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Would a cohort-level approach to cost-effectiveness modelling have led to a different decision in an important NICE appraisal for obesity patients?



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Key message: Cohort-level approach in modelling obesity (NICE TA494) could have led to similar recommendations, however our recreation indicates that the flexibility of the patient-level approach used in the submission was valuable in capturing patient heterogeneity and time-dependency, though reimplementation in a non-NICE framework was necessary for acceptable model execution speed.

Introduction and objectives

In 2017, the National Institute for Health and Care Excellence (NICE) appraised naltrexone-bupropion (NB32) + standard management (SM) versus SM alone for the treatment of adult obesity (TA494) based on a patient-level model originally implemented as a discretely integrated condition event (DICE) simulation¹.

The Evidence Review Group (ERG) had concerns with how the implementation of the model affected its run time, limiting the ERG's ability to simulate sufficient patients and probabilistic sensitivity analysis (PSA) iterations. This may have contributed to the NICE appraisal committee not recommending NB32 as an appropriate use of National Health Service (NHS) and Personal Social Services (PSS) resources, though a reimplemented version of the model in a non-DICE (within VBA) framework was accepted and used to inform Final Appraisal Determination.

This study aims at examining the impact of modelling the same decision problem using a (non-DICE) cohort-level approach.

Methods

Conceptual modelling and literature reviews were conducted to develop a cohort-level model structure (**Figure 1**).

The model was a non-DICE, 40-state, probabilistic, cohort-level Markov model and was specified and constructed in Microsoft Excel®.

Data inputs and assumptions matched the NICE committee preferred approach based on available materials at time of replication.

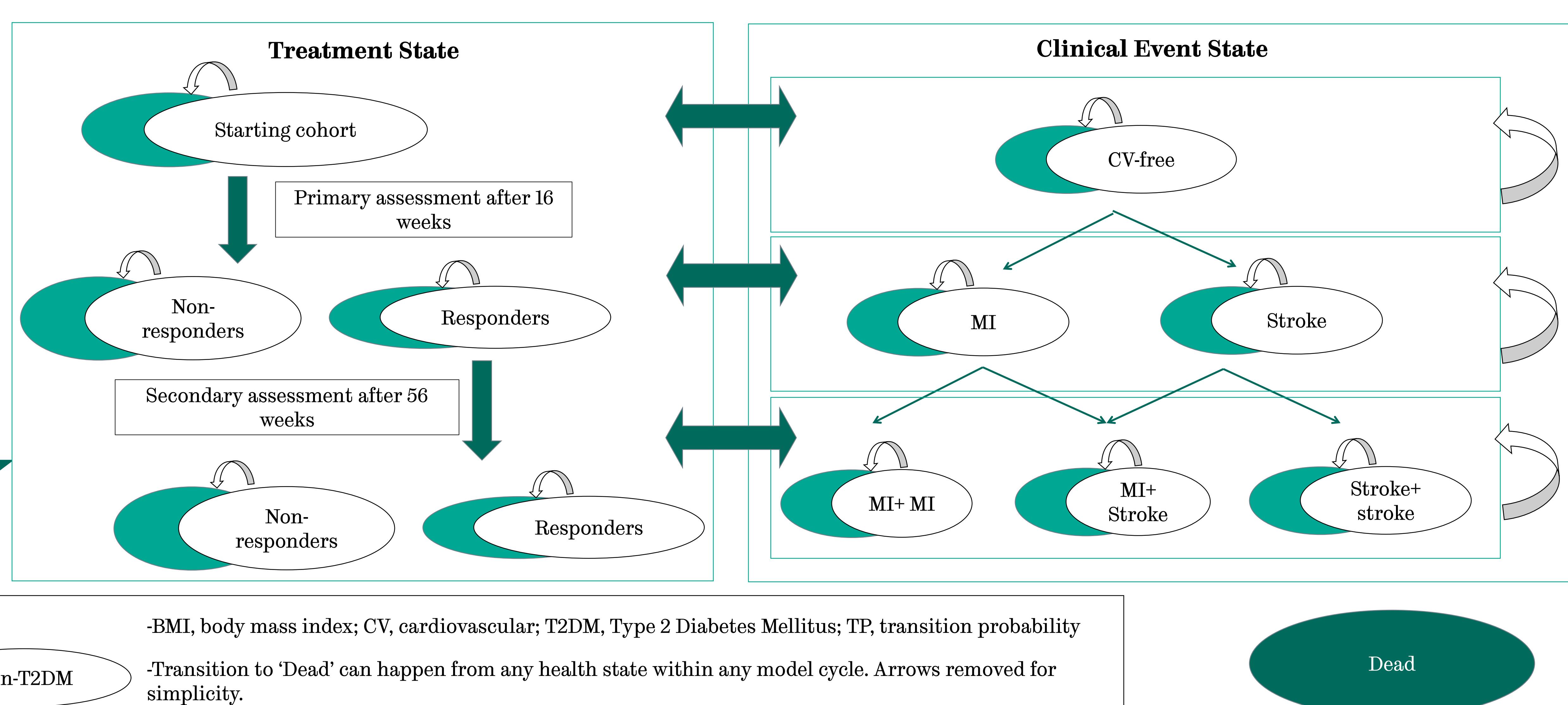
Main model assumptions

Upon model entry, no patients had a history of angina and/or diabetes other than Type 2 Diabetes Mellitus (T2DM).

A 4-week cycle length was applied (based on treatment administration).

Figure 1: *De novo* model schematic

As patients progress, BMI changes per cycle are calculated through calculating BMI trajectory per sub-cohort, and updating TPs and utilities consequently



References: (1) NICE. Naltrexone-bupropion for managing overweight and obesity [TA494]. 2017. Available at: <https://www.nice.org.uk/guidance/ta494/documents/appraisal-consultation-document-2> (accessed 2018 Sep 5); (2) Ara R, Blake L, Gray L, et al. What is the clinical effectiveness and cost-effectiveness of using drugs in treating obese patients in primary care? A systematic review. Health Technol Assess. 2012;16(5); (3) NICE. Guide to the methods of technology appraisal. 2013. Available from: <https://www.nice.org.uk/process/pmg9/chapter/the-appraisal-of-the-evidence-and-structured-decision-making> (accessed 2018 Sep 1).

Disclaimer: Metry completed the dissertation module of his MSc Health Economics and Decision Modelling on placement at BresMed Health Solutions Ltd, under joint supervision with the awarding institution, University of Sheffield. Part of the work presented here

