Meta-analysis: the diagnostic accuracy of critical flicker frequency in minimal hepatic encephalopathy

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Publication data
Submitted 13 August 2012
First decision 6 September 2012
Resubmitted 5 December 2012
Accepted 11 December 2012
EV Pub Online 7 January 2013

As part of AP&T’s peer-review process, a technical check of this meta-analysis was performed by Dr Y. Yuan.

SUMMARY

Background
Minimal hepatic encephalopathy (MHE) reduces quality of life, increases the risk of road traffic incidents and predicts progression to overt hepatic encephalopathy and death. Current psychometry-based diagnostic methods are effective, but time-consuming and a universal ‘gold standard’ test has yet to be agreed upon. Critical Flicker Frequency (CFF) is a proposed language-independent diagnostic tool for MHE, but its accuracy has yet to be confirmed.

Aim
To assess the diagnostic accuracy of CFF for MHE by performing a systematic review and meta-analysis of all studies, which report on the diagnostic accuracy of this test.

Methods
A systematic literature search was performed to locate all publications reporting on the diagnostic accuracy of CFF for MHE. Data were extracted from 2 × 2 tables or calculated from reported accuracy data. Collated data were meta-analysed for sensitivity, specificity, diagnostic odds ratio (DOR) and summary receiver operator curve (sROC) analysis. Prespecified subgroup analysis and meta-regression were also performed.

Results
Nine studies with data for 622 patients were included. Summary sensitivity was 61% (95% CI: 55–67), specificity 79% (95% CI: 75–83) and DOR 10.9 (95% CI: 4.2–28.3). A symmetrical sROC gave an area under the receiver operator curve of 0.84 (SE = 0.06). The heterogeneity of the DOR was 74%.

Conclusions
Critical Flicker Frequency has a high specificity and moderate sensitivity for diagnosing minimal hepatic encephalopathy. Given the advantages of language independence and being both simple to perform and interpret, we suggest the use of critical flicker frequency as an adjunct (but not replacement) to psychometric testing.

Aliment Pharmacol Ther 2013: 37: 527–536
INTRODUCTION

Minimal hepatic encephalopathy (MHE) is an important clinical variant of hepatic encephalopathy (HE), which occurs in up to 60–70% of patients with cirrhosis.\(^1\)\(^,\)\(^2\) The condition comprises a cognitive impairment, observed in patients with cirrhosis who have no clinical evidence of overt hepatic encephalopathy (OHE).\(^3\) It is associated with an increased incidence of road traffic accidents,\(^4\)\(^–\)\(^7\) reduced quality of life and it affects the ability to perform tasks of daily living.\(^8\)\(^,\)\(^9\) It has also been shown to increase the risk of progressing to OHE and inversely correlates with survival in some studies.\(^10\)\(^–\)\(^12\).

Treatment for MHE can improve psychometric performance and health-related quality of life.\(^13\)\(^–\)\(^17\) It is therefore clinically useful that MHE is diagnosed effectively in patients with this condition. A quick, accurate, objective, cost-effective and well-validated diagnostic test is an unmet clinical need and would simplify the early management algorithm for this condition.

Minimal hepatic encephalopathy is not routinely tested, even in specialised cirrhosis clinics. A survey among members of the American Association for the Study of Liver Diseases (AASLD) showed that 72% tested less than half their patients for MHE, despite 84% acknowledging that MHE is a significant problem. Furthermore, 85% said that if clinical staff could perform a quick, accurate test, this would increase the likelihood that people would test for MHE.\(^18\)

There are many diagnostic tests for MHE, but no universal ‘gold standard’ test. The Expert Working Group in 1998 suggested that the psychometric hepatic encephalopathy score (PHES) should be considered the gold standard test.\(^19\) PHES is a selection of five psychometric test batteries that has been validated in Italian,\(^20\) German\(^21\)\(^,\)\(^22\) and Spanish\(^23\) cohorts. However, performing this battery can be time consuming and prone to bias from disturbance, mood and interaction with the tester.

Computer-assisted tests have also been used to diagnose MHE and include the Inhibitory Control Test (ICT),\(^24\)\(^,\)\(^25\) the cognitive drug research system (CDRS),\(^26\) the continuous reaction time test (CRT)\(^27\)\(^,\)\(^28\) and Critical Flicker Frequency (CFF).\(^29\) The CFF has the purported advantages of not being dependent on language, verbal fluency, numeracy or numerics, and therefore studies into its use have been performed in the United States, Europe and Asia.

The CFF was devised originally as an ophthalmological test used to measure visual acuity and to screen for optic nerve lesions.\(^30\) This test measures the frequency at which the patient perceives that a fused/single light becomes a flickering light. The device causes a stepwise decrease in frequency from 60 to 25 Hz. This is done multiple times (usually 8–10) to allow calculation of the mean and standard deviation. It may also be performed in reverse, where the patient determines the frequency at which a flickering light becomes continuous or ‘fused’. This test has the advantage of being able to be carried out by clinical personnel using a portable device with limited running costs.\(^31\)

The CFF has been in limited clinical and research use for a decade, but its diagnostic accuracy has never been subjected to quantitative review. We therefore performed a systematic review and meta-analysis of CFF to fully assess its diagnostic accuracy in detecting MHE and to guide future implementation.

MATERIALS AND METHODS

Search methodology

We searched MEDLINE and EMBASE using OvidSP for articles between January 1948 and November 2012, which reported on the diagnostic accuracy of CFF for MHE. The search terms used were ‘Critical Flicker Frequency’ OR ‘CFF’ in conjunction (AND) with ‘Diagnostic Accuracy’ OR MeSH term ‘Diagnostic techniques and procedures’ OR ‘Sensitivity’ OR ‘Specificity’. Inclusion criteria were adult studies reporting the diagnostic accuracy for CFF in patients with cirrhosis and/or portosystemic bypass. Paediatric studies, studies not related to MHE in the context of cirrhosis or portosystemic bypass, those which did not refer to a gold standard and those not in the English language were excluded. We also obtained primary sources from tracking references from hand searches in review papers and original articles. Only original data were used in the meta-analysis.

Data extraction

Test performance data were extracted as a 2 × 2 table of true negative, true positive, false positive and false negative values directly from tabulated results. If these were not available, they were calculated from reported sensitivity, specificity, positive predictive value and/or negative predictive values; if this was not possible, the authors were contacted for more detailed data; and if this was not possible or there was doubt over the 2 × 2 calculation, the study was excluded from subsequent analysis.

Assessment of study quality

Studies meeting the above criteria were quality assessed using positive scoring in a modified 23-point Standards
for the Reporting of Diagnostic Accuracy Studies (STARD) pro forma. We modified this scoring system from 25 to 23 as two points were not relevant to this study (points 18 and 20). Two readers (FT and MM) independently assessed all included studies according to the prearranged pro forma. An open discussion was then held to determine any disagreement between the readers. Studies were then segregated into ‘low’ or ‘high’ quality depending on whether they met < or > than 50% of the study quality pro forma (Low <12/23, High ≥ 12/23).

Data analysis
The DerSimonian-Laird random effects method was used to produce summary estimates of sensitivity, specificity, likelihood ratios (LR) and diagnostic odds ratio (DOR). Confidence intervals for sensitivity and specificity were calculated using F-distribution method for the binomial proportion. The summary receiver operator curve (sROC) was used to graphically determine performance following testing for correlation between sensitivity and specificity [as logit true positive rate (TPR) vs. logit false positive rate (FPR)] to explore for threshold effects and subsequent assessment for constant DOR using the Moses-Shapiro-Littenburg model. Symmetrical or asymmetrical sROC were used depending on whether the DOR is constant. Heterogeneity was investigated using pre-planned subgroup analysis and calculated by the I² method. Pre-planned subgroups were defined according to: study quality (low or high with 50% quality pro forma cut-off used), type of gold standard test (PHES or non-PHES), study location (Europe or non-European), CFF cut-off (≤ 38 or ≥ 39), whether the study was published in early era or late era (median study year from extracted studies was used as the cut-off), number of patients in the respective studies (<50 subjects or ≥ 50 subjects) (50 = median) and the aetiology of the MHE (cirrhosis or bypass/shunts).

A funnel plot and effective sample size (ESS) regression analysis (the logarithm of the DOR plotted against 1/√ESS) was used to investigate publication bias. 1/√ESS is proportional to root (1/n1 + 1/n2) where n1 = number diseased and n2 number not diseased. Data analyses were performed using the freeware Meta-Disc version 1.4 (Universidad Complutense, Madrid, Spain).

RESULTS
The search strategy identified 265 studies, 209 were excluded, based on title and abstract, while the remaining 56 were read and evaluated. Forty seven were further excluded, based on prestated criteria leaving 9, which were included in the final meta-analysis (Figure 1).
Data from 622 patients were available. Four of nine studies were published in the early era (2002–2008) and five were published in the late era (2009–2011). Five studies were deemed to be of low quality (<12/23) and the other four studies were deemed to be of high quality (≥12/23), according to the predefined pro forma.32, 33 Four studies used PHES as their reference test, whereas five studies used non-PHES as reference tests. Four studies were conducted in Europe, whereas five were performed elsewhere. Six studies used a CFF cut-off value of ≤38 Hz to distinguish an abnormal test, whereas the other three used a cut-off value of ≥39 Hz. The median (range) number of participants was 50 (31–114). Four studies had less than 50 patients and five studies had 50 or more patients.

We grouped studies into MHE caused by cirrhosis and MHE caused by bypass/shunting. Cirrhosis was the cause of the MHE in seven studies, while bypass/shunting (secondary to extra-hepatic portal venous obstruction [EHPVO]) was the cause in the other two.

Finally, we were unable to investigate different aetiologies of cirrhosis (such as alcohol or viral hepatitis) in our meta-analysis, as it was not possible to extract separate 2 × 2 tables for these aetiologies from the studies.

Sensitivity, specificity and diagnostic odds ratio
The pooled sensitivity for all nine studies included in the final meta-analysis was 61% (95% CI: 55–67%), pooled specificity was 79% (95% CI: 75–83%), pooled positive LR was 3.5 (95% CI: 2.0–6.1), pooled negative LR was 0.46 (95% CI: 0.31–0.68) and the pooled DOR was 10.9 (95% CI: 4.2–28.3). (Table 1, Figure 2, Figures S1 and S2).

The Spearman correlation coefficient for logit (TPR) vs. logit (FPR) was 0.38 (P = 0.31), indicating a nonsignificant correlation between the TPR and FPR. The Moses-Shapiro-Littenberg method showed that the DOR was constant [b = −0.018 (P = 0.95)]. A symmetrical sROC was therefore the most appropriate representation of the diagnostic accuracy (Figure 3). It depicted an area under the receiver operator curve (AUROC) (SE) of 0.84 (0.06) and with a Q statistic (SE) of 0.77 (0.06).

Heterogeneity analysis & subgroup analysis
Heterogeneity was detected in all pooled indices. For all nine studies, the heterogeneity (I²) was 87% (sensitivity), 85% (specificity), 72% (positive LR), 85% (negative LR), 74% (DOR).

Subgroup analysis was performed to assess differences in heterogeneity and diagnostic accuracy between the prespecified groups (Table 2). The heterogeneity of the DOR was lower in the high-quality studies (I², 0% vs. 85%). There was less heterogeneity in studies, which used a non-PHES gold standard test (I², 5% vs. 88%). We also found that there was less heterogeneity in the European studies, compared with non-European studies (I², 45% vs. 72%). There was lower heterogeneity in the smaller studies (I², 17% vs. 77%), in the studies with a CFF cut-off ≥39 (I², 15% vs. 82%) and in the studies published in the early era (I², 40% vs. 78%).

Critical Flicker Frequency was more diagnostically accurate in the European studies, DOR = 27.3 (95% CI: 8.2–91.2) compared with a DOR = 5.1 (95% CI: 1.6–15.9) for non-European studies. CFF was also more accurate in diagnosing MHE in studies performed in the early era DOR = 26.6 (95% CI: 7.3–97.4) compared with the late era DOR = 5.9 (95% CI: 1.8–19.4). Smaller studies (<50 patients) had a greater DOR = 32.5 (95% CI: 9.3–113.3), compared with the bigger studies (≥50 patients) DOR = 6.0 (95% CI: 2.1–17.2). There was little difference in the DOR among the quality (low/high), cut-off (≤38/≥39) aetiology (bypass/cirrhosis) and the gold standard (non-PHES/PHES) subgroups. To assess if there was any significant correlation between any of these co-variables and the diagnostic accuracy of CFF (DOR), meta-regression analysis was performed.

Meta-regression & Publication bias
There was no significant correlation between any of the co-variables and the DOR in the univariate meta-regression analysis (Table 3). Owing to only nine studies being included in this meta-analysis, the power of multivariate meta-regression is low and thus this limits the overall value of meta-regression in this meta-analysis.

No significant publication bias was found in our study sample, as the linear regression analysis indicates P = 0.11 (Figure 4).

DISCUSSION
The importance of MHE as a complication of cirrhosis or portosystemic bypass has only been recognised in the last decade and its profound effect on patients with cirrhosis should not be underestimated. Neither, unfortunately, should clinicians’ poor understanding of MHE and hence their reluctance to test for its presence.18 Added to this, there are a number of diagnostic options, but many clinicians are confused as to which to choose. In the absence of other easily implementable clinical alternatives, the PHES battery has been suggested as the current gold standard internationally for MHE diagnosis,19 but patient numeracy, literacy or language skills
| Study name          | Sensitivity (95% CI)/% | Specificity (95% CI)/% | Positive LR (95% CI) | Negative LR (95% CI) | DOR (95% CI) | TP | FP | FN | TN | Reference test | STARD score |
|---------------------|------------------------|------------------------|----------------------|----------------------|--------------|----|----|----|----|----------------|-------------|
| Kircheis 2002       | 56 (35–76)             | 100 (86–100)           | 29.0 (1.8–461.1)     | 0.45 (0.29–0.70)     | 64.3 (3.5–1173.2) | 14 | 0  | 11 | 25 | A               | 14          |
| Romero-Gomez 2007   | 77 (60–90)             | 73 (62–83)             | 2.9 (1.9–4.4)        | 0.31 (0.17–0.58)     | 9.3 (3.7–23.7)  | 27 | 21 | 8  | 58 | PHES           | 14          |
| Montoliu 2007       | 87 (60–98)             | 92 (73–99)             | 10.4 (2.7–39.8)      | 0.15 (0.04–0.53)     | 71.5 (9.0–570.3) | 13 | 2  | 2  | 22 | PHES           | 6           |
| Sharma 2008         | 58 (28–85)             | 100 (85–100)           | 26.5 (1.6–428.1)     | 0.43 (0.23–0.82)     | 61.4 (3.0–1245.8) | 7  | 0  | 5  | 22 | B               | 11          |
| Montoliu 2009       | 88 (62–98)             | 88 (68–97)             | 7.0 (2.4–20.5)       | 0.14 (0.04–0.53)     | 49.0 (7.2–331.8) | 14 | 3  | 2  | 21 | PHES           | 7           |
| Dhiman 2010         | 35 (22–51)             | 92 (82–98)             | 4.6 (1.7–12.7)       | 0.70 (0.56–0.88)     | 6.6 (2.0–21.4)  | 17 | 4  | 31 | 48 | C               | 13          |
| Goel 2010           | 21 (5–51)              | 94 (71–100)            | 3.6 (0.4–31.3)       | 0.84 (0.62–113)      | 4.4 (0.4–47.6)  | 3  | 1  | 11 | 16 | D               | 6           |
| Sharma 2010         | 85 (73–93)             | 58 (43–72)             | 2.0 (1.4–2.9)        | 0.26 (0.14–0.49)     | 7.8 (3.2–19.3)  | 51 | 21 | 9  | 29 | E               | 12          |
| Maldonado-Garza 2011| 36 (20–55)             | 66 (54–77)             | 1.1 (0.6–1.9)        | 0.96 (0.71–1.31)     | 1.1 (0.5–2.7)   | 12 | 24 | 21 | 47 | PHES           | 8           |
| Pooled              | 61 (55–67)             | 79 (75–83)             | 3.5 (2.0–6.1)        | 0.46 (0.31–0.68)     | 10.9 (4.2–28.3) |    |    |    |    |                 |             |

A – Computerised psychometry.
B – NCT A and B or FCT A and B, and/or abnormal P300 ERP.
C – NCT-A, FCT-A, SDT, DST, LTT for time and for error.
D – NCT A and B, FCT A and B, WAIS-P tests (PC, DS, PA, OA, BD).
E – NCT A and B or FCT A and B, and abnormal P300 ERP.
may affect the result.\textsuperscript{47} CFF has been proposed as being unaffected by these issues, but its diagnostic accuracy has never been analysed quantitatively before.

There are a number of broad themes that emerged from the meta-analysis. CFF only had a moderate pooled sensitivity of 61\% (95\% CI: 55–67), but a good specificity of 79\% (95\% CI: 75–83). The symmetrical sROC curve had an AUROC of 0.84, indicating that CFF was effective in discriminating patients with MHE from those without MHE and therefore has potential as a screening tool, either to be used prior to current psychometrics tools, or to be used alongside these tests, but not as a replacement for them given the risk of false negative results.

We used subgroup analysis to compare CFF diagnostic accuracy in MHE caused by portosystemic bypass/shunting (DOR = 13.9), compared with MHE caused by cirrhosis (DOR = 10.6). Further analysis shows that CFF has high specificity of 97\% (95\% CI: 87–100) for MHE in patients with bypass/shunting (secondary to EHPVO),
but a low sensitivity of 39% (95% CI: 20–59). Further studies are needed to inform on whether CFF is effective for diagnosing MHE caused by bypass/shunting, and whether the pathogenesis of MHE in this condition predetermines the best diagnostic modality to choose. It may be that the neurophysiological impairment of the high ammonia states in portosystemic bypass/shunting compared the potentially lower levels of hyperammonaemia, but higher rates of inflammation in patients with cirrhosis affect the performance of CFF in these groups.48, 49

| Table 2 | Assessment of diagnostic accuracy and heterogeneity in subgroup analysis |
|---------|---------------------------------------------------------------|
| Sub groups | No. of studies | Pooled sensitivity (95% CI)/% | Pooled specificity (95% CI)/% | Pooled positive LR (95% CI) | Pooled negative LR (95% CI) | Pooled DOR (95% CI) | $I^2$ (%) DOR |
| All studies | 9 | 61 (55–67) | 79 (75–83) | 3.5 (2.0–6.1) | 0.46 (0.31–0.68) | 10.9 (4.2–28.3) | 74 |
| Era | | | | | | | |
| Early (2002–2008) | 4 | 70 (59–80) | 85 (78–90) | 8.8 (2.0–38.2) | 0.37 (0.26–0.54) | 26.6 (7.3–97.4) | 40 |
| Late (2009–2011) | 5 | 57 (49–64) | 75 (69–81) | 2.6 (1.4–4.9) | 0.58 (0.37–0.91) | 5.9 (1.8–19.4) | 78 |
| Quality assessment | | | | | | | |
| Low | 5 | 54 (44–65) | 81 (74–87) | 5.0 (1.3–18.8) | 0.47 (0.25–0.91) | 14.1 (1.7–117.4) | 85 |
| High | 4 | 65 (57–72) | 78 (71–83) | 2.9 (1.7–4.9) | 0.42 (0.24–0.73) | 8.7 (5.0–15.1) | 0 |
| Gold standard | | | | | | | |
| Non-PHES | 5 | 58 (50–66) | 84 (78–90) | 4.9 (1.7–14.3) | 0.53 (0.35–0.79) | 8.8 (4.4–17.6) | 5 |
| PHES | 4 | 67 (57–76) | 75 (68–81) | 3.3 (1.4–8.0) | 0.31 (0.10–1.00) | 11.7 (1.8–74.5) | 88 |
| Location | | | | | | | |
| Non-Europe | 5 | 54 (46–62) | 76 (70–82) | 2.4 (1.2–4.6) | 0.63 (0.44–0.91) | 5.1 (1.6–15.9) | 72 |
| Europe | 4 | 75 (65–83) | 83 (76–89) | 6.3 (2.3–17.2) | 0.29 (0.16–0.51) | 27.3 (8.2–91.2) | 45 |
| No. of patients | | | | | | | |
| <50 | 4 | 65 (51–77) | 93 (86–97) | 8.1 (3.8–17.1) | 0.32 (0.10–1.08) | 32.5 (9.3–113.3) | 17 |
| ≥50 | 5 | 60 (53–67) | 75 (69–80) | 2.4 (1.4–4.0) | 0.51 (0.32–0.80) | 6.0 (2.1–17.2) | 77 |
| Cut-off | | | | | | | |
| ≤38 | 6 | 57 (49–64) | 80 (75–85) | 4.0 (1.9–8.5) | 0.44 (0.25–0.75) | 12.0 (3.1–45.6) | 82 |
| ≥39 | 3 | 69 (59–78) | 76 (66–84) | 4.1 (0.8–20.6) | 0.48 (0.21–1.06) | 9.1 (3.2–25.6) | 15 |
| Aetiology of MHE | | | | | | | |
| Bypass | 2 | 39 (20–59) | 97 (87–100) | 8.2 (1.1–59.8) | 0.63 (0.29–1.37) | 13.9 (1.0–186.4) | 46 |
| Cirrhosis | 7 | 64 (57–70) | 77 (72–81) | 3.2 (1.9–5.6) | 0.40 (0.25–0.67) | 10.6 (3.7–30.5) | 79 |

We found that the studies published in the early era had a higher DOR than studies published in the late era (DOR: 26.6 vs. 5.9). Although this difference was not significant on univariate meta-regression, it does question whether the high DOR seen in early studies29 might represent an overestimation of diagnostic accuracy, particularly when the test is applied to heterogeneous patient cohorts. We also found that the diagnostic accuracy was
higher in studies published in Europe (DOR: 27.3 vs 5.1). Again, this showed no significance on meta-regression analysis, but may represent an early indicator that CFF may be more accurate in European patients. It remains unclear why this objective, language-independent test would perform better in these subgroups, but it is an important observation and does have implications on the role of CFF as a worldwide screening tool for MHE, particularly when the sensitivity is only 54% in the non-European, compared with 75% in the European subgroups.

We explored the possibility that early studies outperformed later studies due to the problem of publication bias, and we acknowledge that it can be a significant problem with diagnostic accuracy meta-analysis, but we found no statistical evidence of this across our study sample. We are also aware that four of the five non-European studies were published in India and two of them by the same centre. Further studies are regarding the effect of location and race on this diagnostic test, particularly in non-European countries outside of India.

The considerable amount of heterogeneity detected between the studies suggests a need for caution when pooling the diagnostic accuracy measures together. We used subgroup analysis to assess the heterogeneity in prespecified groups, and unsurprisingly found that it was reduced in the 'high quality' studies. The implicit and explicit causes of threshold effect were assessed as another cause of heterogeneity. Most studies used either 38 Hz or 39 Hz as the flicker frequency cut-off value to discriminate between patients with or without MHE; when this was assessed in the meta-regression analysis, there was no correlation between this value and the DOR. We further assessed the implicit causes of threshold effect by calculating the Spearman correlation, which was 0.38 (P = 0.31). While there was no statistical evidence of a threshold effect, a summary ROC (sROC) remains a useful composite measure of the diagnostic accuracy of CFF. The Moses model showed the DOR to be constant, so we investigated further using a symmetrical sROC confirming the good overall accuracy of CFF.

One limitation of this meta-analysis is that the nine studies included had to compare CFF to a reference test. This provides two problems: first, the reference test is not the same for all nine studies; and second, the diagnostic accuracy of this test may be less than 100%. Within the subgroup analysis, we looked at studies that compared CFF with PHES (the current suggested diagnostic gold standard) and studies that referenced to another non-PHES test (Table 1). Attempts were made to assess differences in heterogeneity and diagnostic accuracy between these two groups. The diagnostic accuracy for CFF was slightly higher in the PHES subgroup (DOR: 11.7 vs. 8.8), but also significantly more heterogeneous (DOR: 88% vs. 5%). Although there is some difference in diagnostic accuracy, the meta-regression analysis showed no significance. We can, therefore, cautiously conclude from this meta-analysis that CFF seems to perform comparably to PHES and non-PHES diagnosed MHE. The increase in heterogeneity observed may thus be explained to some extent.

One further limitation common to all diagnostic meta-analyses is the lack of clarity, quality and standardisation in diagnostic studies’ methodology. Studies were assessed for quality using STARD pro forma to quantify the methodology of the study design. We had to exclude two studies at a late stage due to ambiguity between the raw data and the diagnostic accuracy data, which could not be resolved with the authors. Furthermore, it should be noted that Maldonado-Garza and colleagues’ study, for example, was not intended to be a diagnostic study, yet included sufficient data to enable its inclusion in this meta-analysis. We would encourage any further studies designed to assess the diagnostic accuracy of CFF to follow either the STARD or PRISMA checklist.

Despite the limitations, we know that CFF is a simple, affordable test, which is easy to perform. It is thus realistic for departments specialising in the management of patients with cirrhosis or portosystemic bypass/shunting, who are at high risk of MHE, to buy and use this device. The test does not require specialist personnel to conduct and is extremely well tolerated and easily understood by the patients within the studies. The only limitation of its use in the nine studies was in Romero-Gomez and colleagues’ study where nine patients and three controls could not perform the test due to visual impairment or inability to understand the fundamentals involved.

Key to the management algorithm of MHE is a correct, early diagnosis, prompting early effective treatment. Many of the diagnostic tests available at present are time consuming and require trained personnel to perform them. Furthermore, the lack of universal consensus on which diagnostic methods and strategies should be implemented heightens the problem of under-diagnosis and as demonstrated, no uniformly implemented international gold standard diagnostic testing criteria for MHE exist. This meta-analysis has shown
that CFF is a diagnostically accurate test, which could be used as an adjunct to conventional psychometric test batteries, such as PHES, but could only become a replacement screening test if further studies show an improvement in sensitivity.

AUTHORSHIP

Guarantor of the article: S. D. Taylor-Robinson.

Author contributions: All authors contributed to the design of the study and writing of the manuscript. FT and MJM undertook the research and performed the analyses. All authors approved the final version of the manuscript.

ACKNOWLEDGEMENTS

We are also grateful to Mary M. E. Crossey, Sebastiana Atzori, Louise Campbell, Yasmin Pasha and Heather Marcinkowski of Imperial College Healthcare Trust Liver Unit at St Mary’s Hospital London for useful discussions on the manuscript. Declaration of personal interests: None.

Declaration of funding interests: All authors are grateful to the NIHR Biomedical Facility at Imperial College London for infrastructure support. MJWM is supported by the Wellcome Trust, UK. SDT-R is supported by grants from the Wellcome Trust, the British Medical Research Council, United Kingdom National Institute for Health Research (NIHR) and the Alan Morement Memorial Fund.

SUPPORTING INFORMATION

Additional Supporting Information may be found in the online version of this article:

Figure S1. Forest plot for positive and negative likelihood ratios for the nine studies.

Figure S2. Forest plot for diagnostic odds ratio for the nine studies.

REFERENCES

1. Butterworth RF. Editorial: rifaximin and minimal hepatic encephalopathy. Am J Gastroenterol 2011; 106: 317–8.
2. Poordad FF. Review article: the burden of hepatic encephalopathy. Aliment Pharmacol Ther 2007; 25: 3–9.
3. Amiodo P, Montagnese S, Gatta A, Morgan MY. Characteristics of minimal hepatic encephalopathy. Metab Brain Dis 2004; 19: 253–67.
4. Kircheis G, Knoche A, Hilger N, et al. Hepatic Encephalopathy and Fitness to Drive. Gastroenterology 2009; 137: 1706–15.
5. Wein C, Koch H, Popp B, Oehler G, Schauder P. Minimal hepatic encephalopathy impairs fitness to drive. Hepatology 2004; 39: 739–45.
6. Córdoba J, Lucke R. Driving under the influence of minimal hepatic encephalopathy. Hepatology 2004; 39: 599–601.
7. Bajaj JS, Saeki K, Schubert CM, et al. Minimal hepatic encephalopathy is associated with motor vehicle crashes: the reality beyond the driving test. Hepatology 2009; 50: 1175–83.
8. Schomerus H, Hamster W. Quality of life in cirrhotics with minimal hepatic encephalopathy. Metab Brain Dis 2001; 16: 37–41.
9. Groeneweg M, Quero JC, De Bruijn I, et al. Subclinical hepatic encephalopathy impairs daily functioning. Hepatology 1998; 28: 45–9.
10. Hartmann IJC, Groeneweg M, Quero JC, et al. The prognostic significance of subclinical hepatic encephalopathy. Am J Gastroenterol 2000; 95: 2029–34.
11. Romero-Gómez M, Boza F, García-Valdecasas MS, García E, Aguilar-Reina J. Subclinical hepatic encephalopathy predicts the development of overt hepatic encephalopathy. Am J Gastroenterol 2001; 96: 2718–23.
12. Montagnese S, Biancardi A, Schiﬀ S, et al. Different biochemical correlates for different neuropsychiatric abnormalities in patients with cirrhosis. Hepatology 2011; 53: 558–66.
13. Sidhu SS, Goyal O, Mishra BP, Sood A, Chhina RS, Soni RK. Rifaximin improves psychometric performance and health-related quality of life in patients with minimal hepatic encephalopathy (the REME Trial). Am J Gastroenterol 2011; 106: 307–16.
14. Prasad S, Dhiman RK, Duseja A, Chawla YK, Sharma A, Agarwal R. Lactulose improves cognitive functions and health-related quality of life in patients with cirrhosis who have minimal hepatic encephalopathy. Hepatology 2007; 45: 549–59.
15. Bajaj JS, Saeki K, Christensen KM, et al. Probiotic yogurt for the treatment of minimal hepatic encephalopathy. Am J Gastroenterol 2008; 103: 1707–15.
16. Ndahra S, Hasan I, Simadibrata M. The effect of L-ornithine L-aspartate and branch chain amino acids on encephalopathy and nutritional status in liver cirrhosis with malnutrition. Acta Med Indones 2011; 43: 18–22.
17. Shukla S, Shukla A, Mehboob S, Guha S. Meta-analysis: the effects of gut flora modulation using prebiotics, probiotics and symbiotics on minimal hepatic encephalopathy. Aliment Pharmacol Ther 2011; 33: 662–71.
18. Bajaj JS, Etemadian A, Hafeezullah M, Saeki K. Testing for minimal hepatic encephalopathy in the United States: an AASLD survey. Hepatology 2007; 45: 833–4.
19. Ferenci P, Lockwood A, Mullen K, Tarter R, Weissenborn K, Blei AT. Hepatic encephalopathy-definition, nomenclature, diagnosis, and quantification: final report of the Working Party at the 11th World Congresses of Gastroenterology, Vienna, 1998. Hepatology 2002; 35: 716–21.
20. Amiodo P, Campagna F, Olianas S, et al. Detection of minimal hepatic encephalopathy: normalization and optimization of the psychometric hepatic encephalopathy score. A neuropsychological and quantified EEG study. J Hepatol 2008; 49: 346–53.
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21. Schomerus H, Hamster W. Neuropsychological aspects of portal-systemic encephalopathy. Metab Brain Dis 1998; 13: 361–77.

22. Weissenborn K, Ennen JC, Schomerus H, Rücker N, Hecker H. Neuropsychological characterization of hepatic encephalopathy. J Hepatol 2001; 34: 768–73.

23. Romero Gomez M, Cordoba J, Jover R, et al. Normality tables in the Spanish population for psychometric tests used in the diagnosis of minimal hepatic encephalopathy. Med Clin 2006; 127: 246–9.

24. Bajaj JS, Saean K, Verber MD, et al. Inhibitory control test is a simple method to diagnose minimal hepatic encephalopathy and predict development of overt hepatic encephalopathy. Am J Gastroenterol 2007; 102: 754–60.

25. Bajaj JS, Hafeezullah M, Franco J, et al. Inhibitory control test for the diagnosis of minimal hepatic encephalopathy. Gastroenterology 2008; 135: 1591–600.

26. Mardini H, Saxby BK, Record CO. Computerized psychometric testing in minimal encephalopathy and modulation by nitrogen challenge and liver transplant. Gastroenterology 2008; 135: 1582–90.

27. Elsass P, Christensen SE, Ranek L, Theiligard A, Tystrup N. Continuous reaction time in patients with hepatic encephalopathy. A quantitative measure of changes in consciousness. Scand J Gastroenterol 1981; 16: 441–7.

28. Elsass P, Christensen SE, Mortensen EL, Vilstrup H. Discrimination between organic and hepatic encephalopathy by means of continuous reaction times. Liver 1985; 5: 29–34.

29. Kircheis G, Wettstein M, Timmermann L, Schnitzler A, Haussinger D. Critical flicker frequency for quantification of low-grade hepatic encephalopathy. Hepatology 2002; 35: 357–66.

30. Baatz H, Raak P, de Ortueta D, Mirshahi A, Scharioth G. Practical significance of critical fusion frequency (CFF). Chronological resolution of the visual system in differential diagnosis. Ophthalmologe 2010; 107: 715–9.

31. Bajaj JS. Review article: the modern management of hepatic encephalopathy. Aliment Pharmacol Ther 2010; 31: 537–47.

32. Bossuyt PM, Reitsma JB, Bruns DE, et al. The STARD statement for reporting studies of diagnostic accuracy: explanation and elaboration. Clin Chem 2003; 49: 7–18.

33. Bossuyt PM, Reitsma JB, Bruns DE, et al. Towards complete and accurate reporting of studies of diagnostic accuracy: the STARD initiative. Fam Pract 2004; 21: 4–10.

34. Leemis LM, Trivedi KS. A comparison of approximate interval estimators for the Bernoulli parameter. The American Statistician 1996; 50: 63–8.

35. Moses LE, Shapiro D, Littenberg B. Combining independent studies of a diagnostic test into a summary ROC curve: data-analytic approaches and some additional considerations. Stat Med 1993; 12: 1293–316.

36. Higgins JPT, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. BMJ 2003; 327: 557–60.

37. Deeks JJ, Macaskill P, Irwig L. The performance of tests of publication bias and other sample size effects in systematic reviews of diagnostic test accuracy was assessed. J Clin Epidemiol 2005; 58: 882–93.

38. Zamora J, Abraira V, Muriel A, Khan K, Coomarasamy A. Meta-DiSc: a software for meta-analysis of test accuracy data. BMC Med Res Methodol 2006; 6: 31.

39. Romero-Gomez M, Cordoba J, Jover R, et al. Value of the critical flicker frequency in patients with minimal hepatic encephalopathy. Hepatology 2007; 45: 859–85.

40. Montoliu C, Piedrafita B, Serra MA, et al. Activation of soluble guanylate cyclase by nitric oxide in lymphocytes correlates with minimal hepatic encephalopathy in cirrhotic patients. J Mol Med 2007; 85: 237–45.

41. Sharma P, Sharma BC, Puri V, Sarin SK. Minimal hepatic encephalopathy in patients with extrahepatic portal vein obstruction. Am J Gastroenterol 2008; 103: 1406–12.

42. Montoliu C, Piedrafita B, Serra MA, et al. IL-6 and IL-18 in blood may discriminate cirrhotic patients with and without minimal hepatic encephalopathy. J Clin Gastroenterol 2009; 43: 272–9.

43. Dhiman RK, Kurmi R, Thumbarri KK, et al. Diagnosis and prognostic significance of minimal hepatic encephalopathy in patients with cirrhosis of liver. Dig Dis Sci 2010; 55: 2381–90.

44. Goel A, Yadav S, Saraswat V, et al. Cerebral oedema in minimal hepatic encephalopathy due to extrahepatic portal venous obstruction. Liver Int 2010; 30: 1143–51.

45. Sharma P, Sharma BC, Sarin SK. Critical flicker frequency for diagnosis and assessment of recovery from minimal hepatic encephalopathy in patients with cirrhosis. Hepatobiliary Pancreat Dis Int 2010; 9: 27–32.

46. Maldonado-Garza HJ, Vazquez-Elizondo G, Gaytan-Torres JO, Flores-Rendon AR, Cardenas-Sandoval MG, Bosques-Padilla FJ. Prevalence of minimal hepatic encephalopathy in cirrhotic patients. Ann Hepatol 2011; 10 (Suppl. 2): S40–4.

47. Montagnese S, Schiff S, De Rui M, Crosey MM, Amodio P, Taylor-Robinson SD. Neuropsychological tools in hepatology: a survival guide for the clinician. J Viral Hepat 2012; 19: 307–15.

48. Onal IK, Akdogan M, Oztas E, Balci M, Kurt M, Kacar S, et al. Does interleukin-18 play a role in the pathogenesis of hepatic encephalopathy? Hepatogastroenterology 2011; 58: 497–502.

49. Shawcross DL, Wright G, Olde Damink SW, Jalan R. Role of ammonia and inflammation in minimal hepatic encephalopathy. Metab Brain Dis 2007; 22: 125–38.

50. Hanley JA, McNeil BJ. The meaning and use of the area under a receiver operating characteristic (ROC) curve. Radiology 1982; 143: 29–36.

51. Sharma P, Sharma BC, Puri V, Sarin SK. Critical flicker frequency: diagnostic tool for minimal hepatic encephalopathy. J Hepatol 2007; 47: 67–73.

52. Sharma P, Sharma BC. Predictors of minimal hepatic encephalopathy in patients with cirrhosis. Saudi J Gastroenterol 2010; 16: 181–7.