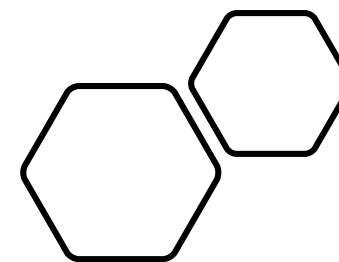


STARD - BLCM

Polychronis Kostoulas



The STARD family

- **STARD**
 - STARD 2015
 - STARDdem
 - STRADAS-paraTB
 - STARD-BLCM
 - STARD for Abstracts



Enhancing the QUALity and Transparency Of health Research

Home

About us

Library

Toolkits

Courses & events

News

Blog

Libra

Your one-stop-shop for writing and publishing high-impact health research

find reporting guidelines | improve your writing | join our courses | run your own training course | enhance your peer review



Library for health research reporting

The Library contains a comprehensive searchable database of reporting guidelines and also links to other resources relevant to research reporting.



Reporting guidelines for main study types

[Randomised trials](#)

[CONSORT](#)

[Extensions](#)

[Observational studies](#)

[STROBE](#)

[Extensions](#)

[Systematic reviews](#)

[PRISMA](#)

[Extensions](#)

[Study protocols](#)

[SPIRIT](#)

[PRISMA-P](#)

STARD-BLCM Checklist

TITLE OR ABSTRACT

ABSTRACT

INTRODUCTION

METHODS

Study design

Participants

Test methods

Analysis

RESULTS

Participants

Test results

DISCUSSION

OTHER INFORMATION





| Section & Topic | No | Item |
|-------------------|----|--|
| TITLE OR ABSTRACT | | |
| | 1 | Identification as a study of diagnostic accuracy, using at least one measure of accuracy (such as sensitivity, specificity, predictive values, or AUC) and Bayesian latent class models |
| ABSTRACT | | |
| | 2 | Structured summary of study design, methods, results, and conclusions (for specific guidance, see <u>STARD</u> for Abstracts) |
| INTRODUCTION | | |
| | 3 | Scientific and clinical background, including the intended use and clinical role of the tests under evaluation |
| | 4 | Study objectives and hypotheses, such as estimation of diagnostic accuracy of the tests for a defined purpose through <u>BLCM</u> |



| | | |
|---------------------|----------|---|
| METHODS | | |
| <i>Study design</i> | 5 | Whether data collection was planned before the tests were performed (prospective study) or after (retrospective study) |
| <i>Participants</i> | 6 | Eligibility criteria and description of the source population |
| | 7 | On what basis potentially eligible participants were identified (such as symptoms, results from previous tests, inclusion in registry) |
| | 8 | Where and when potentially eligible participants were identified (setting, location and dates) |
| | 9 | Whether participants formed a consecutive, random or convenience series |



| | | |
|---------------------|------------|--|
| <i>Test methods</i> | 10 | Description of the tests under evaluation , in sufficient detail to allow replication, and/or cite references |
| | 11 | Rationale for choosing the tests under evaluation in relation to their purpose |
| | 12 | Definition of and rationale for test positivity cut-offs or result categories of the tests under evaluation , distinguishing pre-specified from exploratory |
| | 13 | Whether clinical information was available to the performers or readers of the tests under evaluation |
| <i>Analysis</i> | 14a | <u>BLCM</u> model for estimating measures of diagnostic accuracy |
| | 14b | Definition and rationale of prior information and sensitivity analysis |
| | 15 | How indeterminate results of the tests under evaluation were handled |
| | 16 | How missing data of the tests under evaluation were handled |
| | 17 | Any analyses of variability in diagnostic accuracy, distinguishing pre-specified from exploratory |
| | 18 | Intended sample size and how it was determined |



| | | |
|---------------------|-----------|--|
| RESULTS | | |
| <i>Participants</i> | 19 | Flow of participants, using a diagram |
| | 20 | Baseline demographic and clinical characteristics of participants |
| | 21 | Not applicable: the distribution of the targeted conditions is unknown, hence the use of <u>BLCM</u> |
| | 22 | Time interval and any clinical interventions between the tests under evaluation |
| <i>Test results</i> | 23 | Cross tabulation of the tests' results (or for continuous tests results their distribution by infection stage) |
| | 24 | Estimates of diagnostic accuracy under alternative prior specification and their precision (such as 95% credible/probability intervals) |
| | 25 | Any adverse events from performing the tests under evaluation |
| DISCUSSION | | |



| | | |
|-------------------|----|---|
| DISCUSSION | | |
| | 26 | Study limitations, including sources of potential bias, statistical uncertainty, and generalisability |
| | 27 | Implications for practice, including the intended use and clinical role of the tests under evaluation in relevant settings (clinical, research, surveillance etc.) |
| OTHER INFORMATION | | |
| | 28 | Registration number and name of registry |
| | 29 | Where the full study protocol can be accessed |
| | 30 | Sources of funding and other support; role of funders |

No Gold Standard

Definition of the targeted infection status gets complicated but..."better"

Detectable immune response (ELISA)

Detectable isolation of the pathogen (Culture)



Bayesian Analysis

Model description

Identifiability

Priors – Sensitivity Analysis

Explicit description of the BLCM model

Model Assumptions

Identifiability

Definition and **rationale** of prior information and **sensitivity analysis**

Can the interested reader reproduce your results?



HARMONY

Novel tools for test evaluation and
disease prevalence estimation

Thank you!