

1 **Diagnosing ADHD in adults in randomized controlled studies: A scoping review**

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13 Running title: Diagnosing ADHD in adults

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16 **Abstract**

17 Background: The diagnosis of ADHD in adults is on the rise. Applying the ADHD  
18 diagnosis, which originally was described in children, to adults have involved a  
19 “subjectivization” of some of the diagnostic criteria, i.e., some behavioral features  
20 (signs) in children have become experiences (symptoms) in adults. These issues raise  
21 the question of how ADHD is best diagnosed in adults? Thus, we examined how ADHD  
22 is diagnosed in adults in research.

23 Methods: A review of how ADHD is diagnosed in adults in randomized controlled  
24 studies (RCTs).

25 Results: We include 292 RCTs. We found substantial variation in and no consensus  
26 about the diagnostic method. More than half of the studies did not seem to include an  
27 assessment of general psychopathology, and only in 35% of studies was the ADHD  
28 diagnosis allocated by a psychiatrists or psychologist. More than half of studies  
29 included patients with psychiatric comorbidity.

30 Conclusion: These findings raise concerns about the validity of the ADHD diagnosis in  
31 many of the included RCTs. It is worrying that securing a reasonably accurate diagnosis  
32 is not prioritized in more than half of the studies. If neither clinicians nor researchers  
33 can rely on the basic fact the patients in scientific studies diagnostically resemble the  
34 patients they are facing, scientific studies risk losing their clinical relevance. Since  
35 RCTs can lead to changes in clinical practice, they must be conducted carefully. To  
36 advance research on adult ADHD, the quality of the diagnostic assessment must be  
37 prioritized, requiring comprehensive differential diagnosis by a skilled psychiatrist or  
38 psychologist.

39 **Keywords**

40 Methodology; Diagnostic Criteria; Comorbidity; Psychopathology

41

42 **Introduction** []

43 The number of adults receiving a diagnosis of Attention Deficit/Hyperactivity Disorder  
44 (ADHD) is increasing [1-2]. In the World Federation of ADHD International Consensus  
45 Statement from 2021 it is estimated that ADHD occurs in 2,5% of adults [3]. However,  
46 a recent systematic review and meta-analysis found a global prevalence of symptomatic  
47 adult ADHD of 6.75% in 2020, corresponding to more than 366 million affected adults  
48 globally [4]. This number also includes individuals, who were diagnosed in childhood  
49 and who remained symptomatic in adulthood. However, several longitudinal studies  
50 have shown that most individuals with adult ADHD have not received a diagnosis of  
51 ADHD in childhood [4-6].

52 One factor that has been discussed as a contributing cause to the increasing number of  
53 adults receiving an ADHD diagnosis, in addition to increased clinical awareness, is the  
54 growing visibility of ADHD on social media platforms, where users are exposed to  
55 symptom descriptions and personal accounts that may prompt self-identification and  
56 help-seeking behavior [1, 7].

57 Originally, ADHD was described in children. The scientific origin of ADHD is by many  
58 considered to the work of George F. Still at the turn of the 20th century, but clinical  
59 descriptions of what we today call ADHD can be found a century earlier in the works of  
60 Alexander Crichton [8]. In 1968, the diagnostic category of Hyperkinetic Reaction of  
61 Childhood (or Adolescence) was included in DSM-II, which described the disorder in  
62 terms of “overactivity, restlessness, distractibility, and short attention span, especially in  
63 young children; the behavior usually diminishes in adolescence” [9].

64 In the subsequent versions of the DSM [10-15], the diagnostic criteria for ADHD have  
65 been diluted and become more inclusive. Unlike most other adult mental disorders,  
66 which are defined by a combination of diagnostic criteria targeting behavioral and  
67 experiential anomalies, i.e., signs and symptoms, the diagnosis of ADHD is based on  
68 behavioral features (signs). Initially, this could hardly be any different, as the original  
69 diagnostic criteria specified observable behavioral features in children reported by  
70 adults (e.g., parents or teachers). Thus, the possibility of diagnosing ADHD in adults  
71 has involved what might be called a “subjectivization” of the diagnostic criteria of  
72 ADHD. Instead of basing a diagnostic assessment on reports of observed behavioral  
73 features (signs) from parents or teachers (e.g., of “excessive running or climbing (...)  
74 having difficulty sitting still” [10, p. 41] or of “interrupting, grabbing objects (...)  
75 excessive talking and by an inability to play quietly” [11, p.50], the adult person must  
76 now herself consider if she believes she, e.g., has “difficulty sustaining attention in a  
77 tasks”, is easily distracted by “unrelated thoughts”, “squirms in seat”, feels “restless”,  
78 etc. [14, p.59f.]. She must also reflect upon whether she believes that some of these  
79 features were present in her childhood, making recall bias a crucial issue. This change in  
80 the perception of the diagnostic criteria from being ‘signs’ to being ‘symptoms’ may  
81 have lowered the diagnostic thresholds of ADHD.

82 This change in the perception of the criteria is also reflected in various national  
83 guidelines for diagnosing and managing ADHD in adults. Such guidelines often  
84 recommend applying diagnostic *interviews* for assessing ADHD. Yet, the national  
85 guidelines do not provide the level of evidence for interviews specifically aimed at  
86 diagnosing ADHD, e.g., European guidelines [16], UK guidelines [17, 18], and  
87 Australian guidelines [19]. A recent meta-analysis of self-report diagnostic methods for  
88 ADHD showed that they often yielded false-positive diagnoses [20]. Another systematic

89 review found that methods of diagnosing ADHD in adulthood varied widely with  
90 respect to source of information, diagnostic instruments, diagnostic symptom threshold,  
91 and whether impairment was required for making the diagnosis. Here, sole reliance on  
92 self-reports was linked to low diagnostic persistence estimate [21].

93 The above-described changes in or perception of the diagnostic criteria for ADHD may  
94 explain some of the global increase in prevalence. Moreover, the possibility of  
95 diagnosing ADHD in adults raises several issues, and scholars have stressed the need  
96 for examining the validity of the diagnostic category of ADHD in adults [22, 23].  
97 Among the issues are difficulties in defining what “impaired functioning” is. Many  
98 adults endorse experiences that could perhaps sound like symptoms of ADHD [24] but  
99 if, say, experiences of inattention do not interfere with functioning, such experiences  
100 should not have the status of symptoms of ADHD according to DSM-5 [15].

101 In sum, the diagnostic criteria pose challenges for diagnosing ADHD in adults, since i)  
102 the original diagnostic criteria and tools for assessing ADHD were developed for use in  
103 children [25], ii) retrospective recall of childhood symptoms is notoriously poor [26],  
104 iii) the ADHD criteria were not tested in adults in the DSM-5 field trials, and iv)  
105 collateral information (e.g., from school teachers or parents), which previously was the  
106 foundation for the making the ADHD diagnosis, is difficult, if not impossible, to  
107 retrieve or access in adults [20]. Thus, we decided to examine how research studies have  
108 handled the challenges surrounding the ADHD diagnosis.

109

## 110 **Aim**

111 The aim of this study was to review how ADHD has been diagnosed in adults in  
112 Randomized Controlled Trials (RCTs).

113

114 **Methods**

115 *Search strategy and selection criteria*

116 Following the PRISMA guidelines [27], we conducted a review, focusing on the  
117 methods of diagnosing ADHD in adults in RCTs. We focused on RCTs as they rank  
118 very high in the hierarchy of evidence in evidence-based medicine and thus are likely to  
119 represent high-quality empirical research e.g., [28]. To be clear, we were only interested  
120 in the diagnostic methods and not the findings of these RCTs. We searched the  
121 PubMed, using the following search string 'ADHD OR Hyperkinetic Disorder AND  
122 Adult' on December 5, 2024. We restricted our search to human and RCT, using  
123 PubMed filters. Inclusion criteria were RCT studies with adult samples (participants at  
124 least 18 years old) with a diagnosis of ADHD/Hyperkinetic disorder, studies written in  
125 English, and studies must include a direct patient assessment. Conference abstracts were  
126 excluded as well as studies relying on registry data. Authors IS and JN screened all  
127 titles and abstracts for inclusion in the study. Disagreement was resolved through  
128 consensus between the authors. We chose to only search one database (PubMed),  
129 because the aim was to get an overview of the methodology used to allocate the ADHD  
130 diagnosis in adults in RCTs.

131

132 *Data extraction*

133 We extracted data on diagnostic methods, on whether an assessment of general  
134 psychopathology was made, whether the study included patients with comorbid  
135 disorders in the sample, and on the person allocating the diagnosis (e.g., a medical  
136 doctor, psychologist, trained rater, or unknown).

137 *Categories*

138 The diagnostic methods were categorized into five main groups based on how the  
139 ADHD diagnosis had been established: 1) studies that only used an ADHD specific  
140 interview/rating scale; 2) Studies that only used clinical diagnoses, 3) studies that used a  
141 structured interview for assessing general psychopathology, 4) studies that used a semi-  
142 structured interview for assessing general psychopathology, and 5) studies that used  
143 other approaches. Some of these categories were further subdivided if there was an add-  
144 on to the main diagnostic approach, e.g., an ADHD specific rating scale in addition to a  
145 structured interview for assessing general psychopathology.

146 The categorization process followed a systematic strategy:

147 1) Diagnostic tools: in each of the included studies, we identified the specific diagnostic  
148 instruments used (e.g., structured interviews, self-report scales, clinician-administered  
149 ADHD-specific interviews). If this information was not explicitly stated in the study  
150 itself, we traced it back to a *parent paper* (i.e., an original or referenced study) that  
151 provided details on the diagnostic method used. If studies did not describe assessing  
152 general psychopathology or report procedures that would allow such an assessment, we  
153 concluded that no such assessment had been made.

154 2) Differential diagnosis and hierarchical considerations: The presence of a systematic  
155 differential diagnostic process was determined based on the study's inclusion and  
156 exclusion criteria or if it was explicitly described in the study, e.g., using a method  
157 allowing for differential diagnosis. We assessed whether studies adhered to a classical  
158 diagnostic hierarchy [29, 30], prioritizing organic disorders, followed by schizophrenia  
159 spectrum and bipolar disorders, and then other psychiatric conditions. If a study  
160 explicitly stated that such hierarchical exclusion criteria were applied, it was categorized  
161 accordingly.

162 3) Comorbidity: The handling of psychiatric comorbidities was assessed based on  
163 whether studies allowed participants with additional diagnoses (e.g., anxiety,  
164 depression) beyond ADHD.

165 4) Interviewer Qualifications: The qualifications of the individual conducting the  
166 diagnostic assessment were extracted from the article. We specifically looked for  
167 whether the study specified that a psychiatrist, psychologist, trained rater, or another  
168 professional was responsible for diagnosing participants. If this information was not  
169 available, we categorized it as "unknown."

170

#### 171 *Definitions*

172 In this study, we defined a structured diagnostic interview as an interview consisting of  
173 a set of predetermined questions that should be presented in a definite order. Diagnostic  
174 information is yielded based on the patient's responses to the questions and on the  
175 interviewer's observations (an example of a structured interview for general  
176 psychopathology following this definition is the *Structured Clinical Interview for DSM*  
177 *(SCID-I)* [31]. Structured diagnostic interviews aim at identifying symptoms that meet  
178 diagnostic Criteria [32] and which can result in allocation of a diagnosis. We defined a  
179 semi-structured diagnostic interview for general psychopathology as a conversational  
180 interview, aiming at eliciting psychopathological information but without using  
181 preformulated questions presented in a definite order. The interviewer's questions  
182 function as triggers that encourage the patient to talk, and through his or her comments  
183 and questions, the interviewer steers the interview to obtain the relevant  
184 psychopathological data necessary for allocating a diagnosis [33].

185 **Results**

186 The PubMed search yielded 706 publications. 376 publications were excluded, leaving  
187 330 which were assessed for eligibility. 38 were excluded for not meeting the inclusion  
188 criteria. We ultimately included 292 RCTs (see supplementary material for the list of  
189 the included studies). The study selection can be seen in Figure 1.

190

191 FIGURE 1 ABOUT HERE

192

193 *Diagnostic methods*

194 The diagnostic methods used to allocate ADHD diagnoses to adults in the included  
195 studies are shown in Table 1. Generally, the methods used to diagnose ADHD in adults  
196 varied considerably, and 49.7 % of the studies allocated the ADHD diagnosis without  
197 an assessment of general psychopathology. This group of studies is composed of studies  
198 using only clinical diagnoses, with (29.5%) or without (12.7%) an additional ADHD  
199 specific rating scale and studies using only an ADHD specific interview/rating scale  
200 (7.5%).

201 Among the studies that included an assessment of general psychopathology, the ADHD  
202 diagnosis was allocated either based on this assessment alone or in combination with a  
203 self- or clinician rated scale targeting ADHD. When dividing studies that assessed  
204 general psychopathology into studies using a structured vs. semi-structured interviews,  
205 the vast majority of studies used a structured diagnostic interview (see Table 1 for  
206 details).

207

208 TABLE 1 ABOUT HERE

209

210 *Who allocated the diagnosis?*

211 In 190 studies (65%), the person who conducted the diagnostic assessment was either  
212 not reported, not a psychiatrist or a psychologist, or it was made by a computer.

213

214 *Comorbidity*

215 From the total of 292 studies, 157 studies (53.8%) accepted some kind of psychiatric  
216 comorbidity in their sample. Moreover, 256 of the studies (87.7%) stated that they  
217 adhered to a diagnostic hierarchy, e.g., a diagnosis of an organic condition overrules an  
218 ADHD diagnosis. Simultaneously, most of these studies did not apply a method that  
219 included assessment of whether the patients suffered from mental disorders, which they  
220 claimed would overrule the ADHD diagnosis.

221

222 TABLE 2 ABOUT HERE

## 223 **Discussion**

224 This review examined how the ADHD diagnosis has been allocated in 292 empirical  
225 studies of adult patients. Overall, there was considerable variation in and no consensus  
226 about the method used for diagnosing ADHD. Moreover, the review identified three,  
227 interrelated methodological issues that raise concern about the quality of the allocated  
228 diagnoses in a substantial part of these studies.

229 First, half of the included studies did not describe conducting an examination of general  
230 psychopathology or report procedures that would have allowed for such an assessment,  
231 which is necessary for allocating any diagnosis. In these studies, either no diagnostic  
232 assessment was made (relying solely on clinical diagnoses), or the diagnosis was  
233 allocated based on results from a self- or clinician-rated scale targeting only ADHD,

234 sometimes in combination with a clinical diagnosis. Just stating that a clinical diagnosis  
235 was used without any description of how and who made the clinical diagnosis is not  
236 sufficient as this can cover a wide range of diagnostic methods, diagnosis been made by  
237 untrained staff, different diagnostic traditions, errors etc [34, 35], and provides no  
238 transparency, which is of greatest importance in research [36]. However, without an  
239 assessment of general psychopathology, it is impossible to make a differential-  
240 diagnosis, e.g., ruling out the possibility of other (often more severe) mental disorders  
241 that may present with similar signs or symptoms. Although 87.7% of the studies  
242 asserted that they adhered to a diagnostic hierarchy, this was practically impossible in  
243 most of these studies as they included no assessment of general psychopathology.  
244 Naturally, a general psychopathological assessment is crucial in the case of ADHD,  
245 because none of the diagnostic criteria of ADHD are specific for ADHD and similar  
246 signs and symptoms can be seen in a range of other mental disorders such as substance  
247 use disorder, schizophrenia spectrum disorders, mood disorders, etc. For example,  
248 attention deficits and motor disturbances have been described as parts of the  
249 psychopathology of schizophrenia since Bleuler coined the concept of schizophrenia in  
250 the early 20th century [37]. Also disorders such as depression, anxiety, and trauma-  
251 related conditions can give rise to attentional complaints that mimic ADHD symptoms.  
252 These overlaps can lead to diagnostic confusion, particularly in adult populations, where  
253 developmental history may be less readily available or prone to recall bias. Mølstrøm et  
254 al. [38] highlight this issue in a study of first-admission psychiatric patients,  
255 demonstrating how affective and anxiety symptoms often manifest in non-specific  
256 complaints, including difficulties with concentration and attention. These findings  
257 underscore the importance of a thorough differential diagnostic process that takes into  
258 account the non-specific nature of attentional symptoms and the disease pictures they

259 are embedded in. Thus, it is highly problematic that half of the included studies diagnosed  
260 ADHD apparently without assessing general psychopathology.

261

262 Although structured diagnostic interviews for long have been regarded as a “gold  
263 standard” for diagnosing mental disorders in research, several studies have reported  
264 serious limitations with structured diagnostic interviews. For example, studies  
265 comparing the agreement of diagnoses allocated by a trained rater using structured  
266 interview with best consensus diagnoses allocated by experienced psychiatrists using  
267 semi-structured diagnostic interviews and including all available information (e.g., from  
268 the clinic and relatives) have reported worryingly low overall concordances [39, 40].

269 The authors recommend that structured interviews should only be used in research with  
270 certain precautions, e.g., only by skilled medical doctors or psychologists and not by  
271 for-the-purpose trained raters. In our review, only 12.7% of the studies used a semi-  
272 structured interview to assess general psychopathology (1.4% used only a semi-  
273 structured interview, 2.4% used it in combination with a self- or clinician-rated scale for  
274 ADHD, and 8.9% used it in combination with a structured interview and a ADHD  
275 specific rating scale). The high reliance on structured interviews for assessing general  
276 psychopathology, amounting to a total of 45.9%, may have compromised the validity of  
277 the allocated ADHD diagnoses in these studies.

278

279 Second, only approximately one third of the studies reported that the diagnosis had been  
280 allocated by a medical doctor or a psychologist. This is also a cause for concern,  
281 because significant discrepancies repeatedly have been demonstrated for psychiatric  
282 diagnoses allocated by trained raters vs. clinicians [39, 40]. Moreover, self-rating  
283 measures to diagnose ADHD have a very low positive predictive value, often in the

284 10% range [20]. The reliance on especially trained raters and self-rating scales for  
285 diagnosing ADHD elevates the likelihood of diagnostic errors.

286

287 Third, more than half of the studies included participants that had some kind of  
288 psychiatric comorbidity. Although developmental disorders were removed as an  
289 exclusion criterion for the ADHD diagnosis in DSM-5 [20], other mental disorders still  
290 function as exclusion criteria for making the ADHD diagnosis—i.e., ADHD cannot be  
291 diagnosed if the symptoms occur only during the course of schizophrenia or another  
292 psychotic disorder or if the ADHD symptoms are better explained by other disorders  
293 such mood disorders, anxiety disorder, personality disorders, and substance use  
294 disorder, etc. [15]. The above-described omission of assessment of general  
295 psychopathology in half of the studies makes it impossible to know if the ADHD  
296 symptoms here occurred during the course of another disorder or if they were better  
297 explained by another disorder. In these studies, we cannot conclude that the ADHD  
298 diagnosis were made in accordance with the diagnostic guidelines. Of course, ADHD  
299 can, in some cases, be diagnosed as a comorbid condition [41].

300

301 The overall implication of these methodological issues is that the validity of the ADHD  
302 diagnoses in many of the included RCTs appears to be severely compromised. If these  
303 diagnoses were allocated on insufficient grounds, it has most likely affected the  
304 outcome of these trials, e.g., results of interventions in samples, whose diagnostic status  
305 were assumed to be ADHD but which in fact remain diagnostically unascertained. It  
306 also implies that comparing results across studies in reviews or meta-analyses comes  
307 with a high degree of uncertainty. Here, it may prove useful to exclude studies relying  
308 on insufficient diagnostic methods. For empirical studies researching subjects related to

309 specific disorders, e.g., testing effects of treatment in ADHD, prioritization of careful  
310 allocation of diagnosis is of utmost importance.

311

312 With the sparse knowledge of how ADHD manifests in adults, and the need to rely on  
313 the patients' own descriptions of their behavior as children to diagnose ADHD in adults,  
314 we are, diagnostically speaking, standing on unstable ground. The lack of real-time  
315 external observations of these patients, who are now adults, has transformed some of the  
316 behavioral *signs* of ADHD in children into *symptoms* of ADHD in adults, viz. the  
317 subjectivization of the diagnostic criteria. This change in the perception of some  
318 diagnostic criteria for child vs adult ADHD raises the question as to whether ADHD  
319 diagnosed in childhood and ADHD diagnosed in adulthood is in fact the same disorder.  
320 Most patients, who are diagnosed with ADHD in adulthood, have not been diagnosed  
321 with ADHD in childhood [4-6]. Perhaps some of these adult ADHD patients were  
322 overlooked as children, but a more likely explanation seems to be that many of them did  
323 not attract psychiatric attention as children, because they did not show the same degree  
324 of behavioral manifestations as those children, who were diagnosed with ADHD in  
325 childhood. Again, there is an urgent need to clarify how exactly ADHD presents in  
326 adults and next to establish diagnostic criteria to delineate the disorder from other  
327 conditions that also present with attention- and hyperkinetic phenomena.

328

329 Consequently, it seems premature to include patients with comorbid disorders in the  
330 empirical research studies of ADHD in adults, which nonetheless was the case in more  
331 than half of the studies. Due to the limited knowledge of the ADHD disorder in adults,  
332 the aim must first be to comprehensively examine a sample of ADHD patients without  
333 comorbidities and follow them over time [42]. For now, we do not know if ADHD

334 symptoms in adults are similar in patients with ADHD with or without psychiatric  
335 comorbidities. Psychopathological studies, clarifying the nature of the subjective  
336 experiences of being distracted by “unrelated thoughts” or “feeling restless” etc. in adult  
337 ADHD, may aid differentiating such ADHD symptoms from seemingly similar  
338 symptoms in other mental disorders.

339

340 These diagnostic challenges underscore the importance of transparency and rigor when  
341 conducting empirical studies on ADHD, not the least RCTs, which are considered to be  
342 providing evidence of high quality [43]. Without clear and consistent reporting of  
343 diagnostic methods and procedures, the reliability of findings becomes questionable,  
344 potentially intensifying the difficulties already inherent in studying adult ADHD. As  
345 emphasized in Guidelines for Reporting Health Research: A User’s Manual [36]:  
346 “Poorly conducted trials are a waste of time, effort, and money. The most dangerous  
347 risk associated with poor-quality reporting is an overestimate of the advantages of a  
348 given treatment ... Whatever the outcome of a study, it is really hard for the average  
349 reader to interpret and verify the reliability of a poorly reported RCT. In turn, this  
350 problem could result in changes in clinical practice that are based on false evidence and  
351 that may harm patients.” [36, p. 3]. Transparent reporting is therefore essential, not only  
352 to ensure that RCTs provide reliable and interpretable evidence, but also to safeguard  
353 clinical practice from being guided by potentially flawed “evidence”.

354

355 In conclusion the results of this review point to a worrying shift in the common  
356 understanding of how a psychiatric diagnosis should be allocated in research studies,  
357 with a dwindling awareness of the importance of making as accurate a diagnosis as  
358 possible, which necessarily imply making a comprehensive general psychopathological

359 assessment. If we, both as clinicians and researchers, cannot be reasonably sure that  
360 patients in scientific studies actually suffers from the diagnosis which the study claims  
361 that they do, we cannot rely on the studies findings.

362 Our finding that half of RCTs exhibited little or no interest in securing the validity of  
363 the ADHD diagnosis and that it was unclear who made the diagnosis in 2/3 of the  
364 studies is certainly alarming. The diagnostic assessment is the foundation, which all  
365 subsequent analyses built upon. As long as it remains unclear precisely what disorder is  
366 being examined in scientific studies, the findings of these studies will have limited  
367 value. In this context, it is noteworthy that we reviewed RCTs, and RCTs are considered  
368 high in the scientific evidence hierarchy in evidence-based medicine. Still, many RCTs  
369 had not made an effort to diagnoses *lege artis*, thus rendering the results of their  
370 otherwise comprehensive study questionably.

371

372

### 373 **Author contributions**

374 **I. Studart:** Conceptualization, literature search and selection, data extraction and  
375 writing

376 **M.G. Henriksen:** Conceptualization and writing

377 **J. Nordgaard:** Conceptualization, literature selection, data extraction and writing.

378

### 379 **Data availability statement**

380 Detailed information about dataextraction is available upon request

381

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384

385 **Conflict of interest disclosure**

386 All authors declare no conflict of interests

387

388 **Ethics approval statement**

389 The study is a literature review. All included data and information is already published,

390 thus no ethics approval is needed.

391

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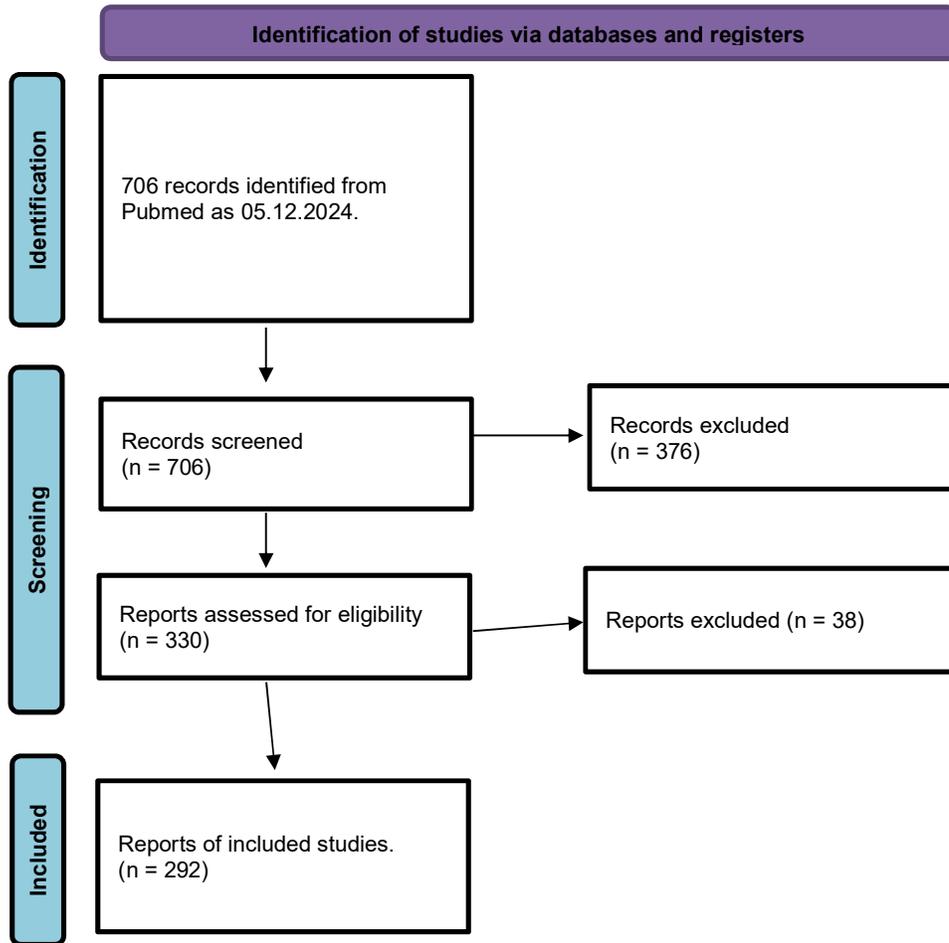
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Figure 1. PRISMA flow diagram for inclusion of papers



556 Table 1. Methods used for allocating ADHD diagnosis and the number of studies using the methods  
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	ADHD specific		Clinical diagnosis		Structured interview		Semi-structured interview		Other
Diagnostic methods	ADHD specific interview or rating scale (either self- or clinician rated)	Clinical diagnosis/ Paper state that patients met DSM or ICD criteria for ADHD	Clinical diagnosis/ Paper state that patients met DSM or ICD criteria for ADHD + ADHD specific rating scale	Structured interview for general psychopathology schedule (SCID or MINI)	Structured interview for general Psychopathology + ADHD specific ratingscale	Structured interview+ semi-structured + ADHD rating scale	Semi-structured interview for general psychopathology (Kiddie SADS)	Semi-structured interview for general psychopathology + ADHD specific ratingscale	Other
Number of papers using the methods (total N=292)	22 (7.5%)	37 (12.7%)	86 (29.5%)	17 (5.8%)	91 (31.2%)	26 (8.9%)	4 (1.4%)	7 (2.4%)	2 (0.7%)

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559 Table 2. Allocation of diagnosis and psychiatric comorbidity  
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<b>Who did the diagnostic interview?</b>	<b>N (%)</b>
Clinician	102 (35%)
Unknown	179 (61%)
Trained rater	10 (3%)
Other (computer allocated diagnosis, confirmed by a neurologist or psychiatrist)	1 (0.5%)
<b>Allow comorbid diagnosis?</b>	
Yes	157 (54%)
No	122 (42%)
Unknown	13 (4%)

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