

## ORIGINAL ARTICLE

# Network meta-analysis of diagnostic test accuracy studies identifies and ranks the optimal diagnostic tests and thresholds for health care policy and decision-making

Rhiannon K. Owen<sup>a,\*</sup>, Nicola J. Cooper<sup>a</sup>, Terence J. Quinn<sup>b</sup>, Rosalind Lees<sup>b</sup>, Alex J. Sutton<sup>a</sup><sup>a</sup>*Department of Health Sciences, University of Leicester, Leicester, UK*<sup>b</sup>*Institute of Cardiovascular and Medical Sciences, University of Glasgow, Glasgow, UK*

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**Abstract**

**Objectives:** Network meta-analyses (NMA) have extensively been used to compare the effectiveness of multiple interventions for health care policy and decision-making. However, methods for evaluating the performance of multiple diagnostic tests are less established. In a decision-making context, we are often interested in comparing and ranking the performance of multiple diagnostic tests, at varying levels of test thresholds, in one simultaneous analysis.

**Study Design and Setting:** Motivated by an example of cognitive impairment diagnosis following stroke, we synthesized data from 13 studies assessing the efficiency of two diagnostic tests: Mini-Mental State Examination (MMSE) and Montreal Cognitive Assessment (MoCA), at two test thresholds: MMSE <25/30 and <27/30, and MoCA <22/30 and <26/30. Using Markov chain Monte Carlo (MCMC) methods, we fitted a bivariate network meta-analysis model incorporating constraints on increasing test threshold, and accounting for the correlations between multiple test accuracy measures from the same study.

**Results:** We developed and successfully fitted a model comparing multiple tests/threshold combinations while imposing threshold constraints. Using this model, we found that MoCA at threshold <26/30 appeared to have the best true positive rate, whereas MMSE at threshold <25/30 appeared to have the best true negative rate.

**Conclusion:** The combined analysis of multiple tests at multiple thresholds allowed for more rigorous comparisons between competing diagnostics tests for decision making. © 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

**Keywords:** Network meta-analysis; Meta-analysis; Diagnostic test accuracy; Multiple tests; Multiple thresholds

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## 1. Background

Evidence-based healthcare evaluations, as endorsed by the National Institute for Health and Care Excellence (NICE) in the UK and similar organizations worldwide, have highlighted the crucial role systematic reviews, including meta-analysis where appropriate, have to play

in the decision-making process to answer clinically relevant questions such as whether a technology works, for whom and how it compares with alternatives [1]. Such evidence-based evaluations are important to the decision-making process within the area of diagnostic test performance as early diagnosis of disease can lead to more successful treatment than if treatment is delayed.

Diagnostic test accuracy is defined by Leeflang et al. [2] as the ability of a test to distinguish between patients with a specified target condition and those without, and the results are usually expressed in terms of sensitivity (i.e., the proportion of people with the condition correctly detected by the test) and specificity (i.e., the proportion of people without the condition correctly detected by the test). The dependence between these outcomes (i.e., sensitivity and specificity) adds an additional complexity that makes evidence synthesis of diagnostic test accuracy data more complicated than for intervention studies. The dependence

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\* Corresponding author. Tel.: +44 0 116 229 7295.

E-mail address: [Rhiannon.owen@le.ac.uk](mailto:Rhiannon.owen@le.ac.uk) (R.K. Owen).

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**What's new?****Key findings**

- We propose a unified model for synthesizing diagnostic test accuracy data of multiple tests at multiple explicit thresholds.
- Building on this model, we incorporate constraints on increasing test-thresholds such that higher test-thresholds have an increased sensitivity but decreased specificity.
- Incorporating constraints on threshold effects has the potential to more appropriately attribute variability between results to genuine threshold effects and better explain heterogeneity between studies.

**What this adds to what was known?**

- Methods for meta-analysing diagnostic test accuracy data are suited to the analysis of single tests.

**What is the implication and what should change now?**

- A network meta-analysis framework allows comparisons to be made between all competing diagnostic tests and consequently inferences regarding the relative ranking of each test can be made for health care decision-making.

between outcomes can occur in one of two ways. For meta-analyses of studies evaluating a single pair of sensitivity and specificity, the dependence between sensitivity and specificity occurs across studies for differing thresholds [3]. For meta-analyses of studies evaluating multiple tests and/or multiple thresholds, and therefore, multiple pairs of sensitivity and specificity, the dependence between outcomes occurs both within and across studies. Another additional issue includes correctly estimating the joint conditional performance of multiple tests when they are used in combination [4]. A number of methods have been proposed to meta-analyze studies reporting single pairs of sensitivity and specificity data including independent meta-analyses of sensitivity and specificity [5], meta-analysis of diagnostic odds ratios [5], summary receiver operating characteristic (sROC) regression modeling [6], hierarchical sROC model [7], and bivariate meta-analysis models [7,8]. In cases where the test threshold is suspected or known to vary between studies, the latter two approaches (which have been shown to be equivalent [9]) are the most appropriate as they allow for this variability in the analysis. However, neither of these incorporates test threshold information explicitly into the analysis, which hampers accurate prediction of test performance at particular thresholds; a fact which limits the clinical applicability of results, at least

in contexts where test threshold can be explicitly specified. Work extending the hierarchical sROC/bivariate approach, allowing for data from multiple thresholds for the same tests from the same study to be synthesized, was described some time ago [10] but is rarely used in practice and suffers from the same limitation that explicit threshold value information is not included in the analysis. Alternative approaches have since been described [3,11–14] including a generalization of the bivariate model to include the use of multivariate random effects [11] and the use of Poisson-correlated gamma frailty models [12]. For both of these approaches, the number of thresholds across all studies has to be identical, which is often impractical in a meta-analysis setting. Multivariate regression models [13] and linear mixed effects models [3] have also been proposed. These methods consist of a two-stage approach; however, estimation of the uncertainty from stage-one of these analyses is ignored and may lead to unrealistic results. More recently, Hoyer et al. [14] proposed a bivariate time-to-event model for interval-censored data incorporating random effects. This approach overcomes the limitations described above and provides a flexible framework to account for various distributions of the underlying diagnostic marker. However, a limitation of this model is the potential constraint of the proportional hazards assumptions when meta-analyzing receiver operating characteristic (ROC) curves from studies that report a single threshold.

In comparison to interventional research, methodology for meta-analyzing diagnostic test accuracy data has relatively recently been adopted by Cochrane. As such, the development of guidance and best practice statements are ever-evolving. However, at present, the preferred Cochrane methods are suited to meta-analyzing diagnostic test accuracy of single tests (using the statistical methods outlined above). From a clinical and decision-making perspective, often interest lies in assessing the performance of multiple diagnostic tests with multiple thresholds in one simultaneous analysis because it addresses clinically relevant questions such as which test at which threshold is most effective or most cost-effective [15]. In interventional research, when multiple competing health care interventions are of interest, network meta-analyses (NMA) have been used extensively to compare and identify the “best” intervention(s). However, methods for evaluating the performance of multiple diagnostic tests are less established. Network meta-analyses of health care interventions are commonly used to synthesise data from several clinical trials in similar patient populations with the aim to evaluate multiple interventions that may not have been compared otherwise. This approach combines both direct information (obtained from head-to-head trials) and indirect information (obtained from trials that share a common comparator) to obtain relative treatment effects for all interventions while maintaining randomization. Combining direct and indirect information in this way assumes an additive relationship between treatment effects. In a diagnostic test accuracy

setting, this framework has previously been adopted to model the difference or relative risk in sensitivity and specificity [16] and the relative diagnostic odds ratio between two tests [16–18]. Throughout this article, NMA are used to describe the synthesis of diagnostic test accuracy data from a network of diagnostic tests that have been evaluated in the same study and thus, the same individuals. Our approach differs to the framework commonly used for health care interventions in that our approach models the absolute sensitivity and specificity of each test incorporating random effects to allow for heterogeneity as well as similarities between data belonging to the same study and the same tests within studies (i.e., to account for multiple thresholds within tests).

Through careful consideration of a motivating example, this article sets out an approach to network meta-analysis of diagnostic test accuracy studies that allows for both the incorporation of multiple tests and multiple explicit threshold values, potentially reported by the same studies. Section 2 introduces the motivating example. Section 3 discusses the visual representation of diagnostic test accuracy data, highlighting where appropriate, key assumptions that are revisited throughout the remainder of this article. Section 4 describes each of the proposed meta-analysis models. Section 5 presents the results from the analysis of the motivating example and Section 6 concludes this article with a discussion.

## 2. Motivating example

Cognitive impairment is highly prevalent in stroke survivors, and it can lead to increased mortality, disability, and institutionalization. Early detection of cognitive impairment is an essential step in the efficient management of patient care. In the UK, screening for cognitive impairment is recommended by governing bodies such as NICE and the Royal College of Physicians (RCP). However, although cognitive assessment is recommended in various clinical guidelines, there is no consensus on how to efficiently diagnose patients, where differing guidelines recommend differing tests and thresholds [19]. A robust synthesis of the evidence in one simultaneous analysis has the potential to provide an evidence base where currently best clinical practice is primarily opinion based.

In a study by Lees et al. [20], the authors investigated the test accuracy of multiple-screening tests for the diagnosis of cognitive impairment and dementia in stroke patients. Cognitive impairment is an umbrella term that encompasses any objective memory and thinking problem. It includes, but is not limited to, the clinical syndrome of dementia. Gold standard assessment for cognitive impairment is a detailed examination of various aspects of cognition (neuropsychological battery [NPB]), although there is no consensus on the “gold” standard antemortem diagnosis of dementia, dementia diagnosis is currently made according to clinical criteria that can often be informed by data from NPB. In

the Lees et al. review, as the index tests of interest were “screening” tests, the authors decided to include both cognitive impairment on NPB and clinical dementia diagnosis as their reference standard. The authors were able to synthesize data from 13 diagnostic test accuracy studies for two key screening tests: Folstein’s Mini-Mental State Examination (MMSE) and Montreal Cognitive Assessment (MoCA). Mini-mental state examination and MoCA examinations are based on a scoring system, where a higher value indicates a more desired response from the participant. Thus a disease-positive response on a lower test threshold would suggest a more severe case of cognitive impairment. The authors focused on two disparate cut points per test for dementia diagnosis: MMSE <25/30, MMSE <27/30, MoCA <22/30, MoCA <26/30. In this article, each of the test-threshold combinations are treated as independent tests. Fig. 1 illustrates the network of comparative studies. The nodes represent each of the test-threshold combinations of interest. The dashed interconnecting lines illustrate that a comparative study examining both tests in the same patient population exists. Typically networks of treatment comparisons are presented alongside NMA of health care interventions, where the interconnecting lines represent direct evidence on the relative differences between treatments. Fig. 1 differs to a network of treatment comparisons for NMA of health care interventions whereby the interconnecting lines represent a comparative study illustrating tests that have been undertaken in the same cohort of patients and thus, there is a within-study dependence between pairs of sensitivity and specificity for these tests. Table 1 shows the extracted data from the original articles. The choice of test thresholds were determined by the most commonly reported test-threshold combinations in the published literature at the time of publication. The reference standard tests included NPB and clinical diagnosis of dementia (and the authors assumed that these were perfect, as do we in this article). In this example, Lees et al. [20] pooled data from studies using both reference tests. Data from 12, 5, 4, and 6 studies were pooled

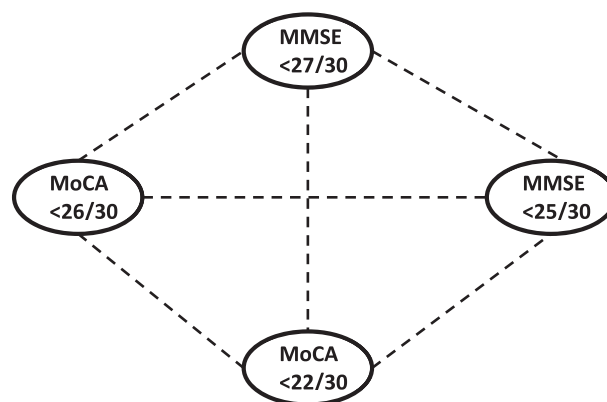


Fig. 1. Network of comparative studies. MMSE, Mini-Mental State Examination; MoCA, Montreal Cognitive Assessment.

**Table 1.** Diagnostic test accuracy data obtained from the original articles

| Study author    | Test     | TP | FP | FN | TN  | Sensitivity (95% CI) | Specificity (95% CI) |
|-----------------|----------|----|----|----|-----|----------------------|----------------------|
| Blake 2002      | MMSE <25 | 19 | 10 | 12 | 71  | 0.61 (0.42, 0.78)    | 0.88 (0.78, 0.94)    |
| Bour 2010       | MMSE <25 | 21 | 29 | 1  | 143 | 0.95 (0.77, 1.00)    | 0.83 (0.77, 0.88)    |
| Cumming 2010    | MMSE <25 | 48 | 22 | 10 | 69  | 0.83 (0.71, 0.91)    | 0.76 (0.66, 0.84)    |
| Cumming 2013    | MMSE <25 | 21 | 4  | 17 | 17  | 0.55 (0.38, 0.71)    | 0.81 (0.58, 0.95)    |
|                 | MMSE <27 | 35 | 10 | 3  | 11  | 0.92 (0.79, 0.98)    | 0.52 (0.30, 0.74)    |
|                 | MoCA <26 | 39 | 12 | 0  | 9   | 1.00 (0.91, 1.00)    | 0.43 (0.22, 0.66)    |
|                 | MoCA <22 | 30 | 5  | 9  | 16  | 0.77 (0.61, 0.89)    | 0.76 (0.53, 0.92)    |
| de Koning 1998  | MMSE <25 | 44 | 51 | 11 | 178 | 0.80 (0.67, 0.90)    | 0.78 (0.72, 0.83)    |
| Dong 2010       | MMSE <25 | 52 | 10 | 8  | 45  | 0.87 (0.75, 0.94)    | 0.82 (0.69, 0.91)    |
|                 | MMSE <27 | 45 | 17 | 3  | 26  | 0.94 (0.83, 0.99)    | 0.60 (0.44, 0.75)    |
|                 | MoCA <22 | 54 | 13 | 6  | 42  | 0.90 (0.79, 0.96)    | 0.76 (0.63, 0.87)    |
| Dong 2012       | MMSE <25 | 28 | 32 | 4  | 65  | 0.88 (0.71, 0.96)    | 0.67 (0.57, 0.76)    |
|                 | MoCA <22 | 53 | 65 | 7  | 114 | 0.88 (0.77, 0.95)    | 0.64 (0.56, 0.71)    |
| Godefroy 2011   | MMSE <25 | 45 | 2  | 19 | 29  | 0.70 (0.58, .81)     | 0.94 (0.79, 0.99)    |
|                 | MMSE <27 | 55 | 12 | 9  | 19  | 0.86 (0.75, 0.93)    | 0.61 (0.42, 0.78)    |
|                 | MoCA <26 | 60 | 20 | 4  | 11  | 0.94 (0.85, 0.98)    | 0.35 (0.19, 0.55)    |
|                 | MoCA <22 | 48 | 4  | 16 | 28  | 0.75 (0.63, 0.85)    | 0.88 (0.71, 0.96)    |
| Grace 1995      | MMSE <25 | 20 | 9  | 26 | 46  | 0.43 (0.29, 0.59)    | 0.84 (0.71, 0.92)    |
| Morris 2012     | MMSE <25 | 21 | 3  | 15 | 10  | 0.58 (0.41, 0.74)    | 0.77 (0.46, 0.95)    |
|                 | MMSE <27 | 30 | 8  | 6  | 5   | 0.83 (0.67, 0.94)    | 0.38 (0.14, 0.68)    |
| Pendlebury 2012 | MMSE <25 | 11 | 3  | 8  | 69  | 0.58 (0.33, 0.80)    | 0.96 (0.88, 0.99)    |
|                 | MMSE <27 | 15 | 15 | 4  | 57  | 0.79 (0.54, 0.94)    | 0.79 (0.68, 0.88)    |
|                 | MoCA <26 | 19 | 39 | 0  | 33  | 1.00 (0.82, 1.00)    | 0.46 (0.34, 0.58)    |
|                 | MoCA <22 | 13 | 11 | 6  | 61  | 0.68 (0.43, 0.87)    | 0.85 (0.74, 0.92)    |
| Salvadori 2013  | MoCA <26 | 78 | 46 | 2  | 29  | 0.97 (0.91, 1.00)    | 0.39 (0.28, 0.51)    |
|                 | MoCA <22 | 73 | 18 | 7  | 58  | 0.91 (0.83, 0.96)    | 0.76 (0.65, 0.85)    |
| Srikanth 2006   | MMSE <25 | 4  | 4  | 4  | 67  | 0.50 (0.16, 0.84)    | 0.94 (0.86, 0.98)    |

Abbreviations: TP, true positives; FP, false positives; FN, false negatives; TN, true negatives; CI, confidence interval; MMSE, Mini-Mental State Examination; MoCA, Montreal Cognitive Assessment.

in separate bivariate meta-analyses for MMSE <25/30, MMSE <27/30, MoCA <26/30, and MoCA <22/30, respectively.

### 3. Illustrating diagnostic test accuracy data

This section highlights some of the key principles of diagnostic test accuracy data through the use of visual representations. Figures 2 and 3 illustrate the relationship between the thresholds of the same study for MMSE and MoCA, respectively. The nodes represent the observed sensitivity and specificity, color coded for each threshold; the corresponding number represents the exact threshold used. The interconnecting lines illustrate the ROC for each study that reports multiple threshold values. One of the properties of an ROC is that higher test thresholds for a positive outcome must, mathematically, have an increased sensitivity but decreased specificity. For the remainder of this article, this property will be referred to as a threshold

assumption. In this example, MMSE and MoCA are scored out of 30 points, with a point deducted for each error. Lower scores, therefore, suggest greater impairment. If the “test positive” threshold is lowered then at the lower threshold the test is more specific and less sensitive, and thus should lie in the lower left-hand side of the ROC space. From Fig. 2, it is evident that there is a large amount of heterogeneity in sensitivities and specificities reported between the studies identified by Lees et al. [20], where data points with lower thresholds for MMSE lie toward the top left hand-side of the ROC space, above that of higher thresholds. Fig. 4 illustrates schematically the relationship between the threshold assumption and that of heterogeneity. Heterogeneity between studies can be described as the difference in ROCs between studies, shifting the ROC in a southeast or northwest direction from the summary ROC. The threshold assumption shifts the observed sensitivity and specificity in a northeast or southwest direction along the study-specific ROC.

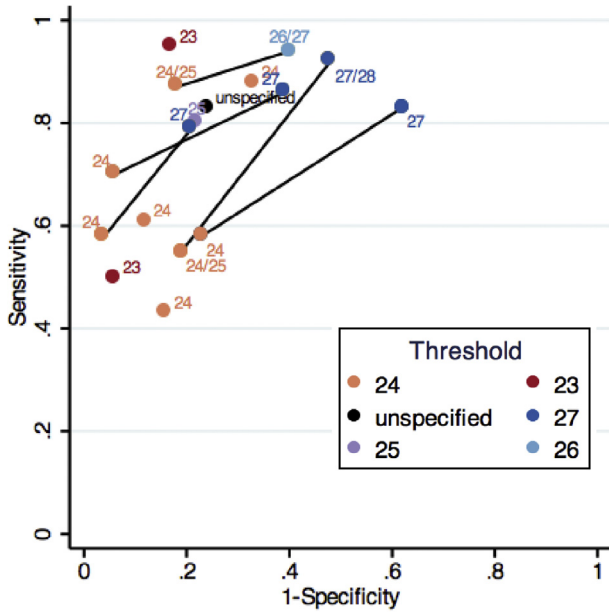


Fig. 2. Relationship of increasing test thresholds within Mini-Mental State Examination (MMSE).

#### 4. Models and estimation

##### 4.1. Network meta-analysis for diagnostic test accuracy studies

Building on the bivariate meta-analysis model described by Reitsma et al. [8] and Sutton et al. [15], let the observed number of true positives,  $tp_i$ , for the  $i$ th observation be considered as a binomial count from a sample of disease-positive individuals,  $pos_i$ . This information allows for estimation of the diagnostic sensitivity,  $sens_i$ , which is associated with the rate of accurate detection of diseased individuals. Likewise, let the number of true negatives,  $tn_i$ , be considered as a binomial count from a sample of disease-negative individuals,  $neg_i$ . This information allows for estimation of the specificity of the test,  $spec_i$ , which is associated with the rate of accurate detection of nondiseased individuals, such that:

$$\begin{aligned} tp_i &\sim \text{Binomial}(sens_i, pos_i) \\ tn_i &\sim \text{Binomial}(spec_i, neg_i) \end{aligned} \quad 1$$

Logistic regression models can be used to specify sensitivity and specificity. Across studies, the sensitivity and specificity of each test are likely to be correlated. To account for this across-study dependence, the (*logit*) sensitivity,  $sens_i$ , and specificity,  $spec_i$ , of observation  $i$ , where observation  $i$  represents each pair of sensitivities and specificities for each study, are drawn from a bivariate normal distribution with mean equal to the pooled test accuracy estimates of sensitivity,  $\mu_{sens}$ , and specificity,  $\mu_{spec}$ , with between-observation covariance matrix  $\Sigma$ :

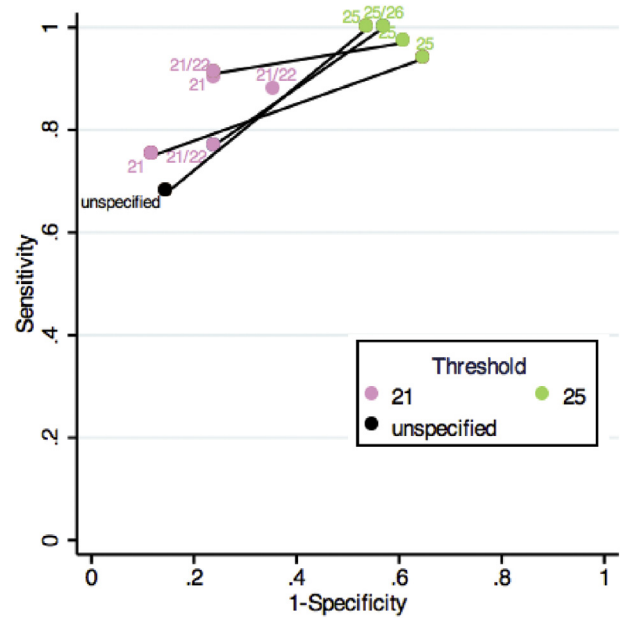


Fig. 3. Relationship of increasing test thresholds within Montreal Cognitive Assessment (MoCA).

$$\begin{aligned} \begin{pmatrix} \text{logit}(sens_i) \\ \text{logit}(spec_i) \end{pmatrix} &\sim \text{Normal} \left[ \begin{pmatrix} \mu_{sens} \\ \mu_{spec} \end{pmatrix}, \Sigma \right] \\ \Sigma &= \begin{pmatrix} \tau_{sens}^2 & \rho \tau_{sens} \tau_{spec} \\ \rho \tau_{sens} \tau_{spec} & \tau_{spec}^2 \end{pmatrix}, \end{aligned} \quad 2$$

where  $\tau_{sens}$  and  $\tau_{spec}$  denote the between-observation standard deviation (SD) in *logit* transformed sensitivity and specificity, and  $\rho$  denotes the between-observation correlation. To model the inherent correlations between multiple sensitivity and specificity data pairs from the same study,

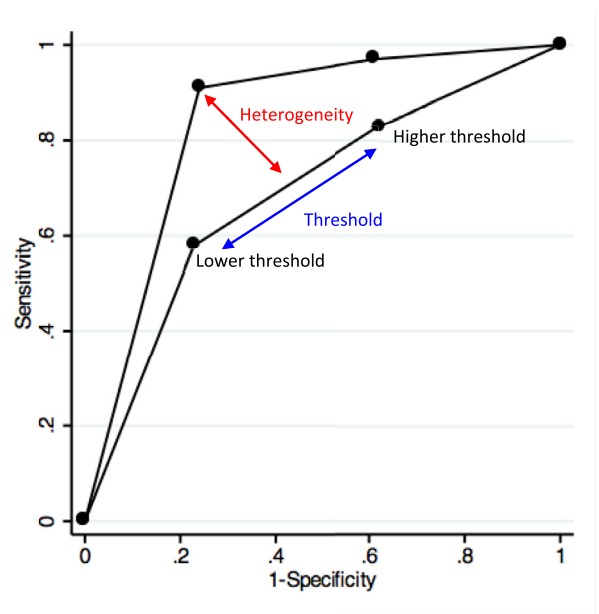


Fig. 4. Schematic relationship of the impact of heterogeneity in test performance due to threshold and between-study heterogeneity. The labels “Lower threshold” and “Higher threshold” apply specifically to the given example.



a variance component model can be used [21] such that diagnostic test-threshold combinations are considered fixed effects, while study and study-specific diagnostic test are considered random effects. Study and study-specific diagnostic test are nested within each observation. The model is given by:

$$\begin{aligned}\mu_{isens} &= \beta_{j,k} + c_{l(i)} + d_{l(i),j(i)} \\ \mu_{ispec} &= \delta_{j,k} + e_{l(i)} + f_{l(i),j(i)}\end{aligned}\quad 3$$

where  $\beta$  and  $\delta$  denote fixed effects of sensitivity and specificity due to diagnostic test,  $j$ , and test-threshold,  $k$ , respectively. Parameters  $c$  and  $d$ , and  $e$  and  $f$ , denote random effects of sensitivity and specificity due to study,  $l$ , and the interaction between study,  $l$ , and diagnostic test,  $j$ , respectively. Noninformative prior distributions were specified for the test-specific and threshold-specific accuracy parameters on the *logit* scale (with more information given in technical Appendix A 1). All models were estimated in a Bayesian framework using Markov Chain Monte Carlo (MCMC) simulation and implemented in the WinBUGS 1.4.3 software [22]. Example WinBUGS code is given in Appendix A 2.

#### 4.2. Network meta-analysis incorporating threshold constraints

Building on the network meta-analysis framework of 4.1, constraints can be specified on the underlying threshold parameters, assuming that overall, higher test thresholds have an increased sensitivity but decreased specificity. We applied these constraints using a method described by Owen et al. [23], where the authors place constraints on increasing doses of an intervention for comparative effectiveness research. In this study, imposing constraints allows information to be

borrowed between thresholds within a test, and thus potentially increases the precision in the estimated test accuracy for decision-making (which is further discussed in Section 6). An alternative approach is to specify the underlying cumulative distribution function of the tests [3,12,14]. This approach is flexible to multiple and different thresholds. However, specifying constraints on the summary estimates of the accuracy measures for increasing test thresholds, as described in this article, makes fewer assumptions about the distributional form of these estimates and rather this approach simply allows the summary estimates to be greater than or equal to, or less than or equal to, the reference threshold.

#### 5. Analysis of the motivating example

In this section, we synthesized diagnostic test accuracy data across all tests and thresholds of interest. This includes MMSE at threshold <25 and <27, and MoCA at threshold <22 and <26. We illustrate estimates of diagnostic test accuracy using a number of models each with different heterogeneity and correlation assumptions, as described in the technical appendix (Appendix A.1). We extend each of these models to incorporate constraints on increasing thresholds.

Table 2 gives the model fit statistics for each of these models. Using measures of the deviance information criterion, it is apparent that there is very little difference in model fit across all models considered (a difference in the deviance information criterion > 5 is considered an important difference). For this reason, the most simplistic model, assuming a common heterogeneity and correlation parameter across models was used for illustration. The results from the remaining models are given in Appendix A.3.

**Table 2.** Model fit statistics

| Model   | Posterior between-observation SD(s) |                 |                 |                 |                          |                  | DIC    |
|---|-------------------------------------|-----------------|-----------------|-----------------|--------------------------|------------------|--------|
|   | Sensitivity                         |                 | Specificity     |                 | Correlation parameter(s) |                  |        |
|   | MMSE                                | MoCA            | MMSE            | MoCA            | MMSE                     | MoCA             |        |
| Assuming common heterogeneity and correlation parameters across tests               |                                     |                 |                 |                 |                          |                  |        |
| Without threshold constraints   | 0.28 (SD: 0.21)                     |                 | 0.17 (SD: 0.13) |                 | −0.17 (SD: 0.70)         |                  | 281.59 |
| With threshold constraints  | 0.29 (SD: 0.21)                     |                 | 0.16 (SD: 0.13) |                 | −0.16 (SD: 0.70)         |                  | 281.23 |
| Assuming common heterogeneity and test-specific correlation parameters              |                                     |                 |                 |                 |                          |                  |        |
| Without threshold constraints   | 0.29 (SD: 0.22)                     |                 | 0.17 (SD:0.13)  |                 | −0.14 (SD: 0.71)         | −0.01 (SD: 0.71) | 281.64 |
| With threshold constraints  | 0.29 (SD: 0.21)                     |                 | 0.17 (SD: 0.13) |                 | −0.20 (SD: 0.70)         | 0.02 (SD: 0.71)  | 281.18 |
| Assuming test-specific heterogeneity and common correlation parameters across tests |                                     |                 |                 |                 |                          |                  |        |
| Without threshold constraints   | 0.39 (SD: 0.27)                     | 0.29 (SD: 0.21) | 0.45 (SD: 0.38) | 0.22 (SD: 0.19) | −0.28 (SD: 0.68)         |                  | 282.94 |
| With threshold constraints  | 0.40 (SD: 0.27)                     | 0.28 (SD: 0.20) | 0.44 (SD: 0.38) | 0.21 (SD: 0.19) | −0.28 (SD: 0.68)         |                  | 282.82 |

Abbreviation: DIC, deviance information criterion; SD, standard deviation.

**Table 3.** Posterior point estimates and 95% credible intervals (CrIs) obtained from a network meta-analysis model assuming a common between-observation standard deviation and correlation parameter

| Test                          | Sensitivity<br>(95% CrI) | Specificity<br>(95% CrI) | Rank best sensitivity<br>(95% CrI) | P (Best)<br>sensitivity | Rank best specificity<br>(95% CrI) | P (Best)<br>specificity |
|-------------------------------|--------------------------|--------------------------|------------------------------------|-------------------------|------------------------------------|-------------------------|
| Without threshold constraints |                          |                          |                                    |                         |                                    |                         |
| MMSE <25                      | 0.72 (0.61, 0.82)        | 0.84 (0.79, 0.89)        | 4 (3,4)                            | 0                       | 1 (1, 2)                           | 0.97                    |
| MMSE <27                      | 0.89 (0.81, 0.95)        | 0.58 (0.45, 0.70)        | 2 (2,3)                            | 0.01                    | 3 (3, 3)                           | 0                       |
| MoCA <22                      | 0.82 (0.70, 0.91)        | 0.77 (0.67, 0.85)        | 3 (2,4)                            | 0                       | 2 (1, 2)                           | 0.03                    |
| MoCA <26                      | 0.97 (0.94, 0.99)        | 0.35 (0.23, 0.48)        | 1 (1,1)                            | 0.99                    | 4 (4, 4)                           | 0                       |
| With threshold constraints    |                          |                          |                                    |                         |                                    |                         |
| MMSE <25                      | 0.73 (0.62, 0.82)        | 0.84 (0.79, 0.88)        | 4 (3,4)                            | 0                       | 1 (1, 2)                           | 0.96                    |
| MMSE <27                      | 0.90 (0.81, 0.95)        | 0.58 (0.44, 0.70)        | 2 (2,3)                            | 0                       | 3 (3, 3)                           | 0                       |
| MoCA <22                      | 0.83 (0.71, 0.91)        | 0.77 (0.67, 0.86)        | 3 (2,4)                            | 0                       | 2 (1, 2)                           | 0.04                    |
| MoCA <26                      | 0.98 (0.94, 0.99)        | 0.35 (0.22, 0.47)        | 1 (1,1)                            | 1                       | 4 (4, 4)                           | 0                       |

### 5.1. Assuming common heterogeneity and correlation parameters across tests

This section illustrates the results from a model assuming a common heterogeneity and correlation parameter across multiple, diverse tests. Section 5.1.1 presents estimates of test accuracy from a model without threshold constraints, and Section 5.1.2 presents results from a model with threshold constraints.

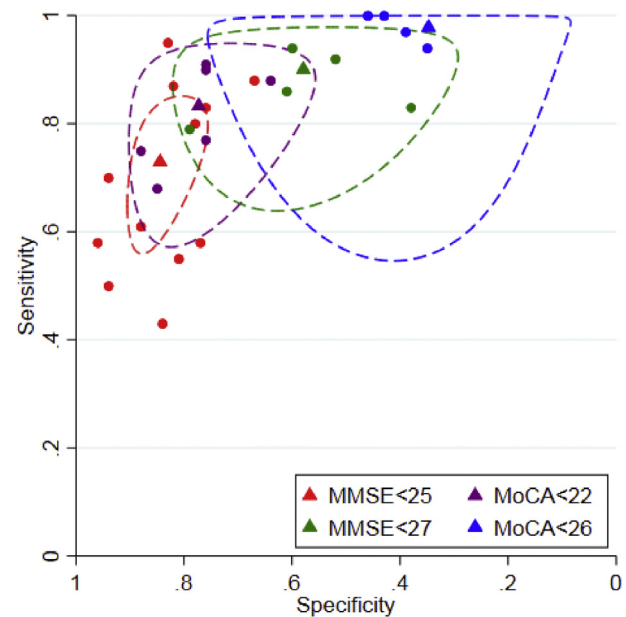
#### 5.1.1. Model without threshold constraints

Table 3 displays the results of the summary test accuracy measures, relative rankings, and probabilities that each test was the most accurate overall in terms of the true positive rate (sensitivity) and true negative rate (specificity). Combining diagnostic test accuracy data for all test-threshold combinations illustrated that MoCA <26 appeared to have the optimal true positive rate (sensitivity: 0.97, 95% credible interval [CrI]: 0.94, 0.99) for 99% of MCMC iterations. There appeared to be little difference between MMSE <27 and MoCA <22 in terms of sensitivity, with each of these tests ranking in second (95% CrI: 2,3) and third place (95% CrI: 2,4), respectively. However, MoCA <22 appeared to have a better true negative rate (specificity: 0.77, 95% CrI: 0.67, 0.85) compared to that of MMSE <27 (specificity: 0.58, 95% CrI: 0.45, 0.70), and subsequently ranked in second place for specificity (rank: 2, 95% CrI: 1, 2). Mini-mental state examination <25 appeared to have the optimal true negative rate overall (specificity: 0.84, 95% CrI: 0.79, 0.89), ranking in first place for 97% of model iterations.

#### 5.1.2. Model with threshold constraints

Incorporating threshold constraints on increasing thresholds marginally reduced the between-observation standard deviation for specificity (Table 2). However, including threshold constraints had very little impact on the posterior point estimates, which mirror those of the unconstrained model (Table 3). Overall, incorporating threshold constraints appeared to marginally increase precision in the effect

estimates. Subsequently, MoCA <26 ranked in first place for sensitivity for 100% of MCMC iterations. Incorporating threshold constraints reduced the variability within studies. The estimated within-study standard deviation for sensitivity was 0.62 (SD: 0.31) compared to 0.63 (SD: 0.30) from the unconstrained model. Similarly, the estimated within-study standard deviation for specificity was 0.42 (SD: 0.21) compared to 0.44 (SD: 0.20) from the unconstrained model. However, the variability within tests within a study marginally increased when incorporating threshold constraints. For sensitivity, the estimated standard deviation from the model incorporating threshold constraints was 0.30 (SD: 0.22) compared to 0.28 (SD: 0.22) from the unconstrained model. For specificity, the estimated standard deviation was 0.27

**Fig. 5.** Posterior point estimates (triangles) and 95% credible regions (dashed lines) in receiver operating characteristic (ROC) space obtained from a model incorporating threshold constraints and assuming a common heterogeneity and correlation parameter across tests. Circles represent individual study estimates. MMSE, Mini-Mental State Examination; MoCA, Montreal Cognitive Assessment.

**Table 4.** Estimated mean difference (95% CrI) in sensitivity (top right) and specificity (bottom left) between each test-threshold combination (row—column) obtained from a model incorporating threshold constraints and assuming a common heterogeneity and correlation parameter across tests

| Test-threshold | MMSE <25           | MMSE <27             | MoCA <22            | MoCA <26          |
|----------------|--------------------|----------------------|---------------------|-------------------|
| MMSE <25       | -                  | 0.17 (0.08, 0.26)    | 0.10 (−0.01, 0.22)  | 0.25 (0.15, 0.35) |
| MMSE <27       | 0.26 (0.15, 0.39)  | -                    | −0.07 (−0.18, 0.03) | 0.08 (0.02, 0.16) |
| MoCA <22       | 0.07 (−0.01, 0.18) | −0.19 (−0.33, −0.06) | -                   | 0.14 (0.07, 0.25) |
| MoCA <26       | 0.49 (0.38, 0.61)  | 0.23 (0.08, 0.37)    | 0.42 (0.31, 0.52)   | -                 |

Above the leading diagonal gives estimates of the mean difference (row—column) in sensitivity (95% CrI), and below the leading diagonal gives estimates of the mean difference in specificity (95% CrI).

(SD: 0.20) compared to a standard deviation of 0.25 (SD: 0.17) for the unconstrained model.

Fig. 5 displays the posterior point estimates and 95% credible regions in ROC space. While MoCA <26 and MMSE <25 rank in first place for sensitivity and specificity, respectively, joint and equal consideration of these diagnostic measures would suggest that MoCA <22 appears to have the optimal diagnostic accuracy overall. Table 4 gives the estimated mean difference in sensitivity (top right) and specificity (bottom left) between each of the tests. In comparison to MoCA <22, the estimated sensitivity gained by receiving the optimal true positive test (MoCA <26) is 0.14 with corresponding 95% CrI (0.07, 0.25). Similarly, the estimated specificity gained from receiving the optimal true negative test (MMSE <25) is 0.07 with corresponding 95% CrI (−0.01, 0.18). As these intervals are close to, or span, the point of no difference, there is no strong evidence to suggest that MoCA <22 loses efficiency in accurate diagnosis.

## 6. Discussion

In this study, we propose a unified network meta-analysis framework for synthesizing diagnostic test accuracy data, which allows for both the incorporation of multiple tests and multiple explicit thresholds of the same tests. We further developed this framework to incorporate constraints on increasing test thresholds such that estimates for higher test thresholds had an increased sensitivity but decreased specificity compared to lower thresholds of the same test. In this way, by departing from the usually conducted separate analyses of different tests, the combined analysis of multiple tests/thresholds allows for more detailed and rigorous comparisons between competing tests. For example, in the dementia example presented, we found that MoCA <26 had the optimum true positive rate, and MMSE <25 had the optimum true negative rate of diagnosing cognitive impairment following a stroke. While joint and equal consideration of sensitivity and specificity suggested that MoCA <22 appeared to have the optimal diagnostic test accuracy overall.

From a decision makers', clinicians', and patients' perspective, interest lies in both the true positive rate (sensitivity) and true negative rate (specificity) of a diagnostic test

to efficiently manage patient care. Therefore, unlike the analysis considered here, consideration to the relative weighting of sensitivity and specificity will be required (i.e., the relative health implications of a false negative compared to a false positive result). To make robust decisions regarding the optimal diagnostic test in terms of clinical effectiveness and/or cost-effectiveness, a fully comprehensive clinical or economic decision model will need to be developed incorporating potential treatment plans and longer term follow-up.

Further extensions to the network meta-analysis framework described in this study could include incorporating meta-regression methods. Both observation and study-level covariates could easily be included in the bivariate component of the model as described by Reitsma et al. [8], with the aim to explain some of the heterogeneity between observations or between studies. Further work could also look to derive a set of inconsistency equations to assess consistency between different sources of information, i.e., comparative vs. noncomparative studies.

In clinical practice, a sequential approach to testing is often used. Thus, stroke patients may receive a brief diagnostic test initially, and those identified as remaining “at risk” may receive further, more comprehensive, testing such as NPB or clinical diagnosis. In this example, it may be reasonable to administer MoCA <26 initially to ensure that the optimal number of true negative patients are identified and discharged from further routine cognitive assessment. Alternatively, MoCA <22 may be used initially to ensure that the maximum number of true positive and true negative patients are identified. The utility of a staged and triaged approach to diagnostic testing in this setting remains unanswered and provides an opportunity for further work, as does the further development of statistical methods to evaluate the performance of sequences of tests taking lack of dependence into account [4,24].

A potential limitation of our approach is that it treats different test and threshold combinations as separate tests for the purposes of the analysis, and thus, full sROC plots are not estimated across different thresholds, as has been done elsewhere [3,10–14]; however, methods to estimate full sROCs have not taken into account explicit threshold values until very recently [14,25] or allow for constraints to be placed on the heterogeneity, which can be attributed to threshold differences. Combining a comparative framework such as the



**Table 5.** Approaches to synthesizing diagnostic test accuracy data of multiple tests

| Reference  | Model description  | Type of model  | Available data  | Number of tests   | Multiple thresholds per test | Imperfect GS |
|--|--|----------------|---|---|------------------------------|--------------|
| Trikalinos, T. A., (2014). Research Synthesis Methods [26].                      | Multinomial model approximated by multivariate normal distribution, modeling the joint TPR and FPR. Correlations between tests are modeled as random parameters.   | Arm-based      | Full cross-tabulations  | Two index tests + GS  | No                           | No           |
| Menten, J. and Lesaffre, E. (2015). BMC Medical Research Methodology [16].       | Hierarchical model that is partly based on contrasts between transformed sensitivity and specificity (similar to that of NMA for interventions)<br><br>Adds in allowance for imperfect GS by modeling response pattern across tests as multinomial: latent class models  | Contrast-based | Full cross-tabulations  | Three index three tests + multiple GS                               | No                           | Yes          |
| Hoyer, A. and Kuss, O. (2016). Statistical Methods in Medical Research [27].     | Multinomial model similar to that of Trikalinos et al. [21] but does not account for correlations between tests because full cross-tabulations are not used. This model can be extended to account for multiple test thresholds.   | Arm-based      | 2 × 2 tables for each test vs. GS only  | Two index tests + GS from each study                                | Yes                          | No           |
| Nyaga, V. N., et al. (2016a,b). Statistical Methods in Medical Research [28,29]. | Two-stage hierarchical model based on <i>logit</i> transformed sensitivity and specificity. Shared random effects are specified to induce study level correlations.<br><br>One-stage approach modeling directly on the probability scale (using beta-binomial distribution) without <i>logit</i> transformation. | Arm-based      | 2 × 2 tables for each test vs. GS only  | 11 tests (in example data set) Between one and six tests per study. | No                           | No           |
| Dimou, N. L., et al. (2016). Statistics in Medicine [30].                        | Multivariate normal distribution with closed form formulae for the within-study covariance matrix (needs full cross-tabulations). Full reporting then used to impute when only 2 × 2 table information is presented. Correlation between tests are estimated and “plugged in”.                                   | Arm-based      | Full cross-tabulation and 2 × 2 tables for each test vs. GS only                              | Two index tests + GS (Not all studies have to report both tests)    | No                           | No           |
| Cheng, W. (2016). Repository Library, Brown University (Doctoral thesis) [31]    | Multinomial model with decomposition of test and study-specific effects (Chapter 2)<br>Multivariate extension of the HSROC model (chapter 3)<br>Beta-binomial marginal and multivariate Gaussian copulas (chapter 4)<br>All models account for study-type specific effects and within study-type random effects  | Arm-based      | Full cross-tabulations, partially crossed-tabulations, 2 × 2 tables for each test vs. GS only | Three index tests + GS  | No<br><br>Yes<br><br>No      | No           |

Abbreviation: GS, Gold standard; TPR, true positive rate; FPR, false positive rate; NMA, network meta-analyses; HSROC, hierarchical summary receiver operating characteristic.

model described in this study, with the flexible approach of modeling the distributional form of multiple test thresholds such as the model described by Hoyer et al. [14,25], whereby both the model parameters and distributional forms could be estimated simultaneously, is an opportunity for further work.

Further, if studies do not report the threshold values used or the thresholds cannot be explicitly expressed, as is often the case for tests involving the interpretation of some sort of diagnostic image etc., then the performance of a test at different thresholds cannot be estimated via the model presented. However, this is not an argument for not using the valuable extra information when thresholds are known, and more generally there is no reason why the same methodology should be used in these two different contexts (despite this being the case historically).

Table 5 describes the similarities and differences between a number of additional approaches in the current literature to synthesize diagnostic test accuracy data from multiple tests [16,26–31]. All of these methods extend the bivariate model of Reitsma et al. [8] to synthesize data for two or more tests. Trikalinos et al. [26], Hoyer and Kuss [27], and Cheng [31] make use of multinomial distributions to model the within-study variation for multiple tests, whereas Dimou et al. [30] make use of multivariate normal distributions on *logit* sensitivities and specificities to account for within-study covariances. If full cross-tabulations, i.e., the full response array across all competing tests, are available, then the correlation between tests can be taken in to account as in Trikalinos et al. [26], Dimou et al. [30], and Cheng [31]. These approaches have the advantage of appropriately modeling the within-study covariance structure; however, partial or full cross-tabulations are required for a sufficient number of studies to adequately model the correlation between tests [31]. A limitation of these approaches is that as the number of competing tests increases, the number of parameters to be estimated by the model rapidly increases, which can result in issues with model convergence [31]. Cheng [31] further considers a multivariate extension of the Hierarchical Summary Receiver Operating Characteristic (HSROC) model by Rutter and Gatsonis [7] and explores the use of beta-binomial marginal and multivariate Gaussian copulas. The author found that use of the beta-binomial marginal and multivariate Gaussian copulas produced less biased estimates of the posterior mean summary points compared to the multivariate extension of the bivariate model and HSROC model; however, this approach appeared to be computationally expensive. Nyaga et al. [28,29] use a two-stage hierarchical model based on the *logit* transformed sensitivity and specificity [28], and a one-stage approach based on the beta-binomial distribution modeling on the probability scale [29]. Both models include shared random effects to induce study level correlations. This approach is most similar to the model we describe in this study. However, our model further incorporates shared random effects on test to account for multiple thresholds and applies constraints on increasing test thresholds. Similarly to the model outlined in this study, many of

the approaches described in Table 5 adopt an arm-based approach and model the absolute measures of test accuracy [26–31]. Menten and Lesaffre [16] developed a contrast-based approach, which models both the direct (or head-to-head) comparisons of diagnostic tests as well as indirect comparisons through a common diagnostic test. This approach directly models the relative *logit* sensitivities and specificities between multiple tests. This model can be further extended to a hierarchical latent class model to account for imperfect gold standards. A contrast-based approach works well if all studies evaluate all diagnostic tests. In the case of mixed reporting, further assumptions regarding the missingness of data are required. In a comparison between the contrast-based approach of Menten and Lesaffre [16] and a hierarchical arm-based approach of Nyaga et al. [28], Nyaga et al. [28] argue that an arm-based approach may be more appealing than contrast-based approaches because it allows for a more straightforward interpretation of the parameters, makes use of all available data resulting in increased precision, and adopts a more natural variance-covariance structure. In choosing an approach to synthesize diagnostic test accuracy data for multiple tests, the user needs to consider a number of factors related to the decision question. The first of which is the available data; if partial or full cross-tabulations are available for a sufficient number of studies, a multivariate approach may be preferred to adequately model the within-study correlation structures [16,26,30,31]. If there are a number of test thresholds of interest, a multivariate approach to the HSROC model [31] or an extension to the multinomial approach as described by Hoyer and Kuss [27] may be most appropriate. However, if there are many competing diagnostic tests, the number of additional parameters to be estimated by a multivariate model may be too large causing issues with model convergence. In this instance, a hierarchical model [28,29], including the model described in this study, may be preferred. Furthermore, in the case of many competing tests and multiple test thresholds, our approach using shared random effects and constraints on increasing test thresholds may be considered. Indeed, a comprehensive comparison of all of the approaches described in Table 5, including an assessment of model performance under different criteria and different reporting structures, is still required and would be a valuable addition to the current literature.

In this study, we evaluated a number of models each with different assumptions regarding the heterogeneity parameters and correlation parameters. The first model assumed common heterogeneity and correlation parameters across tests, the second model assumed common heterogeneity and test-specific correlation parameters, and the third model assumed test-specific heterogeneity parameters and a common correlation parameter. It is worth noting that allowing both heterogeneity and correlation parameters to be test-specific leads to unidentifiability of the covariance matrix, and thus causes issues with model convergence.

In conclusion, this study proposes a number of network meta-analysis models for synthesizing diagnostic test accuracy data. The proposed frameworks allow for the analysis of multiple tests at multiple thresholds together with the option to incorporate constraints on increasing test thresholds. It could be argued that constraints on threshold effects should be applied to all models with explicit threshold information regardless of model fit due to the implicit threshold assumption which by definition must be satisfied (but is not imposed by previous models commonly fitted). Incorporating this information through the use of constraints has the potential to more appropriately attribute variability between results to (genuine) threshold effects and better explain heterogeneity between studies.

## Supplementary Data

Supplementary data related to this article can be found at <https://doi.org/10.1016/j.jclinepi.2018.03.005>.

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