Development of an Evidence-Based Prosthodontic Record: An Action Research Study

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This study sought to develop a database in the field of removable prosthodontics by using a participatory action research method. Data collection consisted of a comprehensive literature review, focus-group discussions, and interviews. Applying action research methods ensures consideration of the needs, perspectives, and expertise of academia in the design and implementation of an evidence/ research-based patient record, and academic educators are well placed to conduct such research. *Int J Prosthodont 2013;26:359–364. doi: 10.11607/ijp.3397*

Although the importance of practice-based re-search has been widely documented, this process advances slowly and faces several barriers, such as validity of the evidence and high costs of providing clinical data.¹ Therefore, strategies need to be implemented that allow powerful data queries on large pools of patient data with relatively low cost and without information and measurement bias. In the field of prosthodontic research, as for other research domains, a university-based dataset can facilitate this process. To this aim, a participatory action research method that links researchers and clinicians synergistically to evaluate and change practices can be appropriate.^{2,3} This study briefly describes the approach of creating an evidence-based patient record in the field of removable prosthodontics. Ultimately, this record could be used to provide the basic elements of a prosthodontic database.

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Materials and Methods

The study was approved by the Ethics Committee of the Université de Montréal, and informed written consent was obtained from each participant. A participatory action research design and the purposive sampling technique were used to select study participants (Table 1).^{2–4}

Data were collected during the different phases of the action research cycle (Fig 1). In the first phase, "problem identification," six focus-group discussions and 13 individual interviews were conducted to develop the criteria for designing the new patient record.

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.

In the next phases, "action planning/acting on the evidence," the new prosthodontic record was designed based on the results of the previous phases. Finally, in the last phases, "action evaluation/interpretation," the completeness and appropriateness of the new prosthodontic record was assessed and necessary modifications were carried out.

The analysis included debriefing, transcription, and thematic analysis.⁵ The interviews and focus groups were all audio-recorded, transcribed verbatim, and coded using computer qualitative software (QDA Miner version 3.2.3, Provalis Research).

Results

The thematic analysis of the collected data yielded several key concepts that are summarized below.

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Characteristics of Study Participants ($n = 14$)	
Participants Tot	al
Sex	
Male	9
Female	5
Academic status	
Full-time professor in removable prosthodontics	5
Part-time clinician in removable prosthodontics	9
Age	
30 to 40 y	4
	6
	4
Teaching experience	
	1
10 10 20 9	4 6
	3

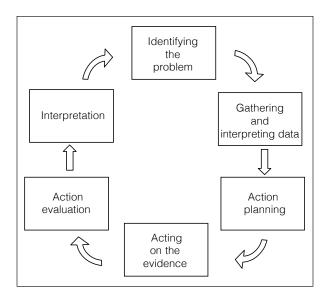


Fig 1 Action research cycle model (adapted from Susman⁷).

 Table 2
 The Content of the Newly Designed Evidence-Based Prosthodontic Record and Related Questionnaires

Prosthodontic record	
Part 1:	
Assessment of potential risk factors of prosthodontic outcomes	Sociodemographic characteristics, medical and dental history, lifestyle habits, dental service use, oral hygiene habits, dental anxiety, and psychologic characteristics
Part 2:	
Oral clinical examinations and assessment of disease-oriented outcomes	Evaluation of current denture, soft and hard tissues, caries, periodontal diseases, denture stomatitis, and alveolar bone resorption
Part 3:	
Assessment of patient-oriented outcomes	Oral health-related quality of life, patient satisfaction, and dental visit satisfaction

Enthusiasm to Change

This theme was evident when the professors and clinical instructors demonstrated their willingness to collaborate in the process.

Empowerment in Practice, Education, and Research

Three types of weaknesses in the current prosthodontic record were identified. (1) Clinical weaknesses: for most interviewees, the actual clinical form did not allow for patient follow-up or for conducting clinical audits. In addition, the participants 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. (3) Research weaknesses: the clinical researchers stated that the existing record was solely clinical.

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.

Expanding Knowledge

Based on an extensive systematic review, the important outcomes of interest in removable prosthodontic research and their data collection instruments were selected and the new research-based prosthodontic record was developed in the action planning phase and evaluated by the research team in the action evaluation and interpretation phases (Table 2, Appendices I to IV).

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Table 1 The Demographic and Academic Characteristics of Study Participants (n = 1/2)

Appendix I: Part of Questionnaire no. 1

	ng?				-	-	-	
0	1	2	3 4	1	5			
H.8. H chewir		of your poste	erior (back) r	natural te	eth in the lef	t side get	in touch during	
0	1	2	3 4	ļ	5			
Before	answering the	following qu	estions, plea	ise pay a	attention to th	ese defini	tions:	
prosthet that re in the u	denture: a rem esis with artific place some of upper and/or lo	ial teeth your teeth ower jaw.	Yes (If yes, r	pro tha the	nplete dentui sthesis with a t replace all c upper and/o	artificial te of your tee r lower jav	eth th in v.	and and
	,			0		,	the table below.	
	proximately whe					B) Which		
			-			/ -		
C) WIII	ich jaw?	D)	Please ment	ion ii you	i were wearii	ig it/them	during mastication	JH.
A: Ag	le	В: Тур	e of denture		C: Jaw		D: Wearing dur mastication	ing
Be	tween 20–34		nplete dentu tial denture	ire	Upper Lower Both Non appli	cable	□ Yes □ No	
	tween 35–49		nplete dentu tial denture	ire	Upper Lower Both	cable	☐ Yes ☐ No	
			nplete dentu	ire	Upper		□ Yes □ No	
	tween 50–64		tial denture		□ Lower □ Both □ Non appli	cable		

Discussion

An evidence-based patient record has several advantages, such as monitoring dental care, facilitating clinical research, and allowing clinical audit in the university-based setting. Furthermore, it will create a research-training environment for clinicians and dental students. It will also raise awareness about evidence-based practice. Previously, 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.⁶ However, in terms of implementation, barriers could be expected, such as students' resistance toward the complexity of the design and lack of knowledge regarding the concept. The authors believe that by providing appropriate support and training these barriers could be resolved.

Appendix II: Part of Complete Removable Prosthesis Questionnaire

	1	eprosthetic –		
	g preprosthetic surgery			
 Extraction Minor soft tis Implant surg 				
		prosthetic		
Suggested treatme	nt (Please choose more		treatments are	needed)
□ New prosthesis □ Reline	C/C C/ Rebase	□ /C □ □ Repair] Implant-suppo	orted prosthesis
	PRC	GNOSIS		
	- Influencing factors			
Anatomical factors			Favorable	Unfavorable
Underlying systemat				
Orofacial problems (Psychosocial factors				
Prosthesis history				
In general the progno		Conditional		Poor
Max	_			

Conclusion

This study showed that the application of action research methods ensures consideration of the needs, perspectives, and expertise of academia in the design and implementation of an evidence/ research-based patient record in the field of removable prosthodontics.

Acknowledgment

The authors reported no conflicts of interest related to this study.

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Appendix III: Part of Partial Removable Prosthesis Questionnaire

Evaluation	of	abutmont	tooth
Evaluation	OT	abutment	teetn

Abutment tooth structure

□ Ideal or minimally compromised abutments. (No preprosthetic therapy is indicated.)

□ Moderately compromised abutments. (Abutments in 1 or 2 sextants have insufficient tooth structure to retain or support intracoronal or extracoronal restorations and they require localized adjunctive therapy.)

□ Substantially compromised abutments. (Abutments in 3 or more sextants have insufficient tooth structure to retain or support intracoronal or extracoronal restorations and they require more substantial localized adjunctive therapy.)

□ Severely compromised abutments. (Abutments in 4 or more sextants have insufficient tooth structure to retain or support intracoronal or extracoronal restorations and they require extensive adjunctive therapy. Abutments have guarded prognoses.)

Occlusal plan	OVD
□ Adequate	□ Adequate
Minor occlusal adjustment	□ To modify: □ To be increased
	□ To be reduced
Occlusal rehabilitation	
Jaw relationship	Posterior support
Class I	□ Adequate
Class II division	□ Inadequate
Class III	
_	
	Ridge deformities
Class I (buccolingual loss of tissue contended and the second se	u pur with a normal apicocoronal height)
-	
□ Class II (apicocoronal loss of tissue with	normai buccolingual contour)
Class III (a combination of buccolingual	and apicocoronal loss)

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Appendix IV: Part of Outcomes Questionnaire

On a scale of 1 to 5, please answer (please circle one).		r that best fits your		Never	Rarely	Sometimes	Often	Always			
B.15. Do you experience di	ifficulty cleaning y	our denture?		1	2	3	4	5			
B.16. Do you experience a	ny bad mouth odo	r with your denture	s?	1	2	3	4	5			
B.17. Do you experience a	ny mouth dryness'	?		1	2	3	4	5			
B.18. Do you experience p	roblems with oral of	continence (droolin	g)?	1	2	3	4	5			
Section C											
C.1. How satisfied do	o you think you wil	I be with a new con	iventional		s?						
0				— 100							
0 = not satisfied at al	0 = not satisfied at all					100 = completely satisfied					
			_		_	_	_				
Section D											
D.1. After talking with th	ne dentist/student,	I know what the co	ondition of	my mout	h is.						
Strongly disagree	Disagree	Uncertain	🗆 Agre	ee D	□ Strongly	agree					
D.2. After talking with the dental health in the nex		I have a good idea	a of what c	changes t	o expect in	n my					
□ Strongly disagree	Disagree	Uncertain	🗆 Agre	e E	□ Strongly	agree					
D.3. The dentist/studen	t told me all I wan	ted to know about r	ny dental	problem(s).						
Strongly disagree	Disagree	Uncertain	□ Agre		☐ Strongly	agree					
D.4. I really felt underst	ood by my dentist	/student.									
Strongly disagree	Disagree	Uncertain	🗆 Agre	e D	□ Strongly	agree					
D.5. I felt that this denti	st/student really kr	new how upset I wa	as about th	ne possih	ility of pair	1					
Strongly disagree	Disagree	Uncertain	Agre		☐ Strongly						
0, 0	0		0		5,	2					

Literature Abstract

Oral bisphosphonate use increases the risk for inflammatory jaw disease: A cohort study

Bisphosphonates (BPs) have been widely used as anti-resorptive agents due to their anti-osteoclatic action. The purpose of this cohort study was to determine whether Danish patients who were treated with BPs and other drugs for the preventive treatment of osteoporosis have an elevated risk for inflammatory jaw-related events, such as osteomyelitis, osteitis, periostitis, or sequestrum, compared with a random sample of the Danish population (the nonexposed group). Results showed that the study sample consisted of 103,562 index subjects and 310,683 control subjects. After adjusting for other factors, including diabetes and chemotherapy, two BPs, alendronate (HR = 3.15, 95% confidence interval: 1.44–6.87) and etidronate (HR = 2.23, 95% confidence interval: 1.15–4.31), were correlated with an elevated risk for inflammatory jaw events. There was no dose-response relationship between oral BPs and inflammatory jaw events. The authors concluded that the oral BPs alendronate and etidronate were correlated with an elevated risk for inflammatory jaw events.

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