Are there researcher allegiance effects in diagnostic validation studies of the PHQ-9? A systematic review and meta-analysis

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ABSTRACT

Objectives To investigate whether an authorship effect is found that leads to better performance in studies conducted by the original developers of the Patient Health Questionnaire (PHQ-9) (allegiant studies).

Design Systematic review with random effects bivariate meta-analysis. Search strategies included electronic databases, examination of reference lists and forward citation searches.

Inclusion criteria Included studies provided sufficient data to calculate the diagnostic accuracy of the PHQ-9 against a gold standard diagnosis of major depression using the algorithm or the summed item scoring method at cut-off point 10.

Data extraction Descriptive information, methodological quality criteria and 2×2 contingency tables.

Results Seven allegiant and 20 independent studies reported the diagnostic performance of the PHQ-9 using the diagnostic scoring method. Pooled diagnostic OR (DOR) for the allegiant group was 64.40, and 15.05 for non-allegiant studies group. The allegiance status was a significant predictor of DOR variation (p<0.0001). Five allegiant studies and 26 non-allegiant studies reported the performance of the PHQ-9 at recommended cut-off point of 10. Pooled DOR for the allegiant group was 49.31, and 24.96 for the non-allegiant studies. The allegiance status was a significant predictor of DOR variation (p=0.015). Some potential alternative explanations for the observed authorship effect including differences in study characteristics and quality were found, although it is not clear how some of them account for the observed differences.

Conclusions Allegiant studies reported better performance of the PHQ-9. Allegiance status was predictive of variation in the DOR. Based on the observed differences between independent and non-independent studies, we were unable to conclude or exclude that allegiance effects are present in studies examining the diagnostic performance of the PHQ-9. This study highlights the need for future meta-analyses of diagnostic validation studies of psychological measures to evaluate the impact of researcher allegiance in the primary studies.

Strengths and limitations of this study

- An original study—the first meta-analysis of diagnostic validation studies of psychological measures to evaluate the impact of researcher allegiance.
- Using rigorous methodology—strict inclusion/exclusion and quality assessment criteria.
- We found that the allegiance effect was a significant predictor of the variation of the diagnostic OR in the meta-regression analysis.
- Substantial variability observed in methodological quality of included studies.
- Based on the observed methodological differences between the independent and non-independent studies, we were unable to conclude or exclude that allegiance effects are present in studies examining the diagnostic performance of the Patient Health Questionnaire (PHQ-9).

Research on allegiance effects has a long tradition in psychotherapy research. In this context, allegiance describes the phenomenon that researchers and clinicians who developed a treatment approach or are for other reasons invested in it tend to find larger effect sizes in favour of their treatment than for comparison groups. This finding has been extensively replicated2 3 and is also robust when the quality of research is controlled for. Researcher allegiance is subject of ongoing debates about the design of efficacy studies as well as implications for policy.4 5 Researcher allegiance is also discussed widely in the literature on experimental as well as evaluation research.6 Since the motivational underpinnings of allegiance effects are potentially far more ingrained into human behaviour and decision making than previously thought,7 they may occur commonly in clinical research in general.

Although it has been suggested that allegiance effects may play a role in the validation of psychological screening and case-finding tools (eg, O’Shea et al., in press), systematic
evaluations of this hypothesis are rare and studies that acknowledge potential allegiance effects in such studies mainly come from forensic psychology and psychiatry backgrounds. Diagnostic validation studies are geared at establishing the sensitivity and specificity of a screening or case-finding tool, which is used in practice to differentiate cases from non-cases or to decide about whether further assessment or treatment is indicated or will be offered. An allegiance effect in such studies would be seen in systematically higher sensitivities or specificities if the original author(s) is (are) part of the team of such a study. Such a bias would have a deleterious affect on practice through promising overoptimistic accuracy of the screening or case-finding tool or in evaluating the cost-effectiveness of the measure in a screening or case-finding context.

The depression module of the Patient Health Questionnaire (PHQ-9) is a widely used depression-screening instrument in non-psychiatric settings. The PHQ-9 was developed by a team of researchers, with its development underwritten by an educational grant from Pfizer US Pharmaceuticals. The PHQ-9 can be scored using different methods, including an algorithm based on Diagnostic and Statistical Manual of Mental Disorders (DSM)-IV criteria and a cut-off based on summed-item scores. The psychometric properties of these two approaches have been summarised in two recently published meta-analyses. The goal of the current review is to investigate, based on an established database of PHQ-9 diagnostic validation studies, whether an allegiance effect is found that leads to an increased sensitivity and specificity in studies that were conducted by researchers closely connected to the original developers of the instrument.

**METHODS**

**Study selection**

Similar search strategies were used in both systematic reviews (for full details, please see Manea et al and Morarity et al). Embase, Medline and PsycINFO were searched from 1999 (when the PHQ-9 was first developed) to August 2013 and September 2013, respectively, using the terms ‘PHQ-9’, ‘PHQ’, ‘PHQs’ and ‘patient health questionnaire’. The search strategy is presented in online supplementary appendix 1. The reference lists of studies fitting the inclusion criteria were manually searched and a reverse citation search in Web of Science was performed. The authors of unpublished studies were contacted and conference abstracts were reviewed in an attempt to minimise publication bias.

The following inclusion-exclusion criteria were used:

- **Population:** adult population. **Instrument:** studies that used the PHQ-9. **Comparison (reference standard):** the accuracy of the PHQ-9 had to be assessed against a recognised gold-standard instrument for the diagnosis of either DSM or International Classification of Disease (ICD) criteria for major depression. Studies were included if the diagnoses were made using a standardised diagnostic structured interview schedule (eg, Mini International Neuropsychiatric Interview (MINI), Structured Clinical Interview for DSM Disorders (SCID)). Unguided clinician diagnoses with no reference to a standard structured diagnostic schedule or comparisons of the PHQ-9 with other self-report measures were excluded. Studies were also excluded if the target diagnosis was not major depressive disorder (MDD, eg, any depressive disorder). **Outcome:** studies had to report sufficient information to calculate a 2×2 contingency table for the algorithm or the recommended cut-off point 10.

- **Study design:** any design. **Additional criterion:** we avoided double counting of evidence by ensuring that only one study of those that reported overlapping datasets in different journals were included in the meta-analysis. Citations with overlapping samples were examined to establish whether they contained information relevant to the research question that was not contained in the included report.

**Quality assessment**

Quality assessment was performed using the Quality Assessment of Diagnostic Accuracy Studies (Revised) (QUADAS-2) tool, a tool for evaluating the risk of bias and applicability of primary diagnostic accuracy studies when conducting diagnostic systematic reviews. It covers the areas of patient selection, index test, reference standard and flow and timing. This tool was adapted for the two reviews and quality assessments were carried out by two independent reviewers for all studies included in the reviews.

**Data synthesis and statistical analysis**

We constructed 2×2 tables for cut-off point 10 and the algorithm scoring method. Pooled estimates of sensitivity, specificity, positive/negative likelihood ratios and diagnostic ORs (DOR) were calculated using random effects bivariate meta-analysis. Heterogeneity was assessed using I² for the DOR, an estimate of the proportion of study variability that is due to between-study variability rather than sampling error. We considered values of ≥50% to indicate substantial heterogeneity. Summary receiver operating characteristic curves (sROC) were constructed using the bivariate model to produce a 95% confidence ellipse within ROC space. Each data point in the sROC space represents a separate study, unlike a traditional ROC plot, which explores the effect varying thresholds on sensitivity and specificity in a single study. We undertook a meta-regression analysis of logit DOR using research allegiance as covariate in the meta-regression model. Analyses were conducted using STATA V.12, with the metan, metandi and metareg user-written commands.

**Allegation rating**

We rated authorship on a paper if any of the developers of the PHQ-9—Kurt Kroenke, MD, Robert L. Spitzer, MD and Janet BW Williams—as an indicator of potential allegiance. We also rated as evidence of allegiance as
acknowledged collaborations with the developers of the PHQ-9, even if they were not listed as coauthors or if the authors acknowledged funding from Pfizer to conduct the study.

RESULTS
Overview of included studies
Thirty-one studies reported the diagnostic properties of the PHQ-9. Two studies were included in the algorithm review. The study selection flow charts can be found in online supplementary appendix 2 (figures 1 and 2). The characteristics of these studies are reported in tables 1 and 2 and the results of the methodological assessment are presented in tables 3 and 4.

Algorithm scoring method
Descriptive characteristics
The descriptive characteristics of the included studies are presented in table 1. Seven individual studies that reported the diagnostic performance of the PHQ-9 using the algorithm scoring method were co-authored by the original developers of the PHQ-9, specifically acknowledged one of the developers and support by an educational grant from Pfizer USA, or were co-authored by the first author of a previous study that had also been co-authored by one of the developers. Twenty non-allegiant studies reported the diagnostic properties of the PHQ-9 using the algorithm scoring method.

Three (43%, 3/7) of the allegiant studies were conducted exclusively in hospital settings. The remaining four studies (67%, 4/7) were conducted in different settings or non-exclusively hospital settings: one in primary care and three in mixed settings: psychosomatic walk in clinics and family practices, outpatient clinics and family practices, and primary care and hospital settings. In the non-allegiant group, 13 (65%, 13/20) studies were conducted in hospital settings. Of the remaining seven studies, six were conducted in primary care settings and one in a community sample.

In both groups (non-allegiant and allegiant studies), the majority of studies validated a translated version of the PHQ-9. Two of the studies authored by developers (28%, 2/7), and eight (40%, 8/20) allegiant studies were conducted in English. The mean prevalence of MDD in the group of allegiant studies was 13.4% (range 6.1%–29.2%); in the non-allegiant group it was 15.5% (range 3.9%–32.4%). The mean age of patients in the PHQ-9 developers group was 45.7; all but one study had a mean age in the range of 40–50 years. In the non-allegiant group, the mean age was 54.6 (range 29.3–75.0), with almost half (8) of the studies reporting a mean age of over 60. The percentage of females in the PHQ-9 developers was 56.8% (range 28.6%–67.8%) and in the non-allegiant group was 59.1 (18%–100%).

All allegiant studies used a self-reported PHQ-9, whereas in seven non-allegiant studies (50%, 6/20) the PHQ-9 was administered by a researcher. Apart from Muramatsu et al., all allegiant studies used the SCID as a gold standard; the non-allegiant studies used a wider range of gold standards including SCAN, CIDI, MINI and C-DIS, although the SCID was also frequently used by the independent studies as well (45%, 9/20 studies).

Four out of the seven allegiant studies (57%) did not include a conflict of interest statement. Also, four (57%) of the allegiant studies acknowledged funding from Pfizer. Only one study acknowledged the collaboration with one of the developers of the PHQ-9.

Of the non-allegiant studies, 12 (60%) did not include a conflict of interest statement. It appears that newer studies were more likely to include a conflict of interest statement, which may reflect a recent change in reporting. Funding was acknowledged by most studies (18/20) and most received funding from academic or and health research institutions. Two studies received funding from pharmaceutical companies—Lundbeck and Pfizer, and one study acknowledged that Pfizer Italia provided the Italian version of PHQ-9 and gave the authors permission to use it.

Diagnostic test accuracy
Pooled sensitivity and specificity was calculated separately for the non-allegiant and allegiant studies. Pooled sensitivity for the allegiant studies of the PHQ-9 was 0.77 (95% CI 0.70 to 0.84), pooled specificity was 0.94 (95% CI 0.90 to 0.97) and the pooled DOR was 64.40 (95% CI 34.15 to 121.43). Heterogeneity was high (I²=78.9%). Figure 1 represents the sROCs for this set of studies.

Pooled sensitivity for the non-allegiant studies was lower compared with the developer authored studies group at 0.48 (95% CI 0.41 to 0.91), pooled specificity was the same at 0.94 (95% CI 0.91 to 0.95). The pooled DOR was approximately four times lower at 15.05 (95% CI 11.03 to 20.52) (see figure 1). Heterogeneity was substantial at I²=68.1%.

The meta-regression analysis for algorithm studies with non-allegiant status as the predictor of the DOR showed that non-allegiant status was a significant predictor of the DOR (p=0.0001) and explained a substantial amount of the observed heterogeneity (51.5%).

Quality assessment
The results of the quality assessment using QUADAS-2 are given in table 3 for the studies reporting on the diagnostic performance of the algorithm scoring method. In the patient selection domain, more non-allegiant studies (65%, 13/20) than allegiant (29%, 2/7) met the criterion for consecutive referrals. There were no marked differences on the other two criteria in this domain (avoid
| Study | Country | Setting | Age (years): Mean (SD) | Female (%) | Sample size (n) | Depression rates: % depressed | PHQ-9 characteristics | Diagnostic standard | COI declaration | Funding | Relationship with original developers |
|-------|---------|---------|------------------------|------------|----------------|-----------------------------|----------------------|-------------------|-----------------|--------|--------------------------------------|
| Diez-Quevedo et al | Spain | Medical and surgical tertiary hospitals | M=43 (SD=14.2) | 45.6% | 1003 | Depressed: 8.2% | Administration: self-report | DSM-III-R SCID | a) No COI declaration | b) Funding acknowledged (academic institutions) | c) Not acknowledged |
| Gräfe et al | Germany | Psychosomatic walk-in clinics and family practices | male=41.9 (SD=13.8) | 67.8% | 528 | Depressed: 29.2% | Language: German | DSM-IV SCID | a) No COI declaration | b) Acknowledged funding from Pfizer | c) Not acknowledged |
| Lowe et al | Germany | Outpatient clinics and family practices | male=41.7 (SD=13.8) | 67.1% | 501 | Depressed: 13.2% | Administration: self-report | DSM-IV SCID | a) COI declaration ‘This study was supported by unrestricted restricted grants from Pfizer Germany and from the medical faculty of the University of Heidelberg Germany, and there are no COI’. b) Acknowledged funding from Pfizer and academic institution | c) Not acknowledged |
| Muramatsu et al | Japan | Primary care and general hospital | male=43.3 (SD=16.4) | 59.5% | 131 | Depressed: 28.2% | Language: Japanese | DSM-IV MINI | a) No COI declaration | b) Acknowledged funding from Pfizer | c) Acknowledged one of the developers of the PHQ-9: ‘The authors acknowledge Dr RL Spitzer’ |
| Navinés et al | Spain | General hospital (patients with chronic HCV) | male=43.4 (SD=10.2) | 28.6% | 500 | Depressed: 6.4% | Administration: self-report | DSM-IV SCID | a) All authors declared that they had no COI. b) Role of funding source declared | c) Not acknowledged |
| Spitzer et al | USA | Primary care | male=46 (SD=17.2) | 66% | 3000 (585 received SCID) | Depressed: 10% | Administration: self-report | DSM-III-R SCID | a) No COI declaration | b) Acknowledged funding from Pfizer. ‘Drs Spitzer and Williams receive honoraria and consulting money from Pfizer, which has supported this work’. | c) N/A |
| Thekkumpurath et al | UK | Hospital (cancer patients) | male=61 | 63% | 782 | Depressed: 6.3% (of the whole sample) | Administration: not stated | DSM-IV SCID | a) COI declaration: ‘Supported by Cancer Research UK’ | b) As in a) | c) Not acknowledged |
| Ayalon et al | Israel | | male=75 (SD=8.1) | 40.5% | 153 | Depressed: 3.9% | Administration: researcher administered | DSM-IV SCID | a) COI declaration: ‘The project was funded by an Investigator’s Initiated Research Grant from Lundbeck International given to Dr Liat Ayalon. Lundbeck International had no other involvement in the project concept of design or in this paper. Per Bech has occasionally over the past 3 years until August 2008 received funding from and has been speaker or member of advisory boards for pharmaceutical companies with an interest in the drug treatment of affective disorders (AstraZeneca, Lilly, H Lundbeck A/S, Lundbeck Foundation and Organon)’. b) Acknowledged funding from Lundbeck International | c) Not acknowledged |
| Study          | Sample characteristics | Sample size and % depressed | PHQ-9 characteristics | Diagnostic standard | a) COI declaration | b) Funding | Relationship with original developers |
|---------------|------------------------|-----------------------------|----------------------|---------------------|-------------------|------------|-------------------------------------|
| Eack et al    | Country: USA<br>Setting: community mental health centres for children<br>Age (years): male=39.20 (SD 9.63)<br>Female: 100%<br>n=50<br>Depressed: 28% | Administration: self-report<br>Language: English | DSM-IV SCID | a) No COI declaration<br>b) Funding acknowledged (academic/health research institutions) |
| Fann et al    | Country: USA<br>Setting: trauma hospital (npatients with traumatic brain injury)<br>Age (years): male=42 (SD=17.9)<br>Female: 29.1%<br>n=135<br>Depressed: 16.3% | Administration: telephone-administered<br>Language: English | DSM-IV SCID | a) No COI declaration<br>b) Funding acknowledged (academic institutions) |
| Gelaye et al  | Country: Ethiopia<br>Setting: general hospital<br>Age (years): 34.9 (SD=11.6)<br>Female: 63.1%<br>n=363<br>Depressed: 12.6% | Administration: researcher-administered<br>Language: Amharic | DSM-IV SCAN | a) No COI declaration<br>b) Funding acknowledged (academic/health research institutions) |
| Gjerdingen et al | Country: USA<br>Setting: community<br>Age (years): male=29.3<br>Female: 100%<br>n=438<br>Depressed: 4.6% | Administration: telephone or self-report<br>Language: English | DSM-IV SCID | a) No COI declaration<br>b) Funding acknowledged (academic/health research institutions) |
| Henkel et al  | Country: Germany<br>Setting: primary care<br>Age (years): not reported<br>Female: 74%<br>n=448<br>Depressed: 10% | Administration: self-report<br>Language: German | DSM-IV CIDI | a) No COI declaration<br>b) Funding acknowledged (academic/health research institutions) |
| Hyphantis et al | Country: Greece<br>Setting: hospital – rheumatology patients<br>Age (years): male=54.2 (SD=13.5)<br>Female: 74%<br>n=213<br>Depressed: 32.4% | Administration: researcher administered<br>Language: Greek | DSM-IV MINI | a) No COI declaration<br>b) No funding acknowledgement |
| Inagaki et al | Country: Japan<br>Setting: general hospital<br>Age whole sample (years): male=73.5 (SD=12.3)<br>Female: 59.3%<br>n=104 out of 511 received MINI<br>Depressed: 7.4% | Administration: researcher administered<br>Language: Japanese | DSM-IV MINI | a) COI declaration: ‘The authors declare that they have no competing interests’. b) Funding acknowledged (academic/health research institutions) |
| Khamseh et al | Country: Iran<br>Setting: diabetes clinic<br>Age (years): male=56.17 (SD=9.60)<br>Female: 51.9%<br>n=185<br>Depressed: 43.2% | Administration: self report<br>Language: Persian | DSM-IV SCID | a) COI declaration: the authors declared no competing interests<br>b) Funding acknowledged (academic/health research institutions) |
| Lamers et al  | Country: The Netherlands<br>Setting: primary care (elderly)<br>Age (years): male=71.4 (SD=6.90)<br>Female: 48.2%<br>n=713<br>Depressed: 10.7% | Administration: self report<br>Language: Dutch | DSM-IV MINI | a) No COI declaration<br>b) Funding acknowledged (academic/health research institutions) |
### Table 1 Continued

| Study                  | Sample characteristics                                                                 | Sample size and % depressed | PHQ-9 characteristics | Diagnostic standard | a) COI declaration               | b) Funding declaration          | c) Relationship with original developers |
|------------------------|----------------------------------------------------------------------------------------|------------------------------|-----------------------|---------------------|----------------------------------|-----------------------------------|----------------------------------------|
| Lotrakul et al<sup>46</sup> | Country: Thailand  
Setting: primary care  
Age (years): male=45.0 (SD=14.30) Female: 73.7% | n=279  
Depressed: 6.8% | Administration: self report  
Language: Thai | DSM-IV  
MINI | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| Persoons et al<sup>36</sup> | Country: Belgium  
Setting: hospital (otorhinolaryngology patients)  
Age (years): male=48.2 (SD=12.9) Female: 65.6% | n=268 (97 received MINI)  
Depressed: 16.5% | Administration: self report  
Language: Dutch | DSM-IV  
MINI | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) and Pfizer Belgium |                          |                         |
| Picardi et al<sup>36</sup> | Country: Italy  
Setting: hospital (dermatology inpatients)  
Age (years): male=37.5 Female: 56% | n=141  
Depressed: 8.5% | Administration: self report  
Language: Italian | DSM-IV  
SCID | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| Stafford et al<sup>37</sup> | Country: Australia  
Setting: hospital (cardiology patients)  
Age (years): male=64.1 (SD=10.3) Female: 66% | n=193  
Depressed: 18% | Administration: self report  
Language: English | DSM-IV  
MINI | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| Thombs et al<sup>38</sup> | Country: USA  
Setting: hospital (outpatients with coronary heart disease)  
Age (years): male=67 (SD=11) Female: 18% | n=1024  
Depressed: 22% | Administration: not stated  
Language: English | DSM  
C-DIS | a) COI declaration ‘None disclosed’  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| Thompson et al<sup>39</sup> | Country: USA  
Setting: patients with Parkinson’s disease  
Age (years): 72.5 (SD=9.6) Female: 42% | n=214  
Depressed: 14% | Administration: self administered  
Language: English | DSM-IV  
SCID | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| Turner et al<sup>40</sup> | Country: Australia  
Setting: stroke patients  
Age (years): 66.7 (SD=13.1) Female: 47.2% | n=72  
Depressed: 18% | Administration: self administered  
Language: English | DSM-IV  
SCID | a) COI declaration: disclosures ‘none’.  
b) Funding acknowledged (academic/health research institutions) |                          |                         |
| van Steenbergen-Weijenburg et al<sup>41</sup> | Country: The Netherlands  
Setting: patients with diabetes  
Age (years): male=61.8 (SD=13.6) Female: 48.7% | n=197  
Depressed: 18.8% | Administration: self administered  
Language: Dutch | DSM-IV  
SCID | a) COI declaration: ‘The authors declare that they have no competing interests’.  
b) Funding acknowledged (academic/health research institutions)—‘this had no influence on the content of this article’. |                          |                         |
| Zuitthoff et al<sup>47</sup> | Country: The Netherlands  
Setting: primary care  
Age (years): male=51 (SD=16.7) Female: 63% | n=1338  
Depressed: 13% | Administration: self report  
Language: Dutch | DSM-IV  
CIDI | a) COI declaration ‘The authors declare that they have no competing interests’.  
b) Funding acknowledged (academic/health research institutions) |                          |                         |

CIDI, Composite International Diagnostic Interview; CIS-R, Clinical Interview Schedule; COI, conflict of interest; DSM, Diagnostic and Statistical Manual of Mental Disorders; MINI, Mini-International Neuropsychiatric Interview; N/A, not available; SCAN, Schedules for Clinical Assessments in Neuropsychiatry; SCID, Structured Clinical Interview for DSM Disorders.
| Study | Sample characteristics | Sample size and % MDD | PHQ-9 characteristics | Diagnostic standard | a) COI declaration | b) Funding | c) Relationship with original developers |
|-------|------------------------|-----------------------|-----------------------|---------------------|-------------------|-----------|----------------------------------------|
| 13. Gräfe et al 23 | Country: Germany Setting: psychosomatic walk-in clinics and family practices Mean age: 41.9 (SD=13.8) Female: 67.8% | n=528 Depressed: 29.2% psychosomatic patients; 6.16% medical patients | Administration: self-report Language: German Cut-offs: 10–14 | DSM-IV SCID | a) No COI declaration | b) Acknowledged funding from Pfizer | c) Not acknowledged |
| 16. Kroenke et al 20 | Country: USA Setting: primary care Mean age: 46 (SD=17) Female: 66% | n=580 7.1% MDD | Administration: self-report Language: English Cut-offs: 9–15 | DSM-IV SCID | a) No COI declaration | b) Acknowledged funding from Pfizer | c) N/A |
| 22. Navinés et al 28 | Country: Spain Setting: general hospital (patients with chronic HCV) Mean age: 43.4 (SD=10.2) Female: 28.6% | n=500 6.4% MDD | Administration: self-report Language: Spanish Cut-offs: 10 | DSM-IV SCID | a) All authors declared that they had no COI | b) Role of funding source declared | c) Not acknowledged |
| 29. Thakkumpurath et al 26 | Country: UK Setting: hospital (cancer patients) Mean age: 61 Female: 63% | n=782 6.3% MDD (of the whole sample) | Administration: not stated Language: English Cut-offs: 5–10 | DSM-IV SCID | a) COI declaration: ‘Supported by Cancer Research UK’ | b) As in a) | c) Not acknowledged |
| 33. Williams et al 49 | Country: USA Setting: secondary care (poststroke) Mean age: unclear Female: unclear | n=316 33.5% MDD | Administration: unclear Language: English Cut-offs: 10 | DSM-IV SCID | a) No COI declaration | b) Funding acknowledged (academic institutions) | c) Not acknowledged |
| 1. Adewuya et al 55 | Country: Nigeria Setting: community (students) Mean age: 24.8 (15–40) Female: 41.2% | n=512 2.5% MDD | Administration: Self-report Language: English Cut-offs: 8–12 | DSM-IV MINI | a) No COI declaration | b) No funding declaration |
| 2. Arroll et al 42 | Country: New Zealand Setting: primary care Mean age: 49 (17–99) Female: 61% | n=2642 6.2% MDD | Administration: not stated Language: English Cut-offs: 8, 10, 12, 15 | DSM-IV SCID | a) No COI declaration | b) Funding acknowledged (academic/health research institutions) |
| 3. Azah et al 62 | Country: Malaysia Setting: primary care Mean age: 38.7 (18–79) Female: 61.7% | n=180 16.6% MDD | Administration: self-report Language: Malay Cut-offs: 5–12 | DSM-IV CIDI | b) No COI declaration | c) Funding acknowledged (academic/health research institutions) |
| 4. Chagas et al 50 | Country: Brazil Setting: secondary care Mean age: not stated Female: 52.7% | n=84 25.5% MDD | Administration: self-report Language: Brazilian Cut-offs: 7–10 | DSM-IV SCID | a) COI declaration ‘None declared’ | b) Funding acknowledged (academic/health research institutions) |

Continued
| Study          | Sample characteristics                                      | Sample size and % MDD | PHQ-9 characteristics                | Diagnostic standard | a) COI declaration | b) Funding | c) Relationship with original developers |
|---------------|-------------------------------------------------------------|-----------------------|--------------------------------------|---------------------|--------------------|------------|----------------------------------------|
| 6. de Lima Osorio et al[60] | Country: Brazil  
Setting: primary care  
Mean age: unclear  
Female: 100% | n=177  
34% MDD | Administration: research assistants  
Language: Brazilian Portuguese  
Cut-offs: 10-15 | DSM-IV SCID | a) No COI declaration  
b) Funding acknowledged (academic institutions) |
| 7. Elderon et al[61] | Country: USA  
Setting: secondary care  
Mean age: unclear  
Female: 18% | n=1022  
18.3% MDD | Administration: self-report  
Language: English  
Cut-offs: 10 | C-DIS | a) COI declaration—‘No disclosures’  
b) Funding acknowledged (academic institutions and industry—AHA Pharmaceuticals Roundtable)—‘The funding organisations had no role in the design or conduct of the study, collection, management, analysis or interpretation of data; or preparation, review or approval of the manuscript’. |
| 8. Fann et al[60] | Country: USA  
Setting: trauma hospital (inpatients with traumatic brain injury)  
Mean age: 42 (SD=17.9)  
Female: 29.1% | n=135  
16.3% MDD | Administration: telephone-administered  
Language: English  
Cut-offs: 10 | DSM-IV SCID | a) No COI declaration  
b) Funding acknowledged (academic institutions) |
| 9. Fine et al[64] | Country: USA  
Setting: primary care (Ohio Army National Guard)  
Mean age: 31 (17-60)  
Female: 29.1% | n=498  
21.5% MDD | Administration: telephone-administered  
Language: English  
Cut-offs: 10, 15 | DSM-IV SCID-I | a) COI—last author disclosed financial and consulting interests (Pfizer not one of them). All other authors declared that they have no COI.  
b) Funding acknowledged—DoD Medical Research. ‘The sponsor had no role in study design, data collection, analysis, interpretation of results, report writing or manuscript submission’. |
| 10. Gelaye et al[61] | Country: Ethiopia  
Setting: general hospital  
Mean age: 34.9 (SD=11.6)  
Female: 63.1% | n=363  
12.6% MDD | Administration: researcher-administered  
Language: Amharic  
Cut-offs: 9-11 | DSM-IV SCAN | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |
| 11. Gilbody et al[67] | Country: UK  
Setting: primary care  
Mean age: 42.5 (SD 13.6)  
Female: 77% | n=96  
37.5 MDD | Administration: not stated  
Language: English  
Cut-offs: 9-13 | DSM-IV SCID | a) COI declaration—last author involved in the development of one of the instruments (CORE-OM), ‘but does not gain financially from its use.  
b) Funding acknowledged (academic/health research institutions) |
| 12. Gjerdingen et al[64] | Country: USA  
Setting: community  
Mean age: 29.3  
Female: 100% | n=438  
4.8% MDD | Administration: telephone or self-report  
Language: English  
Cut-offs: 10 | DSM-IV SCID | c) No COI declaration  
d) Funding acknowledged (academic/health research institutions) |
| 14. Hyphantis et al[62] | Country: Greece  
Setting: hospital—rheumatology patients  
Mean age: 54.2 (SD=13.5)  
Female: 74% | n=213  
32.4% MDD | Administration: researcher administered  
Language: Greek  
Cut-offs: 4-16 | DSM-IV MINI | a) No COI declaration  
b) No funding acknowledgement |

Continued
| Study          | Sample characteristics                  | Sample size and % MDD | PHQ-9 characteristics         | Diagnostic standard | a) COI declaration                                      | b) Funding                      | c) Relationship with original developers |
|---------------|-----------------------------------------|-----------------------|--------------------------------|---------------------|--------------------------------------------------------|----------------------------------|------------------------------------------|
| 15. Khamseh et al<sup>34</sup> | Country: Iran  
Setting: outpatient diabetic clinic  
Mean age: 56.1 (SD=9.6)  
Female: 51.8% | n=185  
43.2% MDD | Administration: self-report  
Language: Persian  
Cut-offs: 10, 13 | DSM-IV SCID | a) COI declaration: the authors declared no competing interests.  
d) Funding acknowledged (academic/health research institutions) |                       |                          |
| 19. Liu et al<sup>53</sup> | Country: Taiwan  
Setting: primary care  
Mean age: not specified  
Female: 60.9% | n=1532  
3.3% MDD | Administration: self-report  
Language: Chinese version  
Cut-offs: 9–11 | SCAN | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 20. Lotrakul et al<sup>46</sup> | Country: Thailand  
Setting: primary care  
Mean age: 45.0 (SD=14.30)  
Female: 73.7% | n=279  
6.8% MDD | Administration: self report  
Language: Thai  
Cut-offs: 7–15 | DSM-IV MINI | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 23. Patel et al<sup>61</sup> | Country: India  
Setting: primary care  
Mean age: 37.5 (18–83)  
Female: 56.4% | n=299  
4.3% MDD | Administration: face-to-face interview  
Language: not specified  
Cut-offs: 7–15 | CIS-R | a) COI declaration— No declaration of Interest  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 24. Phelan et al<sup>58</sup> | Country: USA  
Setting: primary care (elderly)  
Mean age: 78 (SD=7)  
Female: 62% | n=71  
12% MDD | Administration: research assistant  
Language: English  
Cut-offs: 8–12 | DSM-IV SCID | a) COI declaration— no competing interests  
b) Funding acknowledged (academic/health research institutions)  
'b) The funder had no role in the study design, methods, data collection, analysis or interpretation of data, nor any role in the preparation of the manuscript or decision to submit the manuscript for publication'. |                       |                          |
| 25. Rooney et al<sup>52</sup> | Country: UK  
Setting: secondary care (glioma)  
Mean age: 54.2 (SD=12.3)  
Female: 42.6% | n=129  
13.5% MDD | Administration: self-report  
Language: English  
Cut-offs: 8–11 | DSM-IV SCID | a) COI declaration ‘The authors declare that they have no COI’.  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 26. Sherina et al<sup>41</sup> | Country: Malaysia  
Setting: primary care  
Mean age: 30.9 (18–81)  
Female: 100% | n=146  
21.2% MDD | Administration: self-report  
Language: Malay  
Cut-offs: 10 | CIDI | a) COI declaration ‘The authors declare that they have no competing interests’.  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 27. Sidebottom et al<sup>59</sup> | Country: USA  
Setting: community (prenatal)  
Mean age: 23 (SD=5.5)  
Female: 100% | n=745  
3.6% MDD | Administration: interview  
Language: English  
Cut-offs: 10 | DSM-IV SCID | b) COI declaration ‘The authors declare that they have no financial COI’.  
b) Funding acknowledged (academic/health research institutions) |                       |                          |
| 28. Stafford et al<sup>37</sup> | Country: Australia  
Setting: secondary care (cardiac procedures)  
Mean age: 64.14 (38–91)  
Female: 19.2% | n=193  
18.1% MDD | Administration: self-report  
Language: English  
Cut-offs: 10 | DSM-IV MINI | a) No COI declaration  
b) Funding acknowledged (academic/health research institutions) |                       |                          |

Continued
### Table 2  Continued

| Study       | Sample characteristics                                      | Sample size and % MDD | PHQ-9 characteristics                  | Diagnostic standard | a) COI declaration                          | b) Funding                                           | c) Relationship with original developers |
|-------------|-------------------------------------------------------------|-----------------------|----------------------------------------|---------------------|--------------------------------------------|------------------------------------------------------|------------------------------------------|
| 30. Thombs et al<sup>39</sup> | Country: USA  
Setting: hospital (outpatients with coronary heart disease)  
Mean age: 67 (SD=11)  
Female: 18% | n=1024  
22% MDD | Administration: not stated  
Language: English  
Cut-offs: 7–10 | DSM  
C-DIS | a) COI declaration ‘None disclosed’ | b) Funding acknowledged (academic/health research institutions) |                          |
| 32. Watnick et al<sup>31</sup> | Country: USA  
Setting: secondary care (dialysis)  
Mean age: 63 (SD=15)  
Female: 32.3% | n=62  
19% MDD | Administration: self-report  
Language: English  
Cut-offs: 10 | DSM-IV  
SCID | a) No COI declaration | b) Funding acknowledged (academic/health research institutions) |                          |
| 34. Wittkampf et al<sup>42</sup> | Country: The Netherlands  
Setting: primary care  
Mean age: 49.8  
Female: 66.7% | n=664  
12.3% MDD | Administration: self-report  
Language: not specified  
Cut-offs: 10 and 15 | DSM-IV  
SCIDI | a) No COI declaration | b) Funding acknowledged (academic/health research institutions) |                          |
| 35. Zhang et al<sup>32</sup> | Country: Hong Kong  
Setting: secondary care (diabetic outpatients)  
Mean age: 55.1 (SD=9.5)  
Female: 40.8% | n=99  
23.2% MDD | Administration: self-report  
Language: Chinese version  
Cut-offs: 15 | DSM-IV MINI  
CIDI | a) COI declaration—last author acknowledged financial COI. The other authors declare that they have no competing interests. | b) Funding acknowledged (academic/health research institutions) |                          |
| 36. Zuithoff et al<sup>47</sup> | Country: The Netherlands  
Setting: primary care  
Age (years): male=51 (SD=16.7)  
Female: 63% | n=1338  
Depressed: 13% | Administration: self-report  
Language: Dutch | DSM-IV  
CIDI | a) COI declaration ‘The authors declare that they have no competing interests’. | b) Funding acknowledged (academic/health research institutions) |                          |

COI, conflict of interest; DSM, Diagnostic and Statistical Manual of Mental Disorders; MDD, major depressive disorder; N/A, not available; SCID, Structured Clinical Interview for DSM Disorders.
| Study                          | Patient selection: | Patient selection: | Patient selection: | Index test: | Index test: | Index test: | Index test: | Overall risk of bias |
|-------------------------------|--------------------|--------------------|--------------------|-------------|-------------|-------------|-------------|----------------------|
| Allegrant studies             |                    |                    |                    |             |             |             |             |                      |
| Diez-Quevedo et al<sup>22</sup> | ✓                  | ✓                  | ✓                  |             | ?           | ✓           | ✓           | Unclear              |
| Graße et al<sup>23</sup>      | ✓                  | ✓                  | ✓                  |             | ?           | ✓           | ✓           | Unclear              |
| Lowe et al<sup>24</sup>       | ✓                  | ✓                  | ✓                  |             | ✓           | ✓           | ✓           | Low                  |
| Muramatsu et al<sup>27</sup>  | ?                  | ✓                  | ?                  |             | Unclear     | ✓           | ✓           | ?                    |
| Navines et al<sup>28</sup>    | ✓                  | ✓                  | ✓                  |             | ✓           | ✓           | ✓           | Unclear              |
| Spitzer et al<sup>25</sup>    | ✓                  | ✓                  | ✓                  |             | ✓           | ✓           | ✓           | Low                  |
| Thekkumpurath et al<sup>36</sup> | ✓                  | ✓                  | ✓                  |             | ✓           | N/A         | N/A         | Low                  |
| Non-allegant studies          |                    |                    |                    |             |             |             |             |                      |
| Arroll et al<sup>42</sup>     | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | N/A         | N/A                  |
| Ayalon et al<sup>43</sup>     | ?                  | ✓                  | ✓                  |             | Unclear     | ✓           | ?           | Unclear              |
| Eck et al<sup>49</sup>        | ?                  | ✓                  | ?                  |             | Unclear     | ?           | N/A         | N/A                  |
| Fann et al<sup>50</sup>       | ✓                  | ✓                  | ✓                  |             | High        | ✓           | N/A         | N/A                  |
| Gelaye et al<sup>51</sup>     | ?                  | ✓                  | ?                  |             | High        | ✓           | ✓           | ?                    |
| Gjerdingen et al<sup>48</sup> | ✓                  | ✓                  | ✓                  |             | Low         | ?           | N/A         | N/A                  |
| Henkel et al<sup>44</sup>     | ✓                  | ✓                  | ✓                  |             | Low         | ?           | N/A         | N/A                  |
| Hyphantis et al<sup>32</sup>  | ✓                  | ✓                  | ✓                  |             | High        | ✓           | ?           | Unclear              |
| Inagaki et al<sup>33</sup>    | ✓                  | ✓                  | ✓                  |             | High        | ✓           | ?           | ?                    |
| Khamseh et al<sup>34</sup>    | ✓                  | ✓                  | ?                  |             | Unclear     | ✓           | ✓           | ?                    |
| Lamers et al<sup>45</sup>     | ✓                  | ✓                  | ✓                  |             | High        | ✓           | ?           | ?                    |
| Lotrakul et al<sup>46</sup>   | ✓                  | ✓                  | ✓                  |             | High        | ✓           | ✓           | ?                    |
| Persoons et al<sup>35</sup>   | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | ✓           | N/A                  |
| Picardi et al<sup>56</sup>    | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | ?           | Unclear              |
| Stafford et al<sup>57</sup>   | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | N/A         | N/A                  |
| Thombs et al<sup>58</sup>     | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | N/A         | N/A                  |
| Thomson et al<sup>59</sup>    | ?                  | ✓                  | ✓                  |             | Unclear     | ?           | N/A         | N/A                  |
| Turner et al<sup>60</sup>     | ✓                  | ✓                  | ✓                  |             | Low         | ✓           | N/A         | N/A                  |

Continued...
| Study                      | Patient selection: | Avoid case-control/avoid artificially inflated base rate | Avoided inappropriate exclusions | Overall risk of bias | PHQ-9 interpreted blind to reference test | If translated, appropriate translation | If translated, psychometric properties reported | Overall risk of bias |
|---------------------------|--------------------|--------------------------------------------------------|----------------------------------|----------------------|------------------------------------------|----------------------------------------|---------------------------------------------|---------------------|
| van Steenbergen-Wijenburg et al | ?                  | ✓                                                      | ✓                                | Unclear              | ?                                         | ?                                      | ?                                           | Unclear             |
| Zuihoff et al             | ✓                  | ✓                                                      | ✓                                | Low                  | ✓                                        | ✓                                      | ✓                                           | Unclear             |
| Reference test            | Reference test     | Reference test                                        | Reference test                   | Reference test       | Flow/timing                               | Flow/timing                           | Flow/timing                                           | Overall risk of bias |
| Study                     | Reference test     | Reference test                                        | Reference test                   | Reference test       | Flow/timing                               | Flow/timing                           | Flow/timing                                           | Overall risk of bias |
| Allegiant studies         |                    |                                                        |                                  |                      |                                          |                                        |                                             |                     |
| Diez-Quevedo et al        | ✓                  | ✓                                                      | ✓                                | ?                    | Unclear                                  | ✓                                      | ✓                                           | Low                 |
| Gräfe et al               | ✓                  | ?                                                      | N/A                              | N/A                  | Unclear                                  | ✓                                      | ✓                                           | Low                 |
| Lowe et al                | ✓                  | ✓                                                      | N/A                              | N/A                  | Low                                      | ✓                                      | ✓                                           | Low                 |
| Muramatsu et al           | ✓                  | ✓                                                      | ✓                                | Low                  | ✓                                        | ✓                                      | ✓                                           | Low                 |
| Navines et al             | ✓                  | ✓                                                      | ?                                | Unclear              | ✓                                        | ✓                                      | ✓                                           | Low                 |
| Spitzer et al             | ✓                  | ✓                                                      | N/A                              | N/A                  | Low                                      | ✓                                      | x                                           | High                |
| Thekkumpurath et al       | ✓                  | ✓                                                      | N/A                              | N/A                  | Low                                      | ?                                      | ✓                                           | x                   |
| Non-allegiant studies     |                    |                                                        |                                  |                      |                                          |                                        |                                             |                     |
| Arroll et al              | ✓                  | ✓                                                      | N/A                              | N/A                  | Low                                      | ✓                                      | ✓                                           | Low                 |
| Ayalon et al              | ✓                  | ?                                                      | ✓                                | ?                    | Unclear                                  | ✓                                      | ✓                                           | Unclear             |
| Eack et al                | ✓                  | ?                                                      | N/A                              | N/A                  | Unclear                                  | ✓                                      | ✓                                           | Unclear             |
| Fann et al                | ✓                  | ?                                                      | N/A                              | N/A                  | Unclear                                  | ✓                                      | ✓                                           | Unclear             |
| Gelaye et al              | ✓                  | ✓                                                      | ✓                                | Low                  | ✓                                        | ✓                                      | x                                           | High                |
| Gjerdingen et al          | ✓                  | ?                                                      | N/A                              | N/A                  | Unclear                                  | ✓                                      | x                                           | High                |
| Henkel et al              | ✓                  | ?                                                      | N/A                              | N/A                  | Unclear                                  | ✓                                      | x                                           | High                |
| Hyphantis et al           | ✓                  | ✓                                                      | ?                                | ?                    | Unclear                                  | ✓                                      | ✓                                           | x                   |
| Inagaki et al             | ✓                  | ✓                                                      | ?                                | Unclear              | ✓                                        | ✓                                      | x                                           | x                   |
| Khamseh et al             | ✓                  | ✓                                                      | ✗                                | Unclear              | ✓                                        | ✓                                      | ?                                           | Unclear             |
| Lamers et al              | ✓                  | ✓                                                      | ?                                | Unclear              | ✓                                        | ✓                                      | x                                           | High                |

Table 3 Continued
| Study                        | Reference test correctly classifies target condition | Reference test interpreted blind to PHQ-9 | If translated, if appropriate | Psychometric properties reported | Overall risk of bias | Interval of 2 weeks or less | All participants receive same reference test | All participants included in analysis? | Overall risk of bias |
|-----------------------------|-----------------------------------------------------|------------------------------------------|-------------------------------|---------------------------------|----------------------|-----------------------------|------------------------------------------|--------------------------------------|---------------------|
| Lotrakul et al<sup>46</sup> | ✓                                                   | ✓                                        | ✓                             | ✓                               | Low                  | ?                           | ✓                                        | ✗                                    | High                 |
| Persoons et al<sup>35</sup> | ✓                                                   | ✓                                        | ?                             | ?                               | Unclear              | ✓                           | ✓                                        | ✓                                    | Low                  |
| Picardi et al<sup>46</sup>  | ✓                                                   | ✓                                        | ✓                             | ?                               | Unclear              | ✓                           | ✓                                        | ✓                                    | High                 |
| Stafford et al<sup>47</sup> | ✓                                                   | ✓                                        | N/A                           | N/A                             | Low                  | ✓                           | ✓                                        | ✓                                    | High                 |
| Thombs et al<sup>38</sup>   | ?                                                   | ✓                                        | N/A                           | N/A                             | Unclear              | ✓                           | ✓                                        | ✓                                    | Low                  |
| Thompson et al<sup>39</sup> | ✓                                                   | ?                                        | N/A                           | N/A                             | Unclear              | ✓                           | ✓                                        | ✓                                    | High                 |
| Turner et al<sup>40</sup>   | ✓                                                   | ?                                        | N/A                           | N/A                             | Unclear              | ?                           | ✓                                        | ✓                                    | High                 |
| van Steenbergen-Wijenburg et al<sup>41</sup> | ✓                                                   | ✗                                        | ?                             | ?                               | High                 | ✓                           | ✓                                        | ✗                                    | High                 |
| Zuihoff et al<sup>47</sup>  | ✓                                                   | ✓                                        | ?                             | ?                               | Unclear              | ?                           | ✓                                        | ✓                                    | Unclear              |

✓, criterion met; ✗, criterion not met; ?, insufficient information to code whether criterion met; N/A, not applicable; PHQ-9, Patient Health Questionnaire-9.
## Table 4  Quality assessment of included studies in the summed item scoring method cut-off point 10 meta-analysis^{14}

| Study                | Patient selection: Consecutive or random sample | Avoid case-control/avoid artificially inflated base rate | Avoided inappropriate exclusions | Overall risk of bias | PHQ-9 interpreted blind to reference test | Was a threshold prespecified? | If translated, appropriate translation | If translated, psychometric properties reported | Overall risk of bias |
|----------------------|-------------------------------------------------|-------------------------------------------------------|----------------------------------|----------------------|------------------------------------------|-------------------------------|--------------------------------------|---------------------------------------------|---------------------|
| **Legitimate studies** |                                                 |                                                       |                                  |                      |                                          |                               |                                      |                              |                     |
| 13. Gräfe et al^{23} | ✓                                               | ✓                                                     | ✓                                | Low                  | ?                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 16. Kroenke et al^{12} | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 22. Navinés et al^{28} | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 29. Thekkumprath et al^{29} | x                                               | x                                                     | ✓                                | High                 | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 33. Williams et al^{49} | ✓                                               | ✓                                                     | ✓                                | Low                  | ?                                        | ?                             | N/A                                 | N/A                          | Unclear             |
| **Non-legitimate studies** |                                                 |                                                       |                                  |                      |                                          |                               |                                      |                              |                     |
| 1. Adewuya et al^{55} | ✓                                               | ✓                                                     | x                                | Unclear              | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 2. Arroll et al^{42}  | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 3. Azah et al^{52}    | ✓                                               | ✓                                                     | ?                                | High                 | ✓                                        | ✓                             | ✓                                   | N/A                          | Low                 |
| 4. Chagas et al^{50}  | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | ✓                                   | ✓                            | Low                 |
| 6. de Lima Osorio et al^{60} | ✓                                               | x                                                     | ✓                                | High                 | ?                                        | x                             | N/A                                 | N/A                          | High                |
| 7. Elderon et al^{53} | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 8. Fann et al^{50}    | ✓                                               | x                                                     | x                                | High                 | ✓                                        | ✓                             | ✓                                   | N/A                          | Low                 |
| 9. Fine et al^{64}    | ✓                                               | ✓                                                     | ✓                                | Low                  | ?                                        | ✓                             | N/A                                 | N/A                          | Unclear             |
| 10. Gelaye et al^{51} | ?                                               | x                                                     | ?                                | High                 | ✓                                        | ×                             | ✓                                   | ?                            | High                |
| 11. Gilbody et al^{57} | ?                                               | ✓                                                     | ?                                | Unclear              | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 12. Gjerdingen et al^{48} | ✓                                               | ✓                                                     | ✓                                | Low                  | ?                                        | ✓                             | N/A                                 | N/A                          | Unclear             |
| 14. Hyphantis et al^{12} | ✓                                               | x                                                     | ✓                                | High                 | ✓                                        | ✓                             | ?                                   | ?                            | Unclear             |
| 15. Khamseh et al^{24} | ✓                                               | ✓                                                     | ✓                                | Unclear              | ✓                                        | ✓                             | ?                                   | Unclear                      |                     |
| 19. Liu et al^{63}   | ✓                                               | ✓                                                     | ?                                | Unclear              | ✓                                        | ✓                             | ?                                   | ?                            | High                |
| 20. Lotrákul et al^{46} | x                                               | ✓                                                     | ?                                | Unclear              | ✓                                        | ✓                             | ?                                   | Unclear                      |                     |
| 23. Patel et al^{51}  | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | ?                                   | ?                            | Unclear             |
| 24. Phelan et al^{25} | x                                               | ✓                                                     | ✓                                | High                 | ✓                                        | x                             | N/A                                 | N/A                          | High                |
| 25. Rooney et al^{52} | ✓                                               | ✓                                                     | ✓                                | Low                  | ?                                        | x                             | N/A                                 | N/A                          | High                |
| 26. Sherina et al     | ✓                                               | ✓                                                     | x                                | High                 | ✓                                        | ✓                             | ✓                                   | ✓                            | Low                 |
| 27. Sidebottom et al^{69} | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |
| 28. Stafford et al^{17} | ✓                                               | ✓                                                     | ✓                                | Low                  | ✓                                        | ✓                             | N/A                                 | N/A                          | Low                 |

Continued
### Table 4 Continued

| Study | Consecutive or random sample | Avoid case-control/avoid artificially inflated base rate | Avoided inappropriate exclusions | Overall risk of bias | PHQ-9 interpreted blind to reference test | Was a threshold prespecified? | If translated, appropriate translation | Overall risk of bias |
|-------|-------------------------------|----------------------------------------------------------|-----------------------------------|----------------------|------------------------------------------|-------------------------------|----------------------------------------|---------------------|
| 30. Thombs et al<sup>38</sup> | ×                              | ✓                                                        | ?                                 | High                 | ✓ √                                      | ?                            | N/A N/A                                | Unclear             |
| 32. Watnick et al<sup>43</sup> | ?                              | ×                                                        | ✓                                 | High                 | ✓ ✓ ✓                                    | ✓                            | N/A N/A                                | Low                 |
| 34. Wittkamp et al<sup>44</sup> | ✓                              | ✓                                                        | ✓                                 | Low                  | ✓ ?                                      | ?                            | N/A N/A                                | Unclear             |
| 35. Zhang et al<sup>45</sup> | ✓                              | ✓                                                        | ?                                 | Unclear              | ? ✓                                      | ?                            | ? ?                                   | Unclear             |
| 36. Zuithoff et al<sup>47</sup> | ✓                              | ✓                                                        | ✓                                 | Low                  | ✓ ✓ ✓                                    | ✓                            | ✓ ?                                   | Unclear             |

| Reference test: | Reference test: | Reference test: | Reference test: | Reference test: | Flow / timing: | Flow / timing: | Flow / timing: | Flow / timing: | Overall risk of bias |
|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|---------------------|
| Reference test correctly classifies target condition | Reference test interpreted blind to PHQ-9 | If translated, appropriate translation | If translated, psychometric properties reported | Overall risk of bias | Interval of 2 weeks or less | All participants receive same reference test | All participants included in analysis? | Overall risk of bias |
| Allegiant studies | | | | | | | | | |
| 13. Gräfe et al<sup>33</sup> | ✓ | ? | N/A | N/A | Unclear | ✓ ✓ ✓ | Low |
| 16. Kroenke et al<sup>12</sup> | ✓ | ✓ | N/A | N/A | Low | ✓ ✓ ✓ | Low |
| 22. Navinès et al<sup>48</sup> | ✓ | ✓ | ? | ? | Unclear | ✓ ✓ ✓ | Low |
| 29. Thekkumpurath et al<sup>36</sup> | ✓ | ✓ | N/A | N/A | Low | ? | ✓ | Unclear |
| 33. Williams et al<sup>19</sup> | ✓ | ? | N/A | N/A | Unclear | ? | ✓ | Unclear |
| Non-allegiant studies | | | | | | | | | |
| 1. Adewuya et al<sup>55</sup> | ✓ | ✓ | N/A | N/A | Low | ✓ ✓ ✓ | Low |
| 2. Arroll et al<sup>12</sup> | ✓ | ✓ | N/A | N/A | Low | ? | ✓ | Unclear |
| 3. Azah et al<sup>40</sup> | ✓ | ✓ | ✓ | ✓ | Low | ✓ ✓ ✓ | High |
| 4. Chagas et al<sup>50</sup> | ✓ | ✓ | ? | ? | Unclear | ✓ ✓ ✓ | × | High |
| 6. de Lima Osorio et al<sup>60</sup> | ✓ | ? | N/A | N/A | Unclear | ? | ✓ | Unclear |
| 7. Elderon et al<sup>51</sup> | ✓ | ✓ | N/A | N/A | Low | ✓ ✓ ✓ | Low |
| 8. Fann et al<sup>60</sup> | ✓ | ? | N/A | N/A | Unclear | ✓ ✓ ✓ | × | High |
| 9. Fine et al<sup>16</sup> | ✓ | ? | N/A | N/A | Unclear | ? | ✓ | Unclear |
| 10. Gelaye et al<sup>51</sup> | ✓ | ✓ | ✓ | ✓ | Low | ✓ ✓ ✓ | High |
| 11. Gilbody et al<sup>57</sup> | ✓ | ✓ | N/A | N/A | Low | ? | ✓ | Unclear |
| 12. Gjerdingen et al<sup>48</sup> | ✓ | ? | N/A | N/A | Unclear | ✓ | ✓ | High |

Continued
| Study                  | Reference test correctly classifies target condition | Reference test interpreted blind to PHQ-9 | If translated, appropriate translation | If translated, psychometric properties reported | Overall risk of bias | Interval of 2 weeks or less | All participants receive same reference test | All participants included in analysis? | Overall risk of bias |
|-----------------------|------------------------------------------------------|------------------------------------------|----------------------------------------|-------------------------------------------------|---------------------|------------------------------------------|----------------------------------------|----------------------------------------|---------------------|
| 14. Hyphantis et al   | ✓                                                    | ✓                                        | ?                                      | ?                                               | Unclear             | ✓                                        | ✓                                      | ?                                      | Unclear             |
| 15. Khamseh et al     | ✓                                                    | ✓                                        | ✓                                      | ?                                               | Unclear             | ✓                                        | ✓                                      | ?                                      | Unclear             |
| 19. Liu et al         | ✓                                                    | ✓                                        | ✓                                      | ✓                                               | Low                 | ✓                                        | ✓                                      | ?                                      | Unclear             |
| 20. Lotrakul et al    | ✓                                                    | ✓                                        | ✓                                      | ✓                                               | Low                 | ✓                                        | ✓                                      | ?                                      | Unclear             |
| 23. Patel et al       | ✓                                                    | ✓                                        | ✓                                      | ?                                               | Unclear             | ✓                                        | ?                                      | ✓                                      | High                |
| 24. Phelan et al      | ✓                                                    | ✓                                        | N/A                                    | N/A                                             | Low                 | ✓                                        | ✓                                      | ✓                                      | Low                 |
| 25. Rooney et al      | ✓                                                    | ?                                        | N/A                                    | N/A                                             | Unclear             | ✓                                        | ?                                      | ✓                                      | High                |
| 26. Sherina et al     | ✓                                                    | ✓                                        | ✓                                      | ✓                                               | Low                 | ✓                                        | ✓                                      | ✓                                      | Low                 |
| 27. Sidebottom et al  | ✓                                                    | ✓                                        | N/A                                    | N/A                                             | Low                 | ✓                                        | ✓                                      | ✓                                      | High                |
| 28. Stafford et al    | ✓                                                    | ✓                                        | N/A                                    | N/A                                             | Low                 | ✓                                        | ✓                                      | ✓                                      | High                |
| 30. Thoms et al       | ?                                                    | ✓                                        | N/A                                    | N/A                                             | Unclear             | ✓                                        | ✓                                      | ✓                                      | Low                 |
| 32. Watnick et al     | ✓                                                    | ✓                                        | N/A                                    | N/A                                             | Low                 | ✓                                        | ✓                                      | ✓                                      | Low                 |
| 34. Wittkampf et al   | ✓                                                    | ✓                                        | N/A                                    | N/A                                             | Low                 | ✓                                        | ✓                                      | ?                                      | High                |
| 35. Zhang et al       | ✓                                                    | ?                                        | ✓                                      | ✓                                               | Unclear             | ✓                                        | ?                                      | ✓                                      | High                |
| 36. Zuihof et al      | ✓                                                    | ✓                                        | ?                                      | ✓                                               | Unclear             | ✓                                        | ✓                                      | ?                                      | Unclear             |

N/A, not applicable; PHQ-9, Patient Health Questionnaire-9. ✓, criterion met; X, criterion not met; ?, insufficient information to code whether criterion met.
case-control design, avoid inappropriate exclusions). In
the index test domain, the proportion of studies reporting
that the PHQ-9 was conducted blind to the reference
test was comparable between the two groups. There
were differences in this domain for those studies using
a translated version of the test. All non-English allegiant
studies (5/5) used an appropriately translated version
of the PHQ-9, whereas just over a half of the non-allegiant
studies reported this (55%, 6/11). However, the majority
of both sets of studies did not report details of psycho-
metric properties of the translated version. For the refer-
ence test domain, nearly all studies in both groups were
rated as using a reference test that would correctly clas-
sify the condition. While most allegiant studies reported
that the reference test was interpreted blind to the PHQ-9
score (86%, 6/7), this was reported in only 60% (12/20)
of the non-allegiant studies.

The two sets of studies that used translated versions
of the reference test were broadly comparable. There
was a slight indication that the allegiant studies were
more likely to use an appropriately translated version
of the reference test and report data on the psycho-
metric properties of the translated version, although
the numbers for the translated comparison are very
low. There were, however, some more notable differ-
ences on the flow and timing domain. Most allegiant
studies ensured that the time between the index and
reference test was under 2 weeks (86%, 6/7) in compar-
ison to 70% (14/20) of the non-allegiant studies. More
allegiant studies met the criterion for ‘all participants
included in the analysis’ (57%, 4/7) than non-allegiant
studies (25%).

Descriptive characteristics
Table 2 presents the sample characteristics of the 31
PHQ-9 validation studies that reported the psychometric
properties of the PHQ-9 at cut-off point 10 or above.
Five of these studies were coauthored by the original
developers of the instrument or acknowledged collabo-
ration12 23 26 49 or were coauthored by the first author of
a previous study that had also been coauthored by one
of the developers.28 Twenty-six studies were conducted by
independent researchers.

Three (60%, 3/5) allegiant studies26 28 49 and 11
non-allegiant studies (42%, 11/26)30–32 34 37 38 50–54
were conducted in hospital settings.

Three (60%, 3/5) allegiant studies12 26 49 and 13 non-
allegiant studies (13/26)30 37 38 42 48 51–53 55–59
were conducted in English.

The mean prevalence of MDD in the allegiant group
was 13.2% (range 6.1%–33.5%) and in the non-alle-
giant group was 16.1% (range 2.5%–43.2%). The mean
age of patients in the allegiant group studies was 48.1
(range 41.9–61.0) and in the 26 non-allegiant studies
that reported these data was 49.1 (range 23.0–78.0).
The percentage of females in the allegiant studies that
reported these data12 23 26 28 was 56.3% (range 28.6%–
67.8%) and in the non-allegiant group was 64.9% (range
12%–100%).

Three allegiant studies used the self-reported mode
of administration and two of them did not specify how
the PHQ-9 was administered. In nine non-allegiant
studies (34%, 9/26), the PHQ-9 was administered by the
researcher.30–32 48 58 61 All allegiant studies used SCID as
a gold standard; the non-allegiant studies used a wider range of gold standards including SCAN, CIDI, MINI, CIS-R, C-DIS, although the SCID was used in half of the studies (50%, 15/26 studies).

Three allegiant studies (60%) did not include a conflict of interest statement. Two of these studies acknowledged funding from Pfizer. None of the allegiant studies acknowledged collaboration or authorship of one of the developers of the PHQ-9.

Of the non-allegiant studies, 13 (42%) did not include a conflict of interest statement. Similar to the algorithm studies, the newer studies were more likely to include a conflict of interest statement. Funding was acknowledged by most studies (27/31) and most received funding from academic and/or health research institutions. One study acknowledged that the last author involved in the development of one of the instruments (CORE-OM), ‘but does not gain financially from its use’. One study acknowledged funding from industry, AHA Pharmaceuticals Roundtable, but stated that ‘the funding organisations had no role in the design or conduct of the study, collection, management, analysis or interpretation of data; or preparation, review or approval of the manuscript. Fine et al. disclosed that the last author had financial and consulting interests (Pfizer was not cited as one of them).’

### Diagnostic test accuracy

Pooled sensitivity of allegiant studies was 0.87 (95% CI 0.77 to 0.93), pooled specificity was 0.87 (95% CI 0.76 to 0.94) and the pooled DOR was 49.31 (95% CI 25.74 to 94.48)—see table 5. Heterogeneity was moderate (I²=55.1%). Figure 2 represents the sROCs for this group.

Pooled sensitivity of non-allegiant studies was 0.76 (95% CI 0.67 to 0.83), pooled specificity was 0.88 (95% CI 0.85 to 0.91) and the pooled DOR was 24.96 (95% CI 14.81 to 42.08), approximately half that of the allegiant studies (table 2). Heterogeneity was high at I²=81.5%. Figure 2 represents the sROCs for this group.

The meta-regression for the studies using a cut-off point of 10 or above with allegiance status of the predictor showed that allegiance status was a significant predictor of the DOR (p=0.015) and explained 19.0% of observed heterogeneity.

### Quality assessment

The results of the quality assessment using the QUADAS-2 are given in table 4. For the patient selection domain, the two groups of studies were broadly comparable on two items (consecutive or random sample, avoid case-control design). However, all allegiant studies were rated as avoiding inappropriate exclusions (5/5) in contrast to 58% (15/26) of the non-allegiant studies.

On the index test domain, there were a number of differences between the two groups of studies. More of the non-allegiant studies (81%, 21/26) reported that the PHQ-9 was interpreted blind to the reference test compared with 60% (3/5) of the allegiant studies. All

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**Table 5** Pooled estimates of diagnostic properties of the Patient Health Questionnaire-9 at cut-off point 10 and using algorithm scoring method in the non-independent studies groups

| Settings       | No. of studies | No. of patients | Sensitivity (95% CI) | Specificity (95% CI) | Pooled positive likelihood ratio (95% CI) | Pooled negative likelihood ratio (95% CI) | Diagnostic OR (95% CI) | Heterogeneity: I² |
|----------------|----------------|-----------------|---------------------|---------------------|------------------------------------------|-------------------------------------------|----------------------|-------------------|
| Manea et al, 2014 SR–RA group | 7              | 4985            | 0.77 (0.70 to 0.84) | 0.94 (0.90 to 0.97) | 14.97 (8.39 to 26.71)                   | 0.23 (0.17 to 0.31)                     | 64.40 (24.15 to 121.43) | 78.9%             |
| Manea et al, 2014 SR Independent studies | 2             | 9900            | 0.48 (0.41 to 0.55) | 0.94 (0.91 to 0.99) | 8.26 (6.15 to 11.09)                    | 0.54 (0.48 to 0.62)                     | 15.05 (11.03 to 20.52) | 68.1%             |
| Moriarty et al, 2015 SR–RA group | 5              | 6188            | 0.87 (0.77 to 0.93) | 0.87 (0.76 to 0.94) | 7.24 (5.34 to 11.03)                    | 0.14 (0.08 to 0.23)                     | 49.31 (25.74 to 94.48) | 55.1%             |
| Moriarty et al, 2015 SR Independent studies | 2             | 13164           | 0.76 (0.67 to 0.83) | 0.88 (0.85 to 0.91) | 6.12 (5.06 to 8.32)                     | 0.26 (0.19 to 0.37)                     | 24.96 (14.81 to 42.08) | 81.5%             |
Figure 2  Patient Health Questionnaire-9 summed items scoring method at cut-off point 10 summary receiver operating characteristic plot for diagnosis of major depressive disorder in allegiant studies (panel A) and non-allegiant studies (panel B). Pooled sensitivity and specificity using a bivariate meta-analysis. HSROC, hierarchical receiver operating characteristic.

(5/5) allegiant studies were rated as prespecifying the threshold on the PHQ-9 compared with 73% (19/26) of the non-allegiant studies. The two sets of studies were broadly comparable in terms of two items from the reference test domain (correctly classify target condition, reference test interpreted blind). Only one allegiant study used a translated version of the index test or reference test, so it is not possible to comment on differences between the two sets of studies in terms of these items from the index or reference test domains. For the flow and timing domain, the two groups of studies were broadly comparable for two of the criteria (interval of 2 weeks or less, all participants receive same reference test). However, fewer than half of the non-allegiant studies met the criterion for ‘all participants included in the analysis’ (42%, 11/26), whereas all allegiant studies met this criterion.

DISCUSSION

This is to our knowledge the first systematic examination of a possible ‘allegiance’ or authorship effect in the validation of screening or case-finding psychological instrument for a common mental health disorder. We reviewed diagnostic validation studies of the PHQ-9, a widely used depression screening instrument. We found that allegiant studies reported higher sensitivity paired with similar specificity compared with non-allegiant studies. When entered as a covariate in meta-regression analyses, allegiance status was predictive of variation in the DOR for both the algorithm scoring method and the summed-item scoring method at a cut-off point of 10 or above.

Previous research has proposed several possible explanations for the allegiance effect. One possibility is the adventent bias that may serve to inflate the performance of a test when evaluated by those who have developed it. However, before concluding that the differences are due to this, it is important to explore and rule out alternative explanations. First, it is possible that any observed differences are a result of differences in study characteristics of the two sets of studies (e.g., setting, clinical population). Second, differences in the methodological quality of the studies may also account for any differences. These possibilities are examined below.

Difference in study characteristics as potential alternative explanations

The two sets of studies were broadly comparable in terms of gender and the prevalence of depression, so these variables are unlikely to offer an explanation for the differences. While there were some indications from both sets of comparisons that the PHQ-9 may have been researcher-administered more often in the independent studies, it is not immediately clear how this would lead to lowered diagnostic performance.

The diagnostic meta-analyses of the PHQ-913 14 have shown that the sensitivity and DOR of the PHQ-9 tends to be lower in hospital settings for both algorithm and summed-item scoring methods. While the fact that proportionally more non-allegiant algorithm studies were conducted in secondary care could explain the lower sensitivity and DOR values in the algorithm studies, in the studies that reported the cut-off point of or above this would not be the case as proportionally more allegiant studies were conducted in hospital settings.

Similarly, differences in the proportions of studies using translated versions of the PHQ-9 are also unlikely to offer
an obvious explanation of the difference in diagnostic performance, because in the algorithm set of studies more of the allegiant studies used a translated version of the test, but the proportions were in the opposite direction for the studies using a cut-off of 10 or above. We tested this by carrying out a sensitivity analysis restricting the sample to English studies and studies with adequate translation. The allegiance effect was still predictive of DOR variation between allegiance and non-allegiance studies variation in both algorithm (p = 0.00) and summed item scoring at cut-off point of 10 meta-analyses (p = 0.02).

A similar conclusion is also likely to apply to the age of the samples. There were more older adults studies in the non-allegiant than allegiant studies in the algorithm comparison. Depression could be more difficult to identify in older adults due to physical comorbidities that may present with similar symptomatology to depression and could account for the lower diagnostic performance in the non-allegiant studies. However, the non-allegiant samples in the studies that reported the psychometric properties at cut-off point 10 or above had younger samples than the allegiant studies, so this would not support this interpretation.

The SCID was used as the gold standard in nearly all allegiant studies. The fact that some non-allegiant studies used other gold standards could potentially explain the poorer psychometric properties of the PHQ-9 in these studies. The SCID is often regarded as the most valid of the available semi-structured interviews used in depression diagnostic validity studies as the reference standard. If we assume that this is the case and, furthermore, that the PHQ-9 is an accurate method of screening for depression, then the PHQ-9 may be more likely to agree with the SCID than other reference standards. However, when we carried out a sensitivity analysis restricting the sample to SCID-only studies, the allegiance effect was still predictive of DOR variation between allegiance and non-allegiance studies variation in both algorithm (p = 0.01) and summed item scoring at cut-off point of 10 reviews (p = 0.02).

Differences in methodological quality as potential alternative explanations

The quality of the studies was evaluated using the QUADAS-2. Although there were several potential methodological differences between the two groups of studies from the algorithm papers, not all of these offer obvious explanations of the observed differences and some are unlikely as explanations. For example, more allegiant studies ensured that the reference test was interpreted blind to the index test. This is unlikely to account for the observed differences, because a lack of blinding is typically associated with artificially increased diagnostic performance, which is in the opposite direction to the pattern of results observed here. The impact of some other differences is less clear-cut. For example, a higher number of the non-allegiant studies met the criterion for consecutive referrals. For this to provide an explanation of the observed differences, the non-consecutive nature of the referrals in the studies by those who had developed the PHQ-9 would need to have led to the overinclusion of true positives or underinclusion of false negatives given that these studies tended to report higher sensitivity relative to the non-allegiant studies (and vice versa for the independent studies). It is not immediately obvious how this would occur. The allegiant studies were more likely to have met the criterion of ‘included all participants in the analysis’. It is possible that the greater loss of participants from the non-allegiant studies may have artificially reduced the observed diagnostic accuracy, although, again, it is not immediately obvious how this would have affected the true positive and false negative rates. Although there is not an obvious explanation of how these differences in methodological quality could account for the observed differences in diagnostic performance, it is important to recognise that they cannot on that basis be ruled out.

There are, however, two differences in methodological quality among the algorithm studies that are clearer potential alternative explanations. The higher rate of appropriate translations among the allegiant studies is potentially important, because lower diagnostic estimates may be expected from studies that have poorly translated versions of the index test. In the flow and timing domain, more allegiant studies ensured that there was a less than 2-week interval between the index and reference test. This is consistent with lower diagnostic performance in the non-allegiant studies: as the interval increases it is likely that depression status may change and this would lead to lower levels of agreement between the index test and the reference test.

There were also differences on some quality assessment items between the two sets of studies in the summed item scoring method comparison. The threshold was reported as prespecified in all allegiant studies in contrast to approximately three-quarters of the non-allegiant studies. On the face of it, this is unlikely to explain the observed differences, because the use of a prespecified cut-off point is likely to be associated with lower not higher diagnostic test performance. One possibility, however, is that studies that performed poorly at this cut-off point were less likely to be reported by those who had developed the measure. As discussed in more detail in the ‘Limitations’ section, we were unable to explore this possibility through the use of formal tests for publication bias.

All allegiant studies avoided inappropriate exclusions compared with approximately half of the non-allegiant studies. While this is a potential alternative explanation of the differences, it is not immediately obvious how this would explain the differences in diagnostic performance between the two sets of studies. Fewer than half of the non-allegiant studies met the criterion for ‘all participants included in the analysis’, in contrast to all of the allegiant studies met this criterion, but again this difference should usually work against the inclusive studies, not those excluding cases. More of the non-allegiant studies reported that the PHQ-9 was interpreted blind to the
reference test. This does offer a potential explanation, because the absence of blinding may artificially inflate diagnostic accuracy.

LIMITATIONS

The results of this review need to be viewed in light of the limitations of the primary studies that contributed to the review and the review itself. An important consideration is to establish whether any observed differences between the diagnostic performance of the non-allegiant and allegiant studies are better accounted for by study characteristic or methodological differences. Caution, however, is needed in interpreting any differences, because of the small number of allegiant studies in both the algorithm and cut-off 10 or above comparisons. The small number of allegiant studies also meant that we were also unable to explore the potential role of publication bias in the non-allegiant and allegiant studies. At least 10 studies are required to use standard methods of examining publication bias, but the number of allegiant studies in both the algorithm and cut-off 10 or above comparisons were fewer than this. Papers published from August 2013 onwards are not covered in the literature search used and so it potentially misses some more recent studies that would be eligible for inclusion, although it is unlikely that many, if any, new allegiant studies have been published since.

CONCLUSIONS AND IMPLICATIONS FOR FURTHER RESEARCH

The aims of the review was to investigate whether an allegiance effect is found that leads to an increased diagnostic performance in diagnostic validation studies that were conducted by teams connected to the original developers of the PHQ-9. Our analyses showed that diagnostic studies conducted by independent/non-allegiant researchers had lower sensitivity paired with similar specificity compared with studies that were classified as allegiant. This conclusion held for both the algorithm and cut-off 10 or above studies. We explored a range of possible alternative explanations for the observed allegiance effect including both differences in study characteristics and study quality. A number of potential differences were found, although for some of these it is not clear how they would necessarily account for the observed differences. However, there were a number of differences that offered potential alternative explanations unconnected to allegiance effects. In the algorithm studies, the studies rated as allegiant were also more likely to use an appropriate translation of the PHQ-9 and were also more likely to ensure that the index and reference test were conducted within 2 weeks of each other, both of which may be associated with an improvement in observed diagnostic performance of an instrument. The majority of studies in both meta-analyses did not provide clear statements about potential conflict of interest and/or funding; however, the newer studies were more likely to provide such statements, which may reflect increasing transparency in this area of research.

We cannot, therefore, conclude that allegiance effects are present in studies examining the diagnostic performance of the PHQ-9; but nor can we rule them out. Conflicts of interest are an important area of investigation in medical and behavioural research, particularly due to concerns about trial results being influenced by industry sponsorship. Future diagnostic validity in this area should as a matter of routine present clear statements about potential conflicts of interest and funding, particularly relating to the development of the instrument under evaluation. Future meta-analyses of diagnostic validation studies of psychological measures should routinely evaluate the impact of researcher allegiance in the primary studies examined in the meta-analysis.

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