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Fonseca anamnestic index for screening Temporomandibular Disorders - reliability to discriminate muscular from intra-articular disorders

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1 **Fonseca anamnestic index for screening Temporomandibular Disorders - reliability to discriminate**
2 **muscular from intra-articular disorders.**

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16 **Abstract:**

17

18 **Background/ Objective:** Fonseca anamnestic index (FAI) is a simple and quick survey used for screening the
19 presence and severity of Temporomandibular Disorders (TMD). The presented study aimed to screen the FAI
20 accuracy to discriminate different types of TMD: intra-articular (AD), Masticatory Muscular Disorder (MMD),
21 or the presence of both typologies.

22 **Methods:** The existence of a pattern in the FAI based on the frequency of answers was evaluated and supported
23 by other variables: sex, age, medical diagnosis and Visual Analog Scale of health-related quality of Life
24 (VASLife). The non-parametric Chi-square test (χ^2) or Fisher's exact test were used to assess the existence of
25 associations between these variables. In the pairs of variables where such association was identified, its
26 intensity was measured by Cramér's V Coefficient. The prediction if FAI could be a good decision tool for
27 distinguish the type of TMD was assessed through logistic regression models (ordinal and multinomial).

28 **Results:** The higher FAI score was associated with questions related with temporomandibular joint (TMJ)
29 pain, TMJ clicks and person anxiety. Severe cases classified by FAI are correlated with typology of Both
30 (AD+MMD). Moreover, the female patients presented more moderate and severe cases in FAI and also
31 correlated with the presence of AD+MMD. The logistic model showed low accuracy to distinguish the TMD
32 typology (~70%).

33 **Conclusion:** FAI is a good initial methodology in TMD diagnosis, however integrated in a logistic regression
34 model for distinguish the typology of TMD has proved to be insufficient. It is expected that the combination
35 of this survey with other outcomes will make the model more accurate.

36

37 **Keywords:**

38 Fonseca Anamnestic Index; Intra-articular temporomandibular Disorders; Masticatory muscle
39 temporomandibular disorders; Multinomial logistic regression; Patient-reported questionnaire;
40 Temporomandibular Disorders.

41

42 **Introduction:**

43 Temporomandibular disorders (TMD) are a set of musculoskeletal and/or articular conditions that affect
44 respectively the masticatory musculature and/or the temporomandibular joint (TMJ) complex. TMD is the
45 most common nondental cause of orofacial pain and have a negative impact on the patient's daily life [1].
46 Epidemiologically it is known that this disease affects mainly females (70-85%) [2]. TMD present a
47 multifactorial etiology and due to its complexity represents a real challenge for clinicians in terms of a correct
48 diagnosis [3]. The two main origins of pain in this region are associated with intra-articular or masticatory
49 muscle changes. Actually, the diagnosis of TMD is largely based on the patient's symptoms, as pain in TMJ
50 and surrounding muscles, difficulty in opening the mouth, and other complaints such as the presence of
51 clicking in the joint, malocclusion and headaches. Clinical observation evaluates different parameters such as
52 the presence of joint inflammation (synovitis), measurement of mouth opening and laterality of jaw
53 movements, dental occlusion, the presence of clicks and crepitus in the joint and muscle tenderness. Definitive
54 diagnosis is normally performed with medical imaging support, using computed tomography (CT), magnetic
55 resonance imaging (MRI) or minimally invasive diagnostic interventions [4].

56 The Fonseca Anamnestic Index (FAI) is a TMD patient-reported questionnaire, quick and easy to administer,
57 based in signs and symptoms with 10 questions, used in recent years to classify the severity of TMD [5-8].

58 The final score obtained can be interpreted using a classification table that assigns each individual one of four
59 possible categories of severity: no TMD ($0 \leq \text{FAI} \leq 15$ points); mild TMD ($20 \leq \text{FAI} \leq 40$ points); moderate TMD
60 ($45 \leq \text{FAI} \leq 65$ points) and severe TMD ($70 \leq \text{FAI} \leq 100$ points) [5]. However, it is unknown if the score obtained
61 by this survey can contribute to a correct TMD diagnosis regarding three possible typologies: Articular
62 Disorder (AD), Masticatory Muscle Disorder (MMD) or both. The aim of this study is to identify patterns in
63 the FAI, together with the characteristics of the patients, allow us to assess the robustness of this questionnaire
64 as an aid in the clinical diagnosis of TMD.

65

66 **Methods:**67 **Study Design**

68 A retrospective study was conducted in a private health institution in Portugal (Instituto Português da Face),
69 including patients diagnosed with TMD from January of 2019 to March 2022. This study was approved by the
70 *Instituto Português da Face* ethics committee (IPF/08/22). All enrolled patients gave their informed consent
71 in writing, following current legislation. The inclusion criteria was: (1) age >18 years; (2); full response to
72 FAI; (3) clinical diagnosis of TMD. The exclusion criteria included was: (1) a history of facial trauma or other
73 orofacial disorder; (2) severe medical problems or impaired cognitive capacity; (3) pregnant or breastfeeding
74 women. All patients were examined by the same doctor. The information was recorded and stored in a database
75 (EUROTMJ). Confidentiality of information is ensured through anonymity. Demographic data for all patients
76 like: sex and age was registered. As an initial diagnosis of the presence of a TMD, the patients were instructed
77 to answer the FAI. The survey was applied in Portuguese, which is already validated in the literature [9], and
78 was subsequently translated into English. The FAI is a Likert scale questionnaire based in 10 questions with
79 three points/levels ("No", "Sometimes", "Yes") (Table S1). In FAI, answers are scored as follows: no - 0 points,
80 sometimes - 5 points, yes - 10 points. The final score (0-100) was classified into the following categories: no
81 TMD ($0 \leq \text{FAI} \leq 15$ points); mild TMD ($20 \leq \text{FAI} \leq 40$ points); moderate TMD ($45 \leq \text{FAI} \leq 65$ points) and severe
82 TMD ($70 \leq \text{FAI} \leq 100$ points). Additionally, the Visual Analog Scale (VAS) has been used in the valuation of
83 health-related quality of Life (VASLife) with the question: "If you could give a life impact score to your TMJ
84 problem in a 0 to 10 scale, where 0 means no impact and 10 means the maximum impact possible, what would
85 be your score?"[10]. The identification of the type of temporomandibular disease (MMD, AD, Both) was
86 performed by the clinician through medical evaluation and MRI to assess intra-articular derangements. To
87 access the MMD derangements, muscle tenderness was measured using a 0-3 classification as defined in
88 Research Diagnostic Criteria for Temporomandibular Disorders (RDC/TMD) [4, 11].

89

90 **Statistical Analysis**

91 The initial methodology was to identify the existence of a pattern in the FAI based on the frequency of answers
92 in each of its three levels (No; Sometimes; Yes) in the ten questions (Table S1). This analysis was supported
93 by a descriptive study of the following variables: sex, age, medical diagnosis and score in VASLife. The mean
94 was presented as the location measure accompanied by its standard deviation (SD) in the form $\text{mean} \pm \text{SD}$. The
95 normality in the distributions of the FAI and VASLife (BevilaquaGrossi et al, 2006) was determined. Given
96 the absence of normality, Spearman's correlation coefficient (r_s) was used to determine the correlation
97 between the FAI and VASLife scales. The classification of the correlation was determined in accordingly with
98 Davis [12]. Subsequently, based on each patient's FAI score, bivariate contingency tables were created
99 containing the absolute frequency in each of the possible combinations of categories in the following pairs of
100 variables: TMD severity vs sex/diagnosis; diagnosis vs FAI levels for each of the items. The non-parametric
101 Chi-square test (χ^2) and Fisher's exact test were used to assess the existence of associations between these
102 variables. In the pairs of variables where such association was identified, its intensity was measured by
103 Cramér's V Coefficient (ϕ_c). To compare more than two groups was used the non-parametric Kruskal-Wallis

104 test and when there was statistical significance the Mann-Whitney (W) test was used for pair-wise comparison.
 105 Finally, the extent to which the FAI could be a good decision tool for distinguish the type of TMD was assessed
 106 through logistic regression models (ordinal and multinomial) [13]. To avoid bias, the final sample (171
 107 patients) was subdivided into two data sets: one for training (adjustment) with 70% of the information and
 108 another for testing (validation and prediction) with the remaining. The response variable considered in logistic
 109 regression models (ordinal and multinomial) was type of TMD, constituted by mutually exclusive classes (each
 110 patient is assigned to only one class), with the following ordering based on the complexity of the TMD
 111 typology: MMD<AD<Both. Moreover, the independent variables of the regression models were: FAI, SEX,
 112 Age and VASLife. The following models were considered: Model 1: $TMD \sim \beta_1 FAI + \varepsilon$; Model 2: $TMD \sim$
 113 $\beta_1 FAI + \beta_2 SEX + \varepsilon$; Model 3: $TMD \sim \beta_1 FAI + \beta_2 SEX + \beta_3 Age + \varepsilon$; Model 4: $TMD \sim \beta_1 FAI +$
 114 $\beta_2 SEX + \beta_3 Age + \beta_4 VASLife + \varepsilon$. For the models considered, the presence of multicollinearity of the
 115 predictors was tested through the Variance Inflation Factor (VIF). In a first approach, ordinal logistic
 116 regression models were considered. If the assumption of proportional hazards by Brant [12] is not violated the
 117 choice was ordinal logistic regression models, otherwise the multinomial logistic regression models were
 118 adopted. For this last class of models the independence of irrelevant alternatives was tested by the Hausman-
 119 McFadden test (p-values \approx 1) [17]. The model that presented the lowest Akaike Information Criterion(AIC)
 120 value and the highest Nagelkerke's pseudo R² value was adopted as a selection criterion. The accuracy of the
 121 model was also presented as well as its degree of agreement expressed by Kappa coefficient (Poor<0.00;
 122 Slight:0.00-0.20; Fair:0.21-0.40; Moderate:0.41-0.60; Substantial:0.61-0.80; Almost Perfect:0.81-1.00) and
 123 classified according to Landis and Koch [14].
 124 The significance level set was 5% and all statistical treatment and graphical representation was performed in
 125 the R programming language [15].

126 **Results:**

127 A total of 541 patients (80% of female), with a mean age of 39.543 ± 15.951 years, who answered the Fonseca
 128 questionnaire were included in the study. FAI score was 58.822 ± 21.313 . Initially, an assessment of the
 129 frequency of answers and mean score of the different FAI questions was performed. The items 7, 6, 10 had the
 130 highest mean score on the FAI, which reflects a higher concentration of answers at the last level of the scale
 131 (Yes) (Figure 1). In opposition, the items with the lowest mean score were 1, 9 and 2, respectively, reflected
 132 by the less frequent answers in the highest score (Yes) (Figure 1).

133 In second part of study, 171 patients with clinical diagnosis (between 18 and 90 years old), were included. 30
 134 patients were diagnosed with MMD and 33 with AD and 108 with both. Patients had a mean age of
 135 38.444 ± 16.172 years, 140 of whom were female (82%). Female patients had a higher mean age (39.200) than
 136 male patients (35.032), although without statistical significance (Mann-Whitney (W) = 2513.500; p= 0.169).
 137 The average pain impact on patients' lives (VASLife) was 6.525 ± 2.423 , with females having higher averages

138 than males (6.781 vs 5.433, $W=2579.500$, $p= 0.003$). The global FAI mean was 60.380 ± 21.337 , being
 139 differentially expressed between females and males (62.071 vs 52.742, respectively, $W=2744.500$, $p= 0.021$).
 140 Initially, an analysis was performed between the two scales, VASLife and FAI. A moderate positive
 141 correlation between the two scales was verified ($r_s=0.358$; $p<0.001$), i.e. an increase in the FAI score is
 142 accompanied by an increase in the VASLife scale.

143 Subsequently, the existence of an association between the type of diagnosis (MMD, AD, Both) and the level
 144 of the FAI (No, Sometimes, Yes) was assessed. Once the conditions for applicability of the Chi-Square test
 145 (χ^2), were verified, i.e. “expected values $e_{ij} > 1$ and $80\% e_{ij} > 5$ ”, it was determined that there was a statistically
 146 significant relationship between these variables ($\chi^2 = 46.413$, $df=4$, $p<0.001$) (Table 1). Cramér's V coefficient
 147 assumed a value of 0.116 and is classified as moderate ($CI_{95\%}^{p_c}:[0.078;0.147]$).-The relative frequency of
 148 diagnosis per item of the FAI at each level was then analyzed (Table S2). This results seems to indicate that:
 149 a) in MMD diagnosis the items that present higher relative frequency was the items 4,5,6 and 8; b) the AD and
 150 Both diagnosis presented the higher relative frequency in items 6 and 7.. Furthermore, Both (MMD+AD)
 151 diagnosis presented the highest FAI scores comparatively to MMD and AD (Kruskal-Wallis chi-squared =
 152 16.734, $df = 2$, $p<0.001$; Both vs AD, $p=0.001$; and Both vs MMD, $p=0.019$) (Figure 2).

153 Analogously, the analysis was carried out considering the sex and diagnosis of the patient with the severity of
 154 the FAI (no severe, mild, moderate, severe). Once the conditions for the applicability of the χ^2 were violated,
 155 we used Fisher's exact test, which seems to indicate that there were statistically significant relationships
 156 between these pairs of variables ($p=0.050$; $p<0.001$ respectively, Table 2). Cramér's V coefficient assumed the
 157 values of 0.208 ($CI_{95\%}^{p_c}:[0.009;0.335]$) and 0.245 ($CI_{95\%}^{p_c}:[0.092;0.325]$) classifying the intensity as strong
 158 and very strong [16].

159 The frequency distribution of the number of diagnosed cases and the FAI score by type of TMD according to
 160 sex was then checked (Figure 3A and B). In the case of female patients there is a prevalence of diagnosis of
 161 MMD+AD (Both) (69%), while in the opposite sex AD (52%) is the most prevalent. In Figure 3B in females
 162 FAI score was significantly higher in Both diagnosis comparatively to MMD (Kruskal-Wallis chi-squared =
 163 7.337, $df = 2$, $p= 0.026$; Both vs MMD, $p=0.046$), while in men this profile was not verified and it is not
 164 possible to draw the same conclusion. Finally, we sought to assess to what extent the FAI could be a good
 165 predictor of the type of TMD diagnosis using the ordinal logistic regression model. Once the absence of
 166 multicollinearity in the predictors confirmed by the VIF (FAI:1.174;SEX:1.070;AGE:1.028;VASLife:1.208),
 167 the null hypothesis of proportionality of risks was rejected in Models 1 to 4 (p -values 0.051; 0.000; 0.011;
 168 0.010, respectively) leading to the approach by multinomial logistic regression. The predictors AGE and
 169 VASLife did not show any statistical significance (Model 3: AD:AGE p -value=0.307; Both:AGE p -
 170 value=0.357; Model 4: AD:AGE p -value=0.305; Both:AGE p -value=0.337;AD:VASLife p -value=0.783;
 171 Both:VASLife p -value=0.125) and the choice between Models 1 and 2 was made. Analysis of deviance table
 172 revealed that both predictors in models 1 and 2 are statistically significant (Model 1: FAI, likelihood-ratio χ^2
 173 =15.764, $df=2$, p -value= <0.001 ; Model 2: FAI, likelihood-ratio $\chi^2 =12.132$, $df=2$, p -value <0.001 , SEX

174 likelihood-ratio $\chi^2 = 17.694$, $df=2$, $p\text{-value} < 0.001$). Since Model 2 has a lower AIC value relatively to Model
 175 1 (196.732 vs 204.601) and higher Nagelkerke pseudo R^2 (0.301 vs 0.213), this model was chosen. The
 176 accuracy of the model is 0.667 ($CI_{95\%}: [0.580, 0.754]$) and the level of agreement expressed by the Kappa
 177 coefficient is 0.230 being classified as fair [14]. In the model test the accuracy value was 0.629 ($CI_{95\%}: [0.449,$
 178 $0.785]$) with a Kappa agreement level of 0.187. The Figure 3C represents the adjusted logistic model.
 179 The adjusted Model 2 can be expressed by the following set of equations:

$$180 \quad \log \left[\frac{P(Y = AD)}{P(Y = MMD)} \right] = 0.618 - 0.012FAI + 1.202SEX.Male \quad (\text{Equation 1})$$

$$181 \quad \log \left[\frac{P(Y = Both)}{P(Y = MMD)} \right] = -0.463 + 0.027FAI - 0.560SEX.Male \quad (\text{Equation 2})$$

182 Discussion:

183 TMD continue to represent a clinical challenge in diagnosis due to complex muscle and intra-articular
 184 involvement. Thus, a precise diagnosis of TMD is crucial and has been the object of a large number of studies.
 185 The RDC/TMD classification continues to be the most widely used in clinical practice, contributing
 186 significantly to a standardization of diagnosis [17, 18]. However, at a practical level, it has implementation
 187 disadvantages, being time-consuming, difficult in data collection and requiring extensive clinical experience
 188 [19, 20]. On the other hand, FAI is a simple questionnaire to implement and is a useful initial tool to distinguish
 189 the presence of a TMD and the degree of severity. This tool has been the subject of study in various scientific
 190 studies [5-8]. However, the accuracy of FAI to distinguish the possible origin of TMD, muscular or intra-
 191 articular, is still unknown. Thus, this study aimed to identify the behavior of the FAI in a set of patients
 192 diagnosed with different TMD.

193 Primarily we found that the items relatively to clicks in TMJ (item 7), pain in TMJ area (item 6) and muscle
 194 surrounding (item 3) are among the biggest contributors to higher values on the FAI . Indeed, pain seems to
 195 have a strong impact on FAI, corroborated by the moderate correlation of FAI with the VASLife scale.
 196 Interestingly, being an anxious person can also be crucial for high FAI scores (item 10). On the other hand,
 197 the difficulty in items related to mandibular movements, either laterally or vertically, as well as the difficulty
 198 in touching the teeth were the least determinant factors (items 2, 9 and 1) . Importantly, it has also been shown
 199 that higher FAI scores are correlated with a more complex diagnosis with the simultaneous presence of muscle
 200 and intra-articular changes (Both) . In fact, it is possible to verify a greater number of moderate and severe
 201 cases of the FAI, when muscular changes are verified simultaneously with intra-articular alterations.
 202 Interestingly in MMD diagnosis was found a pattern of higher relative frequency in the items related with pain
 203 (headache, neck pain and TMJ pain) and parafunctional habits (clenching or grinding). On other hand, in AD
 204 and Both was verified a higher predominantly of positive answers in items related with pain and noise in TMJ
 205 area. This results, showed an alteration in higher relative frequency of FAI items when are present intra-
 206 articular derangements. At the same time, a different distribution of severity of cases in females and males was

207 demonstrated. In females, MMD and MMD+AD were the TMD types with the lowest and highest FAI scores,
208 respectively. In addition, there was a higher distribution of cases with both diagnoses in females. This trend
209 was not equally demonstrated in males. Previous studies have pointed that female sex has a higher prevalence
210 of TMD, around 80% [2, 21]. Although not entirely certain, hormonal imbalance in females may be related to
211 increased susceptibility to TMD [22]. Beyond these data, this study also showed that there is a growing trend
212 towards more severe cases in females. A logistic model was conducted and following conclusions were drawn:
213 (i) there is a higher probability of a patient being diagnosed with MMD, although low, have lower FAI final
214 scores; (ii) the probability of a patient being diagnosed with AD with low FAI scores is much higher in men
215 compared to women, a difference that is attenuated as FAI scores increase; (iii) when a patient is diagnosed
216 with both AD and MMD simultaneously, the FAI scores show an increasing behavior, which is identical in
217 both genders (parallel lines).

218 However, the present model with FAI as predictor showed an accuracy <70%, which means that for every 100
219 diagnoses made, at most, 70 are expected to be correctly classified. In clinical terms, a higher accuracy is
220 desirable. Additionally, the number of diagnoses between the three groups has different values, with the
221 MMD+AD (107) three times higher compared to the other diagnoses (~30). In addition, the number of final
222 diagnoses (171) represents around 32% of the total number of records (539). The authors consider that a higher
223 number of cases with final diagnosis may allow a better illustration of the role of FAI in the distinction of
224 disease typology.

225 In conclusion, FAI is an important tool in the diagnosis of TMD, however a more complex model is needed to
226 more accurately distinguish the type of TMD. In the future, it is expected that other complementary measures
227 and scales will be incorporated to strengthen the model. This study also demonstrated a differential behavior
228 of the FAI between the two sexes. Thus, clinicians should take sex into consideration when using this tool.

229 **Ethics committee and informed consent:**

230 The current research was approved by an independent ethics committee and subjects gave their informed
231 consent before they were enrolled in the study.

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294

295 **Tables:**

296 **Table 1. Contingency table relatively to the type of diagnosis and the level of Fonseca Anamnestic Index**
 297 **(FAI).** MMD-Masticatory Muscle Disorder; AD-Articular Disorder; χ^2 - Chi-Square test; df- degrees of
 298 freedom, O_{ij} - observed value in row i (i=MMD, AD, Both) and column j (j=No, Sometimes, Yes); e_{ij} - expected
 299 value in the row i (i=MMD, AD, Both) and column j (j=No, Sometimes, Yes);

300

FAI	No	Sometimes	Yes	χ^2; df; p-value
Diagnostic				
MMD	$O_{11}=83$ $e_{11}=70.351$	$O_{12}=109$ $e_{12}=97.018$	$O_{13}=108$ $e_{13}=132.632$	46.413; 4; <0.001
AD	$O_{21}=108$ $e_{21}=77.386$	$O_{22}=113$ $e_{22}=106.719$	$O_{23}=109$ $e_{23}=145.895$	
Both	$O_{31}=210$ $e_{31}=253.263$	$O_{32}=331$ $e_{32}=349.263$	$O_{33}=539$ $e_{33}=477.474$	

301

302 **Table 2. Contingency table regarding sex and type of diagnosis compared to the severity level of the**
 303 **Fonseca Anamnestic Index (FAI) determined by Fisher-Exact test.** F- Female; M- Male; MMD-
 304 Masticatory Muscle Disorder; AD-Articular Disorder.

305

Sex	Severity				p-value
	No Severe	Mild	Moderate	Severe	
F	1	26	54	59	0.050
M	0	12	12	7	
Diagnostic					<0.001
MMD	0	8	17	5	
AD	0	13	13	7	
Both	1	17	36	54	

306

307 **Figures legends:**

308 **Figure 1. Distribution of the scores by levels of the Fonseca Anamnestic Index (FAI).** The 2ndcolumn
 309 corresponds to the sample mean and standard deviation (SD) for each FAI item. The 3rd and remaining columns
 310 show the concentration of the score by FAI level. Higher concentrations are accompanied by a more intense
 311 violet/purple coloration.

312

313 **Figure 2. Dispersion of the Fonseca Anamnestic Index (FAI) by type of diagnosis.** The asterisk inside the
314 boxplot symbolizes the mean score. MMD-Masticatory Muscle Disorder; AD-Articular Disorder.

315

316 **Figure 3. Analysis of the type of temporomandibular disorders (TMD) as a function of the Fonseca**
317 **Anamnestic Index (FAI) taking sex into account.** A. Percentage of the number of TMD cases by sex. B.
318 Dispersion of FAI score by type of diagnosis and sex. C. Graphical representation of the multinomial logistic
319 regression model.

320

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