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Title page

**The accuracy and suitability of eating disorder screening tools for Binge
Eating Disorder and Bulimia Nervosa in a primary care setting: a systematic
review and narrative summary**

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Abstract

Background

BED & BN screening tools review

27 Despite available screening tools for eating disorders (EDs), the accuracy and
28 suitability of these in identifying Binge eating disorder (BED) and Bulimia nervosa (BN)
29 in a primary care setting are undetermined, despite BED/BN being the most common
30 EDs.

31 **Aim**

32 To evaluate the accuracy and suitability of ED screening tools for BED/BN in a primary
33 care setting.

34 **Design and setting**

35 A systematic review with narrative synthesis in a primary care setting.

36 **Method**

37 Six databases were searched, including MEDLINE, PsycINFO, and Embase. Two
38 independent reviewers screened studies for inclusion. Studies were included that
39 assessed the accuracy and/or suitability of screening tools for BED/BN in primary care.
40 Quality was assessed using the Mixed Methods Appraisal Tool. A narrative summary
41 was created after integrating the data using a convergent segregated approach.

42 **Results**

43 Four studies met inclusion criteria. The included studies reported on BEDS-7, EDE-Q
44 and SCOFF screening tools. No studies reported on the accuracy of screening tools
45 for BED and suitability for BN. BEDS-7 and EDE-Q screening tools reported variations
46 in their suitability in primary care. The main barrier to implementation in primary care
47 was time constraints and a lack of trust in screening. SCOFF showed high sensitivity
48 (97.88%–100%) for BN but had lower specificity (89.6%–94.4%), increasing false
49 positives.

50 **Conclusion**

51 ED screening tools face feasibility and accuracy concerns for BED/BN in a primary
52 care setting. Further research is needed to validate screening tools' accuracy and
53 suitability in a primary care setting for BED and BN in the general population.

54 **Keywords**

55 Binge eating disorder, bulimia nervosa, primary health care, general practice,
56 screening, systematic review, suitability, accuracy, eating disorders, identification

57 **How this fits in**

58 This research is highly relevant to general practice, where early identification is crucial
59 for early intervention. Given that there is an increase in prevalence of BED/BN, with
60 primary care providers often serving as the first point of contact or presentation for
61 individuals with BED/BN, understanding the accuracy and suitability of screening tools
62 for these eating disorders within the primary care setting is essential.

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Introduction

72 Eating disorders (EDs) are serious mental health conditions characterised by
73 disturbed eating patterns, a focus on body image, and a preoccupation with food,
74 weight, or shape (1). The most common EDs are binge-type EDs, such as Bulimia

75 Nervosa (BN) and Binge Eating Disorder (BED), affecting approximately 4.2% of the
76 global population (2,3) with suggestions of a much higher real prevalence (3). Both
77 BED and BN are characterised by recurrent binge episodes, during which individuals
78 consume a large amount of food in a short period of time without the ability to control
79 the behaviour (1). BN is further characterised by compensatory behaviours such as
80 self-induced vomiting, excessive exercise, or laxative use to counteract the potential
81 effects of the binge episode, such as weight gain (1). While BN and Anorexia Nervosa
82 (AN) share some common characteristics, such as a focus on weight, when looking at
83 eating disorder symptoms dimensionally instead of categorically, BED and BN are
84 suggested to share clinical presentations more than AN and BN (4).

85

86 Delayed identification of BED/BN could lead to reduced quality of life (5), with an
87 increased risk of mortality and comorbidities such as obesity or depression (6–9), sub-
88 optimal use of the healthcare system, and presents an economic burden (10–12).
89 Hence, early identification of BED/BN is crucial. However, poorly performing screening
90 tools can lead to overdiagnosis and overmedicalisation (13). Despite primary care
91 being the hub for early identification (14), detection rates of BED/BN in primary care
92 remain low (15).

93

94 Several screening tools have been developed to identify eating disorders (EDs),
95 including BED and BN (16). The most commonly used screening tools, both in
96 research and practice, are the EDE-Q (17) and the SCOFF (18). However, there is an
97 inconsistency in the validity of screening tools. One systematic review reported that
98 SCOFF have high sensitivity in young females with anorexia nervosa (AN) when
99 compared to clinical interviews (19). When compared to EDE-Q as a benchmark,

BED & BN screening tools review

100 SCOFF had lower sensitivity and specificity than Eating Disorders Screen for Primary
101 Care and Screen for Disordered Eating (20). However, the convergence between
102 EDE-Q and clinical interviews was low to moderate, as it both overestimated and
103 underestimated AN and BN (21,22). This questions the efficacy of EDE-Q being used
104 as a comparator in earlier studies.

105

106 While there is some evidence for good sensitivity of these screening tools for detecting
107 EDs (23–25), mainly AN, more evidence is needed to understand screening accuracy
108 in other EDs, such as BED and BN, and the potential suitability of screening tools for
109 key stakeholders within the primary care setting. Feltner et al. conducted a systematic
110 review which aimed to address the general accuracy of ED screening tools in a
111 comprehensive review on primary care and the US (26). However, measures of
112 suitability and accuracy of these tools within primary care for BED and BN were not
113 reported on. In addition, studies were limited to the US. Hence, building on Feltner et
114 al.'s work and the gap identified by the authors, this systematic review aims to evaluate
115 the accuracy and suitability of ED screening tools for BED/BN in a primary care setting.

116

117

Methods

118 Protocol

119 The protocol for this review was prospectively registered with the PROSPERO
120 database of systematic reviews (Registration Number: CRD42024595253). This
121 systematic review was conducted in accordance with the Preferred Reporting Items
122 for Systematic Reviews and Meta-Analyses (PRISMA) statement (27).

123

124 Data sources and search strategy

BED & BN screening tools review

125 The search was conducted in the following databases in August 2024: CINAHL
126 Complete (EBSCO), Cochrane Central Register of Controlled Trials (CENTRAL),
127 Embase, MEDLINE, PsycINFO, Global Health (OVID). Keyword search included
128 binge*", "bulimi*", "general practi*", "primary healthcare", "screening*", and
129 "questionnaire*" (see full search strategy in Supporting Information Appendix A). Due
130 to the difference in MESH term availability across databases, some adaptations were
131 made to ensure standardisation. Backwards and forward citation was implemented.
132 Grey literature was considered for search; however, it was not included due to the
133 limited time and resources of this study, and was judged to be unlikely to meet the
134 quality required for inclusion in this review.

135

136 **Selection criteria**

137 Original qualitative, quantitative and mixed-method research that reported on the
138 accuracy and/or the suitability of EDs screening tools for BED and BN in a primary
139 care setting in peer-reviewed sources was included in this review (See Table 1. for
140 definitions). No restriction was applied for language, year of publication, country or ED
141 screening tool. The primary care setting included professions such as general
142 practitioners (GPs), dentists, community pharmacists, optometrists, nurse
143 practitioners and psychological wellbeing practitioners (PWPs) and patients of all ages
144 with BED or BN. Studies with a true screening population sample were included.
145 However, studies with low prevalence testing in primary care were also included to
146 allow for a more generic understanding of the current situation. Studies using a
147 diagnostic population were excluded. Studies focusing on secondary and tertiary care
148 settings were excluded. Studies where non-primary care HCPs were administering the
149 screening tools were excluded. Studies reporting exclusively on Anorexia Nervosa

150 (AN), Other specified feeding or eating disorders or Avoidant/restrictive food intake
151 disorder were excluded.

152 **Table 1.** Definition of terms

| | |
|--------------------|---|
| Accuracy | Any objective measurement of a tool's ability to accurately detect BED/BN. |
| Suitability | Views and opinions of health care practitioners (HCPs) or patients on the usage of screening tools within the primary care context and/or on BED/BN |

153

154 **Study selection**

155 First, title and abstracts were screened by two independent reviewers (JE, RY) using
156 EndNote. Potentially relevant studies were subsequently retrieved and screened in full
157 text (JE, RY). After full-text screening, potential articles were assessed for inclusion
158 by a third and fourth independent reviewer (SK, JS). Disagreements were resolved via
159 discussion.

160

161 **Data extraction**

162 Data from the included studies were extracted on the study-level by JE and RY.
163 Extracted data included title, authors, year and country of publication, design,
164 diagnostic and screening tool used, recruitment setting, sample characteristics and
165 accuracy and suitability results.

166

167 **Quality Assessment**

168 The quality of the papers was assessed by RY, SK and JS independently using the
169 Mixed Methods Appraisal Tool (MMAT) (28). The MMAT assesses quality using a set
170 of questions depending on the study design. We identified the design of the studies
171 included for data analysis and then chose the corresponding measure.

172 **Data Analysis**

173 Due to the amount and type of studies included, a meta-analysis was neither
174 appropriate nor feasible for data synthesis, hence we deviated from our original
175 protocol. Extracted data was synthesised using a convergent segregated approach as
176 described by the Joanna Briggs Institute (29), and a narrative summary was created
177 to describe the accuracy and suitability of EDs' screening tools reporting separately
178 for BED and BN.

179

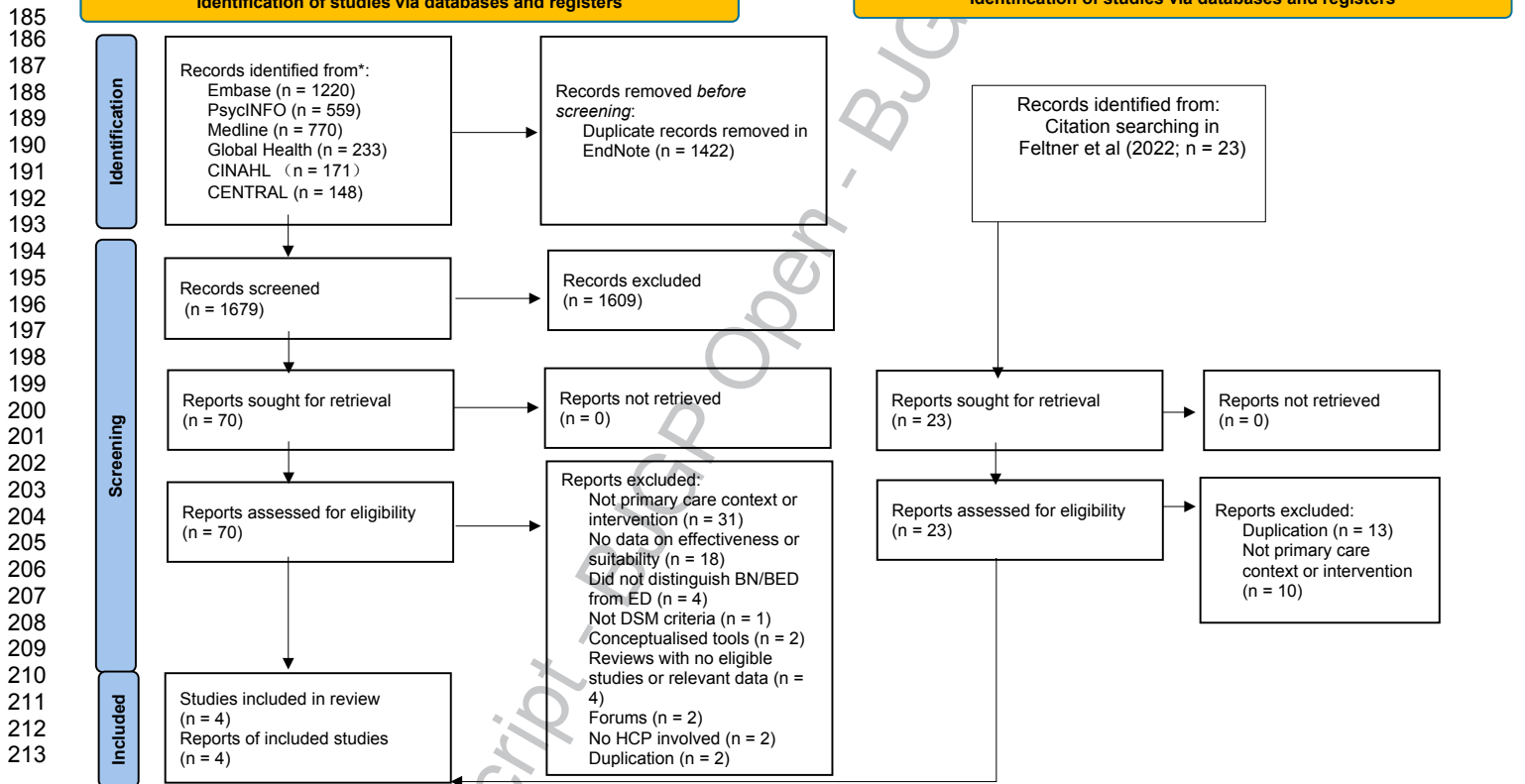
180

Results

181 **Study Selection**

182 The search results identified 3101 records, of which 1422 were duplicated. 70 studies
183 were eligible for full-text screening, of which four studies were included in data
184 analysis. See Figure 1. for detailed selection process.

BED & BN screening tools review



214 **Figure 1.** Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA) flow diagram.

215 **Overview of Studies**

216 The included studies were conducted in the UK (30), US (31,32) and Spain (33). Three
217 studies reported on the accuracy of screening tools using sensitivity and specificity
218 (30,32,33). Two studies reported on suitability (31,32). Two accuracy studies
219 compared screening tools to clinical interviews. See Table 2. for further details.

220

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221 **Table 2.** Characteristics of included studies

| Reference | Location | Tool administrator | Population | Gender (%) | Tool | Standard procedure | Condition | Data collection | Data analysis | COSMIN Dimensions |
|--------------------------|----------|---|---|--|--------|--|---------------|---|--|-------------------|
| McClure et al. 2020 (31) | USA | Nurse practitioners (N=3), Registered nurse (N=1) | Adolescent family clinic attendees without ED diagnosis (N = 7) | PCP gender unavailable Patients: M (57.1%), F (42.9%) | EDE-Q | No comparison | BED | Screening: in-person survey during clinic visit; PCP views: pre- and post-implementation survey | Qualitative Analysis, not further specified | Not reported |
| Herman et al. 2017 (32) | USA | N/A | Primary care physicians serving adults (N=122), Psychiatrists (N=123) | PCPs: M (69.7%), F (30.3%) Psychiatrist: M (62.6%), F (37.4%) | BEDS-7 | No comparison | BED | Online surveys of PCPs | Descriptive statistics | Not reported |
| Luck et al. 2002 (30) | UK | Primary care professional | Sequential general practice attendees (N=341) | F (100%) | SCOFF | Clinical diagnostic interview based on DSM-4 (10-15 minutes) | BN, AN, EDNOS | Cross-sectional questionnaires (2 minutes) | Comparison with clinical interview diagnosis | Not reported |

BED & BN screening tools review

| | | | | | | | | | | |
|---------------------------------|-------|---------------------------|--|----------|---------------|---|---------------|--|--|--------------|
| Garcia-Campayo et al. 2005 (33) | Spain | Primary care professional | General practice attendees with a probable ED diagnosis assessed by GP (N=203) | F (100%) | Spanish SCOFF | Clinical Assessment in Neuropsychiatry (SCAN) | BN, AN, EDNOS | Cross-sectional questionnaires (2 minutes) | Comparison with clinical interview diagnosis | Not reported |
|---------------------------------|-------|---------------------------|--|----------|---------------|---|---------------|--|--|--------------|

222

223 **Risk of Bias/Quality**

224 All studies were assessed to have medium quality and a medium risk of bias (See Tables 3a and 3b).

225 **Table 3a.** Quality appraisal of included studies (Mixed methods)

| Author (year) | 5.1. Is there an adequate rationale for using a mixed methods design to address the research question? | 5.2. Are the different components of the study effectively integrated to answer the research question? | 5.3. Are the outputs of the integration of qualitative and quantitative components adequately interpreted? | 5.4. Are divergences and inconsistencies between quantitative and qualitative results adequately addressed? | 5.5. Do the different components of the study adhere to the quality criteria of each tradition of the methods involved? |
|--------------------------|--|--|--|---|---|
| McClure et al. 2020 (31) | Yes | Yes | Yes | Yes | Can't tell |

226

227

BED & BN screening tools review

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229 **Table 3b.** Quality appraisal of included studies (Quantitative descriptive)

| Author (year) | 4.1. Is the sampling strategy relevant to address the research question? | 4.2. Is the sample representative of the target population? | 4.3. Are the measurements appropriate? | 4.4. Is the risk of nonresponse bias low? | 4.5. Is the statistical analysis appropriate to answer the research question? |
|---------------------------------|--|---|--|---|---|
| Luck et al. 2002 (30) | Yes | Can't tell | Yes | Can't tell | Yes |
| Garcia-Campayo et al. 2005 (33) | Yes | Can't tell | Yes | Can't tell | Yes |
| Herman et al. 2017 (32) | Yes | No | Yes | Can't tell | Yes |

231 **Narrative Synthesis**

232 **Binge Eating Disorder (BED)**

233 Herman et al. (32) explored the use of the BEDS-7, a condition-specific screening
234 tool, among both HCPs in primary care and psychiatrists, with over 75% of
235 respondents considering BEDS-7 to be either “very” or “somewhat” valuable and easy
236 to use. According to HCPs, the primary reason for usage was to identify and initiate
237 discussion on binge eating, while forgetfulness was the main reason for failing to use
238 BEDS-7. Despite positive responses about usage, limited conclusions can be made
239 about the suitability of BEDS-7 for BED. McLure et al. (31) reported similar findings
240 on the usefulness of EDE-Q, a general ED screening tool, among HCPs in primary
241 care in identifying BED risk factors in adolescents. However, the suitability of EDE-Q
242 to the primary care setting was reported to be only adequate. While some HCPs
243 acknowledged the potential positive benefits of using EDE-Q on adolescents’ health,
244 others were unsure about the potential effects that EDE-Q integration into practice
245 could have on patient care delivery. The main concerns about the implementation of
246 screening into practice were stigma, lack of trust in screening, HCPs, and insufficient
247 screening time. Only one HCP reported confidence in integrating EDE-Q without
248 disrupting patient care. Overall, while a condition-specific tool is reported to be more
249 useful and easier to use than a general ED screening tool, both have limitations in
250 their implementation in practice. Hence, no firm conclusions can be made about the
251 suitability of BEDS-7 and EDE-Q due to limited data and a limited sample.

252

253 Regarding accuracy, no studies reported on specificity and sensitivity measures for
254 BED. Two studies (30,33) reported accuracy for Eating Disorder Not Otherwise
255 Specified (EDNOS), which included BED in the DSM-IV. However, due to the separate

256 categorisation for BED in DSM-5, the statistical results did not allow for conclusions
257 regarding the accuracy of screening for BED.

258 **Bulimia Nervosa (BN)**

259 No studies reported on the suitability of screening tools for BN or the primary care
260 setting. Studies focusing on accuracy reported on sensitivity, specificity, and
261 applicability in different patient populations. Both Garcia-Campayo et al. (33) and Luck
262 et al. (30) reported high sensitivity of the SCOFF questionnaire (97.88%; 100%) for
263 detecting BN, which might be due to the population used in the study. However,
264 differences between SCOFF versions are noted with Garcia-Campayo et al. (33)
265 reporting high specificity (94.4%) for the Spanish version with a cut-off at >2 positive
266 responses, while Luck et al. (30) reported a lower specificity (89.6%) for the English
267 SCOFF. However, this data did not separate the outcome for AN and BN; thus, we
268 cannot make a strong conclusion for BN. Luck et al. (30) reported a positive predicted
269 value of 24.4% (12.9%-39.5%) for SCOFF. Garcia-Campayo et al. (33) argued that
270 this predictive value could be due to low ED prevalence in primary care. In two EDNOS
271 cases, non-disclosure of symptoms resulted in missed cases. Hence, the authors (30)
272 summarised that overidentification in this case could be acceptable, as patients who
273 do not meet BN diagnostic criteria are harder to detect, and perhaps further
274 questioning is needed after a positive screening result instead of automatic referral.

275

276 Despite both condition-specific and general ED screening tools being considered
277 useful by HCPs for BED, feasibility concerns limited their perceived suitability in
278 primary care. Similarly, with screening accuracy for BN, SCOFF demonstrated strong
279 sensitivity but lower specificity, indicating a high risk of false positives. The lack of
280 DSM-5-specific validation for BED screening and the limited heterogeneity in BN

281 screening studies highlight the need for further research to refine tool applicability
282 across diverse primary care populations.

283

284

Discussion

285 Summary

286 This systematic review aimed to explore the accuracy and suitability of ED screening
287 tools on BED/BN in a primary care setting. Our synthesis highlights that screening
288 tools for BED are perceived as useful by HCPs but face feasibility challenges due to
289 limited data on their accuracy. SCOFF showed high sensitivity and limited specificity
290 for BN with no data on suitability. Overall, there was a limited amount of evidence
291 available to draw strong conclusions. This could be due to limited funding for ED
292 research (34), debates around mental health screening (35) and the focus on
293 secondary care in this field due to prioritisation of AN and low-weight BN treatment.

294

295 Strengths and limitations

296 To the authors' knowledge, this is the first systematic review exploring the accuracy
297 and suitability of ED screening tools for BED/BN in a primary care setting. A rigorous,
298 systematic approach was used to address this, combining qualitative and quantitative
299 literature, generating an in-depth understanding. However, the small number of eligible
300 studies, all of which had a medium risk of bias, lowered the reliability of conclusions.
301 The evidence was not the most balanced, with only the accuracy or suitability of tools
302 for each condition, not allowing a comparison of the tools' performances for BN or
303 BED. Furthermore, the differences in populations used to test the tools, combined with
304 limited reporting of COSMIN dimensions, did not allow us to draw strong conclusions.
305 However, it did highlight significant gaps in this field. A notable limitation was the

306 reliance on studies using DSM-IV criteria for BN, which do not fully align with the
307 current DSM-5 diagnostic criteria. Future research should prioritise validating existing
308 tools under the DSM-5 criteria and explore their use in a primary care setting, to
309 simulate how screening can be realistically implemented in HCPs' routines. This could
310 improve the understanding of the accuracy and suitability of screening tools for a wider
311 population.

312

313 **Comparison with existing literature**

314 Our findings suggest that even if a screening tool is accurate, its implementation into
315 practice might be limited. This is similar to the literature on primary care delivery, which
316 suggests that limited consultation times available for GPs pose a barrier to
317 implementing additional screening (36). Furthermore, Johnston et al (37) reported
318 HCP concerns about follow-up actions after a positive screen due to potential
319 differences in patient expectations around BED/BN management and the available
320 pathways in their systematic review exploring the feasibility of eating disorders in
321 primary care. This was further supported by Bryant et al (38), whose systematic
322 review recommended that a clear post-positive screen procedure needs to be in place
323 for effective referral and treatment. Hence, further investigation is needed into the
324 practical integration of BED/BN screening into primary care.

325

326 Findings from this review suggest that SCOFF might not be suitable for BED/BN and
327 for the primary care setting. This is in line with the literature, which suggests that
328 SCOFF was developed without using a co-design framework (39), potentially making
329 it less suitable for BED/BN and primary care. However, not using co-design in the
330 development of tools is not unheard of, as a systematic review reported limited input

331 from individuals with lived experiences on clinical tool design, administration or
332 evaluation (40). Stakeholder engagement is crucial in the development process of
333 screening tools as it can improve accuracy, suitability, engagement and
334 implementation into practice as seen in PHQ-9 (41). Hence, lived experience of
335 BED/BN is important to be included in general EDs screening tools to improve
336 suitability and potentially accuracy (42).

337

338 Our findings highlight that SCOFF produced mixed results regarding its accuracy on
339 BN identification in primary care. In clinical samples, SCOFF has been largely
340 sensitive (69.6%-100%) and specific (73.6%-89.6%) (43–45). However, this
341 decreased in general population samples, where sensitivity ranges from 53.7% to
342 77.4%, and specificity from 60.5% to 93.5% (46–48) which is consistent with our
343 findings. Kutz et al. (19) commented that SCOFF was initially developed and validated
344 in care-control studies in specific and homogeneous populations with a high
345 prevalence of ED, such as young women with AN and BN. This suggests that SCOFF
346 might not be effective for BED in primary care in a general population, especially
347 considering the varying demographics of general practice populations, variations in
348 clinical strategy and governance between practices. Given that primary care settings
349 are the first point of contact with healthcare and are the main hubs to identify BED and
350 BN early, additional validation of SCOFF in more heterogeneous samples is needed
351 (19,38).

352

353 **Implications for Research and/or Practice**

354 Our findings and the existing literature show that further research is needed to better
355 understand the accuracy and suitability of ED screening tools for BED/BN in primary

356 care. However, it is important to highlight that it is the psychometric properties of
 357 existing tools that need improvement, as suggested by Hay et al. (24), rather than
 358 developing new tools. Based on the findings of this review, a summary of
 359 recommendations is presented by the authors in Table 4.

360 **Table 4.** Recommendations based on the review

| Recommendation | Explanation |
|---|--|
| <p data-bbox="539 596 688 659">Validation Studies</p> <p data-bbox="266 701 516 974">Further understanding of currently available condition-specific and general ED screening tools</p> | <p data-bbox="824 596 1357 903">The sensitivity and specificity of the BEDS-7 and EDE-Q, as well as the accuracy of the SCOFF for BED, should be evaluated in a primary care setting. Further empirical evidence is needed in the form of, for example, randomised controlled trials or large data set studies to measure the health benefits of screening for BED and BN.</p> |
| | <p data-bbox="539 907 799 940">Suitability studies</p> <p data-bbox="824 907 1357 1075">There is a need to assess the needs of stakeholders (primary care staff and lived experience) to address gaps in the suitability of existing screening tools for BED and BN in primary care.</p> |
| <p data-bbox="344 1117 721 1180">Exploring and Addressing Implementation Barriers</p> | <p data-bbox="824 1079 1357 1213">Further research should explore practical strategies to overcome barriers to screening implementation for eating disorders.</p> |
| <p data-bbox="272 1255 792 1318">Development and Implementation of Screening Protocols</p> | <p data-bbox="824 1218 1357 1352">development of a clear post-positive screening protocol is recommended to tackle potential false-positives and barriers to screening implementation</p> |

361

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368 **Ethical approval**

369 Not applicable.

370 **Data availability**

371 Materials and data used for the conduct of this research are available from the study
372 authors on request.

373 **Competing interests**

374 The authors declare no competing interests.

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378 Attribution (CC BY) licence to any Author Accepted Manuscript version arising from
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380

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BED & BN screening tools review

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