A Computationally Simply Central Monitoring Procedure was Proposed and Effectively Applied to Empirical Trial data with Known Fraud

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PII: S0895-4356(16)30398-5
DOI: 10.1016/j.jclinepi.2017.03.018
Reference: JCE 9364

To appear in: Journal of Clinical Epidemiology

Received Date: 31 August 2016
Revised Date: 27 February 2017
Accepted Date: 31 March 2017

Please cite this article as: van den Bor RM, Vaessen PW, Oosterman BJ, Zuihoff NP, Grobbee DE, Roes KC, A Computationally Simply Central Monitoring Procedure was Proposed and Effectively Applied to Empirical Trial data with Known Fraud, Journal of Clinical Epidemiology (2017), doi: 10.1016/j.jclinepi.2017.03.018.

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Role of the funding source
Funding was provided by Julius Clinical Ltd. No individuals besides the authors were involved in the study design; in the collection, analysis and interpretation of data; in the writing of the report; or in the decision to submit the article for publication.

Conflicts of Interest
None

Trial information
The methodology proposed in this paper is illustrated using data from the Second European Stroke Prevention Study (ESPS2). ESPS2 was initiated in 1988 and supported by a grant from Boehringer Ingelheim.

Acknowledgments
The authors thank Boehringer Ingelheim for the provision of the data, NA Hilkens for useful discussion, and M Hugenholtz and P Nowak for their assistance in generating pilot data.

Abstract
Objective: Central monitoring of multi-center clinical trials becomes an ever more feasible quality assurance tool, in particular for the detection of data fabrication. More widespread application, across both industry-sponsored as well as academic clinical trials, requires central monitoring methodologies that are both effective and relatively simple in implementation.
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