Université de Montréal

Development and Evaluation of a Research-based Prosthodontic Clinical Record

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Mémoire présenté à la Faculté des études supérieures en vue de l'obtention du grade de Maîtrise ès Sciences (M.Sc.)
En sciences bucco-dentaires

Avril 2012

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Université de Montréal Faculté des études supérieures

Ce mémoire intitulé:

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RÉSUMÉ

Introduction: Bien que l'importance de transférer les données de la recherche à la pratique a été largement démontrée, ce processus est toujours lent et fait face à plusieurs défis tels que la conceptualisation des évidences, la validité interne et externe de la recherche scientifique et les coûts élevés de la collecte de grandes quantités de données axées sur le patient. Les dossiers dentaires des patients contiennent des renseignements valables qui donneraient aux chercheurs cliniques une opportunité d'utiliser un large éventail d'informations quantitatives ou qualitatives. La standardisation du dossier clinique permettrait d'échanger et de réutiliser des données dans différents domaines de recherche.

<u>Objectifs:</u> Le but de cette étude était de concevoir un dossier patient axé sur la recherche dans le domaine de la prosthodontie amovible à la clinique de premier cycle de l'Université de Montréal.

Méthodes: Cette étude a utilisé des méthodes de recherche-action avec 4 étapes séquentielles: l'identification des problèmes, la collecte et l'interprétation des données, la planification et l'évaluation de l'action. Les participants de l'étude (n=14) incluaient des professeurs, des chercheurs cliniques et des instructeurs cliniques dans le domaine de la prosthodontie amovible. La collecte des données a été menée à l'aide d'une revue de littérature ciblée et complète sur les résultats en prosthodontie ainsi que par le biais de discussions de groupes et d'entrevues. Les données qualitatives ont été analysées en utilisant QDA Miner 3.2.3.

Résultats: Les participants de l'étude ont soulevé plusieurs points absents au formulaire actuel de prosthodontie à la clinique de premier cycle. Ils ont partagé leurs idées pour la conception d'un nouveau dossier-patient basé sur 3 objectifs principaux: les objectifs cliniques, éducatifs et de recherche. Les principaux sujets d'intérêt en prosthodontie amovibles, les instruments appropriés ainsi que les paramètres cliniques ont été sélectionnés par le groupe de recherche. Ces résultats ont été intégrés dans un nouveau formulaire basé sur cette consultation. La pertinence du nouveau formulaire a été évaluée

par le même groupe d'experts et les modifications requises ont été effectuées. Les participants de l'étude ont convenu que le cycle de recherche-action doit être poursuivi afin d'évaluer la faisabilité d'implémentation de ce dossier modifié dans un cadre universitaire.

<u>Conclusion:</u> Cette étude est une première étape pour développer une base de données dans le domaine de la prothodontie amovible. La recherche-action est une méthode de recherche utile dans ce processus, et les éducateurs académiques sont bien placés pour mener ce type de recherche.

<u>Mots-clés:</u> Recherche-action, Prosthodontie, Prothèses amovibles, La médecine dentaire fondée sur des données probantes

ABSTRACT

<u>Introduction:</u> Although the importance of research translating into practice has been widely recognized, this process is still slow and faces several barriers such as conceptualizations of evidence, internal and external validity of the evidence and high costs of providing large amounts of patient-based outcome data. Patient's dental records contain valuable information that would give clinical researchers an opportunity to use a wide range of quantitative or qualitative information. Standardization of clinical record would allow the interoperability and reusability of data in different research fields.

Objectives: The aim of this study was to design a research-based patient record in the field of removable prosthodontics in the undergraduate clinic of the "Université de Montréal."

<u>Methods:</u> This study used action research methods with 4 sequential steps: problem identification, gathering and interpreting data, action planning, and action evaluation. Study participants included professors, clinical researchers, and clinical instructors in the field of removable prosthodontics. Data collection consisted of a comprehensive literature review on prosthodontic outcomes as well as focus-group discussions and interviews. The qualitative data were analysed using QDA Miner 3.2.3.

Results: The study participants raised several concerns about the deficiencies of the existing patients' prosthodontic record in the undergraduate clinic. They shared their ideas for designing a new patient record based on 3 key objectives: clinical, educational, and research objectives. The prosthodontic outcomes of interest and appropriate instruments as well as the clinical parameters were selected by the research group and were integrated into a new research-based record. The appropriateness of the new record has been evaluated by the same panel of experts and the necessary modifications have been carried out. The study participants agreed that the action research cycle should be continued to evaluate the feasibility of the implementation of this redesigned record in the university-based setting.

<u>Conclusion:</u> This study is a beginning effort to develop a database in the field of removable prosthodontics. Action research is a useful research method in this process, and academic educators are well placed to conduct such research.

<u>Keywords:</u> Action research, Prosthodontics, Removable prosthesis, Evidence-based dentistry

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LIST OF ABBREVIATIONS

WHO GODB World health organization global oral data bank

COHRI Consortium for oral health-related informatics

HMDB Hospital morbidity database

GPRD General practice research database

CCPDS Centralized cancer patient data system

SEER Surveillance, epidemiology and end results

POMR Problem-oriented medical record

SOAP Subjective, objective, assessment, and plan

CDMS Clinical data management system

JCR Journal citation reports

DEDICATION

To my dear mother who trained me love of life,

To my lovely father who taught me love of knowledge,

To my dear husband who supports me to courageously face disappointments.

ACKNOWLEDGMENTS

In a journey of this duration there are many people to acknowledge. First, I wish to acknowledge my supervisor, Dr. Elham Emami who was so helpful and supportive during my master program. I wish that this master thesis can make her proud. Secondly, I would like to thank Dr. Claude Lamarche for his thoughtful comments and timely feedback.

As well, I would like to thank Dr. Pierre de Grandmont, Dr. Gilles Gauthier, and Dr. Louis de Koninck for their contribution and help in this study. Of equal importance, I would like to thank the interviewed individuals in this project. This study will never be performed without their participation and help. I would also like to thank Dr. Robert Durand for his helpful comments throughout the project.

Finally, I am thankful to my husband for all his support. Without his love, patience, and encouragement, this dissertation would not exist. Thank you, Farjad!

CHAPTER I

LITERATURE REVIEW

1.1. INTRODUCTION

Research in oral health or any health-related disciplines requires access to several sources of health information including dental/medical records, hospital record databases, epidemiological databases, disease and vital registration systems, and health statistics ^{1, 2}. Consequently, standardized databases are needed for the interoperability and reusability of data in different research fields such as epidemiology, clinical research, and oral health-care services research ³⁻⁶. In fact the use of databases is a cost-effective and methodologically sound approach that will facilitate conducting clinical and comparative effectiveness research ⁷. For example, clinical researchers could identify the most effective and most efficient interventions, treatments, and services by having access to information from patient clinical records/charts, and tissue or data repositories ^{6, 8, 9}. Health services researchers could assess the quality of health-care services using large databases of health-care information made available by health-care providers, institutions, and governmental agencies.

The collection of health-related data by academic institutions, university hospitals, and clinics in a systemic and standardized way and establishment of a university-based dataset would be a major advantage for any institution as it would enable the recording of large amounts of information across a wide range of diagnoses, treatment plans, interventions, and outcomes ^{8, 9}. To our knowledge, such a research-based recording system in dental academic institutions is still unexplored. This is mainly due to the extensive planning phase, high cost, and lack of the availability of clinical scientists. Therefore, innovative approaches are needed to develop comprehensive clinical recording systems to provide data and to support clinical and oral health-care services research. This task needs the collaboration of both researchers and clinicians in a way that integrates clinical care and

clinical research. This chapter consists of a review of the literature offering background knowledge on this topic.

1.2. CLINICAL DATABASE

1.2.1. Definition

A clinical database is an accurate dataset concerning clinical practice, recorded in an organized way and connected to outcome descriptors ¹⁰. The concept of clinical database has been used to remove the difficulties in creating, designing, and keeping complex information systems for varying amounts of data ⁹. It allows retrieving and accessing organized information about patients' histories and clinical findings ¹¹⁻¹³.

Databases are composed of fields, records, and files. A field can be defined as a single piece of information; a record is defined as a set of fields; and a file is defined as a group of records ¹⁴.

1.2.2. Characteristics

In all databases, there is information that is collected and stored as data elements. This information is retrieved from admission forms, history sheets, and reports of laboratory results, operations, and consultations ^{13, 15}. In general, clinical information can be collected in one of two ways ¹⁰:

- 1) As a part of the patient care procedure
- 2) As a separate information file to be entered into the database

Collection of clinical information in the first way has advantages over the second way. Firstly, the data collection is prospective. Secondly, the quality control of collected information can be improved by using quality control measures as a part of the patient care process. Finally, the cost of data gathering can be decreased by imposing the financial duty on the patient care ¹⁰. Clinical data collection should be well organized, easy to access, and

should not interfere with the patient care process ^{4, 12, 16}. Since these clinical data are important for health-care decision making, they should be recorded precisely ^{7, 17}.

Clinical databases can be in paper form or electronic form. Although paper-based records are the most used method of recording patient information in hospitals and health centers, electronic-based records are more advantageous ¹⁸. This format of clinical records is more efficient than the paper format because it ensures that the same information is not recorded on multiple occasions. In addition, such records can be shared more quickly across various health-care settings ¹⁹⁻²¹. Due to an increase in the amount of data, health care organizations increasingly want to replace their manual systems with reliable electronic systems to reduce the possibility of human error ²².

Data element collection depends mainly on the function of the database ¹⁰. As an example, research databases contain descriptors that are helpful for examining research hypotheses. Similarly, databases that are designed for administrative objectives comprise information that affects administrative decisions ¹⁰.

Each clinical database has a distinctive focus that can be as narrow as a specific therapy or as broad as a whole medical/dental record. Clinical database focus is usually one of the following two types $^{10, 23}$:

- 1) Disease or population specific; or
- 2) Procedure, intervention, or health technology specific.

In general, in a successful clinical database there exists a balance between the focus and function of the database. Although this concern seems clear, many clinical databases have imprecise focus, and consequently their function changes ¹⁰. During the database collection procedure, considerations include what data is collected, how it is recorded, and for how long it is registered ^{24, 25}. Often gathering too much or too little data hinders achieving the long-term goals of the database.

The function of a database determines the requirements of the data retrieval process ¹⁰. The retrieval process should be simple and should enable the user to have access to information in a suitable format.

1.2.3. Objectives

1.2.3.1. Objectives in Patient Care

The patient care process requires using clinical databases in different ways. Clinical databases assist health-care professionals to document observations, diagnostic decisions and health-care treatments, and to exchange information. Also, these databases help physicians to better understand diseases and treatment procedures. Furthermore, clinical databases can be used to improve diagnostic processes. Finally, they are a functional tool to improve the outcome of the delivered patient care by predicting expected outcomes with alternative therapies ^{7, 12, 17, 24, 26}.

Clinical databases can gather information in response to patient care needs in different forms including patient-oriented analyzing and patient-group reporting. The main function of patient-oriented analyzing is creating information related to health care, concerning each individual patient. The management of the patient's health-care process (including planning and controlling), the prognosis procedure, and the critical review of a completed treatment are the principal aims of patient-oriented analyzing ^{27, 28}.

In patient-oriented analyzing, there may be problems due to inadequate clarity of data presentation and incomplete collected data. For this reason, data recording has to be complete, accessible, and unambiguously connected to a single patient.

The purpose of patient-group reporting is creating helpful information about a predefined group of patients. These data consist of measures describing quantitative attributes (e.g., the duration of a treatment in terms of the mean, the standard deviation, etc.) or the frequency of certain conditions (difficulties, diagnoses, etc.). The other typical

aim of patient- group reporting is recording essential information on cost benefit structures and on details of the institution's work processes. In addition, identifying problems and reviewing intervention outcomes require monitoring the quality indicators of the institution's work processes. Problems often occur in patient-group reporting due to incomplete data recording, which leads to the cost and quality of patient care not being accurately reported. Therefore, patient-group reports must be precise and comprehensible based on reliable and valid collected data ^{27, 28}.

1.2.3.2. Objectives in Education

A clinical database can be a helpful tool for the education and training of health-care professionals. It provides a useful tool for evaluation of students' actions. Furthermore, it has a key role in providing an excellent example of clinical problems and an explanation of courses of diseases ^{8, 26}.

1.2.3.3. Objectives in Research

Clinical databases provide enormous amounts of patient data. These data are useful resources for different types of studies including observational and epidemiological studies, as well as clinical trials ^{1, 8, 29}. Clinical databases help observational studies by providing organized information on a large sample of patients.

Epidemiological studies are conducted in specified populations by studying the distribution and determinants of health-related states or events ^{26, 30}. Precise clinical databases relevant to the diagnosis and demographic characteristics of patients can be used to determine incidence and prevalence rates, risk factors, and chronological trends ^{23, 26, 31, 32}.

In clinical research, the role of a clinical database is to gather all relevant information with the goal of improving health. It can also be used to evaluate new diagnostic or therapeutic procedures. In addition, it can be a useful resource for planning an

original study based on previous observations. Finally, new scientific insights can be obtained through the analysis of collected data ³³.

In general, clinical databases contribute in expanding clinical research in three different ways ^{1, 4, 8, 10, 26, 34-36}:

- 1. They create precise information for the evaluation of the course of diseases, with the purpose of detecting the starting points for generalization.
- 2. The use of clinical databases during the preparation of scientific clinical studies may facilitate the selection of patients with defined characteristics (e.g., all female patients with burning mouth syndrome) for a specific study. This selection, in turn, forms the basis for a scientific study that must be designed and documented separately.
- 3. For research studies, clinical databases can provide requested data for each patient participating in the study.

Furthermore, clinical databases contribute to the quality of clinical trials by ensuring the continuity and consistency of observations made by different examiners over the years 10, 37, 38.

In health outcomes research, clinical databases help to provide data to cover themes of outcome studies such as safety, effectiveness, efficiency, and timeline ^{39, 40}. Over the last several decades, medical research and technology have been improved to prevent, to diagnose, and to treat diseases, while questions are increasingly being asked about these technologies and their effectiveness in clinical practice ³⁹⁻⁴¹. The main goal of health outcome research is to improve care and to prevent diseases. It plays an important role in shaping healthcare decisions and policies ⁴¹. The main problem in conducting health outcome research is unorganized databases and unknown data quality. An organized clinical database can provide scientific evidence related to the decisions taken by all who participate in patient health care ³⁹⁻⁴¹.

1.2.3.4. Objectives in Administration

Clinical databases play an important administrative role in health-care institutions. Administrative uses of clinical databases have an effect on all phases of health-care delivery ²⁶. They allow health-care institutions to select and design efficient work processes as well as to receive the correct reimbursement for their services. Clinical databases are used to generate reports and to improve office management ⁴². In addition, in the event of legal proceedings, adequate medical documentation in clinical databases can have positive implications for health-care institutions ²⁶.

1.2.3.5. Objectives in Health Care Quality Management

Clinical databases provide suitable information for medical audit and for systematic quality monitoring ^{8, 9, 17, 43}. They have been frequently used in health technology assessment. In other words, clinical databases are essential elements to support clinical and health-care policy decisions ^{10, 26, 44-47}.

1.2.4. Examples of Successful Databases

A number of oral health and dental databases as well as several medical databases have been developed and have been successfully implemented in different organizations and institutions.

The World Health Organization Global Oral Data Bank (WHO GODB) was established in 1969 to fill the gap of data on oral health status and oral disease process. This oral health information system is categorized into the following interrelated subsystems ^{48,} ^{49,}

- Epidemiological surveillance
- Service coverage of the population
- Service records and reporting
- Administration and resource management
- Quality of care

Oral health program monitoring and outcome evaluation

The GODB databank has met all the requirements for the collection of complete and precise oral health information ^{48, 50}. In addition, the GODB gathers relevant information from sources that are not in the scientific literature, e.g., reports prepared for the ministries of health ⁴⁹. Over the years, this databank has been used as the main international reference for global oral health epidemiology. Currently, the WHO GODB includes 1,850 data sets on dental caries from 178 countries. Standard criteria and methodology are checked by WHO for each dataset before it is accepted into the GODB. Implementation of preventive oral care strategies and programs will be improved by the collected data in the GODB ^{48, 49}.

The Consortium for Oral Health-Related Informatics (COHRI) was created in 2007 to establish an oral health data repository that can accept and integrate data from different dental data sources ⁵¹. The COHRI will support users' research or decision-making needs by allowing exploring and extracting information. This database is the result of collaboration between 20 dental schools using the same Electronic Health Record platform ⁵¹. The COHRI will provide useful data for such diverse clinical research studies as randomized prospective clinical trials, retrospective case control studies, cross sectional studies, and cohort studies. It will help outcome assessment for patient care, and will allow exploring the relationships between oral health and systemic diseases. It also allows measuring student clinical performance, determining the accuracy of treatment planning, and validating educational outcomes.

The Hospital Morbidity Database (HMDB) is a database conducted by the Canadian Institute for Health Information since 1960 ⁵². The HMDB contains administrative data elements, clinical data elements, and demographic data elements. The aim of this databank is recording, processing, and analyzing diagnoses and procedures for all hospital separations (number of discharges and deaths). In addition, it collects data for federal agencies such as Statistics Canada.

The General Practice Research Database (GPRD) is the biggest computerized database in the world, consisting of anonymous patient data. Data in the GPRD have been collected by contributing general practices throughout the United Kingdom continuously since 1987. To date, it has collected information on approximately 3 million patients ⁵³. Information collected from GPs includes: demographic data, medical diagnosis, treatment outcomes, miscellaneous patient care information (e.g., smoking status, height), and laboratory results. As the GPRD contains demographic and clinical details of patients, it can be used as a functional resource to conduct pediatric drug utilization and pharmacovigilance studies as well as to study rare adverse events ^{53, 54}.

Another example of a clinical database is the Duke University Medical Center Databank that was created in 1969 in order to improve health care for patients with cardiovascular diseases. The collected information in the Duke databank focuses on therapies or specialized care for these specific patients. In this university medical center databank, the patient care procedures are followed prospectively and so they can be linked to long-term outcomes. Using the Duke databank, physicians can learn from previous experience, resulting in improvement of patient care. Furthermore, this databank is recognized as a useful resource for research studies and administrative functions ^{10, 55-59}.

The Centralized Cancer Patient Data System (CCPDS) and Surveillance, Epidemiology and End Results (SEER) are two more examples of successful clinical databases in the United States ^{10, 60-63}. The CCPDS and SEER are used to collect high-quality data on cancer cases in American hospitals. Data for SEER and CCPDS have been collected since 1973 and 1977, respectively.

1.2.5. High-Quality Clinical Databases

The need for development of high-quality clinical databases for use as a reliable resource in clinical practice, evaluation research, clinical audit, and evaluation health

technologies is well recognized ⁶⁴. The quality of data is a main concern for each database user and determines the value of the clinical database ^{42, 43, 65-68}.

Data quality is determined by a great number of diverse attributes ^{42, 69}. However, the definitions of these data quality attributes are often unclear or unavailable ⁶⁷. According to the literature ⁶⁸⁻⁷², the most important cited data quality attributes are "accuracy" and "completeness." Data accuracy is the amount of conformity to the truth of the data recorded in the database. Data completeness is the extent to which all required data are registered ⁴².

Wyatt and Sullivan (2005) proposed seven criteria for evaluating the quality of information of a clinical database ⁷³ (Table 1.1). Black and Payne (2003)presented quality assessment criteria based on database coverage and accuracy ⁴³. According to these criteria, the assessment of database coverage should be based on: representativeness of the registered population, completeness of the recruitment of each eligible individual, completeness of the collected data for each individual, and finally collection of all the necessary variables in the database ^{9, 43}. Database accuracy is evaluated by the following criteria: clear and unambiguous definition of data variables, standardization in data collection and data coding, and independent observation of the outcomes ^{9, 42, 43}. The completeness and accuracy of clinical databases can be increased by developing organized and structured clinical documentation ^{8, 9}.

Table 1.1: Quality criteria for patient data (adapted with permission from Wyatt and Sullivan 73)

Criterion	How to test it?
Accurate	Comparison with a gold standard source of data.
Complete	Percent of missing data at a given point.
Timely	Delay from the event the data describes to its availability for use on the information system.
Relevant	Amount that data alter decisions or actions of the user; the impact of leaving an item out of the dataset.
Appropriately represented	Degree of structuring and coding of items.
Relevant detail included	If data are detailed enough to support decisions.
Relevant context included	Is there enough context to support appropriate interpretation of data?

1.3. CLINICAL DOCUMENTATION: HEALTH-CARE PATIENT RECORD

1.3.1. Definition and Content

The patient record is a collection of information concerning patients and their health care, gathered during the patient's health-care procedure. It includes a variety of important elements such as patient history, administration of drugs and therapies, test results, the details of clinical findings for each patient, and summarized reports ^{8, 12, 17, 24}. The data retrieved directly from patients and from clinical records and laboratory results may be grouped as patient findings ²⁴. The essential requirement of health-care providers is maintaining the complete and accurate medical record, which is generally considered as a certification prerequisite as well. The patient record provides data for continuity of care and is crucial in the occurrence of a malpractice insurance claim ^{74, 75}. According to Wyatt (1994) there are four main categories in each health-care record ²⁴ (Figure 1.1).

- 1) Identifiers: Contain identity numbers, sociodemographic data, and supplementary information if necessary, including family doctor, health insurance information, etc.
- 2) Patient findings: Include observations and history data, which can be subjective data (such as symptoms) and objective data (such as clinical signs, X-ray results, and laboratory results).
- 3) Hypotheses: Include assessment and plans like diagnosis, problem list, and possible explanations.
- 4) Actions: Contain therapy and follow-up.

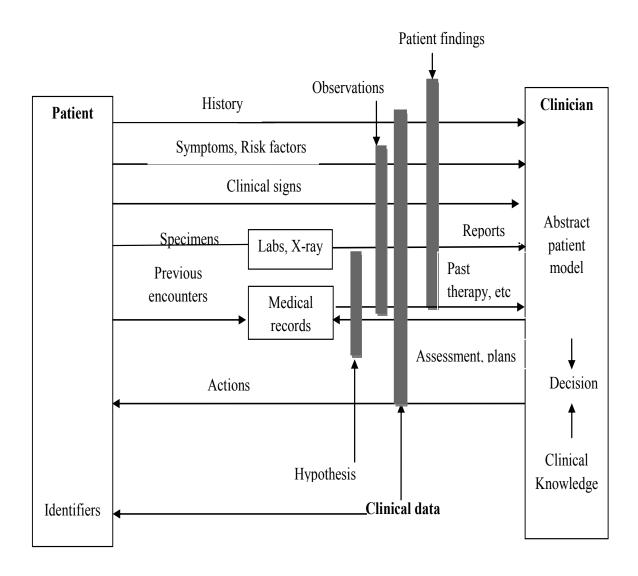


Figure 1.1: Categories of clinical data (adapted with permission from Wyatt ²⁴)

1.3.2. Functions

Recording what happened between physician and patient in a reliable, legible, and objective manner is the principal function of a patient health-care record ⁷⁶. Medical records maintain the continuity of care by documenting the data required to follow up the patient care for the treating clinician or other health-care providers ^{76, 77}. Clinical patient records are used not only for supporting patient care, but also for secondary clinical purposes including clinical audit, research, epidemiology, and resource allocation ^{4, 15} (Figure 1.2). Additionally, the patient care record could become a legal document and could be used in litigation ^{76, 77}.

1.3.3. Standardization

The most important advantage of standardized patient records is the ability to gather enormous amounts of organized information containing diagnosis and interventions, which creates a useful source for analysis ^{8,9}.

Development and implementation of evidence-based standards in clinical records is mandatory because:

- Good records monitor patients' health status and health care ⁷⁸
- Accurate record-keeping guarantees a suitable and systematic flow of the treatment plan/s
- Clinical records are reliable tools which systematize all important parameters and aspects of short- and long-term results ^{15, 79-81}
- High quality patient records influence the effectiveness of care ⁷⁸

Furthermore, standards are necessary for sharing and reusability of data in clinical research including mechanisms of human disease, epidemiology, behavior studies, outcomes, and health services research ⁴.

Richmond et al. (2007) examined two hundred dental records of edentulous patients attending the University of Manchester School of Dentistry ⁸². They found that only 67.8%

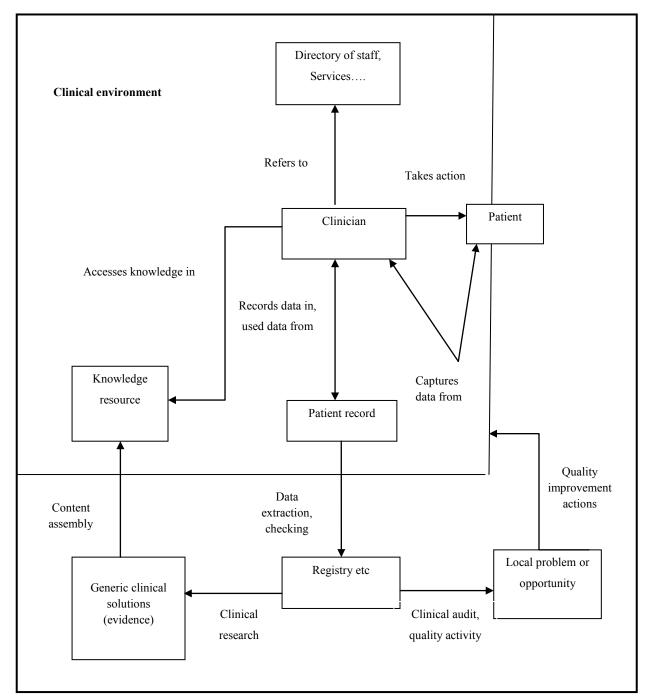


Figure 1.2: Information flows in clinical and non-clinical environment (adapted with permission from Wyatt and Sullivan ⁷³)

of the written records were rated as good. Helminen et al. (1998) assessed the quality of oral health record-keeping in public oral health-care services in Finland ⁸³. They reported that the notes on the patients' occlusion and temporomandibular joint function were omitted in 63% of cases and that 89% of dental records contained missing information about oral soft tissues. An audit of 184 Worcestershire general dental practitioners revealed a wide variation between dentists' clinical record-keeping ⁸⁴.

Martin Garcia et al. (2008) analyzed dental records in two steps. First, they used 46 criteria to evaluate quality of 50 dental records ⁸⁵. The results showed that no record was without error. Then, in order to correct the deficiencies detected in the first evaluation, they produced a new model of dental record which was implemented for two years. Re-audit showed that appropriate fulfilment was reached in 29 criteria while in the first phase the standards had been reached in only 12 criteria ⁸¹. In another research study, Osborn et al. (2000) found a noticeable discrepancy between the American Dental Association guidelines and dentists' perception of dental record adequacy ⁸⁶.

A retrospective audit of 316 dental records was conducted in 13 selected institutional providers of dental care in New York City in order to assess the level of documentation presented in the dental record ⁶⁴. At the time of the initial audit, more than 50% of these institutions were unable to present all the data requested. Few of the audited dental records were without deficiencies. Also, in most institutions the documented result of the intra-oral examination was lacking. A plan to correct the identified deficiencies was implemented. The follow-up audit showed significant improvement in the level of recorded information.

In fact, standards have a key role in improving the effectiveness and quality of clinical documentation. Implementation of standardized clinical documentation will minimize difficulties encountered in data extraction. Evaluation of available standards in clinical documentation is the basic requirement for developing an optimum database ^{13, 15, 87, 88}

According to Weed (1968), the father of the problem-oriented medical record, one example of well-organized record is the "SOAP" model ⁸⁹. The SOAP format derives from the problem-oriented medical record (POMR) as a guide for health-care providers to collect and organize the patient data ⁹⁰. The POMR actually focuses on the idea that the most useful way of organizing clinical data is through the concept of patients' problems and their evolution ⁹¹. Consequently, decisions taken during patient care in POMR are related to those problems. In this way, health-care providers can have access to the patients' problems, the frequency of the problems, and the received treatments. The POMR provides a functional patient form with standardized data that can adapt without difficulty to new information technologies. Additionally, easy accessibility of the standardized collected data in the POMR encourages continuing assessment of the health-care plan ⁸⁹⁻⁹¹.

Using the "SOAP" note, the basic POMR was developed ⁸⁷. The SOAP format can make a structure for documenting clear and concise data to improve communication among health-care professionals, as well as to improve recording of the patient's concerns and health issues ⁹⁰. The SOAP format includes ⁸⁹⁻⁹⁴:

- S: Subjective, is the patient's perception and outcome. Items to be included in this
 section are patients' concerns, any allergies, medical history, and complete patient
 history.
- O: Objective, is the observation of the clinician. Information in this category includes observations of patient clinical examinations, and laboratory and X-ray test results.

- A: Assessment, is the diagnosis, measurement, and evaluation. This part is the conclusion reached based on the subjective and objective sections, which should be complete in order to dictate the plan.
- P: Procedures and plan, are the treatment plans, treatment received, and follow-up/s. This part is divided into three sections. One section is all the medications, or devices recommended by the physician for the patient. The next section is a list of the expected results of the considered diagnostics. The third section is used for recording all referrals or consultations.

Documentation without a standardized structure may be time consuming to complete and confusing.

1.4. CLINICAL DATA MANAGEMENT SYSTEM

A clinical data management system or CDMS is a generic name for a tool or set of tools to manage and organize the data in a computer ^{33, 95}. The most important task of CDMS is collecting and storing the enormous quantities of scientific data, which is the key to continuing research and clinical trials. A CDMS can be used as an individual unit or can be part of a set of clinical trials management tools. The CDMS can generally be in two different forms, paper-based and electronic data capturing system. In paper-based systems, the data are filled out by hand. Then, the data on forms is transferred to the clinical data management systems tool. In this phase, there are two ways for data entry. In single data entry, there is one data entry operator while in double data entry, two different data entry operators enter the data separately. Then, the system compares the entered data and in the case of value conflicts, verification should be done. The data in the CDMS are then transferred for data validation ^{26, 96-98}. In electronic data capturing systems, the data are directly uploaded on the CDMS and they can then be viewed by the data validation team ^{95, 99}

1.4.1. Utilization of Clinical Data Management System

The most important use of a CDMS is to collect, to verify, to code, and to prepare the organized data for further statistical analysis. In addition, this tool or set of tools helps to conduct diverse clinical research studies ^{33, 100}. CDMS can be a useful resource for various groups such as organizations carrying out research studies, clinical research sites that are participating in studies, and finally external organizations (labs). Also, the accurate collected patient data in the CDMS can be used by statisticians for statistical analysis to verify the usefulness of the conducted clinical studies ¹³. Generally, multiple uses of clinical data are ensured if the responsibilities of the data management system and analysis questions are determined earlier. As an example, in an individual patient's care the results of all examinations are recorded, while for clinical study, according to the study question, special relevant characteristics are recorded ^{26, 29, 47}.

CHAPTER II

METHODOLOGY

2.1. PROBLEMATIC, HYPOTHESIS, OBJECTIVES

Patients' dental records contain valuable information that can give clinical researchers an opportunity to access and use a wide range of quantitative or qualitative information for conducting clinical and oral health outcomes research ^{8, 9}. However, reusing clinical data can be complicated for a number of reasons including the inaccurate and unstructured nature of the oral health care record, the poor maintenance of accurate and complete clinical data over time, lack of outcome of interest as well as heterogeneity and diversity of interests between clinical researchers, and finally technical issues related to the infrastructures ^{1-6, 78, 93, 101}. Thus, an effective and efficient record-keeping system could facilitate this process and overcome some of these problems. An organized and research-based dental record could be a reliable tool to keep and organize important clinical parameters and aspects of short- and long-term outcomes ^{15, 79-81}. The aim of this study was to design and develop a clinical and research-adapted record in the field of removable prosthodontics.

2.1.1. Hypothesis

Our hypothesis was that the prosthodontic clinical record in the removable department could be redesigned to adequately respond to clinical and research objectives.

2.1.2. Objectives

Our long-term objectives are itemized as follows:

Educational objectives

- 1. To create a research training environment for clinicians and dental students with the goals to:
 - Introduce an evidence-based approach in clinical practice
 - Provide support and education in research
- 2. To assess clinicians' response to and acceptance of the research-based prosthodontic clinical record
- 3. To provide a useful model for other departments and institutions

Clinical and patient-based objectives

- 1. To improve oral health outcomes for patients
- 2. To facilitate auditing of clinical services

Research objectives

- 1. To access a wide range of research data
- 2. To conduct different types of clinical research

2.2. RESEARCH METHODOLOGY AND METHOD

2.2.1. Study Design

A qualitative approach and action research method were used to conduct this study. Action research is a method in which participants help one another by working in a group. This methodology is usually employed in educational studies where the research process assists the participants (clinical professors and clinical researchers) to carry out a needs assessment, document the process, analyze the data, and make decisions to achieve their objectives. As demonstrated in figure 2.1, an action research study has several phases:

identifying the problem, gathering and interpreting data, action planning, acting on evidence, and action evaluation and interpretation ¹⁰²⁻¹¹².

This research methodology can improve and strengthen the links among evidence, research, and practice ^{108, 109}. According to Bell (1999), the action research approach is attractive to both practitioners and clinical researchers because the research will inform and influence their practice ¹¹³. In this type of research, the focus of collaboration is on the interaction between a group of practitioners and a research team.

A practitioner is a person who understands the field from working 'from the inside' and from professional experiences. He/she knows the historical background of the institution and has the knowledge and experiences provided by working within a clinical setting and dealing with its related issues. The nature of any collaboration between the practitioner and researcher is variable. It depends essentially on its preliminary goals and ranges from simple periodic participation to intensive active involvement ^{108, 113-115}.

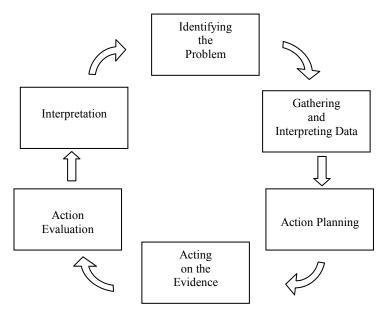


Figure 2.1: Action research cycle model (adapted from Susman, 1983)¹¹⁶

2.2.2. Study Participants

This study used purposive sampling to recruit participants. Purposeful sampling is based on selecting participants who are information-rich ^{117, 118}.

Participants in this study included:

- Five full-time professors (two clinical researchers and three clinicians) in the field of prosthodontics in the Department of Removable Prosthodontics at Université de Montréal.
- Nine part-time professors (clinicians) in the Department of Removable Prosthodontics at Université de Montréal.

2.2.3. Data Collection and Study Phases

Data were collected between December 2010 and October 2011. In total, 6 focus groups and 10 interviews of 60 to 120 minutes duration in French and English were conducted and audio-recorded by a trained interviewer at the Faculty of Dentistry of the Université de Montréal

Different phases of this study comprised (Figure 2.2.):

Phase 1: Problem identification

Focus group discussions as well as semi-structured, open-ended, individual (face-to-face) interviews were carried out to assess interviewees' experiences and perceptions about prosthodontic clinical records and to develop criteria for designing new research-based records.

During individual interviews and focus groups, several questions were asked to gather information about the:

• Person's knowledge and perception on ideal clinical records

- Record-keeping needs, barriers, and difficulties
- Opinions on how to implement a standardardized research-based record.

These questions were used to guide the interview in such a way that the interviewers and interviewees could address issues as they arose. The interviews continued until a saturation level was reached, i.e., no new information could be obtained with more and new interviews ¹¹⁹.

Through discussions with the other participants, everyone in the focus group was able to reflect to the experiences and opinions of the others. The participation of several and different types of professionals during the discussions enabled new perspectives to be introduced ^{120, 121}.

Phase 2: Collection of evidence-based data

An extensive literature review was carried out to identify the main outcomes of interest in removable prosthodontic dental records as well as the standardized data sources that could be linked to answer specific research questions.

We searched systematic reviews in the journals indexed in the most recent edition of the Journal Citation Reports (JCR; 2010), available at http://isiknowledge.com/jcr. We considered all journals under the subject "Dentistry, Oral Surgery & Medicine" that had prosthodontics as the main field of interest. Periodicals about oral implants were also considered, as they likely contain a substantial number of articles about dental prostheses. Table 2.1 shows the titles and respective impact factors for the journals considered.

The Ovid Medline electronic database was searched from 1950 to December week 3 2010 and complemented by hand searching for published randomized controlled trials in English in the field of removable prosthodontics in the six main prosthodontic journals.

In order to obtain reports of systematic reviews, we searched the Medline employing abbreviated titles combined by the "OR" Boolean term. We compiled an initial list of search terms, including index terms and text words. We created groupings of words that combined together with the Boolean term "OR." The results were combined together with Boolean "AND."

Titles were combined as follows: (dental prosthesis OR exp dental implants OR dental prosthesis, implant-supported OR dental prosthesis repair OR denture OR denture bases OR denture liners OR exp denture, complete OR denture design OR denture OR denture, overlay OR denture, partial OR denture, partial, immediate OR denture, partial, removable OR denture, partial, temporary) AND ("clinical implant dentistry & related research".jn. OR "clinical oral implants research".jn. OR "implant dentistry".jn. OR "international journal of prosthodontics".jn. OR "journal of oral rehabilitation".jn. OR "journal of prosthetic dentistry".jn.) AND (exp "Outcome and Process Assessment (Health Care)" OR exp Treatment Outcome).

Table 2.1. List of the prosthodontics or oral implantology journals in the JCR - 2010.

Full Title	Abbreviated Title (ISO)	Impact Factor
Clinical Implant Dentistry and Related	Clin Implant Dent Relat	2.803
Research	Res	
Clinical Oral Implants Research	Clin Oral Implants Res	2.756
Implant Dentistry	Implant Dent	1.455
International Journal of Prosthodontics	Int J Prosthodont	1.423
Journal of Oral Rehabilitation	J Oral Rehabil	1.462
Journal of Prosthetic Dentistry	J Prosthet Dent	1.309

The search yielded 178 papers. Looking at the title and abstract, 117 articles were discarded and consequently 61 articles remained for evaluation at the full text stage. These papers were analyzed to reject articles that did not meet the selection criteria. Five papers were rejected at this stage because they were not clinical trials. The final number of paper included in the review was 56 articles (Figure 2.3).

After determining the main treatment outcomes in removable prosthodontics, we defined validated measurement tools for measuring the prosthodontic outcomes by an extensive literature review. As an example, after defining bone resorption as a treatment outcome, we reviewed the literature to find the most appropriate tools to measure this outcome. We considered the following criteria in the choice of these tools: accuracy, pertinence, and relevancy.

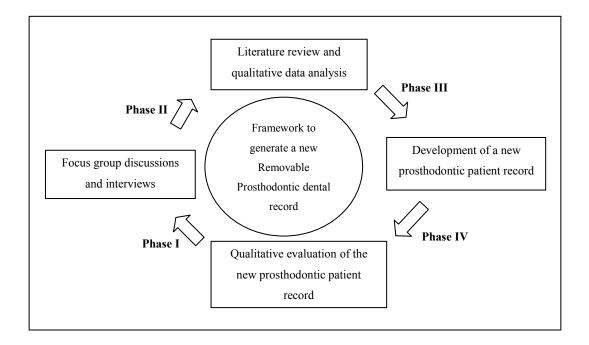


Figure 2.2: Study phases

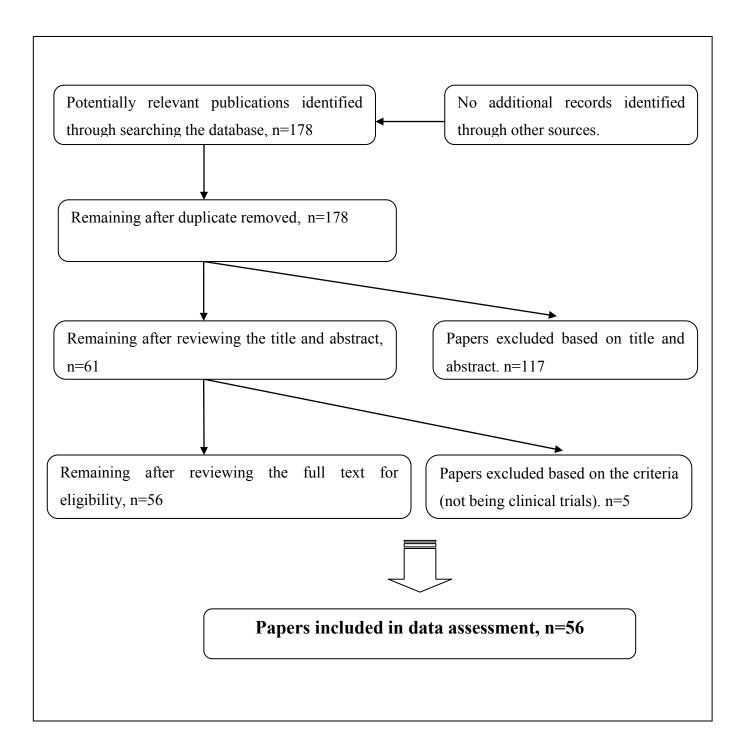


Figure 2.3: Flow chart of search strategy

Phase 3: Action planning

A) Development of new prosthodontic dental record

Based on the literature review and data gathered during qualitative evaluation, the current undergraduate prosthodontic patient record was redesigned and restructured.

B) Evaluation of the new prosthodontic dental record

The completeness and appropriateness of the new dental record was evaluated by complementary interviews with the same panel of experts and those individuals participating in the initial interviews. Necessary modifications were carried out according to experts' opinions. Any differences among members of the group were resolved through discussion with the members of the action research team.

2.2.4. Study Analyses

The analyses included debriefing, transcription, thematic analyzing, and interpretation of interviews and focus groups. The debriefings assessed the data collection and encapsulated the main findings. The computer qualitative software QDA Miner (version 3.2.3, 2009, Provalis Research Corp, Montreal, QC, Canada)

was used to index the transcript and to assign codes to each segment. The interviewers assessed the interview transcripts and described the emerging themes and key points to action research group participants ¹²²⁻¹²⁴. We used triangulation to establish more credibility in the qualitative results. Triangulation is an approach that combines more than one research strategy to make sure that the presented results of the study are true and clear ¹²⁵⁻¹²⁹. In this study, researcher triangulation was used, a process that included different researchers reviewing the data at the different stages. Two members of the research team read all transcripts and defined the themes separately. When their opinions differed, the two research team members came to agreement through discussion in order to avoid individual

interpretation and bias as well as to achieve validity and reliability of the analyzed data ¹²⁵⁻¹³⁰.

2.3. ETHICAL CONSIDERATIONS

This study was approved by the Université de Montréal Research Ethics Board. An informed consent was obtained from all individuals participating in this study.

2.4. CANDIDATE'S ROLE IN THE PROJECT

The candidate developed the protocol for this study, conducted the different phases of the study, gathered and analyzed data, and designed a new dental record in the field of removable prosthodontics.

The candidate's abstract of this research project has been accepted in the 90th General Sessions & Exhibitions of the International Association of Dental Research. The candidate will present the results of this research project in June 2012. The candidate will submit the article included in the chapter III of this master thesis for publication in a prosthodontic journal.

CHAPTER III

RESULTS

3.1. THE MANUSCRIPT

DEVELOPMENT OF AN EVIDENCE-BASED PROSTHODONTIC RECORD: AN ACTION RESEARCH STUDY

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Keywords: Action research, Prosthodontics, Removable prosthesis, Evidence-based dentistry.

DEVELOPMENT OF AN EVIDENCE-BASED PROSTHODONTIC RECORD: AN ACTION RESEARCH STUDY

ABSTRACT

<u>Objectives</u>: The aim of this study was to design a research-based patient record in the field of removable prosthodontics in the undergraduate clinic of the "Université de Montréal."

<u>Methods:</u> This study used action research methods with 4 sequential steps: problem identification, gathering and interpreting data, action planning, and action evaluation. Study participants included professors, clinical researchers, and clinical instructors in the field of removable prosthodontics. Data collection consisted of a comprehensive literature review on prosthodontic outcomes as well as focus-group discussions and interviews. The qualitative data were analysed using QDA Miner 3.2.3.

Results: The study participants raised several concerns about the deficiencies of the existing patients' prosthodontic record in the undergraduate clinic. They shared their ideas for designing a new patient record based on 3 key objectives: clinical, educational, and research objectives. The prosthodontic outcomes of interest and appropriate instruments as well as the clinical parameters were selected by the research group and were integrated into a new evidence-based record. The appropriateness of the new record was evaluated by the same panel of experts and the necessary modifications were carried out. The study participants agreed that the action research cycle should be continued to evaluate the feasibility of the implementation of this redesigned record in the university-based setting.

<u>Conclusion:</u> This study is a beginning effort to develop a database in the field of removable prosthodontics. Action research is a useful research method in this process, and academic educators are well placed to conduct such research.

INTRODUCTION

Although the importance of practice-based research has been widely discussed and documented, this process is still slow and faces several barriers such as conceptualizations of evidence, internal and external validity of the evidence, and high costs of providing large amounts of clinical and patient-based outcome data.^{1, 2} Furthermore, retrieval of data to conduct clinical and comparative healthcare research requires access to clinical records.³⁻⁷ However, these records are often inaccurate or have an unstructured format that makes the use, interoperability, and interpretation of the data difficult or impossible.^{3-5, 8, 9} Several audits in different countries have reported that the quality of dental records is not optimal and many dental records are missing information. 10-15 Therefore, in order to overcome some of these barriers and to decrease the gap between research and practice, we need to implement strategies that support the process of record keeping and access to these records. In the field of prosthodontic research as for other research domains, a university-based dataset can facilitate this process because it will make it possible: 1) to monitor patients' oral health status and prosthodontic care, ¹⁶ 2) to collect large amounts of information across a wide range of interventions and to aggregate a wide range of quantitative and qualitative data, 8, 9, 17-19 and 3) to systematize all important parameters and aspects of short- and longterm oral health outcomes. 10, 20-22 In addition, through this strategy, the clinicians will be exposed to a research-supportive environment that will promote evidence-based dentistry. In this regard, some initiatives have been undertaken to develop a database and to improve the quality of oral health recording systems. As an example, since 2005, several dental schools in North America have formed a Consortium for Oral Health-Related Informatics to develop a data repository that can be shared within these universities.²

To our knowledge a specific dataset to facilitate conducting prosthodontic research in university-based settings has not yet been introduced. Therefore, the objective of this research project was to create an evidence-based patient record in the field of removable prosthodontics that can be used as the basic elements of a prosthodontic database. This record will promote the standardization of the clinical recording system, and will allow dental students to become familiar with research tools and instruments. Additionally, it will provide data for prosthodontic clinical and health-care services research in university-based settings.

MATERIALS AND METHODS

Study Design and Study Participants

This study used a qualitative approach and action research (AR) methods. In health-care settings, action research is a recognized form of experimental research that involves a collaborative process between researchers and clinicians to evaluate and to change practices with the goal of improving performance quality.²³⁻²⁹ Wadsworth's³⁰ process was chosen as an application of AR. This approach is unique since it links researchers and clinicians synergistically to perform a cycle of activities including problem identification, gathering and interpreting data, action planning, action on the evidence, and action evaluation and interpretation (Figure 1).^{24-29, 31-33} Accordingly, this study comprised several phases to reach its objectives.

The study was approved by the Ethics Committee of the Université de Montréal, and an informed written consent was obtained from each participant.

The purposive sampling technique was used to select study participants who were "information-rich," and had experience in research, teaching, and clinical activities in the field of removable prosthodontics.^{34, 35} These included two full-time clinician-scientists, three full-time academic prosthodontists (including the head of the removable prosthodontics unit), and nine part-time clinical instructors in the undergraduate removable

prosthodontics clinic at the Université de Montréal. The demographic and academic characteristics of the study's participants are presented in Table 1.

Study Phases and Data Collection

Data were collected during the different phases of the action research cycle. In the first phase, "Action reflecting/ Problem identification," we conducted 6 focus-group discussions and 13 individual interviews of 60 to 120 minutes' duration between December 2010 and October 2011. All focus-group discussions and interviews were audio-recorded and transcribed by a trained interviewer. Using an interview guide, the study participants were invited to give their views about strengths and limitations of the existing clinical record in the undergraduate clinic and to provide their comments on improvement. ^{36, 37} This phase helped to develop the criteria for designing new research-based prosthodontic records.

In the second study phase, "Gathering and interpreting data," a systematic review was conducted to identify the main reported outcomes in the field of removable prosthodontics. Electronic databases were searched from 1950 to December 2010 and complemented by hand searching the six main prosthodontic journals as reported in the Journal Citation Reports (2010). The abstract of this systematic review has been presented elsewhere, and the related manuscript is under preparation.³⁸ According to the results of this systematic review, and the full-time professors' research profile and interests, the prosthodontic outcomes of interest were selected. Then, the validated measurement instruments for measuring these selected outcomes were identified.

In the third phase, "Action planning/Action on the evidence," based on the literature review and analysis of the data gathered during the focus groups and interviews, the new prosthodontic record was designed.

Finally, in the last phase, "Action evaluation and interpretation," the action research team members evaluated the completeness and appropriateness of the new prosthodontic record based on the criteria developed during phase I and II. Necessary modifications were carried out according to the study participants' comments. In order to minimize the effect of any particular perspective, and to increase the credibility and validity of the results, triangulation techniques were used. 39-42

Data Analysis

The analysis included debriefing, transcription, thematic analysing, and shared interpretations. The interviews and focus groups were all audio-recorded, transcribed verbatim, and coded. The computer qualitative software QDA Miner (version 3.2.3, 2009, Provalis Research Corp, Montreal, QC, Canada) was used to index the transcript and to assign codes to each segment. In this study, we used thematic analyses, which help to identify, to analyse, and to report the themes within collected data. These processes include text reading and dividing into themes and categories. ⁴³⁻⁴⁵ To ensure the credibility and transferability of the data, two researchers in the study team conducted a detailed analysis of the identified code and themes independently, and all research team members reviewed the data at the different stages to check and validate their interpretations. ^{39-42, 46} As required by qualitative research, the saturation level was considered to estimate the necessary sample size and was achieved by the 6 focus-group discussions and 13 individual interviews.

RESULTS

The thematic analysis of the collected data during the action research process yielded several key concepts and themes, which are summarized below.

Enthusiasm to change and collaboration in planning

This key concept was evident when the professors and clinical instructors agreed to participate in the study, and demonstrated their willingness to collaborate in the improvement of the existing patient prosthodontic record in the undergraduate prosthodontic clinic.

Empowerment in practice, education, and research

This concept was manifested during the first phase of the study. The research participants expressed their views about the weaknesses of the existing patient record. The full-time professors criticized the existing form more than did the clinical instructors. They mentioned that the patient record was outdated and had been developed by a clinician in the removable prosthodontics undergraduate clinic approximately 30 years ago. The head of the clinic mentioned some minor modifications and add-ons that had been made later. These modifications were made to reflect the notion of implant dentistry in the patient record. Three types of weakness in the current prosthodontic patient record were observed by research participants: 1) Clinical weaknesses: for most interviewees, the actual clinical form did not allow following up the patients or conducting clinical audit. In addition, the interviewees mentioned that the clinical form was totally theoretical in format. 2) **Educational weaknesses:** most of the professors expressed that the information gained by the actual patient record did not allow the students to develop clinical decision-making skills. They would prefer a student training based on a clinical decision-making model and hypothetico-deductive approach.⁴⁷ Furthermore, they identified a mismatch between theoretical and practical training. They suggested to include in the new patient record several validated measurement tools to help students define patient complaints, to verify clinical parameters, to make a diagnosis and treatment plan, and finally to reassess the patient. In addition, the participants mentioned that the students' clinical knowledge should be expanded by teaching them the most important prosthodontic clinical outcomes. 3) Research weaknesses: The clinical researchers stated that the existing patient

prosthodontic record was solely clinical, and was not designed to address research needs. The faculty professors pointed out that research is not attractive to young dental students because clinical training is the main objective of their education. Their wish was to enable active research activities in the undergraduate clinic and to motivate students to be involved in future research careers. Furthermore, they wanted the new patient record to allow gathering of a wide range of data, and conducting different types of clinical research, as well as providing support and education in research.

Barriers to change

Combining research and clinical training was found to be difficult in the undergraduate clinic because of several barriers such as deficient infrastructure, lack of time in the clinical sessions, and lack of research training for clinical instructors. Clinical instructors were more positive regarding the clinical, educational, and research capacity of the clinic and the potential role of a patient recording system in building capacity. Most of the instructors mentioned that by following patient record elements, the students were capable of doing intra and extra oral examinations, asking their supervisors relevant questions, and discussing patients' needs and treatment. The majority of clinical instructors suggested having an easy-to-use, organized clinical form, and to be realistic in the design of the new format in terms of time needed and the clinical setting. They mentioned that the prosthodontic record should facilitate the clinical examination without posing a burden on students and educators. They believed that these records could be used as a trustworthy resource for research needs. However, they mentioned that the reliability of the records would depend on the aptitude of the students and the clinical instructors. In this study the female study participants were more flexible towards accepting the new changes and the older clinicians were less interested in the clinical research activities.

Expanding knowledge

During the problem identification phase, the research team members noted that an extensive systematic review was needed to select the important outcomes in removable

prosthodontic research. The results of this systematic review indicated that there was a significant increase in the selection of patient-based outcomes between each 10-year time interval since 1990.³⁸ The research participants agreed on the importance of the identified outcomes in the design of new prosthodontic forms.

In the phase of action planning, the new evidence/research-based prosthodontic record was developed, and then was evaluated by the full-time professors. This new prosthodontic record consists of three parts: the first part comprises a series of assessments on the potential risk factors of prosthodontic outcomes. These include assessment of sociodemographic characteristics, medical and dental history, life style habits, dental service use, oral hygiene habits, dental anxiety, and psychological characteristics. The second part constitutes the oral clinical examination form and the assessment of diseaseoriented outcomes such as caries, periodontal diseases, denture stomatitis, and alveolar bone resorption. Finally, the third part includes assessment of patient-oriented outcomes such as oral health quality of life, patient satisfaction, and dental visit satisfaction. The fulltime professors recommended evaluating the dental record occasionally according to the relevant literature in removable prosthodontics. The study participants agreed that the action research cycle should be continued to evaluate the feasibility of the implementation of this redesigned record in the university-based setting. Several professors expressed their willingness to add a theoretical course and to introduce the new prosthodontic patient record to the undergraduate students.

DISCUSSION

Through this action research study, we have successfully designed and developed an evidence/research-based patient record in the field of removable prosthodontics in the undergraduate clinic of the Université de Montréal.

Our experience showed that applying action research methods ensures the needs, perspectives, and expertise of the participants in the design and implementation of the new prosthodontic record. This patient record will help gather accurate and organized research data in the university-based setting. The research team believed that this approach could create a research-training environment for clinicians and dental students, and will raise awareness about evidence-based prosthodontics. According to the Allison and Bedos survey⁴⁸ on the utility of clinical research in Canada, although the vast majority of dental practitioners were interested in research, the implementation of evidence-based health care was found to be difficult.

Nowadays it is recognized that dentistry, as is the case with other health-care fields, requires integrating the concept of evidence-based health. However, the challenges of how to bridge research and dental practice still exist. Research education could be a solution to this dilemma. In this regard, many dental schools have implemented an educational approach that helps promote evidence-based thinking in the context of clinical training.^{49, 50} This approach allows behavioural changes in the students and clinicians, favors their dedication to scientific research, expands their knowledge, and increases their willingness to support and invest in evidence-based health care. This would help future dentists to use research evidence to offer the optimal care for their patients.⁵¹⁻⁵³

Our results demonstrated that there exists diversity between the views of full-time professors and those of clinical instructors concerning students' research and clinical training. This diversity could be explained by the fact that the professors have different obligations and needs based on their career profiles. All tenure-track faculty members are expected to conduct research activities to achieve their tenure. By contrast, clinical instructors don't have any research responsibility, and they do not generally perform any

independent research. Furthermore, their training and education is mostly limited to practice and they don't receive formal training in education and research. 49, 50, 53

According to Chapnick et al. (1998), part-time dental instructors have the greatest effectiveness while acting as a facilitators rather than educators.⁵⁴ In our study, the full-time professors mentioned that the part-time clinical instructors are important keys in success or failure of the implementation of a new educational/research tool in the clinic. Since these instructors are an important element in success, the following possible strategies could be considered. These include to build the base of the research knowledge and to provide instructional content and methodology by standardizing the medical and dental health record system and facilitating the exchange of information.

For example, the Consortium for Oral Health-Related Informatics (COHRI)² is the result of collaboration of 20 dental schools to develop a monograph that could serve as a guide in the acquisition of an oral health information system. The COHRI provides standardized data for diverse clinical research studies by assembling the largest oral health database ever created. It also allows measuring student clinical performance, determining the accuracy of the treatment planning, and validating the educational outcomes.² Similarly. some initiatives have been undertaken to improve the prosthodontic clinical recording system. For example, the American College of Prosthodontists has developed a classification system to provide a framework for the organization of clinical observations in removable prosthodontics. The use of this framework facilitates clinicians' communication, clinical outcomes assessment and research. 55, 56 The implementation of this new clinical prosthodontic record will allow powerful data queries on large pools of patient data with relatively low cost and without information and measurement bias. Furthermore, using this record in the undergraduate clinic will enable dental students to improve their clinical and research knowledge. In fact, it will raise research awareness and the essential basics of education.^{3, 7} In addition, this newly designed record will help to maintain high standards of record-keeping, which is a necessary part of quality dental care.⁵⁷ Although this new prosthodontic record is designed for the removable prosthodontics clinic, it can also be used as a template for other dental specialities or clinical fields.

Any research methodology has its limitations. Since this study only reflects the perceptions and the experiences of the clinicians and faculty team members in the field of prosthodontic research, it may not be generalizable and applicable to many other disciplines without additional refinement. In terms of the implementation, we could anticipate barriers for various reasons such as the students' and clinicians' resistance toward complexity of the design, lack of knowledge about the concept, and the deficient infrastructure. However, we believe that once the clinicians and students become familiar with the new prosthodontic record and receive appropriate support and training, these barriers could be resolved.³ This phase should include post-implementation surveys and interviews with dental students, clinical instructors, and faculty members to evaluate the feasibility and efficacy of this patient record in the clinical setting. Then, the research cycle may start to find solutions for any potential barriers. Finally, evaluation studies should be conducted to assess the long-term impact of this record on clinical, educational, and research capacities.

CONCLUSION

This study is a beginning effort to develop an evidence/research-based patient record in the field of removable prosthodontics. The study shows how action research can be useful in this process and demonstrates that academic educators are well placed to conduct such research. Developing a prosthodontic patient record has several advantages, such as monitoring prosthodontic care, collecting a large amount of valid data, and facilitating clinical research and clinical audit in the university-based setting.

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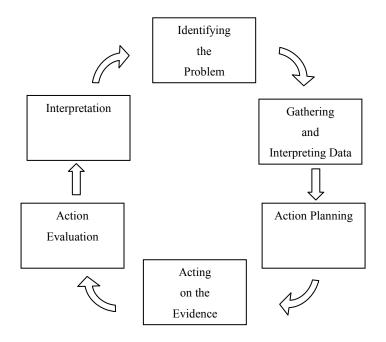


Figure 1: Action research cycle model (adapted from Susman1983)⁵⁸

Table 1. Characteristics of study participants (N=14)

Participants	Numbers	
Gender		
Male	9	
Female	5	
Academic Status		
Full-time professor	5	
Part-time clinician	9	
Age		
30-40 years	4	
41-50 years	6	
51+ years	4	
Teaching Experience		
10 years	1	
10-20 years	4	
21 - 30 years	6	
31≥ years	3	

CHAPTER IV

DISCUSSION AND DIRECTIONS FOR FUTURE RESEARCH

The measurement of oral health outcomes in clinical settings relies on the maintenance of an accurate and valid clinical record. This action research study started by identifying the limitations and deficiencies of the existing prosthodontic patient record in the field of removable prosthodontics at the undergraduate clinic of Université de Montréal. Through the research cycle and active participation of faculty members and clinicians, ideas for the improvement and design of an innovative research-based patient record system arose. Finally, an evidence-based patient record was tailored and driven by clinicians and faculty members. This study is significant because of its evidence-based nature and its action research design that empowers clinical researchers and clinicians to recognize clinical, educational, and research deficits in the patient record system and helps them to meet their needs. In this chapter, we will briefly discuss this project.

5.1. THE CHOICE OF STUDY DESIGN

In this study, a qualitative approach and an action research method were selected as the study design. In general, there are three research approaches in the field of health and oral health science: quantitative, qualitative, and mixed methods ¹³¹. A researcher's choice of measurement approach depends on several factors, including experience and personal training of the researcher, the audience, the type of outcome, and most importantly, the research question ^{131, 132}.

Nowadays, qualitative research approach has found its place within dentistry by generating new hypotheses and providing important information to answer questions, that cannot be found using quantitative techniques ^{110, 132-136}. For example, programs, services, and dental treatments could be all examined by qualitative methods ^{137, 138}.

Action research was introduced by psychologist Kurt Lewin in the mid 1940s ^{106,} ¹³⁹⁻¹⁴¹. He described action research as "proceeding in a spiral of steps, each of which is

composed of planning, action and the evaluation of the result of action" ¹⁴²⁻¹⁴⁵. In contrast to traditional dental research which is validated by statistical presentation, action research uses a qualitative process (e.g., interview, focus group discussion, etc.) and qualitative analysis to identify what people really think, believe, and do ¹⁴⁶. According to Bell (1999), action research is a useful research approach for clinical scientists who have identified a problem during the course of their practice and who would like to find solutions to improve their practice ¹¹³.

As the goal of our study was an "action" (developing a new prosthodontic patient record), the information gathered within the research cycle helped us in decision making about goals and objectives for the desired record. Furthermore, this research methodology will facilitate the transfer of this research to the clinic because clinicians formed part of the research team rather than being the tools of an outside researcher. They will continue to collaborate to implement and evaluate the study results, since they are responsible and self-decision-makers ^{147, 148}.

The action research method has been widely used in the field of oral health. Drewry and Chu (1995) conducted an action research project to gather information on issues related to dental care access ¹⁴⁶. Reeson and Jepson (2005) used action research to improve collaboration and communication between dental undergraduates and dental technicians to increase the quality of dental care for patients in removable prosthodontics ¹¹³.

In this study, the focus groups and interviews were used to capture opinions from the professors. The important advantages of the methods used include capturing data that may be difficult to understand as opinions and beliefs, and helping to generate a wide range of information and innovative ideas by creating synergy between the participants. In addition, in the focus group, the interviewees discuss the issues at the same time. Thus, data collection is less time consuming and the process helps participants to be more comfortable to express their beliefs ^{117, 149, 150}.

5.2. THE CHOICE OF STUDY SETTING

This study was conducted in a university-based setting because of the important role of universities and educators in translating research and research findings into dental education. Promoting and strengthening research skills in clinicians and students, as well as training them how to apply this new knowledge to their practices are considered to be the main roles of dental schools and their faculty members ¹⁵¹. Furthermore, the students should be able to perform evidence-based clinical decision-making, which has important impact on the practice of dentistry ^{152, 153}. They should also learn to the importance of having a documentation record that allows patient treatment follow-up. Therefore, this study will help to implement new learning strategies in the dental education setting. This will increase the ability for scientific thinking among dental students ^{154, 155}.

5.3 THE CHOICE OF STUDY TEAM MEMBERS AND PARTICIPANTS

The use of purposeful sample strategies to select the team members and interviewees allowed us not only to collect information-rich data but also ensured variation in the views expressed. The active involvement of researchers, academic clinicians, and clinical instructors in this study served several critical functions:

- 1) Following the action research cycle, the research team members have accumulated considerable knowledge about how the idea for the new dental record emerged and how it was developed. They have also gained considerable knowledge in teaching and explication of the new concept to students.
- 2) The research team members validated that the new dental record is implementable and created value for it.

3) These educators will play an active role in the implementation process of the study results in the clinic.

5.4. STUDY OBJECTIVES AND FINDINGS

Nowadays it is recognized that dentistry, like other health-care fields, requires integrating the concept of evidence-based health. However, there still exist challenges in how to bridge research and dental practice. Research education could be a solution to this dilemma. In this regard, many dental schools have implemented an educational approach that helps promote evidence-based thinking in the context of clinical training ^{151, 156}. This approach allows behavioral changes in the students and clinicians and increases their willingness to support and invest in evidence-based health care. This would help future dentists to use research evidence to offer the optimal care for their patients ^{157, 158}.

One of the tools of this approach is to build the base of research knowledge and to provide instructional content and methodology by standardizing the medical and dental health recording system and facilitating the exchange of information. As an example, the Consortium for Oral Health-Related Informatics (COHRI) ⁵¹ is the result of collaboration of 20 dental schools to develop a monograph that could serve as a guide in the acquisition of an oral health information system. The COHRI provides standardized data for diverse clinical research studies by assembling the largest oral health database ever created. It also allows measuring student clinical performance, determining the accuracy of treatment planning, and validating educational outcomes ⁵¹. Similarly, some initiatives have been undertaken to improve the prosthodontic clinical recording system. For example, the American College of Prosthodontists has developed a classification system to provide a framework for the organization of clinical observations in removable prosthodontic. This framework has potential benefits including: improving professional communication, increasing diagnostic and treatment stability, and standardizing criteria for outcomes assessment and research ^{159, 160}.

Based on this concept, the main aim of this action research project was to create an evidence-based recording system, to increase research and educational capacities as well as to improve treatment outcomes in the undergraduate removable prosthodontic clinic.

During the research action cycles, the professors expressed their perceptions about the limitations, weaknesses, and strengths of the current prosthodontic record. In addition, the participants were asked about their suggestions to realize a comprehensible and feasible evidence-based clinical record.

The study results indicated that the research team had the desire to develop a clinical tool that could respond to clinical, educational, and research needs. Accordingly, the research team members expressed the need for a practical clinical guide for oral examination, diagnosis, and treatment decision making, which will improve oral health outcomes for patients and will enhance clinical audit. Furthermore, this tool should simplify the collection and retrieval of clinical parameters for research.

In addition, the research team expressed their belief that this approach could create a research training environment for clinicians and dental students and will introduce an evidence-based approach. The focus group discussions and interviews with study participants allowed reflection on the effectiveness and design of this new tool. This precious and rich information was then translated into the action planning phase.

Our findings showed that the current format was solely clinical and it has not been designed to address research needs. In addition, the acquisition of clinical information was judged by the experts in the field to be incomplete and outdated. Thus, redesigning a new format to overcome these weaknesses and limitations was found to be necessary.

However, the research team members were concerned about conditions leading to unsuccessful implementations of the new model, and gave insights about the conditions for

success, as well as for failure. The full-time professors mentioned that the part-time dental instructors are important in succeeding or failing of the implementation of the new record in the clinic, therefore it would be essential to ascertain their perceptions of the process and their willingness to adapt to these new changes ¹⁵³. In this regard, the interview results with both full-time and part-time professors demonstrated that there exists diversity between the opinions of the full-time professors and those of part-time clinical instructors in regard to the formatting of the prosthodontic clinical record. The full-time professors expressed that the new clinical form should not only provide interesting research elements, but also it should simultaneously respond to clinical needs. However, a number of clinical instructors insisted that the present form is adequately precise and a useful tool for clinical examination. Many believed that the new form will be a time-consuming task. For that reason they preferred a brief dental record form with simplified presentation, which would facilitate clinical examination without putting a burden on them and on students. This diversity could be explained by the fact that the professors have different obligations and needs based on their career profile. All tenure-track faculty members are expected to perform research activities to achieve their tenure. By contrast, clinical instructors do not have research responsibility and do not generally perform independent research. Furthermore, their training and education is mostly limited to the undergraduate level and they don't receive formal training in education. According to L. Chapnick and A. Chapnick (1998), part-time dental instructors have the greatest effectiveness while acting as a facilitators rather than educators ¹⁶¹.

In order to overcome the above mentioned barriers, the following possible strategies could be considered. A close communication between clinical researchers and clinical instructors should be encouraged. Additionally, continuing education and workshops led by academic professors for the clinicians would be helpful. Such a close communication would also assist to deliver more high-quality and evidence-based care to the patients ¹⁵¹.

The systematic literature review in this project helped to identify the most important treatment outcomes of interest in removable prosthodontics research and to assess their evolution and their quality over time. The study results indicated that many current studies of prosthodontic treatment use patient-based outcomes and that there has been a significant increase in the selection of patient-based outcomes in each 10-year time interval since 1990 ¹⁶². This could be explained by the fact that although in prosthodontics technical skills are extremely important, they are not the main predictors of patient satisfaction with treatment. So, the use of patient-based measurement instruments has increased over the last decades. These tools make it possible to measure the social, psychological, and physical impacts of oral diseases and their associated treatments ¹³². Thus, in the design of the prosthodontic dental record, we considered patient-reported outcomes as well as clinically measured objective outcomes.

The newly designed clinical record is made of three parts. The first will allow the assessment of risk factors. These include assessment of sociodemographic characteristics, medical and dental history, life style habits, dental service use, oral hygiene habits, dental anxiety, and psychological characteristics (Appendix I). The second part consists of the oral clinical examination form and the assessment of disease-oriented outcomes such as caries, periodontal diseases, denture stomatitis, and alveolar bone resorption (Appendices II and III). Finally, the third part includes assessment of patient-oriented outcomes such as oral health quality of life, patient satisfaction, and dental visit satisfaction (Appendix IV).

Although the development of the new prosthodontic records with clinical, educational, and research characteristics was successful and feasible, we believe that to ensure a balance in education, clinical care, and research productivity within faculties, we have to recognize and resolve fundamental problems. The fact that dental research is a prerequisite for dental education should be integrated into the curriculum of the dental education setting for undergraduate students. Dental schools should inform dental students

about the importance of research that can be incorporated into the process of clinical decision making ¹⁵¹. In this way, dental schools will develop reflective and technically capable practitioners in the future ^{154, 156}. Furthermore the faculty should create favorable research milieus by allocating resources such as sufficient scientific faculty and research infrastructure.

5.5. STUDY RELEVANCE AND PRACTICAL IMPLICATIONS

To the best of our knowledge, this study is among the first to create a prosthodontic evidence-based recording system in a dental academic institution. Development of a standardized dental record that integrates patient dental care data with clinical research data is a high priority in clinical research. The process of collecting prospective clinical data is laborious, time consuming, and expensive. The implementation of this new clinical prosthodontic record will allow powerful data queries on a large patient databank, with relatively low cost and without information and measurement bias ².

Furthermore, using this record in the undergraduate clinic will enable dental students to improve their clinical and research knowledge. Importantly, we will raise research awareness and the essential basic of education ². In addition, this newly designed record will help to maintain high standards of record-keeping, which is a necessary part of quality dental care ⁶⁴. Although this new prosthodontic record is designed for the removable prosthodontic clinic, it can also be used as a template for other dental departments within the Université de Montréal as well as other universities.

5.6. STUDY LIMITATIONS

Qualitative research, like other research methods, has particular criteria to be valid and reliable. According to Lincoln and Guba (1985), these criteria include credibility (vs.

internal validity) and transferability (vs. external validity) ¹²⁶. Furthermore, when the results are trustworthy and believable for research participants and readers, credibility in qualitative research is obtained.

In this study we attempted to increase study validity through means such as triangulation (using different data sources, methods of data collection, and different team members to compare the results), precise coding of data, checking the interpretations of collected data with the participants, and controlling the effects of the interaction between the research team and the study participants ^{117, 130}. However, in focus groups it is likely that the whole discussion may be dominated by a few individuals. Thus, this can result in inaccurate and biased results ¹⁶³.

Transferability refers to the extent that the research result can be applied to the other settings or groups ¹⁶⁴. The procedure to increase transferability in qualitative research includes: enough reporting of the sampling strategy, the methods, and the results in order to allow others to reproduce the study in a similar or different setting ^{117, 130}.

Since, this study only reflects the perceptions and the experiences of the clinicians and faculty team members in the field of prosthodontic research, it may not be generalizable and applicable to many other disciplines without additional refinement.

In terms of implementation, we could expect barriers such as students' and clinicians' resistance toward complexity of the design, lack of knowledge about the concept, and the existing infrastructure. However, believe that once the clinicians and students become familiar with the new prosthodontic record and receive appropriate feedback, these blocking limits will be resolved.

5.7. FUTURE RESEARCH

At this stage of the action research cycle, the new designed prosthodontic record is at an early, crude form. The implementation phase should be piloted in an academic session in the undergraduate clinic. This phase should include post-implementation surveys and interviews with dental students, clinical instructors, and faculty members to evaluate the feasibility and efficacy of this patient record in the clinical setting. Then, the research cycle may start to find solutions for any potential barriers. Finally, evaluation studies should be conducted to assess the long-term impact of this record on clinical, educational, and research capacities.

CHAPTER V

CONCLUSIONS

The results of this action research study suggest that:

- 1. To ensure the translation of the research into clinical training, the teamwork of clinical researchers and clinicians is essential.
- 2. Action research is a useful and feasible research method in the university setting. Through this method, we have developed a new patient dental care record that reflects the needs of those who are interested in removable prosthodontic research, training, and practice.
- 3. Applying an evidence-based approach in the patient clinical record systems in the field of removable prosthodontics is practical and has real potential.
- 4. A prosthodontic clinical record should respond to both educational and research objectives while improving the quality of care for patients.
- 5. Clinicians and students should be informed, educated, and motivated to assure the successful implementation of this new prosthodontic patient record.

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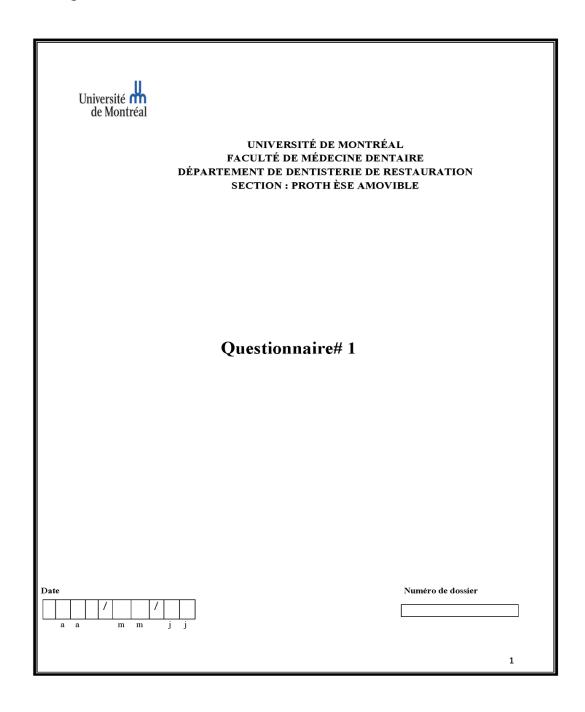
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APPENDICES

APPENDIX I: QUESTIONNAIRE #1



Section A: Questionnaire sociodémographique	
A.1. Êtes vous ☐ Mâle ou ☐ Femelle ?	
A.2. Date de naissance: Année	
A.3. Quels sont les trois (3) premières lettres/chiffr	es de votre code postal?
A.4. Êtes-vous né(e) au Canada? ☐ Oui ☐ Non	(dans quel pays êtes-vous né(e)?):
A.4.1. Si non, quand avez-vous immigré a	u Canada? Année : ШШШШ
A.5. Êtes-vous une personne autochtone? 🗖 Oui	□ Non
Si Non, S'il vous plaît, allez à la question A.7 Si Oui ➤ A.5.1. Êtes-vous: ☐ Indien nord-améric	ain 🗆 Métis 😊 Inuit (Esquimau)
A.6. Êtes-vous membre d'une bande indienne ou d Si Oui ➤ Quelle bande indienne ou Première nation	
A.7. Quelle est votre origine ethnique (veuillez en	choisir plus d'une, si elles sont pertinentes)?
☐ Africain (Afrique /Afro-américain)	☐ Amérique du nord (Canadien-français,
☐ Asie de l'Est (S'il vous plaît cocher une case:	Canadien-anglais, Américain, Mexicain) Amérique du Sud (par exemple Latine / Hispano-
☐ Chinois, ☐ Japonais, ☐ Coréen) ☐ Européen (par exemple Slaves, Germaniques,	américain)
Anglo-saxon, Scandinave, Grecque)	☐ Asie du sud (par exemple des Indes orientales, Pakistanais, Sri lankais)
☐ Autochtones / Natif américain	☐ Asie du sud est (par exemple au Cambodge, Indonésie, Laotien, Vietnamien)
☐ Moyen-Orient /Afrique du nord (par exemple Afg Israéliens, Palestinien, Syrien, Tunisien, Turc)	ghan, Algérien, Marocain, Égyptien, Iranien, Irakien,
☐ Autre ➤ Veuillez préciser :	
En référant à la liste ci-dessus:	
A.8. Quel est l'origine culturelle et géo ethnique de	e vos parents?
Mère?Père?	<u> </u>
A.9. Quel est l'origine culturelle et géo ethnique de	e vos grands- parents?
Mère du père? Mère de Père du père? Père de	
	2

☐ Française	☐ Anglais	e 🗆	Arabe □I	Espagnole	□ Ita	lienne 🗖 Autre	::
A.11. Quel est ☐ Moins que : ☐ \$20,000 à \$ ☐ \$40,000 à : ☐ \$60,000 à \$ ☐ Ne sais pas	\$10,000 par at 5 29,999 par at \$49,999 par at 5 69,999 par at	nnée nnée nnée	□ \$30 □ \$50 □ \$70	,000 à \$19 ,000 à \$39 ,000 à \$59 ,000 ou plu fère ne pas	,999 pa ,999 pa is par a	ar année ar année nnée	
4 12 Oval act	. 10 marramir da :	rratus us ámo	2	_	_		
A.12. Quel est ☐ Moins que : ☐ \$30,000 à \$ ☐ \$70,000 à \$ ☐ \$110,000 à ☐ Ne sais pas	\$10,000 par a 649,999 par an 689,999 par an \$139,999 par	nnée née née	□ \$10 □ \$50 □ \$90 □ \$14	,000 to \$29 ,000 à \$69, ,000 à \$109 0,000 ou pl fère ne pas	,999 pa: 9,999 p lus par	r année ar année an	
A.13. Habitez-	-vous						
	· ous.			- D (0)			
□ Seul (e) □	l Avec d'autre	s adultes	J En famille	☐ Prefere	e ne pas	répondre	
A.14. Quel est	votre statut m	natrimonial	,		•		mme
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre≽ Vei A.15. Quel a é	votre statut n (jamais marié éparé (e) uillez préciser té le plus haut	natrimonial' (e) et aucun Divorcé (conjoint(e) e)	de fait) Veuf (ve) Préfère ne	Marié(pas rép	(e) ou vivant cor	
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre≻ Vei A.15. Quel a é un et encerclez	votre statut n (jamais marié éparé (e) iillez préciser té le plus haut z l'année si ap	natrimonial' (e) et aucum Divorcé (niveau de s plicable, s'i	conjoint(e) e)	de fait) □ Veuf (ve) Préfère ne e vous avez	Marié(pas rép	(e) ou vivant cor	
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre > Ver A.15. Quel a é un et encerclez ☐ Pas d'école	votre statut n (jamais marié léparé (e) uillez préciser té le plus haut z l'année si ap	natrimonial' (e) et aucum Divorcé niveau de : plicable, s'i	conjoint(e) (e)	de fait) Veuf (ve) Préfère ne e vous avez	Marié(pas rép compl	(e) ou vivant cor ondre été? (Veuillez er	
marié(e) □ S □ Autre≯ Vet A.15. Quel a é un et encerclez □ Pas d'école □ École prima	votre statut n (jamais marié iéparé (e) uillez préciser té le plus haut z l'année si ap	natrimonial' (e) et aucun Divorcé (niveau de : plicable, s'i P 2 3	conjoint(e) [e)	de fait) Veuf (ve) Préfère ne e vous avez	Marié(pas rép compl	(e) ou vivant cor ondre été? (Veuillez er	n encerclez
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre ➤ Vei A.15. Quel a é un et encerclez ☐ Pas d'école ☐ École prima	votre statut n (jamais marié léparé (e) luillez préciser té le plus haut z l'année si ap laire: 1	natrimonial' (e) et aucun Divorcé (niveau de splicable, s'i	conjoint(e) [e)	de fait) Veuf (ve) Préfère ne e vous avez	Marié(pas rép compl	e) ou vivant cor ondre été? (Veuillez ei	n encerclez
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre > Vei A.15. Quel a é un et encerclez ☐ Pas d'école ☐ École prima ☐ École secon ☐ Collège (CI	votre statut n (jamais marié léparé (e) lillez préciser té le plus haut z l'année si ap aire: 1 ndaire: 7 EGEP/Technic	natrimonial' (e) et aucun Divorcé (iniveau de : plicable, s'i 2 3 8 9 que):	conjoint(e) e)	de fait) Veuf (ve) Préfère ne je vous avez s répondre 11	Marié(pas rép compl	(e) ou vivant cor ondre été? (Veuillez ei 7 Certificat d'éq	n encerclez
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre > Vei A.15. Quel a é un et encerclez ☐ Pas d'école ☐ École prima ☐ École secon ☐ Collège (CI ☐ Programme	votre statut n (jamais marié léparé (e) luillez préciser té le plus haut z l'année si ap naire: 1 EGEP/Technic	natrimonial' (e) et aucun Divorcé niveau de : plicable, s'i 2 3 8 9 que):	conjoint(e) e)	de fait) Veuf (ve) Préfère ne e vous avez s répondre 5 11 /programm	Marié(pas rép compl 6 12	(e) ou vivant cor ondre été? (Veuillez en 7 Certificat d'éq	n encerclez
A.14. Quel est ☐ Célibataire marié(e) ☐ S ☐ Autre ➤ Vei A.15. Quel a é un et encerclez ☐ Pas d'école ☐ École prima	votre statut n (jamais marié léparé (e) luillez préciser té le plus haut z l'année si ap naire: 1 EGEP/Technic	natrimonial' (e) et aucun Divorcé niveau de : plicable, s'i 2 3 8 9 que):	conjoint(e) e)	de fait) Veuf (ve) Préfère ne e vous avez s répondre 5 11 /programm	Marié(pas rép compl 6 12	(e) ou vivant cor ondre été? (Veuillez en 7 Certificat d'éq	n encerclez

Section B: Modes de vie
B.1.Selon votre activité physique, vous considéreriez-vous comme : Très actif Moyennement actif Légèrement actif Pas du tout actif B.2. Au cours de la dernière semaine, combien de jours avez-vous fait un total de 30 minutes ou plus d'activité physique qui a été suffisante pour élever votre rythme respiratoire. (Veuillez encercler une réponse) (Cela peut inclure le sport, l'exercice et la marche rapide ou du vélo pour le loisir ou pour se déplacer, mais ne devrait pas inclure les travaux ménagers ou l'activité physique faisant partie de votre travail.)
0 1 2 3 4 5 6 7
B.3. Êtes-vous motorisé (voiture, camionnette, moto, etc)?
B.4. Quel est le moyen de transport le plus courant que vous utilisez ?
☐ Ma propre voiture ☐ Covoiturage ☐ Taxi ☐ Transports publics ☐ Marche / Vélo
B.5. Avez-vous déjà fumé un total de 100 cigarettes ou plus durant votre vie?
☐ Oui ☐ Non (Si non, veuillez passer à la ☐ Je ne sais pas ☐ Je préfère ne pas répondre question B.12)
B.6. Y a-t-il déjà eu une période durant laquelle vous avez fumé des cigarettes régulièrement (au moins une fois par semaine)?
☐ Oui ☐ Non ☐ Je ne sais pas ☐ Je préfère ne pas répondre
B.7. Quel âge aviez-vous la première fois quand vous avez commencé à fumer des cigarettes régulièrement? L. Ans B.8 Fumez-vous encore des cigarettes?
☐ Oui ☐ Non (Si non, s'il vous plaît passez à la ☐ Je ne sais pas ☐ Je préfère ne pas répondre deuxième partie de cette question)
B.8.1. Si non, quel âge aviez-vous lorsque vous avez cessé de fumer régulièrement :
B.9. Pendant les périodes où vous fumiez, combien de cigarettes aviez-vous l'habitude de fumer? (généralement 1 paquet contient 20 cigarettes) \bot # de cigarettes
☐ Par jour ☐ Par semaine ☐ Par mois ☐ Je ne sais pas ☐ Je préfère ne pas répondre
4

□ Oui :	De l'âge L De l'âge L De l'âge L	l à l'âge l l l No l à l'âge l l l à l'âge l l l l l l l l l l l l l l l l l l l	on □ Je ne s	sais pas 🔲 Je p	oréfère ne pas répo		
différents n	noments de vie. Essaye	avoir une idée sur le type votre vie. Nous voudric z s'il vous plait de vous il y a lieu).	ons que vous p	ensiez à quatre ar	mées différentes		
De (âge)	À (âge)	Nombre de cigarettes par jour	Marque la plus fumée	Type lo	e plus fumé		
				☐ Avec filtre	□ Roulé		
				☐ Sans filtre	☐ Je ne sais pas		
				☐ Avec filtre	☐ Roulé		
				☐ Sans filtre	☐ Je ne sais pas		
				☐ Avec filtre	☐ Roulé		
				☐ Sans filtre	☐ Je ne sais pas		
				☐ Avec filtre	☐ Roulé		
				☐ Sans filtre	☐ Je ne sais pas		
B.12. Au cours des 12 derniers mois, avez-vous bu de la bière? (S'il vous plaît ne pas inclure pière sans alcool.) Non (passez à la question suivante) Oui B.12.a) Combien de fois avez-vous bu de la bière en été? Jamais							

☐ Jamais ☐ 1-2 fois par semaine ☐ 1fois par jour ☐ 6 fois ou plus par jour	☐ 1fois par mois ou moins☐ 3-4 fois par semaine☐ 2-3 times fois par jour	2-3 fois par mois5-6 fois par semaine4-5 fois par jour
B.12.c) Chaque fois que habituellement?	vous avez bu de la bière ,	quelle quantité en buviez-vous
☐ Moins que 1 cannette ou l☐ 1 à 3 cannettes ou bouteill☐ Plus de 3 cannettes ou bou	es de 12 onces	
B.13. Combien de fois avez-	-vous bu du vin ou des panachés	de vin?
☐ Jamais (Passez à la quest☐ 1-2 fois par semaine☐ 1fois par jour☐ 6 fois ou plus par jour	ion B.14.)	ine 5-6 fois par semai
B.13.a) Chaque fois que v buviez- vous habituellement Moins de 5 onces ou moir 5 à 12 onces ou 1 à 2 tasse Plus de 12 onces ou plus	r? ns de 1 tasse es	anachés de vin , quelle quantité en
B.14. Combien de fois avez-	-vous bu de la liqueur ou des coc	ektails?
☐ Jamais ☐ 1-2 fois par semaine ☐ 1fois par jour ☐ 6 fois ou plus par jour	☐ 1fois par mois ou moins☐ 3-4 fois par semaine☐ 2-3 times fois par jour	2-3 fois par mois5-6 fois par semaine4-5 fois par jour
B.14.a) Chaque fois que voi	us avez bu de la liqueur ou des c	cocktails, quelle quantité en buviez-
vous habituellement?		

ı	Non		
Si oui, s'il vous plaît expliquer la raison:			
C.2. S'il vous plaît vérifiez si vous avez une des maladies suivantes et vous avez eu cette (ces) maladie(s).	pour con	nbien o	de temps
iste de maladies	Oui	Non	Depuis
Maladie osseuse			
Ostéoporose			
Diabète			
Problème de thyroïde			
Problème parathyroïdien			
Maladie hépatique (hépatite: virus A, B, C, cirrhose, etc)			
Maladie cardiaque (infarctus, angine, problèmes valvulaires)			
Problèmes sanguins (hémophilie, des saignements prolongés, anémie)			
Maladies vénériennes (V.D.)			
Maladie rénale			
Sclérodermie			
Arthrite, arthrose			
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires			
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale			vez pris
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois.			_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois.	G Oui	ous en a	vez pris
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole)	Oui	ous en a	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicaments dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax)	Oui	ous en a	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicaments dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne)	Oui	ous en a	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicaments dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine)	Oui	ous en a	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicaments dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa)	Oui	ous en a	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. Iiste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine) L'hormonothérapie (œstrogène, Contraceptifs)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine) L'hormonothérapie (æstrogène, Contraceptifs) Antibiotiques (Clindamycine, l'érythromycine)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine) L'hormonothérapie (æstrogène, Contraceptifs) Antibiotiques (Clindamycine, l'érythromycine) Psychothropes (Chlorpromazine, la phénothiazine, Carbonate de lithium)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. Iiste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine) L'hormonothérapie (œstrogène, Contraceptifs) Antibiotiques (Clindamycine, l'érythromycine) Psychothropes (Chlorpromazine, la phénothiazine, Carbonate de lithium) Les anti-hypertenseurs (diurétiques, bêta-bloquants, la méthyldopa)	Oui	Non	_
Arthrite, arthrose Maladie mentale Asthme, Problèmes pulmonaires Dyskinésie buccale C.3. Veuillez vérifier si vous prenez présentement un de ces médicament dans les 6 derniers mois. liste de médicaments Médicaments thyroïdiens (lévothyroxine, Méthimazole, Carbimazole) Les bisphosphonates (Fosamax) Antiépileptiques (Carbamazépine, Phénytoïne) Immuno-suppresseurs (Méthotrexate, l'azathioprine, Pénicillamine) Anticoagulants (Héparine, Warfarine, Pradaxa) Corticoïdes (Prednisolone, Bétaméthasone, l'hydrocortisone) Anti arthritiques (Colchicine) Anti allergiques (Chlorphéniramine) L'hormonothérapie (æstrogène, Contraceptifs) Antibiotiques (Clindamycine, l'érythromycine)	Oui	Non	_

C.4. Avez tumeur?		ou une radiothérapie pour traiter un cancer ou une	
Si oui, da	ans quelle partie?		
C.5. Faite	es-vous l'usage de drogues?	□ Oui □ Non	
Si oui,	depuis		
Section D: Uti	ilisation des services dentaires	ş.	
D.1. Const	ultez-vous souvent un professionnel d	dentaire?	
☐ Plus d' traitement	une fois par an pour un contrôle ou un t on une fois par an pour un contrôle ou t	nn	u
D. 2. Qua réponse)	and pensez-vous que vous avez besoin	n d'une visite chez le dentiste? (Veuillez cocher une	
☐ De 1 se	oins d'une semaine emaine à moins de 1 mois nois à moins de 3 mois	☐ De 3 mois à moins de 6 mois ☐ 6 mois ou plus ☐ Je ne sais pas	
D. 3. À c	quand remonte la dernière fois que vo	ous avez vu un professionnel dentaire?	
□ 1 an à i	oins de 1 an moins de 2 ans a moins de 3 ans	☐3 ans à moins de 4 ans ☐4 ans à moins de 5 ans ☐5 ans ou plus ☐ Jamais	
S'il vous p	plaît expliquer pourquoi ?		
D. 4. Com	nbien de fois avez-vous visité le denti	tiste durant les 12 derniers mois?	
D.5. Où a	eu lieu votre dernière visite dentaire	? (Veuillez cocher une réponse)	
☐ Pratic	que dentaire privée (y compris un spé	écialiste) ☐ École/Université dentaire / ☐ Hôpital	
☐ Dentui	rologiste	base militaire Autre site:	
D.6. Pour	quoi préférez-vous voir un dentiste p	privé? (S'il vous plaît cochez une réponse)	
☐ La qua clinique p ☐Autres_	oublique	attendre Le traitement n'est pas disponible à la blique accessible La continuité des soins	
		8	

D.7. Qui pourrait influencer votre décision de subir des soins ou des traitements dentaires? (S'il
vous plaît vérifiez les réponses qui s'appliquent:) ☐ Mon dentiste ☐ Mon hygiéniste ☐ Ma famille ☐ Moi
Section E: Habitudes d'hygiène buccodentaire
E.1. En moyenne, combien de fois par jour <u>brossez</u>-vous les dents (ou nettoyez-vous vos prothèses)? LILI Fois
E.2. En moyenne, combien de fois par semaine utilisez-vous <u>la soie dentaire</u> ?
☐ Fois ☐ Non applicable
E.3. En moyenne, combien de fois par semaine utilisez-vous <u>un bain de bouche</u> ?
E.4. Est-ce que votre eau du robinet est fluorée? 🗆 Oui 👚 Non 🗖 Ne sais pas
Section F: Anxiété dentaire
F.1. Si vous deviez vous rendre demain chez le dentiste, comment vous sentiriez-vous à ce sujet? J'aimerais qu'elle soit une expérience relativement agréable. Je ne me soucierais pas d'une façon ou l'autre. Je serais un peu inquiet à ce sujet. Je craindrais que ce soit désagréable et douloureux. J'aimerais très peur de ce que le dentiste pourrait faire F.2. Lorsque vous attendez votre tour au cabinet dentaire, comment vous sentez-vous? Détendu(e)

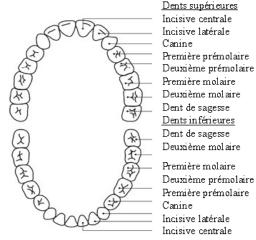
	s instruments do			oyer vos dents. Pendant que a pour nettoyer vos dents a					
☐ Détendu(e) ☐ Tellement anxieux malade.	☐ Un peu i (se) que je devid			Tendu(e) S de sueur ou me sens presqu	oucieux (se) ne physiquement	t			
Section G: Caractéristiqu	ue psychologiqu	ıe							
Voici un certain nombre d'é énoncé et encerclez le numé réponse juste ou fausse: Votre	éro approprié à	côté de	l'éno	ncé qui correspond à votr					
0=Faux 1=Plutôt f	aux 2=	Neutre	9	3= Plutôt vrai	4= Vrai				
G.1. J'ai un contact facile, que rencontre des gens.	uand je	0 1 2	3 4	G.8. Je trouve difficile de conversation.	e démarrer une	0 1	1 2	3	4
G.2. Je fais souvent des histochoses sans importance.	oires sur des	0 1 2	3 4	G.9. Je suis de mauvaise	humeur.	0 1	1 2	3	4
G.3. Je parle souvent à des é	etrangers.	0 1 2	3 4	G.10. Je suis une personn	e renfermée.	0 1	1 2	3	4
G.4. Je me sens souvent mal	lheureux.	0 1 2	3 4	G.11. Je veux plutôt distances avec les autres.	t garder mes	0 1	1 2	3	4
G.5. Je suis souvent irritable	÷.	0 1 2	3 4	G.12. Je me retrouve s pour quelque chose.	souvent inquiet	0 1	1 2	3	4
G.6. Je me sens souvent inh interactions sociales.	ibé (e) dans les	0 1 2	3 4	G.13. Je suis souvent dép	rimé (e).	0 1	1 2	3	4
G.7. J'ai une vision sombre	des choses.	0 1 2	3 4	G.14. Pour socialiser, comment débuter la conv		0 1	1 2	3	4
						10			

Section H: Histoire de la perte des dents

- H.1. Combien de dents avez-vous actuellement dans votre bouche (y compris les dents avec des couronnes dessus? Les dents du haut:
- H.2. Dans le schéma suivant, veuillez tracer un cercle autour des dents que vous avez actuellement en bouche et une croix (X) en travers des dents que vous avez perdues.

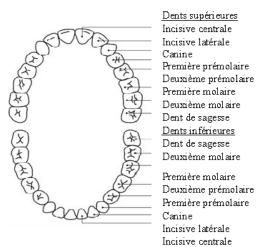
 Droit

 Gauche



H.3. Veuillez tracer un cercle autour des dents que vous avez perdues quand vous étiez entre 20-34 ans:

Droit

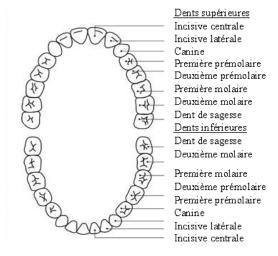


Gauche

11

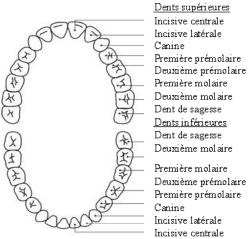
H.4. Veuillez tracer un cercle autour des dents que vous avez perdues quand vous étiez entre 35-49 ans:

Droit Gauche



H.5. Veuillez tracer un cercle autour des dents que vous avez perdues quand vous étiez entre 50-64 ans:

Droit



12

Gauche

H.6. Veuillez tracer un cercle autour des dents que vous avez perdues quand vous étiez ≥65 ans: Droit Gauche Dents supérieures Incisive centrale Incisive latérale Canine Première prémolaire Deuxième prémolaire Première molaire Deuxième molaire Dent de sagesse Dents inférieures Dent de sagesse Deuxième molaire Première molaire Deuxième prémolaire Première prémolaire Canine Incisive latérale Incisive centrale H.7. Combien de paires de dents naturelles postérieures (en arrière) dans le côté droit font contact pendant 1a mastication? 2 3 H.8. Combien de paires de dents naturelles postérieures (en arrière) dans le côté gauche font contact pendant la mastication? 2 3 5 Avant de répondre aux questions suivantes, veuillez prêter attention à ces définitions: Prothèse partielle: une prothèse amovible avec des dents artificielles qui remplacent certaines de vos dents naturelles perdues, à la mâchoire supérieure et / ou inférieure. Prothèse complète: une prothèse amovible avec des dents artificielles qui remplacent l'ensemble de vos dents naturelles perdues, à la mâchoire supérieure et / ou inférieure. 13

A . Â	D. Toma de deution	C: Mâchoire	D: Portez-vous les
A:Âge quand vous avez commencé à porter un dentier	B: Type de dentier	C: Macnoire	prothèses pour mastiquer
☐ Avant 20 ans	☐ Prothèse complète ☐ Prothèse partielle	☐ Supérieure ☐ Inférieure ☐ Les deux ☐ Non applicable	Oui Non
☐ Entre 20-34	☐ Prothèse complète ☐ Prothèse partielle	☐ Supérieure ☐ Inférieure ☐ Les deux ☐ Non applicable	Oui Non
☐ Entre 35-49	☐ Prothèse complète ☐ Prothèse partielle	☐ Supérieure ☐ Inférieure ☐ Les deux ☐ Non applicable	Oui Non
□ Entre 50-64	☐ Prothèse complète ☐ Prothèse partielle	☐ Supérieure ☐ Inférieure ☐ Les deux ☐ Non applicable	Oui Non
☐ 65 et plus	☐ Prothèse complète ☐ Prothèse partielle	☐ Supérieure ☐ Inférieure ☐ Les deux ☐ Non applicable	Oui Non

APPENDIX II : PROTHÈSE COMPLÈTE AMOVIBLE

Date a a m m j j		Numéro de dossier					
PROTHÈSES COMPLÈTES AMOVIBLES							
Histor	ique des Prothèses						
Avez-vous déjà porté des prothèses complètes?							
	Motif de consultation						
☐ Inconfort, blessures ☐ Besoin de dents ☐ Renouvellement de l'ancienne	□ Esthétique non-satisfaisante □ Mastication non-satisfaisante □ Rétention non-satisfaisante □ Inconfort, blessures □ Besoin de réparation □ Nouvelle prothèse suite à l'extraction des						
Attentes du patient concernant les nouvelles	prothèses:						
Évalu	ation de la prothèse actu	elle					
Espace libre d'inocclusion	☐ Adéquat (3-7mm)	☐ Inadéquate (>7mm or <3mm)					
DVO	☐ Acceptable	Inacceptable: □Réduite □Excessive					
Occlusion	□ Balancée	☐ Glissement (contact irrégulier)					
Rétention de la prothèse supérieure (résistance à la traction verticale)	☐ Résistance adéquate	☐ Pas de résistance					
Stabilité de la prothèse supérieure 1) Déplacement latéral	☐ Déplacement lat ≤5mm	☐ Déplacement lat >5mm					
2) Mvt. de bascule prononcé	☐ Mvt.de bascule minimal	☐ Mvt.de bascule prononcé					
Stabilité de la prothèse inférieure							
1) Déplacement, bouche ouverte	☐ Reste en place	☐ Déplacement perceptible					
2) Mouvement de la langue	☐ Pas de mouvement	☐ Mouvement prononcé					
3) Mvt. antéro-postérieur	☐ Mouvement minimal	☐ Mouvement prononcé					
		1					

Niveau d'hygiène 1) Combien de fois par jour/semaine vous nettoyez vos prothèses?	lfois par jour lfo	ois par semaine	
2) Comment nettoyez-vous vos prothèses?	☐ Rinçage ☐ Brossage ☐ Tremper dans un nettoya	ant pour dentiers	
3) Retirez-vous vos prothèses pendant la nuit?	Prothèse Supérieure	*1 —	Oui Non
4) Utilisez-vous un rince- bouche?	□Oui ☐ fois par jour ☐	☐ Non	
	TISSUS M	IOUS	
	EXAMEN	ORAL	
Visag	ge	Longueur de la Salive	
Symétrie Facial from Symétrique Triangu Asymétrique Carré Ovoïde Autres	alaire	Favorable Moyenne Défavorable Classe II (insuf Classe III (excellent) Classe III (exc	ffisante)
Voile du pa	ılais	Insertions freinales	
Favorable	Défavorable	Favorable Défavorable	
□ Classe I (rétention idéal	-	☐ Classe I (idéal) ☐ Classe III (requie	ert une
☐ Classe II (rétention bon	ne) (rétention faible)	☐ Classe II (moyen) correction chirur	
			2

			Lang	ue					
Positi	on de la lan	gue			Taille de l	a langue			
Favoi					Favorable □ Classe I	e (normale)			
□ Clas	v orable sse I (rétrude sse II (tendu lée vers le ha	e tirée vers l'arrière et				ble II (changement d III (excessiveme		e fonction)	
		Insertions musculai	ires mandi	ibulaire	es		Contr	ôle	
□ Type A (insertions adéquates) □ Type B (insertions adéquates exception mentonnier proche de la crête alvéolaire) □ Type C (insertions adéquates exception mentonnier et génioglosse proches de la crête alvéolaire) □ Type D (ins.muq. en post seulement) □ Type E (Pas d'ins.muq., joue/lèvre déplace la langue)									
		États de la muque	euse			Ту	pe de muqu	euse	
Code	•	Description		Max	Mand	Туре	Max	Mand	
1		omalie de la muqueus	e			Mobile:			
2	Perlèche	1 1 1	_				_	_	
3	Plaques bi	anches de la muqueuse	e			a) Antérieu	re 🗆		
	Stomatite	41.74				h) Poetériei	ıra 🗆		
	Degré 0: n	ion de Bergendal et Isa nuqueuse normale (ros				b) Postérier Non mobil			
	Degré 0: n Degré 1: é	ion de Bergendal et Isa				Non mobil Épaisse			
4	Degré 0: n Degré 1: é Degré 2: é Classifica	on de Bergendal et Isa nuqueuse normale (ros rythème léger				Non mobil Épaisse Inflammée	e 🗆		
4	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: Classe III:	on de Bergendal et Isa nuqueuse normale (ros rythème léger rythème sévère tion de Newton	sée isée ou			Non mobil Épaisse	e		
	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: L Classe III: localisée a	on de Bergendal et Isa- nuqueuse normale (ros rythème léger rythème sévère tion de Newton nflammation localisée Inflammation généralis Inflammation généralis	sée isée ou			Non mobil Épaisse Inflammée	e -		
5	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: Classe III: localisée a Ulcération	on de Bergendal et Isa- nuqueuse normale (ros rythème léger rythème sévère tion de Newton nflammation localisée Inflammation généralis Inflammation généralis vec hyperplasie papilla	sée isée ou aire			Non mobil Épaisse Inflammée	e -		
5 6	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: L Classe III: localisée a Ulcération Hyperplasie	on de Bergendal et Isac nuqueuse normale (ros rythème léger rythème sévère tion de Newton nflammation localisée Inflammation généralis Vec hyperplasie papilla aphteuse e d'origine prothétique (d	sée isée ou aire			Non mobil Épaisse Inflammée	e -		
5 6 7	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: Classe III: localisée a Ulcération Hyperplasie Hyperkérat	on de Bergendal et Isa- nuqueuse normale (ros- rythème léger rythème sévère tion de Newton inflammation localisée inflammation généralis vec hyperplasie papilla aphteuse e d'origine prothétique (dosse	sée isée ou aire épulis)			Non mobil Épaisse Inflammée	e -		
5 6	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: Classe III: localisée a Ulcération Hyperplasie Hyperkérat	on de Bergendal et Isac nuqueuse normale (ros rythème léger rythème sévère tion de Newton nflammation localisée Inflammation généralis Vec hyperplasie papilla aphteuse e d'origine prothétique (d	sée isée ou aire épulis)			Non mobil Épaisse Inflammée	e -		
5 6 7 8	Degré 0: n Degré 1: é Degré 2: é Classifica Classe II: Classe III: localisée a Ulcération Hyperplasie Hyperkérat Ulcération Glossite	on de Bergendal et Isa- nuqueuse normale (ros- rythème léger rythème sévère tion de Newton inflammation localisée inflammation généralis vec hyperplasie papilla aphteuse e d'origine prothétique (dosse	sée isée ou aire épulis)			Non mobil Épaisse Inflammée	e -		

		TISSUS OSSEUX	
Tubéro	osité	Forme du Palais dur	Rapport inter-crêtes
Présente Rétentiv Non réte	ntive	 □ Plat □ Rond □ En forme de V □ En forme de U □ Profond 	☐ Classe I (normal) ☐ Classe II ☐ Class III
		Espace inter-arche	
□ Cla	asse II (limité) prrection chirur	gicale est-elle indiquée? 🗆 🗅 🕻	Oui □ Non
Mandibule		Si présent, une réduction chirurgindiquée? □ Oui	gicale est-elle □ Non
Maxillaire		ATM	<u> </u>
Symptômes Symptomatique du visage ou de la range de tête ou de la range de la	mastication □ I des migraines	Grincement 🗆 Claquement 🗆 Fa	atigue ou de la tension au niveau des muscles ors de l'ouverture ou de fermeture buccale

Hauteur de la crête	1	г			7								
Résiduelle		Résorption	1 de l'os a	dvéolaire	e								
☐ Type I: 21mm ou plus	Mandibule			Maxilla	aire								
☐ Type II: 16-20mm	□ Dentée			□ Der									
☐ Type III: 11-15mm	□ Post extrac		on Larrondie)										
☐ Type IV: 10mm ou moins	☐ Classe 4 (c		rête en lame de 🔲 Classe 4 (peu de perte en										
1 Type IV. Tollan ou mells	couteau)	crête plate)			geur) a ss e 5/6 (p	erte en largeur et							
	☐ Classe 6 (c	crête de form	e		hauteur)	-							
	négative)												
Quantitá d'as t		ıen radiologi		1444 41	/1 A)								
Quantité d'os (A-E)		- V	ualité d'	os (1-4)								
777													
à B C	D E		1	2	3	4							
0 0 0	0 0												
A B	C D E		1	2	3	4							
Max 🗆 🗆		□ Max											
Mand 🗆 🗆		□ Mand											
	<u> </u>												
	Prothèses impla	nto-portées											
Prothèses actuelles adéquates		que et chirur	gical:	□ Oui	Ĺ	□ Non							
Hauteur osseuse disponible: L													
Distance suffisante entre les d	eux trous mentonnies	rs:	□ Oui		□ Non	ı							
Nombre d'implants souhaités:		Type d'imp	lants:										
Longueur: LL mm		Emplacemen	nt:										
													

	PLAN DE TRAIT	EMENT		
	A: pré-prothétique	2		
Conditions nécessitant une chi	rurgie pré-prothétique			
□ Extraction□ Interventions mineures su□ Chirurgie implantaire	ır les tissus mous			
	B: prothétique			
Fraitement suggéré (Veuillez c	hoisir plus de un, si plusieur	s traitements sont	nécessaires)	
	C/C □ C/		Prothèse-impla	nto-portées
-			Troniese imp	nto portees
□ Regarnissage/ Rebasage	☐ Réparation ☐ Imi	médiates		
	PRONOSTI	(C		
	To de disconnecto	1		
	Facteurs influençants		Favorable	Défavorable
Facteurs anatomiques Maladie systémique sous-jacent	2			
Considération orofaciale (sympt				
Facteurs psychosociaux Antécédents prothétiques				
En général, le pronostic est Max	Bon □	Réservé		Pauvre
	_			
Mand				
Mand				
Mand Signature du patient:		Date:		
Mand				

Compagnie	Quantité	Moule	Couleur	Résine acrylique	Porcelaine	Deg
Sup. Ant.						
Inf. Ant.						
Post. Inf.						
Post. Sup.						
Date:		Signat NSENTEI		ent:		_
Je comprends les a proposées et je com Je comprends que l traitements supplém Je comprends que l en ce sens, ils peuve	prends aussi les ind e prix ci-dessus en entaires sont requi	convénients de st une estimat s par ma cond ffectué par les	e ne faire auc ion du coût ition pendan s étudiants de	un traitement. réel et peut ê t le traitement.	tre modifié si	des
		•		et à acquitter	les frais enco	urus
Je m'engage à respe par le traitement de						

APPENDIX III: PROTHÈSE PARTIELLE AMOVIBLE

Date				Numéro de dossier								
	/											
y y m m	d d											
PROTHÈSE PARTIELLE AMOVIBLE												
	Histo	rique des	Prothèses									
Avez-vous déjà porté des	prothèses ?	□ Oui		Non								
Quel type:	□ P /		□ / P	□ P / P								
Nombre de prothèses: L Depuis combien de temps	I : └──│└─── ans											
Âge de la (des) prothèse (s	s) actuelle (s): Max: L			ns								
Cause de la perte dentaire	□ Maladies	parodontal	es 🗆 Caries	☐ Traumatisme	□ A	utres						
	Clas	ssification	de Kennedy									
		MAND			MAX	MAND						
Classe I			Classe IV									
Classe II			Classe V									
Classe III Pour toute modification, v	zouillaz datarminar la r	numára at 1	Classe VI	Postáriouro)do mod	lificatio	n?						
MAX: Numéro:		vpe :	• •	rosterieure)de mod	umcano	11 (
MAND: Numéro:	•											
	*.	ур										
	□ Besoin de réj nent de l'ancienne pro	paration thèse sans _l	□ Nouvelle	non-satisfaisante prothèse suite à l'e	extractio	n des						
Attentes du patient conce	ernant les nouvelles pro	othèses:										
	Évaluatio	n de la p	rothèse actuelle	e								
Rétention de la prothèse	☐ Excellente (résist au déplacement vert	tical) rés	Bonne (une istance modérée au blacement vertical)			ment						
Stabilité de la prothèse	☐ Excellente (stable mouvement de rotat et les appuis occlusa sont ajustés)	tion cer	☐ Bonne (stable, mais certains appuis sont pas ajustés) ☐ Mauvaise (déplacée provide mouvement rotationnel)									
Châssis métallique	□ Bien-ajusté	app	Adaptation des ouis occlusaux et chets inacceptable	☐ Fracturé								
Adaptation de la base acrylique à la muqueuse	☐ Bien-ajustée		Légère imprécision	n 🗖 Adaptation	n inacce	ptable						
						1						

EXAMEN ORAL

Évaluation des dents piliers

Veuillez enregistrer chaque élément pour toutes les dents piliers selon l'annexe. Veuillez arrondir chaque mesure au plus bas millimètre entier.

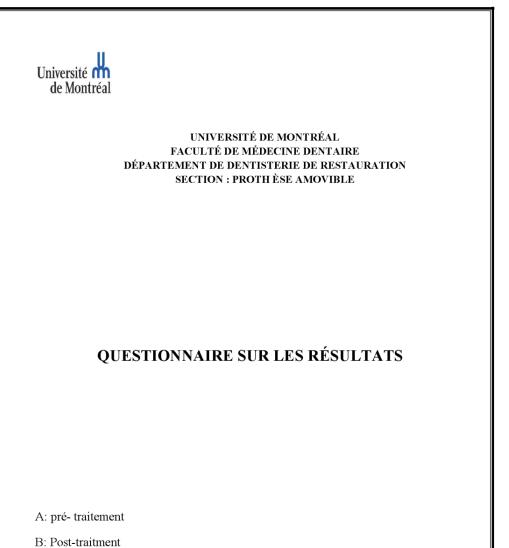
Max Dent pilier			•															-
	MB	В	DB	MB	В	DB	MB	В	DB	MB	В	DB	MB	В	DB	МВ	В	DB
	ML	L	DL	ML	L	DL	ML	L	DL	ML	L	DL	ML	L	DL	ML	L	DL
IP																		
PP																		
SAS																		
	M	В	D	M	В	D	M	В	D	M	В	D	M	В	D	M	В	D
Carie	L		o	L		O	L		o	L	L,	0	L		0	L		o
											Н			_				
Mobilité Relation Couronne/ Racine	□Acc □Ina			□Acc			□Acc			□Acc		table ptable	□Ac. □Ina		table ptable			table ptable
Mand Dent pilier								_			_	-					_	
Dent piner	МВ	В	DB	МВ	В	DB	МВ	В	DB	MB	В	DB	M B	В	DB	МВ	В	DB
				ML	L	DL	ML	L	DL	ML	L	DL	M	L	DL	ML	L	DL
	ML	L	DL				MIL	_					L	_		I NIL		
IP	ML	L	DL				ML						L			ML		
IP PP	ML	L	DL										L			ML		
	ML	L	DL										L			MIL		
PP SAS	M	В	D	M	В	D	M	В	D	M		D	M	В	D	M	В	D
PP		В			В								L	В	D O			D O
PP SAS	M	В	D	M	В	D	M		D	M		D	M	В		M		
PP SAS	M	В	D	M	В	D	M		D	M		D	M	В		M		
PP SAS Carie	M	B	D O	M	eepta	D O	M	B	D O	M L	В	DO	M L	В	0	M L	В	

En général, l'hygiène orale est En général, le taux des caries e			i Moyenne □ Modéré	□ Mauvaise □ Faible
	Évaluation des de	ents piliers		
Structure des dents piliers □ État du pilier idéal ou peu c	ompromis. (Pas de ti	raitement pré p	rothétique indiqué	.)
☐ État du pilier modérémen insuffisante pour retenir ou sup				
☐ État du pilier considérab insuffisante pour retenir ou su considérables.)				
☐ État du pilier sévèrement insuffisante pour retenir ou su Piliers ont un pronostic réservé	ipporter des restaui			
	Occh	usion]	
Plan occlusal		DVO		
□ Adéquat□ Un traitement adjuvant le	ocalisé (meulage)	☐ Adéquat ☐ À modif	e ier : □Augmentée	□ Réduite
Rétablir entièrement le plan	occlusal	8 4	47. •	
□ Classe I	□II	Support pos Adéquat Inadéqua		
	Malformatio	ns de la crête		
☐ Classe I (Perte vestibulo-lin	guale du contour tiss	sulaire avec un	e hauteur apico-co	ronaire normale)
_ Classe I (I cite vestibulo-illi			_	
☐ Classe II (Perte tissulaire ap	ico-coronane avec t			

	PLAN	DE TRA	ITE	ME	NT					
	A	: pré-proth	étiqı	ue						
Conditions nécessitant un traiteme Traitement endodontique Traitement de restauration: Traitement parodontal: Conditions nécessitant un traiteme Extraction Interventions mineures sur le	□ D.O (□ Soins : ent pré-p s tissus m	dentisterie op hygiéniques (prothétique c	ératoi détart hirur	re) rage			□ Courom □ Chirurgi			
Traitement Suggéré (Veuillez choisir plus de un, si plusieurs traitements sont nécessaires) □ Nouvelles prothèses : □ P/P □ P/ □ /P □ Prothèse-implanto-portée □ Regarnissage □ Rebasage □ Réparation										
	C	lassification of	de Ke	nnedy						
Classe I	MAX	MAND	Clas	se IV			MAX	MAND		
Classe III			Clas	se V se VI						
Pour toute modification, veuillez det MAX: Numéro: MAND: Numéro:	terminer l	e numéro et l Type: Type:				stérieure)de m	odificatio	on?		
		PRONO	STI	С						
	Fact	eurs influenç	ants			Favorable	Défav	orable		
Facteurs anatomiques Maladie systémique sous-jacente Considération orofaciale (symptô Facteurs psychosociaux Antécédents prothétiques Hygiène orale	mes d'A'	ГМ)				0 0 0 0				
En général, le pronostic est Max Mand		Bon		I	Réservé		Pauv	re		
					<u> </u>			4		

Signature du patient: Signature de l'étudiant: Signature du clinicien:			Date: Date: Date:					
	CHOIX DES DENTS							
ompagnie	Quantité –	Moule	Couleur	Résine acrylique	Porcelaine	Degré		
Sup. Ant.								
Inf. Ant.								
Post. Inf.								
Post. Sup.								
	es avantages et les in comprends aussi les i					ement		
	ue le prix ci-dessus démentaires sont requ			•		si des		
	ue le traitement sera euvent avoir besoin d			dans le cadre	e de leur format	ion et		
Je m'engage à r	especter les rendez-v	•		és et à acquit	ter les frais enc	ourus		
par le traitement	Date: Signature du patient:							
•			Signature o	iu patient				
•			Signature o	ти рацени				

APPENDIX IV: QUESTIONNAIRE SUR LES RÉSULTATS



Numéro de dossier

1

ion A: Qualité de vie					
A.1. En général, diriez-vous que votre santé est?					
☐ Excellente ☐ Très bonne ☐ Bonne		Juste		☐ Maı	ıvaise
A.2. En général, diriez-vous que la santé de votre bouche est	•				
☐ Excellente ☐ Très bonne ☐ Bonne		Juste		□ Mau	vaise
A.3. Questions sur la qualité de vie reliée à la santé bucco-de	ntaire	:			
The Quarter on the quarter of the control of the co					
Sur une échelle de 0 à 4, veuillez encerclez le numéro qui convient le mieux à votre réponse (veuillez encercler un).	Jamais	Presque jamais	Occasionnellemen	Assez souvent	Très souvent
1. Avez-vous éprouvé de la difficulté à mastiquer des aliments à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?		1	2	3	4
2. Les aliments sont-ils restés coincés entre vos dents ou dans vos	0	1	2	3	4
prothèses ? 3. Avez-vous eu l'impression que vos prothèses étaient mal ajustées?	0	1	2	3	4
4. Avez-vous eu de la douleur au niveau de la bouche ?	0	1	2	3	4
5. Avez-vous éprouvé de la difficulté à consommer certains types d'aliments à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
6. Avez-vous remarqué des points sensibles dans votre bouche?	0	1	2	3	4
7. Vos prothèses ont-elles été inconfortables ?	0	1	2	3	4
B. Vous êtes-vous fait du souci à cause de problèmes buccaux ?	0	1	2	3	4
9. Vous êtes-vous senti(e) mal à l'aise à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
10. Avez-vous évité de consommer certains types d'aliments à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses?		1	2	3	4
11. Avez-vous été incapable de manger avec vos prothèses à cause de problèmes avec celles-ci?	0	1	2	3	4
12. Avez-vous dû interrompre un repas à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
13. Avez-vous été perturbé à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses?	0	1	2	3	4
14. Avez-vous été légèrement incommodé(e) à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
15. Vous êtes vous abstenu(e) de sortir à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
16. Vous êtes-vous senti(e) plus intolérant(e) envers votre famille ou votre conjoint(e) à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4

17. Avez-vous été irritable au milieu d'un groupe à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos	0	1	2	3	4
problemes dus a votre defittion, a i etat de votre bouche ou a vos					
18. Avez-vous été incapable d'apprécier la compagnie des autres à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4
19. Avez-vous pensé que la vie était généralement moins satisfaisante à cause de problèmes dus à votre dentition, à l'état de votre bouche ou à vos prothèses ?	0	1	2	3	4

A.4. Durant les 12 derniers mois, vous absentiez-vous du travail ou de vos activités normales en raison de la nécessité d'avoir des soins dentaires, y compris des examens dentaires, ou à cause de problèmes au niveau de votre bouche?

🗖 Oui .	Heures (MIN: 0.5)	(MAX: 99.5)	☐ Non
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Section B: La satisfaction du patient

Sur une échelle de 0 à 4, veuillez encerclez le numéro qui convient le mieux à votre réponse (veuillez encercler un).	Très satisfait	Satisfait	Ni satisfait ni insatisfait	insatisfait	Très insatisfait
B.1. En général, comment est votre satisfaction concernant votre prothèse supérieure?	0	1	2	3	4
B.2. En général, comment est votre satisfaction concernant votre prothèse inférieure?	0	1	2	3	4
Rétention B.3. Comment est votre satisfaction concernant la rétention de votre prothèse supérieure?	0	1	2	3	4
B.4. Comment est votre satisfaction concernant la rétention de votre prothèse inférieure?	0	1	2	3	4
Confort B.5. Comment est votre satisfaction concernant le confort fonctionnel de vos prothèses?	0	1	2	3	4
Esthétique B.6. Comment est votre satisfaction concernant l'aspect de vos prothèses?	0	1	2	3	4
Aptitude à parler B.7. Comment est votre satisfaction concernant la parole avec vos prothèses?	0	1	2	3	4

Sur une échelle de 1 à 4, veuillez encerclez le numéro qui convient le mieux à votre réponse (veuillez encercler un).	Souvent	Fréquemment	Occasionnellement	Rarement
Stabilité B.8. Ressentez-vous un déplacement de la prothèse supérieure en mangeant et/ou en parlant?	1	2	3	4
B.9. Ressentez-vous un déplacement de la prothèse inférieure en mangeant et/ou en parlant?	1	2	3	4
Aptitude à mordre B.10. Avez-vous des problèmes à mordre avec vos prothèses?	1	2	3	4
Aptitude à mâcher B.11. Avez-vous des problèmes à mâcher avec vos prothèses?	1	2	3	4
B.12. Trouvez-vous des particules alimentaires sous vos prothèses?	1	2	3	4
B.13 Ressentez-vous une différence de goût des aliments à cause de vos prothèses?	1	2	3	4
B.14. Est-ce que la prothèse influence votre choix des aliments?	1	2	3	4
Sur une échelle de 1 à 4, veuillez encerclez le numéro qui convient le mieux à votre réponse (veuillez encercler un).	Souvent positive	Positive	Négative	souvent négative
Hygiène Prothétique B.15. Avez-vous de la difficulté à nettoyer vos prothèses?	1	2	3	4
B.16. Ressentez-vous une mauvaise haleine à cause de vos prothèses?	1	2	3	4
B.17. Ressentez-vous une sécheresse buccale?	1	2	3	4
B.18. Avez-vous des problèmes d'hyper sialorrhée (hyper salivation) ?	1	2	3	4

Section C: Attentes de satisfaction à l'égard de la prothèse totale

C.1. Quelle serait votre satisfaction avec de nouvelles prothèses conventionnelles?			
0 ————	100		
0= Pas du tout satisfait	100= complètement satisfait		

4

ec	ction D: Satisfaction par la visi	te dentaire			
	Information- communication				
	D.1. Après avoir discuté avec le d	entiste/étudiant (e), □ En désaccord	J'ai pris conna ☐ Incertain	nissance de mo	n état buccal. ☐ Fortement d'accord
	D.2. Après avoir discuté avec le seraient apportées à mon état de s		· // U		odifications qui
	☐ Fortement en désaccord	☐ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	D.3. Le dentiste/étudiant (e) m'a mon (mes) problème(s) dentaire(s	* *	le tout ce que	je désirais sav	oir à propos de
	☐ Fortement en désaccord	☐ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	Compréhension-acceptation				
	D.4. Je me sentais vraiment comp	ris par mon dentiste	e/l'étudiant (e).		
	☐ Fortement en désaccord	☐ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	D.5. J'ai senti que ce dentiste/étu possibilité de la douleur.	ıdiant (e) savait vr	aiment comme	nt j'ai été cont	trarié (e) par la
	☐ Fortement en désaccord	☐ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	D.6. J'ai senti que ce dentiste/étud Fortement en désaccord Compétence technique	diant (e) m'a accept ☐ En désaccord	é en tant que po ☐ Incertain	ersonne. D'accord	☐ Fortement d'accord
	D.7. Le dentiste/étudiant (e) a été	complet en faisant	la procédure.		
	☐ Fortement en désaccord	☐ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	D.8. Le dentiste/étudiant (e) était Fortement d'accord		me soignait. rtain □ En d	ésaccord □	Fortement en désaccord
	D.9. J'ai été satisfait (e) de ce que	le dentiste/étudiant	(e) a fait.		
		□ En désaccord	☐ Incertain	☐ D'accord	☐ Fortement d'accord
	`	e) semblait savo □ En désaccord	oir ce qu'il □ Incertain	faisait dura	nt ma visite □ Fortement d'accord
					5

APPENDIX V: ANNEXE PROTHÈSE COMPLETE AMOVIBLE

ANNEXE: PROTHÈSE COMPLETE AMOVIBLE

Évaluation fonctionnelle des prothèses dentaires 1-4

Espace libre d'inocclusion

L'espace entre les surfaces occlusales des dents maxillaires et mandibulaires lorsque la mandibule est en position de repos physiologique.

La dimension verticale d'occlusion (DVO)

la dimension verticale d'occlusion est la hauteur de l'étage inférieur de la face lorsque les dents des maxillaires supérieur et inférieur sont en intercuspidation maximale.

Occlusion

Le patient est prié de se détendre et de fermer doucement sur ces dents postérieures plusieurs fois à partir d'une position de légère ouverture (20 mm). l'occlusion est jugée satisfaisante (quand il y a un contact régulier entre les dents et un retour compatible avec la position d'intercuspidation) ou insatisfaisante (quand il y a un contact irrégulier entre les dents et un retour non conforme avec la position d'intercuspidation ou un glissement supérieur à 4 mm).

Rétention de la prothèse supérieure

(Résistance à la traction verticale): l'ouverture buccale est de 20 mm. Notez si la prothèse tombe. Pendant que la bouche est encore ouverte, la prothèse est saisie par le pouce et l'index au niveau des prémolaires et une force descendante est appliquée. La présence ou l'absence d'une résistance appropriée est enregistrée.

Stabilité de la prothèse supérieure

(Le déplacement latéral): La stabilité en rotation est évaluée en saisissant la prothèse supérieure dans la région prémolaire avec le pouce et l'index et en appliquant une force de rotation dans le plan d'occlusion. Nous considérons le déplacement de 5 mm ou moins (environ 2,5 mm de chaque côté de la ligne médiane) comme satisfaisante.

(Basculement prononcé): une force légère est appliquée sur les premières molaires simultanément aux côtés droit et gauche. Essayez de pencher en direction antéro-postérieure avec le pouce et l'index placés en arrière et en avant simultanément et jugez de la présence ou l'absence d'un mouvement de bascule prononcé.

Stabilité de la prothèse inférieure

(Déplacement): L'ouverture buccale est de 20 mm et la langue est en position de repos. L'assise prothétique est vérifiée avec les doigts.

(Mouvement prononcé): Le patient est invité à bouger sa langue de manière que la pointe repose doucement au niveau des angles de la bouche, celle-ci est ouverte de 20 mm. Vérifiez l'assise de la prothèse avec les doigts. Le jugement est fait au troisième essai.

(Le mouvement antéropostérieur): la prothèse supérieure est déposée. La prothèse inférieure est maintenue sur la crête avec un doigt et le pouce sur les incisives et une tentative est faite pour déplacer la prothèse avec la langue en position de repos.

Niveau d'hygiène ⁵

TISSUS MOUS:

Longueur de la lèvre supérieure⁶⁻⁹

Favorable: Normale

Défavorable:

A) Longue: une lèvre longue découvre très peu les dents antérieures.

B) Courte: une lèvre supérieure courte affiche la base prothétique

Salive⁷⁻¹⁰

Classe I: Normale en quantité et en qualité.

Classe II: Insuffisante, xérostomie.

Classe III: Abondante et très visqueuse.

Voile du palais^{6, 9, 10}

<u>Classe I</u>: Idéal pour la rétention. (le voile du palais est presque horizontal. Plus de 5mm de tissus mobiles disponibles pour l'endiguement postérieur).

<u>Classe II</u>: Une bonne rétention est généralement possible. (le voile du palais s'incline vers le bas en formant un angle de 45⁰ avec le palais dur. 1 à 5 mm de tissus mobiles disponibles pour l'endiguement postérieur).

<u>Classe III</u>: La rétention est généralement faible. (le voile du palais s'incline vers le bas en formant un angle de 75⁰ avec le palais dur. Moins de 1 mm de tissus mobiles disponibles pour l'endiguement postérieur).

Insertions freinales¹⁰

Classe 1: Haute au maxillaire ou basse à la mandibule tout en respectant le sommet de la crête.

Classe 2: Moyenne

<u>Classe 3</u>: Le frein empiète sur le sommet de la crête et peut interférer avec le joint prothétique.

Une correction chirurgicale peut être nécessaire.

Langue

Position de la langue (Classification selon Wright)

Normale: La langue occupe le plancher buccal et est délimitée par les dents mandibulaires.

Classe I: La langue est rétractée. Le plancher de la bouche est tiré vers le bas et s'expose en arrière dans la région molaire.

Classe II: la langue est très tendue, tirée vers l'arrière et enroulée vers le haut.

Taille de la langue¹⁰ (Classification selon House)

<u>Classe I</u>: Taille, développement et fonction normaux; les dents sont présentes en nombre suffisant pour maintenir une forme et une fonction normales.

Classe II: Les dents ont été absentes assez longtemps pour entrainer un changement de forme et de fonction.

Classe III: Langue excessivement large. Toutes les dents ont été absentes pour une longue période autorisant un développement anormal de la taille de la langue.

Insertion musculaire à la mandibule¹¹

Type A: Base d'insertion muqueuse sans empiètement musculaire excessif durant le fonctionnement normal dans toutes les régions.	Type B: Base d'insertion muqueuse dans toutes les régions excepté au vestibule labial. L'insertion du muscle mentonnier est proche du sommet de la crête alvéolaire.
Type C: Base d'insertion muqueuse dans toutes les régions excepté les vestibules antérieurs buccal et lingual de canine à canine. L'insertion du muscle génio-glosse et du muscle mentonnier est proche du sommet de la crête alvéolaire.	Type D: Base d'insertion muqueuse uniquement dans la région linguale postérieure. La base muqueuse dans toutes les autres régions est détachée.
Type E: Pas de muqueuse at	achée dans toutes les régions

État de la muqueuse¹²

Code	Description	Définition
1	Pas d'anomalies de la muqueuse	
2	Chéilite angulaire	Inflammation et fissures au niveau des angles ou des commissures labiales
3	Plaques blanches de la muqueuse	Zones blanches sur la muqueuse tel que le lichen plan, leucoplasie, candidose ou muguet.
4	Stomatite prothétique Classification de Bergendal and Isacsson ^{13, 14}	
	<u>Degré 0</u> : muqueuse rosâtre normale	
	Degré 1: érythème léger	
	<u>Degré 2</u> : érythème sévère	
	Classification de Newton ¹⁵	
	<u>Classe I</u> : Hyperémie localisée: des zones d'inflammation localisées dans un tissu palatin normal, qui se trouvent généralement autour des orifices des conduits des glandes muqueuses palatines.	Inflammation sous-prothétique
	<u>Classe II</u> : Hyperémie diffuse: inflammation généralisée de la région de support de la prothèse dentaire.	
	<u>Classe III</u> : Granulaire : surface palatine hyperplasique, qui pourrait être généralisée ou restreinte aux régions médianes.	
5	Ulcération aphteuse	Ulcération qui est passagère et douloureuse
6	Hyperplasie d'origine prothétique (épulis)	Tissu hyperplasique associé au bord prothétique
7	Hyperkératose	L'épaississement d'une zone de l'épithélium de la bouche
8	Ulcération traumatique ou non spécifique	Ulcération d'origine inconnue incluant les lésions traumatiques.
9	Glossite	Inflammation de la langue. Ceci devrait exclure la langue géographique.
10	Autres-Précisez	États de la muqueuse qui ne sont pas mentionnés dans ce tableau (fistule, sinus,)

TISSU OSSEUX

Relation maxillo- mandibulaire 11

<u>Classe I</u> (la plus favorable): La relation maxillo- mandibulaire autorise une position dentaire en occlusion normale, les dents étant supportées par la crête résiduelle.

<u>Classe II</u>: La relation maxillo-mandibulaire requiert une position dentaire en dehors du rapport inter crête normal pour pouvoir assurer l'esthétique, la phonétique et l'occlusion. (Ex: les dents antérieures ou postérieures ne sont pas supportées par la crête résiduelle; le chevauchement antérieur vertical et/ou horizontal dépasse les critères de l'occlusion entièrement balancée).

<u>Classe III</u>: La relation maxillo-mandibulaire requiert une position dentaire en dehors du rapport inter crête normal pour pouvoir assurer l'esthétique, la phonétique et l'occlusion. (Ex. Les dents en articulé croisé antérieur ou postérieur ne sont pas supportées par la crête résiduelle).

Espace inter-arche^{7, 8}

Favorable:

Classe I: Espace inter-arche idéal

Défavorable:

Classe II: Espace inter- arche excessif entrainant de faibles stabilité et rétention prothétiques.

<u>Classe III</u>: Espace inter-arche limité. Un espace inter-arche insuffisant pour accueillir les dents artificielles renforce la stabilité des prothèses tant que la surface occlusale des dents est proche de la crête en réduisant l'effet de levier, mais diminue la rétention.

ATM^{16, 17}

Symptômes

Êtes-vous conscient de serrer des dents?

Êtes-vous conscient de grincer des dents?

Ressentez vous de la fatigue ou de la tension au niveau des muscles du visage ou de la mastication?

Ressentez-vous la douleur lors de l'ouverture ou de fermeture buccale?

Avez-vous de maux de tête ou des migraines ?

Signes

Claquement et /ou crépitation durant le mouvement de la mâchoire.

Limitation de l'ouverture buccale

palpation latérale et postérieure de l'ATM en mouvement.

Déviation de la mandibule sur ouverture de la bouche

Hauteur de la crête résiduelle¹¹

<u>Type I</u>: 21mm ou plus (la plus favorable), la hauteur de l'os résiduel est de 21mm mesurée à la plus faible hauteur verticale de la mandibule.

<u>Type II</u>: 16-20mm, la hauteur de l'os résiduel est de 16 à 20 mm mesurée à la plus faible hauteur verticale de la mandibule.

<u>Type III</u>: 11-15mm, la hauteur de l'os résiduel est de 11 à 15 mm mesurée à la plus faible hauteur verticale de la mandibule.

<u>Type IV</u>: 10mm ou moins, la hauteur de l'os résiduel est de 10 mm ou moins, mesurée à la plus faible hauteur verticale de la mandibule.

Résorption de l'os alvéolaire 18-21

Mandible

Classe1:Dentée

Classe2: Post extraction

<u>Classe 3</u>: Forme bien arrondie de la crête avec peu de perte de hauteur et de largeur.

<u>Classe 4</u>: Perte de largeur, mais une bonne hauteur- " crête en lame de couteau ".

<u>Classe 5</u>: Perte de hauteur et de largeur. Le niveau de la crête est au plancher buccal. (crête de forme plate)

<u>Classe 6</u>: Perte de hauteur et de largeur avec une perte de l'os basal. Crête en dessous du niveau du plancher buccal. (crête de forme négative)

Maxillaire

Classe 1:Denté

Classe 2: Post extraction

<u>Classe 3</u>: Forme bien arrondie de la crête. La papille rétro incisive est au palatin de la crête, le reste de la gencive palatine est au palatin de la crête osseuse postérieure.

<u>Classe 4</u>: quelque perte en largeur. la papille incisive est sur la crête ainsi que le reste de la gencive palatine.

<u>Classe 5/6</u>: perte de hauteur et de largeur. La papille est dans le sulcus labial et le reste de la geneive palatine est dans le sulcus buccal.

Examen radiologique 22-24

Quantité d'os

- A: La majorité de l'os alvéolaire est présent.
- B: La résorption de l'os alvéolaire est modérée
- <u>C</u>: La résorption de l'os alvéolaire est avancée . Seulement l'os basal est présent.
- $\underline{\mathbf{D}}$: Faible résorption de l'os basal.
- <u>E</u>: Résorption extrême de 1'os basal.

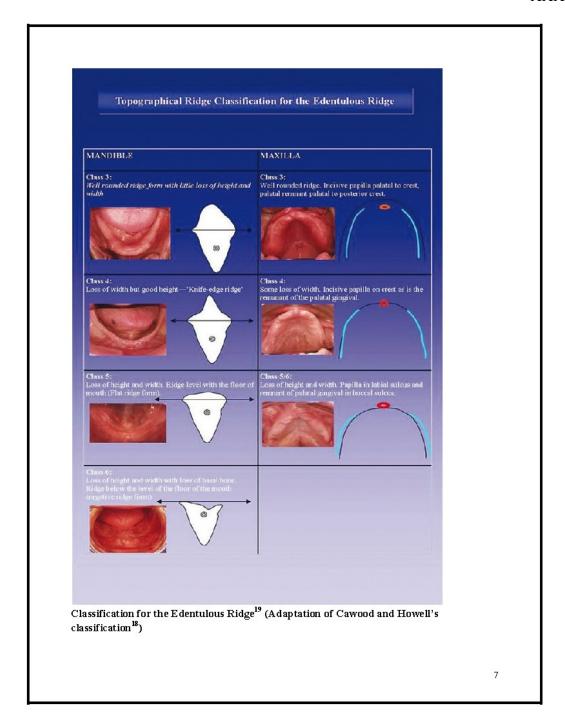
Qualité de l'os

<u>Qualité 1</u>: Presque toute la mâchoire est composée d'os compact homogène.

Qualité 2: Une couche épaisse d'os compact entoure un noyau d'os trabéculaire dense.

Qualité 3: Une couche fine d'os cortical entoure un noyau d'os trabéculaire dense.

Qualité 4: Une couche fine d'os cortical entoure un noyau d'os trabéculaire peu dense.



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ANNEXE: PROTHÈSE PARTIELLE AMOVIBLE

Classification de Kennedy 1-4

Classe I: Zones édentées bilatérales situées postérieurement par rapport aux dents naturelles restantes.

Classe II: Zone édentée unilatérale située postérieurement par rapport aux dents naturelles restantes.

Classe III: Édentement intercalaire de tout type (sauf antérieur unique)

Classe IV: Édentement antérieur avec un seul segment édenté et traversant la ligne médiane.

Classe V: une situation édentée dans laquelle les dents sont présentes antérieurement et postérieurement mais la délimitation antérieure n'est pas adaptée pour une fonction d'appui (telle que l'incisive latérale).

Classe VI: une situation édentée dans laquelle les dents limites sont capables d'assurer le support total de la prothèse requise.

Évaluation de la prothèse partielle actuelle^{5, 6}

Rétention de la prothèse

La rétention est classé comme "excellente" (résistance au déplacement vertical); "bonne" (une résistance modérée au déplacement vertical), ou "mauvaise" (pas de résistance au déplacement vertical).

Stabilité de la prothèse

La stabilité est classé comme "excellente" (stable au mouvement de rotation et les appuis occlusaux sont ajustés); "bonne" (stable, mais certains appuis sont pas ajustés), ou "mauvaise" (déplacée par mouvement rotationnel).

Châssis métallique

Le châssis métallique est classé comme "bien ajusté", "Adaptation des appuis occlusaux et crochets inacceptable" ou "fracturés".

Adaptation de la base acrylique à la muqueuse

L'adaptation de la base acrylique à la muqueuse est classé comme "bien ajustée", " légère imprécision" ou " adaptation inacceptable".

IP (Indice de plaque)^{7, 8}

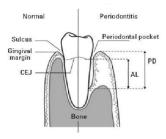
0= Pas de plaque

1= une pellicule de plaque adhérente à la gencive marginale libre et à la surface dentaire adjacente. La plaque pourrait être révélée par l'usage de la sonde sur la surface dentaire.

2= Accumulation modérée de dépôts mous dans la poche gingivale, ou sur la dent et la gencive marginale qui pourrait être vus à l'œil nu.

3= Abondance de matière molle dans la poche gingivale, ou sur la dent et la gencive marginale.

PP (**Profondeur de poches**)⁹ :Score du Sondage (**Profondeur en millimètres**). Il est mesuré à partir du bord de la gencive à la base de la poche clinique.



SAS (Saignement au sondage)¹⁰

0= Pas de saignement

1= Saignement

Carie¹¹⁻¹⁴

- 1. Surface saine: Toute surface ne présentant ni caries ni obturations.
- 2. Lésion active: Toute surface bien définie et qui présente une décoloration jaunâtre ou brun clair. La lésion est ramollie ou de consistance coriace lorsqu'elle est sondée avec une pression modérée et peut être recouverte de plaque.
- 3. Lésion inactive: Toute surface bien définie, noir brunâtre ou noire. La surface est lisse, brillante et dure au sondage.
- 4. Surface obturée: considérée principalement comme le résultat du traitement. Les récidives de caries sont marquées comme des caries actives.
- 5. Surface délabrée: Elle se définit comme suit: surface dentaire entièrement détruite et totalement absente. Lorsque cinq ou plusieurs surfaces sont détruites, la dent est considérée comme un fragment radiculaire.

Mobilité dentaire 15, 16

- 0=Pas de mobilité
- 1= Jusqu'à 1 mm de mouvement dans le sens horizontal.
- 2= Plus de 1 mm de mouvement dans le sens horizontal.
- 3= Mouvement horizontal et vertical excessifs.

Structure des dents piliers¹⁷

Etat du pilier	ideal	ou peu	compromis.

□Pas de traitement pré prothétique indiqué.

État du pilier modérément compromis.

- ☐ Piliers dans 1 ou 2 sextants* ont une structure dentaire insuffisante pour retenir ou supporter des restaurations intracoronaires ou extracoronaires.
- ☐ Piliers dans 1 ou 2 sextants requièrent un traitement adjuvant localisé (e.g. , parodontal, endodontique, ou procédures orthodontiques).

État du pilier considérablement compromis.

- ☐ Piliers dans 3 sextants ont une structure dentaire insuffisante pour retenir ou supporter des restaurations intracoronaires ou extracoronaires.
- ☐ Piliers dans 3 sextants requièrent davantage de traitements localisés considérables (e.g. , parodontal, endodontique, ou procédures orthodontiques).

État du pilier sévèrement compromis.

- ☐ Piliers dans 4 sextants ou plus ont une structure dentaire insuffisante pour retenir ou supporter des restaurations intracoronaires ou extracoronaires.
- □ Piliers dans 4 sextants ou plus requièrent un traitement adjuvant localisé (e.g. , parodontal, endodontique, ou procédures orthodontiques).
- ☐ Piliers ont un pronostic réservé.
- * Un sextant est une subdivision de l'arcade dentaire. Les arcades dentaires maxillaire et mandibulaire pourraient être subdivisées en six zones ou sextants. Dans le maxillaire supérieur, le sextant postérieur droit s'étend de la dent 14 à la dent 18, le sextant postérieur gauche s'étend de la dent 24 à la dent 28, et le sextant antérieur s'étend de la dent 13 à la dent 23. A la mandibule, le sextant postérieur droit s'étend de la dent 44 à la dent 48, le sextant postérieur gauche s'étend de la dent 34 à la dent 38, et le sextant antérieur s'étend de la dent 33 à la dent 43.

		Maxillaire				
Dents	18-14	13-23	24-			
			28			
Droite				Gauche		
Dents	48-44	43-33	34-			
			38			
	Mandibule					

Occlusion¹⁷

Types de malformations de la crête¹⁸ (classification de Seibert)

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APPENDIX VII: ANNEXE QUESTIONNAIRE#1 ET QUESTIONNAIRE SUR LES RÉSULTATS

ANNEXE: QUESTIONNAIRE#1

Section B: Modes de vie¹

Section C: Antécédents médicaux²⁻²¹

Section F: Anxiété dentaire ^{22, 23}

Section G: Caractéristiques psychologiques^{24, 25}

ANNEXE: QUESTIONNAIRE SUR LES EFFETS

Section A: Qualité de vie

- A.1. En général, diriez-vous que votre santé est^{26, 27}?
- A.2. En général, diriez-vous que la santé de votre bouche est^{28, 29}?
- A.3. OHIP-EDENT 30
- A.4. Durant les 12 derniers mois, vous absentiez-vous du travail ou de vos activités normales en raison de la nécessité d'avoir des soins dentaires, y compris des examens dentaires, ou à cause de soucis avec votre bouche³¹?

Section B: La satisfaction du patient

(**B.1. B.7.**^{32, 33})

- B.1. En général, comment est votre satisfaction concernant votre prothèse supérieure?
- B.2. En général, comment est votre satisfaction concernant votre prothèse inférieure?
- B.3. Comment est votre satisfaction concernant la rétention de votre prothèse supérieure?
- B.4. Comment est votre satisfaction concernant la rétention de votre prothèse inférieure?
- B.5. Comment est votre satisfaction concernant le confort fonctionnel de vos prothèses?
- B. 6. Comment est votre satisfaction concernant l'aspect de vos prothèses?
- B.7. Comment est votre satisfaction concernant la parole avec vos prothèses?

(**B.8. B.11**.^{34, 35})

- B.8. Ressentez-vous un déplacement de la prothèse supérieure en mangeant et/ou en parlant?
- B.9. Ressentez-vous un déplacement de la prothèse inférieure en mangeant et/ou en parlant?

- B.10. Avez-vous des problèmes à mordre avec vos prothèses?
- B.11. Avez-vous des problèmes à mâcher avec vos prothèses?

(**B.12.**_**B.14**. ^{36, 37})

- B.12. Trouvez-vous des particules alimentaires sous vos prothèses?
- B.13. Ressentez-vous une différence de goût des aliments à cause de vos prothèses?
- B.14. Est-ce que la prothèse influence votre choix d'aliments?

(B.15. B.18. 38)

- B.15. Avez-vous de la difficulté à nettoyer vos prothèses?
- B.16. Ressentez-vous une mauvaise haleine à cause de vos prothèses?
- B.17. Ressentez-vous une sécheresse buccale?
- B.18. Avez-vous des problèmes d'hypersialorrhée (hypersalivation)?

Section C: Attentes de satisfaction à l'égard de la prothèse totale

C.1. Quelle serait votre satisfaction avec de nouvelles prothèses totales conventionnelles³⁹?

Section D: Satisfaction par la visite dentaire $^{40,\,41}$

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