

1 **Title Page**

2

3 **Title:** Prevalence of computer vision syndrome and its risk factors in a Spanish university
4 population.

5 **Running short title:** Computer vision syndrome in a university.

6

7 **Authors:** Sara Ortiz-Toquero (PhD)¹, Irene Sanchez (PhD)¹, Alicia Serrano (OD)¹ and
8 Raul Martin (PhD)¹

9

10 **Affiliation:**

11 1. Universidad de Valladolid, Instituto Universitario de Oftalmobiología Aplicada
12 (IOBA), Paseo de Belén, 17 - Campus Miguel Delibes, Valladolid 47011, España;
13 Universidad de Valladolid, Departamento de Física Teórica, Atómica y Óptica,
14 Paseo de Belén, 7 - Campus Miguel Delibes, Valladolid 47011, España; Optometry
15 Research Group, IOBA Eye Institute, School of Optometry, University of Valladolid,
16 Valladolid, Spain.

17

18 **Note:** American Journal Experts has reviewed the English grammar of this manuscript
19 and may be verified on the AJE website using the verification code 045D-1145-345C-
20 79EF-6057.

21

22 **Tables:** 4

23 **Figures:** 4

24 **Date of manuscript submission:** 2nd March 2024

25 **Name and full mailing address for correspondence:** Sara Ortiz-Toquero. Paseo de
26 Belén, 7 - Campus Miguel Delibes, Valladolid 47011, Spain. E-mail:
27 sortizt@ioba.med.uva.es. Tel: +34 983 181851

28

29 **Conflicts of Interest and Source of Funding:** The authors declare no conflict of interest.
30 This study was supported by an investigator-initiated study grant from Alcon (IIT#
31 73376825).

32 **Abstract**

33 **Objectives:** To determine the prevalence of digital eye strain or computer vision
34 syndrome (CVS) and its risk factors in a university population (University of Valladolid,
35 Spain).

36 **Methods:** An anonymous cross-sectional online survey was conducted in a university
37 population [staff (lecturers and administrative employees) and students (undergraduate,
38 master's and PhD)], including two validated questionnaires (OSDI and CVSS17) and
39 questions about sociodemographic data and VDT use. The prevalence and risk factors for
40 CVS (CVSS17 \geq 29) (multivariate logistic regression model) were calculated.

41 **Results:** 1009 participants responded to the survey (35.2 \pm 15.2 years; 64.1% women). The
42 mean OSDI and CVSS17 questionnaire scores were 18.9 \pm 15.6 and 31.5 \pm 6.4,
43 respectively, and 35.4% of the respondents had dry eye symptoms (OSDI $>$ 22). The total
44 prevalence of CVS was 65.4% (95%CI 62.1-68.3%). Undergraduate students showed the
45 highest CVS prevalence (72.6%; $P<$ 0.01), which was significant. Additionally, women,
46 participants younger than 36 years old, CL wearers and subjects with dry eye symptoms
47 reported a statistically higher CVSS17 score ($P\leq$ 0.01). In the multivariate model,
48 significant factors associated with the presence of CVS ($P\leq$ 0.03) were female sex
49 (OR=2.10; 95%CI 1.54-2.88), dry eye symptoms (OSDI $>$ 22) (OR=16.98; 95%CI 10.36-
50 27.84), VTD use \geq 6 hours daily (OR=1.96; 95%CI 1.09-3.52) and being an undergraduate
51 student (OR=2.23; 95%CI 1.54-3.24).

52 **Conclusion:** A high prevalence (65.4%) of CVS was found among the Spanish university
53 population, with the undergraduate student group having the highest prevalence (72.6%).
54 Female gender, more than 6 hours/day of VDT use, being an undergraduate student and
55 dry eye symptoms significantly increased the risk of CVS in the university population.

56 **Key words:** Computer vision syndrome; university population; dry eye; prevalence.

57 In recent decades, new forms of digital displays have been developed and become
58 widespread throughout the world, making them indispensable in every personal,
59 professional, or academic activity. The use of visual display terminals (VDTs), such as
60 computers, laptops, smartphones, tablets, or e-readers, increases the visual demands on
61 the user and can lead to several ocular symptoms related to digital displays that can
62 adversely affect both quality of life¹ and productivity.²

63

64 This group of eye- and vision-related problems associated with prolonged use of VDTs
65 is described as digital eye strain or computer visual syndrome (CVS) by the American
66 Optometric Association.³ CVS involves asthenopic symptoms (eye strain, tired eyes or
67 headache), ocular surface symptoms (eye dryness, ocular redness, and itching or tearing,
68 among others), visual disturbances (double vision or blurred vision), and extraocular
69 discomfort (head, neck or back pain).³⁻⁵ It is estimated that CVS could affect 80% of
70 adults who use digital devices for at least 2 hours daily.³ This condition could have a
71 global prevalence of 60 million people worldwide with an incidence of one million new
72 cases each year.⁴ In eye care practice, a diagnosis of CVS mainly depends on subjective
73 patient answers to different validated questionnaires, one of which is the 17-item
74 Computer-Vision Symptom Scale questionnaire (CVSS17).^{6,7}

75

76 It is well known that VDT users exhibit decreased blinking frequency and amplitude,
77 leading to an increase in ocular surface exposure, tear evaporation and alteration of
78 meibomian gland secretion,^{8,9} which contribute to the development of dry eye
79 symptomatology,⁹ and close to 50% of VDT users suffer from dry eye disease (DED).¹⁰
80 It has also been reported that contact lens (CL) wear may increase or exacerbate the
81 presence of CVS symptoms.¹¹ In addition, a greater effort is required from the

82 accommodative system because of the greater number of working hours with the VDT
83 placed at a close distance and a higher angle of visualization, which can cause fatigue
84 symptoms, especially in cases of uncorrected refractive error or previous accommodation
85 anomalies.⁸

86

87 It is estimated that from 2000 to the present, the number of internet users has increased
88 by 1,392%, which means that 67.9% of the world's population is a user of some digital
89 device.¹² Given the worldwide rise in the use of technology, VDT use has created an
90 unprecedented revolution in learning or teaching strategies in higher educational
91 institutions, especially after the COVID-19 pandemic.¹³ For this reason, the detection and
92 prevention of CVS in this population to guarantee the visual health of the academic
93 population is crucial and challenging. Eye care practitioners should be aware of the
94 potential impact of VDT use on CVS and its related factors to respond with the most
95 effective preventive strategies.^{6,7}

96

97 Therefore, the aim of this study was to determine the prevalence of CVS and its risk
98 factors among students and staff at the University of Valladolid (Spain) to provide
99 evidence-based epidemiology data to help eye care providers improve visual health in
100 university population.

101

102 **Methods**

103 **Population and design of the study**

104 An anonymous cross-sectional online survey was emailed to the total population
105 (approximately 25700 people) of the University of Valladolid (Spain), including students
106 (undergraduate, master's, and PhD degrees) and staff (lecturers and administrative

107 employees), during the 2022/2023 academic year. The online questionnaire was designed
108 using Microsoft Forms and hosted on a secure network server of the University of
109 Valladolid to determine the symptomatology related to CVS. Before answering the
110 questionnaire, all participants gave their consent to be enrolled in the study. The study
111 was approved by the Human Sciences Ethics Committee of Valladolid Area-Este Clinic
112 Hospital (Castilla y Leon Public Health System-SACYL) and followed the tenets of the
113 Declaration of Helsinki and the standards of Good Clinical Practice. The online
114 questionnaire was sent on the mid-November 2022 and was left open for five weeks for
115 completion. Participants cannot modify their answers after sending questionnaire.

116

117 **Questionnaire design**

118 The online questionnaire was organized into three sections with all compulsory questions.
119 The first section collected sociodemographic data (sex, age, and connection with the
120 university), systemic and ocular health (systemic and/or ocular disorders,
121 pharmacological treatment and ocular surgery), current optical correction and their
122 preference when using VDTs (spectacles and/or CL wear), VDT exposure (daily hours of
123 use of digital devices both inside and outside the classroom or work office), use of
124 artificial tears and information on breaks taken when working with VDTs.

125

126 The second section included the validated CVSS17 questionnaire^{6,7} that was designed to
127 provide a patient-reported measure of CVS over the preceding month among VDT
128 users.^{6,7} This questionnaire is composed of 17 questions, which provide a total score
129 ranging from 17 to 53 points (the higher the score, the greater the patient's CVS
130 symptomatology). The CVSS17 provides information on approximately 15 different
131 symptoms, including a symptom's severity and frequency and the subject's opinion. CVS

132 can be classified into 5 levels according to the total score, with level 1 corresponding to
133 an absence of symptoms and level 5 to very severe discomfort related to the use of digital
134 screens.⁶ Participants with CVSS17 total scores ≥ 29 points (levels 3, 4 and 5) were
135 considered to have CVS symptoms.

136

137 The third section of the questionnaire included the OSDI questionnaire to determine the
138 symptomatology related to DED.¹⁴ It includes 12 items that evaluate the frequency of
139 symptoms over the preceding week. The OSDI questionnaire is structured into three main
140 domains: ocular symptoms (5 questions), vision-related daily function (4 questions) and
141 environmental triggers (3 questions). The OSDI score ranges between 0 and 100, where
142 higher scores represent a greater severity of symptoms, and is classified as follows: no
143 symptoms (score ≤ 12), mild (score 13-22), moderate (score 23-32), and severe symptoms
144 (score 33-100).¹⁵ Additionally, participants were grouped into two categories, non-DED
145 symptomatology (OSDI ≤ 22) and DED symptomatology (OSDI > 22), following
146 recommendations to provide comparable results with previous epidemiological DED
147 reports.¹⁶

148

149 **Statistical Analysis**

150 Statistical analyses were performed using SPSS for Windows software (version 27.0;
151 SPSS, Inc., Chicago, IL, USA) and Microsoft Office Excel (Microsoft Corp. Washington,
152 USA). The normal distribution of the variables was assessed with the Kolmogorov–
153 Smirnov test. Mean, standard deviation, and percentages were used to describe the data
154 when appropriate. Continuous variables were assessed with the Kruskal–Wallis and
155 Mann–Whitney U tests, and categorical variables were assessed with the chi-square test.
156 Descriptive analysis of CVS prevalence (according to CVSS17 levels 1 to 5 and CVSS17

157 score <29 and ≥ 29) was performed on the total sample and on groups based on the main
158 study variables of university group [students (undergraduate, master's, and PhD students)
159 and university staff (lectures and administrative employees)], sex (women and men), age
160 (analysed in both two [≤ 35 and >35 years old] age groups and five [18-25; 26-35; 36-45;
161 46-55 and >55 years] age groups), CL wear and dryness symptomatology (participants
162 with [OSDI >22] and without [OSDI ≤ 22] dryness symptomatology). CVSS17 levels and
163 CVS prevalence were reported as percentages with 95% confidence intervals (95% CIs)
164 calculated using bootstrapping through random repetition of 1,000 samples.

165

166 The odds ratios (ORs) along with Wald X^2 tests and 95% CIs were calculated with
167 multivariate logistic regression analysis to assess the relationships of university group
168 (undergraduate, master's, or PhD students; lecturers; and administrative employees), sex,
169 age (≤ 35 and >35 years old), CL wear, VDT use, and dryness symptomatology as
170 independent variables with CVS (CVSS17 ≥ 29 points). All statistical analyses were
171 considered significant at $P < 0.05$.

172

173 **Results**

174 **Descriptive data of the overall sample**

175 One thousand nine participants responded to the survey (response rate of 3.9%) and were
176 included in the study. Table 1 summarizes the sociodemographic characteristics of the
177 overall sample. The mean age was 35.2 ± 15.2 years (range 18 to 69 years). Of the total
178 participants, 64.1% ($n=647$) were women ($P < 0.001$). The average hours of daily VDT
179 use to study or work in an office were 5.8 ± 2.2 (ranging from 0 to 14). On the other hand,
180 the average number of hours of personal VDT use was 3.5 ± 2.0 (ranging from 0 to 12).

181 The mean OSDI score was 18.9 ± 15.6 (0 to 90.9), and the percentage of respondents with
182 dry eye symptoms (OSDI >22) was 35.4% (95% CI 32.5%-38.2%). Participants were
183 classified according to their role at the university as lecturers (n=271, 26.9%),
184 administrative employees (n=208, 20.6%), undergraduate students (n=405, 40.1%),
185 master's degree students (n=53, 5.3%) and PhD students (n=72, 7.1%).

186 **CVSS17 scores**

187 The mean CVSS17 score in the overall sample was 31.5 ± 6.4 (95% CI 31.1-31.9).
188 Lecturers had the lowest CVSS17 score, with statistically significant differences from
189 administrative staff and undergraduate students ($P < 0.01$; Figure 1). Additionally, women,
190 participants younger than 36 years old, CL wearers and subjects with dry eye symptoms
191 had statistically higher CVSS17 scores ($P < 0.03$; Figure 1).

192

193 **Overall CVS prevalence**

194 According to the CVSS17 score, 7.8% (95% CI 6.1%-9.5%) of the participants were
195 classified at level 1, 26.8% (95% CI 23.9%-29.5%) at level 2, 38.1% (95% CI 34.9%-
196 41.1%) at level 3, 22.5% (95% CI 19.9%-25.0%) at level 4 and 4.9% (95% CI 3.5%-
197 6.3%) at level 5 ($P < 0.001$). The total prevalence of CVS (CVSS17 score ≥ 29) was 65.4%
198 (95% CI 62.1%-68.3%), and 34.6% (95% CI 31.7%-37.9%) of the sample did not have
199 CVS (CVSS17 score < 29) ($P < 0.001$).

200

201 **CVS prevalence in university groups**

202 Statistically significant differences in the percentages of CVS level and symptomatic
203 CVS in each university group were found (Figure 2). A higher percentage of students
204 were classified as levels 3, 4 and 5 ($P = 0.01$) and had CVS (70.2%, 95% CI 66.0%-74.0%;
205 $P < 0.01$) compared with university staff. A detailed analysis among the different

206 university groups found a similar trend, with a significantly different proportion of CVS
207 levels ($P<0.01$) and symptomatic CVS ($P<0.01$) (Figure 3). Undergraduate students
208 showed the highest CVS prevalence (72.6%), followed by master's students (67.9%),
209 with lecturers having the lowest prevalence of CVS (54.6%). Table 2 summarizes the
210 results of the university groups.

211

212 **CVS prevalence by sex**

213 Women showed a higher proportion of CVS levels 3, 4 and 5 ($P<0.01$; Figure 2) and a
214 higher CVS prevalence ($P<0.01$; 73.7%, 95% CI 70.2%-77.0%) compared with men
215 (50.6%, 95% CI 45.3%-55.5%).

216

217 **CVS prevalence by age**

218 A similar trend was found according to age groups, where participants younger than 35
219 years old showed a higher proportion of CVS levels 3, 4 and 5 ($P<0.01$; Figure 2) and a
220 higher percentage of CVS (69.7%, 95% CI 65.8%-73.4%; $P<0.01$). A detailed analysis
221 of the age groups found that all groups showed a greater than 50% prevalence of
222 symptomatic CVS ($P<0.01$, Table 3). The percentage of CVS decreased with increasing
223 age ($P<0.01$), from a prevalence of 72.2% in the youngest group (18-25 years) to 59.9%
224 in the group older than 55 years (Table 3).

225

226 **CVS prevalence by CL wear**

227 CL wearers showed a higher percentage of CVS levels 3 and 5 ($P=0.04$; Figure 2) and a
228 significantly higher CVS prevalence (73.2%, 95% CI 66.8%-79.0%; $P=0.01$) than non-
229 CL wearers (63.4%, 95% CI 60.1%-66.8%).

230

231 **CVS prevalence by dry eye symptomatology**

232 Participants with dryness symptomatology (OSDI score >22) showed a statistically higher
233 percentage of CVS levels 3, 4 and 5 ($P<0.01$; Figure 2). Additionally, participants with
234 dryness symptomatology showed a statistically significant difference in CVS prevalence
235 ($P<0.01$). In this analysis, a substantial percentage of participants without CVS symptoms
236 did not have dryness symptomatology (94.5%, 95% CI 92.0%-96.6%), but a slightly
237 higher percentage of participants with CVS (51.2%, 95% CI 47.2%-55.3%) showed
238 dryness symptomatology (Figure 2).

239

240 **CVS prevalence by other study variables**

241 Finally, no statistically significant differences ($P\geq 0.35$; Table 4) between the CVS and
242 non-CVS groups were found for systemic diseases, systemic medication, medication
243 affecting the ocular surface, ocular surgeries or the time breaks taken by participants when
244 using VDTs. In turn, the proportion of respondents using VDTs for ≥ 6 hours daily and
245 the use of artificial tears were significantly higher in the group with significant CVS
246 symptoms ($P\leq 0.03$).

247

248 **Risk factors for CVS**

249 According to the multivariate adjusted model, significant factors associated with the
250 presence of CVS ($P\leq 0.03$) were female sex (OR=2.10), dry eye symptoms (OSDI >22)
251 (OR=16.98), VTD use ≥ 6 hours daily (OR=1.96) and being an undergraduate student
252 (OR=2.23) (Figure 4). Although there was a trend of a higher prevalence of CVS in CL
253 users and subjects ≤ 35 years of age, there was a nonsignificant association between CL
254 wear (OR=1.18, 95% CI 0.80-1.77; $P=0.42$) and age group (OR=0.93, 95% CI 0.56-1.54;
255 $P=0.78$) and the risk of CVS in the multivariate model.

256 **Discussion**

257 This is the first study conducted considering different university population groups
258 (lecturers, administrative employees, and undergraduate, master's, and PhD students) to
259 determine the prevalence and associated factors of CVS in a large sample of 1009 subjects
260 of a highly educated population.

261

262 A high prevalence of CVS of 65.4% was found in the university population assessed, with
263 undergraduate students having the highest prevalence (72.6%) and lecturers having the
264 lowest prevalence (54.6%; $P < 0.01$). These results are consistent with other reports that
265 found a similar CVS prevalence in Spanish university students [between 73.8%¹⁷
266 (undergraduate and postgraduate students) and 76.6%¹⁸ (undergraduate and master's
267 degrees) with the CVS-Q questionnaire] and in other countries and degrees. For example,
268 a CVS prevalence between 69.1%¹⁹ (in medical students; CVS-Q questionnaire) and
269 77.1%¹⁹ (in undergraduate students; CVS-Q questionnaire) was described in the USA,
270 80%²⁰ (in medicine and engineering students; nonvalidated questionnaire) in India, and
271 82.5%²¹ (in medical students who wear spectacles; CVS-Q questionnaire) in Paraguay,
272 and the highest prevalence reported was 90%²² (undergraduate students from 5
273 universities; nonvalidated questionnaire) in Malaysian students.

274

275 The prevalence of CVS in the university student population was significantly higher than
276 the prevalence in lecturers (54.6%) and administrative employees (67.3%). However, the
277 CVS prevalence in lecturers and administrative staff is similar to the previously reported
278 prevalence in other professional populations, such as Italian office workers (67.2%; CVS-
279 Q questionnaire)²³ and Spanish health care workers (56.8%; CVS-Q questionnaire).²⁴
280 Therefore, these results suggest that undergraduate students have a greater risk of

281 developing CVS, as the multivariate logistic model confirmed that students have more
282 than double the probability of suffering from CVS (OR=2.19; 95% CI 1.51-3.17; p<0.01)
283 than the rest of the university population assessed (Figure 4). However, although the
284 prevalence of CVS varies depending on the population assessed and the questionnaire
285 used, all previous reports described an elevated prevalence of CVS in undergraduate
286 university students, which could develop into a major public health problem in
287 developing countries, contributing to reduced academic performance and work
288 productivity with a negative impact on the quality of life of VDT users.^{1,25}

289

290 According to the multivariate logistic model, being an undergraduate student is not the
291 only risk factor for developing CVS in the assessed Spanish university population.
292 Women (OR=2.10; 95% CI 1.54-2.88; p<0.01) also had more than double the probability
293 of suffering from CVS than men (Figure 4). This higher risk for women has been
294 previously described,^{17,23,26-28} and other authors have found similar (OR=1.78; 95% CI
295 1.35-2.34 in female professionally active computer users²⁸) or higher (OR=2.95; 95% CI
296 2.14-4.08¹⁷ in female university students and OR=3.42; 95% CI 1.94-6.04²³ in female
297 office workers) female-associated risks of CVS. Some authors suggest that this higher
298 risk is related to the higher prevalence of DED in women,¹⁶ and the relationship between
299 female sex and DED is well known.¹⁶ Therefore, special attention should be given to
300 female VDT users by eye care practitioners. Nevertheless, because most of the
301 participants (64.1%) in this survey were women [with a higher percentage in all studied
302 groups (Table 2) and age groups (Table 3)], further research to assess the reasons for the
303 higher risk of CVS in females should be conducted.

304

305 Moderate or severe dry eye symptoms (OSDI >22) represent the highest risk of CVS
306 (OR=16.98; 95% CI 10.36-27.84; p<0.01; Figure 4). The percentage of participants with
307 dryness symptomatology (OSDI score >22) was significantly higher (p<0.01) in the CVS
308 group (51.2%) than in the non-CVS group (5.4%). These results are consistent with recent
309 literature reports that have found a significant relationship between dry eye symptoms
310 and VDT use.^{10,29} Additionally, undergraduate students with CVS had higher OSDI
311 scores, and the higher the OSDI value is, the greater the risk of CVS (OR=1.20; 95% CI
312 1.17-1.24; p<0.01).¹⁷ According to the OSDI score, 35.4% of the university population
313 assessed presented moderate or severe dry eye symptoms. It is also well known that the
314 prevalence of DED increases in people over 50 years of age,¹⁶ and therefore, older
315 participants could be expected to exhibit CVS. However, the results showed a slightly
316 higher CVS prevalence in younger VDT users (Table 3), with no statistically significant
317 risk factor (OR=0.93, 95% CI 0.56-1.54; P=0.78), consistent with a previous report¹⁸ that
318 found lower CVS among older undergraduate students (22-29 years) than among younger
319 undergraduate students (18-19 years).

320

321 A higher prevalence of DED symptomatology was found compared with that of a recent
322 epidemiology report in a large population (n=1,077) in Spain that found dryness
323 symptoms (OSDI >22) in 15.5% (95% CI 13.2%-17.6%) of the population, suggesting a
324 higher risk of DED symptomatology in the university population.³⁰ Other reports (in
325 office workers in New York) also found a higher prevalence (29.9%) of dryness
326 symptomatology.¹² However, in a 2016 meta-analysis, the global prevalence of DED in
327 office workers was estimated to be 49.5% (ranging from 9.5% to 87.5%), although the
328 authors highlighted the necessity of implementing common DED diagnostic criteria in
329 research to allow for a more relevant estimation of DED prevalence.¹⁰

330

331 This high prevalence of dry eye symptoms may be largely because the use of VDTs
332 increases the vicious cycle of dry eyes.^{16,31} The use of VDTs decreases blink rates and
333 increases incomplete blinks, resulting in tear film instability, increased tear evaporation
334 and hyperosmolarity.³¹ Given that VDT use is a consistent risk factor for DED^{10,29}, eye
335 care practitioners should pay special attention to VDT users with preexisting symptoms
336 of dryness, as they are at an increased risk of suffering from CVS. According to the
337 current literature, management strategies for digital display-induced dry eye to improve
338 tear film stability and decrease tear evaporation, such as blink animation programmes,
339 oral intake of omega-3 fatty acids, the 20-20-20 rule, instillation of high-viscosity
340 artificial tears or adjustable chairs with ergonomic training, are usually recommended.^{31,32}

341

342 Moreover, study findings that spending more than 6 hours/day with digital devices
343 doubles the probability of suffering from CVS are common.^{5,23} The results of this study
344 are consistent with those of other studies; a CVS OR of 1.96 (95% CI 1.09-3.52; P=0.03)
345 was found in participants who used VDTs more than 6 hours daily. Additionally, the CVS
346 group spent significantly more time in front of digital screens for study/work-related tasks
347 and personal use than the non-CVS group (P<0.01; Table 4), and although in both groups
348 most of the participants (over 90%) spent more than 6 hours/day in front of VDTs, this
349 percentage was significantly higher in the CVS group (P=0.03; Table 4).

350

351 Additionally, it is extensively recognized that CL wear is one of the leading risk factors
352 for DED,¹⁶ with reports suggesting a DED prevalence up to four times higher in CL
353 wearers.¹⁷ Considering that dryness symptomatology is one of the main causes of CVS,
354 CL wear may also be related to CVS. However, although a significantly higher number

355 of CL wearers (22.7%) was found in the CVS group than in the non-CVS group (15.8%),
356 CL wear was not a significant risk factor for CVS (OR=1.18; 95% CI 0.80-1.77; P=0.42).
357 Our results are in line with those published by Meyer et al.³³ who found that CL wearers
358 do not experience symptoms of digital eye strain at higher frequency or severity than non-
359 CL wearers. These authors emphasize that the digital strain symptoms may not be directly
360 linked to the surface sensation of the CL itself but may result from several simultaneous
361 factors, such as, blink pattern, non-ergonomic environment, taking breaks or binocular
362 vision factors.³³ In contrast, other studies have reported that CL wear is a risk factor for
363 CVS (OR=1.97; 95% CI 1.16-2.20; P=0.01)¹⁷ in a sample of 851 university students and
364 that regular CL wear (OR=4.85; 95% CI 1.25-18.8; P=0.02) could also increase CVS after
365 6 h of computer work.¹¹

366

367 The main limitation of this study is the cross-sectional study design, which limits the
368 inference of causality of the results but allow to demonstrate the association between CVS
369 and the identified risk factors due to the large number of participants. Moreover, in the
370 current study, no eye care clinical examinations were conducted, and all the variables
371 analysed were self-reported, so future studies including an eye exam could be necessary
372 to improve the description of the relationship between CVS and refractive or ocular
373 surface patient characteristics. Additionally, the survey did not include questions about
374 stress, hours of sleep, neck, and shoulder pain, ergonomic or environmental conditions
375 (indoor air quality, lighting conditions, etc.) during VDT use, which could lead to
376 measurement bias. In future studies, it would be interesting to compare the prevalence of
377 CVS in university students and staff between different countries using the same
378 methodological procedure and validated questionnaires to improve knowledge about
379 CVS. These findings could inform preventive strategies and provide information on the

380 implications of CVS in university populations, especially among younger subjects.
381 Finally, eye care practitioners should be aware of the relevance of triaging questionnaires
382 and CVS risk factors when exploring university populations and/or patients who use
383 VDTs for long periods for study, work, or personal tasks.

384

385 **Conclusions**

386 This study investigated the prevalence of CVS and its associated factors in a large Spanish
387 university population. CVS was found to have a prevalence of 65.4%, with undergraduate
388 students having the highest prevalence (7 out of 10 students). Female gender, more than 6
389 hours/day of VDT use, being an undergraduate student and presenting dry eye symptoms
390 significantly increased the risk of CVS. These results highlight the importance of
391 preventing CVS in university populations and the paramount role of eye care practitioners
392 in reducing eye complications related to CVS in patients who use VDTs for long periods
393 and present CVS risk factors.

394

395 **References**

396

397 1. Hayes JR, Sheedy JE, Stelmack JA, et al. Computer use, symptoms, and quality of
398 life. *Optom Vis Sci.* 2007;84:738-744.

399 2. Daum KM, Clore KA, Simms SS, et al. Productivity associated with visual status
400 of computer users. *Optometry.* 2004;75:33-47.

401 3. American Optometric Association, Computer Vision Syndrome. Available at:
402 [https://www.aoa.org/patients-and-public/caring-for-your-vision/protecting-your-](https://www.aoa.org/patients-and-public/caring-for-your-vision/protecting-your-vision/computer-vision-syndrome)
403 [vision/computer-vision-syndrome.](https://www.aoa.org/patients-and-public/caring-for-your-vision/protecting-your-vision/computer-vision-syndrome) Accessed June 29, 2023.

404 4. Singh S, McGuinness MB, Anderson AJ, et al. Interventions for the management
405 of computer vision syndrome: a systematic review and meta-analysis.
406 *Ophthalmology.* 2022;129:1192-1215.

407 5. Talens-Estarellles C, García-Marqués JV, Cervino A, et al. Use of digital displays
408 and ocular surface alterations: A review. *Ocular Surface.* 2021;19:252-265.

409 6. González-Pérez M, Susi R, Barrio A, et al. Five levels of performance and two
410 subscales identified in the computer-vision symptom scale (CVSS17) by Rasch,
411 factor, and discriminant analysis. *PLoS One.* 2018;13:e0202173.

412 7. González-Pérez M, Susi R, Antona B, et al. The Computer-Vision Symptom Scale
413 (CVSS17): development and initial validation. *Invest Ophthalmol Vis Sci.*
414 2014;55:4504-4511.

415 8. Rosenfield M. Computer vision syndrome: a review of ocular causes and potential
416 treatments. *Ophthalmic Physiol Opt.* 2011;31:502-515.

417 9. Uchino M, Yokoi N, Uchino Y, et al. Prevalence of dry eye disease and its risk
418 factors in visual display terminal users: The Osaka study. *Am J Ophthalmol.*
419 2013;156:759-66.

- 420 10. Courtin R, Pereira B, Naughton G, et al. Prevalence of dry eye disease in visual
421 display terminal workers: A systematic review and meta-analysis. *BMJ Open*.
422 2016;6:e009675.
- 423 11. Tauste A, Ronda E, Molina MJ, et al. Effect of contact lens use on Computer
424 Vision Syndrome. *Ophthalmic Physiol Opt*. 2016;36:112-119.
- 425 12. World Internet users statistics and 2019 World population stats. Available at:
426 <https://www.internetworldstats.com/stats.htm>. Accessed February 20, 2024.
- 427 13. Talens-Estarellles C, García-Marqués JV, Cervino A, et al. Online Vs In-person
428 education: evaluating the potential influence of teaching modality on dry eye
429 symptoms and risk factors during the COVID-19 pandemic. *Eye Contact Lens*.
430 2021;47:565-572.
- 431 14. Schiffman RM, Christianson MD, Jacobsen G, et al. Reliability and validity of the
432 Ocular Surface Disease Index. *Arch Ophthalmol*. 2000;118:615-621.
- 433 15. Miller KL, Walt JG, Mink DR, et al. Minimal clinically important difference for
434 the ocular surface disease index. *Arch Ophthalmol*. 2010;128:94-101.
- 435 16. Stapleton F, Alves M, Bunya VY, et al. TFOS DEWS II Epidemiology Report.
436 *Ocul Surf*. 2017;15:334-365.
- 437 17. Talens-Estarellles C, García-Marqués JV, Cerviño A, et al. Dry eye-related risk
438 factors for digital eye strain. *Eye Contact Lens*. 2022;48:410-415.
- 439 18. Cantó-Sancho N, Sánchez-Brau M, Ivorra-Soler B, et al. Computer vision
440 syndrome prevalence according to individual and video display terminal exposure
441 characteristics in Spanish university students. *Int J Clin Pract*. 2021;75: e13681.
- 442 19. Wang C, Joltikov KA, Kravets S, et al. Computer vision syndrome in
443 undergraduate and medical students during the COVID-19 pandemic. *Clin*
444 *Ophthalmol*. 2023;17:1087-1096.

- 445 20. Logaraj M, Madhupriya V, Hegde S. Computer vision syndrome and associated
446 factors among medical and engineering students in Chennai. *Ann Med Health Sci*
447 *Res.* 2014;4:179-185.
- 448 21. Coronel-Ocampos J, Gómez J, Gómez A, et al. Computer visual syndrome in
449 medical students from a private university in Paraguay: a survey study. *Front*
450 *Public Health.* 2022;10:935405.
- 451 22. Reddy SC, Low CK, Lim YP, et al. Computer vision syndrome: a study of
452 knowledge and practices in university students. *Nepal J Ophthalmol.* 2013;5:161-
453 168.
- 454 23. Cantó-Sancho N, Porru S, Casati S, et al. Prevalence and risk factors of computer
455 vision syndrome—assessed in office workers by a validated questionnaire. *PeerJ.*
456 2023;11:e14937.
- 457 24. Artime-Ríos E, Suárez-Sánchez A, Sánchez-Lasheras F, et al. Computer vision
458 syndrome in healthcare workers using video display terminals: an exploration of
459 the risk factors. *J Adv Nurs.* 2022;78:2095-2110.
- 460 25. Anbesu EW, Lema AK. Prevalence of computer vision syndrome: a systematic
461 review and meta-analysis. *Sci Rep.* 2023;13:1801.
- 462 26. Portello JK, Rosenfield M, Bababekova Y, et al. Computer-related visual
463 symptoms in office workers. *Ophthalmic Physiol Opt.* 2012;32:375-382.
- 464 27. Ranasinghe P, Wathurapatha WS, Perera YS, et al. Computer vision syndrome
465 among computer office workers in a developing country: An evaluation of
466 prevalence and risk factors. *BMC Res Notes.* 2016;9:150.
- 467 28. Toomingas A, Hagberg M, Heiden M, et al. Risk factors, incidence and persistence
468 of symptoms from the eyes among professional computer users. *Work.*
469 2014;47:291-301.

- 470 29. Sánchez-Valerio MDR, Mohamed-Noriega K, Zamora-Ginez I, et al. Dry eye
471 disease association with computer exposure time among subjects with computer
472 vision syndrome. *Clin Ophthalmol.* 2020;14:4311-4317.
- 473 30. Martin R. Symptoms of dry eye related to the relative humidity of living places.
474 *Cont Lens Anterior Eye.* 2023;46:101865.
- 475 31. Kamøy B, Magno M, Nøland ST, et al. Video display terminal use and dry eye:
476 preventive measures and future perspectives. *Acta Ophthalmol.* 2022;100:723-
477 739.
- 478 32. Talens-Estarellles C, García-Marqués JV, Cerviño A, et al. Ocular surface
479 predisposing factors for digital display-induced dry eye. *Clin Exp Optom.*
480 2023;106:373-379.
- 481 33. Meyer D, Rickert M, Kollbaum P. Ocular symptoms associated with digital device
482 use in contact lens and non-contact lens groups. *Cont Lens Anterior Eye.*
483 2021;44:42-50.
- 484
- 485

486 **Figure Legends**

487

488 **Figure 1.** Summary of differences in CVSS17 scores between university role, sex, age,
489 CL wear and DED symptoms groups. Mean, maximum, minimum and Mann–Whitney U
490 test p values are presented for each variable. Und-St=undergraduate students; MSc-
491 St=master students; PhD-St=Doctorate students; CL=contact lens; y=years.

492

493 **Figure 2.** Summary of CVSS17 levels and computer vision syndrome (CVS) prevalence
494 by each group analysed. The 95% confidence interval (CI) bars are represented. The χ^2 P
495 values for each CVS level (1 to 5) and for groups with or without CVS are shown.

496

497 **Figure 3.** Summary of CVSS17 levels and computer vision syndrome (CVS) prevalence
498 by university population groups. The 95% confidence interval (CI) bars are represented.
499 The χ^2 P values for each CVS level (1 to 5) and for groups with or without CVS are
500 shown.

501

502 **Figure 4.** Association of computer vision syndrome (CVS) with main risk factors. The
503 results of the multivariate logistic regression model for CVS versus non-CVS are shown
504 for the independent variables of sex, dry eye symptoms, VTD use ≥ 6 hours daily and
505 university groups. CI =confidence interval; OR =odds ratio.