

PONTIFÍCIA UNIVERSIDADE CATÓLICA DO RIO GRANDE DO SUL
PROGRAMA DE PÓS-GRADUAÇÃO EM ODONTOLOGIA
MESTRADO EM ODONTOLOGIA

BÁRBARA THEREZA DE FREITAS KOPPE

**VALIDAÇÃO DO HISTÓRICO DE TRATAMENTO ENDODÔNTICO
AUTORREPORTADO EM UMA SUBPOPULAÇÃO URBANA DO SUL DO BRASIL:
UM ESTUDO MULTICÊNTRICO**

Porto Alegre
2019

PÓS-GRADUAÇÃO - STRICTO SENSU



Pontifícia Universidade Católica
do Rio Grande do Sul

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ESCOLA DE CIÊNCIAS DA SAÚDE
CURSO DE ODONTOLOGIA
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Dissertação apresentada como requisito obrigatório para obtenção do título de mestre na área de Endodontia pelo Programa de Pós-Graduação do Curso de Odontologia da Escola de Ciências da Saúde da Pontifícia Universidade Católica do Rio Grande do Sul

Orientador: Prof. Dr. Maximiliano Schünke Gomes

Porto Alegre, fevereiro de 2019

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Aprovada em: ____ de _____ de _____

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Dedico este trabalho à minha mãe e
ao meu pai, meus primeiros e mais
importantes mestres na vida.

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LISTA DE ABREVIATURAS

DCNT – Doença crônica não-transmissível

DCV – Doença cardiovascular

DM – Diabetes mellitus

OMS – Organização Mundial da Saúde

DP – Doença periodontal

PA – Periodontite apical

TE – Tratamento endodôntico

HTEAR – Histórico de tratamento endodôntico autorreportado

RESUMO

Testes diagnósticos e de rastreamento são ferramentas essenciais para determinar medidas acurazes de doenças, entender suas etiologias e seus padrões de transmissão. Para que um teste tenha níveis de validação adequados, sua performance deve ser comparada em uma variedade de populações e cenários. Medidas de saúde autorreportadas já se comprovaram como testes eficazes para avaliar diversas condições de saúde e doença, tanto gerais quanto orais. Especialmente no campo da Endodontia, esses testes podem ser úteis na substituição da necessidade de radiografias para estudos epidemiológicos. O histórico de tratamento endodôntico autorreportado (HTEAR) já foi utilizado como método para identificar a experiência de um indivíduo com doença e tratamento endodônticos, mas apenas em populações masculinas e com elevado nível de educação. O objetivo deste estudo multicêntrico foi quantificar a validade do HTEAR para identificação da presença de tratamento endodôntico (TE) e periodontite apical (PA) em uma subpopulação de pacientes iniciando tratamento em universidades de duas diferentes cidades do sul do Brasil. A variável de exposição principal (HTEAR) foi coletada através de questionário respondido por 228 participantes, e os desfechos TE e PA foram avaliados através da análise de radiografias panorâmicas. Os dados coletados incluíram número total de dentes e número de dentes com TE e PA para cada participante. Foram calculados os valores de acurácia, sensibilidade, especificidade, valores preditivos positivos e negativos (VPP e VPN), eficiência e razões de verossimilhança positiva e negativa (RVP e RVN). A amostra final foi composta por 198 indivíduos para análise de TE e 192 para análise de PA, após exclusões. Os valores para HTEAR foram: acurácia (TE=0,858; PA=0,474); sensibilidade (TE=0,954; PA=0,739); especificidade (TE=0,671; PA=0,250); VPP (TE=0,850; PA=0,454); VPN (TE=0,882; PA=0,530); eficiência (TE=0,812; PA=0,494); RVP (TE=2,899; PA=0,985); RVN (TE=0,068; PA=1,004). HTEAR pode ser considerado um método válido para identificação da presença de TE, mas não de PA nesta população. A maior parte dos valores das medidas que foram calculados variaram nesta população, em comparação com estudos anteriores, demonstrando que o método do HTEAR se mostrou dependente do contexto em que foi aplicado. Os presentes resultados, portanto, apontam para a necessidade de mais estudos em outros cenários populacionais sobre a acurácia do HTEAR, para

que este possa vir a ser utilizado em futuros estudos epidemiológicos de larga escala que incluam variáveis endodônticas.

Palavras-chave: tratamento endodôntico, periodontite apical, autorrelato, validação

ABSTRACT

Diagnostic and screening tests are essential tools for determining accurate estimates of diseases, understanding their etiology and transmission patterns. In order to have appropriate validity, a test's performance should be compared in a variety of populations and settings. Self-reported health status has proved to be a low cost, effective test to assess various general and oral conditions. Especially in the field of Endodontics, they can be useful in replacing the need for radiographs in epidemiological studies. Self-reported history of endodontic treatment (SRHET) has been used as a method of identifying an individual's experience with endodontic disease and treatment, but only in highly educated male populations. The aim of this multicenter study was to evaluate the validity of SRHET for endodontic treatment (ET) and apical periodontitis (AP) in a subpopulation of first-time patients at universities in two different cities in the South of Brazil. Main exposure SRHET was collected through questionnaire for 228 participants and outcomes ET and AP were assessed by analysis of panoramic radiographs. Data collected included total number of teeth and number of teeth with ET and/or AP for each participant. Validities of SRHET for presence of ET and AP were calculated separately through values of accuracy, sensitivity, specificity, positive and negative predictive values (PPV and NPV), efficiency and positive and negative likelihood ratios (PLR and NLR). Final sample comprised 198 individuals for ET analysis and 192 for AP analysis, after exclusions. The SRHET values were: accuracy (ET=0.858; AP=0.474); sensitivity (ET=0.954; AP=0.739); specificity (ET=0.671; AP=0.250); PPV (ET=0.850; AP=0.454); NPV (ET=0.882; AP=0.530); efficiency (ET=0.812; AP=0.494); PLR (ET=2.899; AP=0.985); NLR (ET=0.068; AP=1.004). SRHET is a valid method for predicting ET, but not for AP in this population. Values of most measures calculated varied in this population in comparison to previous studies. SRHET thus showed to be context-dependent and should be tested in a variety of other populational settings in order to allow its use in future large-scale epidemiological studies including endodontic variables.

Keywords: endodontic treatment, apical periodontitis, self-report, validation

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1 INTRODUÇÃO

As doenças crônicas não-transmissíveis (DCNTs) têm sido as maiores causadoras de morte na população mundial nas últimas décadas, sendo responsáveis por 38 milhões (67,8%) das 56 milhões de mortes no mundo em 2012. Destas, 82% foram provocadas pelas quatro principais DCNTs: doenças cardiovasculares (DCV), diabetes mellitus (DM), câncer e doenças respiratórias crônicas. A Organização Mundial da Saúde (OMS) relata que esses números vêm aumentando – no ano de 2000, houve 31 milhões de mortes causadas por essas enfermidades – e estima que em 2030, o total chegará a 52 milhões de mortes por DCNTs (WHO, 2014).

Contribuindo com a relevância desses dados, destaca-se ainda que cerca de 42% de tais eventos ocorrem em indivíduos com menos de 70 anos e, destes, 82% se dão em populações de países de baixa e média renda, como o Brasil. Além disso, as DCNTs provocam altos índices de morbidade nas populações, demandando altos investimentos dos serviços de saúde pública. Tais gastos, somados ao impacto provocado pela elevada prevalência das mortes prematuras, diminuem a força de trabalho e, conseqüentemente, dificultam o desenvolvimento socioeconômico e a erradicação da pobreza nesses países (WHO, 2010, 2014).

Tendo esses números alarmantes em vista, a OMS publicou, no ano de 2013, um plano de ação global, visando à prevenção das DCNTs e de seus eventos consequentes. Os objetivos foram estipulados principalmente com base no combate aos fatores de risco mais conhecidos para DCNTs (fumo, dieta não-balanceada, inatividade física e uso abusivo do álcool). Dentre esses objetivos estão a redução no consumo de sal e de bebidas alcoólicas, a redução na prevalência de atividade física insuficiente, diminuição do fumo, contenção do aumento na prevalência de diabetes e obesidade e redução dos índices de morte prematura por DCNTs (WHO, 2013a).

Entretanto, a própria OMS afirma que a eliminação completa dos principais fatores de risco comportamentais seria capaz de evitar apenas três quartos de todas as mortes causadas por DCNTs (WHO, 2013b). Isso significa que 25% dessas mortes podem ainda possuir outros fatores de risco, ainda não bem elucidados. Recentemente, o papel da inflamação crônica vem sendo avaliado como possível contribuinte para o desenvolvimento de DCNTs, especialmente doenças

35 cardiovasculares e diabetes mellitus. Estudos na área médica têm investigado a
36 associação da inflamação crônica com riscos aumentados para aterosclerose (bem
37 como suas consequências – doença coronariana e acidente vascular cerebral) e
38 resistência à insulina, levando à síndrome metabólica e diabetes mellitus tipo 2
39 (DREGAN et al., 2014).

40 Seguindo essa linha, patologias bucais como doença periodontal (DP) e
41 periodontite apical (PA), que são doenças crônicas infecto-inflamatórias, também
42 têm sido investigadas como possíveis novos fatores de risco para DCNTs. Na área
43 odontológica, Mattila e colaboradores foram os primeiros a identificar uma
44 associação entre saúde oral e infarto agudo do miocárdio, mesmo após ajuste para
45 idade, sexo, classe social, fumo, concentrações lipídicas sanguíneas e presença de
46 diabetes (MATTILA et al., 1989). Posteriormente, outros estudos que avaliaram
47 doença periodontal, tanto prospectivos (BECK et al., 1996; DESTEFANO et al.,
48 1993) quanto retrospectivos (MORRISON; ELLISON; TAYLOR, 1999), conseguiram
49 observar associação entre DP e DCVs. Ainda em relação à DP, ficou igualmente
50 estabelecida a relação com diabetes mellitus (IACOPINO, 2001).

51 Como a periodontite apical (PA) divide aspectos em comum com a DP, faz
52 sentido que também fosse investigada como fator de risco para DCV e DM. Ambas
53 DP e PA são doenças infecciosas que apresentam microbiota semelhante (em sua
54 maioria bactérias anaeróbias Gram-negativas) e causam elevação em marcadores
55 inflamatórios sistêmicos (COTTI et al., 2011a).

56 No que se refere às DCVs, sabe-se que se originam de disfunção endotelial
57 que leva à formação de placas ateroscleróticas (MÜLLER; ANDREA, 2000).
58 Processos inflamatórios crônicos e doenças infecciosas, como são PA e DP, por sua
59 vez, são capazes de provocar tais danos endoteliais e consequente aterosclerose,
60 bem como causam eventos cardiovasculares agudos, incluindo ruptura da placa
61 aterosclerótica e trombose (ROSS, 1999). Alguns estudos já demonstraram,
62 inclusive, que microrganismos causadores de infecções endodônticas, como
63 *Porphyromonas endodontalis*, estão presentes em placas ateroscleróticas
64 (CHHIBBER-GOEL et al., 2016) e são capazes de invadir culturas de células
65 endoteliais de artérias coronarianas (DORN et al., 2002). Além disso, uma recente
66 metanálise mostrou que a PA está associada à elevação de marcadores
67 inflamatórios sistêmicos, os quais são indicadores de risco para DCV (GOMES et al.,
68 2013).

69 Apesar desses aspectos, ainda há poucos estudos que se dedicaram a avaliar
70 a associação entre PA e DCV. Cotti e colaboradores (COTTI et al., 2011b)
71 encontraram indícios de disfunção endotelial incipiente em homens adultos jovens
72 com PA, através da avaliação de baixos níveis de reserva de fluxo endotelial e
73 aumento dos níveis de interleucina-2 circulante. Em uma população semelhante, foi
74 observada associação entre lesões periapicais de origem endodôntica e eventos
75 cardiovasculares (infarto do miocárdio, angina pectoris e cardiopatia isquêmica
76 crônica) (CAPLAN et al., 2006). Da mesma forma, porém em população de meia-
77 idade, outro estudo identificou a carga endodôntica (soma da quantidade de
78 elementos dentários com tratamento endodôntico e/ou com PA) como preditor de
79 risco independente para eventos cardiovasculares (angina, infarto do miocárdio e
80 morte por evento cardiovascular) (GOMES et al., 2016). Pasqualini e colaboradores
81 também observaram este tipo de associação em uma população de adultos de meia-
82 idade (PASQUALINI et al., 2012).

83 Em um trabalho no qual o histórico de tratamento endodôntico (TE) foi
84 autorreportado, foram observados resultados semelhantes. Os participantes com
85 maior histórico de tratamento endodôntico autorreportado tiveram maior chance de
86 desenvolver DCVs do que aqueles que não reportaram histórico de tratamento
87 endodôntico (CAPLAN et al., 2009).

88 Apesar de ainda não existirem estudos que avaliem prospectivamente se o
89 tratamento endodôntico de dentes com PA é capaz de reduzir o risco de eventos
90 cardiovasculares – algo necessário para confirmação da PA como fator de risco para
91 DCVs –, já foi sugerido que essa redução possa ocorrer. Em um estudo de 2014,
92 foram avaliadas retrospectivamente as imagens de tomografias computadorizadas,
93 bem como a carga aterosclerótica da aorta abdominal através de um escore de
94 cálcio, de 531 pacientes com um total de 11.191 dentes. Os resultados mostraram
95 que houve aumento da carga aterosclerótica com o aumento de dentes com PA,
96 mas não com TE. Pacientes com TE tiveram menor carga aterosclerótica que
97 aqueles sem TE. Dentes com TE e com PA simultaneamente não tiveram relevância
98 na carga aterosclerótica dos pacientes participantes (PETERSEN et al., 2014).

99 Já em relação à diabetes mellitus, diversos estudos também sugerem
100 associação entre DM e maior prevalência de PA, com lesões periapicais maiores do
101 que em pacientes controle e com pior prognóstico de reparo pós-TE. Por outro lado,
102 lesões crônicas de PA não tratadas podem contribuir para maior dificuldade no

103 controle glicêmico de diabéticos. Assim como para as DCVs, entretanto, são
104 necessários mais estudos epidemiológicos prospectivos para comprovar a relação
105 efetiva entre DM e PA (SEGURA-EGEA; MARTÍN-GONZÁLEZ; CASTELLANOS-
106 COSANO, 2015).

107 Tendo em vista que cada vez mais pesquisas apontam para que a PA possa
108 ser um fator de risco para DCNTs, principalmente DCVs e DM, estudos
109 epidemiológicos de grande escala tornam-se extremamente importantes. Segundo a
110 OMS, a obtenção de informações sistemáticas referentes à identificação e à
111 prevalência de fatores de risco são essenciais no planejamento da prevenção de
112 doenças e promoção de saúde. Em seu método de “STEPS” para vigilância de
113 doenças crônicas, o “step” ou passo número 1 a ser realizado é a coleta de
114 informações sobre saúde e doença através de questionários. O uso de questionários
115 é a forma mais rápida e com menor custo para obtenção de dados sobre populações
116 (GOMES et al., 2012; PETERSEN; BAEZ, 2013; PITIPHAT et al., 2002).

117 Questionários vêm sendo um meio de coleta de dados em estudos
118 epidemiológicos desde a Segunda Guerra Mundial e evoluíram significativamente
119 até os dias de hoje, especialmente pela publicação de guias como o *Survey Methods*
120 pela OMS (CANADA, 2010). É um método utilizado para a avaliação autorreportada
121 de diversas questões de saúde-doença, tais como diabetes (BERGMANN et al.,
122 2004; HUERTA et al., 2009; JACKSON et al., 2014), dieta (RIMM et al., 1992),
123 hipertensão arterial sistêmica (ALONSO et al., 2005; HUERTA et al., 2009; TAYLOR
124 et al., 2010; TORMO et al., 2000), doenças cardiovasculares (BERGMANN et al.,
125 2004; HELIÖVAARA et al., 1993), respiratórias, musculoesqueléticas e psiquiátricas
126 (HELIÖVAARA et al., 1993), hipercolesterolemia (HUERTA et al., 2009; TAYLOR et
127 al., 2010), artrite reumatoide (WONG et al., 2004) e câncer (BERGMANN et al.,
128 2004). Outros aspectos menos comuns também já foram avaliados, como uso de
129 entorpecentes (Ecstasy) (YACOUBIAN JR; WISH, 2006) e violência doméstica
130 (HALPERN et al., 2006).

131 Tal como quaisquer outros tipos de testes diagnósticos, questionários devem
132 ser submetidos a avaliações sobre sua qualidade como método de identificação de
133 condições de saúde-doença. Para que isto ocorra, devem ser realizados estudos
134 para análise dos valores de sua validade ou acurácia, ou seja, a capacidade de
135 distinção entre aqueles que têm (sensibilidade) e aqueles que não tem a doença
136 (especificidade). Este dado é bastante importante no rastreamento de doenças em

137 populações. Além disso, devem ser analisados os valores preditivos positivos e
138 negativos para que se possa responder a uma questão mais presente na prática
139 clínica: “se o paciente tiver resultado positivo no teste, qual a probabilidade de ter a
140 doença?”, sendo o mesmo válido para o caso negativo. Outros valores importantes
141 que devem ser calculados são as razões de verossimilhança (ou *likelihood ratios*)
142 positivas e negativas, que representam a probabilidade de se encontrar o resultado
143 positivo ou negativo em pessoas doentes quando comparado a pessoas não-
144 doentes (GORDIS, 2014; HULLEY et al., 2003).

145 Diversos trabalhos relatam formas de se avaliar a qualidade de testes
146 diagnósticos (BOSSUYT et al., 2003a; DAYA, 1996; IRWIG et al., 2002; VETTER;
147 SCHOBER; MASCHA, 2018), sendo que uma das maiores considerações é dada
148 sobre a capacidade de generalização ou transferência dos resultados para diversas
149 populações. Como os sujeitos são normalmente uma das principais fontes de
150 variabilidade entre os testes, sugere-se que, para superar este obstáculo, sejam bem
151 descritas a população incluída no estudo, sua forma de seleção, incluindo critérios
152 de inclusão e exclusão, o local, a época e a situação da coleta dos dados
153 (BOSSUYT et al., 2003a; DAYA, 1996; HULLEY et al., 2003; IRWIG et al., 2002).
154 Além disso, é indicado que os estudos sejam reprodutíveis, e assim possam
155 realizados em diversos locais, tempos e com variadas populações, com o objetivo de
156 fornecer dados suficientes para que subsequentes metanálises possam ser
157 realizadas (BOSSUYT et al., 2003b; HULLEY et al., 2003; IRWIG, 1994; IRWIG et
158 al., 2002; LEE et al., 2015).

159 No campo da Odontologia, pode-se considerar que a avaliação de doenças
160 através de questionários está ainda nos estágios iniciais deste processo, sendo
161 relativamente recente, já que remete às últimas três décadas. O primeiro estudo,
162 realizado em 1986 na Finlândia, conseguiu confirmar a eficácia do uso de
163 questionário para a autoavaliação sobre o número de dentes e presença de próteses
164 dentárias (KÖNÖNEN; LIPASTI; MURTOLOMAA, 1986). No ano de 1991, dois
165 trabalhos deste tipo realizaram validação para saúde oral, obtendo bons resultados
166 também para contagem de dentes (DOUGLASS; BERLIN; TENNSTEDT, 1991) e
167 uso de próteses dentárias (PALMQVIST; B; ARNBJERG, 1991). Desde então,
168 diversos estudos confirmaram tais resultados (AXELSSON; HELGADÓTTIR, 1995;
169 BUHLIN et al., 2002; GILBERT; DUNCAN; KULLEY, 1997; PITIPHAT et al., 2002;
170 RAMOS; BASTOS; PERES, 2013), enquanto outros expandiram a avaliação para

171 presença de doença periodontal (ABBOOD et al., 2016; BLICHER; JOSHIPURA;
172 EKE, 2005; BUHLIN et al., 2002; COBURN et al., 2015; CYRINO et al., 2011;
173 DIETRICH et al., 2005; EKE et al., 2013; GENCO et al., 2007; JOSHIPURA;
174 PITIPHAT; DOUGLASS, 2002; PITIPHAT et al., 2002; RAMOS; BASTOS; PERES,
175 2013), cárie (PITIPHAT et al., 2002; SILVA et al., 2014), tratamento endodôntico
176 (FRANCISCATTO et al., 2019; GOMES et al., 2012; PITIPHAT et al., 2002), dor
177 orofacial (NILSSON; LIST; DRANGSHOLT, 2006) e agenesia dentária (BAELUM et
178 al., 2011).

179 Apesar de alguns estudos terem encontrado resultados insuficientes para
180 validação de DP (BLICHER; JOSHIPURA; EKE, 2005; BUHLIN et al., 2002;
181 DIETRICH et al., 2005; PITIPHAT et al., 2002; RAMOS; BASTOS; PERES, 2013),
182 cárie (PITIPHAT et al., 2002) e agenesia dentária (BAELUM et al., 2011)
183 autorreportadas, a maioria verificou como tendo validade o método de questionário.
184 Para DP há inclusive uma revisão sistemática com metanálise confirmando a
185 possibilidade do uso do diagnóstico autorreportado em grandes estudos
186 epidemiológicos (ABBOOD et al., 2016). Outros corroboram com tais bons
187 resultados, tanto para DP (COBURN et al., 2015; CYRINO et al., 2011; EKE et al.,
188 2013; GENCO et al., 2007; HEATON et al., 2017; JOSHIPURA; PITIPHAT;
189 DOUGLASS, 2002; LAMONTE et al., 2014), quanto para cárie (LEVIN; SHPIGEL;
190 PERETZ, 2013; SILVA et al., 2014), dor orofacial (NILSSON; LIST; DRANGSHOLT,
191 2006), tratamento endodôntico (FRANCISCATTO et al., 2019; GOMES et al., 2012;
192 PITIPHAT et al., 2002) e retenção de restaurações com uso de retentor intracanal
193 pós-tratamento endodôntico (VON STEIN-LAUSNITZ et al., 2018).

194 De uma forma geral, percebe-se que questionários de saúde tendem a ter
195 piores resultados para detecção de doenças transitórias ou agudas e melhores para
196 doenças crônicas (BERGMANN et al., 2004; HELIÖVAARA et al., 1993). Os
197 resultados também tendem a ser melhores para especificidade do que para
198 sensibilidade, ou seja, identificam mais adequadamente saúde do que doença
199 (FARMER et al., 2017; HUERTA et al., 2009).

200 Ainda assim, considerando que o diagnóstico para histórico e necessidade de
201 tratamento endodôntico envolve necessariamente o uso de exames de imagem, o
202 que aumenta consideravelmente seus custos, a ferramenta do questionário torna-se
203 fundamental para a realização de estudos epidemiológicos nessa área. Alguns
204 estudos já confirmaram a validade e utilizaram questionários autorreportados para

205 identificação de histórico de tratamento endodôntico, sendo a maioria dos
206 participantes homens brancos, com elevado nível de escolaridade (CAPLAN et al.,
207 2009; FRANCISCATTO et al., 2019; GOMES et al., 2012; JOSHIPURA et al., 2006;
208 PITIPHAT et al., 2002). Há um estudo que conseguiu bons resultados para validação
209 de DP autorreportada em uma amostra brasileira (CYRINO et al., 2011) e apenas
210 um que tenha feito o mesmo para histórico de tratamento endodôntico
211 (FRANCISCATTO et al., 2019). Este último trabalho, entretanto, foi composto por
212 uma amostra extremamente específica, composta de policiais militares, em sua
213 maioria homens, com consultas regulares ao dentista.

214 Nos trabalhos em que o histórico de tratamento endodôntico autorreportado foi
215 utilizado, isto ocorreu em substituição aos exames de imagem normalmente
216 realizados (radiografias ou tomografias computadorizadas), que seriam o padrão-
217 ouro em imagens para avaliação da presença de TE e PA. Nos casos em que há
218 presença de radiopacidade referente à existência de dentes com TE, pode-se
219 sugerir que o paciente, conseqüentemente, tenha um histórico de doença pulpar ou
220 periapical prévia. Entretanto, esta hipótese ainda deve ser validada, tendo apenas
221 dois trabalhos avaliado a possibilidade da ocorrência desta relação através do
222 histórico de TE (FRANCISCATTO et al., 2019; GOMES et al., 2012).

223 Este trabalho, portanto, teve como objetivo validar o histórico de tratamento
224 endodôntico autorreportado como método de identificação de indivíduos que
225 realizaram TE e/ou apresentam PA em uma população do sul do Brasil, composta
226 por pacientes que estivessem iniciando tratamento em cursos de Odontologia de
227 uma universidade particular da cidade de Porto Alegre, Rio Grande do Sul e de uma
228 universidade pública da cidade de Santa Maria, Rio Grande do Sul.

229 **2 ARTIGO**

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231 a South Brazilian urban population. *Community Dentistry and Oral Epidemiology*,
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235

236 **MULTICENTER VALIDATION OF SELF-REPORTED HISTORY OF ENDODONTIC**
237 **TREATMENT IN A SOUTH BRAZILIAN URBAN POPULATION**

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239

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266 **ABSTRACT**

267

268 The aim of this multicenter study was to evaluate the accuracy of self-reported
269 history of endodontic treatment (SRHET) as a mean of identifying patients who have
270 undergone endodontic treatment (ET) and present with apical periodontitis (AP) in an
271 urban subpopulation of first-time patients at two different universities in South Brazil.
272 Main exposure SRHET was collected through questionnaire for 228 participants, and
273 outcomes ET and AP were assessed by the analysis of panoramic radiographs.
274 Collected data included total number of teeth and number of teeth with ET and with
275 AP for each participant. Validities of SRHET for the presence of ET and AP were
276 calculated separately through values of accuracy, sensitivity, specificity, positive and
277 negative predictive values (PPV and NPV), efficiency and positive and negative
278 likelihood ratios (PLR and NLR). Final sample comprised 198 individuals for ET
279 analysis and 192 for AP analysis, after exclusions. The SRHET values were:
280 accuracy (ET=0.858; AP=0.474); sensitivity (ET=0.954; AP=0.739); specificity
281 (ET=0.671; AP=0.250); PPV (ET=0.850; AP=0.454); NPV (ET=0.882; AP=0.530);
282 efficiency (ET=0.812; AP=0.494); PLR (ET=2.899; AP=0.985); NLR (ET=0.068;
283 AP=1.004). SRHET is a valid method for predicting ET, but not for AP in this
284 population. The values of most validation measures varied in this population
285 compared to previous studies. Present findings thus suggest that the accuracy of
286 SRHET is dependent on the population profile, which encourages further validation
287 research of this test in different scenarios.

288

289 **Key-words:** endodontic treatment, apical periodontitis, self-report, validation

290 Introduction

291 Diagnostic and screening tests are essential tools for determining accurate
292 estimates of diseases, understanding their etiology and transmission patterns.
293 Consequently, they enable health professionals to provide appropriate treatment for
294 patients, contributing to the development of customized preventive public health
295 programs according to community profiles¹. Several studies have been published
296 over the years specifying directions on how to evaluate the quality of diagnostic
297 tests²⁻⁵. Besides having proper validity, one of their main concerns is the possible
298 variability of test accuracy between settings, which can hinder the transferability or
299 generalization of its applicability to different populations.

300 The Standards for Reporting of Diagnostic Accuracy (STARD) initiative,
301 alongside other studies, has suggested that the description of the study population
302 and participant recruitment, the inclusion and exclusion criteria, and the settings,
303 locations and time of data collection are some of the ways to overcome this issue^{2,3,5}.
304 Furthermore, the test should be reproducible and its performance should be
305 compared in a variety of populations and settings, paving the way for subsequent
306 meta-analytic studies that can improve the test's external validity⁵⁻⁸. This is also
307 relevant because many studies do not report considerations on the calculation of
308 sample sizes, and having small numbers of participants can result in imprecise
309 estimates of the overall accuracy of the test⁹.

310 Among many diagnostic and screening tests, self-reported health status has
311 proved to be a low cost, effective way to assess various general¹⁰⁻¹⁴ as well as oral
312 conditions¹⁵⁻²², hence being extremely useful for epidemiological studies. In contrast,
313 the diagnosis of endodontic diseases is usually deemed as costly in regards to time,
314 money and equipment, exposes the patient to radiation and requires the work of
315 qualified professionals. Considering that these characteristics can make it difficult to
316 perform epidemiological studies in the field of endodontics, the self-reported history
317 of endodontic treatment (SRHET) has been used as a simplified way to assess
318 patient's experience of endodontic disease and treatment^{19,23-26}. In these studies,
319 SRHET was used to the detriment of the usual method of imaging exams, such as
320 periapical and panoramic radiographs or cone beam computed tomography (CBCT).
321 When radiopaque evidence of endodontic treatment (ET) is present in these images,
322 they are thought to represent one's history of pulp and/or periapical pathology, even
323 though this assumption has yet to be validated.

324 However, the few studies that have used SRHET so far comprised participants
325 with a specific profile (white males with high education and regular access to dental
326 care) from the United States and Brazil^{19,23–26}. Only two tested the possibility of apical
327 periodontitis (AP) being related to SRHET^{25,26}. At this time, SRHET has yet to be
328 tested in other varieties of populational contexts. The aim of this multicenter study
329 was therefore to evaluate the validity of SRHET as a method for identifying patients
330 who have undergone ET and present with AP in a subpopulation of patients seeking
331 and initiating treatment at the school of Dentistry of one private and one public
332 university in two southern Brazilian cities.

333

334 **Methodology**

335 This study was approved by the Ethics Committee of the School of Health
336 Sciences of the Pontifical University of Rio Grande do Sul and by the Ethics
337 Committee of the Faculty of Dentistry of the Santa Maria Federal University (CAAE
338 #91678518.4.1001.5336). Every participant read and signed an informed consent
339 before entering the study. Participants were recruited in the period between August
340 and November 2018 in the cities of Porto Alegre (Dentistry Course, School of Health
341 Sciences, Pontifical University of Rio Grande do Sul) and Santa Maria (Faculty of
342 Dentistry, Federal University of Santa Maria), in the state of Rio Grande do Sul,
343 Brazil. All collected data were de-identified before analysis.

344 Patients were invited to participate in the study after arriving for an appointment
345 in the triage discipline of the Dentistry undergraduate course and before having any
346 contact with students. Inclusion criteria comprised adults with 18 years or more, who
347 understood and signed informed consent and had a clinical indication of a panoramic
348 radiograph.

349 After reading and signing the informed consent, participants answered the
350 questionnaire, which included sociodemographic questions (age, gender, education
351 level, income), dentist visits-related questions (“Have you ever been to the dentist?”
352 and “When was the last time you visited the dentist?”) and the main exposure
353 question (SRHET) - “Have you ever had root canal treatment?”. Possible answers to
354 that question were “yes” or “no”. Sociodemographic answers such as monthly
355 household income were collected in categories as done in a national oral health
356 survey²⁷(≤ R\$ 250; 251 to 500; 501 to 1500; 1501 to 2500; 2501 to 4500; 4501 to
357 9500; ≥ 9501) and then stratified. The same applies to level of education which was

358 collected as “none, elementary school (incomplete), elementary school (complete),
359 high school (incomplete), high school (complete), graduate school (incomplete),
360 graduate school (complete), post-graduation (incomplete), post-graduation
361 (complete)”, according to a national survey²⁸, before stratification. After completing
362 the questionnaire, participants followed the course’s protocol, having their clinical
363 examination performed by the students and a panoramic radiograph taken for
364 diagnostic and treatment planning purposes.

365 The digital panoramic radiographs were used to obtain the main outcomes ET
366 and AP. The image exams were analyzed by a specialist in endodontics, blinded to
367 SRHET data and previously calibrated to assess the following parameters, according
368 to Gomes *et al.*²⁵:

369 - ET absent = 0, when absence of radiopaque materials inside root canals was
370 observed.

371 - ET present = 1, when presence of radiopaque materials inside one or more
372 root canals was observed.

373 - AP absent = 0, when integrity of the periapical lamina dura was observed, as
374 well as a periodontal ligament width ≤ 2 mm.

375 - AP present = 1, when lack of integrity of the periapical lamina dura, as well as
376 a radiolucency compatible with bone destruction, and periodontal ligament width
377 > 2 mm was observed in one or more roots.

378 The conditions for analysis of the exams were standardized, where radiographs
379 were viewed in a darkened room, in a 20-inch widescreen monitor (LG E2050T; LG,
380 Seoul, South Korea) with an Intel HD Graphics 4600 (Intel, Santa Clara, CA) video
381 card, 32-bit color and 1600x900 resolution.

382 The calibration process comprised the analysis of 40 panoramic radiographs
383 not included in the study sample, evaluated by two examiners. Both specialists
384 independently evaluated the images twice, with a 45-day hiatus between the first and
385 second evaluations. Intra and interexaminer kappa agreement levels were calculated.
386 After calibration, the results for interexaminer agreement were kappa = 0.922 (ET)
387 and kappa = 0.800 (AP). Intraexaminer results were kappa = 0.973 (ET) and kappa =
388 0.956 (AP).

389 Parameters for ET and PA were analyzed for every tooth in every participant,
390 and total number of teeth were recorded. When limitations of the panoramic
391 radiograph, such as overlapping or severe distortions of the image, made it

392 impossible for the examiner to determine parameters for ET or AP for a tooth, that
393 unit was considered “undefined”. If an individual had 10% or more of all teeth with
394 “undefined” parameters, then that subject was excluded from the sample analysis for
395 that outcome (ET or AP). Participants with less than 10 teeth were also excluded.
396 The initial sample was calculated to include 162 patients, based on a previous
397 study²⁵ and considering a power of 80% and $\alpha = 5\%$.

398 After exclusions, for the remaining participants, the validity of the main exposure
399 (SRHET) was calculated separately in relation to both outcomes (presence of ET and
400 AP) obtained from radiographic analysis. Values for accuracy, sensitivity, specificity,
401 positive and negative predictive values, efficiency, and positive and negative
402 likelihood ratios were assessed. Pearson correlation (r) was also calculated between
403 SRHET and ET, SRHET and AP, and ET and AP.

404

405 **Results**

406 At the end of the study, out of the 228 individuals that answered the
407 questionnaire, 30 were excluded for reasons shown in Figure 1. Furthermore, 6
408 patients were also excluded from AP analysis for having 10% or more of remaining
409 teeth with undefined parameters for AP. Consequently, final sample size comprised
410 198 participants for ET and 192 for AP.

411 Sociodemographic and dental characteristics of the sample are shown in Table
412 1. The majority of participants were female (65.7%), with a mean BMI index of 27.76
413 ± 4.99 (overweight), and low levels of education (67.2% had never been to school or
414 had never completed a High School degree). The monthly household income was
415 also low, with the majority of participants’ families earning between R\$ 1501 and R\$
416 2500. The mean age was 49.55 ± 13.58 , with the youngest participant being 18 and
417 the oldest 78 years old. Almost all individuals had already gone to the dentist before
418 in their lives (98.9%), with the most recent visit being had within the past year or two
419 (76.8%). Comparisons between all sociodemographic and dental characteristics of
420 the sample in both cities (Porto Alegre – POA and Santa Maria – SM) are also
421 presented in Table 1.

422 The evaluations of panoramic radiographs showed a mean number of teeth of
423 23.34 ± 5.76 per patient for the sample. Total number of teeth evaluated was 4639, of
424 which 347 (7.5%) presented ET and 161 (3.5%) presented AP. Most individuals had
425 more than 1 tooth with ET present (37.4%), as opposed to one (28.8%) or no teeth

426 with ET (33.8%). In contrast, for the outcome AP, a slight majority had no teeth with
 427 AP (54.2%), while 24.0% had one and 21.8% had more than one teeth with AP. In
 428 total, 131 participants presented at least 1 tooth with ET (66.2%) and 88 had 1 or
 429 more teeth with AP (45.8%). The overwhelming majority of individuals (147 or 74.2%)
 430 responded positively to SRHET.

431 The contingency table for SRHET in relation to both ET and AP is shown in
 432 Table 2. Among that majority who said they had a history of ET, 125 (63.1%) actually
 433 presented with ET in radiographic evaluation, thus representing true positives. Also,
 434 22 (11.1%) of them did not present ET (false positives). Among those who reported
 435 negative SRHET, analysis of radiographs showed 45 (22.8%) participants who did
 436 indeed not have ET (true negatives), while 6 (3.0%) presented ET (false negatives).
 437 In regard to the 192 analyzed for AP, 143 (74.5%) said yes to SRHET: 65 (33.8%) of
 438 them presented AP (true positives) and 78 (40.6%) did not (false positives). Among
 439 the ones who responded negatively to having history of ET, 26 (13.6%) did not
 440 present AP in radiographic evaluations (true negatives), while 23 (12.0%) had 1 tooth
 441 or more with AP (false negatives).

442 Table 3 presents the results of accuracy, sensitivity, specificity, positive
 443 predictive value, negative predictive value, efficiency, positive likelihood ratio, and
 444 negative likelihood ratio for SRHET in relation to the presence of ET and AP. Good
 445 results were found for ET, especially for sensitivity (.954), accuracy (.858), positive
 446 (.850) and negative (.882) predictive values. On the other hand, the opposite was
 447 found for AP with overall low values, particularly for specificity (.250), positive
 448 predictive value (.454) and accuracy (.474).

449 Finally, Table 4 shows the contingency table for the presence of ET and PA.
 450 Out of 126 (65.6%) who had ET, almost half (61/48.4%) presented with AP as well,
 451 while the other half did not have any teeth with AP (65/51.6%). Among those who
 452 had no teeth with ET (66/34.4%), 39 (59.1%) also had no teeth with AP and 27
 453 (40.9%) had at least 1 tooth with AP. The Pearson correlation calculated between ET
 454 and AP was $r = 0.072$ ($p=0.322$), between SRHET and AP was $r = -0.013$ ($p=0.857$)
 455 and between SRHET and ET was $r = 0.677$ ($p<0.001$).

456

457 Discussion

458 This study has confirmed previous findings^{19,25,26} that SRHET is an accurate
 459 method for the screening of patients who have undergone ET. On the other hand, it

460 has also confirmed that SRHET is not predictive of the presence of AP in this
461 population, which agrees with the results found in other studies with different
462 populations^{25,26}. The main novel finding of this study is that the SRHET method
463 showed to be context-dependent, since variations in the population profile modified
464 the accuracy of the instrument.

465 Unlike the aforementioned preceding studies^{19,25,26}, the subjects in this sample
466 were mostly women, with low levels of education and monthly household income,
467 and these characteristics are somewhat representative of such aspects in Southern
468 Brazilian populations. For instance, women are known to seek more frequently for
469 health treatment and have more regularly scheduled appointments than men^{29,30} and
470 thus are more likely to be initiating treatment in two universities. Additionally, just as
471 seen in our sample, the majority (55%) of Southern Brazilians has not completed a
472 high school degree²⁸ and has a mean monthly household income of R\$ 2373³¹. Since
473 education is inversely related to risk of obesity³², this could also explain the high BMI
474 found in our study, indicating that most participants are either overweight or obese
475 (69.6%). Finally, while the number of people that has been to the dentist at least one
476 time in their lives is actually high for the South more developed region of Brazil
477 (95.4%)²⁷, in our findings this rate was even higher (98.9%). The high frequency of
478 consultation, with the last visit having been in the past year or two (76.8%), is in
479 accordance with Southern Brazilian populations as well (85%)²⁷.

480 This sociodemographic profile may explain why so many participants (66.2%)
481 had one or more teeth with ET present, which is much higher than the prevalence
482 found by Franciscatto *et al.* (37.5%)²⁶, Gomes *et al.* (35.8%)²⁵ and Pitiphat *et al.*
483 (34.5%)¹⁹ in other subpopulations. Low socioeconomic status (education and
484 income) is related to higher risk of caries³³, which can consequently lead to ET need,
485 with or without AP. The high frequency of visits to the dentist of the subjects of our
486 study will therefore likely have those needs identified and then ET performed. The
487 prevalence of AP in this population was much higher than in the study of Franciscatto
488 *et al.* and Gomes *et al.* as well (45.8% versus 24.3%²⁶ and 17.8%²⁵), and the same
489 reasoning could be applied.

490 The prevalence of ET and AP, considering the tooth as a unit, also mostly
491 differed from those found in previous studies. The prevalence of ET (347 or 7.5%)
492 was slightly higher and the prevalence of AP (161 or 3.5%) was lower than the
493 results from a different Brazilian population³⁴, but both values were lower in

494 comparison to the ones reported in a systematic review with studies from various
495 countries³⁵. Both of these studies comprised a much larger sample in number of
496 teeth than our study, which can be considered as a limitation to epidemiological
497 comparisons regarding number of teeth.

498 Another highly contrasting aspect we found when comparing present results to
499 those from previous studies that attempted to validate SRHET before is the
500 proportion of individuals that responded positively to having a history of ET. While in
501 our study that response was given by an overwhelming majority of 74.2% of
502 participants, in previous results that percentage was of only 46.3%²⁶, 39.7%²⁵ and
503 31%¹⁹. These importantly different numbers, alongside the high prevalence of
504 patients with ET, could explain the fact that we found a highly significant value for
505 sensitivity of the SRHET (0.954) test in identifying individuals with history of ET in
506 comparison to the three studies (the other values were 0.960²⁶, 0.915²⁵ and 0.900¹⁹).
507 On the other hand, our number of negative SRHET responses (25.8% versus
508 53.7%²⁶, 60.3%²⁵ and 60.3%¹⁹) and the prevalence of patients who actually did not
509 have any ET (33.8% versus 62.5%²⁶, 64.3%²⁵ and 65.5%¹⁹) was so much lower than
510 the other studies that it may have interfered with our true negative analysis and
511 specificity numbers, which were lower than previously reported (0.671 versus
512 0.835²⁶, 0.891²⁵ and 0.921¹⁹).

513 The same reasoning could be applied to the analysis of the AP outcome. The
514 prevalence of patients with at least one tooth with AP was also higher than the
515 previous studies that tried to validate SRHET as a predictor for AP (45.8% versus
516 24.3%²⁶ and 17.8%²⁵). Both our and the other studies' results found SRHET to be
517 insufficient for the screening of patients with AP, but the results for accuracy,
518 sensitivity, specificity, positive and negative predictive values, efficiency and positive
519 and negative likelihood ratios were overall lower in our findings (Table 3). This is
520 most likely explained by the more even distribution of true and false positives and
521 negatives in the sample of this study (Table 2), which in consequence fails to present
522 good results for the validation of SRHET as a method to identify AP. The *r* values
523 results confirm this, showing that the only correlation found was between SRHET and
524 ET.

525 We aimed to select first-time patients of both universities in order to avoid
526 having the patient get the SRHET information from students. However, with the
527 frequent visits to the dentist profile of this sample, it is apparent that individuals had a

528 good knowledge of their oral conditions. Additionally, the fact that SRHET is more
529 accurate in predicting ET than AP is expected, considering that ET is a relatively
530 costly, complex treatment that often requires more than one long appointment and
531 that is frequently indicated after an episode of acute pain or swelling. These factors
532 alone can cause sufficient impact on a persons' life to have them remember having
533 ET performed. In contrast, AP is a commonly asymptomatic disease that is only
534 found through radiographic and clinical examinations.

535 ET is also a treatment that can be indicated in cases without the presence of
536 AP, such as symptomatic irreversible pulpitis or trauma. However, if AP is present,
537 after ET being done it would be expected to heal and consequently disappear from
538 periapical images in radiographic exams. These reasons would increase true
539 positives for SRHET and ET and false positives for SRHET and AP, which was
540 observed in this study (Table 2). In the case of a negative answer for SRHET, it could
541 also mean an increase in false negatives for AP, if it was present and asymptomatic.
542 Furthermore, AP is a disease that can be treated not only through ET, but also by
543 tooth extraction, which can happen to both endodontically treated or non-treated
544 teeth and can thus affect the incidence of false positive and false negative results for
545 ET and AP. Tooth loss is consequently an important confounder and that is why we
546 included only patients with 10 or more teeth.

547 Besides excluding individuals with less than 10 teeth, we also excluded the
548 ones with 10% or more of remaining teeth with undefined parameters for ET or AP
549 evaluation because of image distortions or overlapping of structures, since this is
550 common for panoramic radiographs, which were the ones we used. The six patients
551 excluded from AP analysis for this reason were mainly from distortions in the anterior
552 region, restricted to anterior teeth. Even though these exclusions mean a reduction in
553 the sample, they are important to strengthen results. Still, our sample size was
554 greater than the calculated 162 that were needed based on a previous study²⁵.

555 Regarding radiographic examination it is important to discuss that we used
556 panoramic radiographs in this study because they are still most commonly indicated,
557 cost-effective, low-radiation imaging exams in used our universities. While complete
558 series of periapical radiographies and especially cone beam computed tomography
559 can significantly improve the detection of apical periodontitis³⁶, the only true gold
560 standard in detecting actual presence of a periapical lesion would be a histological
561 assessment^{37,38}. Therefore, despite its limitations, panoramic radiographs do have

562 good accuracy for detecting AP lesions while being more radioprotective than
563 CBCT³⁹. They are still a reasonable method that has been used in many other
564 epidemiological studies regarding the prevalence of periapical radiolucency and
565 endodontic treatment³⁵. Another important aspect is that this is a cross-sectional
566 study, meaning we cannot identify the stage of development or healing of periapical
567 disease, which can only be done in longitudinal studies^{35,38}.

568 Even considering all the differences in characteristics of populations, high
569 values (0.850 - 0.954) of accuracy, sensitivity, positive and negative predictive values
570 were still found for SRHET and ET in this sample as they were in males with high
571 levels of education and lower prevalences of ET^{19,25,26}. In comparison, cold and
572 electric pulp vitality tests, broadly used in daily clinical practice, have less significant
573 or equal values than those of SRHET and ET⁴⁰. However, despite the higher
574 prevalence of AP in this sample, SRHET showed even weaker numbers for the
575 prediction of this disease (Table 3) than it had in the only previous studies that
576 evaluated SRHET and AP^{25,26}. Notwithstanding the fact that SRHET has apparently
577 been sensible to population and settings changes, it is likely that the only way of
578 identifying AP through questionnaire is to add a series of questions related to
579 endodontic risk factors in the same manner that was suggested for periodontal
580 disease⁴¹.

581

582 **Conclusions**

583 SRHET has confirmed to be an accurate method for identifying individuals with
584 experience of ET but not for the presence of AP in this population. Values of
585 accuracy, sensitivity, specificity, positive and negative predictive values, efficiency,
586 positive and negative likelihood ratios and Pearson correlation r values for ET and
587 AP, SRHET and ET and SRHET and AP have shown to be sensible to characteristics
588 of the population in which SRHET is applied. More studies in other contexts should
589 be performed to maximize data in the validation of SRHET as a diagnostic tool that
590 would allow the insertion of endodontic variables in large-scale epidemiological
591 studies. Further investigations should be dedicated to find a specific set of questions
592 that could predict the presence of AP in individuals.

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595

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References

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- 602 1. Gordis L. *Epidemiology*. 5th edition. Philadelphia: Elsevier Saunders; 2014.
- 603 2. Daya S. Study Design for the Evaluation of Diagnostic Tests. *Semin Reprod*
604 *Endocrinol*. 1996;14(2):111-118.
- 605 3. Bossuyt PM, Reitsma JB, Bruns DE, et al. Towards complete and accurate
606 reporting of studies of diagnostic accuracy: the STARD initiative. *Br Med J*.
607 2003;326:41-44.
- 608 4. Vetter TR, Schober P, Mascha EJ. Diagnostic Testing and Decision-Making.
609 *Anesth Analg*. 2018;127(4):1.
- 610 5. Irwig LM, Bossuyt PM, Glasziou PP, Gatsonis C, Lijmer JG. Designing Studies
611 to Ensure that Estimates of Test Are Transferable. *Br Med J*.
612 2002;324(7338):669-671.
- 613 6. Bossuyt PM, Reitsma JB, Bruns DE, et al. The STARD Statement for Reporting
614 Studies of Diagnostic Accuracy: Explanation and Elaboration. *Clin Chem*.
615 2003;49(1):7-18.
- 616 7. Irwig L. Guidelines for Meta-analyses Evaluating Diagnostic Tests. *Ann Intern*
617 *Med*. 1994;120(8):667.
- 618 8. Lee J, Kim KW, Choi SH, Huh J. Systematic Review and Meta-Analysis of
619 Studies Evaluating Diagnostic Test Accuracy : A Practical Review for Clinical
620 Researchers – Part II . Statistical Methods of. *Korean J Radiol*.
621 2015;16(6):1188-1196.
- 622 9. Bachmann LM, Puhan MA, ter Riet G, Bossuyt PM. Sample sizes of studies on
623 diagnostic accuracy: literature survey. *Br Med J*. 2006;332:1127-1129.
- 624 10. Rimm EB, Giovannucci EL, Stampfer MJ, Colditz GA, Litin LB, Willett WC.
625 Reproducibility and Validity of an Expanded Self-Administered Semiquantitative
626 Food Frequency Questionnaire among Male Health Professionals. *Am J*
627 *Epidemiol*. 1992;135(10):1114-1126.
- 628 11. Alonso A, Beunza JJ, Delgado-Rodríguez M, Martínez-González MA.
629 Validation of self reported diagnosis of hypertension in a cohort of university
630 graduates in Spain. *BMC Public Health*. 2005;5:94.
- 631 12. Wong AL, Harker JO, Mittman BS, et al. Development and evaluation of a
632 patient self-report case-finding method for rheumatoid arthritis. *Semin Arthritis*
633 *Rheum*. 2004;34(1):484-499.
- 634 13. Huerta JM, Tormo MJ, Egea-Caparrós JM, Ortolá-Devesa JB, Navarro C.
635 Accuracy of self-reported diabetes, hypertension and hyperlipidemia in the
636 adult Spanish population. DINO study findings. *Rev Española Cardiol*.
637 2009;62(2):143-152.

- 638 14. Jackson JM, DeFor TA, Crain AL, et al. Validity of Diabetes Self-Reports in the
639 Women's Health Initiative. *Menopause*. 2014;21(8):861-868.
- 640 15. Eke PI, Dye BA, Wei L, et al. Self-reported measures for surveillance of
641 periodontitis. *J Dent Res*. 2013;92(11):1041-1047.
- 642 16. Silva AER, Menezes AMB, Assunção MCF, et al. Validation of self-Reported
643 information on dental caries in a birth cohort at 18 years of age. *PLoS One*.
644 2014;9(9):1-8.
- 645 17. Ramos RQ, Bastos JL, Peres MA. Diagnostic validity of self-reported oral
646 health outcomes in population surveys : literature review Validade diagnóstica
647 de agravos. *Rev Bras Epidemiol*. 2013;16(3):716-728.
- 648 18. Nilsson I-M, List T, Drangsholt M. The Reliability and Validity of Self-reported
649 Temporomandibular Disorder Pain in Adolescents. *J Orofac Pain*.
650 2006;20(2):138-144.
- 651 19. Pitiphat W, Garcia RI, Douglass CW, Joshipura KJ. Validation of self-reported
652 oral health measures. *J Public Health Dent*. 2002;62(2):122-128.
- 653 20. Farmer J, Ramraj C, Azarpazhooh A, Dempster L, Ravaghi V, Quiñonez C.
654 Comparing self-reported and clinically diagnosed unmet dental treatment
655 needs using a nationally representative survey. *J Public Health Dent*.
656 2017;(c):1-7.
- 657 21. Axelsson G, Helgadóttir S. Comparison of oral health data from self-
658 administered questionnaire and clinical examination. *Community Dent Oral*
659 *Epidemiol*. 1995;23(6):365-368.
- 660 22. Palmqvist S, B S, Arnbjerg D. Self-assessment of dental conditions: validity of
661 a questionnaire. *Community Dent Oral Epidemiol*. 1991;19(5):249-251.
- 662 23. Joshipura KJ, Pitiphat W, Hung HC, Willett WC, Colditz GA, Douglass CW.
663 Pulpal inflammation and incidence of coronary heart disease. *J Endod*.
664 2006;32(2):99-103.
- 665 24. Caplan DJ, Pankow JS, Cai J, Offenbacher S, Beck JD. The relationship
666 between self-reported history of endodontic therapy and coronary heart
667 disease in the Atherosclerosis Risk in Communities Study. *J Am Dent Assoc*.
668 2009;140(8):1004-1012.
- 669 25. Gomes MS, Hugo FN, Hilgert JB, et al. Validity of self-reported history of
670 endodontic treatment in the Baltimore longitudinal study of aging. *J Endod*.
671 2012;38(5):589-593.
- 672 26. Franciscatto GJ, Koppe BTF, Hoppe CB, et al. Validation of self-reported
673 history of root canal treatment in a South Brazilian subpopulation. *Braz Oral*
674 *Res*. 2019;in press.
- 675 27. BRASIL. MINISTÉRIO DA SAÚDE. *SB Brasil 2010: Pesquisa Nacional de*
676 *Saúde Bucal - Resultados Principais*. Brasília; 2012.

- 677 28. Instituto Brasileiro de Geografia e Estatística. *Pesquisa Nacional Por Amostra*
678 *de Domicílios Contínua: Educação 2017*. Rio de Janeiro; 2018.
679 https://biblioteca.ibge.gov.br/visualizacao/livros/liv101576_informativo.pdf.
- 680 29. Instituto Brasileiro de Geografia e Estatística. *Pesquisa Nacional Por Amostra*
681 *de Domicílios: Acesso e Utilização de Serviços de Saúde - 2003*. Rio de
682 Janeiro; 2005. <https://biblioteca.ibge.gov.br/visualizacao/livros/liv6194.pdf>.
- 683 30. Nascimento GG, Weber CM, Silva DDF. Oral health care in primary care
684 attention: experience from two facilities of Porto Alegre. *Rev Fac Odontol P*
685 *Alegre*. 2011;52(1/3):19-24.
686 [http://seer.ufrgs.br/index.php/RevistadaFaculdadeOdontologia/article/view/304](http://seer.ufrgs.br/index.php/RevistadaFaculdadeOdontologia/article/view/30416/24246)
687 [16/24246](http://seer.ufrgs.br/index.php/RevistadaFaculdadeOdontologia/article/view/30416/24246).
- 688 31. Instituto Brasileiro de Geografia e Estatística. *Pesquisa Nacional Por Amostra*
689 *de Domicílios Contínua: Rendimento de Todas as Fontes 2017*. Rio de
690 Janeiro; 2018.
- 691 32. Monteiro CA, Conde WL, Popkin BM. Independent Effects of Income and
692 Education on the Risk of Obesity in the Brazilian Adult Population. *J Nutr*.
693 2001;131(3):881-886.
- 694 33. Costa SM, Martins CC, Bonfim M de LC, et al. A systematic review of
695 socioeconomic indicators and dental caries in adults. *Int J Environ Res Public*
696 *Health*. 2012;9(10):3540-3574.
- 697 34. Berlinck T, Tinoco JMM, Carvalho FLF, Sassone LM, Tinoco EMB.
698 Epidemiological evaluation of apical periodontitis prevalence in an urban
699 Brazilian population. *Braz Oral Res*. 2015;29(1):1-7.
- 700 35. Pak JG, Fayazi S, White SN. Prevalence of periapical radiolucency and root
701 canal treatment: A systematic review of cross-sectional studies. *J Endod*.
702 2012;38(9):1170-1176.
- 703 36. Estrela C, Bueno MR, Leles CR, Azevedo B, Azevedo JR. Accuracy of Cone
704 Beam Computed Tomography and Panoramic and Periapical Radiography for
705 Detection of Apical Periodontitis. *J Endod*. 2008;34(3):273-279.
- 706 37. Kanagasingam S, Lim CX, Yong CP, Mannocci F, Patel S. Diagnostic accuracy
707 of periapical radiography and cone beam computed tomography in detecting
708 apical periodontitis using histopathological findings as a reference standard. *Int*
709 *Endod J*. 2017;50(5):417-426.
- 710 38. Huuonen S, Ørstavik D. Radiological aspects of apical periodontitis. *Endod*
711 *Top*. 2002;1:3-25.
- 712 39. Nardi C, Calistri L, Grazzini G, et al. Is Panoramic Radiography an Accurate
713 Imaging Technique for the Detection of Endodontically Treated Asymptomatic
714 Apical Periodontitis? *J Endod*. 2018;44(10):1500-1508.
- 715 40. Salgar AR, Singh SH, Podar RS, Kulkarni GP, Babel SN. Determining
716 predictability and accuracy of thermal and electrical dental pulp tests: An in vivo

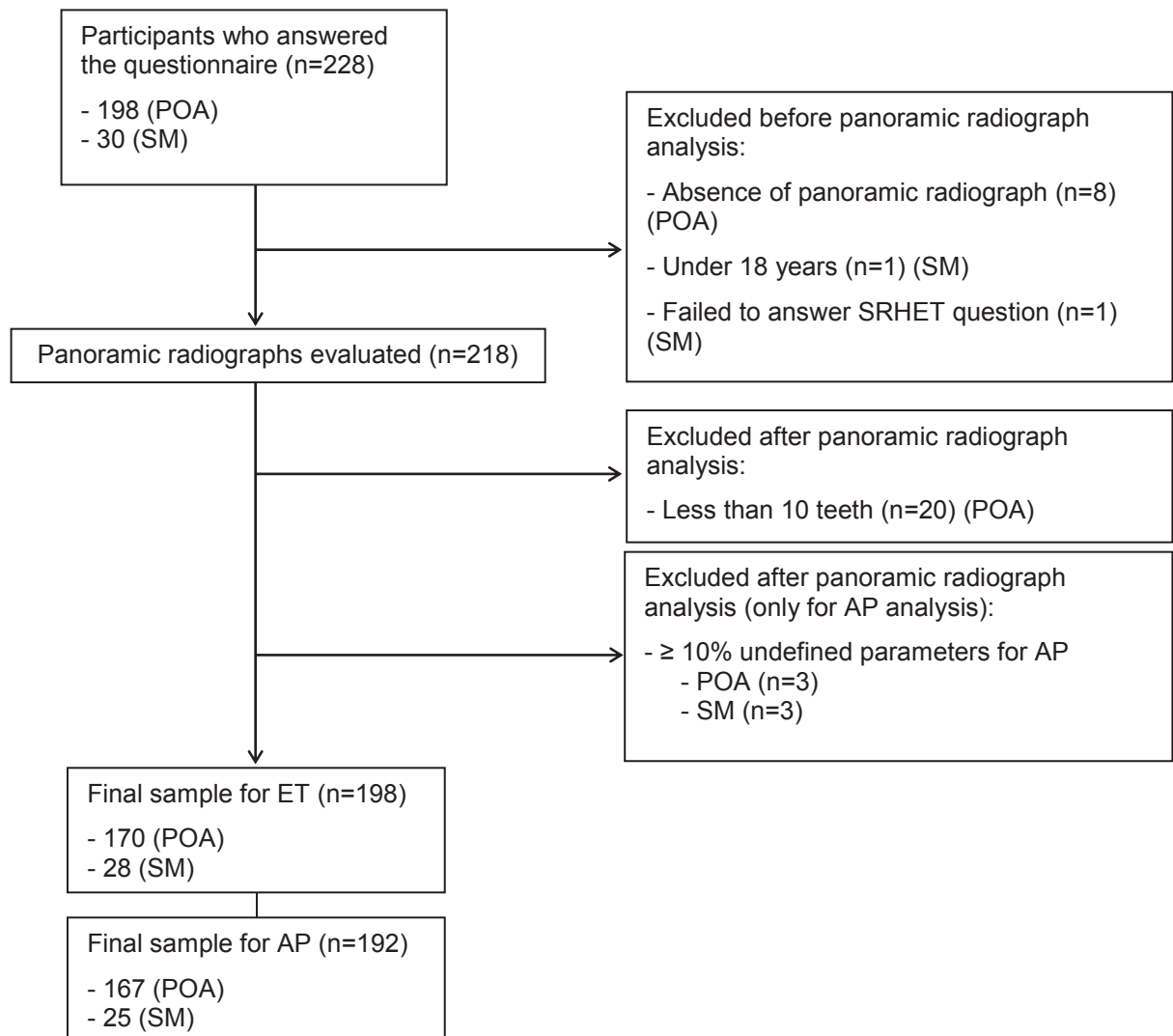
717 study. *J Conserv Dent*. 2017;20(1):46.

718 41. Blicher B, Joshipura K, Eke P. Validation of self-reported periodontal disease: a
719 systematic review. *J Dent Res*. 2005;84(10):881-890.

720 **Figure 1 - Sample selection flowchart**

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722



723 **Table 1** - Sociodemographic and dental characteristics of the sample (n(%) or Mean±SD)

Variable	POA	SM	Total
Sociodemographic			
Age (years) (n=198)	50.54 ±	43.57 ±	49.55 ± 13.58
POA (n=170) / SM (n=28)	13.13	14.54	
Gender (n=198)			
POA (n=170) / SM (n=28)			
Male	59 (35.0)	9 (32.0)	68 (34.3)
Female	111 (65.0)	19 (68.0)	130 (65.7)
Monthly household income (R\$) (n=188)			
POA (n=165) / SM (n=23)			
0 to 500	4 (2.4)	2 (8.7)	6 (3.2)
501 to 1500	46 (27.9)	10 (43.5)	56 (29.8)
1501 to 2500	66 (40.0)	9 (39.1)	75 (39.9)
2501 to 4500	34 (20.6)	2 (8.7)	36 (19.1)
≥ 4501	15 (9.1)	0 (0.0)	15 (8.0)
Education (n=198)			
POA (n=170) / SM (n=28)			
None to High School (incomplete)	75 (44.1)	16	133 (67.2)
High School (complete) to Graduate School (incomplete)	63 (37.1)	6	42 (21.2)
Graduate school (complete) to Post-graduation (complete)	32 (18.8)	6	23 (11.6)
BMI (n=194)			
POA (n=168) / SM (n=26)	27.96 ± 5.15	26.48 ± 3.64	27.76 ± 4.99
Normal weight (18.5–24.9)	49 (29.2)	10 (38.5)	59 (30.4)
Overweight (25.0–29.9)	72 (42.8)	13 (50.0)	85 (43.8)
Obesity (≥ 30.0)	47 (28.0)	3 (11.5)	50 (25.8)
Dental			
Total number of teeth evaluated	3940 (84.9)	699 (15.1)	4639
Number of teeth per patient (n=198)			
POA (n=170) / SM (n=28)	23.18 ± 5.85	24.96 ± 5.02	23.34 ± 5.76
SRHET (n=198)			
POA (n=170) / SM (n=28)			
Yes	128 (75.3)	19 (67.9)	147 (74.2)
No	42 (24.7)	9 (32.1)	51 (25.8)
Number of teeth with ET (n=4639)	299 (7.6)	48 (6.9)	347 (7.5)
Number of teeth with ET per patient (n=198)			
POA (n=170) / SM (n=28)	1.76 ± 2.28	1.71 ± 1.80	1.75 ± 2.21
0	60 (35.3)	7 (25.0)	67 (33.8)
1	46 (27.1)	11 (39.3)	57 (28.8)
> 1	64 (37.6)	10 (35.7)	74 (37.4)
Number of teeth with AP (n=4639)	141 (3.6)	20 (2.9)	161 (3.5)
Number of teeth with AP per patient (n=192)			
POA (n=167) / SM (n=25)	0.84 ± 1.22	0.80 ± 1.08	0.84 ± 1.20
0	93 (55.7)	11 (44.0)	104 (54.2)
1	35 (21.0)	11 (44.0)	46 (24.0)
> 1	39 (23.3)	3 (12.0)	42 (21.8)

"Have you ever gone to the dentist before?" (n=189)

POA (n=161) / SM = (n=28)

Yes	161 (100.0)	26 (92.9)	187 (98.9)
No	0 (0.0)	2 (7.1)	2 (1.1)

Time since last visit to the dentist (n=177)

POA (n=150) / SM (n=27)

< 1 year to 2 years	113 (75.3)	23 (85.2)	136 (76.8)
3 years or more	37 (24.7)	4 (14.8)	41 (23.2)

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727 **Table 2** - Contingency table for the presence of ET and AP in relation to SRHET. Results
728 shown as n(%).

SRHET	Endodontic treatment			Apical periodontitis		
	Yes	No	Total	Yes	No	Total
Yes	125 (85.0)	22 (14.9)	147 (74.2)	65 (45.5)	78 (54.5)	143 (74.5)
No	6 (11.8)	45 (88.2)	51 (25.8)	23 (46.9)	26 (53.1)	49 (25.5)
Total	131 (66.2)	67 (33.8)	198 (100)	88 (45.8)	104 (54.2)	192 (100)

729

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731 **Table 3** - Values of Accuracy, Sensitivity, Specificity, Positive Predictive Value, Negative
732 Predictive Value, Efficiency, Positive Likelihood Ratio, and Negative Likelihood Ratio for
733 SRHET in Relation to the Presence of ET and AP

	ET (n = 198)	PA (n = 192)
Accuracy	.858	.474
Sensitivity	.954	.739
Specificity	.671	.250
Positive predictive value	.850	.454
Negative predictive value	.882	.530
Efficiency	.812	.494
Positive likelihood ratio	2.899	0.985
Negative likelihood ratio	0.068	1.044

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735

736 **Table 4** - Contingency table correlating the presence of ET and AP; n(%). n = 192

Presence of ET	Presence of AP		Total
	Yes	No	
Yes	61 (48.4)	65 (51.6)	126 (65.6)
No	27 (40.9)	39 (59.1)	66 (34.4)
Total	88 (45.8)	104 (54.2)	192 (100)

737

738 3 CONSIDERAÇÕES FINAIS

739

740 O histórico de tratamento endodôntico autorreportado (HTEAR) é uma
741 ferramenta que tem se mostrado acuraz para a identificação de indivíduos com
742 experiência de tratamento endodôntico em diferentes populações. O HTEAR se
743 apresenta, portanto, como uma alternativa viável para a substituição das radiografias
744 em estudos epidemiológicos em Endodontia. Por outro lado, este teste não tem se
745 demonstrado capaz de identificar a presença de periodontite apical. Para possibilitar
746 a apuração desta doença através de instrumentos de autorrelato parece ser
747 necessário desenvolver um conjunto maior de perguntas sobre fatores de risco
748 endodôntico.

749 A maior parte dos valores calculados para os testes de validação (acurácia,
750 sensibilidade, especificidade, valores preditivos positivos e negativos, eficiência,
751 razões de verossimilhança positivas e negativas e correlações de Pearson) do
752 HTEAR variaram nesta população, especialmente em relação à presença de PA, em
753 comparações com amostras de diferentes perfis sociodemográficos. Dessa forma, o
754 HTEAR parece ser sensível a mudanças nas características da população e deve,
755 portanto, ser testado em outros contextos populacionais para permitir seu uso em
756 futuros estudos epidemiológicos de larga escala que incluam variáveis endodônticas.

757 **4 REFERÊNCIAS BIBLIOGRÁFICAS**

758

759 ABBOOD, H. M. et al. Validity of Self-Reported Periodontal Disease: A Systematic
760 Review and Meta-Analysis. **Journal of Periodontology**, v. 87, n. 12, p. 1474–1483,
761 2016.

762 ALONSO, A. et al. Validation of self reported diagnosis of hypertension in a cohort of
763 university graduates in Spain. **BMC public health**, v. 5, p. 94, 2005.

764 AXELSSON, G.; HELGADÓTTIR, S. Comparison of oral health data from self-
765 administered questionnaire and clinical examination. **Community Dentistry and**
766 **Oral Epidemiology**, v. 23, n. 6, p. 365–368, 1995.

767 BAELUM, V. et al. The validity of self-reported dental agenesis. **European Journal**
768 **of Oral Sciences**, v. 119, n. 4, p. 282–287, 2011.

769 BECK, J. et al. Periodontal disease and cardiovascular disease. **J. Periodontol.**, v.
770 67, n. 10 Suppl, p. 1123–1137, 1996.

771 BERGMANN, M. et al. Agreement of self-reported medical history: comparison of an
772 in-person interview with a self-administered questionnaire. **European Journal of**
773 **Epidemiology**, v. 19, n. 5, p. 411–416, 2004.

774 BLICHER, B.; JOSHIPURA, K.; EKE, P. Validation of self-reported periodontal
775 disease: a systematic review. **Journal of Dental Research**, v. 84, n. 10, p. 881–890,
776 2005.

777 BOSSUYT, P. M. et al. Towards complete and accurate reporting of studies of
778 diagnostic accuracy: the STARD initiative. **British Medical Journal**, v. 326, p. 41–
779 44, 2003a.

780 BOSSUYT, P. M. et al. The STARD Statement for Reporting Studies of Diagnostic
781 Accuracy: Explanation and Elaboration. **Clinical Chemistry**, v. 49, n. 1, p. 7–18,
782 2003b.

783 BUHLIN, K. et al. Validity and limitations of self-reported periodontal health.
784 **Community dentistry and oral epidemiology**, v. 30, n. 6, p. 431–7, 2002.

785 CANADA. **Report on the Findings of the Oral Health Component of the**
786 **Canadian Health Measures Survey, 2007-2009**. [s.l.] Health Canada, 2010.

787 CAPLAN, D. J. et al. Lesions of Endodontic Origin and Risk of Coronary Heart
788 Disease. **Journal of Dental Research**, v. 85, n. 11, p. 996–1000, 2006.

789 CAPLAN, D. J. et al. The relationship between self-reported history of endodontic
790 therapy and coronary heart disease in the Atherosclerosis Risk in Communities
791 Study. **Journal of the American Dental Association**, v. 140, n. 8, p. 1004–1012,
792 2009.

793 CHHIBBER-GOEL, J. et al. Linkages between oral commensal bacteria and
794 atherosclerotic plaques in coronary artery disease patients. **NPJ Biofilms and**
795 **Microbiomes**, v. 2, n. 1, p. 7, 2016.

- 796 COBURN, B. W. et al. Performance of self-reported measures for periodontitis in
797 rheumatoid arthritis and osteoarthritis. **Journal of periodontology**, v. 86, n. 1, p. 16–
798 26, 2015.
- 799 COTTI, E. et al. Can a chronic dental infection be considered a cause of
800 cardiovascular disease? A review of the literature. **International Journal of**
801 **Cardiology**, v. 148, n. 1, p. 4–10, 2011a.
- 802 COTTI, E. et al. Association of endodontic infection with detection of an initial lesion
803 to the cardiovascular system. **Journal of Endodontics**, v. 37, n. 12, p. 1624–1629,
804 2011b.
- 805 CYRINO, R. M. et al. Evaluation of Self-Reported Measures for Prediction of
806 Periodontitis in a Sample of Brazilians. **Journal of Periodontology**, v. 82, n. 12, p.
807 1693–1704, 2011.
- 808 DAYA, S. Study Design for the Evaluation of Diagnostic Tests. **Seminars in**
809 **Reproductive Endocrinology**, v. 14, n. 2, p. 111–118, 1996.
- 810 DESTEFANO, F. et al. Dental disease and risk of coronary heart disease and
811 mortality. **British Medical Journal**, v. 306, n. 6879, p. 688–91, 1993.
- 812 DIETRICH, T. et al. The accuracy of individual self-reported items to determine
813 periodontal disease history. **European Journal of Oral Sciences**, v. 113, n. 2, p.
814 135–140, 2005.
- 815 DORN, B. R. et al. Invasion of vascular cells in vitro by *prophyromonas endodontalis*.
816 **International Endodontic Journal**, v. 35, p. 366–371, 2002.
- 817 DOUGLASS, C.; BERLIN, J.; TENNSTEDT, S. The validity of self-reported oral
818 health status in the elderly. **Journal of Public Health Dentistry**, v. 51, n. 4, p. 220–
819 222, 1991.
- 820 DREGAN, A. et al. Chronic inflammatory disorders and risk of type 2 diabetes
821 mellitus, coronary heart disease, and stroke : A population-based cohort study.
822 **Circulation**, v. 130, n. 10, p. 837–844, 2014.
- 823 EKE, P. I. et al. Self-reported measures for surveillance of periodontitis. **Journal of**
824 **dental research**, v. 92, n. 11, p. 1041–7, 2013.
- 825 FARMER, J. et al. Comparing self-reported and clinically diagnosed unmet dental
826 treatment needs using a nationally representative survey. **Journal of Public Health**
827 **Dentistry**, p. 1–7, 2017.
- 828 FRANCISCATTO, G. J. et al. Validation of self-reported history of root canal
829 treatment in a South Brazilian subpopulation. **Brazilian Oral Research**, in press,
830 2019.
- 831 GENCO, R. J. et al. Validity of self-reported measures for surveillance of periodontal
832 disease in two western New York population-based studies. **The Journal of**
833 **periodontology**, v. 78, n. 7 Suppl, p. 1439–1454, 2007.
- 834 GILBERT, G.; DUNCAN, R.; KULLEY, A. Validity of self-reported tooth counts during
835 a telephone screening interview. **Journal of public health dentistry**, v. 57, n. 3, p.

- 836 176–80, 1997.
- 837 GOMES, M. S. et al. Validity of self-reported history of endodontic treatment in the
838 Baltimore longitudinal study of aging. **Journal of Endodontics**, v. 38, n. 5, p. 589–
839 593, 2012.
- 840 GOMES, M. S. et al. Can apical periodontitis modify systemic levels of inflammatory
841 markers? A systematic review and meta-analysis. **Journal of Endodontics**, v. 39, n.
842 10, p. 1205–1217, 2013.
- 843 GOMES, M. S. et al. Apical periodontitis and incident cardiovascular events in the
844 Baltimore Longitudinal Study of Ageing. **International Endodontic Journal**, v. 49, n.
845 4, p. 334–342, 2016.
- 846 GORDIS, L. **Epidemiology**. 5th edition. Ed. Philadelphia: Elsevier Saunders, 2014.
- 847 HALPERN, L. et al. A protocol to diagnose intimate partner violence in the
848 emergency department. **Journal of trauma**, v. 60, n. 5, p. 1101–1105, 2006.
- 849 HEATON, B. et al. A Clinical Validation of Self-Reported Periodontitis Among
850 Participants in the Black Women’s Health Study. **Journal of Periodontology**, v. 88,
851 n. 6, p. 582–592, 2017.
- 852 HELIÖVAARA, M. et al. Reliability and validity of interview data on chronic diseases.
853 The Mini-Finland Health Survey. **Journal of Clinical Epidemiology**, v. 46, n. 2, p.
854 181–191, 1993.
- 855 HUERTA, J. M. et al. Accuracy of self-reported diabetes, hypertension and
856 hyperlipidemia in the adult Spanish population. DINO study findings. **Revista**
857 **Española de Cardiología**, v. 62, n. 2, p. 143–152, 2009.
- 858 HULLEY, S. B. et al. Delineando a pesquisa clínica: uma abordagem epidemiológica.
859 In: 2ª edição ed. Porto Alegre: Artmed, 2003. p. 203–224.
- 860 IACOPINO, A. Periodontitis and diabetes interrelationships: role of inflammation.
861 **Annals of periodontology / the American Academy of Periodontology**, v. 6, n. 1,
862 p. 125–137, 2001.
- 863 IRWIG, L. Guidelines for Meta-analyses Evaluating Diagnostic Tests. **Annals of**
864 **Internal Medicine**, v. 120, n. 8, p. 667, 1994.
- 865 IRWIG, L. M. et al. Designing Studies to Ensure that Estimates of Test Are
866 Transferable. **British Medical Journal**, v. 324, n. 7338, p. 669–671, 2002.
- 867 JACKSON, J. M. et al. Validity of Diabetes Self-Reports in the Women’s Health
868 Initiative. **Menopause**, v. 21, n. 8, p. 861–868, 2014.
- 869 JOSHIPURA, K. J. et al. Pulpal inflammation and incidence of coronary heart
870 disease. **Journal of Endodontics**, v. 32, n. 2, p. 99–103, 2006.
- 871 JOSHIPURA, K. J.; PITIPHAT, W.; DOUGLASS, C. W. Validation of self-reported
872 periodontal measures among health professionals. **Journal of public health**
873 **dentistry**, v. 62, n. 2, p. 115–121, 2002.
- 874 KÖNÖNEN, M.; LIPASTI, J.; MURTOMAA, H. Comparison of dental information

- 875 obtained from self-examination and clinical examination. **Community dentistry and**
876 **oral epidemiology**, v. 14, n. 5, p. 258–260, 1986.
- 877 LAMONTE, M. J. et al. Accuracy of Self-Reported Periodontal Disease in the
878 Women's Health Initiative Observational Study. **Journal of Periodontology**, v. 85, n.
879 8, p. 1006–1018, 2014.
- 880 LEE, J. et al. Systematic Review and Meta-Analysis of Studies Evaluating Diagnostic
881 Test Accuracy : A Practical Review for Clinical Researchers – Part II . Statistical
882 Methods of. **Korean Journal of Radiology**, v. 16, n. 6, p. 1188–1196, 2015.
- 883 LEVIN, L.; SHPIGEL, I.; PERETZ, B. The use of a self-report questionnaire for dental
884 health status assessment: A preliminary study. **British Dental Journal**, v. 214, n. 5,
885 p. 1–4, 2013.
- 886 MATTILA, M. S.; VALTONEN, V. V.; RASI, V. P.; KESÄNIEMI, Y. A.; SYRJÄLÄ, S.
887 L.; JUNGELL, P. S.; ISOLUOMA, M.; HIETANIEMI, K. ;JOKINEN, M. J. Association
888 between dental health and acute myocardial infarction. **British Medical Journal**, v.
889 298, n. March, p. 779–781, 1989.
- 890 MORRISON, H. I.; ELLISON, L. F.; TAYLOR, G. W. Periodontal disease and risk of
891 fatal coronary heart and cerebrovascular diseases. **Journal of Cardiovascular Risk**,
892 v. 6, n. 1, p. 7–11, 1999.
- 893 MÜLLER, M. M.; ANDREA, G. Markers of Endothelial Dysfunction. **Clinical**
894 **Chemistry and Laboratory Medicine**, 2000. Disponível em:
895 <[https://www.degruyter.com/view/j/cclm.2000.38.issue-](https://www.degruyter.com/view/j/cclm.2000.38.issue-2/cclm.2000.013/cclm.2000.013.xml)
896 [2/cclm.2000.013/cclm.2000.013.xml](https://www.degruyter.com/view/j/cclm.2000.38.issue-2/cclm.2000.013/cclm.2000.013.xml)>
- 897 NILSSON, I.-M.; LIST, T.; DRANGSHOLT, M. The Reliability and Validity of Self-
898 reported Temporomandibular Disorder Pain in Adolescents. **Journal of Orofacial**
899 **Pain**, v. 20, n. 2, p. 138–144, 2006.
- 900 PALMQVIST, S.; B, S.; ARNBJERG, D. Self-assessment of dental conditions: validity
901 of a questionnaire. **Community Dentistry and Oral Epidemiology**, v. 19, n. 5, p.
902 249–251, 1991.
- 903 PASQUALINI, D. et al. Association among oral health, apical periodontitis, CD14
904 polymorphisms, and coronary heart disease in middle-aged adults. **Journal of**
905 **Endodontics**, v. 38, n. 12, p. 1570–1577, 2012.
- 906 PETERSEN, J. et al. The association of chronic apical periodontitis and endodontic
907 therapy with atherosclerosis. **Clinical Oral Investigations**, v. 18, n. 7, p. 1813–1823,
908 2014.
- 909 PETERSEN, P.; BAEZ, R. **Oral Health Suverys Basic Methods**. WHO, 2013.
- 910 PITIPHAT, W. et al. Validation of self-reported oral health measures. **Journal of**
911 **public health dentistry**, v. 62, n. 2, p. 122–128, 2002.
- 912 RAMOS, R. Q.; BASTOS, J. L.; PERES, M. A. Diagnostic validity of self-reported oral
913 health outcomes in population surveys : literature review Validade diagnóstica de
914 agravos. **Rev Bras Epidemiol**, v. 16, n. 3, p. 716–728, 2013.

- 915 RIMM, E. B. et al. Reproducibility and Validity of an Expanded Self-Administered
916 Semiquantitative Food Frequency Questionnaire among Male Health Professionals.
917 **American Journal of Epidemiology**, v. 135, n. 10, p. 1114–1126, 15 maio 1992.
- 918 ROSS, R. Inflammation or Atherogenesis. **The New England Journal of Medicine**,
919 v. 340, n. 2, p. 115–126, 1999.
- 920 SEGURA-EGEA, J.; MARTÍN-GONZÁLEZ, J.; CASTELLANOS-COSANO, L.
921 Endodontic medicine: Connections between apical periodontitis and systemic
922 diseases. **International Endodontic Journal**, v. 48, n. 10, p. 933–951, 2015.
- 923 SILVA, A. E. R. et al. Validation of self-Reported information on dental caries in a
924 birth cohort at 18 years of age. **PLoS ONE**, v. 9, n. 9, p. 1–8, 2014.
- 925 TAYLOR, A. et al. Comparing self-reported and measured high blood pressure and
926 high cholesterol status using data from a large representative cohort study.
927 **Australian and New Zealand Journal of Public Health**, v. 34, n. 4, p. 394–400,
928 2010.
- 929 TORMO, M. J. et al. Validation of self diagnosis of high blood pressure in a sample of
930 the Spanish EPIC cohort: overall agreement and predictive values. EPIC Group of
931 Spain. **Journal of epidemiology and community health**, v. 54, n. 3, p. 221–6,
932 2000.
- 933 VETTER, T. R.; SCHOBBER, P.; MASCHA, E. J. Diagnostic Testing and Decision-
934 Making. **Anesthesia & Analgesia**, v. 127, n. 4, p. 1, 2018.
- 935 VON STEIN-LAUSNITZ, M. et al. Patients' self-report on post-retained restoration is
936 more valuable than expected! Explorative analysis of an 11-year follow-up. **Acta**
937 **Odontologica Scandinavica**, v. 0, n. 0, p. 1–6, 2018.
- 938 WHO. **Global status report on noncommunicable diseases 2010**.
- 939 WHO. **Global action plan for the prevention and control of noncommunicable**
940 **diseases 2013-2020**.
- 941 WHO. **Key Facts on Noncommunicable diseases**. Disponível em:
942 <<http://www.wpro.who.int/mediacentre/factsheets/fs20130311/en/>>. Acesso em: 30
943 set. 2017b.
- 944 WHO. **Global status report on noncommunicable diseases 2014**World Health.
- 945 WONG, A. L. et al. Development and evaluation of a patient self-report case-finding
946 method for rheumatoid arthritis. **Seminars in Arthritis and Rheumatism**, v. 34, n. 1,
947 p. 484–499, 2004.
- 948 YACOUBIAN JR, G.; WISH, E. Exploring the validity of self-reported Ecstasy use
949 among club rave attendees. **Journal of Psychoactive Drugs**, v. 38, n. 1, p. 31–34,
950 2006.