cases worldwide.² The disease stems from cell-mediated autoimmune destruction of the pancreatic β -cells that produce insulin. Depending on the degree of β -cell destruction, T1DM patients can experience reduced or absence of insulin secretion, as indicated by low or negligible levels of plasma C-peptide. The chronic hyperglycemic status can lead to long-term damage, dysfunction, and failure of different organs in DM patients, especially the eyes, kidneys, nerves, heart, and blood vessels.³ Treatment modalities for T1DM include subcutaneous injections of insulin, maintenance of a healthy diet, and regular exercise.⁴

Periodontal disease (PD) is a condition resulting from the infection and inflammation of the tooth-supporting tissues.⁵ In the early stages of the disease, gingival inflammation may cause the gingival tissues to appear erythematous, or edematous, and potentially result in bleeding. Worse, gingivitis can progress into periodontitis that presents as attachment loss, alveolar bone resorption, and tooth mobility.⁶ PD is one of the most common causes of tooth loss.⁷ DM is associated with an increase in the prevalence, severity, and progression of periodontitis.^{8,9} DM patients are three times more likely to suffer from periodontitis as compared with nondiabetics.¹⁰ Furthermore, 10% of children with T1DM were reported to have higher rates of attachment loss and bone loss compared with their nondiabetic peers despite having comparable plaques score.¹¹ Moreover, a recent study showed that children with DM have twice the number of periodontal sites that eventually developed into periodontitis as compared with non-DM children.¹²

The bidirectional association between DM and PD has been established in previous literature.^{13,14} DM is known as one of the modifying factors of periodontitis that can accelerate the progress of PD¹⁴ as DM can impair the periodontal tissue growth and matrix formation with fibroblasts, osteoblasts, and osteoclasts.^{15,16} In addition, increased thickness of the gingival basement membrane in DM patients could also impair the vasculature of the periodontal tissues.^{17,18} It has also been postulated that DM complicates PD by tipping the balance of oral microbiota, resulting in the dominance of periodontal pathogens.^{19,20} However, more in-depth studies are needed to ascertain the differences in the biofilm of diabetics and nondiabetics. The latest evidence indicated that chronic inflammation in PD can aggravate complications of DM by worsening glycemic control.

The American Dental Association recommendation for diabetic patients is to receive medical follow-up on a 3-monthly basis.²¹ During these visits, evaluation of hemoglobin A1c (HbA1c) and reassessment of diabetes management are recommended. However, T1DM patients with unsatisfactory glycemic control should be arranged for more frequent follow-ups to enhance their adherence to the treatment regime. During this quarterly visit, T1DM patients would be able to obtain the maximum benefits from the other multidisciplinary team of specialists, thus reinforcing good self-care practices, such as routine dental visit.^{22,23} Apart from having known risk of PD, poor oral hygiene was observed in youngsters with T1DM especially those with uncontrolled HbA1c, which eventually could increase the risk of future oral disease(s).²⁴

In 2020, a study by Moore et al on professional health care workers in pediatric diabetes care teams showed that 76.2% of them were aware that periodontitis is a possible complication of diabetes. However, in 2022 Siddiqi and Zafar recorded a contradicting finding in which they found that the majority of medical practitioners (89%) were aware of the bidirectional association and knew that the glycemic index of patients with DM and suffering PDs could be improved by providing periodontal therapy.²⁵ However, as low as 4.8% had received training in recognizing patients who require dental care. This study reinforced the need for further training in this area to provide holistic care to DM patients.^{26,27} There is an urgent need for accurate and reliable means of surveillance, detection, and diagnosis of periodontitis among children suffering from T1DM. In addition to that, referral to dental care professionals for appropriate and timely management is crucial, therefore appropriate screening tools for medical professionals to initiate referral of patients with DM is needed.²⁸ Hence, it is vital to establish an effective system for nondental health providers to identify individuals in need of dental care.²⁹ However, with limited resources available for regular screening and timely examination by dental practitioners, other options should be explored.

In 2022, a study by Mohd Said from Malaysia recommended the use of validated simplified digital periodontal health screening software for identification of PD at early stage in dental practice.³⁰ This identification tool appeared simple and appropriate to be used for screening of PDs. This showed an effort have been made to initiate the link from the general dental practice to specialist care. Nevertheless, the initiation from the medical practitioner has yet to be established. Hence, there is a need of questionnaires is of detecting populations at risk of oral diseases, which can be initiated from the medical professionals. On top of that, the accuracy and effectiveness of using a questionnaire when compared with the clinical examination ranged from moderate to high in various studies, thus indicating its potential to be can be administered to instil awareness, facilitate early detection, and predict the disease.³¹ Therefore, in this context, a selfself-reported perception and the clinical parameters could contribute to the prevention and earlier diagnosis of PD, especially in individuals requiring complex clinical care.³²

Therefore, the use of adapted self-report measures for PD can be considered a low-cost alternative for the early detection and prevention of PD in DM patients. It can be beneficial for epidemiological studies and population surveillance of the periodontal condition.³³ This study aimed to compare the self-perceived periodontal health status using a guided question-naire (GQ) with the clinical measurement by dental professionals among children and adolescents with T1DM. The study also set out to determine if the GQ is a valid tool to be used by nondental professionals to evaluate periodontal health status.

Materials and Methods

This two-part study was a cross-sectional study conducted from October 2020 to May 2022 at two centers, Universiti Teknologi MARA (UiTM), Selangor, and the University of Malaya (UM), Kuala Lumpur. Content validity index for item (I-CVI) and face validity was conducted to validate the questionnaire. A reliability study of 20 participants and intraexaminer training and calibration was performed prior to the study.

All children and adolescents below 18 years old with T1DM diagnosis at UM and UiTM were invited to participate in the study. They must be able to communicate in English and or Bahasa Melayu. However, those undergoing active orthodontic therapy or using any antibiotics or medications in the last 3 months that might cause gingival alteration, such as drug-induced gingival enlargement, were excluded.

The sample size was determined using Epi-Info StatCal software based on the total number of the eligible T1DM patient $(n = 166)^{34}$ and the prevalence of diabetes test knowledge was 50.4% $(11.6/23 \times 100)$.³⁵ Considering an attrition rate of 10% with a 95% confidence level and an acceptable margin of error of 5.6%, the final sample calculated was 118 (108 + 10%). A total of 113 T1DM patients were recruited. The participants and their parent(s) or caregiver(s) answered the questionnaire during their follow-ups. The questionnaire would be answered by parents of children below 16 years old. Subsequent appointments were arranged for the participants to undergo a dental examination.

Definition

Body Mass Index

Body mass index (BMI) was calculated as weight in kilograms divided by height in meters squared, i.e., $BMI = weight (kg)/height2 (m^2)$. The calculated BMI was plotted on the Centers for Diseases and Prevention (CDC) BMI-to-age chart for the respective gender³⁶ (**-Table 1**).

Table 1 Body Mass II	ndex classification
----------------------	---------------------

BMI classification	Definition
Underweight	<5th percentile
Normal weight	<85th centile
Overweigh	≥85th but <95th centile
Obesity	\geq 95th centile

Abbreviation: BMI, body mass index.

Blood Pressure

In this study, blood pressure (BP) was reported as systolic (SBP) and diastolic (DBP) percentiles for age/sex/height. The classification is based on the individual's age plotted against the percentile of height in SBP or DBP (mm Hg) of the respective gender.³⁷ The height percentile was obtained using the CDC height-for-age percentile of the respective gender. Based on American Academy of Pediatrics, BP categories and stages are as follows³⁸ (**►Table 2**).

Lipid Profile

According to the International Society for Pediatrics and Adolescent Diabetes and the American Diabetes Association recommended low-density lipoprotein cholesterol (LDL-C) of < 100 mg/dL (2.6 mmol/L) in youth with DM.³⁹

High-density lipoprotein cholesterol (HDL-C), total cholesterol, triglycerides can be categorized according to NHLBI 2011.⁴⁰

Hemoglobin A1c

HbA1c is a glycoprotein formed by a direct reaction between blood glucose and hemoglobin. It is routinely in clinical research and clinical practice to evaluate diabetes control. For children, adolescents, and young adults \leq 25 years old. With access to comprehensive care, HbA1c of < 53 mmol/dL (7.0%) is recommended.⁴¹

In children aged 1–13 y	bld
BP classification	Definition
Normal	<90th percentile
Elevated	\geq 90th percentile to <95th percentile, or 120/80 mm Hg to <95th percentile (whichever is lower)
Stage 1 hypertension (HTN)	\geq 95th percentile to <95th percentile + 12 mm Hg, or 130/80–139/89 mm Hg (whichever is lower)
Stage 2 HTN	\geq 95th percentile $+$ 12 mm Hg or \geq 140/90 mm Hg (whichever is lower)
In children aged \ge 13 y c	bld
BP classification	Definition
Normal	<120/<80 mm Hg
Elevated	120/< 80–129/< 80 mm Hg
Stage 1 HTN	130/80–139/89 mm Hg
Stage 2 HTN	≥140/90 mm Hg

Table 2 Blood pressure classification

Abbreviations: BP, blood pressure; HTN, hypertension.

Study Instrument

Guided Questionnaire

Self-reported questionnaires adopted from two recent studies were translated and validated to be used in this study.^{42,43} The GQ consisted of 14 items with 8 items on patients' baseline characteristics (Part A) and 6 items on symptoms of PD (Part B). The GQ was used to screen for any PD comorbidity among the study participants.

Clinical Examination

Simplified Basic Periodontal Examination

The simplified basic periodontal examination (sBPE) codes formed the basis of the assessment for patients under 18 years old. The examined teeth included one tooth from each sextant, i.e., the upper right six (tooth 16), the upper right one (tooth 11), upper left six (tooth 26), lower left six (tooth 36), lower left one (tooth 31), and lower right six (tooth 46). The WHO 621 probe with a light probing force of 20 to 25 g was used for this assessment. The sBPE codes were as follows: 0, healthy; 1, bleeding on gentle probing; 2, calculus present and/or plaque retention factors; 3, the presence of 4- to 5-mm pocket; and 4, the presence of 6 mm or more pocket; and *, furcation.⁴⁴

In children between 12 and 17 years with erupted permanent teeth, the full range of sBPE codes (0-4) was used. For children aged between 7 and 11 years with mixed dentition, the sBPE codes 0,1, and 2 were used. In the full primary dentition, sBPE similar to the one used for children as young as 3 years of age was performed.⁴⁵ Apart from that, plaque index (PI)⁴⁶ and gingival index (GI)⁴⁷ were also assessed.

Statistical Analysis

Data analysis was performed using the Statistical Package for the Social Sciences Version 20.0. Association between the GQ and the clinical examination was assessed using the Pearson chi-square test and Fisher's exact test. A *p*-value < 0.05 was considered statistically significant. Diagnostic tests (sensitivity, specificity, accuracy, positive predictive value, negative predictive value, and receiving operating characteristic curve) were performed to measure the performance of each question with the basic periodontal examination (BPE) as the reference for the periodontal evaluation. For this purpose, the periodontal status was dichotomized as "0" for healthy and "1 and above" for having PD.

Results

Questionnaire I-CVI value 0.96 indicate high value of all items. And, following changes made based on expert opinion, a good face validation could be seen. Intraexaminer training and calibration (intraclass correlation coefficient revealed a good value of 85%) with acceptable reliability Cronbach's α of 0.77. **Table 3** outlines the sociodemographic, clinical parameters, and oral hygiene care characteristics of the participants. From the 113 participants, 24.8% (n = 28) presented with stage 1 hypertension, whereas another 18.6%

Table 3 Sociodemographic, clinical parameters, and oralhygiene care of participants

SexMale56(49.6)Female57(50.4)Age7(14.2)7-12 y39(34.5)13-18 y58(51.3)Race27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (24.8)Stage 1 hypertension28(24.8)Stage 1 hypertension28(24.8)Illucose21(18.6)EBG ⁴¹ (47.8)(47.8)Jucose12(16.6)Indian54(56.6)Elevated12(18.6)Stage 1 hypertension54(36.6)Bodicose12(16.6)Impaired fasting glucose12(16.6)Diabetes fasting glucose12(16.6)Diabetes fasting glucose(47.8)(47.8)Diabetes fasting glucose(47.8)(47.8)Diabetes fucose tolerance(31.6)(55.8)BMI ³⁶ (50.2)(55.8)Diabetes glucose tolerance(53.9)(55.8)Diabetes glucose27(23.9)Diabetes glucose(53.9)(55.8)BMI ³⁶ (50.2)(50.2)HbA1c ⁴¹ (27.9)(23.9)Controlled9(8.0)Uncontrolled104(92.0)Lipid profile24.00(26.5)Borderline24.00(26.5)Borderline24.00(26.5)Borderline24.00	Variables	n	(%)
Male56(49.6)Female57(50.4)Age16(14.2)7-12 y39(34.5)13-18 y58(51.3)Race27(23.9)Malay55(48.7)Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (24.8)Yormal64(56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)Mormal fasting glucose54(47.8)Impaired fasting glucose12(16.6)Diabetes fasting glucose12(16.6)Diabetes fasting glucose12(16.6)Diabetes fasting glucose12(25.8)Diabetes fasting glucose(47.8)(47.8)Impaired fasting glucose11(9.7)RBC ⁴¹ 11(9.7)KBC ⁴¹ (55.8)(55.8)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance27(23.9)Diabetes glucose tolerance27(23.9)Diabetes glucose tolerance(56.9)(56.9)BMI ³⁶ (27.9)(52.8)Controlled9(8.0)Uncontrolled104(92.0)HbA1c ⁴¹ (24.9)(25.9)Controlled24(21.2)HbA1c ⁴¹ (24.9)(26.9)Borderline24(21.9)Hoderline24<	Sex		
Female57(50.4)Age<< 6 y	Male	56	(49.6)
Age< 6 y	Female	57	(50.4)
< 6 y16(14.2)7-12 y39(34.5)13-18 y58(51.3)Race(34.7)Malay55(48.7)Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (48.7)Normal64(56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)PBG ⁴¹ (47.8)(47.8)glucose12(10.6)Impaired fasting glucose12(10.6)Diabetes fasting glucose11(9.7)RBG ⁴¹ (41.6)(55.8)Normal glucose39(34.5)Diabetes fasting glucose11(9.7)Impaired glucose tolerance39(35.8)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance(47.9)(41.6)Diabetes glucose tolerance(55.8)(55.8)Diabetes glucose tolerance(60.2)(55.8)Diabetes glucose tolerance(61.2)(62.2)Doesity7(63.2)(62.2)Obesity9(8.0)(62.2)Uncontrolled9(8.0)Uncontrolled9(8.0)Uncontrolled104(92.0)High41(36.3)Missing18(15.9)	Age		
7-12 y39(34.5)13-18 y58(51.3)Race(51.3)Malay55(48.7)Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (18.6)Elevated21(18.6)Stage 128(24.8)hypertension54(47.8)glucose12(10.6)glucose12(10.6)glucose47(41.6)glucose11(9.7)Diabetes fasting glucose11(9.7)RBG ⁴¹ (55.8)Normal glucose39(34.5)Diabetes fasting glucose11(9.7)Impaired glucose tolerance63(55.8)Diabetes glucose tolerance63(50.2)Dobesity7(23.9)Obesity104(92.0)HbA1c ⁴¹ 5(24.2)Uncontrolled9(8.0)Uncontrolled9(8.0)Uncontrolled9(8.0)Uncontrolled104(92.0)HbA1c ⁴¹ 5(24.2)High41(36.3)Missing18(15.9)	<6 y	16	(14.2)
13-18 y58(51.3)RaceMalay55(48.7)Chinese27(23.9)Indian30(26.5)Others10(0)Blood pressure ³⁸ (56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)PBG ⁴¹	7–12 y	39	(34.5)
RaceMalay55(48.7)Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)PBG ⁴¹ 54(47.8)glucose12(10.6)Impaired fasting glucose12(10.6)Diabetes fasting glucose47(41.6)Diabetes fasting glucose11(9.7)Normal glucose39(34.5)Diabetes fasting glucose39(34.5)Diabetes glucose tolerance63(55.8)BMI ³⁶ 11(9.7)Underweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(23.9)Uncontrolled9(8.0)Uncontrolled9(8.0)Uncontrolled9(8.1)High41(36.3)Missing18(15.9)	13–18 y	58	(51.3)
Malay55(48.7)Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)RBG ⁴¹ (78.8)(10.6)Impaired fasting glucose54(10.6)Jucose12(10.6)glucose47(41.6)RBC ⁴¹ (9.7)(10.6)Normal glucose39(34.5)Childerance39(34.5)Diabetes fasting glucose39(34.5)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance27(23.9)Overweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(5.2)HbA1c ⁴¹ (9.7)Controlled9(8.0)Uncontrolled104(92.0)Lipid profile104(92.0)High41(36.3)	Race		
Chinese27(23.9)Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (76.6)Elevated21(18.6)Stage 1 hypertension28(24.8)FBG ⁴¹ 28(47.8)glucose12(10.6)glucose47(41.6)glucose11(9.7)Diabetes fasting glucose39(34.5)KBG ⁴¹ 11(9.7)Normal glucose39(34.5)Diabetes fasting glucose63(55.8)Diabetes glucose tolerance11(9.7)Diabetes glucose tolerance27(23.9)Overweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(6.2)HbA1c ⁴¹ 104(92.0)Lipid profile104(92.0)High30(26.5)Borderline30(26.5)Borderline31(35.3)Borderline30(26.5)Sorderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(26.5)Borderline30(2	Malay	55	(48.7)
Indian30(26.5)Others1(0.9)Blood pressure ³⁸ (56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)FBC ⁴¹ 28(24.8)rbgc 4112(10.6)glucose12(10.6)glucose47(41.6)glucose39(34.5)rbgertension39(34.5)cloreance39(34.5)Diabetes fasting glucose39(55.8)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance11(9.7)Healthy weight68(60.2)Overweight114(9.7)Healthy weight68(60.2)Overweight104(23.9)Obesity7(6.2)HbA1c ⁴¹ 104(92.0)Lipid profile104(92.0)Lipid profile24(21.2)High41(36.3)Missing18(15.9)	Chinese	27	(23.9)
Others1(0.9)Blood pressure3864(56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)FBC4112(10.6)glucose54(47.8)glucose12(10.6)glucose47(41.6)glucose39(34.5)Cherance63(55.8)Diabetes fasting glucose39(34.5)Impaired glucose tolerance39(34.5)Diabetes glucose tolerance63(55.8)BMI3611(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(6.2)HbA1c ⁴¹ (27.0)(50.7)Controlled9(8.0)Uncontrolled10492.00Hiph ofile24(21.2)High18(15.9)	Indian	30	(26.5)
Blood pressure ³⁸ Normal 64 (56.6) Elevated 21 (18.6) Stage 1 hypertension 28 (24.8) FBC ⁴¹ 28 (24.8) FBC ⁴¹ 54 (47.8) glucose 12 (10.6) glucose 47 (41.6) glucose 11 (9.7) Objectes fasting glucose 39 (34.5) RBC ⁴¹ 53 (55.8) Normal glucose 63 (55.8) Diabetes glucose tolerance 63 (50.2) Diabetes glucose tolerance 27 (23.9) Overweight 114 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ (92.0) (11.0) Uncontrolled 9 (8.0) Uncontrolled 104 (92.0) Upid profile 104 (92.0) Upid profile </td <td>Others</td> <td>1</td> <td>(0.9)</td>	Others	1	(0.9)
Normal64(56.6)Elevated21(18.6)Stage 1 hypertension28(24.8)FBC ⁴¹ 28(47.8)Normal fasting glucose54(47.8)Impaired fasting glucose12(10.6)Diabetes fasting glucose47(41.6)Diabetes fasting glucose39(34.5)Normal glucose tolerance39(34.5)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance63(60.2)Overweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(6.2)HbA1c ⁴¹ (97.0)(8.0)Lipid profile104(92.0)Lipid profile104(92.0)High41(36.3)Missing18(15.9)	Blood pressure ³⁸		
Elevated21(18.6)Stage 1 hypertension28(24.8)FBC ⁴¹ 28(24.8)FBC ⁴¹ 54(47.8)glucose12(10.6)glucose12(10.6)glucose47(41.6)glucose11(9.7)cloerance39(34.5)tolerance63(55.8)tolerance63(55.8)Diabetes glucose tolerance63(60.2)Diabetes glucose tolerance63(60.2)Overweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(6.2)HbA1c ⁴¹ (97.0)Lipid profile9(8.0)Lipid profile104(92.0)Lipid profile24(21.2)High41(36.3)Missing18(15.9)	Normal	64	(56.6)
Stage 1 hypertension28(24.8)FBC ⁴¹ Normal fasting glucose54(47.8)Impaired fasting glucose12(10.6)Diabetes fasting glucose47(41.6)Diabetes fasting glucose11(9.7)RBC ⁴¹ 11(9.7)Normal glucose tolerance39(34.5)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance63(55.8)Diabetes glucose tolerance63(60.2)Diabetes glucose tolerance27(23.9)Obesity7(6.2)Overweight104(92.0)HbA1c ⁴¹ (92.0)Lipid profile104(92.0)Lipid profile24(21.2)Acceptable30(26.5)Borderline24(21.2)High18(15.9)	Elevated	21	(18.6)
FBG ⁴¹ S4 (47.8) Normal fasting glucose 12 (10.6) Impaired fasting glucose 47 (41.6) Diabetes fasting glucose 47 (41.6) RBG ⁴¹ (47.8) (41.6) Normal glucose 11 (9.7) tolerance 39 (34.5) Diabetes glucose tolerance 63 (55.8) BMI ³⁶ (60.2) (9.7) Underweight 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ (92.0) (104 Uncontrolled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile 104 (92.0) Lipid profile 104 (26.5) Borderline 30 (26.5) Borderline 24 (21.2) High 41 (36.3)	Stage 1 hypertension	28	(24.8)
Normal fasting glucose54(47.8)Impaired fasting glucose12(10.6)Diabetes fasting glucose47(41.6)RBG ⁴¹	FBG ⁴¹		
Impaired fasting glucose12(10.6)Diabetes fasting glucose47(41.6)RBG ⁴¹	Normal fasting glucose	54	(47.8)
Diabetes fasting glucose 47 (41.6) RBG ⁴¹	Impaired fasting glucose	12	(10.6)
RBG ⁴¹ (9.7) Normal glucose tolerance 11 (9.7) Impaired glucose tolerance 39 (34.5) Diabetes glucose tolerance 63 (55.8) BMI ³⁶ (57.8) (57.8) Underweight 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ (92.0) (8.0) Uncontrolled 104 (92.0) Lipid profile T T Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Diabetes fasting glucose	47	(41.6)
Normal glucose tolerance11(9.7)Impaired glucose tolerance39(34.5)Diabetes glucose tolerance63(55.8)BMI ³⁶ (57.8)Underweight11(9.7)Healthy weight68(60.2)Overweight27(23.9)Obesity7(6.2)HbA1c ⁴¹ (92.0)Lipid profile104(92.0)Lipid profile104(92.0)High30(26.5)Borderline24(21.2)High18(15.9)	RBG ⁴¹		
Impaired glucose tolerance 39 (34.5) Diabetes glucose tolerance 63 (55.8) BMI ³⁶ 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ 9 (8.0) Uncontrolled 104 (92.0) Lipid profile T T Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Normal glucose tolerance	11	(9.7)
Diabetes glucose tolerance 63 (55.8) BMI ³⁶ Underweight 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile TG ⁴⁰ Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Impaired glucose tolerance	39	(34.5)
BMI ³⁶ (9.7) Underweight 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹	Diabetes glucose tolerance	63	(55.8)
Underweight 11 (9.7) Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ (6.2) (6.2) Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile	BMI ³⁶		
Healthy weight 68 (60.2) Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ 7 (6.2) Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile TG ⁴⁰ (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Underweight	11	(9.7)
Overweight 27 (23.9) Obesity 7 (6.2) HbA1c ⁴¹ (6.2) Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile	Healthy weight	68	(60.2)
Obesity 7 (6.2) HbA1c ⁴¹ Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile TG ⁴⁰ Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Overweight	27	(23.9)
HbA1c ⁴¹ Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile	Obesity	7	(6.2)
Controlled 9 (8.0) Uncontrolled 104 (92.0) Lipid profile	HbA1c ⁴¹		
Uncontrolled 104 (92.0) Lipid profile TG ⁴⁰ Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Controlled	9	(8.0)
Lipid profile TG ⁴⁰ Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Uncontrolled	104	(92.0)
TG ⁴⁰ 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Lipid profile		
Acceptable 30 (26.5) Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	TG ⁴⁰		
Borderline 24 (21.2) High 41 (36.3) Missing 18 (15.9)	Acceptable	30	(26.5)
High 41 (36.3) Missing 18 (15.9)	Borderline	24	(21.2)
Missing 18 (15.9)	High	41	(36.3)
	Missing	18	(15.9)

Variables	n	(%)
TC ⁴⁰		
Acceptable	31	(27.4)
Borderline	34	(30.1)
High	30	(26.5)
Missing	18	(15.9)
LDL-C ³⁹		
Acceptable	48	(42.5)
Borderline	25	(22.1)
High	22	(19.5)
Missing	18	(15.9)
HDL-C ⁴⁰	-	
Low	13	(11.5)
Borderline	6	(5.3)
Acceptable	75	(66.4)
Missing	19	(16.8)
Duration of diabetes		
< 5 y	66	(58.4)
5–10 y	38	(33.6)
> 10 y	9	(8.0)
Frequency of brushing		
Morning	17	(15.0)
Morning/before going to sleep	72	(63.7)
Morning/before going to sleep and after eating food	22	(19.5)
Use flossing	1	(0.9)
Others/adjunct	1	(0.9)

Table 3 (Continued)

Abbreviations: BMI, body mass index; FBG, fasting blood glucose; HbA1c, hemoglobin A1c; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; RBG, random blood glucose; TC, total cholesterol; TG, triglycerides.

(n = 21) had elevated BP. Despite only 6.2% of the participants being obese, as high as 92% of them recorded uncontrolled HbA1c. Similarly, 19.5% of the participants were found to have high LDL-C despite normal body weight.

Further analysis revealed that 40.9 and 53.8% of participants with healthy BMI exhibited high LDL-C and low HDL-C, respectively (**-Table 4**). In addition, uncontrolled HbA1c was observed among 91.2% of participants despite normal BMI. A higher percentage of uncontrolled HbA1c was also reported among participants with acceptable LDL-C (87.5%), acceptable HDL-C (90.7%), and less than 5 years of DM (90.9%). However, all these associations were not statistically significant (p > 0.05) (**-Table 5**).

• Table 6 shows that 27.5% of participants who reported bleeding from gingiva upon brushing, flossing, and eating were found to have BPE during the clinical examination. This finding

indicated a high sensitivity (50%) and specificity of the questionnaire (94.5%). In addition, 90% of T1DM presented with gingival bleeding were at risk of having PD (p < 0.001).

As presented in **-Table 7**, the mean PI and GI were 0.37 ± 0.31 and 0.27 ± 0.32 , respectively. In other words, 48.7% of the participants had healthy periodontal status. Among 47.8% of participants with unhealthy gingiva, 7% of them were found to have periodontitis.

Discussion

PD is the sixth most common complication among DM patients.⁴⁸ As T1DM patients face an increased risk of PD,⁴⁹ it is vital to improve the awareness of medical professionals and patients on the prevention and identification of oral diseases. Medical-dental coordinated care needs to be strengthened for this purpose. The use of self-reported questionnaires can increase T1DM patients' awareness and self-perception of oral diseases. During most medical consultations, physicians and other nonoral health professionals often overlook the need for oral clinical examination.³² Hence, the use of a self-reported questionnaire can also evoke their attention to DM patients at risk of PD. Following that, health care providers can refer the patients to a dentist for further management to prevent oral diseases.⁵⁰

In this study, it was found that respondents who answered "yes" to the question "Do you have bleeding when brushing, flossing, or eating food?" were associated with a high score of BPE compared with those who answered "no." The sum of sensitivity plus specificity for this item was 144%, which is considered as a "good validity."³³ Agreed by study in Japan population, inclusion of the question on gum bleeding would improve the predictive performance of the questionnaire as it is less confusing than other items.⁵¹ This finding also echoed a few other studies in which the term gum bleeding should be used rather than gingivitis when interacting with patients ^{52–54} as not all may be familiar with the dental term of gingivitis, especially children and adolescents. In addition, Elhassan et al (2017) also suggested that patients could relate to bleeding better than the appearance of swollen gum.⁵⁵

According to the Nelson's validity classification, self-reported PD (painful gums, tooth mobility, and people's opinions whether they have gums disease) can be classified as having moderate to high validity.^{33,56} However, in our study, the items "Do you think you have gum disease," "Swelling, red, or painful gums for no apparent reason," and "Loosening or shifting of teeth in the affected area" were found to have low sensitivity of 16.7, 13, and 3%, respectively. The reason probably due to this set of questionnaire has good predictive ability for periodontitis, especially in the severe cases rather than gingivitis alone, as only 3.5% patients in our study presented with periodontitis is high, the predictive performance of similar self-reported questions presented reported to be more accurate.⁵⁷

In this study, the questionnaire on the perception of oral health for participants aged less than 16 years old (72.6%) was answered by their parents or caregiver. With a low sensitivity (18.7%) and high specificity (86%), it can be

Variables	Underw	veight	Healthy	weight	t Overweight		Obesity		p-Value
	n	(%)	n	(%)	n	(%)	n	(%)	
LDL-C									
Acceptable	2	(4.2)	33	(68.8)	10	(20.8)	3	(6.3)	0.180 ^a
Borderline	2	(8.0)	17	(68.0)	6	(24.0)	0	(0.0)	
High	1	(4.5)	9	(40.9)	10	(45.5)	2	(9.1)	
HDL-C									
Low	0	(0.0)	7	(53.8)	4	(30.8)	2	(15.4)	0.392 ^a
Borderline	1	(16.7)	3	(50.0)	2	(33.3)	0	(0.0)	
Acceptable	4	(5.3)	48	(64.0)	20	(26.7)	3	(4.0)	

Table 4 Association between low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, and body mass index

Abbreviations: BMI, body mass index; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol. ^aFisher's exact test.

Table 5 Association between selected variables and hemoglobin A1c

Variables	Controlled	ontrolled Unco		Uncontrolled		
	n	(%)	n	(%)		
BMI						
Underweight	1	(9.1)	10	(90.9)	1.000 ^a	
Healthy weight	6	(8.8)	62	(91.2)		
Overweight	2	(7.4)	25	(92.6)		
Obesity	0	(0.0)	7	(100.0)		
LDL-C						
Acceptable	6	(12.5)	42	(87.5)	0.270 ^a	
Borderline	2	(8.0)	23	(92.0)		
High	0	(0.0)	22	(100.0)]	
HDL-C	-					
Low	1	(7.7)	12	(92.3)	1.000 ^a	
Borderline	0	(0.0)	6	(100.0)		
Acceptable	7	(9.3)	68	(90.7)		
Duration of T1DM	•					
< 5 y	6	(9.1)	60	(90.9)	0.647 ^a	
5–10 y	2	(5.3)	36	(94.7)		
> 10 y	1	(11.1)	8	(88.9)		

Abbreviations: BMI, body mass index; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; T1DM, type 1 diabetes mellitus. ^aFisher's exact test.

regarded as having parents may have good perception of the child's oral health rather than poor perception. This finding agrees with previous studies in which parental perception of their children's oral status was found to be superior to the clinical findings.^{58–61} In addition, parental perceptions of oral health are often dependent on clinical conditions with recognizable symptoms, such as dental caries with toothache rather than other less obvious oral conditions such as gum problems, malocclusion, or dental trauma.⁶² Similarly, in an earlier study by Cyrino et al (2011), the patient's perception of health was found to be better than the actual presentation of the disease.⁶³ The remaining 27.4% of the questionnaire

answered by 16- and17-year-olds showed slightly higher sensitivity (27.3%) and specificity (100%). More studies reported that younger individuals may not be able to identify PD.^{32,64} However, contradicting to the statement, adult age more than 60 were less likely to report gingival bleeding symptom correctly compared with less than 40 years old, which probably due to more serious manifestation occur rather than gingival bleeding.⁶⁵

According to a new classification of PD, 3.5% or four patients in this study had periodontitis.⁶⁶ All four participants answered "yes" for gum bleeding even though only three of them claimed that they have gum disease. One of them with an advanced stage

Questions	Answer	n (%)	SS	SP	ACC	PPV	NPV	AUC	p-Value
Part A 1.Do you think you have gum disease?	Yes No	14 (12.8) 95 (87.2)	16.7	90.9	54.1	64.3	52.6	0.538	0.495
2.Overall, how do you rate your teeth and gum health?	Good Bad	91 (83.5) 18 (16.5)	20.4	87.3	54.1	61.1	52.7	0.538	0.492
3.Have you ever had gum treatment for gum disease, such as scaling either above or below gum?	Yes No	19 (17.4) 90 (82.6)	22.2	87.3	55.0	63.2	53.3	0.547	0.393
4.Have you ever had loose teeth without injury?	Yes No	2 (1.8) 107 (98.2)	1.9	98.2	50.5	50.0	50.5	0.500	0.998
5.Have you ever been told by a dental professional that you have gum disease?	Yes No	1 (0.9) 108 (99.1)	0.0	98.2	49.5	0.00	50.0	0.491	0.870
6.During the past 3 mo, have you ever noticed that your gum doesn't look good?	Yes No	7 (6.4) 102 (93.6)	11.1	98.2	55.0	85.7	52.9	0.546	0.403
7.Did you use dental floss or "other devices" for tooth cleaning in the last 7 d?	Yes No	9 (8.3) 100 (91.7)	11.1	94.5	53.2	66.7	52.0	0.528	0.611
8.Did you use mouthwash or other dental rinses for "dental problems" treatment in the last 7 d?	Yes No	13 (11.9) 96 (88.1)	14.8	90.9	53.2	61.5	52.1	0.529	0.606
Part B Do you have the following symptoms?									
1. Bleeding when brushing, flossing, or eating food	Yes No	30 (27.5) 79 (72.5)	50.0	94.5	72.5	90.0	65.8	0.723	<0.001
2. Swelling, red, or painful gums for no apparent reason	Yes No	9 (8.3) 100 (91.7)	13.0	96.4	55.0	77.8	53.0	0.547	0.401
3. Teeth look longer, and the smile appears more "toothy"	Yes No	3 (2.8) 106 (97.2)	0.0	94.5	47.7	0.0	49.1	0.473	0.623
4. Bad breath/ halitosis/foul mouth odor	Yes No	11 (10.1) 98 (89.9)	14.8	94.5	55.0	72.7	53.1	0.547	0.400
5. Loosening or shifting of teeth in the affected area	Yes No	4 (3.7) 105 (96.3)	3.7	96.4	50.5	50.0	50.5	0.500	0.995
6. Pus oozing between the teeth	Yes No	0 (0.0) 109 (100.0)	0.0	100.0	50.5	0.0	50.5	0.500	1.000

Table 6 Frequency of responses for periodontal health questionnaire and diagnostic tests for each question in relation to simplified basic periodontal examination

Abbreviations: ACC, accuracy in percentage; AUC, area under the curve; NPV, negative predictive value in percentage; PPV, positive predictive value in percentage; SP, specificity in percentage; SS, sensitivity in percentage.

(stage III) of gum disease answered "yes" to people's opinions on whether they have gum disease. The individual also answered "yes" for bad gum condition, painful swollen gums, and tooth mobility. This means that, our findings agree that more value was seen pertaining to self-reported PD if severe stage periodontitis was encountered.⁶⁷ Thus, this questionnaire would be able to exclude healthy individuals from periodontal clinical examination at a cheaper cost and would be an alternative of gold standard periodontal examination in cost-limited epidemiology survey.⁶⁸ More importantly, this can expedite the early diagnosis and treatment of PD.

The HbA1c of most participants showed uncontrolled T1DM as many of them did not achieve the target HbA1c of < 53 mmol/mol (< 7.0%) for children and adolescents who have access to comprehensive care.⁴¹ In this study, almost

half of the patients were diagnosed with gingivitis (47.8%). However, there was no significant association between HbA1c and gingivitis (p = 0.271). In contrast, a significant correlation was detected between PI and sBPE (p < 0.00). This shows that periodontal condition was more associated with the presence of visible plaque rather than the underlying metabolic status. According to a recent study, HbA1c in children and adolescent has a low correlation with the gingival condition compared with adult DM patients.⁶⁹

Even though the prevalence of T1DM in children and adolescents is increasing,⁷⁰ we faced certain challenges in recruiting eligible patients during the pandemic in view of limited follow-up appointments available during the lock-down period. Furthermore, more than 10 patients or their parents declined to participate.

Variables	n	(%)	Mean	Standard deviation
Index		•	•	•
Plaque index	109	96.5	0.37	0.31
Gingival index	109	96.5	0.27	0.32
Missing	4	3.5		
Periodontal status				
Healthy	55	48.7		
Gingivitis	54	47.8		
Missing	4	3.5		
Periodontitis				
Nonperiodontitis	105	92.9		
Periodontitis	4	3.5		
Missing	4	3.5		

Table 7 Clinical parameters of the periodontal health status of participants

Conclusion

This study concluded that the GQ has a high potential to be used in identifying T1DM children at a greater risk for PD and in need of an oral examination. It can be adopted as a tool for nondental health care providers to screen for PD before making appropriate dental referrals for the patients. In the long term this will ensure a seamless and coordinated care pathway for DM patients requiring dental care

Ethical Approval Statement

Ethical approval was obtained from the Research Ethics Committee of UiTM (REC/07/2020(MR/169) and from Medical Ethics Committee, Faculty of Dentistry, UM (DF RD2018/0110 (L). UiTM Research Ethics Committee and Medical Ethics Committee, UM operates in accordance with the International Council for Harmonization Good Clinical Practice Guidelines, Malaysia Good Clinical Practice Guidelines, and the Declaration of Helsinki. Written informed consent was obtained from the parents and assent was obtained from the child for participation and for the purpose of publication. The research was performed in accordance with a named standard.

Data Availability Statement

Data available on request due to restrictions. The data presented in this study are available on request from the corresponding author. The data are not publicly available due to confidentiality issue.

Funding

This work was supported by Post Graduate Research Funding from University Teknologi MARA Sungai Buloh.

Conflict of Interest None declared.

Acknowledgment

I would like to offer my special thanks to Dr A'isyah Nabila Uzaimi, Dr Sarah Athirah Rizal, Nurse Hazean Abd Talib, Nurse Suzilah Jumati from UiTM, and Nurse Noor Azleen Ambak from UM for their assistance with the collection of my data. I would also like to thank Universiti Teknologi MARA Sungai Buloh (UiTM) for funding the study.

References

- Mattson JS, Cerutis DR. Diabetes mellitus: a review of the literature and dental implications. Compend Contin Educ Dent 2001;22 (09):757–760, 762, 764 passim, quiz 773
- 2 Liese AD, D'Agostino RB Jr, Hamman RF, et al; SEARCH for Diabetes in Youth Study Group. The burden of diabetes mellitus among US youth: prevalence estimates from the SEARCH for Diabetes in Youth Study. Pediatrics 2006;118(04):1510–1518
- ³ Harding JL, Pavkov ME, Magliano DJ, Shaw JE, Gregg EW. Global trends in diabetes complications: a review of current evidence. Diabetologia 2019;62(01):3–16
- 4 Chiang JL, Maahs DM, Garvey KC, et al. Type 1 diabetes in children and adolescents: a position statement by the American Diabetes Association. Diabetes Care 2018;41(09):2026–2044
- 5 Irfan UM, Dawson DV, Bissada NF. Epidemiology of periodontal disease: a review and clinical perspectives. J Int Acad Periodontol 2001;3(01):14–21
- 6 Kinane DF, Berglundh T, Lindhe J. Host-parasite interaction in periodontal disease. 4th ed. Clinical Periodontology and Implant Dentistry. Blackwell Munksgaard; 2003
- 7 Papapanou PN. Periodontal diseases: epidemiology. Ann Periodontol 1996;1(01):1-36
- 8 Nelson RG, Shlossman M, Budding LM, et al. Periodontal disease and NIDDM in Pima Indians. Diabetes Care 1990;13(08):836–840
- 9 Thorstensson H, Hugoson A. Periodontal disease experience in adult long-duration insulin-dependent diabetics. J Clin Periodontol 1993;20(05):352–358
- 10 Mealey BL, Ocampo GL. Diabetes mellitus and periodontal disease. Periodontol 2000 2007;44:127–153
- 11 Cianciola LJ, Park BH, Bruck E, Mosovich L, Genco RJ. Prevalence of periodontal disease in insulin-dependent diabetes mellitus (juvenile diabetes). J Am Dent Assoc 1982;104(05):653–660

- 12 Lalla E, Cheng B, Lal S, et al. Diabetes mellitus promotes periodontal destruction in children. J Clin Periodontol 2007;34(04):294–298
- 13 Preshaw PM, Bissett SM. Periodontitis and diabetes. Br Dent J 2019;227(07):577–584
- 14 Casanova L, Hughes FJ, Preshaw PM. Diabetes and periodontal disease: a two-way relationship. Br Dent J 2014;217(08):433–437
- 15 Liu R, Desta T, He H, Graves DT. Diabetes alters the response to bacteria by enhancing fibroblast apoptosis. Endocrinology 2004; 145(06):2997–3003
- 16 Liu R, Bal HS, Desta T, Behl Y, Graves DT. Tumor necrosis factoralpha mediates diabetes-enhanced apoptosis of matrix-producing cells and impairs diabetic healing. Am J Pathol 2006;168(03): 757–764
- 17 Frantzis TG, Reeve CM, Brown AL Jr. The ultrastructure of capillary basement membranes in the attached gingiva of diabetic and nondiabetic patients with periodontal disease. J Periodontol 1971;42(07):406–411
- 18 Listgarten MA, Ricker FH Jr, Laster L, Shapiro J, Cohen DW. Vascular basement lamina thickness in the normal and inflamed gingiva of diabetics and non-diabetics. J Periodontol 1974;45(09): 676–684
- 19 Mandell RL, Dirienzo J, Kent R, Joshipura K, Haber J. Microbiology of healthy and diseased periodontal sites in poorly controlled insulin dependent diabetics. J Periodontol 1992;63(04):274–279
- 20 Mashimo PA, Yamamoto Y, Slots J, Park BH, Genco RJ. The periodontal microflora of juvenile diabetics. Culture, immunofluorescence, and serum antibody studies. J Periodontol 1983;54(07):420–430
- 21 Ukai T, Ichikawa S, Sekimoto M, Shikata S, Takemura Y. Effectiveness of monthly and bimonthly follow-up of patients with wellcontrolled type 2 diabetes: a propensity score matched cohort study. BMC Endocr Disord 2019;19(01):43
- 22 Fisher E, Lazar L, Shalitin S, et al. Association between glycemic control and clinic attendance in emerging adults with type 1 diabetes: a tertiary center experience. J Diabetes Res 2018; 2018:9572817
- 23 Smith A, Harris C. Type 1 diabetes: management strategies. Am Fam Physician 2018;98(03):154–162
- 24 Babatzia A, Papaioannou W, Stavropoulou A, et al. Clinical and microbial oral health status in children and adolescents with type 1 diabetes mellitus. Int Dent J 2020;70(02):136–144
- 25 Siddiqi A, Zafar S. The two-way link between diabetes mellitus and periodontal disease: medical-healthcare professionals' clinical practice. Biomed J Sci Tech Res 2022;44:32793
- 26 Moore J, Csikar J, Kang J, Tugnait A, Campbell F, Clerehugh V. Awareness, practices, training, and confidence of paediatric diabetes care teams in relation to periodontitis. Pediatr Diabetes 2020;21(02):384–389
- 27 Matrooshi KA, Raeesi SA, Tawfik AR, et al. Knowledge of physicians about the interrelationship between diabetes mellitus and periodontitis in the United Arab Emirates. Eur J Dent 2023;17 (01):219–226
- 28 Borgnakke WS. IDF diabetes atlas: diabetes and oral health a two-way relationship of clinical importance. Diabetes Res Clin Pract 2019;157:107839
- 29 Myers-Wright N, Cheng B, Tafreshi SN, Lamster IB. A simple selfreport health assessment questionnaire to identify oral diseases. Int Dent J 2018;68(06):428–432
- 30 Mohd-Said S, Mohd-Norwir NA, Ariffin AN, et al. Validation of a simplified digital periodontal health screening module for general dental practitioners. Healthcare (Basel) 2022;10(10): 1916
- 31 Montero E, La Rosa M, Montanya E, et al. Validation of selfreported measures of periodontitis in a Spanish population. J Periodontal Res 2020;55(03):400–409
- 32 Perdoncini NN, Furquim CP, Bonfim CMS, Soares GMS, Torres-Pereira CC. Self-perception of periodontal health status among individuals with Fanconi anemia. Hematol Transfus Cell Ther 2021;43(04):453–458

- 33 Abbood HM, Hinz J, Cherukara G, Macfarlane TV. Validity of selfreported periodontal disease: a systematic review and metaanalysis. J Periodontol 2016;87(12):1474–1483
- 34 Fuziah MZ, Hong JY, Zanariah H, et al. A national database on children and adolescent with diabetes (e-DiCARE): results from April 2006 to June 2007. Med J Malaysia 2008;63(Suppl C):37–40
- 35 O'Neil KJ, Jonnalagadda SS, Hopkins BL, Kicklighter JR. Quality of life and diabetes knowledge of young persons with type 1 diabetes: influence of treatment modalities and demographics. J Am Diet Assoc 2005;105(01):85–91
- 36 CDC. Individual growth charts. Accessed May 5, 2022 at: http:// www.cdc.gov/growthcharts/charts.htm
- 37 Blood pressure levels for boys and girls by age and height percentile. Accessed May 5, 2022, at: https://www.nhlbi.nih.gov/ files/docs/guidelines/child_tbl.pdf
- 38 Flynn JT, Kaelber DC, Baker-Smith CM, et al; Subcommittee on screening and management of high blood pressure in children. Clinical practice guideline for screening and management of high blood pressure in children and adolescents. Pediatrics 2017;140 (03):e20171904
- 39 Kim G, DeSalvo D, Guffey D, et al. Dyslipidemia in adolescents and young adults with type 1 and type 2 diabetes: a retrospective analysis. Int J Pediatr Endocrinol 2020;2020:11
- 40 Janet MExpert Panel on Integrated Guidelines for Cardiovascular Health and Risk Reduction in Children and Adolescents National Heart, Lung, and Blood Institute. Expert panel on integrated guidelines for cardiovascular health and risk reduction in children and adolescents: summary report. Pediatrics 2011;128(Suppl 5, Suppl 5):S213–S256
- 41 DiMeglio LA, Acerini CL, Codner E, et al. ISPAD Clinical Practice Consensus Guidelines 2018: glycemic control targets and glucose monitoring for children, adolescents, and young adults with diabetes. Pediatr Diabetes 2018;19(Suppl 27):105–114
- 42 Han TM, Ismail R, Yaacob M, et al. Validation of self-reported questionnaires to screen periodontal disease among diabetes patients: a study at selected health centers in Kuantan. Malays Fam Physician 2016;11(01):9
- 43 Han TM Effectiveness of medical-dental coordinated care for diabetes patients with periodontal disease at selected public primary care clinic in Kuantan, Pahang [Doctoral Dissertation]. 2018
- 44 Clerehugh V, Kindelan S. Guideline for periodontal screening and management of children and adolescent under 18 years of age. British Society of Periodontology and The British Society of Paediatric Dentistry. 2021
- 45 Rapp GE, Garcia RV, Motta AC, Andrade IT, Bião MA, Carvalho PB. Prevalence assessment of periodontal disease in 3-6 year old children through PSR-a pilot study. J Int Acad Periodontol 2001;3 (03):75–80
- 46 Silness J, Loe H. Periodontal disease in pregnancy. Ii. Correlation between oral hygiene and periodontal condition. Acta Odontol Scand 1964;22:121–135
- 47 Loe H, Silness J. Periodontal disease in pregnancy. I. Prevalence and severity. Acta Odontol Scand 1963;21:533–551
- 48 Löe H. Periodontal disease. The sixth complication of diabetes mellitus. Diabetes Care 1993;16(01):329–334
- 49 Mealey BL. Periodontal disease and diabetes. A two-way street. J Am Dent Assoc 2006;137(Suppl):26S-31S
- 50 Zainal Abidin Z, Zainuren ZA, Noor E, Mohd Nor NS, Mohd Saffian S, Abdul Halim R. Periodontal health status of children and adolescents with diabetes mellitus: a systematic review and meta-analysis. Aust Dent J 2021;66(Suppl 1):S15–S26
- 51 Iwasaki M, Usui M, Ariyoshi W, et al. Validation of a self-report questionnaire for periodontitis in a Japanese population. Sci Rep 2021;11(01):15078
- 52 Kallio P, Nordblad A, Croucher R, Ainamo J. Self-reported gingivitis and bleeding gums among adolescents in Helsinki. Community Dent Oral Epidemiol 1994;22(5 Pt 1):277–282

- 53 Goulão B, MacLennan GS, Ramsay CR. Have you had bleeding from your gums? Self-report to identify gingival inflammation (The SING diagnostic accuracy and diagnostic model development study). J Clin Periodontol 2021;48(07):919–928
- 54 Nadanovsky P, Dos Santos APP, Bloch KV. Prevalence of selfreported gingival bleeding in a representative sample of the Brazilian adolescent population. J Clin Periodontol 2018;45(08): 952–958
- 55 Elhassan A, Alfakry H, Peeran S. Reasons to seek periodontal treatment in a Libyan community. Dent Med Res 2017;5(02):38–42
- 56 Nelson DE, Holtzman D, Bolen J, Stanwyck CA, Mack KA. Reliability and validity of measures from the Behavioral Risk Factor Surveillance System (BRFSS). Soz Praventivmed 2001;46(Suppl 1):S3–S42
- 57 Reiniger APP, Londero AB, Ferreira TGM, da Rocha JM, Moreira CHC, Kantorski KZ. Validity of self-reported measures for periodontitis surveillance in a rural sample. J Periodontol 2020;91(05):617–627
- 58 Al-Batayneh OB, Al-Khateeb HO, Ibrahim WM, Khader YS. Parental knowledge and acceptance of different treatment options for primary teeth provided by dental practitioners. Front Public Health 2019;7:322
- 59 Hamasha A, Rasheed S, Aldosari M, Rajion Z. Parents knowledge and awareness of their children's oral health in Riyadh, Saudi Arabia. Open Dent J 2019;13:236–241
- 60 Butani Y, Gansky SA, Weintraub JA. Parental perception of oral health status of children in mainstream and special education classrooms. Spec Care Dentist 2009;29(04):156–162
- 61 Daly JM, Levy SM, Xu Y, et al. Changes in parental perceptions of their care of their children's oral health from age 1 to 4 years. J Prim Care Community Health 2019;10:2150132719836908
- 62 Gomes MC, Clementino MA, Pinto-Sarmento TC, et al. Parental perceptions of oral health status in preschool children and associated factors. Braz Dent J 2015;26(04):428–434

- 63 Cyrino RM, Miranda Cota LO, Pereira Lages EJ, Bastos Lages EM, Costa FO. Evaluation of self-reported measures for prediction of periodontitis in a sample of Brazilians. J Periodontol 2011;82(12): 1693–1704
- 64 Vered Y, Sgan-Cohen HD. Self-perceived and clinically diagnosed dental and periodontal health status among young adults and their implications for epidemiological surveys. BMC Oral Health 2003;3(01):3
- 65 Romano F, Perotto S, Bianco L, Parducci F, Mariani GM, Aimetti M. Self-perception of periodontal health and associated factors: a cross-sectional population-based study. Int J Environ Res Public Health 2020;17(08):2758
- 66 Sanz M, Tonetti MS. New classification of periodontal and peri implant disease. Eur Federation Periodontol 2019. Accessed May 5, 2022 at: https://www.efp.org/fileadmin/uploads/efp/Documents/ Campaigns/New_Classification/Guidance_Notes/report-02.pdf
- 67 Nguyen VTN, Furuta M, Zaitsu T, et al. Periodontal health predicts self-rated general health: a time-lagged cohort study. Community Dent Oral Epidemiol 2022;50(05):421–429
- 68 Chatzopoulos GS, Tsalikis L, Konstantinidis A, Kotsakis GA. A twodomain self-report measure of periodontal disease has good accuracy for periodontitis screening in dental school outpatients. J Periodontol 2016;87(10):1165–1173
- 69 Lipski J, Burchardt D, Duda-Sobczak A, Wyganowska M. The assessment of the relationship between the severity of gingivitis and the glycosylated hemoglobin levels in adolescent and adult patients with type 1 diabetes. Postepy Hig Med Dosw 2021; 75:868–872
- 70 Mobasseri M, Shirmohammadi M, Amiri T, Vahed N, Hosseini Fard H, Ghojazadeh M. Prevalence and incidence of type 1 diabetes in the world: a systematic review and meta-analysis. Health Promot Perspect 2020;10(02):98–115