

Evidence tables III Prognostic and economic evaluation

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Why work on evidence tables?

- Need for a common definition
- Provides a real opportunity for joint working between guideline agencies and between GIN and other international groups
- Potentially could save significant duplication of effort across all guideline development agencies

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Work so far...

- 1. WG agreed definition of ETs at Lyon Conference 2005
 - Evidence tables are methodological and outcome summaries that present data from a number of related studies. These answer a well defined question in a consistent format and aim to demonstrate overall trends in the evidence and enable the process of making recommendations.
- 2. Development of a minimum data set for summarising studies that evaluate an intervention and a definitive template for describing them
 - Presented in a workshop at the Toronto conference in August 2007.
 - Now accepted for publication by Quality & Safety in **Health Care**



Work so far...

- 3. Development of a template for summarising studies addressing a diagnostic question
 - The results of the evaluation study were presented at the Helsinki Conference in October 2008
- 4. Agreement of the specification for the registry of summarised studies to enable G-I-N members to share their wok with others
 - Discussed at the Lisbon Conference 2009
 - G-I-N Board approved funding 2010



What we are doing now...

• Developing templates for:

- summarising economic evaluation studies
- summarising prognostic studies.

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Draft template for prognostic studies

Ton Kuijpers Hans de Beer

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Introduction

- Prognosis: foreseeing, predicting, or estimating the probability or risk of future conditions
- In medicine, prognosis commonly relates to the probability or risk of an individual developing a particular state of health (an outcome) over a specific time, based on his or her clinical and non-clinical profile

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Introduction



 Outcomes are often specific events, such as death or complications, but they may also be quantities, such as disease progression, (changes in) pain, or quality of life

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Introduction

Explanatory studies

• Studies investigating whether a single variable (such as a tumour or other biomarker) may be prognostic.

Prediction rules

Studies aimed at predicting outcomes from multiple variables.

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Introduction



Examples of prediction rules

- the Apgar score to determine the prognosis of newborns
- cardiovascular risk profiles to predict heart disease in the general population
- prenatal testing to assess the risk that a pregnant woman will give birth to a baby with Down's syndrome

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Introduction

Consecutive phases in multivariable prognostic research

- Development studies
- · Validation studies
- Impact studies

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Introduction



Prediction rules: development studies

 Development of a multivariable prognostic model, including identification of the important predictors, assigning relative weights to each predictor, and estimating the model's predictive performance through calibration and discrimination and its potential for optimism using internal validation techniques, and, if necessary, adjusting the model for overfitting.

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Introduction

Prediction rules: validation studies

 Validating or testing the model's predictive performance (eg, calibration and discrimination) in new participants. This can be narrow (in participants from the same institution measured in the same manner by the same researchers though at a later time, or in another single institution by different researchers using perhaps slightly different definitions and data collection methods) or broad (participants obtained from various other institutions or using wider inclusion criteria).

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Prediction rules: impact studies

 Quantifying whether the use of a prognostic model by practising doctors truly improves their decision making and ultimately patient outcome, which can again be done narrowly or broadly

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References

- Moons KG, Royston P, Vergouwe Y, Grobbee DE, Altman DG. Prognosis and prognostic research: what, why, and how? BMJ 2009;338:b375.
- Royston P, Moons KG, Altman DG, Vergouwe Y. Prognosis and prognostic research: Developing a prognostic model. BMJ 2009;338:b604..
- Altman DG, Vergouwe Y, Royston P, Moons KG. Prognosis and prognostic research: validating a prognostic model. BMJ 2009;338:b605.
- Moons KG, Altman DG, Vergouwe Y, Royston P. Prognosis and prognostic research: application and impact of prognostic models in clinical practice. BMJ 2009;338:b606.

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Finding existing templates



- Literature search carried out to identify existing templates for summarising prognostic studies
- Conducted by Robin Harbour and colleagues at the The Scottish Intercollegiate Guidelines Network SIGN), Edinburgh

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Finding existing templates

- Asking G-I-N member organisations for prognostic templates they are currently using
- Contacting Cochrane Prognosis Methods Group by CBO (Ton Kuijpers / Hans de Beer)

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Background information



Bibliographic citation	Use Vancouver style (Authors. Title. Journal name. Publication Date; Volume (Issue):Page Numbers) Insert the link to the publication.
Sources of funding and competing interest	Report: • The source of funding cited in the paper: give name(s) of organisation or corporation. Specify if possible the source type (public research funds, NGO, government, Academic/university healthcare industry or other) • Competing interests: Write "Stated" or "Not Stated" and specify if any
Setting	Multicenter, Location/Country(ies), Healthcare setting
Objective(s) of the study	Report, as cited by author(s), the objective(s) of the study including both primary and secondary aims, if applicable

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Questions addressed	Mention the questions really addressed (e.g. include all questions evenif only one is relevant for you at the moment)
Type of prognostic study	Specify whether the study relates to what prognostic factors are associated with outcome (explanatory study), or to what groups of prognostic factors best predicted outcome (outcome prediction study)

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Methods

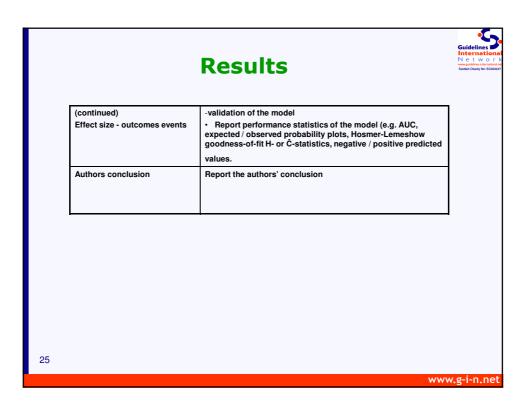


Study design (cited by author or actual)	Specify the study design: cross sectional study, cohort study, case control study, other (give details). Precise if it's the design cited by author(s).
Eligibility criteria	Describe the: Eligibility criteria i.e. inclusion-exclusion criteria, Methods of selection of participants
Follow-up	Specify the follow-up moments, periods of recruitment, and data collection if applicable
Outcome events	Describe the outcome events identified by author(s), both primary and secondary ones
Prognostic factors and potential confounders	Describe the potential prognostic factors (in case of an outcome prediction study) or the central prognostic factor, potential confounders and effect modifiers (in case of an explanatory study)

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Results Report numbers of individuals at each stage of study: Numbers numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed Number of patients excluded and reasons (i.e. give reasons for non-participation at each stage) Patients characteristics Describe the actual population involved in the study: Give characteristics of study participants (e.g. demographic, clinical, social) and information on exposures and potential confounders · Indicate number of participants with missing data for each variable of interest Report Follow-up time: average/ median and range if applicableØ Highlight discrepancies between groups Outcome events data Report all available figures with 95% confidence intervals when available:Report numbers of outcome events or summary measures 23

Results Effect size - outcomes events Explanatory Report <u>all available figures</u> with 95% confidence intervals when available: Report unadjusted univariate estimates and their precision Report adjusted multivariate estimates and their precision Specify category boundaries when continuous variables were categorized Outcome prediction studies - development of the model · Report unadjusted univariate estimates and their precision Report adjusted multivariate estimates and their precision Report performance statistics of the model (AUC, expected / observed probability plots, Hosmer-Lemeshow goodness-of-fit H- or Ĉ (ik krijg hier een hokje te zien)-statistics, negative / positive predicted values. 24 www.g-i-n.ne





Draft template for economics studies

Rob Cook

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Finding existing templates



- Literature search carried out to identify existing economic templates
- Conducted by Andreas Gerber and colleagues at the Institute for Quality and Efficiency in Health Care (IQWiG), Cologne

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Literature search strategies

PubMed

cost-effectiveness* [ti] OR economic evaluation* [ti] OR cost-utility-analy* [ti] OR cost-benefit* [ti] OR decision-making model* [ti]) AND (data extraction sheet* OR data extraction template)

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Literature search strategies



The Cochrane Library (Wiley), including partial data banks

Search strategy: data extraction sheet* OR data extraction template*

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Literature search strategies

Hand search (Web-sites)

- HTA-Institutions
- HTA-Handbooks/ Methods manuals
- HTA-reports containing "economic evaluation OR costeffectiveness" in the title

All: Full text screening)

- Institutes with focus on health economic evaluation
- WHO
- · EUnetHTA Core model

Google

Internet portals:

Karlsruher Virtueller Katalog

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Evidence tables used by member organisations



CADTH
CCOHTA
CDR
HTA (Germany)
KCE (Belgium)
Ludwig Boltzman Institut
NCCHTA (UK)

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Background information



Name of person completing template	
Date of completion	
Bibliographic citation	Use Vancouver style (Authors. Title. Journal name. Publication Date; Volume (Issue):Page (Numbers) Hyperlink to publication
Sources of funding and competing interests	Quote the source of funding if mentioned in the paper. If possible, specify the source type (public research funds, Government, NGO, academic/university, healthcare industry, other). Quote any declared competing interests, or write "None stated"

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Methods



Study design	Specify the form of economic evaluation being used(CUA, CEA, CBA, etc).
Viewpoint	State the viewpoint of the analysis.
Country/Countries	List the country/countries for economic evaluation
Time horizon	State the time horizon for both costs and benefits
Population	Describe the eligible population and the population used for effect/cost data
Interventions	Describe the alternatives being compared
Primary outcome measure(s)	State the primary outcome measure(s)
Source of effectiveness estimates	Identify the source (meta analysis, RCT, etc)
Methods used to value benefits	Identify the methods used (stated WTP, revealed WTP, conjoint analysis etc)
Utilities and benefits	Specify where utilities or benefits came from (literature values, elicited in the study etc).

Methods continued



Source of estimates of quantities of resources used	Describe the methods use to identify relevant resource use (survey, lit values, expert consultation, formal consensus methods etc)
Methods used to estimate unit costs	Describe the methods used to identify relevant unit costs (survey, lit values etc) Include cost year
Methods used to estimate unit costs	Describe the methods used to identify relevant unit costs (survey, lit values etc) Include cost year
Adjustment for inflation	Was adjustment for inflation performed if unit costs stemmed from different years?
Discounting	Was discounting performed? Were effects and cost discounted with the same rate(s)?
Approach to sensitivity analysis	Describe the methods used for sensitivity analysis (one way, two way, probabilistic etc)
Modelling and analysis	Detail any model used (Markov, Discrete Event Simulation etc)

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Results



Costs and outcomes	Presented relevant costs and outcomes in both disaggregated and aggregated form
Subgroup analyses	Were subgroup analyses performed? Are the based on the results of RCTs etc.?
Incremental analysis results	Present the results of IA
Sensitivity analysis results	Present the results of SA

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Critical appraisal



Authors conclusion	Report authors' conclusions verbatim
Validity of results	Comment on the strengths and weaknesses of the study. If weaknesses were identified were these likely to have made a difference to the results of the economic evaluation?
Other / addendum (optional)	

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