Cerebrospinal fluid lactate: Is it a reliable and valid marker to distinguish between acute bacterial meningitis and aseptic meningitis?

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See related research by Huy et al., http://ccforum.com/content/14/6/R240

Abstract

Cerebrospinal fluid (CSF) lactate assay has been a subject of research since 1925. A systematic review by Huy and colleagues in the previous issue of Critical Care summarizes data from 25 studies evaluating the role of CSF lactate in the differential diagnosis between acute bacterial and aseptic meningitis. The authors concluded that CSF lactate is a good single indicator and a better marker compared with conventional markers. But concerns remain because of poor quality of included studies, lack of proper 'gold standard', and limited applicability. More studies with a rigorous design are needed to determine definitively whether CSF lactate assay is a reliable and valid marker to distinguish between acute bacterial meningitis and aseptic meningitis.

Threats to internal validity

The quality of a meta-analysis can be only as good as the included studies ('GIGO' principle: garbage in, garbage out). The quality of studies included in the review by Huy and colleagues is somewhat unsettling. Specifically, reported blinding of lactate assay in only 3 (13%) studies and consecutive or random recruitment of participants in 12 (50%) is a matter of concern. It is possible that other studies blinded the lactate assay without reporting the fact, but this matter remains speculative in the absence of confirmation from the study authors. Compromised quality of original studies threatens the validity of the conclusions.

Lack of a proper 'gold standard' for AM or VM is a vexing problem in this area of research. When an imperfect standard is used to evaluate a diagnostic test, distortions occur in the commonly used measures of test performance, like sensitivity/specificity [2]. Distorted measures carry the error in their meta-analysis. This review suffers from this error.
Two comments on the comparison of CSF lactate assay with conventional CSF markers are warranted. First, clinicians diagnose AM on the basis of a pattern of a combination of findings on conventional CSF markers (CSF total number of leukocytes, CSF glucose, CSF/plasma glucose quotient, and CSF protein), not on individual markers. Thus, it is clinically relevant to compare this pattern and the CSF lactate assay and to determine whether the assay adds significantly to the pattern. However, the authors did not address this question; this was probably because they did not have access to the individual patient data that are necessary to perform this comparison. The review, therefore, fails to answer this question.

Second, the authors assert a lower accuracy of individual CSF markers compared with the CSF lactate test based on point estimates of AUC. It is not clear whether the observed differences in AUC are statistically significant. An objective assessment of this is possible by using the Hanley test [3] and by calculating the confidence interval around the estimates.

**Applicability**
CSF lactate assay is not available in most centers in developing countries and rural settings. This and the fact that many patients receive antibiotics before lumbar puncture compromise the applicability of the findings.

**Summary**
The review is a worthwhile contribution to the field, has a sound methodology, and provides a summary of the results from published data. Clearly, a meta-analysis of individual patient data and more studies are required to determine definitively whether CSF lactate assay is a reliable and valid marker to distinguish between BM and AM.

**Abbreviations**
AM, aseptic meningitis; AUC, area under the curve; BM, bacterial meningitis; CSF, cerebrospinal fluid; VM, viral meningitis.

**Competing interests**
The authors declare that they have no competing interests.

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