

# The prevalence of smartphone, nicotine, and alcohol addiction among university students with temporomandibular disorders

Received: 2 May 2025

Accepted: 7 January 2026

Published online: 17 January 2026

Cite this article as: Kaynak B.A., Yüzbaşıoğlu Ü., Ekici E. *et al.* The prevalence of smartphone, nicotine, and alcohol addiction among university students with temporomandibular disorders. *BMC Oral Health* (2026). <https://doi.org/10.1186/s12903-026-07698-9>

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1 **The prevalence of smartphone, nicotine, and alcohol**  
2 **addiction among university students with**  
3 **temporomandibular disorders**

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17 **Word Count**

18 **Total:** 4166 Words

**Abstract:** 209 Words

19 **Number of tables:** 6 Tables

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24 **Funding:** This study was conducted without receiving any financial  
25 support

26 **Conflicts of interest:** No declaration of conflicts of interest.

27 **Acknowledgments:** None.

28 **Clinical Trial Registration declaration:** Not applicable.

## 29 **Abstract**

30 **Background:** This cross-sectional study aimed to assess the prevalence of  
31 smartphone, nicotine, and alcohol addiction among university students  
32 with temporomandibular dysfunctions (TMDs) and compare these findings  
33 with a control group.

34 **Methods:** A total of 909 university students participated. The presence of  
35 TMDs was determined according to the Diagnostic Criteria for TMDs and  
36 participants categorized as TMDs and control groups. TMD severity was  
37 assessed via the Fonseca Anamnestic Index (FAI). Nicotine addiction was  
38 evaluated using the Fagerström Test for Nicotine Dependence (FTND),  
39 alcohol dependence with the Addiction Profile Index Risk Scanning Scale  
40 (APIRSS), smartphone addiction with the Smartphone Addiction Index-  
41 Short Form (SPAI-SF), and depression and anxiety levels using the Patient  
42 Health Questionnaire-4 (PHQ-4).

43 **Results:** Nicotine ( $p=0.009$ ), alcohol ( $p=0.004$ ), and smartphone  
44 ( $p<0.001$ ) addictions were more prevalent in students with TMDs. PHQ-4,  
45 SPAI-SF, APIRSS, and FTND scores were significantly higher in this group  
46 ( $p<0.001$ ;  $p=0.003$  for FTND). Among nicotine-addicted participants,  
47 those with TMDs had higher FTND scores than controls ( $p<0.007$ ). FAI  
48 scores were positively correlated with SPAI-SF ( $r=0.283$ ,  $p<0.001$ ), FTND  
49 ( $r=0.147$ ,  $p<0.001$ ), APIRSS ( $r=0.123$ ,  $p<0.001$ ), and PHQ-4 ( $r=0.48$ ,

50  $p < 0.001$ ) scores. PHQ-4 scores were correlated with SPAI-SF ( $r = 0.328$ ,  
51  $p < 0.001$ ), APIRSS ( $r = 0.125$ ,  $p < 0.001$ ), and FTND ( $r = 0.120$ ,  $p < 0.001$ ).

52 **Conclusions:** The findings emphasize the need for a multidimensional  
53 treatment approach in TMDs that consider coexisting addictions.

54 **Keywords:** Temporomandibular Joint Disorders; Nicotine Addiction;  
55 Smartphone Addiction; Alcoholism; Depression; Smoking.

56

## 57 **Background**

58 Temporomandibular disorders (TMDs) constitute a group of conditions  
59 involving different regional and systemic symptoms, including pain and  
60 limitation at the temporomandibular joint (TMJ), with a high prevalence,  
61 ranging from 31% to 75% in the adult population [1, 2]. TMDs have a  
62 multifactorial etiology involving numerous factors including sex, age,  
63 parafunctional habits, psychological issues, and lifestyle [3]. It has been  
64 established that lifestyle is of paramount importance in the management  
65 of TMDs. Consequently, there has been a recent proliferation of literature  
66 examining the relationship between lifestyle and TMDs severity, with an  
67 emphasis on adopting multidisciplinary approaches.

68 Addictions, which play a significant role in the etiology of most diseases  
69 and often complicate treatment, represent a key area of investigation.

70 Among the most prevalent addictions in today's society are nicotine and  
71 alcohol [4]. The observed association between the severity of depression  
72 and anxiety and the prevalence of TMDs may predispose individuals with

73 TMDs to the development of these addictions [5, 6]. A systematic review  
74 reveals a consensus on this point. Miettinen et al.[7] showed that the  
75 regular consumption of alcohol and tobacco products was associated with  
76 increased TMDs symptom severity. Similarly, another study reported  
77 higher TMDs-related pain in smokers compared with non-smokers [8].  
78 Furthermore, with the advent of technological advancement, smartphone  
79 addiction has been added as a global health problem [4, 9]. Although  
80 researchers have postulated a correlation between long-term smartphone  
81 use and TMDs, there is currently no conclusive evidence in the literature  
82 to substantiate this hypothesis. The underlying rationale for this  
83 proposition is the alteration in posture that occurs with prolonged  
84 smartphone use. Individuals who use smartphones for an extended period  
85 exhibit a forward head posture, which places excessive stress on the  
86 cervical spine [9]. This prolonged flexion of the cervical spine, coupled with  
87 the increased weight of the head, can negatively affect the neck muscles,  
88 ligaments, and bony structures [9]. It is therefore hypothesized that the  
89 negative effects on the neck may have consequences for the TMJ [9]. Pei  
90 et al.[10] reported that individuals with smartphone addiction exhibited  
91 more severe and frequent pain and symptoms related to TMDs[10].

92 Demonstrating the effect of addictions, which are highly prevalent in  
93 society, on TMDs will facilitate the development of proactive approaches  
94 for the management of TMDs. The relationship between addiction and  
95 TMDs has been previously investigated in different countries, including  
96 China and Finland [7, 10]. Smartphone, nicotine, and alcohol addiction  
97 related to TMDs may be different between Turkish university students and

98 other populations because addictive behavior is influenced by various  
99 factors, including age and cultural characteristics. Therefore, our primary  
100 purpose was to investigate the prevalence of smartphone, nicotine, and  
101 alcohol addiction among Turkish university students with TMDs.  
102 Furthermore, we sought to compare these findings with those from a group  
103 of healthy individuals. Additionally, we aimed to identify the relationship of  
104 nicotine, smartphone, and alcohol addiction with depression and TMDs  
105 severity. The hypothesis of this study was that Turkish university students  
106 with TMDs would be more likely to exhibit addictive behaviors regarding  
107 nicotine, alcohol, and smartphone use.

## 108 **Methods**

### 109 **Ethical Statement**

110 This study was approved by the Toros University Non-Interventional  
111 Clinical Research Ethics Committee (Protocol Number: 2024/06-09). Prior  
112 to their participation in the study, all subjects were provided with  
113 comprehensive information regarding the nature and scope of the  
114 investigation. Their consent was obtained in an informed manner.  
115 Specifically, both written and verbal consent were required, with written  
116 consent documented through signed forms and verbal consent recorded in  
117 the presence of a witness. Throughout the course of the study, the research  
118 team adhered to the ethical standards set forth in the Declaration of  
119 Helsinki. Additionally, the study was conducted in accordance with the  
120 STROBE (Strengthening the Reporting of Observational Studies in

121 Epidemiology) statement for case-control studies and the STROBE  
122 Checklist for observational studies.

### 123 **Sample Size**

124 A power analysis was conducted to determine the optimal sample size  
125 using the Calculator.net software. A total of 3.744 students were  
126 registered, and a prevalence of TMDs of 31% was used as a reference point.  
127 It was determined that a minimum of 303 individuals per group were  
128 required to achieve a power of 0.95 and a type I error rate of 0.05 [2].

### 129 **Participants**

130 The present study was conducted with a sample of 18-25 years old  
131 university students from a variety of academic departments at Toros  
132 University. The evaluation form was administered to a sample of 909  
133 students between June 24, 2024, and July 22 of 2024. Participant  
134 recruitment, initial screening, and administration of sociodemographic and  
135 questionnaire assessments were conducted by two physiotherapists with 5  
136 years of experience (ÜY, EE). The presence of TMDs was evaluated by a  
137 maxillofacial surgeon with 26 years of expertise (BAK) in accordance with  
138 the Diagnostic Criteria for TMDs (DC/TMD) and the standards of the  
139 International Association for Dental, Oral and Craniofacial Research  
140 (IADR). Participants were then classified into TMD and control groups [11,  
141 12]. Participants who had undergone radiotherapy to the head, neck, or  
142 shoulder; had a history of surgery or orthodontic treatment related to the  
143 maxillofacial and neck area; had used analgesic and anti-inflammatory  
144 drugs or received any treatment related to TMDs in the previous year; or

145 had experienced generalized joint damage affecting the head, neck, or  
146 shoulder region or major trauma were excluded from the study.  
147 Additionally, the study excluded individuals with a history of acute or  
148 chronic inflammatory diseases, as well as those diagnosed with  
149 radiculopathy, myelopathy, systemic, metabolic, neurological, or immune  
150 diseases, post-traumatic stress disorder, or congenital conditions (Figure-  
151 1).

### 152 **Assessment of the Temporomandibular Joint Symptom Severity**

153 The severity of TMDs was evaluated using the Turkish version of the  
154 Fonseca Anamnestic Index (FAI). The FAI was reported to be sufficiently  
155 valid and reliable in the Turkish population (ICC: 0.739-0.897) by Kaynak  
156 et al. [13]. The FAI, which assesses the severity of TMDs according to signs  
157 and symptoms, is a simple questionnaire consisting of 10 specific  
158 questions. In this questionnaire, each question includes 3 answer options  
159 (Yes-Sometimes-No). In the questionnaire scoring, 'Yes' option is scored as  
160 10 points, 'Sometimes' option is scored as 5 points, and 'No' option is  
161 scored as 0 points. According to the total score of the questionnaire; 0-15  
162 points: No TMDs, 20-40 points: Mild TMDs, 45-65 points: Moderate TMDs,  
163 70-100 points: Refers to severe TMDs [13].

### 164 **Assessment of the Depression and Anxiety Symptoms**

165 Depression and anxiety symptoms were evaluated with the four-item short  
166 version of the Patient Health Questionnaire (PHQ-4). Demirci and Ekşi [14]  
167 conducted the study for Turkish adaptation of this scale and the Cronbach's  
168 alpha value was reported as 0.85. The four-item scale is described as the

169 shortest composite measurement tool to assess depression and anxiety  
170 disorders [15]. The questionnaire is a 4-point Likert-type scale with each  
171 item scored as (0) never, (1) few days, (2) more than half of the days, (3)  
172 almost every day [14]. The total score of the questionnaire ranges from 0  
173 to 12, higher scores present increased the prevalence of symptoms  
174 associated with depression and anxiety.

### 175 **Assessment of Nicotine Addiction**

176 Participants' cigarette/nicotine addiction levels were evaluated using the  
177 Turkish version of the Fagerström Test for Nicotine Dependence (FTND)  
178 [16]. The FTND comprises six items, each of which is assigned a distinct  
179 score. The scoring system ranges from 0 to 2 points for very low nicotine  
180 addiction, 3-4 points for low nicotine addiction, 5 points for moderate  
181 nicotine addiction, 6 to 7 points for high nicotine addiction, and 8 to 10  
182 points for very high nicotine addiction [16].

### 183 **Assessment of Alcohol Addiction**

184 Alcohol consumption and dependence were evaluated with the Addiction  
185 Profile Index Risk Scanning Scale (APIRSS). The Cronbach's Alpha  
186 coefficient of the questionnaire, whose validity and reliability study was  
187 conducted by Ögel et al. [17] was determined as 0.70. Each question of  
188 the APIRSS, which comprises six items, is scored between 0 and 2. The  
189 scale score ranges between 0 and 12, and scores 3 and above indicate an  
190 increased level of alcohol addiction [17].

### 191 **Assessment of Smartphone Addiction**

192 The level of smartphone addiction was evaluated using the Turkish version  
193 of the Smartphone Addiction Scale-Short Form. The validity and reliability  
194 study of the Turkish version of SPAI-SF in university students was  
195 conducted by Noyan et al. [18] and the Cronbach's alpha coefficient of the  
196 scale was reported as 0.867. The SPAI-SF, developed for evaluating the  
197 risk of smartphone addiction in adolescents and young adults, is a 6-point  
198 Likert-type scale with 10 items [18]. Participants were asked to choose one  
199 of 6 alternative answers to each question: 'strongly disagree', 'disagree',  
200 'somewhat disagree', 'somewhat agree', 'agree', and 'strongly agree'. Each  
201 item is assigned a rating on a scale of 1 to 6, with a total score ranging  
202 from 10 to 60 [18]. It is reported that the risk of addiction rises as the score  
203 increases [18].

#### 204 **Statistical Analysis**

205 The data of the study were analyzed with a statistical software program  
206 (SPSS software version 26, IBM Corporation, Armonk, NY, USA). The  
207 conformity of the variables to normal distribution was analyzed using visual  
208 (histogram and probability graphs) and analytical methods (Kolmogorov-  
209 Smirnov/Shapiro-Wilk tests). Since the variables were not normally  
210 distributed, the findings are presented as median (interquartile range). The  
211 differences of the parameters between the two groups were analyzed using  
212 the Chi-Square and the Mann-Whitney U tests. The correlation of the  
213 severity of TMDs with depression and smartphone, nicotine, and alcohol  
214 addiction participants with TMDs was evaluated using Spearman's  
215 correlation analysis. The results of the correlation analysis were classified

216 as follows: <0.20: "Poor", 0.21-0.40: "Fair", 0.41-0.60: "Moderate", 0.61-  
217 0.80: "Good", and >0.80: "Very Good" agreement. For the relevant  
218 statistical analysis methods, any p value less than 0.05 was accepted as  
219 statistically significant. Multiple logistic regression analysis was conducted  
220 to identify significant predictors of TMDs presence, dependency levels,  
221 TMDs severity, and psychological parameters. Predictors were selected  
222 using a backward stepwise method, with removal set at  $p > 0.10$  and  
223 reentry at  $p < 0.05$ . Model fit was evaluated using the Hosmer-Lemeshow  
224 goodness-of-fit test, where  $p < 0.05$  indicated a significant difference  
225 between observed and predicted values [19]. Odds ratios (ORs) for TMDs  
226 presence were calculated by simultaneously considering all parameters.  
227 Nagelkerke's  $R^2$  was used to estimate the proportion of variance in TMDs  
228 presence explained by the model, with values  $>0.75$  considered very good,  
229 0.50-0.75 good, 0.25-0.50 moderate, and  $\leq 0.25$  poor predictive power[20].  
230 The model's classification accuracy, including sensitivity and specificity for  
231 correctly identifying participants with and without TMDs, was determined  
232 using a  $2 \times 2$  classification table.

## 233 **Results**

234 A total of 909 volunteers, 641 with TMDs and 268 asymptomatic  
235 individuals, participated in the study. Of the participants, 610 (67.1%) were  
236 female and 299 (32.9%) were male. The overall prevalence of TMDs-related  
237 symptoms was 70.5%. There were no significant differences between  
238 groups in terms of age ( $p=0.446$ ), body weight ( $p=0.757$ ), height  
239 ( $p=0.287$ ), and body mass index ( $p=0.579$ ) parameters (Table 1).

240 The prevalence of nicotine ( $p=0.009$ ), alcohol ( $p=0.004$ ), and smartphone  
241 ( $p<0.001$ ) addiction was statistically significantly higher in individuals with  
242 TMDs in comparison to the control group (Table 2). The prevalence of  
243 nicotine addiction in individuals with TMDs was 32%, while it was 23.9%  
244 in the control group. Similarly, the prevalence of alcohol addiction was  
245 statistically significantly higher in individuals with TMDs (22.3%) in  
246 comparison to the control group (14.6%). Moreover, the prevalence of  
247 screen addiction was higher in individuals with TMDs (44.7%) than in  
248 asymptomatic individuals (26.1%) (Table 2).

249 The median (interquartile) values of the scales assessing TMDs symptoms,  
250 depression, and addiction severity are provided in Table 3. The PHQ-4  
251 ( $p<0.001$ ), SPAI-SF ( $p<0.001$ ), APIRSS ( $p<0.001$ ), and FTND ( $p=0.003$ )  
252 scores were significantly higher in individuals with TMDs compared to  
253 controls (Table 3).

254 In Table 4 only individuals with smoking, alcohol and smartphone addiction  
255 were analysed with reference to FTND, APIRS, SPAI-SF cut-off values.  
256 According to the results of these analyses the FTND total scores of the  
257 individuals with TMDs were found to be higher than those of the  
258 asymptomatic individuals ( $p=0.007$ ). However, APIRSS ( $p=0.757$ ) and  
259 SPAI-SF ( $p=0.119$ ), showed no statistically significant difference between  
260 the individuals with TMDs and the control group regarding the level of  
261 addiction (Table 4).

262 Results of the correlation analysis between TMDs severity and depression  
263 and addiction in participants with TMDs are given in Table 5. A moderate

264 positive correlation was found between the FAI and PHQ-4 scores  
265 ( $r=0.489$ ,  $p<0.001$ ). The FAI scores had poor to fair correlation with the  
266 SPAI-SF ( $r=0.283$ ,  $p<0.001$ ), FTND ( $r=0.147$ ,  $p<0.001$ ), and APIRSS  
267 ( $r=0.123$ ,  $p<0.001$ ) scores. The PHQ-4 exhibited poor to fair correlation  
268 with the SPAI-SF ( $r=0.328$ ,  $p<0.001$ ), APIRSS ( $r=0.125$ ,  $p<0.001$ ), and  
269 FTND ( $r=0.120$ ,  $p<0.001$ ) scores. A significant positive and fair correlation  
270 was observed between the APIRSS and the FTND ( $r=0.377$ ,  $p<0.001$ )  
271 (Table 5).

272 The variables included in the multiple logistic regression analysis were  
273 TMDs severity, symptoms of depression and anxiety, and levels of  
274 smartphone, alcohol, and nicotine dependence (Table 6). The results  
275 indicated that symptoms of depression and anxiety, as well as smartphone  
276 addiction, were significantly associated with the presence of TMDs  
277 diagnosed according to the DC/TMD criteria. The odds ratio for TMDs  
278 presence was 1.29 for depression and anxiety symptoms and 1.11 for  
279 smartphone addiction. Nicotine dependence ( $p = 0.515$ ) and alcohol  
280 dependence ( $p = 0.854$ ) were not statistically significant predictors.

281 The Hosmer-Lemeshow goodness-of-fit test was not statistically  
282 significant, indicating consistency between the observed data and the  
283 fitted model. The potential predictors explained 95.8% of the total log  
284 likelihood for TMDs presence, reflecting excellent predictive capacity  
285 (Nagelkerke's  $R^2 = 0.958$ ). The model demonstrated 99.6% sensitivity,  
286 98.8% specificity, and 99.4% overall accuracy in predicting TMDs  
287 presence.

## 288 **Discussion**

289 This study, which was conducted with a large sample of 909 participants,  
290 is the first study in Turkey to examine the phenomena of nicotine, alcohol,  
291 and smartphone addiction in individuals with TMDs and to compare these  
292 addictive behaviors in individuals with TMDs with those observed in  
293 asymptomatic controls. Another strength of the study is the use of valid  
294 and reliable scales in the evaluation of the parameters. The findings of the  
295 study show that individuals with TMDs have higher prevalence and degree  
296 of addiction to nicotine, alcohol, and smartphone. The results of the  
297 correlation analyses conducted in our study demonstrated a statistically  
298 significant positive correlation of TMDs severity with depression and  
299 smartphone, alcohol, and nicotine addiction. Furthermore, the results  
300 indicated a statistically significant positive fair correlation between  
301 depression severity and smartphone addiction, as well as alcohol and  
302 nicotine addiction, albeit of a weak magnitude. The prevalence of  
303 symptoms related to TMDs was found to be 70.5% in Turkish university  
304 students.

305 Nicotine and alcohol dependence levels of individuals with TMDs were  
306 found to be higher than those of asymptomatic controls. It is also observed  
307 that depression and anxiety levels of individuals with TMDs are higher than  
308 those of the control group. The results of our correlation analysis are  
309 similar with those of prior studies examining the depression and addiction  
310 components in individuals with TMDs confirming this interaction [7, 10]. It  
311 should be noted that the strength of the correlations is generally poor to

312 fair, except for the relationship between TMDs and depression.  
313 Consistently, the regression model identified PHQ-4 and SPAI-SF scores as  
314 significant independent predictors of TMDs presence. Each one-point  
315 increase in depressive/anxiety symptom scores increased the odds of TMDs  
316 by 1.29-fold, and each one-point increase in smartphone addiction scores  
317 increased the odds by 1.11-fold. As expected, higher levels of depression  
318 were associated with an increase in all addiction types examined in the  
319 present study. The enhancing effect of depression on addictive behavior  
320 has previously been explained by researchers based on the tendency to  
321 compensate for the effects of depression and the observation that  
322 individuals experience greater withdrawal when depressed [21]. Research  
323 highlights that up to 60% of adults with depression have smoked at some  
324 point in their lives and that the presence of higher depressive symptoms  
325 has been found to be associated with an elevated likelihood of engaging in  
326 smoking behavior [21]. Considering the multifaceted and biopsychosocial  
327 etiological origin of TMDs, the primary reason for nicotine and alcohol  
328 dependence in individuals with TMDs may be higher levels of depression  
329 and anxiety compared to the control group. Furthermore, one of the most  
330 commonly reported symptoms of TMDs is pain in more than one area and  
331 chronic pain syndrome is found in more than 50% of individuals with TMDs  
332 [22]. The prevalence of co-occurrence of chronic pain and depression was  
333 recorded as 59.1% [23]. Similarly, 35% of individuals with chronic pain  
334 have been shown to have increased levels of anxiety [24]. The frequent co-  
335 occurrence of chronic pain and depression symptoms is attributed to  
336 neuroplastic changes in the brain [25]. It has been suggested that

337 increased alcohol and nicotine addiction constitutes risk factors for sleep  
338 bruxism, which is related to dopaminergic system dysfunction [26].  
339 Research has shown that psychoactive substances such as alcohol and  
340 nicotine have an effect on the central nervous system by altering the  
341 functional properties of neurotransmitters and receptors [7]. In light of this  
342 information, another reason for the difference in nicotine and alcohol  
343 addiction levels between the two groups may be that TMDs trigger chronic  
344 pain, increasing depression and anxiety levels, and individuals resort to  
345 alcohol and cigarettes as a strategy to cope with chronic pain.

346 Although the association between psychological symptoms and TMDs is  
347 well established in the literature, recent perspectives on this relationship  
348 highlight the need to move beyond simple correlations. De la Torre Canales  
349 et al. proposed a biopsychosocial model in which emotional and behavioral  
350 factors are integral to both the onset and maintenance of TMDs [6]. More  
351 recently, Souza et al. emphasized that psychological factors should be  
352 regarded not merely as associated variables but as central drivers within  
353 multifactorial models [27]. In this context, the findings align with this  
354 theoretical framework, demonstrating that psychological symptoms are  
355 significant contributors in the multivariate logistic regression model  
356 predicting TMDs. Furthermore, Sangalli et al. highlight the importance of  
357 integrating brief behavioral interventions and psychological therapies  
358 (e.g., cognitive behavioral therapy, stress management) into treatment  
359 strategies for patients with TMDs, particularly when psychological  
360 comorbidities are present [28]. Therefore, the results support the growing  
361 body of evidence indicating that psychological assessment and

362 management should be regarded as essential components of both etiologic  
363 understanding and therapeutic planning for TMDs.

364 When individuals with nicotine, alcohol, and smartphone addiction were  
365 analyzed according to the FTND, APIRSS, and SPAI-SF cut-off scores, the  
366 addiction level of individuals with TMDs who were addicted to nicotine was  
367 higher than that of asymptomatic controls. When individuals with alcohol  
368 and smartphone addiction were analyzed, results show that the two groups  
369 have similar addiction levels. In contrast, neither alcohol addiction (APIRS)  
370 nor nicotine dependence (FTND) emerged as significant predictors in the  
371 regression model, with odds ratios of 0.96 and 0.91, respectively,  
372 indicating no meaningful association with TMDs presence after controlling  
373 for other variables. In order to determine the relationship between  
374 smoking and TMDs, Sanders et al.[29] dichotomized age as adults over 30  
375 and young adults under 30 and found that smoking was associated with the  
376 risk of TMDs in young adults. Miettinen et al.[7] reported that increased  
377 frequency in daily smoking and alcohol consumption is associated with  
378 TMDs symptoms in young adults. The results of our study are in parallel  
379 with those of two previous studies conducted in a similar age range and  
380 population. The observed higher nicotine and alcohol dependence among  
381 individuals with TMDs may be partly attributable to underlying anxiety,  
382 which has been identified as a driver of substance use. Recent research by  
383 Saracutu et al. has highlighted the mediating role of bruxism, particularly  
384 awake bracing, in the relationship between anxiety and TMDs-related  
385 symptoms [30, 31]. The main reason for the high level of addiction among  
386 nicotine addicts in the TMDs group, while the level of addiction among

387 alcohol and smartphone addicts was similar to the control group, may be  
388 due to the fact that cigarette consumption is a risk factor for parafunctional  
389 oral behavior, poor oral hygiene, and oral diseases [32]. Studies suggesting  
390 that non-smokers have better oral and dental health care than smokers,  
391 that university students are unaware of good oral hygiene practice  
392 procedures, and that inappropriate oral behaviors are associated with  
393 painful TMDs support this view [32].

394 The level of smartphone addiction was higher in individuals with TMDs  
395 compared to controls. The prevalence of mobile device usage, particularly  
396 among adolescents and young adults, has reached approximately 75% of  
397 the global population. This has led to the emergence of a new phenomenon,  
398 text neck syndrome, which has been described as “the pain of the 21st  
399 century” [33]. In this syndrome, prolonged use of mobile devices such as  
400 smartphones, tablets, and personal computers has been identified as a  
401 contributing factor to the development of overuse injuries in the cervical  
402 vertebrae [33]. The degree of neck flexion increases during the use of  
403 mobile devices. Prolonged and repetitive flexion of the neck leads to a  
404 forward posture of the head, which results in impaired postural control,  
405 imbalances in muscle strength, and compression of spinal structures [34].  
406 Moreover, it has been reported that the prevalence of jaw, neck, and  
407 shoulder pain increases with the duration and frequency of mobile device  
408 use [35]. The high prevalence of smartphone addiction observed in  
409 individuals with TMDs can be attributed to the detrimental effects of  
410 prolonged sitting posture and static loading on the cervical joints of the  
411 cranio-cervico-mandibular system.

412 The overall prevalence of symptoms related to TMDs in the 909 university  
413 students included in the study was 70.5% which aligns with a systematic  
414 review on the epidemiology of TMDs, and it is known that the prevalence  
415 of TMDs in the general population varies between 1% and 75% [1]. The  
416 results of this study (70.5%) are similar to those reported by Karthik et al.  
417 (80.6%)[36] and Kaynak et al. (62%) [37], both of which evaluated the  
418 prevalence of TMDs among university students. In our study, which  
419 included university students, 71.3% of the individuals with TMDs were  
420 female. These results are consistent with the literature, and it is reported  
421 that TMDs are significantly more prevalent among individuals within the  
422 20-40 age range [1, 38]. Additionally, many studies have observed that the  
423 prevalence of TMDs is 2-4 times higher in females compared to males [1,  
424 36, 37]. Although the elevated prevalence of TMDs in women is believed to  
425 be attributable to a complex interplay of biological, psychological, and  
426 social factors, including regular hormonal fluctuations and variations in  
427 connective tissue and muscle architecture, the observation that TMDs are  
428 susceptible to influence by reproductive hormones, with symptom  
429 exacerbation occurring during the reproductive years between the ages of  
430 20 and 40, may be the primary determinant of its higher prevalence in the  
431 female gender [36, 39].

432 These findings underscore the interplay between mood, addictive  
433 behaviors, and TMD symptomatology, reinforcing the value of a  
434 biopsychosocial approach to understanding and managing TMDs. Notably,  
435 the results of this study align with the fundamental principles outlined in  
436 the International Association for Dental, Oral and Craniofacial Research

437 (IADR) standard of care for TMDs management [11]. These standards  
438 emphasize patient-centered approaches, acknowledge the multifactorial  
439 biopsychosocial etiology of TMDs, and prioritize outcomes related to  
440 quality of life over mere symptom resolution.

441 There are some limitations to our study. Firstly, addiction rates may be  
442 different in different types of TMDs, such as myogenic, arthrogenic, and  
443 mixed type, which should be confirmed by clinical examination.  
444 Participants in our study were 18-25 years old university students, which  
445 may have the potential to affect the generalizability of the study results to  
446 different age groups. Furthermore, the individuals with TMDs in the  
447 present study were recruited from a university population rather than a  
448 treatment-seeking clinical population, thereby providing community-based  
449 data. This population may also have exhibited abnormal clinical or  
450 psychological outcomes due to academic pressure, as previously reported  
451 by Sangalli et al. [40]. Consequently, the prevalence rates reported in this  
452 study may differ from those observed in clinical, treatment-seeking  
453 populations. In this study, TMDs was defined as a clinical diagnosis based  
454 on DC/TMD criteria; however, emerging literature advocates for a re-  
455 conceptualization that acknowledges the heterogeneity of TMDs  
456 presentations, including the role of bruxism—particularly awake bracing—  
457 which has been described as the “medical gateway for dentistry” and  
458 proposed as a mediator between psychological factors, substance use, and  
459 TMD symptomatology. Notably, bruxism was not directly assessed in the  
460 present study [11, 41]. Given this, the absence of bruxism-related measures  
461 may have limited the ability to fully elucidate the mechanisms linking

462 anxiety, addictive behaviors, and TMDs in this sample. Future research  
463 should address these limitations by enrolling treatment-seeking clinical  
464 populations, directly assessing bruxism and other parafunctional activities,  
465 and incorporating potentially relevant sociodemographic variables such as  
466 income level to provide a more comprehensive understanding of the  
467 relationship between psychological factors, addictive behaviors, and  
468 TMDs.

## 469 **Conclusion**

470 In conclusion, the prevalence of nicotine, alcohol, and smartphone  
471 addiction are higher in individuals with TMDs in comparison to  
472 asymptomatic participants. Increased TMDs severity is associated with  
473 higher depression and higher levels of addiction to nicotine, alcohol, and  
474 smartphone. Depression levels have an influence on addiction behavior;  
475 however, this effect is limited. Clinicians should be aware of this  
476 relationship between TMDs severity and addiction when managing the  
477 treatment of individuals with TMDs.

## 478 **Declarations**

479 **Data availability:** The datasets produced or scrutinized in the present  
480 study can be obtained upon reasonable request from the corresponding  
481 author.

482 **Funding:** This study was conducted without receiving any financial  
483 support

484 **Conflicts of interest:** No declaration of conflicts of interest.

485 **Acknowledgments:** None.

486 **Ethics statement:** All the authors complied with the ethical policies  
487 outlined on the journal's author guidelines page.

488 **Ethics approval and consent to participate:** This study was approved  
489 by the Toros University Non-Interventional Clinical Research Ethics  
490 Committee with the decision number (2024/06-09). This study has been  
491 approved and meets all criteria for ethical standards defined in the 1964  
492 Declaration of Helsinki and all amendments made after. All participants  
493 provided written informed consent prior to participation.

494 **Consent to publish declaration:** Not applicable.

495 **Clinical Trial Registration declaration:** Not applicable.

496 **CRedit authorship contribution statement:** BAK: Methodology,  
497 Supervision, Project administration, Writing - Original Draft/ ÜY: Writing -  
498 Original Draft, Writing - Review & Editing, Investigation,  
499 Conceptualization/ EE: Writing - Original Draft, Writing - Review & Editing,  
500 Investigation, Visualization/ ST: Methodology, Supervision, Project  
501 administration, Conceptualization.

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681 **Figure Legend**682 **Figure 1.** Flowchart of the study683 **Tables**684 **Table 1.** Demographic characteristics of the participants.

<b>Parameter</b>		TMD group (n=641)	Control group (n=268)	<i>p</i>
Age (years)		21(20-22)	21(20-22)	0.446
Weight (kg)		64 (55-76)	64.50(55-80)	0.757
Height (m)		1.68 (1.62-175)	1.70 (1.63-1.78)	0.287
BMI (kg/m <sup>2</sup> )		22.59 (20.30-25.32)	22.39 (20.06-25.24)	0.579
Sex	Female	457 (%71.3)	153 (%57.1)	<b>&lt;0.001**</b> a
	Male	184 (%28.8)	115 (%42.9)	

686 Data was given as Median (Interquartile range) and percentiles. kg, kilogram; m, meter;  
 687 BMI, body mass index. a; \*\* p<0.001, Chi-Square Test.

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**Table 2.** Comparison of addiction rates in TMD and control groups.

<b>Parameters</b>		TMD group	Control group	p
		(n=641)	(n=268)	
Tobacco Addiction	Yes	205 (32.0%)	64 (23.9 %)	<b>0.009*</b>
	No	436 (68.0%)	204 (76.1%)	
Alcohol Addiction	Yes	143 (22.3%)	39 (14.6%)	<b>0.004*</b>
	No	498 (77.7%)	229 (85.4 %)	
Smartphone Addiction	Yes	286 (44.7%)	70 (26.1%)	<b>&lt;0.001*</b>
	No	354 (55.3%)	198 (73.9%)	

\*p<0.05, Chi-Square Test. Data was given as percentile.

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**Table 3.** Comparison of scale score between TMD and control groups.

<b>Parameters</b>	TMD group (n=641)	Control group (n=268)	<i>p</i>
FAI	40 (25-55)	10 (5-15)	<b>&lt;0.001*</b>
PHQ-4	5 (4-8)	3 (1-4)	<b>&lt;0.001*</b>
SPAI-SF	30 (23-38)	24 (19-33)	<b>&lt;0.001*</b>
APIRS	0 (0-2)	0 (0-1)	<b>&lt;0.001*</b>
FTND	0 (0-2,5)	0 (0-0)	<b>0.003*</b>

753 \*p<0.05, Mann-Whitney U Test. Data was given as median (interquartile range). FAI,  
754 Fonseca Anamnestic Index; FTND, Fagerström Test for Nicotine Dependence; APIRS,  
755 Addiction Profile Index Risk Scanning Scale; SPAI-SF, Smartphone Addiction Index-Short  
756 Form; PHQ-4, Patient Health Questionnaire

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**Table 4.** Comparison of addiction scale scores of addicted individuals according to the presence of TMD.

<b>Parameters</b>	TMD group	Control group	p
FTND (n=269)	4 (3-6)	3 (2-5)	<b>0.007*</b>
APIRS (n=182)	4 (4-6)	4 (3-6)	0.757
SPAI-SF (n=356)	39 (35-45)	38 (35-42)	0.119

790 \*p<0.05, Mann-Whitney U Test. Data was given as median (interquartile range). FTND,  
791 Fagerström Test for Nicotine Dependence; APIRS, Addiction Profile Index Risk Scanning  
792 Scale; SPAI-SF, Smartphone Addiction Index-Short Form.  
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**Table 5.** Correlation between depression, addiction and TMD severity

<b>Parameters</b>		PHQ-4	SPAI-SF	APIRS	FTND
FAI	Correlation Coefficient	<b>0.489*</b>	<b>0.283*</b>	<b>0.123*</b>	<b>0.147**</b>
	<i>P</i>	<0.001	<0.001	<0.001	<0.001
PHQ-4	Correlation Coefficient		<b>0.328*</b>	<b>0.125*</b>	<b>0.120**</b>
	<i>P</i>		<0.001	<0.001	<0.001
SPAI-SF	Correlation Coefficient			<b>0.084*</b>	0.047
	<i>P</i>			0.011	0.156
APIRS	Correlation Coefficient				<b>0.377**</b>
	<i>P</i>				<0.001

827 \* p<0.05; \*\*, p<0.001. FAI, Fonseca Anamnestic Index; FTND, Fagerström Test for  
828 Nicotine Dependence; APIRS, Addiction Profile Index Risk Scanning Scale; SPAI-SF,  
829 Smartphone Addiction Index-Short Form; PHQ-4, Patient Health Questionnaire-4.  
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843 **Table 6.** Multivariable logistic regression model

Parameters	B	<i>p</i>	Odds Ratio (95% C.I)	Sensitivit y	Specificit y	Accurac y	R <sup>2</sup>
TMD According to DC/TMD				99.6 %	98.8%	99.4 %	95.8 %
FAI	1.098	<.00 1	3.00 (2.28- 3.93)				
PHQ-4	.261	.021	1.29 (1.04- 1.61)				
SPAI-SF	.104	.003	1.11 (1.04- 1.18)				
APIRS	-.032	.854	.96 (0.69-1.35)				
FTND	-.094	.515	.91 (0.68-1.20)				

844 TMD: Temporomandibular Disorders; FAI, Fonseca Anamnestic Index; PHQ-4, Patient  
845 Health Questionnaire-4; SPAI-SF, Smartphone Addiction Index-Short Form; Addiction  
846 Profile Index Risk Scanning Scale; FTND, Fagerström Test for Nicotine Dependence;  
847 APIRS.

**EXCLUSION CRITERIA**

- Underwent radiotherapy to the head, neck, or shoulder
- History of surgery related to the maxillofacial or neck area
- History of orthodontic treatment related to the maxillofacial or neck area
- Use of analgesic drugs
- Use of anti-inflammatory drugs
- Received any treatment related to TMDs in the previous year
- Generalized joint damage affecting the head, neck, or shoulder
- Major trauma history
- Acute inflammatory diseases
- Chronic inflammatory diseases
- Radiculopathy
- Myelopathy
- Systemic diseases
- Metabolic diseases
- Neurological diseases
- Immune diseases
- Post-traumatic stress disorder
- Congenital conditions

Eligible Participants  
(n=909)

**ASSESSMENTS**

- Presence of TMD
- Smartphone Addiction
- Nicotine addiction
- Alcohol addiction
- TMD severity
- Depression and Anxiety