

University of Birmingham Research at Birmingham

Weighted variogram analyses for estimating withinpatient variance components using routine data from biomarker monitoring programmes

Baldwin, Simon; Sitch, Alice; Takwoingi, Yemisi; Deeks, Jon

Document Version Peer reviewed version

Citation for published version (Harvard):

Baldwin, S, Sitch, A, Takwoingi, Y & Deeks, J 2020, 'Weighted variogram analyses for estimating within-patient variance components using routine data from biomarker monitoring programmes', Methods for Evaluation of medical prediction Models, Tests and Biomarkers (MEMTAB) 2020, Leuven, Belgium, 10/12/20 - 11/12/20.

Link to publication on Research at Birmingham portal

General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- •Users may freely distribute the URL that is used to identify this publication.
- •Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- •User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- •Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

Download date: 02. May. 2024

39.

Weighted variogram analyses for estimating within-patient variance components using routine data from biomarker monitoring programmes

S. Baldwin^{1,2}, A. Sitch^{1,2}, Y. Takwoingi^{1,2}, J. Deeks^{1,2}
¹Test Evaluation Research Group, Institute of Applied Health Research, University of Birmingham, Edgbaston, Birmingham, B15 2TT, UK;
²National Institute for Health Research Birmingham Biomedical Research Centre, University Hospitals Birmingham NHS Foundation Trust and University of Birmingham, UK

Correspondence: S. Baldwin

Diagnostic and Prognostic Research 2021, 5(Suppl 1):39.

Background: Measurement error in biomarkers is best estimated in Biological Variability Studies (BVS) where individuals have repeated measures both at the same and at different time points. However, BVS are not always feasible. We investigate whether measurement error can be estimated using routine data from biomarker monitoring programmes by application of a method known as the variogram.

We aimed to demonstrate the potential of the variogram using open-source monitoring programme data of serum albumin measurements on stage 2-4 primary biliary cirrhosis patients.

Methods: Variation in measurements from patients over time includes three components: true differences at baseline between-patients; true changes from baseline within-patients ('signal'); and measurement error ('noise'). The variogram considers differences within-patients computed between baseline and each follow-up point; the variances of these differences increase at a rate dependent on the magnitude of the within-patient variability. We grouped measurements by year, and assigned weights according to the closeness of the actual time to the midpoint using a Gaussian kernel approach. Weighted variances of differences in serum albumin were calculated per time. The variogram is a plot of weighted variance of differences (y-axis) against time (x-axis), with a fitted line estimated by linear regression, weighted according to sample size. Extrapolation of the fitted line to intersect the y-axis was used to estimate the measurement error ('noise').

Results: The measurement error ('noise') estimate was 0.10 (gm/dL)². (Figure 1) 'Signal' first surpassed 'noise' at five years; the variance of differences at one year was estimated almost entirely 'noise'. Such results from weighted variogram analyses could be used to help define optimal measurement timings for monitoring programmes.

Conclusions: Weighted variogram analyses have potential for application where health status changes are unlikely; care should be exercised in implementation, particularly related to bias from dropout. **Keywords:** Variogram, variability, monitoring, measurement error

